BEFORE THE NEVADA STATE BOARD OF PHARMACY

NEVADA STATE BOARD OF PHARMACY,

CASE NOS. 21-069-RPH-N 21-069-PH-N

Petitioner,

V.

NATHAN DAHL, RPH, Certificate of Registration No. 17735, and

DAHL'S SPECIALTY PHARMACY OF CARSON, License No. PH03605,

Respondent.

STIPULATED FACTS

Brett Kandt, General Counsel for Petitioner the Nevada State Board of Pharmacy (Board), and Respondents Respondent Nathan Dahl, R.Ph., Certificate of Registration No. 17735, and Respondent Dahl's Specialty Pharmacy of Carson, License No. PH03605, by and through counsel, William J. Stilling, Esq. HEREBY STIPULATE AND AGREE THAT:

- The Board has jurisdiction over this matter because, at the time of the events
 herein, Respondent Nathan Dahl, R.Ph. (Dahl), Certificate of Registration No. 17735, was a
 pharmacist registered by the Board, and Respondent Dahl's Specialty Pharmacy of Carson
 (Dahl's Pharmacy), License No. PH03605, was a pharmacy licensed by the Board.
 - 2. Dahl owned and operated Dahl's Pharmacy at the time of the events herein.
- 3. On or about February 3, 2021, D.L. was prescribed dalfampridine 5mg to be compounded in capsule form, 120 capsules, to be taken orally, 1 capsule 4x per day, with five (5) refills. The prescription was transmitted to Dahl's Pharmacy and designated as Prescription No. 5070945.
- 4. Between February 3 and May 12, 2021, each month Dahl compounded, filled and dispensed Prescription No. 5070945, 5mg dalfampridine, 120 capsules, for D.L. D.L. was Dahl's only patient taking dalfampridine during this period of time.

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- On or about May 12, 2021, Dahl dispensed Prescription No. 5070945 which contained a mixture of capsules compounded on April 20, 2021, lot #4202021, and May 10, 2021, lot #5102021.
- Between May 12 and May 23, 2021, D.L. ingested Prescription No. 5070945 as directed by her physician.
- 7. On or about May 15, 2021, D.L. was taken to the emergency room with symptoms she thought might have resulted from taking dalfampridine. She returned home.
- On or about May 23, 2021, D.L. was taken to the emergency room and was transported via Care Flight from Carson City to Renown Regional Medical Center.
- On June 1, 2021, D.L.'s daughter advised Dahl of D.L.'s condition and questioned whether Prescription No. 5070945 was correctly compounded. Dahl stated that he would have the capsules tested.
- On June 2, 2021, D.L.'s daughter provided Dahl with ten (10) dalfampridine capsules to be tested.
- On June 3, 2021, Dahl left D.L.'s daughter a voicemail message representing that lab results indicated the correct dosage of 5mg dalfampridine per capsule.
- 12. On or about June 9, 2021, D.L.'s daughter provided Segedy with fifty (50) dalfampridine capsules from Prescription No. 5070945.
- 13. Dahl sent and ARL Bio Pharma in Oklahoma (ARL) received 10 (ten) capsules from lot #5102021 on June 16, 2021, for testing.
- 14. On June 16, 2021, Segedy sent two (2) of the suspect capsules to ARL for a complete assay test of the dalfampridine potency. ARL received those capsules on June 18, 2021.
- 15. On or about July 12, 2021, the Board received a certificate of analysis from ARL confirming that one of the capsules tested at 19.48 mg of dalfampridine, 4 times the prescribed dosage, and the other capsule contained 5.3439 mg of dalfampridine.

- 16. Inasmuch as it cannot be determined which batch the high potency capsule came from, lot #4202021 or lot #5102021, Dahl erroneously compounded at least one capsule of the drug on or about April 20, 2021, or May 10, 2021, at 4 times the prescribed dosage.
- 17. On or about July 21, 2021, Dahl's Pharmacy ceased to do business and permanently closed.

The undersigned parties stipulate to these facts solely for the purpose of the administrative hearing in this matter based on negotiations by counsel to facilitate the efficient conduct of the hearing and in the context of administrative proceedings. The parties do not intend to stipulate to or admit to these facts in any other context or proceeding.

AGREED:

and the state of	T	
Signed this	day of December, 2021	S

Signed this $\frac{1}{2}$ day of December, 2021

NATHAN DAHL, RPH,

Certificate of Registration No. 17735

BRETT KANDT, ESQ.

General Counsel

Nevada State Board of Pharmacy

Signed this 157 day of December, 2021

DAHL'S SPECIALTY PHARMACY OF CARSON, License No. PH03605

APPROVED AS TO FORM AND

CONTENT this ___ day of December, 2021

WILLIAM J, STILLING, ESQ.

Counsel for Respondents



06/09/2024

NEVADA STATE BOARD OF PHARMACY

985 Damonie Ranch Pkwy, Ste 206, Reno, NV 89521 (775) 850-1440 • 1-800-364-2081 • FAX (775) 850-1444 • Web Page: bop.av.gov

Complaint Form

opyridine Compound Error
City, State, Zip: Gardnerville, NV 89460
Business N/A
Date of Birth
Telephone Number: 775-770-3554
City, State, Zip: Reno, NV 89503
Prescription Number: 5070945
1007 N Curry Street Suite C Carson City, NV 89703

STATEMENT OF COMPLAINT: Type or neatly print your complaint below. Be as concise as possible. Use reverse side if necessary. Make copies and attach any documents you have which support your allegation(s). After completing your statement of complaint, please sign and date it. <u>The Board does not have jurisdiction over complaints involving rudeness</u>, customer service and/or pricing/billing disputes.

William writing this complaint on behalf of my mother, D currently on life support in ICU. On Wednesday 5/12 at 11:42am DeAnna picked up her compounded Dalfampridine (4-Aminoprydine) from Dahls Pharmacy located in Carson City, She did not start the new bottle of medication until Saturday 5/15, ironically the day she had her first episode of overdose like symptoms resulting in being taken by ambulance to the emergency room. She was very confused as to why this had happened, and thought maybe she had taken two of the 5mg capsules a tad too close together. Feeling the best choice would be to not take the full recommended dosage of 5mg pills 4x a day, she continued the week with only taking two 5mg pills 2x a day instead. Throughout the week her walking ability declined, as if the pills could have been making it worse. On Sunday 5/23 she began to feel the same overdose like symptoms prompting my father Lance to take her to the emergency room. There, she went in to convulsions for 2 hours before suffering two seizures and still convulsing afterwards. She was put on life support and careflighted from Gardnerville to Reno. My mother went in to Status Epilepticus (non stop seizures) that was resistant to all medications the hospital tried for days. It wasnt until sedating her with extreme drugs and more days passing that the seizures finally subsided. After trying to figure out the cause of this I decided to contact Dahls Pharmacy. On Tuesday 6/1 I spoke with the owner and pharmacist Nathan and he said that he would send out a pill to be tested and also gave me the ingredients and formulation for the medication. He offered to even come pick up the pills from me. Please understand that by signing and submitting this form to the Board of Pharmacy, you are authorizing and allowing this Board's staff to access your medical history and records, including pharmacy records, as needed to investigate your complaint. If you would like to limit what the Board's staff can review, you must inform us of those limitations in writing.

Posted 4:22/2021



06/08/2021

NEVADA STATE BOARD OF PHARMACY

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Complaint Form

Complainant Name: Dalfampridine aka 4-Amin Address: Jobs Peak Drive	City, State, Zip: Gardnerville, NV 89460
Telephone Numbers: Home	Business N/A
Patient Name: D	Date of Birth:
Physician Name: Dr. Timothy Louie	Telephone Number: 775-770-3554
Address: 645 N Arlington Ave #655	City, State, Zip: Reno, NV 89503
Drug Prescribed: Dalfampridine	Prescription Number: 5070945
Pharmacy Name & Address: Dahls Specialty Pharmacy	
Pharmacist/Staff: Nathan Dahl	

STATEMENT OF COMPLAINT: Type or neatly print your complaint below. Be as concise as possible. Use reverse side if necessary. Make copies and attach any documents you have which support your allegation(s). After completing your statement of complaint, please sign and date it. <u>The Board does not have inrisdiction over complaints involving rudeness, customer service and/or pricing/billing disputes.</u>

me. Later in the day he left a voicemail message stating that he spoke with the lab and they would need 10 pills, again offering to come to me to retrieve them. Wednesday morning on 6/2 we dropped the pills into Dahl's drop box and later I called to make sure they were received. Nathan said that he would be overnighting them to his lab in Texas and would have results the next day in the afternoon. On Thursday 6/3 Nathan left a voicemail stating that the lab results showed the pills were fine, but that he wanted to get all of the pills back from me to test and would be crediting my mother's account back. He again offered to come to me and retrieve the pills again, sounding adament and somewhat panicked. I called back leaving a message, expressing gratefulness, not realizing that something seemed off said that I would get him the pills on Monday 6/7. I know someone that works at Dahls and Nathan had her send me a text message today 6/8 just before 8am asking when we are going to get the pills to the pharmacy. He also had her message me another day to call him back. All of this seems off and has been making my father and I rather uncomfortable. Upon researching the medication I came across an NIH study of a woman whose dalfampridine, that was supposed to be 10mg, was compounded incorrectly at 100mg. All of her symptoms, trauma, hospitalization, etc. was like reading about every single thing my mother has gone through. At this time, a tracheostomy vent was placed yesterday and a stomach tube is being placed today. Tonight marks 16 days of unbearable tragedy for our family, and we need answers as to why this happened to her. I want to express gratitude ahead of time to everyone who will be

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Data 06/08/2021

N VADA STATE BOARD C. PHARMACY

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Complaint Form

Address: Jobs Peak Drive	City, State, Zip: Gardnerville, NV 89460
Telephone Numbers: Home	Business N/A
Patient Name: D. L.	Date of Birth:
Physician Name: Dr. Timothy Louie	Telephone Number: 775-770-3554
Address: 645 N Arlington Ave #655	City, State, Zip: Reno, NV 89503
Drug Prescribed: Dalfampridine	Prescription Number: 5070945
Pharmacy Name & Address: Dahls Specialty Pharmacy	
Pharmacist/Staff: Nathan Dahl	

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Nathan said that he would be overnighting them to his lab in Texas and would have results the next day in the afternoon. On Thursday 6/3 Nathan left a voicemail stating that the lab results showed the pills were fine, but that he wanted to get all of the pills back from me to test and would be crediting my mother's account back. He again offered to come to me and retrieve the pills again, sounding adament and somewhat panicked. I called back leaving a message, expressing gratefulness, not realizing that something seemed off said that I would get him the pills on Monday 6/7. I know someone that works at Dahls and Nathan had her send me a text message today 6/8 just before 8am asking when we are going to get the pills to the pharmacy. He also had her message me another day to call him back. All of this seems off and has been making my father and I rather uncomfortable. Upon researching the medication I came across an NIH study of a woman whose dalfampridine, that was supposed to be 10mg, was compounded incorrectly at 100mg. All of her symptoms, trauma, hospitalization, etc. was like reading about every single thing my mother has gone through. At this time, a tracheostomy vent was placed yesterday and a stomach tube is being placed today. Tonight marks 16 days of unbearable tragedy for our family, and we need answers as to why this happened to her. I want to express gratitude ahead of time to everyone who will be helping us get these answers. We do not want anyone else to suffer like we have, especially my mother, who at this point, will never be the same again.

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Certificate of Analysis

CLIENT: Nevada State Board of Pharmacy

DESCRIPTION: Dalfampridine-4-Aminopyridine

LOT #: Suspected from lot 42021

FORMULATION ID: Not Provided

ARL#: 765171

DATE RECEIVED: 06/18/2021

STORAGE: 20°C to 25°C

Test	Method	Specifications	Results	Date Tested
Assay - 4-Aminopyridine	HPLC	Report Result	389,5% (19.4765mg / 1Capsule)	07/09/2021

Notes

The potency method(s) used for testing passed system suitability requirements per ARL SOP AMP-012 for non-cGMP analysis. Product specific method validation is not available for this sample and specification(s) are for informational purposes only. Client should verify the specification and analyte reported are correct for the compounded formulation.

07/12/2021

Katie Coats - Chemist II

Date



Certificate of Analysis

CLIENT: Nevada State Board of Pharmacy DESCRIPTION: Dalfampridine-4-Aminopyridine

LOT#: Suspected from lot 5102021

ARL#: 765182

FORMULATION ID: Not Provided

DATE RECEIVED: 06/18/2021

STORAGE: 20°C to 25°C

Test	Method	Specifications	Results	Date Tested
Assay - 4-Aminopyridine	HPLC	Report Result	106.9% (5.3439mg / 1Capsule)	07/08/2021

Notes

The potency method(s) used for testing passed system suitability requirements per ARL SOP AMP-012 for non-cGMP analysis. Product specific method validation is not available for this sample and specification(s) are for informational purposes only. Client should verify the specification and analyte reported are correct for the compounded formulation.

07/12/2021

Katie Coats - Chemist II

Date

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QUF-414-V1, results reported above relate only to the sample that was tested. Report may only be reproduced in full.

Statement of Events

I received a phone call from D L L daughter, C informing me that D had been hospitalized. C asked for information about the compounded capsules we dispensed to D reviewed the log record and prescription record with Christy, explaining each ingredient and what it is used for.

I received a second call from Calinquiring about the possibility of testing to see what was in the compounded capsules. I told her I knew of a lab that did that type of testing and that we would need at least 10 capsules to send for analysis. I offered to send a delivery driver to collect the capsules. Calinquiring about the capsules of the capsules dropped off into our afterhours drop box.

I contacted Eagle Labs and asked if they could test for the active chemical. I was told that they could. While we were on the phone, I reviewed the Certificate of Analysis for the active chemical in this compound and the logs of this compound and the compounds we had dispensed in the past. I was told that based on the Certificate of Analysis of the active chemical and based on our logs for the capsules they should be correct. I reached out to Capsules and communicated the above information. I followed up our conversation with a voice mail message asking Capsules if she would be interested in having all the capsules tested by the lab to ensure the accuracy of every capsule. I did not receive any further communication from Capsules.

I followed up with Eagle Labs and was told that they would in fact not be able to test the actual capsules, only the raw chemical on its own. After receiving this information, I cancelled the shipment to Eagle Labs.

I began researching to find a lab that would be able to test the individual capsules for the active ingredients. Letco Med provided me with the contact information for ARL Labs. I contacted ARL Labs and was told that they could perform the chemical identity tests on the capsules. I sent all 10 capsules to ARL Labs. At that time, I was told that ARL Labs should be done with testing in one to two days.

Further discussion with ARL Labs determined that the capsules would need to be evaluated by their forensic lab as they had been ingested by a person. I was told that this process would take at least ten days. After asking about expedited testing I was able to shorten the process to five days for an additional fee.

Shortly after finalizing testing details with ARL Labs, I received a phone call from Monica with the Nevada State Board of Pharmacy requesting that I cancel my testing. The Board of Pharmacy was going to have the testing performed and we would split the cost. As requested, I contacted ARL Labs and cancelled the testing on the sample I submitted for testing.

Nathan Dahl

Received via email from Nate Dall on 7/1/2021.

Moniea Segeoly

Dahls007



ARL Bio Pharma 840 Research Parkway, Ste. 546 Oklahoma City, OK 73104 (800) 393-1595

6/16/2021

Nevada State Board of Pharmacy Attn: Monica Segedy 431 West Plumb Lane Reno, NV 89509 P: (775) 850-1440

Dear Ms. Segedy,

ARL Bio Pharma Inc. is devoted to providing the highest quality analytical work and problem solving available in the industry. This commitment is supported by the implementation of an ISO 17025 compliant system, FDA registration and DEA license. ARL has completed GMP/GLP studies and customized problem solving to over 1,000 clients.

Attached you will find the requested quotation. We appreciate the opportunity to consult with you about your testing needs and look forward to hearing from you. Please feel free to contact me with any questions or comments associated with this proposal or any future work.

Thank you,

Michael Darnaby Client Services Representative ARL Bio Pharma 840 Research Parkway, Ste. 546 Oklahoma City, OK 73104 T: 405.271.1144

E: mdarnaby@arlok.com



ARL Bio Pharma 840 Research Parkway, Ste. 546 Oklahoma City, OK 73104

(800) 393-1595

Project Overview

Attempt to identify the major pharmaceutical ingredient in two capsules: one labeled as "Pill Box" in a plastic baggie; one labeled as Dalfampridine 5mg. Quantitative analysis will be performed on each capsule should 4-aminopyridine (Dalfampridine) be identified.

Summary of Charges*

Price	Includes	
\$3,500	 Chain of Custody Electronic Documentation of Sample Analysis Certificate of Analysis 	

FOR RESEARCH PURPOSES ONLY - Work conducted under non-cGMP conditions

Payment Terms

Payment to be made upon receipt of signed quote and material to be tested.

Turnaround Time

10 business days

Sending a signed copy of this quotation to ARL certifies that: (1) all information provided in this quotation is true and correct; (2) you have reviewed the Terms and Conditions attached to this quotation; (3) you agree to be bound by the Terms and Conditions; and (4) if you are submitting this quotation on behalf of a company or other entity, you have the authority to bind that company or entity to the Terms and Conditions.

ARL Bio Pharma, Inc.	Nevada State Board of Pharmacy Attn: Monica Segedy
840 Research Parkway, Suite 546	431 West Plumb Lane
Oklahoma City, OK 73104	Reno, NV 89509
T: 405.271.1144 F: 405.271.1174	T: (775) 850-1440
E: mdarnaby@arlok.com	
Submitted by:	Accepted:
Michael Darnaby	
Date: 6/16/2021	Ms. Monica Segedy
	Date:

^{*}Identification and quantification are not guaranteed. Fees will apply regardless of outcome. This work is not intended to be a deformulation. Samples will be disposed of 30 days after the completion of testing. Prior arrangements must be made to return DEA controlled substances to licensed facilities.



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Conduct of the Services. ARL Bio Pharma, Inc., an Oklahoma corporation ("ARL.") will perform testing, prepare a Certificate of Analysis, and all other services agreed to by ARL and Client (collectively, the "Services") in accordance with generally prevailing industry standards of professional conduct. For non-compendial testing. the specification(s) are for informational purposes only For analytical testing, the analyte is reported as it was calculated to derive the result. Client shall verify that the specification and analyte reported are correct for the formulation For Services to be performed pursuant to a Quote. ARL will perform the Services in accordance with the standards set forth in the Quote For cGMP Services, a Quality Agreement may be executed by Client and ARL in addition to the Quote In such instance, ARL will perform the Services in accordance with the Quote. cGMP, and the Quality Agreement ARI will not be required to perform any Services in accordance with cGMP unless a Quality Agreement exists

ARL makes no representations or warranties regarding the release of any Client product. The test results and underlying data of the test results are insufficient to determine whether to release any pharmaceutical products for distribution. The test results and underlying data of the test results only relate to the sample that was tested.

Test Material. Client is responsible for selecting the samples or other materials ("Test Material") that Client sends to ARL for Services in compliance with all applicable laws, regulations, and rules of the relevant governmental regulatory authorities. Client will provide ARL (at no cost to ARL) sufficient amounts of Test Material necessary to perform each test, as well as such data and other information as may be necessary or useful for ARL to perform the Services and to apprise ARI of the stability, proper storage, and safe handling requirements with respect to the Test Material, including a Safety Data Sheet (SDS) or equivalent documentation Client will promptly send to ARL any additional Test Material requested by ARL for completion of the Services

Terms and Conditions

Thirty (30) days following the completion of Services, ARL will discard any remaining Test Material unless Client advises ARL prior to the expiration of the thirty (30) day period that Client wants the remaining Test Material returned and provides ARL with instructions and payment for the return of the remaining Test Material. Client will not use, nor cause another person or entity to use, any Test Material for human or animal consumption or use

Change in Scope. Chent may request a change in scope of any Services, but ARL must agree to such change prior to implementing the change, and ARL may revise the fee for the Services affected by the change in scope

Cancellation of Routine Testing.
Client may cancel a routine test at any time prior to ARL's commencement of the routine test. In such event, ARL, in its sole discretion, may charge a cancellation fee of \$20 per canceled test for any testing canceled by the Client after ARL's receipt of the relevant Test Material

Cancellation of On-Going Studies.
Client may cancel any on-going studies performed by ARL at any time without cause upon fifteen (15) business days prior written notice to ARL. In such event, Client shall pay ARL for all Services rendered through the effective date of termination, together with any additional expenses incurred by ARL in connection with the termination of the study, including those which were previously committed to by ARL for completion of the study.

Personnel. To the best of ARL's knowledge, none of its employees who will participate in testing have been debarred, or are under consideration to be debarred, by the Food and Drug Administration from working in or providing Services to any pharmaceutical or biotechnology company under the Generic Drug Enforcement Act of 1992, as amended

Inspections. Once per year, upon thirty (30) days advance notice to ARL. Client or its designated representative, if such

representative is reasonably acceptable to ARL, may visit ARL's facilities to observe the testing. The visit must be during normal business hours and occur at a mutually agreeable time.

Test Records and Reports. ARL will keep complete and accurate records of each test for five (5) years after completion of the test

Fees. Chent shall promptly pay all fees for services when due and payable All payments must be in US Dollars If Client requests a rush for the performance of any Service, ARL may in its sole discretion, add a surcharge to the rushed Services For Services not specified in a Quote, such as routine testing, each new Client must prepay all charges for at least sixty (60) days after commencing business with ARL in order to establish a purchase history Prepayments can be made via check, credit card, or wire transfer After the sixty (60) day period. Client may request a credit review Once ARL establishes a credit limit for Client, ARL will invoice Client for Services and Client must pay each invoice within fifteen (15) days of the date of the invoice

For Services performed pursuant to a Quote. Client must pay the amounts specified in the Quote. The pricing of each Quote is valid for mnety (90) days from the date of the Quote. Client shall pay all invoices and other amounts due under the Quote within thirty (30) days of receipt of the relevant invoice unless otherwise specified in the Quote. Any changes in the fees must be mutually agreed to by the parties in a written amendment to the Quote.

All fees for all Services, whether or not performed pursuant to a Quote, must be paid by the applicable due date. All fees not paid will bear interest at a rate of one and one-half percent (15%) per month from the applicable due date until paid If Client does not pay each invoice when due, ARL may elect to suspend any Services, including, but not limited to, any testing that may be in progress, delaying the start of new testing, and withholding reports or other deliverables



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Additionally, Client shall reimburse ARL for all costs related to collection of unpaid fees, including reasonable attorneys' fees and costs

Subcontractors. ARL may outsource or use contractors for any or all Services

Confidentiality. If the parties have executed a confidentiality agreement prior to the commencement of Services, that confidentiality agreement will control the disclosure of confidential information between the parties through the performance of Services. If the parties have not executed such an agreement, these Terms and Conditions control the exchange of Confidential Information between the Parties

In the event there is no confidentiality agreement between the parties, the parties anticipate that they may exchange proprietary and confidential information (the "Confidential Information") related to the performance of Services All Confidential Information must be identified in writing as confidential Each party will use commercially reasonable efforts to maintain the other party's Confidential Information in confidence and will employ reasonable procedures to prevent its unauthorized publication or disclosure to third parties No party may use the other party's Confidential Information for any purpose other than performance of the Services

Warranties. Client warrants that it owns all rights, title, and interest in and to all Test Material and intellectual property related thereto, and that ARI s use of any and all such Test Material in connection with the Services does not infringe any copyrights, patent rights, trade secrets, or other intellectual property rights of any third party. Client also warrants that it will comply with all applicable laws, regulations, and rules of the relevant governmental regulatory authorities related to the sale, distribution, final product release, or other use of any Test Material.

ARL warrants to Client that all Services provided to Client will be in accordance with generally prevailing industry standards of professional conduct and comply with all applicable laws, regulations, and rules of the relevant governmental regulatory authorities 1f Services are performed pursuant to a Quote, ARL also warrants that the Services will conform to the specifications in the Quote These warranties of ARL are made only to Client, are not transferable, and do not extend to the benefit of any other person or entity. OTHER THAN THE FOREGOING WARRANTIES, THE SERVICES ARE SOLD AND PROVIDED "AS IS," WITHOUT WARRANTY OF ANY KIND, WHETHER STATUTORY, EXPRESS. OR IMPLIED THE WARRANTIES PROVIDED IN THIS PARAGRAPH ARE ARL'S SOLE AND EXCLUSIVE WARRANTIES WITH RESPECT TO THE SERVICES AND IN LIEU OF ALL OTHER WARRANTIES, WHETHER STATUTORY, EXPRESS. OR IMPLIED ALL OTHER WARRANTIES ARE EXPRESSLY DISCLAIMED, INCLUDING. WITHOUT LIMITATION, ANY IMPLIED WARRANTY OF MERCHANTABILITY, FITNESS FOR A PARTICULAR PURPOSE, NON-INFRINGEMENT, AND RESULTS OBTAINED (INCLUDING, WITHOUT LIMITATION, ANY CLAIM OF INACCURATE, INVALID, OR INCOMPLETE RESULTS), WHETHER ARISING BY STATUTE, OTHER SOURCES OF LAW, OR FROM COURSE OF PERFORMANCE OR DEALING, OR USAGE OF TRADE

Limitation of Liability. ARL WILL NOT BE LIABLE FOR PENALTIES OR LIQUIDATED DAMAGES, OR SPECIAL, INDIRECT, INCIDENTAL. CONSEQUENTIAL COLLATERAL, PUNITIVE, EXEMPLARY, OR OTHER DAMAGES OR LOSSES OF ANY TYPE OR KIND (INCLUDING, WITHOUT LIMITATION, LOSS OF USE AND LOST PROFITS) REGARDLESS OF WHETHER ANY SUCH LOSSES OR DAMAGES ARE CHARACTERIZED AS ARISING FROM BREACH OF CONTRACT, BREACH OF WARRANTY, TORT STRICT LIABILITY, OR OTHERWISE, EVEN IF ARL IS ADVISED OF THE POSSIBILITY OF SUCH LOSSES OR DAMAGES, OR SUCH LOSSES OR DAMAGES ARE FORESEEABLE

NOTWITHSTANDING ANYTHING TO THE CONTRARY IN THESE TERMS AND CONDITIONS, CLIENT'S SOLE AND EXCLUSIVE REMEDY FOR ARL'S BREACH OF WARRANTIES SET FORTH IN THESE TERMS AND CONDITIONS WILL BE, AT ARL'S SOLE AND ABSOLUTE DISCRETION: (i) RE-PERFORMANCE OF THE SERVICES AFFECTED BY THE BREACH OF WARRANTY AT ARL'S SOLE COST AND EXPENSE, OR (ii) REFUND OF THE SERVICE FEES PAID TO ARL BY CLIENT FOR THE SERVICES AFFECTED BY THE BREACH OF WARRANTY FOR ALL OTHER CLAIMS ASSERTED BY CLIENT AGAINST ARL RELATED TO THE SERVICES, THE APPLICABLE QUOTE (IF ANY), OR THESE TERMS AND CONDITIONS (INCLUDING CLAIMS FOR INDEMNIFICATION). ARL'S MAXIMUM LIABILITY FOR ANY DAMAGES OR LOSSES. REGARDLESS OF THE FORM OF ACTION OR PROCEEDING, WILL NOT EXCEED THE TOTAL SERVICE FEES PAID BY CLIENT FOR THE SERVICES GIVING RISE TO THE DAMAGES OR LOSSES

Indemnification. Subject to the Limitation of Liability contained herein, ARL shall indemnify Client and its respective directors, officers, employees. and agents (collectively, the "Client Indemnitees") from and against any losses, damages, fines, and liabilities, including attorney fees and litigation expenses (collectively, "Damages"), incurred by the Client Indemnitees as a result of any third-party claims. demands, suits, actions, or causes of action (collectively, "Claim") arising from. (i) ARL's breach, violation, noncompliance, or non-performance of these Terms and Conditions or Quote (if applicable), and (ii) ARL's gross negligence or willful misconduct in the performance of Services ARL will pay any Damages subject to the Limitation of Liability set forth herein which, by final judgment, after exhaustion of all reasonable appeals, may be assessed against the Client Indemnitees in connection therewith



ARL Bio Pharma 840 Research Parkway, Ste. 546 Oklahoma City, OK 73104 (800) 393-1595

Client shall indemnify and defend ARL and its respective directors, officers, employees, and agents (together, the "ARL Indemnitees") from and against any third-party Claim, and any Damages resulting from such Claim, against an ARL Indemnitees arising from (1) Client's breach, violation, noncompliance, or non-performance of these Terms and Conditions or Quote (if applicable). (ii) Client's gross negligence or willful misconduct, (iii) the marketing, labeling, recall, manufacture. distribution, use, sale, or other disposition by Client or any distributor, customer, sublicensee, or representative of Client, of any Test Material, product. process, technology, or other material or information that Client provides to ARL (collectively, the "Client Supplied Materials and Technology"). (iv) any assertion that the Client Supplied Materials and Technology or an ARL Indemnitee's use of the Client Supplied Materials and Technology infringes the know-how, trade secrets, patent rights, copyrights, or other intellectual property rights or confidential information rights of a third party If Client breaches its duty to defend an ARL Indemnitee against such a third-party Claim, Client shall reimburse that ARL Indemnitee for the reasonable attorney fees and litigation expenses incurred by that ARL Indemnitee in defending the Claim, and the reasonable attorney fees and litigation expenses incurred in recouping the defense attorney fees and litigation expenses from Client

Ownership. ARI, will exclusively own all techniques, methods, processes, models, tools, assays, test results, and the underlying data of the test results that are developed, generated, conceived, or utilized in the performance of the Services

Licenses. Client grants to ARL a nonexclusive, irrevocable, fully paid-up, worldwide license (including the right to sublicense to any subcontractor for that subcontractor's performance of any Services) to use and duplicate any proprietary technology and Test Material disclosed to ARL solely to the extent necessary to perform the Services ARL grants to Client a non-exclusive, irrevocable, fully paid-up, worldwide license (including the right to sublicense) to use, duplicate, and disseminate the test results and underlying data of the test results that are disclosed by ARL to Client in connection with the Services

Controlling Terms. In the event that there is any conflict between these Terms and Conditions and the Quote, the terms in the Quote will apply

Independent Contractor. The business relationship of the parties is that of independent contractors and not of partners, joint venturers, employers, employees, or any similar kind of relationship

Force Majeure ARL will not be hable for any delay or failure of performance. including, without limitation, failure to perform a Service, where such delay or failure arises or results from any cause beyond ARL's reasonable control, including, but not limited to, flood, fire, explosion, natural catastrophe, military operations, war, computer or other equipment failure, severe weather, earthquake, tomado, or other act of God, power loss or reduction, labor disputes of any kind (whether relating to its own employees or others), embargos, governmental regulation, or an inability or delay in obtaining materials In the event of any such delay or failure of performance, ARI will have additional time to perform the Services as reasonably necessary under the circumstances

Applicable Law, Jurisdiction, and Venue. The Services, these Terms and Conditions, and any applicable Quote are governed by, and construed in accordance with, the laws of the State of Oklahoma, USA, without regard to any choice of law principle that would dictate the application of the law of another jurisdiction. Venue of all disputes regarding the Services, these Terms and Conditions, or an applicable Quote must be brought in the District Court for the State of Oklahoma, Oklahoma County Each party waives any right to or option for a trial by jury

Shortened Statute of Limitations. Any claim against ARL for breach of warranty, or any other claim related to the Services, a Quote, or these Terms and Conditions (including a claim for indemnification), must be brought within one (1) year from the date the cause of action arose

Entire Agreement. These Terms and Conditions and the Quote (if any) constitute the complete, final, and exclusive expression of the agreement between the parties, superseding any and all previous agreements and understandings, whether oral or written

Modification and Waiver. No modification or waiver of the provisions of these Terms and Conditions or a Quote will be valid or binding on either party unless set forth in a writing signed by both parties. No waiver of any term, right, or condition of these Terms and Conditions or a Quote may be construed or deemed to be a waiver or continuing waiver of any such term, right, or condition on any subsequent occasion, or a waiver of any other term, right, or condition

Severability If any of the provisions of these Terms and Conditions or an applicable Quote are deemed to be invalid or prohibited under applicable law, such provisions will be ineffective to the extent of such invalidity or prohibition, without invalidating the remainder of such provision or the remaining provisions of these Terms and Conditions or the Quote

Voluntary Agreement Each party represents that they have carefully read and understand all provisions, terms, and aspects of these Terms and Conditions and the applicable Quote (if any), and have knowingly and voluntarily agreed to be bound by them Each party also represents that they have had the opportunity to review these Terms and Conditions and the applicable Quote (if any) with legal counsel of such party's choice

Revised 6/2018



9901 S. Wilcrest Drive Houston, TX 77099-5132 800.331.2498 | pccarx.com

SOLD TO: 00014667

DAHL'S PHARMACY OF FERNLEY NATHAN DAHL P.O. BOX 5160 FALLON, NV 89407 UNITED STATES OF AMERICA

Thank you for being a member. Total Savings: 0.00

INVOICE

Invoice: 12713362

Page: 1 of 1

Invoice Date:	08/17/20	Print Date:	08/17/20
Order Date:	08/17/20	Ship Date:	08/17/20
Sales Order:	2689346	PO Number:	
BOL:	7169974		
Ship Via:			
Credit Terms:	10P		
Salesperson:	00060034 0006	0075	

SHIP TO: 14667-1

DEA: FD5763203

DAHL'S PHARMACY OF FERNLEY NATHAN DAHL 805 EAST MAIN ST. FERNLEY, NV 89408 UNITED STATES OF AMERICA

Item#	UM	Description	Shipped	B/Ord	Price	Total Discount	Total Ne
50-5030-10	GM	Description DALFAMPRIDINE USP(4-AMINOPYRIDINE)	Shipped 1.0	0.0	210.00	0.00	210.00
Site: LV							
	_				n	et Savinge: 0.00	

Please remit to

PCCA PO Box 734687 Dallas, TX 75373-4687

Billing questions call 1-800-221-8768

You are responsible for properly disclosing and appropriately reflecting all discounts and price reductions in cost reports and claims submitted to federal health care programs (including Medicare and Medicaid) and other reimbursing agencies, for maintaining documentation and records of such discounts and price reductions, and for providing such records to government representatives on request, in accordance with all applicable laws and regulations. The prices shown on this invoice may be subject to subsequent rebates.



9901 S. Wilcrest Drive Houston, TX 77099-5132 800.331.2498 | pccarx.com

SOLD TO: 00014667

NATHAN DAHL P.O. BOX 5160 FALLON, NV 89407 UNITED STATES OF AMERICA

DAHL'S PHARMACY OF FERNLEY

INVOICE

Invoice: 12713362 Page: 1 of 1

	The state of the s		
voice Date:	08/17/20	Print Date:	08/17/20
Order Date:	08/17/20	Ship Date:	08/17/20
Sales Order:	2689346	PO Number:	
BOL:	7169974		
Ship Via:			
redit Terms:	10P		
alesperson:	00060034 00060	0075	
Ship Via:	10P	0075	

SHIP TO: 14667-1

DEA: FD5763203

DAHL'S PHARMACY OF FERNLEY NATHAN DAHL 805 EAST MAIN ST. FERNLEY, NV 89408 UNITED STATES OF AMERICA

Thank you for being a member. Total Savings: 0.00

Item#	UM	Description	Shipped	B/Ord	Price	Total Discount	Total Ne
50-5030-10	GM	Description DALFAMPRIDINE USP(4-AMINOPYRIDINE)	Shipped 1.0	0.0	210.00	0.00	Total Ne 210.00
Site: LV							

Please remit to PCCA PO Box 734687 Dallas, TX 75373-4687

Billing questions call 1-800-221-8768

Product Savings: 0.00 Invoice #: 12713362 Net Line Total: 210.00 Cust #: 00014667 Freight: 9.89 0.00 Freight Discount: Sales Tax/GST: 0.00 USD INVOICE TOTAL: 219.89

You are responsible for properly disclosing and appropriately reflecting all discounts and price reductions in cost reports and claims submitted to federal health care programs (including Medicare and Medicaid) and other reimbursing agencies, for maintaining documentation and records of such discounts and price reductions, and for providing such records to government representatives on request, in accordance with all applicable laws and regulations. The prices shown on this invoice may be subject to subsequent rebates

/10/2021 2:33:09 FM age 1	1.138.0m/m	221396						Filles Rx 50	5110121 170945
alfampridine (4-aminopyr	udinej sing s	LOW RELEASE	#1 CAPSU	ILES 5 MG	CAPSU	LE .			102021016
Tall Man:								Cet 07	(0 222,122
Flavor;					S	chedule:	4902		Activo 🗸 .
Description: Quantity made: 100 CAPS		Batch yield:	100 000		D	OCA ID.		Formula	ID: 4902 ID: 21396
equality mass. 100 On 6		ity remaining:	100.000			CCA ID:		Lug I	D. 21030
Date made: 5/10/2021		•		ricing cal				g	
Lot number: 05102021@ eyond use date: November 6	. 2021	2:28 PM	Es	timated pri	ce	\$30.00	as of		
Pharmaclet: NATHAN DA	s effer compound	ding date					Time t	make:	0
Technician: - <none></none>		_					i iiie ti	Jillake.	٠
Packaging:									
Equipment:									
Labeling: bility information:									
redients		Sch.	Quantity	used	QS /	Balance	1		
DALFAMPRIDINE USP POWDER		70111		0.5 GMS	П 0.				
Lot il: C195958 Chemical Code:	Mig: Volume	Potency:	Exp. date:	3/31/2025		hlsr:		AWP:	60.00
Purity:	(mag-			Bar code cho	cked:	Each CAPS	contains O.	005 GMS or 0.5	%
METHOCEL EAM PREMIUM (HYP	ROWELLOSEL	ISP) 1		NDC: 10 GMS	[] 10	.001 g	С	hemInvID: 13	86
Lot #: 149262/E Chemical Code:	Mig: MEDISC	A	Exp. date:	7/31/2022		nisr. MEDI			
Purity:	Volume	Potency:	QS emo	7.00		Fach CAPS		WP: GMS or 10%	\$26.00
				NDC: 519:	27-1188-0	00		neminviD: 13	46
AVICEL PH-105 POWDER Lot #: 1805020016	Mg:	L	Exp. date:	.5 GMS 2/3/2022	□ 11 W	.500 g	0		
Charried Code: Purby:	Volume:	Potency:	QS amo	nt			A	WP;	\$9.31
CCC				NDC: 6296	xxx: √ 31-2007-0	Each CAPS o		15 GMS or 11.5 eminviD: 127	
Capsule #1-Clear Locking of Lol #:	CAPSULE	L		00 CAPS	D	100c	985		-
Chemical Codet	Mfg: Volume:	Potency;	Exp. date: Q5 amos	rt	Wh	isi.	^	WP:	\$0.00
Purity:				Bar code ched NDC:	ked:	Each CAPS o	ontains 1 C	APS or 100% ominviD:	
				NDC,	-		Cit	GININVID:	
Instructions & Notes									
Log Originally made as: 100 I	DALFAMPRID	NE (4-AMINOPY	RIDINE) 5	AG SLOW R	RELEAS	E#1 CAF	SULES	5 MG CA	PSULE
nule ID: 4902 Salculated lot number: 0510202	1608 Beyond	usa date: 11/6/2	021						
MILL & MOTELLICATIONS.						. 4		II antico	eve or
dd food color to Active mical is a consistent co	(Darrampri	oine) and mo	ined wit	h the foo	d cold	e to ens	uie a	ii active	avl.
nicai is a consistent co	ioi and con	ipietely com	JINGU WI	11 110 100	u con	<i>.</i>		ner	2 10
dd Avicel, Methocel, an	id Acitve (D	alfampridine) to a zip	lock bag	and	seal.		- P. S.	I CT V
									with them
sing rolling bar go back	and forth	over the zip lo	ock bag	until all p	owde	r is the	same	color to	W
re complete dispersion	of active.						1/01	:1-1n	A HAND
	محمد محمد	a acastila m	ahina a	nd anaon	culat	ueind	VU	in the	11014
to the total				nu encap	Sulate	using	Size 1	+1 capsu	les.
		M: 5/10/2021 2:2	1.59 PM		1	110	th	ue of	NYTYP
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entered: 6/10/2021 2:27:59 PM	Last modific	o d	Date:	/	()		118	hing	NOA
Dump powder from zip lo entered: 6/10/2021 2:27:59 PM ked by:	Lagsmodific	ber	Date:	V /	7).	1 3	fes	ting	OV QA
entered: 6/10/2021 2:27:59 PM	Last-modific	bera	Date:	y / 1	7).	3	fes	ting	or QA
entered: 6/10/2021 2:27:59 PM	Last modific	ber	Dats:	y /	7).	8	fer	ting you	OV QA Compor

	1980 Formula VV 2021 11:31:45 AM	orksheet (=								1/19/21	
-	AMPRIDINE (& AUDINO	DPYRIDINE) 5M	0 SLOW RELEAS	SEGIC/	PSULLS 5 MG	CAPSI	ILE		1 1 1 0°	उधायका व	LI III.
								*		12020218	
T111	dan:										
Do	Flaver:					8	Schedule:	4902	F	Active 🗹	. 1
1	ntity made: 100 C	APS	Batch yiel Qty remainin	d: 100.	000		CCA ID:		Formula Log I	D; 4902 D; 21248	
	Date made: 4/20/20		any romaning	g. 100.	Pricing c		11 11 11 11 11 11 11 11	the lo	q		
Boyo	Lot number: 042020 nd use date: Novem 18 Pharmacist: NATHA	121@2 ber 5, 2020 0 days efter como	11:09 cunding date	AM	Estimated p	rice	\$30.00	as of			
	Pharmacist: NATHA Technician: - <non< td=""><td>N DAHL '</td><td></td><td></td><td></td><td>1</td><td></td><td>Time to</td><td>make:</td><td>0</td><td></td></non<>	N DAHL '				1		Time to	make:	0	
	NDC1: Packaging: Equipment		-								
ын	Labeling:										3
redi	ents		Sch.	Quai	ntity used	QS I	(Balance	2)			
	AMPRIDINE POWDER			Tour 1	0.5 GMS	0 0	500 g				
Lote	Charried Code:	Mfg: Vokume:	Potency:		data: Semount	W	hisr:				1.0
nce.	dy.	177	-3.5		Bar code d NDC:	hedead: 🗹	Éach CAPS	contains 0.0 Ch	CS GMS or 0.53 eminvio:	4	
	OCEL (R) E4M PREMI		ELLOSE L L	row.	10 GMS		.002 g				
	Chemical Code:	Wifg: Volume:	Potency:		dale: 11/5/2020 9 mount	VV	hlsr:			1	
THE PL	101.				Ber code di NDC: 51	1927-1 188-1	Each CAPS	contains 0.1 Cha	GMS or 10% eminviD: 106	4 .	81
	EL PH-105 POWDER	9.860	L	P.m.	11.5 GMS	D 11	.500 g	-		la:	
	1805020016 Chemical Code:	Mig: Volume	Potency:		date: 2/3/2022 Senount	W	hlsr. LETC		MP;	\$9.31	
KIT.	37:				Bar code ch NDC: 62	ocked: [V]	Each CAPS	centains 0.1	15 GMS or 11.6 eminviD: 127	1,000,000	1.
	ULE #1-CLEAR LOCK!		·L	-	100 CAPS		100 CM	Qs.			
Lot #	Chamical Code:	Mfg; Volume:	Potanoy:	Exp. d	Sate:	Wh	lsr.	7			
Pur					Ber code che NDC:	ecked:	Each CAPS o		PS or 100% minvID:		
ins	ructions & Notes									,	
mula	Originally made as: 1 ID: 4902 ated lot number: 0420				5 5MG SLOW	RELEAS	E#1 CAF	SULES	5 MG CAI	PSULE -	
FOR	KULA INSTRUCTIONS	8:			12 v	1277					3
. Ac	ld food color to A nical is a consist	Active (Dalfa ent color and	mpridine) and i completely o	mix w	ith mortar a ed with the	ind per food c	stle to e olor.	ensure	all activ	e	7
	d Avicel, Method										1
	ing rolling bar go re complete disp			zip lock	bag until a	ll pow	der is th	ne sam	e color	to	7
	mp powder from			e mach	ine and en	capsul	ate usir	ng size	#1 cap	sules.	
	THE PERSON OF THE		A 725 5 1 2 8 0 3 2 2 3			70					17.0
			*								
		A11 1	Bied: 4/20/2021 11	:31:42 AM			vi				1
ante ad t	red: 4/20/2021 11:09:41 /:	LESS HAX	MINOUDEVIEWE! !!	Da							,
- 1						,					

egged Formula Worl 12021 1:18:40 PM	ishaal (inti	D 1220506			Filler 4/191	121
ge 1					Rx 507094	15
Fampridine (4-aminopy	ridine) sing :	SLOW RELEASE	en capsules 5 mg c	APSULE	Let 03012021	
all Men:					- 14 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	-
Flavor:				Schedule: 4	902 Active	171
Description: Quantity made: 100 CAP	9	Dateb worlds	100 000	THE STATE OF	Formula ID: 4902	500
aunity made. 100 O/II		Batch yield: City remaining:		PCCA ID: Route of admin:	Log ID: 2090	6
Date made: 3/1/2021 Lot number: 03012021			Pricing cale	ulations from th	e log	
ryond use date: August 28.	2021	1:18 PM	Estimated price	e \$30.00 as	of	
Pharmacist NATHAN		nding date		-		
Technician: - <none></none>		_		in	ne to make: 0	
Packaging: Equipment:					0.7	_
Labeling:						
bility Informa tion: redients		Sch.	Quantity used	QS (Balance)		
DALFAMPRIDINE USP POW	DER	SCII.	0.5 GMS	0.500 g		-
Lot 8: C195988 Chemical Codes	Mig: Volumes	Datasas	Exp. date: 3/31/2025	Whisr:	AWP: 50	
Purity:	VOMEN	Petency:	Bat code ch NDC:	ected: Each CAPS or	entains 0.005 GMS or 0.5% CheminvID: 1386	,00
METHOCEL EAM PREMIUM (F			10 GMS	10.000 g		_
Lot #: 149262/E Chemical Code:	Mig: MEDI	Potancy:	Exp. date: 7/31/2022 Q3 enour2	Whisr: MEDIS	AWP: \$26.	.50
Purity: por.			Bar code ch	ocket 2 Each CAPS co	ntains 0.1 GMS or 10% CheminviD: 1346	>
AVICEL PH-108 POWDER		L	11.5 GMS	☐ 11.500 g	CHEMINIO. 1340	-
Lot #: 1805020018 Chemical Code:	Milg: Volume:	Potency	Exp. date; 2/3/2022	Whisr: LETCO	AWP: 59.	74
Purity:	7		200,000,000	ched: Each CAPS co	nitains 0.115 GMS or 11.5% CheminviD; 1275	
CAPSULE #1-CLEAR LOCKING	CAPRILE		100 CAPS	91-2007-01	CheminvID; 1275	_
Lot #: Chemical Code:	Mfg:		Exp. data:	Whisr.	₽ 	
Purity:	Volume	Polandy:	QS amount	cked: Each CAPS cor	AWP: 50.0	00
			NDC:	AND LEGITOR S CO.	CheminviD:	-
Instructions & Notes	-30.00					
Log Originally made as: 10 nuta ID: 4902	O DALFAMPRI	DINE (4-AMINOP	YRIDINE) 5MG SLOW	RELEASE #1 CAPS	SULES 5 MG CAPSUL	E
Calculated lot number: 03012	2021@4 Beyon	nd use date: 8/28/	2021			
rmula instructions: Idd food color to Activ	a (Delfemn	ridine) and m	ix with mortar and	pestle to ensi	ure all active	
mical is a consistent	color and co	impletely com	bined with the for	od color.	75 - 407 (33-2-7-7)	
dd Avicel, Methocel,						.7
lsing rolli ng bar g o ba ure com plete dis pers	ick and forth	over the zip	lock bag until all p	oowder is the s	same color to	
	Laste bon o	nta canoula m	achine and enca	osulate using s	size #1 capsules	2
ump powder from zip				La contract (Section)	The state of the s	
entered: 3/1/2021 1:18:28 PM		Hed: 3/1/2021 1:18	237 PM	1		
				a)		

3	/1/2021 1:18:10 PM	200			Filles 3/1	21
	age 1	2030904			12x 5070	945
1	usampridine (C-almopyridine) elic	BLOW RELEASE	M CAPSULES 6 MG	CAPSULE		0370215
ı					Let 630	
	Tell Wan:				وط د م	יו גטגומ
	Player: Description:			Schedule:	4902	Activa 🗸
I	Quantity made: 100 CAPS	Batch yield:	100,000	PCCA ID:	Formula	ID: 4902 ID: 20904
	Date made: 3/1/2021	Ony remaining:	100.000	Route of admin:		D. 20904
	Lot number 03012021@3		Prieing ca	iculations from	the log	
	Pharmacist: NATHAN DAHL	12:11 PA	Estimated pr	rice \$30.00	as of	
	Pharmacist: NATHAN DAHL Technician: - <none> NDC1:</none>				Time to make:	0
	Packaging: Equipment:					
	Labeling: ullity information:					
	redlents	Sch.	Quantity used	QS' (Balance	e)	
	DALFAMPRIDINE USP POWDER		0.5 GMS	□ 0.501 g	5)	-
	Lot 5: C195988 Mig: Chemical Code: Volume:	Polency:	Exp. date: 3/31/2025	Whisr;	AWP:	447
	Puty:		Ber code d	hadred: VI Each CAPS	contains 0,005 GMS or 0.	\$0.00
	METHOCEL EAM PREMIUM (HYPROMELLOSS	1110ch 1	NDC:		CheminviD: 1	386
	of 6: 149202/E Mitg: MEDI Chemical Code: Volume:		10 GMS Exp. date: 7/31/2022 08 emount	Whisr. MED	ISCA AWP:	\$26.90
			Bar code ch	ecked Each CAPS	contains 0.1 GMS or 10%	
	VICEL PH-105 POWDER of \$: 1805020018 Mfg: Charlest Code: Volume:	L Potansy:	11.5 GMS Exp. date: 2/3/2022 OS emount	11.500 g Whisr: LETC	CheminyiD: 1:	y U
	Purity:	10.50	Bar code ch	ocked: M Each CAPS	contains 0.115 GMS or 11	\$9,31
	APSULE 51-CLEAR LOCKING CAPSULE		NDC: 82	991-2007-01	CheminviD: 12	75
	ol #: Mfg:		100 CAPS Exp. date:	Whist.	400	
	Chamical Code: Volume Furity:	Potency:	QS amount		AWP;	\$0.00
			Bar code che NDC:	ecked: Eech CAPS of	contains 1 CAPS or 100% CheminvID;	
	Instructions & Notes					
	Log Originally made as: 100 DALFAMPRII uda ID: 4902 siculated lot number: 03012021@3 Beyon MULA INSTRUCTIONS:			RELEASE #1 CAP	SULES 5 MG CAI	PSULE
	dd food color to Active (Dalfampi nical is a consistent color and co				sure all active	
	dd Avicel, Methocel, and Acitve (Dalfampridine)	to a zip lock ba	g and seal.		
	sing rolling bar go back and forth re complete dispersion of active		ck bag until all p	owder is the	same color to)
		do concula ma	chine and ancar	naista usina	size #1 cans	iles
1	imp powder from zip lock bag or	ili capaule illa	Crime and encap	Journale Using	oize ir i oapsi	uico.

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egged Formula Work 3/2021 11:52:05 AM	shoot (===	EG20761			Filled	2/3/21
ge 1					BX 50	70945
Fampridine (4-alinopy)	ridine) sug	SLOW RELEASE (M CAPSULES 6 MG C	APSULE	Lot 12	092020 D4 1
					Lot oz	C3702123 #
Il Man:	-			22.00	0.62	
Flavor: Description:				Schedule:	4902 Formula	Active V
uantity made: 100 CAPS	3	Batch yield:	100.000	PCCA ID:	-00.000	ID: 20761
D-11 0D-004		Qty remaining:	100.000	Route of admin:		
Date made: 2/3/2021 Lot number: 02032021@	23		Pricing cale Estimated price	culations from 8 \$30.00		
wond use date: August 2. 2		11:46 AM	Countained price	25 \$50.00	a5 UI	
. Pharmacist: NATHAN I	SAHL	——			Time to make:	0
Technician: . NDC1:		-			Time to make.	0
Packaging:						
Equipment						
Lebeling:						
ility Information:		2.0	ALCOHOL: CONTRACT	22 43 2		
edients	en .	Sch.	Quantity used	QS (Balan	ce)	
DALFAMPRIDINE USP POWD Let 0: C195988	Mig:		0.6 GMS Exp. date: 3/31/2025	0.500 g Whisr:	-	
Chemical Code: Purity:	Volume	Potency:	OS amount		AWP:	20.00
*			Bar code et NDC:	sector Ench CA	PS contains 0.005 GMS (CheminviD	× 0.5%
ETHOCEL EAM PREMIUM (H			10 GMS	□ 10.002 g		. 1500
of #: 149282/E Chemical Code:	Milg: MED	Fotonoy:	Exp. date: 7/31/2022 Q8 emount	Whisr: ME	DISCA AWP:	
Purky:	. 15 2000	1 4		ecked IVI Each CAS	S contains D.1 GMS or 1	\$26.90
			NDC: 51	927-1186-00	CheminviD	1346
VICEL PH-105 POWDER of #: 1805020018	Mfg:	L	11.5 GMS Exp. date: 2/3/2022	U 11.501 g Whisr. LE	rco	
Chemical Code: Purity:	Volume	Potency:	QS amount	William EC	AWP:	\$9,31
r any.			Bar code the	cked: Esch CAP	3 contains 0.115 GMS of CheminyiD:	11.5%
APSULE SI-CLEAR LOCKING	CAPSULE	L	100 CAPS	D /00	CheminviD:	12/5
ol #: Chemical Code:	Mfg:	Datasas	Exp. date:	Whisr:	Sal s	
Putty:	Volume	Potency:	QS amount	4.4T 5.4.616	AWP:	\$0.00
			Ear code che NDC:	existi Li Each CAP	S contains 1 CAPS or 10 CheminviD:	0%
Instructions & Notes						
Log Originally made as: 10	O DALFAMPR	IDINE (4-AMINOPY	RIDINE) 5MG SLOW	RELEASE #1 C	APSULES 5 MG	CAPSULE
nda ID: 4902 alculated lot number: 02032	021@3 Bevo	and use date: 8/2/20	121			
MULA INSTRUCTIONS:				No.	and the same	
id food color to Activ	e (Dalfamp	ridine) and mix	with mortar and	pestle to er	isure all activ	<i>l</i> e
nical is a consistent of	color and o	ompletely com	bined with the foo	od color.		
		(D-1/2	Var a min lands ha			
ld Avicel, Methocel, a	and Active	(Danamphoine) to a zip lock ba	g and seal.		4
			ante boo until all r	ouder is the	came color	to
sing rolling bar go bac re complete dispersi	on of active). Over the zip it	ow has mini all t	OWUGI IS LIR	same color	10
mp powder from zip			achine and encar	sulate usin	g size #1 car	sules.
intored: 2/3/2021 11:48:35 AM		ified: 2/3/2021 11:46				2000
nd by:	Lest mou	med Borava 11.70	Date:/_/	_		
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Logged Formula Wor 129/2020 2:37:59 PM	ksheet (zaca	ra) [[7]				Filler	12/1/20	
Page 1 .		1420N32				Ry 1	613266	
DALFAMPRIDME (4-AMINOP	VRIDINE) SMG	SLOW RELEASE	M CAPSULES 5 MG	CAPSULE			11052020 a	051
						Lot 1	20920202	a 4 !
Tall Man:				2.10100				
Description:	20			Schedule:	4902		Active VID: 4902	
Quantity made: 100 CAF		Batch yield: Qty remaining:		PCCA ID: Route of admin:		Log	ID: 20422	
Date made: 12/9/2020	0	and contamination	Pricing cal	culations from	the loc			
Beyond use date: June 7, 2	021	12:36 PM	Estimated pri	ce \$30.00	as of			
Pharmacist: NATHAN Technician: NDC1:	lays after compour DAHL	nding date		19	Time to	make:	0	
Packaging: Equipment:								
Lebeling: Stability information:								
Ingredients		Sch.	Quantity used	QS (Balan	ce)			
DALFAMPRIDINE USP POW Lot 6: C195989 Chemical Codes	VDER Mig: Votass	Polanoy:	0.5 GMS Exp. date: 3/31/2025 OS emount	O.501 g Whisr:		AWP:	\$0,00	
Balmor Purty:			Ear code o NDC:	thackent D Each CAS	S contains (0,005 GMS o	r 0.5%	
METHOCEL EAM PREMIUM Lot #: 149282/E Charles Code:	(HYPROMELLOSE Mfg: MEDI		10 GMS Exp. date: 7/31/2022	10.001 g Whisr: ME	DISCA	AWP:		-
Educa: Pully:			Bar code o	hecked	S contains 0	0.7175.5	\$26.90 0% 1346	
AVICEL PH-108 POWDER Lot \$: 1805020018 Chemical Codiz Purity:	Mig: Volume:	L. Patency:	11.5 GMS Exp. date: 2/3/2022 QS amount	Whise LET		AWP:	\$9.31	-
Believe			Bar code ch NDC: 63	10ckod: M Each CAP 2691-2007-01	S contains 0.	115 GMS cr	11.5%	
CAPSULE #1-CLEAR LOCKIN	NG CAPSULE Mfg: Volume	L Polancy:	100 CAPS Exp. date: QS emount	□ /co Whisr:	CAPS			-
Balance.	your.	rouncy.		acked: Each CAPS	S contains 1	AWP: CAPS or 100	\$0.60	
			NUC:		Ci	reminviD:		-
Log Instructions & Notes								
New Log. Originally made as: 1 Formula ID: 4902 Calculated lot number: 1205 FORMULA INSTRUCTIONS: 1. Add food color to Action	92020@4 Beyon	nd use date: 6/7/2	021 x with mortar and	i pestle to er				
2. Add Avicel, Methocel	, and Acitve	(Dalfampridine	e) to a zip lock ba	g and seal.			,	2
. Using rolling bar go bansure complete dispen			lock bag until all	powder is the	e same	color	to	-41
Dump powder from zi	in lock bag o	nto capsule m	achine and enca	psulate using	g size i	#1 can	sules.	
Date entered: 12/9/2020 12:36:22		fied: 12/9/2020 12:			7 7 7 7 7		C. St. Sel.	
			- CAST V. S	^				
				1)				

		1470181	323(633)22		alled 101	100
GO 1	MOINE PAGE	OWEST	El al Fall		Rx 161326	
LFAMPRIDINE (4-AMINOPYR	udine) SMG SL	OW RELEASE	#1 CAPSULES 6 M	G CAPSULE	Lot 09232	
T-11 0.6					Let 11052	NC 22 11
Tall Man: Flavor:		8		Cabadata	4000	
Description:				Schedule:	4902 Formula II	Active ✓ 0:4902
Quantity made: 100 CAPS		Batch yield:		PCCA ID:	100 700 700	20184
Date made: 11/5/2020	Q	ty remaining:		Route of admin	and the second s	
Lot number: 11052020@			Estimated	price \$30.00		
seyond use date: May 4, 2021	s after compound	2:02 PM		200.00	000,	
Pharmacist: NATHAN D.	AHL _				Time to make:	0
NDC1:	-	_				
Packaging: Equipment:						
5.5.5						
Labeling: bility information:						
redients		Sch.	Quantity used	QS (Balan	ce)	
DALFAMPRIDINE USP POWDE			0.5 GMS	□ 0.500 g		
Let #: C195988 Chanical Code:	Mig: Volume:	Potency.	Exp. date: 3/31/202	Whisr.	AWP:	***
Purity;	- street	i small	. 22,000	checked: [V] Each CAU	PS contains 0.005 GMS or 0.	\$0.00
			NDC:		CheminviD: 1	
METHOCEL EAM PREMIUM (HY) Lot 0: 149262/E	Mig MEDISC		10 GMS Exp. date: 7/31/202	2 10.001 g Whisr, ME	DISCA	
Chemical Code:	Volume	Potency:	Q8 amount:	7	AWP:	\$26.90
nox			Bar code NDC:	thecked: [V] Each CAI 51927-1185-00	S contains 0.1 GMS or 10% CheminylD: 13	348
AVICEL PH-105 POWDER	1	L	11.5 GMS	11.501 g		
Lot #: 1805020018 Chantest Code:	Mig: Volume	Potency:	Exp. date: 2/3/2022 QS amount	Whisr: LE	TCO AWP:	\$931
Purity:			Bar code	checked: Esch CAP	S contains 0.115 GMS or 11. CheminviD: 12	.5%
CAPSULE #1-CLEAR LOCKING	CAPSULE		100 CAPS	The same of the sa		75
Lot#:	Mfg:		Exp. date:	Whisr:	1. 7	
Chemical Code: Purity:	Volume	Pciency:	Q6 arrount		AWP:	\$0,00
			NDC:	checked: Each CAP	S contains 1 CAPS or 100% CheminviD:	
g Instructions & Notes						3.00
w Log Originally made as: 100	DALFAMPRIDI	NE (4-AMINOP	YRIDINE) 5MG SLO	W RELEASE #1 C	APSULES 5 MG CA	PSULE
rmula ID: 4902 Calculated lot number: 110520	20@5 Beyond	use date: 5/4/2	021			
PHILI A INSTRICTIONS.				ad partle to o	nausa all antius	
Add food color to Active	(Danampro	line) and mi	x will mortal at	iu pestie to ei	isure an active	
emical is a consistent co	nor and com	ihierally cour	Dillien Mini nie i	oou coloi.		
Add Avicel, Methocel, an	nd Aditya (N	alfamnridina) to a zip lock h	ag and seal		
						3
Jsing rolling bar go back	k and forth o	ver the zip I	ock bag until al	powder is the	e same color to)
ong roung but go such	n of active.	AC. 14740 2.000	7.445 7.41	2 d 2 t 2 t 2 t 2		
sure complete dispersion						
Dump powder from zip k	ock bag onto	capsule m		apsulate usin	g size #1 caps	ules.

1

ogged Formula Worksh 23/2020 3:29:17 PM		1619803		K.			lles 9		
ge 1			4000			R	x 1617	3266	
LFAMPRIDINE (4-AMINOPYRID	NNE) 5MG	SLOW RELEASE	M CAPS	ULES 5 MG C	APSULE			0276200	
							Lot 09	2320206	05 Br
all Man:						4000			
Playor: Description:					Schedula:	4902	Formula	Active ✓ ID: 4902	
Quantity made: 100 CAPS		Batch yield:	100.000)	PCCA ID:		Log	ID: 19808	
Date made: 9/23/2020		Qty remaining:	100.000		Route of admin:	46 - 1-			
Lot number: 09232020@5			E	stimated price	ulations from \$30.00		9		
eyond use date: March 22, 202	ffer compo	3:09 PM unding date		comatoa prio	000.00	40 01			
Pharmacist NATHAN DAI	HL Compa				-	Time to	make:	0	
Technician: NDC1:							771-51		
Packaging: Equipment:									
Labeling:									
ability Information:									
gredients		Sch.	Quan	tity used	QS' (Balan	ce)			
DALFAMPRIDINE USP POWDER			2	0.5 GMS	□ 0.501 g				
Lot 6: C195988 Chemical Code:	Mig: Volume	Petancy:		late: 3/31/2025	Whisr:		AWP:	\$0.00	
Purity:	- semen			Bar coda che	eched: Each CA	PS contains	0.005 GMS	or 0.5%	
HENIOACI EMI BORNIBA (INC	POINT I O	on tions 1		NDC:	□ 10.001 g		CheminviD	: 1366	-
METHOCEL EAM PREMIUM (HYP Lot #: 149282/E	Mig: ME	DISCA		10 GMS ate: 7/31/2022	Whisr: ME	DISCA			
Chemical Code: Purity:	Volumes	Polency:	QS	emount	😝		AWP:	\$20.90	
izer.				NDC: 619	ecked: Each CAI 927-1186-00	S contains	O.1 GMS or 1 CheminvID	: 1348	
AVICEL PH-105 POWDER	1	L	Laws or	11.5 GMS	☐ 11.500 g				
Lot #: 1805020018 Chemical Code:	Mfg: Volume:	Polancy:		ale: 2/3/2022 emount:	Whisr. LE	ico	AWP:	\$9.31	
Purity:	Williams.	6.400.00		Bar code che	cked: 1 Each CAI	S contains	0.115 GMS o		
	ADOLES				91-2007-01		CheminviD:	1275	_
CAPSULE #1-CLEAR LOCKING C	Mfg:		Exp. d	100 CAPS	Whisr.	SHE'S			
	Volume	Polancy:	QB	emount			AWP:	\$0.00	
Chemical Code: Pusitic				Bar code che NDC:	cked: Each CAF		1 CAPS or 10 Cheminvib:		
Chemical Code: Purity: anon:						_			

2 Add Avicel, Methocel, and Acitve (Dalfampridine) to a zip lock bag and seal.

3 Using rolling bar go back and forth over the zip lock bag until all powder is the same color to ensure complete dispersion of active.

4 Di	imp powder from zip	lock bag onto	capsule	machine a	and encapsulate	using size	#1 capsules.
------	---------------------	---------------	---------	-----------	-----------------	------------	--------------

Date entered: 9/23/2 Checked by:	2020 3:09:07 PM	Last modified:	9/23/2020 3:09:07	PM Date:	 2
	1				
					1

Logged Formula Worksheet (2000) 8/20/2020 11:04:27 AM Page 1			122/20/
DALFAMPRIDINE (4-AMINOPYRIDINE) EMG SLOW RE	LEASE # CAPSULES 5 MG		
Tall Man: Flavor: Description: Quantity made: 100 CAPS Batcl	n yleld: 100.000 aining: 100.000	Tiller	# 20 au 2/18/20 his let Active 1/1 Formula ID: 4902 Log ID: 19542
Date made: 8/20/2020 Lot number: 08202020@3 Beyond use date: February 16, 2021 180 days after compounding date Pharmacist: NATHAN DAHL Technician: NDC1: Packaging: Equipment:		lculations from the log	
Laboling:			
Stability information:	h Ountibured	OC /Bolomon	
Ingredients Sc DALFAMPRIDINE USP POWDER Let #: C185980 M/g: Chemical Code: Volume: Potentic Estator:	0.5 GMS Exp. date: 3/31/2025 GS arount	nocked: Each CAPS contains 0.0	XWP: \$0.00 DOS GMS or 0.5% reminvID: 1386
METHOCEL EAM PREMIUM (HYPROMELLOSE USP) L Lot 6: 149262/E MSg: MEDISCA Chanical Code: Volume: Potency Purity:	Bar code ch	ecked Each CAPS contains 0.1	WP: \$26,90 GMS or 10% eminviD: 1346
AVICEL PH-108 POWDER L Lot #: 1805020018 Mfg: Charminal Code: Volume: Potency Purity:		☐ 11.501 g Whisr: LETCO A school: ☑ Each CAPS contains 0.1	WP: \$9,31
	NDC: 62		eminvID: 1275
CAPSULE 61-CLEAR LOCKING CAPSULE L Lot #: Milg: Chemical Code: Volume: Potency Party:	100 CAPS Exp. date: Q5 arrount Ear code div	ocked: Each CAPS contains 1 C	
	RDC:	Cne	eminvi0:
og Instructions & Notes			
Vew Log Originally made as: 100 DALFAMPRIDINE (4-A) formula ID: 4902 Calculated lot number: 08202020@3 Beyond use date ORMULA INSTRUCTIONS: Add food color to Active (Dalfampridine) a nemical is a consistent color and completel	e: 2/16/2021 Ind mix with mortar and	I pestle to ensure al	
Add Avicel, Methocel, and Active (Dalfam)			
Using rolling bar go back and forth over the name complete dispersion of active.	e zip lock bag until all p	powder is the same	color to
Dump powder from zip lock bag onto caps ate entered: 8/20/2020 10:53:58 AM Last modified: 8/20/2 ected by:	Marie II. The Control of the Control	psulate using size # ——A	1 capsules.
		7.	

8/20/2020 10:53:10 AM			F:11	les 8/18/20	
Page 1	TV)4240		6	2× 1613266	
DALFAMPRIDINE (4-AMINOPYRIDINE) SMG SL	OW RELEASE	M CAPSULES 5 MG	CAROUNE	of 08022020	002 #100
Tall Man:			L	of 08022020	003 4 20
Flavor:			Schedule:	4902	Adiva 🗹
Quantity made: 100 CAPS		***	02.000	Formula II	
[] [] : 그램 이번에 가고 있다. 남자 이상에 생겼는데 되는데 기업이 생활하셨다. 생활하는데 보다 다음을 받는데 없는데 되었다.	Batch yield: y remaining:		PCCA ID: Route of admin:	Log II	D: 19540
Date made: 8/20/2020	y temaning.		lculations from	the lon	
Lot number: 08202020@2 Beyond use date: February 16, 2021	10:11 A	Estimated p	rice \$30.00		1
Pharmacist: NATHAN DAHL	ng date	"			- 21
Technician:				Time to make:	0
NDC1: Packaging:		-			
Equipment					
Labeling:					
Stability Information:					
Ingredients	Sch.	Quantity used	QS (Balance	9)	
DALFAMPRIDINE USP POWDER Lot #: C195986 Mfg:		0.5 GMS	0.500 g		1
Chemical Code: Volume:	Potency:	Exp. date: 3/31/2025 03 emount	Whisr:	AWP:	\$0.00
Purity: Estance:		Bar code ci NDC:	nocked: Each CAPS	contains 0 005 GMS or 0.5	
METHOCEL EAM PREMIUM (HYPROMELLOSE US	P) L	10 GMS	☐ 10.001 g	CheminviD: 13	366
Lot #: 149282/E Mfg: MEDISCA Chomical Code: Volume:	Potency:	Exp. date: 7/31/2022	Whisr: MED		20000
Purity:	rumay.		ecked A Each CAPS	AWP; conlains 0.1 GM3 or 10%	\$26,60
		NDC: 51	927-1188-00	CheminviD: 13	46
AVICEL PH-105 POWDER Lot #: 1805020016 Rfg:	L	11.5 GMS Exp. date: 2/3/2022	☐ 11.500 g Whisr: LETC	20	
Chamical Code: Volume: Purity:	Potency:	QS emount	77	AWP:	\$9,31
Edanox		Bar code ch NDC: 62	ecked: 2 Each CAPS 991-2007-01	contains 0.115 GMS or 11.5 CheminviD: 12:	5% 75
CAPSULE #1-CLEAR LOCKING CAPSULE	L	100 CAPS	0_100	2A/25	
Lot #: Mfg: Chemical Code: Volume:	Potency:	Exp. date: QS errount	Whisr.	AWP:	\$0.00
Purity:	0.44	Bar code ch	ecked: Each CAPS	contains 1 CAPS or 100%	00.00
		NDC:		ChaminviD:	-
og Instructions & Notes					
New Log Originally made as: 100 DALFAMPRIDIN	E (A AMINODY	DIDINE SUC SLOW	DELEASE #1 CA	DOLU EC E MO CAT	Delli E
Formula ID: 4902			NELCOOL #1 CA	SUCCES 3 ING CAP	JOLE
Calculated tot number: 08202020@2 Beyond u FORMULA INSTRUCTIONS:					
Add food color to Active (Dalfampridi	ne) and mix	with mortar and	pestle to ens	sure all active	
nemical is a consistent color and comp	pletely comb	ined with the fo	od color.		
			leas been		
Add Avicel, Methocel, and Active (Da	inamphoine) to a zip lock ba	g and seal.		-
Using rolling bar go back and forth or	er the zin la	ock bag until all i	powder is the	same color to	
nsure complete dispersion of active.	rei alo zip i	on bag and an	2011461 10 4115		
nsure complete dispersion of active.					
Dump powder from zip lock bag onto	capsule ma	chine and enca	psulate using	size #1 capsu	ules.
[1] [1] [1] [1] [1] [1] [1] [1] [1] [1]	8/20/2020 10:11				2.0
ecked by:	0/20/20/20 TO.11	Date://	_ \lambda		
			9		
					:

Monica Segedy

From:

nate dahl <natedahl@dahlsrx.com>

Sent:

Friday, June 11, 2021 12:49 PM

To:

Monica Segedy

Subject: Attachments: Dalfampridine 5mg Caps Dalfampridine 5mg caps.pdf

Monica

Here is the RX, Formula log form, and lab certificate of analysis.

Each chemical was barcode scanned into PK Software prior to weighing it which is confirmed on the formula. The weights were then recorded electronically directly from our balance into the PK Software and recorded by the software. I have triple checked each barcode on the chemicals to ensure each one is correct for each chemical involved in this compound and they are all accurate.

If you need anything else please reach out. Thank you.

Nathan Dahl

PharmD, RPH

Dahl's Pharmacy Compounding and Specialty

1007 N Curry Street

Carson City, NV 89703

Phone: 1-844-914-9067

Fax: 775-885-8547

Website: dahlspharmacy.com

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Monica Segedy

From:

Monica Segedy

Sent:

Friday, June 11, 2021 1:15 PM

To:

'nate dahl'

Subject:

RE: Dalfampridine 5mg Caps

Just one more thing:

I see that the log you sent me is for 100 capsules. However, 120 capsules were dispensed. Where did the other 20 capsules come from? I will need the same paperwork for those 20.

Thank You!

Monica S. Segedy Investigator Nevada State Board of Pharmacy msegedy@Pharmacy.nv.gov (775) 850-1440



NOTICE: This information is provided as a courtesy on behalf of the Nevada State Board of Pharmacy. This information does not constitute legal advice and does not establish an attorney-client relationship. This information does not override the specific provisions of Nevada law as applied to a particular set of facts.

From: nate dahl [mailto:natedahl@dahlsrx.com]

Sent: Friday, June 11, 2021 12:57 PM

To: Monica Segedy <msegedy@pharmacy.nv.gov>

Subject: Re: Dalfampridine 5mg Caps

Monica

Yes I will reach out to them right now and get that. I will send it over as soon as I get it. Thank you.

On Fri, Jun 11, 2021 at 12:50 PM Monica Segedy <msegedy@pharmacy.nv.gov> wrote:

Thank you! Could you also send the report from the lab you used to test the drug in Texas.

Monica S. Segedy

Investigator

Nevada State Board of Pharmacy

msegedy@Pharmacy.nv.gov

(775) 850-1440



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From: nate dahl [mailto:natedahl@dahlsrx.com]

Sent: Friday, June 11, 2021 12:49 PM

To: Monica Segedy <msegedy@pharmacy.nv.gov>

Subject: Dalfampridine 5mg Caps

Monica

Here is the RX, Formula log form, and lab certificate of analysis.

Each chemical was barcode scanned into PK Software prior to weighing it which is confirmed on the formula. The weights were then recorded electronically directly from our balance into the PK Software and recorded by







Update 6/5

Little change in the clinic exam today. We will plan on getting MRI brain scans any cortical changes. There is concern patient may have overdosed on 4-aminopyridine. Which is a CNS stimulant. However most case reports the patients usually recover after a few days. Seizure subsides.

Plan:

- 1. D/c EEG
- 2. Discontinue propofol -6/2
- 3. Discontinue Versed 6/3
- 4. Continue Vimpat 200 mg twice daily
- 5. Continue Keppra 1500 twice daily
- 6. Decrease Depakote 500 mg 3 times daily
- 7. Zonisamide 200 mg twice daily

The evaluation of the patient, and recommended management, was discussed with the resident staff. I have performed a physical exam and reviewed and updated ROS and Plan today (6/5/2021). In review of yesterday's note (6/4/2021), there are no changes except as documented above.

This chart was partially generated using voice recognition technology and sound alike word replacement may be present, best efforts were made to make the chart accurate.

Michael Medeiros, MD Board Certified Neurology, ABPN (t) 775-982-2994

Back to the previous page

Monica Segedy

From:

Sent:

Wednesday, June 09, 2021 4:56 PM

To: Subject: Monica Segedy Re: Capsules

Attachments:

image001.png; Screenshot_20210609-165129_Chrome.jpg

Hey Monica,

Attached is a 6/5 update on my mother's Renown Chart from Neurologist Dr. Medeiros stating the concern of a 4-aminopyridine (Dalfampridine) overdose. Please let me know if there is anything else I may provide to help.

Thank you



On Wed, Jun 9, 2021, 3:45 PM Monica Segedy mscgcdy@pharmacy.nv.gov wrote:

HI CISS

Thank you so much for meeting with me today, I know you have a lot going on so I really appreciate it!

Here is a copy of the receipt as well as pictures of the items you turned over to me.

Wishing for the best,

Monica

Monica S. Segedy

Investigator

Nevada State Board of Pharmacy

msegedy@Pharmacy.nv.gov

(775) 850-1440

D Dalfampridine Timeline

May 12th picked up new prescription

13th-4 pills

14th-4 pills

15th-4 pills

16th-2 pills

17th-2 pills

18th-4 pills

19th-4 pills

20th-4 pills

21st-4 pills

22nd-4 pills

23rd-4 pills

- -2 pills soaked in vinegar/ home test
- -10 pills to Dahls
- -18 pills with family
- -50 pills to Nevada Pharmacy Board

Total 120 pills



Monica Segedy

From:

Carlo W <

Sent:

Thursday, June 10, 2021 9:22 AM

To:

Monica Segedy, Yenh Long

Subject:

DeAnna Lopes Timeline

Attachments:

Screenshot_20210610-085943_Samsung Notes.jpg

Good morning Monica and Yehn,

I was able to take some time to really think and work out dates and numbers for all of the 120 pills. I had originally assumed there were 100 pills when Nathan was giving me formulas out of the book. I understood more yesterday when Yehn explained the compound process with different batches and I looked at the bottle quantity. We will still be looking around to see if she had some separately for work. But by going off of my math she took everything accordingly, as well as half dosed for 2 days after her first episode. It's very hard to know precisely since she is unconscious and unable to converse about all of this. We do know that she is always extremely vigilant with taking things correctly. I hope that this can be of some help if needed.



RECEIPT

June 9, 2021

On 6/9/2021 I received from C W one amber vial containing 45 capsules, labeled RX 5070946 dated 5/10/21, 5mg Dalfampridine and prescribed to D Additionally, I received five (5) capsules in a zip lock bag marked "pill box." These five capsules are believed to be 5mg Dalfampridine from Lopes' prescription filled in April, 2021.

Monica S. Segedy Investigator Nevada State Board of Pharmacy

Monica Segedy

From:

C W < MARKET STATE >

Sent:

Monday, June 14, 2021 12:44 PM

To:

Monica Segedy

Subject:

Updates? DeAnna Lopes

Good afternoon Monica,

Just touching base to see about the progress of lab testing. We are at a hard fork right now with my mother being on antiepileptics and trying to wake her up. The results will be extremely beneficial in the approaches being used to treat her condition. As of now she is still in a Vegetative State per basic recovery steps with TBI (Traumatic Brain Injury). At this time she is having an EEG done to monitor if there is any current seizure or epileptic activity before moving on to next steps. Please let me know when you have information.

Thank you

Case

Monica Segedy

From:

C AND WILL OF MARKETS AND MARKETS DE MONTHER DE MONTHER

Sent:

Friday, June 11, 2021 12:57 PM

To: Subject: Monica Segedy Re: Capsules

Attachments:

image001.png; Screenshot_20210611-124959_Chrome.jpg

Unfortunately they said that they had no way to test it by blood. I was requesting they do that from the first day every day for over a week. However, on an EEG of her brain shows that she has Toxic/Metabolic Encephalopathy. By certain readings in the EEG is how they are able to look at certain parts of the brain that are affected by Encephalopathy. Attached is the reading of an EEG performed on my mother from Neurologist Dr. Ania on 6/4. In all studies of overdose of 4-Aminopyridine (Dalfampridine) Toxic Encephalopathy is the result. Also included in this email is a link to a case study that I came across that is extremely similar to what my mother is going through.

https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4009212/

On Fri, Jun 11, 2021, 11:17 AM Monica Segedy <msegedy@pharmacy.nv.gov> wrote:

Thank you Christy. Is there any information you can provide from the Doctor about testing for toxins in her blood?

Monica S. Segedy

Investigator

Nevada State Board of Pharmacy

msegedy@Pharmacy.nv.gov

(775) 850-1440



NOTICE: This information is provided as a courtesy on behalf of the Nevada State Board of Pharmacy. This information does not constitute legal advice and does not establish an attorney-client relationship. This information does not override the specific provisions of Nevada law as applied to a particular set of facts.

INTERPRETATION:

This is an abnormal continuous video electroencephalogram recording in a sedated and or encephalopathic patient. A moderate to severe. toxic / metabolic encephalopathy is suggested. Continuous triphasic waves and frontal sharps (can be more prominent on the left frontal region). Frequent runs of sharply contoured Frontal Intermittent Rhythmic Delta Activity (FIRDA), frequent runs of generalized periodic and rhythmic sharps with triphasic morphology, suggestive of frequent but brief, non convulsive seizures. The findings suggest persistent areas of significantly increased underlying cortical irritability, and/or underlying structural abnormalities. Clinical and radiological correlation is recommended.

Updates provided to Dr. Medeiros.

Rolando Ania, MD
Section Chief Epilepsy and Neurodiagnostics.
Clinical Assistant Professor of Neurology University of Nevada, Reno School of Medicine.
Diplomate in Neurology, Epilepsy, and Electrodiagnostic Medicine.

Office: 775-982-2970 Fax: 775-982-2973

71-Ma toxicalogu apolat Bahlsooss

To: (Name, Title, Address	s, etc.)	Office: N-BOP-RENO
		File #: 21-069-PH-N File Title: Aah! S Pharn Date: 6-15-2021
Type Of Action:		
Property Seized:	Returned To Owner:	Other:
Amount/Quanity Descri	ption Of Items	Purpose
144 CO	psule of Dalfam	pridine in
	ules of Dalfan	
	A	
	1	
	4.00	
ecalved By (Signature)	Witne	essed By (Signature)
(INCOM	. 1//	e And Title (Print)
ame And Title(Print)		Vestigator, NBOP

2 capsules retained from each original container of 45 and 5, for Submission to ALL Bir Prasm Lab for testing.

DI-101-0 Donate Property Rothern

Monica Segedy

From:

Monica Segedy

Sent:

Friday, June 11, 2021 12:51 PM

To:

'nate dahl'

Subject:

RE: Dalfampridine 5mg Caps

Thank you! Could you also send the report from the lab you used to test the drug in Texas.

Monica S. Segedy Investigator Nevada State Board of Pharmacy msegedy@Pharmacy.nv.gov (775) 850-1440



NOTICE: This information is provided as a courtesy on behalf of the Nevada State Board of Pharmacy. This information does not constitute legal advice and does not establish an attorney-client relationship. This information does not override the specific provisions of Nevada law as applied to a particular set of facts.

From: nate dahl [mailto:natedahl@dahlsrx.com]

Sent: Friday, June 11, 2021 12:49 PM

To: Monica Segedy <msegedy@pharmacy.nv.gov>

Subject: Dalfampridine 5mg Caps

Monica

Here is the RX, Formula log form, and lab certificate of analysis.

Each chemical was barcode scanned into PK Software prior to weighing it which is confirmed on the formula. The weights were then recorded electronically directly from our balance into the PK Software and recorded by the software. I have triple checked each barcode on the chemicals to ensure each one is correct for each chemical involved in this compound and they are all accurate.

If you need anything else please reach out. Thank you.

Nathan Dahl

21-11-9 Parison for I oh Re Baysodar

PharmD, RPH

Dahl's Pharmacy Compounding and Specialty

1007 N Curry Street

Carson City, NV 89703

Phone: 1-844-914-9067

Fax: 775-885-8547

1000

Website: dahlspharmacy.com

CONFIDENTIALITY NOTICE. The contents of this email message and any attachments are intended solely for the addressee(s)







840 Research Parkway, Sulte 546 Oklahoma City, OK 73104 Ph: (800) 393-1595 Fax: (405) 271-1174

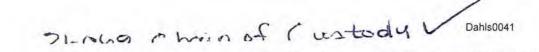
CHAIN-	OF-CUSTODY FORM
LIMS#:	Project ID:
	LAR LISE ONLY

	(CLIENT INF	ORMATION	1	
Company: NBOP			Address: 985 Nan Special Instructions:	nonthe Kar	ich, Res
Contact: Montea S. S	Segedi	1	Potency & Dal	Com asid	1:010
Phone: A series of the series of	计算证据外		boceuch & base	-cirripiria	rac .
775-856-14	140 ext				
Sample Name/Concentration:	S	AMPLE IN	FORMATION		
Dalfam priding	e-4-An	ninopyr	Description: 2 Car	psulea	
Lot/Batch#: See evide	rce Bo	19	Container: Ziplock e	victence l	29
Prepared By: Vanica	Gesed	u	Date of Preparation: Lo/	612026	-
Persons n	elinouishing and	receiving samp	-CUSTODY	nation:	777
Signature, Or	rganization, Date	Mime and Reason	on for Transfer (Start with Box Number	r 1 below)	
Relinquished By	Organization N/A	6/9/21 1:150	2. Monica Seagdy	NBO P	69121 1:15P
Reason for Transfer:	1.	1	1 3.3		
Relinquished By	Organization	Date/Time	Received by	Organization	Date/Time
3. Monica Sagedy	NBOP	6/16/21 10:30A	4.	Organization	Daterrine
Reason for Transfer					
Relinquished By 5.	Organization	Date/Time	Received by	Organization	Date/Time
Reason for Transfer.	1	1	L		
Relinguished By	Organization	Date/Time	Received by	Organization	Date/Time
7.			6.		
Reason for Transfer:					
Relinquished By	Organization	Date/Time	Received by	Organization	Date/Time
9.			10.		
Reason for Transfer.					
Relinquished By	Organization	Date/Time	Received by	Organization	Date/Time
11.			12.		
Reason for Transfer		t .	P. Contraction of the Contractio		

QUF-137-V1 Chain Of Custody (9/28/07)

Page 1 of 1

1/6/2017





Nevada State Board of Pharmacy

985 Damonte Ranch Parkway, Suite 206 • Reno, NV 89521 (775) 850-1440 • FAX (775) 850-1444 E-mail: bkandt@pharmacy.nv.gov • Web Page: bop.nv.gov

June 16, 2021

VIA U.S. MAIL AND ELECTRONIC MAIL TO mdarnaby@arlok.com

Michael Darnaby Client Services Representative ARL Bio Pharma, Inc. 840 Research Parkway, Suite 546 Oklahoma City, OK 73104

RE: Request for records - Case #21-069

Dear Mr. Darnaby:

The Nevada State Board of Pharmacy (Board) hereby requests the following records related to a Board investigation, Case No. 21-069:

Any records and or reports with regard to the testing of Dalfampridine capsules submitted on or about the first week in June 2021 to ARL Bio Pharma, Inc. by Nathan Dahl, RPH (Nevada Certificate of Registration No. 17735), owner/operator of Dahl's Pharmacy located Carson City, NV (Nevada Pharmacy License No. PH03605).

This request is made pursuant to NRS 639.090, NRS 639.236-.239, NRS 639.289, NAC 639.245, NAC 639.482-.489 and/or NAC 639.935(4).

If you have any questions, please do not hesitate to contact me at 775-850-1440 or bkandt@pharmacy.nv.gov.

Best regards,

Brett Kandt General Counsel

Nevada State Board of Pharmacy

21-069 Records Records Fro

Carson Pharmacy LLC d/b/a Dahl's Specialty Pharmacy of Carson 1007 North Curry Street Carson City, Nevada 89701

Date: 7/21/21

NV State Board of Pharmacy Dave Wuest, Executive Secretary 985 Damonte Ranch Pkwy #206 Reno, NV 89521

Dear Mr. Wuest:

This letter will serve as official notification that the above referenced pharmacy with State Board license PH03605 closed its entire pharmacy business on 7/21/21.

All prescription records, controlled & non-controlled substances and required DEA record keeping were transferred to Sierra Specialty Pharmacy and Sierra Compounding Pharmacy 9738 S Virginia St Suite F, Reno, NV 89511.

Sincerely,

Nathan Dahl, Owner



Carson Pharmacy LLC d/b/a Dahl's Specialty Pharmacy of Carson 1007 North Curry Street Carson City, Nevada 89701

Date: 7/21/21

NV State Board of Pharmacy Dave Wuest, Executive Secretary 985 Damonte Ranch Pkwy #206 Reno, NV 89521

Dear Mr. Wuest:

This letter will serve as official notification that the above referenced pharmacy with State Board license PH03605 closed its entire pharmacy business on 7/21/21.

All prescription records, controlled & non-controlled substances and required DEA record keeping were transferred to Sierra Specialty Pharmacy and Sierra Compounding Pharmacy 9738 S Virginia St Suite F, Reno, NV 89511.

Sincerely,

Nathan Dahl, Owner



RXNBR: 5070945 PATIENT: L DOCTOR: LOUIE, TIMOTHY DRUG: DALFAMPRIDINE 5MG SLOW RLS SCANNED: 02-03-2021 P.001/001 (FAX)7757703555 02/03/2021 C9:53 REUHATOLOGY SMMG Neurology Saint Mary's Timothy Louis, tab LIC #: 8282 . DEA #: BL4183202 NPI #: 1407805401 . ALT #: Medical Group 045 N Arlington Ave Sto 655 Reno, NV 895034452 Tel: (775) 778-3554 • Fax: (775) 770-9555 NAME DOB MEETING ADDRESS PEAK DR CARDNERVILLE, NV 69460 DATE 02/01/2021 datampridine (bulk) powder Disp: 120 (One Hundred Twenty) Caseule Ski: take 6 milliaram by oral route 4 times every day compand there as sistalled release per Dr. Verice 2/3/2/ COMMENTS Compounded drug capeules Retal Dispense Only As Written

LOUIS PEAK DAY AGNOWENDILE TWO SEASON TO AS 172700

OAL TARRIPHIE SAND AGNOWENDILE TWO SEASON TO AS 172700

TAKE ONE CAPBULE BY MOUTH FOUR TIMES A DAY

OF LOUIS, TARGITHY DEA: BL—4183202 SUS: NVG202

R45 H ARLHOTON 6955 Reng, NV 89503

Logged Formula	Worksheet (uardan)
5/10/2021 2:33:09 PM	



5/10/2021 2:33:09 PM ###\f	36	Rx 507.09	45
DALFAMPRIDINE (4-AMINOPYRIDINE) 5MG SLOW RELE	EASE #1 CAPSULES 5 MG	CAPSULE	ž.
Tall Man: Flavor: Description: Quantity made: 100 CAPS Batch y Qty remain Date made: 5/10/2021 Lot number: 05/10/2021@8 Beyond use date: November 6, 2021 2::	ning: 100.000	Schedule: 4902 PCCA ID: Route of admin: alculations from the I	Formula ID: 4902 Log ID: 21396
Pharmacist: NATHAN DAHL Technician: - <none> NDC1: Packaging: Equipment:</none>		Time	to make: 0
Labeling: Stability information:			
Ingredients Sch.	Quantity used	QS (Balance)	
DALFAMPRIDINE USP POWDER Lot #; C195988 Mig: Chenical Coda: Volume: Potency: Puthy:	0.5 GMS Exp. date; 3/31/2025 08 amount	0.500 g Whisr:	AWP: \$0.00
Batanos:	Bar code o NDC:	shocked; Each CAPS contains	0.005 GMS or 0.5% CheminviD: 1380
METHOCEL EAM PREMIUM (HYPROMELLOSE USP) L Lot #; 149262/E Mfg: MEDISCA Chamical Code: Volume: Patency:	10 GMS Exp. date: 7/31/2022 OS emount:	10.001 g	AWP: \$26,90
Parity,		hecked: Each CAPS contains	
AVICEL PH-105 POWDER L Let #: 1805020016 Mfg: Chemical Code: Volume: Patency:	11.5 GMS Exp. date: 2/3/2022 QS emount:	11.500 g Whisr LETCO	AWP: \$9.31
Purity:	Bar code d NDC: B	hocked: Each CAPS contains 2691-2007-01	0.115 GMS or 11.5% ChemInvID: 1275
CAPSULE #1-CLEAR LOCKING CAPSULE L Lot #: Mfg: Chemical Code: Volume: Potency:	100 CAPS Exp. date: OS emount:	U 100 Caps	AWP: \$0.00
Balanca:		hacked: Each CAPS contains	
Log Instructions & Notes			
New Log Originally made as: 100 DALFAMPRIDINE (4-AMI Formula ID: 4802 Calculated lot number: 05102021@6 Beyond use date: FORMULA INSTRUCTIONS: I. Add food color to Active (Dalfampridine) an chemical is a consistent color and completely 2. Add Avicel, Methocel, and Acitve (Dalfampri	11/6/2021 d mix with mortar an combined with the fo	d pestle to ensure ood color.	
B. Using rolling bar go back and forth over the ensure complete dispersion of active.			e color to
Dump powder from zip lock bag onto capsu	le machine and enca	apsulate using size	#1 capsules.
Date entered: 5/10/2021 2:27:59 PM Last modified: 5/10/20	021 2:27:59 PM Date; 5, 10)	21	
2)			



PCCA USA SS01 South Wilcrest Drivo Houston, TX 77069 Tel:281.933.6948 Fax: 281.933.6627

PCCA Canada 744 Baransway Drive London, ON N5V 5J2 Tel: 800.668.9453 Fax: 519.455.0890

PCCA Australia Unit 1, 73 Beauchamp Road Matraville, NSW 2036 Tel: 02.9316.1500 Fax: 02.9318.7422

CERTIFICATE OF ANALYSIS

RODUCT: TEM NUMBER: DALFAMPRIDINE USP (4-AMINOPYRIDINE)

OT NUMBER: IFG DATE: EXPIRATION:

50-5030 C195988 04/01/2020 03/31/2025 CAS: MW:

E04-24-5 84.11000000000 FORMULA: C5H6N2

TEST	SPECIFICATIONS	RESULTS		
Assay	98.0-102.0 %	100.6 % Anhydrous basis		
Clarity	% transmittance of 1 cm layer of 5% w/v solution in water at 650 nm should be not less than 95.0%	97.7% T		
Color Indox	Absorbance of 1 cm layer of 5% w/v solution in water at 420 nm should be not more than 0.1AU	0.019 Au		
Dosenption	WHITE POWDER	pass		
Identificiation	paus	Identification per USP		
Melling Point	157-161 c	159.9 c		
Organie impurities	Isonicolinamide: NMT 0.15%; Dalfampridine Related Compound A: NMT 0.10%; Any Individual unspecified impurity: NMT 0.10%; Total Impurities: NMT 0.50%	Isonicotinamide: BQL (<0.0201%); Dalfampridine Related Compound A; BQL (<0.0052%); Any Individual unspecified Impunity: BQL (<0.0048%); Total Impunities; 0.026%;		
Particle Size	pass	D90: 159.57 microns		
pH	10.5-12.5 1% solution	11.33 1% solution		
Related Compound B	<=0.0075 %	0.005 % BQL (<0.005%)		
Related Compound C	<=0.0075 %	0.005 % BQL (<0.005%)		
Residual Solvents	pass	Mathanol: Not detected; Dichloromethano: Not detected;		

Tosted By: Eria Percock Reviewed By: Kierra Briscoe

QC APPROVED: 2020.08.04 PRINT DATE: 2021.03.15 Page 1 of 2

The above test results have been obtained by our supplier or in our quality control isboratory. This analysis is not to be construed as a warranty, expressed or implied.



PCCA USA 9901 South Wilcrest Drive Houston, TX 77099 Tel:281.933.6948 Fax: 231.933.6827

FCCA Canada 744 Baransway Drivo London, ON N5V 5J2 Tel: 800.668,9453 Fax: 519.455.0590

PCCA Australia Unit 1, 73 Beauchamp Read Matravillo, NSW 2006 Tel: 02.9316.1500 Fax: 02.9318.7422

CERTIFICATE OF ANALYSIS

DALFAMPRIDINE USP (4-AMINOPYRIDINE)

PRODUCT:
TEM NUMBER:
OT NUMBER:
MFG DATE:
EXPIRATION: 50-5030 C195986 04/01/2020 03/31/2025

CAS: MW:

504-24-5 94,1100000000

FORMULA: C5H6N2

TEST	SPECIFICATIONS	RESULTS	
Rosidua on Ignition	C=0.3 %	0.05 %	
Solubility	SOLUBLE IN WATER, METHANOL, ACETONE AND ETHANOL	pass	
Nater	<=0.3 %	0.07 %	
	End of Report		

Tosted By: Erin Percock Reviewed By: Kierra Briscoc QC APPROVED: 2020.06.04 PRINT DATE: 2021.03.15 Page 2 of 2

The above test results have been obtained by our supplier or in our quality control laboratory. This analysis is not to be construed as a warranty, expressed or implied.

7.3a Ingredients used for non-sterile compounds - quality and storage

- Each chemical, when received, must be matched to the corresponding invoice to ensure the accurate chemical was sent.
- The expiration date must be verified, and it must be ensured the chemical has adequate dating to meet the needs of the pharmacy.
- Each chemical will be logged into PK Software with its corresponding Name, Lot#, Expiration
 Date, Package Date if available, and will be automatically assigned and labeled with a bar code
 created by the PK Software for scan verification purposes throughout the compounding
 procedure.
- Each chemical will be verified as Hazardous or Non-hazardous and the storage conditions will be determined, the ingredient will then be placed into inventory based on those parameters.
- No chemical shall be placed into inventory without the above conditions being met.

7.4a Compounding Procedure for Non-sterile preparations

- Ensure all PPE is on and adequate for the compound being made.
- Ensure the compounding area, powder-hood, balance, and all items needed for the compound are available, functioning, calibrated, and clean.
- Pull up formula in PK Software and ensure all ingredients are on hand in sufficient quantity to complete the entire formula.
- Pull all necessary ingredients to make formula and place them in the powder-containment her d
 in the same order the appear in the formula.
- Verify all expiration dates are current for all ingredients being used and will not expire before the beyond use date of the compound.
- Begin with the first Ingredient in the formula and bar code scan it to the PK Software to ensure the correct ingredient is being used. If PK Software does not confirm the correct ingredient is being used do not proceed.
- If PK Software, using the bar code scan verification, confirms the correct ingredient proceed to weigh or measure the ingredient using appropriate means to ensure accuracy and consistency. If measuring weight use only the balance contained within the powder hood. If measuring volume use only calibrated graduated cylinders.
- If the measurement is weight use the balance in the powder hood and record the final weight to the formula using PK Software. If the measure is volume use the appropriate graduated cylinder and put the measurement into the formula using PK Software.
- · Once all ingredients have been measured out accurately, save and print the formula,
- Formula logs are to be stored in a file cabinet under the month and year they were completed.
- Proceed combining the ingredients as instructed on the formula using Good Manufacturing Processes per industry standards.

UPARADOMINIST IN ARRISTMEN	COLDINATE PARA			- Rx	201017
LEFAMPRIDINE (4-AMINOP)	(RIDINE) 5MG	SLOW RELEASE	E#1 CAPSULES 5 MG	CAPSULE Filled	4/19/21 480
A. S. C.				filled	5/10/21 #70
Tell Man:					1.01.01 1.20
Flavor: Description:				Schedule: 4902	Active V
Quantity made: 100 CAP	S	Batch yield:	: 100.000	PCCA ID:	Formula ID: 4902 Log ID: 21248
	2	Qty remaining:		Route of admin:	CON 15. E 12.10
Date made: 4/20/2021				lculations from the le	og I
Lot number: 04202021 Beyond use date: November		44.00.4	Estimated p	rice \$30.00 as of	-
180 d	avs after compo	unding date 11:09 A	UVI		
Pharmacist: NATHAN	DAHL			Time	o make: 0
Technician: - <none< td=""><td></td><td></td><td></td><td>11110</td><td>o make.</td></none<>				11110	o make.
Packaging:					
Equipment					
Lobelines					
Labeling:					
redients		Sch.	Quantity used	QS (Balance)	
DALFAMPRIDINE POWDER		3011.	0.5 GMS	- 0.500	
Lot#:	Mfg:		Exp. date:	Whisr.	
Chemical Code: Punity	Volume:	Patency:	QS emount		
noe:			Bar code ct NDC:		
METHOCEL (R) E4M PREMIUM	CR (HYPROM	ELLOSE L	10 GMS	☐ 10.002 g	CheminvID:
Lot #: C177547	Mfg:	GW	Exp. date: 11/5/2020	Whisr:	
Chemical Code: Punty	Volume	Potency:	QS amount		
non.	4		Bar code ch NDC: 51	echod Each CAPS contains 0	1 GMS or 10% CheminviD: 1084
AVICEL PH-105 POWDER		L	11.5 GMS	☐ 11.500 g	TOTAL
Lot#: 1805020018	Mfg:	24.000	Exp. date: 2/3/2022	Whisr. LETCO	
Chemical Code: Furity:	Volume	Potency:	OS emount		AWP: \$9,31
not			Bar code ch NDC: 62	ected: Each CAPS contains (0.115 GMS or 11.5% CheminviD: 1275
CAPSULE #1-CLEAR LOCKING	CAPSULE '	L	100 CAPS	□ (∞)	TOTAL TETO
Lot#:	Mfg:	• • • • • • • • • • • • • • • • • • • •	Exp. date:	Whisr:	
Chemical Code: Punty:	Volume:	Potency:	QS emount	A Lincoln	
nor			Bar code ch NDC:		CAPS or 100% heminviD:
Instructions & Notes					
w Log Originally made as: 10	O DAL FAMOD	DINE (4 AMINOR	VOIDINE! ENO CLOSE	DELEASE #4 CADOLUIS	C E MO OADOU F
mula ID: 4902	UNLPAWIT	IDINE (SANIMOP	INDINE) SWG SLOW	NELEMOE #1 CAPOULE	SO DING CAPSULE .
Calculated lot number: 04202		nd use date: 10/17	7/2021		
FORMULA INSTRUCTIONS:	the mate	mamplalia at an al	malar saible manufacture	and markle to see	a
1. Add food color to Ad					e all active
chemical is a consister	TE color and	completely c	ombined with the	1000 color.	
			and an order		
Add Avicel, Methoce	and Acit	ve (Dalfampric	tine) to a zip lock	bag and seal.	
			at at language and the		TTT4.v2v
3. Using rolling bar go	back and fo	orth over the z	tip lock bag until a	all powder is the sa	me color to
ensure complete dispe	rsion of ac	tive.			
		777			
4. Dump powder from :	zip lock bar	a onto capsule	machine and en	capsulate using si	ze #1 capsules.
" hamen nem			All all all		- III Televinen
a enternet: 4/20/2021 14:00:41 A	M Lest mod	NRad: 4/20/2021 11:	:31:42 AM	4.5	
	M Lest mod	Sified: 4/20/2021 11:	:31:42 AM Data: 4 / 20/	21	
is entered: 4/20/2021 11:09:41 A cited by:	M Lest mod	ilfied: 4/20/2021 11	:31:42 AM Date: 4 / 20/	21	
	M Last mod	iffied: 4/20/2021 11	:31:42 AM Date: <u>4 / 20/</u>	21	

----- Forwarded message ------

From: Colin Brumlow < cbrumlow@eagleanalytical.com >

Date: Mon, Jun 14, 2021 at 9:00 AM

Subject: RE: New Eagle Client Registration: Dahl's Specialty Pharmacy

To: nate dahl < natedahl@dahlsrx.com >

Good morning Nate,

We do not currently have a method developed for Dalfampridine so we are unable to do potency testing or offer Identification on the finished product.

Thank you, Colin

Colin Brumlow, CPhT| Technical Sales Specialist

Eagle | 11111 South Wilcrest Dr., S1000 | Houston, TX 77099 Ph: 800.745.8916 ext.1133 Fax: 713.570.2350

cbrumlow@eagleanalytical.com

http://eagleanalytical.com/

Exhibit 3



From: nate dahl < natedahl@dahlsrx.com > Sent: Friday, June 11, 2021 7:02 PM

Sent. Friday, June 11, 2021 7.02 FM

To: Colin Brumlow < cbrumlow@eagleanalytical.com >

Subject: Re: New Eagle Client Registration: Dahl's Specialty Pharmacy

Is there any way to do any kind of identification on these capsules? I just need to confirm they have dalfampine in them. The concentration of not critical right now.

Nate

On Fri, Jun 11, 2021 at 4:42 PM Colin Brumlow < cbrumlow@eagleanalytical.com wrote:

Good afternoon Nate,

We don't have the capabilities of doing Identification on the compounded capsules because we don't have a method. If the bulk API needed Raw Material Identification we are able to do that per USP we would just need to order the reference standard.

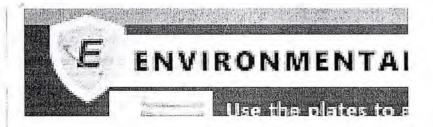
Thank you, Colin

Colin Brumlow, CPhT| Technical Sales Specialist

Eagle | <u>11111 South Wilcrest Dr.</u>, S1000 | Houston, TX 77099 Ph: 800.745.8916 ext.1133 Fax: 713.570.2350

cbrumlow@eagleanalytical.com

http://eagleanalytical.com/



From: nate dahl < natedahl@dahlsrx.com > Sent: Friday, June 11, 2021 4:59 PM

To: Colin Brumlow < cbrumlow@eagleanalytical.com >

Subject: Re: New Eagle Client Registration: Dahl's Specialty Pharmacy

The compounded capsule is what I need verification on.

On Fri, Jun 11, 2021 at 1:31 PM Colin Brumlow <<u>cbrumlow@eagleanalytical.com</u>> wrote:

Good afternoon Nate,

Quick question – I should have clarified with you during the conversation are you needing Raw Material Identification done on the API or are you wanting Identification done on the compounded capsules?

Thank you, Colin

Colin Brumlow, CPhT| Technical Sales Specialist

Eagle | <u>11111 South Wilcrest Dr.</u>, S1000 | Houston, TX 77099 Ph: 800,745.8916 ext.1133 Fax: 713.570.2350

cbrumlow@eagleanalytical.com

http://eagleanalytical.com/



Thank you very much for reaching out to me. I have contacted Eagle Analytical as I need to get a potency test on some dalfampridine 5 mg Sustained Release capsules to verify our processes are yielding correct results. Can you help me with this?

On Wed, Jun 2, 2021 at 1:18 PM Colin Brumlow < cbrumlow@eagleanalytical.com wrote:

Good afternoon Nathan,

An account for EagleTrax has been created for you this afternoon. You should be receiving two emails shortly. The first email will contain your username (NDahl) and the second email will have a link prompting you to confirm your email and create your password. After this is done you'll be able to log into EagleTrax.

If you have any questions once logged into EagleTrax please feel free to contact me via email or at the below phone number. I'm also including an attachment that has information to help you navigate through EagleTrax.

If you plan on doing any USP <71> sterility testing, you will need to get USP <71> Method Suitability performed once per formulation to validate your sterility results. A brief explanation of Method Suitability is the process of inoculating your sample with 6 challenge microorganisms to ensure the ability to see the "bugs" in the product. This assures that you do not have any bacteriostatic properties in your compound that may yield false sterility results. Once that is on file, all of your USP <71> Sterility testing will be fully compliant with USP <797>. To determine the total volume for method suitability, please reference the following equation: (table 2 x table 3) 6 = total volume for Method Suitability. Your USP <71> Method Suitability sample may be in total volume or split up in separate containers.

Once a sample has been submitted via EagleTrax you'll be able to print the sample submission form. The sample and its submission form should be mailed to –

Eagle Analytical Services

11111 S. Wilcrest Dr. #S1000

Houston, TX 77099

Please let me know if you have any further questions or need help determining sample size!

Thank you, Colin

https://eagleanalytical.com/shop/

Please contact me if you have any additional questions.

From: Nathan Dahl < marketing@eagleanalytical.com>

Sent: Wednesday, June 2, 2021 1:48 PM

To: info@eagleanalytical.com; Colin Brumlow <cbrumlow@eagleanalytical.com>

Subject: New Eagle Client Registration: Dahl's Specialty Pharmacy

Name: Nathan Dahl

Company / Pharmacy: Dahl's Specialty Pharmacy

Type of Facility: 503A

Email: <u>natedahl@dahlsrx.com</u>

Address: <u>1007 N Curry Street</u>

City: Carson City

US State: NV Zip: 89703

Country: US

Phone: 1-844-914-9067

PCCA Member # / Customer #: 17091
Desired EagleTrax Username: (updated)

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Nathan Dahl

PharmD, RPH

Dahl's Pharmacy Compounding and Specialty

1007 N Curry Street

Carson City, NV 89703

Phone: 1-844-914-9067

Fax: 775-885-8547

Website: dahlspharmacy.com

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Nate Dahl Dahl's Pharmacy

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Nate Dahl Dahl's Pharmacy

Timesheet for Brett Kandt -

Nathan Dahl, R.Ph., and Dahl's Specialty Pharmacy of Carson, CASE NOS. 21-069-RPH-N/21-069-PH-N

DATE

TIME

6/16/2021

0.50

Confer with staff and draft request for records to ARL Bio Pharma in Oklahoma (ARL). in Case 21-069.

7/27/2021

1.50

Review investigative file and confer with staff; legal research; draft Notice of Intended Action and Accusation in Case 21-069.

7/28/2021

1.75

Review investigative file; legal research; draft Notice of Intended Action and Accusation in Case 21-069.

8/19/2021

2.75

Review investigative file and confer with staff; revise Notice of Intended Action and Accusation in Case 21-069.

8/24/2021

1.25

Review investigative file and consult staff regarding notice of closure of Dahl's Specialty Pharmacy of Carson; finalize and file Notice of Intended Action and Accusation in Case 21-069.

9/9/2021

0.50

Correspond with L. Stone regarding inquiry on Case 21-069.

9/13/2021

0.25

Confer with opposing counsel regarding representation and request for discovery in Case 21-069.

9/17/2021

0.25

Confer with opposing counsel regarding representation and request for extension of time to file Answer and Notice of Defense in Case 21-069.

9/24/2021

1.00

Review discovery production and provide to opposing counsel in Case 21-069.

9/27/2021

1.25

Review Answer and Notice of Defense; confer with staff and review investigative file in Case 21-069.

9/29/2021 0.50

Confer with opposing counsel and provide 6-3-21 voicemail from N. Dahl to D.L.'s daughter.

10/1/2021 0.75

Confer with opposing counsel regarding merits of case and factual allegations and potential resolution in Case 21-069.

10/4/2021 1.50

Prepare for hearing in Case 21-069; confer with staff and witnesses and prepare proposed exhibits.

10/6/2021 1.75

Prepare for hearing in Case 21-069; confer with staff and witnesses and prepare proposed exhibits.

11/8/2021 0.75

Confer with opposing counsel regarding issues of fact and evidence; draft proposed stipulated facts in Case 21-069.

11/24/2021 2.25

Confer with opposing counsel regarding issues of fact and evidence; prepare for hearing in Case 21-069; provide documentation of attorney's fees and costs to opposing counsel.

12/1/2021 1.00

Hearing in Case 21-065-CS-N

TOTAL 20.50 hours x \$65.00/hour = \$1352.50

Timesheet for Monica S. Segedy

Nathan Dahl, RPh, Case No. 21-069-PH-N

DATE	TIME	ACTIVITY
6/09/2021	3.00 Hours	Interviewed Christy Welch at Renown Medical Center Requested records from Dahls Pharmacy
6/11/2021	2.50 Hours	Received and reviewed Pharmacy records Consulted with Yehn Long re: PCCA Analysis of compounded drug. Discussion with Dahl
6/15/2021	5.00 Hours	Requested additional records from Dahl's Consulted with Yehn Long re: PCCA analysis of additional batch in question. Discussions with Dahl Interview of Dr. Tim Louie Met with Christy Welch @ Renown
6/16/2021	6.00 Hours	Discussions with Dahl regarding lab reports Contacted ARL Bio Pharma and Eagle Labs Consulted with General Counsel and request for records from ARL. Shipped pills to ARL, OKC Site visit to Dahl's Pharmacy – Carson City Meeting with Dahl and assessment of Dalfampridine remaining in the Pharmacy
6/21/2021	4.00 Hours	Service of request for records on ARL Bio Pharama Interview of Michael Darnaby – ARL Bio Pharma Received additional records from Dahl – reviewed records for compounded drugs 8/2020 to 5/2021 Review of and Analysis of Dahl formula worksheets
630/2021	1.50 Hours	Interview of Dahl regarding contradicting statements
7/01/2021	2,00 Hours	Review of statement received from Dahl prepared summary of obstruction and unprofessional conduct of Dahl. Review of voicemails provided by Welch
7/12/2021	0.50 Hours	Reviewed ARL Certificates of Analysis
7/14/2021	0.50 Hours	Contact with Christy Welch regarding test results
7/19/2021	1.0 Hours	Contacted Dahl and advised of the lab results Discussion about selling the pharmacy
7/26/2021	1.0 Hours	Discussion with new Pharmacist at Dahls

9/29/2021	2.50 Hours	Review of Accusation and Respondent's Answer Review of Case file
		Discussion with General Counsel
10/4/2021	1.00 Hours	Met with General Counsel – discussed case
		Contacted Witnesses
11/24/2021	1.00 Hours	Met with General Counsel – discussed case
12/1/2021	2.00 Hours	Reviewed case file
		Witness at hearing

TOTAL: 33.50 hours x \$50.00/hour = \$1675.00

Timesheet for Yenh Long -

Nathan Dahl, R.Ph., and Dahl's Specialty Pharmacy of Carson, CASE NOS. 21-069-RPH-N/21-069-PH-N

Date

Time

6/08/2021

1.00

Spoke with complainant regarding complaint submitted.

6/09/2021

1.00

Discussed case and evidence with BOP investigator and Executive Secretary.

6/15/2021

1.00

Reviewed new documents received from respondent with BOP investigator.

6/16/2021

0.50

Discussed with investigator official request for records from ARL Bio Pharma in Oklahoma (ARL).

07/12/2021

0.50

Reviewed ARL lab results with investigator and Executive Secretary

7/28/2021

2.00

Reviewed investigative file and draft of Notice of Intended Action and Accusation.

8/10/2021

2.00

Reviewed investigative file, factual allegations, timeline of events of the case with BOP investigator.

8/12/2021

0.50

Reviewed new draft of factual allegations.

8/24/2021

1.00

Reviewed revised draft of Notice of Intended Action and Accusation in Case 21-069.

TOTAL 9.5 hours x \$65.00/hour = \$617.5

Timesheet for Kris Mangosing -

Nathan Dahl, R.Ph., and Dahl's Specialty Pharmacy of Carson, CASE NOS. 21-069-RPH-N/21-069-PH-N

DATE

TIME

9/24/2021

2

Prepared Discovery request – Compile, redact, bate stamp

9/28/2021

1

Prepared Documents for Board Meeting Material

TOTAL 3 hours x \$31.86/hour = \$95.98

Timesheet for Shirley Hunting –

DAHL'S PHARMACY - NATHAN DAHL - Case No. 21-069-N

Date	Hours	Activity
08/25/2021	1	Prepared Accusations for filing/mailing.

Total Hours

Rate 38.77

Total Costs 38.77

Timesheet for Dave Wuest -

Nathan Dahl, R.Ph., and Dahl's Specialty Pharmacy of Carson, CASE NOS. 21-069-RPH-N/21-069-PH-N

Date

Time

6/09/2021

1.00

Discussed case and evidence with BOP investigator and Deputy Executive Secretary.

6/15/2021

1.00

Reviewed new documents received from respondent with BOP investigator.

6/16/2021

0.50

Discussed with investigator official request for records from ARL Bio Pharma in Oklahoma (ARL).

07/12/2021

0.50

Discussed lab results with ARL staff

7-21-2021

0.50

Reviewed ARL lab results with investigator and Deputy Executive Secretary

7/28/2021

2.00

Reviewed investigative file and draft of Notice of Intended Action and Accusation.

8/10/2021

2.00

Reviewed investigative file, factual allegations, timeline of events of the case with BOP investigator.

8/11/2021

0.50

Reviewed new draft of factual allegations.

8/24/2021

1.00

Reviewed revised draft of Notice of Intended Action and Accusation in Case 21-069.

TOTAL 9 hours x \$65.00/hour = \$585

ARL Bio Pharma, Inc 840 Research Pkwy, Ste 546 Oklahoma City, OK 73104 Phone (405) 271-1144 www.arlok.com

Invoice

Date	Invoice #
6/21/2021	M10062

Bill To

Nevada State Board of Pharmacy Attn David Wuest 431 West Plumb Lane Reno, NV 89509

Project		P.O. No.	Terms	Due Date
			Credit Card	6/21/2021
Quantity	Description		Rate	Amount
1	Box" in a plastic baggie; one labeled as Dalfampridine 5mg. Quantitative analysis will be performed on each capsule 4-aminopyridine (Dalfampridine) be identified		3,500.00	3,500 00

Total

\$3,500 00

Payments/Credits

-\$3.500.00

Balance Due

\$0.00

Payment 2 Payment 2 Balan (NCM MC)

Case: 21-069

CERTIFICATE OF SERVICE

I certify that I am an employee of the Nevada State Board of Pharmacy, and that on this 22nd day of October, 2021, I served a true and correct copy of the foregoing document by Certified U.S. Mail to the following:

URI ASSA KRUTILIN, RPH W PEBBLE RD LAS VEGAS, NV, 89123

OMNICARE OF LAS VEGAS 1525 E SUNSET ROAD SUITE 16 LAS VEGAS, NV 89119

SHIRLEY HUNTING

NEVADA STATE BOARD OF PHARMACY



985 Damonte Ranch Pkwy Suite 206, Reno, Nevada 89521 (775) 850-1440 • 1-800-364-2081 • FAX (775) 850-1444 • Web Page: bop.nv.gov

November 2, 2021

Uri Assa Krutilin W Pebble Rd Las Vegas, NV 89123

Re: Uri Assa Krutilin and Case No. 18-052-RPH-S

Dear Uri Assa Krutilin:

The hearing for case number 18-052-RPH-S has been scheduled for Wednesday, 12/1/2021 9:00:00 AM PST or soon thereafterat the following location:

Home2 Suites Las Vegas Strip South 7740 Las Vegas Blvd. South Las Vegas, NV 89123

Pursuant to NRS 241.033 and 241.034, please be advised that the hearing is a public meeting, and the Board may, without further notice, take administrative action against you if the Board determines that such administrative action is warranted after considering your character, alleged misconduct, professional competence, or physical or mental health. The Board at its discretion may go into closed session to consider your character, alleged misconduct, professional competence, or physical or mental health. You may attend any closed session, have an attorney or other representative of your choosing present during any closed session, and present written evidence, provide testimony, and present witnesses relating to your character, alleged misconduct, professional competence, or physical or mental health during any closed session.

If you have any questions, please feel free to contact the board staff.

Sincerely,

Kristopher Mangosing

Assistant Board Coordinator

Timesheet for Courtney K. Lee, Esq. Nevada State Board of Pharmacy v. Omnicare/Krutilin, RPH – Case No. 18-052

DATE	TIME	DESCRIPTION				
8/23/2021	2.5	Review report of investigation; draft accusation				
9/15/2021	1.0	Finalize accusation, to Reno for final review				
11/9/2021	0.5	Telephone conference with Brian Convery, employee of Omnicare regarding potential settlement				
11/9/2021	1.0	Draft proposed stipulation and order for Omnicare only				
11/30/2021	1.0	Review case file in preparation for hearing (estimated)				
12/1/2021	1.0	Hearing (estimated)				
TOTAL	7	7 hours x \$56 rate per hour = \$392.00				

Total attorney's fees and investigative costs:

\$ 392.00 (Courtney K. Lee's Timesheet)

\$1,357.00 (Dena McClish's, Investigator, Timesheet)

\$1,749.00

Exhibit 2

Timesheet for Investigator Dena McClish Nevada State Board of Pharmacy v. Omnicare/Krutilin – Case No. 18-052

DATE	TIME	DESCRIPTION
6/28/2018	3.25	Visit Omnicare, obtain paperwork
7/30/2018	5.0	Review documents
7/31/2018	5.25	Review employee timecards for Omnicare, review Omnicare records
8/1/2018	3.75	Formulate spreadsheet regarding records
8/2/2018	1.0	Review spreadsheet, emails, records review
8/6/2018	3.25	Organize timecards, review spreadsheet, emails
8/7/2018	2.5	Drafted report of investigation
8/20/2018	1	Review report of investigation
8/27/2018	3.5	Revise report of investigation
9/12/2018	1.0	Final report of investigation, sent to Reno for review
TOTAL	29.5	29.5 hours x \$46 rate per hour = \$1,357.00

NEVADA STATE BOARD OF PHARMACY,

Petitioner,

STIPULATION AND ORDER

٧.

Case No. 97-008A-RPH-S

SIMA MOGHADAM, R.PH, Certificate of Registration No. 11250

Res	po	nd	en	t.
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The Nevada State Board of Pharmacy and Respondent Sima Moghadam, hereby stipulate as follows:

- That Amended Notice of Intended Action and Accusation was sent to Respondent and that Respondent received said Notice.
- 2. That Respondent is fully aware of his right to be served with an Accusation and have a hearing on the matters alleged in the Amended Notice of Intended Action, his right to reconsideration, his right to appeal, and any and all other rights which may be accorded to him pursuant to the Nevada Administrative Procedure Act and the Nevada Pharmacy Act.
- 3. That Respondent hereby freely and voluntarily waives his rights to a hearing, reconsideration, appeal, and any and all other rights that may be accorded to him by the Nevada Administrative Procedure Act and the Nevada Pharmacy Act.
- That Respondent admits the truth of the matters alleged in the Amended
 Notice of Intended Action and Accusation.
- That cause for disciplinary action against the Respondent exists pursuant to the provisions of Nevada Revised Statutes (NRS) 639.210 (4), Nevada
 Administrative Code (NAC) 639.945(1) (d) and (i).

- 6. That the admissions made herein are for the purpose of this proceeding only and shall have no force or effect in any other case or proceeding.
- That based upon the Amended Notice of Intended Action and Accusation and the foregoing admissions, it is stipulated that the following penalty is imposed:

Respondent shall pay to the State of Nevada a fine of \$150.00 personal or certified check, or money order made payable to "State of Nevada, Office of the Treasurer", and an administrative fee of \$100.00 personal or certified check, or money order made payable to "Nevada State Board of Pharmacy". All moneys shall be mailed to and received by the Board office no later than ten days from the receipt of the Board's decision and order in this matter.

Respondent's failure to timely comply with the above-listed conditions shall result in a suspension of Respondent's license for a period of one year, effective immediately upon the date of the failure to comply with this Stipulation.

If this Stipulation is not accepted in its entirety by the Nevada State Board of Pharmacy, it shall have no effect whatsoever.

I have fully considered the charges and allegations contained in the Amended Notice of Intended Action. I understand my right to a hearing, as well as my right to reconsideration, appeal, and any and all other rights accorded me pursuant to the Nevada Administrative Procedure Act and the Nevada Pharmacy Practice Act, including my right to be represented by counsel at my own expense. I hereby freely and voluntarily waive all these rights and agree to the terms of this Stipulation.

DECISION AND ORDER

The Nevada State Board of Pharmacy hereby adopts the foregoing Stipulation as its decision and hereby orders that the foregoing Stipulation be made effective. This decision and order shall be effective on the 6th day of August 1997.

DATED

arry L/Pinson, President

Nevada State Board of Pharmacy

NEVADA STATE BOARD OF PHARMACY,

Petitioner,

NOTICE OF INTENDED ACTION

AND ACCUSATION

V.

SIMA MOGHADAM, R.PH,

Certificate of Registration No. 11250,

Case No. 97-008A-RPH-S

outmoute of Regionation No. 11200

SMITH'S PHARMACY #385, Certificate of Registration No. PH0867; Case No. 97-008B-PH-S

Respondents.

COMES NOW Keith W. Macdonald, in his official capacity as Executive Secretary of the Nevada State Board of Pharmacy, and makes the following that will serve as both a notice of intended action under Nevada Revised Statutes (NRS) 233B.127(3) and as an accusation under NRS 639.241.

1.

The Nevada State Board of Pharmacy has jurisdiction over this matter and these Respondents for the following reasons:

- A. Respondent Sima Moghadam, is a registered pharmacist with the Board (Certificate of Registration Number 11250).
- B. Respondent Smith's Pharmacy (Smith's #385) is a registered pharmacy with the Board (Certificate of Registration Number PH0867), located at 2211 North Rampart Boulevard, Las Vegas, Nevada.

II.

On or about November 19, 1996, Respondent Smith's #385 received a prescription for Tylenol #3 with 30mg Codeine 100 tablets. Respondent Moghadam misfilled the prescription with Tylenol #4 with 60mg Codeine 100 tablets.

Respondent Moghadam labeled a stock bottle of Tylenol #4 with a printed label that indicated Tylenol #3.

IV.

The patient ingested three Tylenol #4 tablets. She took the first tablet and went to bed. The patient woke up several hours later and took a second tablet because she was not feeling well. The patient became very groggy and could not stay awake. She went to the bathroom and fell, injuring herself with broken glass and became bruised. At that time, the patient does not remember clearly, however she took her other medication and must have taken the third Tylenol #4 tablet.

FIRST CAUSE OF ACTION

V

In incorrectly labeling and dispensing a prescription drug, Respondent Moghadam violated Nevada revised Statutes (NRS) 639.210(4) and Nevada Administrative Code

(NAC) 639.945 (1) (d) and (i).

SECOND CAUSE OF ACTION

VI

In owning and operating the pharmacy in which the above acts and violations occurred, Respondent Smith's #385 violated NRS 639.210(4) and 639.945 (1) (i) and (2).

WHEREFORE it is requested that the Nevada State Board of Pharmacy take appropriate disciplinary action with respect to the certificates of registration of the Respondents.

Signed this 25th day of April, 1997.

Keith W. Macdonald, Executive Secretary

Nevada State Board of Pharmacy

NOTICE TO RESPONDENT

You have the right to show the Nevada State Board of Pharmacy that your conduct, as alleged above, complies with all lawful requirements regarding your certificate of registration. To do so, you must mail to the Board within 15 days of your receipt of this Notice of Intended Action and Accusation a written statement showing your compliance.

NEVADA STATE BOARD OF PHARMACY,

Petitioner,

STIPULATION AND ORDER

٧.

Case No. 97-027A-RPH-S

SIMA MOGHADAM, R.PH, Certificate of Registration No. 11250

Res	pond	ent.
-----	------	------

The Nevada State Board of Pharmacy and Respondent Sima Moghadam, hereby stipulate as follows:

- That Amended Notice of Intended Action and Accusation was sent to Respondent and that Respondent received said Notice.
- 2. That Respondent is fully aware of his right to be served with an Accusation and have a hearing on the matters alleged in the Amended Notice of Intended Action, his right to reconsideration, his right to appeal, and any and all other rights which may be accorded to him pursuant to the Nevada Administrative Procedure Act and the Nevada Pharmacy Act.
- 3. That Respondent hereby freely and voluntarily waives his rights to a hearing, reconsideration, appeal, and any and all other rights that may be accorded to him by the Nevada Administrative Procedure Act and the Nevada Pharmacy Act.
- That Respondent admits the truth of the matters alleged in the Amended
 Notice of Intended Action and Accusation.
- That cause for disciplinary action against the Respondent exists pursuant to the provisions of Nevada Revised Statutes (NRS) 639.210 (4), Nevada
 Administrative Code (NAC) 639.945(1) (d) and (i).

- 6. That the admissions made herein are for the purpose of this proceeding only and shall have no force or effect in any other case or proceeding.
- 7. That based upon the Amended Notice of Intended Action and Accusation and the foregoing admissions, it is stipulated that the following penalty is imposed:

Respondent shall pay to the State of Nevada a fine of \$300.00 personal or certified check, or money order made payable to "State of Nevada, Office of the Treasurer", and an administrative fee of \$100.00 personal or certified check, or money order made payable to "Nevada State Board of Pharmacy". All moneys shall be mailed to and received by the Board office no later than ten days from the receipt of the Board's decision and order in this matter.

Respondent's failure to timely comply with the above-listed conditions shall result in a suspension of Respondent's license for a period of one year, effective immediately upon the date of the failure to comply with this Stipulation.

8. If this Stipulation is not accepted in its entirety by the Nevada State Board of Pharmacy, it shall have no effect whatsoever.

I have fully considered the charges and allegations contained in the Amended Notice of Intended Action. I understand my right to a hearing, as well as my right to reconsideration, appeal, and any and all other rights accorded me pursuant to the Nevada Administrative Procedure Act and the Nevada Pharmacy Practice Act, including my right to be represented by counsel at my own expense. I hereby freely and voluntarily waive all these rights and agree to the terms of this Stipulation.

DATED

Sima Moghadam, R.Ph

DECISION AND ORDER

The Nevada State Board of Pharmacy hereby adopts the foregoing Stipulation as its decision and hereby orders that the foregoing Stipulation be made effective. This decision and order shall be effective on the 22nd day of October 1997.

DATED

fry L. Pinson, President

Nevada State Board of Pharmacy

NEVADA STATE BOARD OF PHARMACY,

Petitioner,

NOTICE OF INTENDED ACTION

AND ACCUSATION

V

SIMA MOGHADAM, R.PH, C

Certificate of Registration No. 11250,

Case No. 97-027A-RPH-S

SMITH'S PHARMACY #385, Certificate of Registration No. PH0867;

Case No.

97-027B-PH-S

Respondents.

COMES NOW Keith W. Macdonald, in his official capacity as Executive Secretary of the Nevada State Board of Pharmacy, and makes the following that will serve as both a notice of intended action under Nevada Revised Statutes (NRS) 233B.127(3) and as an accusation under NRS 639.241.

1.

The Nevada State Board of Pharmacy has jurisdiction over this matter and these Respondents for the following reasons:

- A. Respondent Sima Moghadam, is a registered pharmacist with the Board
 (Certificate of Registration Number 11250).
- B. Respondent Smith's Pharmacy (Smith's #385) is a registered pharmacy with the Board (Certificate of Registration Number PH0867), located at 2211 North Rampart Boulevard, Las Vegas, Nevada.

11.

On or about April 16, 1997, Respondent Smith's #385 received a prescription for Altace 5mg. The Mr. C had taken this medication before, still had some at home from his previous prescription, and did not begin taking the new medication until May 8, 1997. Respondent Moghadam misfilled the prescription with Hytrin 5 mg.

Mr. C ingested the first dose of the misfilled Hytrin 5 mg. and called his wife, who had already gone to work, and stated he thought he was going to die. Mr. C's wife took the medication to another pharmacy to find out what it was and was told it was Hytrin 5 mg., not Altace 5 mg. as the prescription container read.

IV.

Mr. C suffered temporary discomfort (headache, weakness, tachycardia, disorientation) and anxiety. When Mrs. C brought the error to the attention of the staff of Smith's #385, whe was told that the error may have occurred "because both tablets are red." Mrs. C was dissatisfied with Smith's #385's response.

FIRST CAUSE OF ACTION

V.

In incorrectly filling, labeling and dispensing a prescription drug, Respondent Moghadam violated Nevada revised Statutes (NRS) 639.210(4) (16) and Nevada Administrative Code (NAC) 639.945 (1) (d) and (i).

SECOND CAUSE OF ACTION

VI.

In owning and operating the pharmacy in which the above acts and violations occurred, Respondent Smith's #385 violated NRS 639.210(4) (16) and NAC 639.945 (1) (d) (i) and (2).

WHEREFORE it is requested that the Nevada State Board of Pharmacy take appropriate disciplinary action with respect to the certificates of registration of the Respondents.

Signed this 22nd day of September, 1997.

Keith W. Macdonald, Executive Secretary

Nevada State Board of Pharmacy

NOTICE TO RESPONDENT

You have the right to show the Nevada State Board of Pharmacy that your conduct, as alleged above, complies with all lawful requirements regarding your certificate of registration. To do so, you must mail to the Board within 15 days of your receipt of this Notice of Intended Action and Accusation a written statement showing your compliance.

NEVADA STATE BOARD OF PHARMACY,

Petitioner,

FINDINGS OF FACT,

CONCLUSIONS OF LAW,

AND ORDER

SIMA MOGHADAM, R.Ph,

٧.

Certificate of Registration #11250,

Case No. 03-049-RPH-S

SUNRISE MEDICAL PHARMACY, Certificate of Registration #PH01698,

Case No. 03-049-PH-S

Respondents.

THIS MATTER was heard by the Nevada State Board of Pharmacy (hereinafter Board) at its regular meeting on April 14, 2004, in Las Vegas, Nevada. The Board was represented by Louis Ling, General Counsel, and Respondent Sima Moghadam appeared and represented herself and Respondent Sunrise Medical Pharmacy. Board staff presented the testimony of Sylena Dean, and Ms. Moghadam testified. Based on testimony, Respondents' admissions, and documents presented as well as the public records in the possession and control of the Board, the Board issues the following Findings of Fact, Conclusions of Law, and Order:

FINDINGS OF FACT

1. Board Staff presented the testimony of Sylena Dean. Respondents made limited admissions regarding the facts of the matter and provided additional testimony by way of explanation and mitigation at hearing. Based upon the admissions, testimony, and evidence taken at hearing, the following are found to be the facts in this matter.

- 2. On May 2, 2003, Ms. Dean took her four-month-old son Gabriel to his pediatrician where he was diagnosed with a digestive problem and was given two prescriptions. One of the prescriptions was for Reglan (metoclopram 5 mg/5 ml syrup) with directions for use of, "1 cc QID AC 30 HS."
- 3. After the appointment with the pediatrician, on May 2, 2003, Ms. Dean took Gabriel's prescriptions to Sunrise Medical Pharmacy to have them filled. At the time, Ms. Moghadam owned and was the managing pharmacist for Sunrise Medical Pharmacy, located at 3006 South Maryland Parkway #101 in Las Vegas.
- 4. Ms. Moghadam filled Gabriel's prescription with the directions for use of,
 "TAKE ONE TEASOPOONFUL BY MOUTH FOUR TIMES DAILY 30 MINUTES
 BEFORE MEALS AND BEDTIME." Thus, the directions for use created and approved
 by Ms. Moghadam were five times as much Reglan as had been ordered.
- 5. Ms. Dean testified that when she picked up the medication, the only counseling she received from Ms. Moghadam was to follow the directions on the label and to have a nice day. Ms. Moghadam presented no convincing evidence that more occurred during the counseling transaction than was remembered by Ms. Dean. When Ms. Dean arrived home, she gave Gabriel a first dose of the Reglan as directed on the incorrect label. Ms. Dean gave Gabriel two more doses thereafter according to the directions on the incorrect label.
- 6. On May 3, 2003, Gabriel began having seizures. Ms. Dean described the seizures as Gabriel's eyes rolling back in his head and his body becoming rigid. Ms. Dean called 911, and Gabriel was taken to the hospital. At the hospital, it was determined that the Reglan prescription label was for teaspoonfuls, not cc's as actually

directed, and that Gabriel had been taking a five times overdose. The physicians who treated Gabriel found that he had a "positive toxic reaction secondary to Reglan."

Gabriel was given Benadryl and Cogentin and the seizures subsided. Gabriel was hospitalized for 24 hours for treatment and observation.

- 7. Ms. Dean related the understandable concerns and fears that a mother would be expected to feel when her infant son began having unexplained and serious seizures. Fortunately, Ms. Dean testified that her understanding was that Gabriel suffered no permanent harm as a result of the Reglan overdose.
- 8. At hearing, Ms. Moghadam explained that she and her husband no longer own Sunrise Medical Pharmacy. She further testified that at the time the error was made regarding Gabriel's Reglan prescription, the pharmacy's computer system automatically defaulted to "teaspoons" for any prescription involving a liquid drug. Ms. Moghadam explained that after learning of the error regarding Gabriel's prescription, she changed the default setting in the pharmacy's computer so that all labels for liquid prescriptions were produced in milliliters rather than teaspoonfuls. Ms. Moghadam could not remember specifically providing counseling to Ms. Dean, and she testified that when the pharmacy was busy and because she was the only pharmacist at the pharmacy, she might not counsel a patient as she knew she should have. Ms. Moghadam acknowledged that the dosage contained on the label she produced on Gabriel's Reglan prescription exceeded the maximum recommended dose for an infant. Ms. Moghadam provided no testimony regarding her previous disciplinary actions. Finally, Ms. Moghadam testified that she is presently not practicing pharmacy and has no intention to practice pharmacy in the future.

CONCLUSIONS OF LAW

- The Nevada State Board of Pharmacy has jurisdiction over this matter because Respondent Moghadam is a pharmacist licensed by the Board and Respondent Sunrise Medical Pharmacy is a pharmacy licensed by the Board.
- In dispensing Gabriel's Reglan prescription with incorrect directions for use,
 Ms. Moghadam violated NRS 639.210(4) and NAC 639.945(1)(d) and (i).
- 3. In failing to counsel Ms. Dean in a way that properly or adequately advised Ms. Dean that the dose of Reglan was excessive, Ms. Moghadam violated NRS 639.210(4) and NAC 639.707(1)(a) and (3)(a) and (e) and 639.945(1)(i).
- 4. In being repeatedly negligent in the filling of prescriptions, as evidenced by Ms. Moghadam's actions in Case Nos. 97-008A-RPH-S, 97-027A-RPH-S, and the present action, Ms. Moghadam violated NRS 639.210(4) and (16).
- The Fourth Cause of Action against Sunrise Medical Pharmacy is dismissed because Ms. Moghadam no longer owns the pharmacy.

ORDER

Based upon the foregoing, the Board hereby orders the following:

- 1. Ms. Moghadam shall pay a total fine of \$5,000.00 (\$3,000.00 for the First Cause of Action, \$2,000.00 for the Second Cause of Action, and no fine for the Third Cause of Action) by cashier's or certified check or money order made payable to "State of Nevada, Office of the Treasurer" to be received by the Board's Reno office within 90 days of the effective date of this Order.
- 2. Within nine months of the effective date of this Order, Ms. Moghadam shall provide to the Board's Reno office evidence of the successful completion of six

continuing education units regarding pediatric dosing of prescription drugs and six continuing education units regarding the prevention of prescription errors. The twelve continuing education units are in excess of and cannot be counted as satisfying the mandatory continuing education units required to be completed for renewal of Ms. Moghadam's license.

- 3. Ms. Moghadam may not commence employment as a pharmacist until she has notified the Board's Reno office in writing of where and when she would like to work as a pharmacist. Upon receiving such notice from Ms. Moghadam that she was returning to the practice of pharmacy, Ms. Moghadam's license shall be placed on probation for a period of three years from the first day on which she returns to the practice of pharmacy. During the period of her probation, Ms. Moghadam will comply with all laws, whether federal or state and whether statute or regulation, regarding the practice of pharmacy.
- 4. The failure by Ms. Moghadam to comply with any term of this Order shall result in the immediate suspension of her registration until all terms applicable to her are fully complied with and will result in further discipline, up to and including revocation of her license. Furthermore, any failure to pay any fine, fee, or cost ordered herein will also result in such legal action as Board staff determines to be necessary to collect the unpaid fine, fee, or cost.

Signed and effective this <u>13</u> day of May, 2004.

arry L. Pinson, President

Nevada State Board of Pharmacy

NEVADA STATE BOARD OF PHARMACY,

Petitioner,

٧.

NOTICE OF INTENDED ACTION AND ACCUSATION

SIMA MOGHADAM, R.Ph. Certificate of Registration No. 11250

Case Number 03-049-RPH-S

SUNRISE MEDICAL PHARMACY Certificate of Registration No: PH01698

Case Number 03-049-PH-S

Respondents.

COMES NOW Keith W. Macdonald, in his official capacity as Executive

Secretary of the Nevada State board of Pharmacy, and makes the following that will

serve as both a notice of intended action under Nevada Revised Statutes (NRS)

233B.127(3) and as an accusation under NRS 639.241.

1.

The Nevada State Board of Pharmacy has jurisdiction over this matter because Respondent Sima Moghadam is a pharmacist licensed by the Board and Respondent Sunrise Medical Pharmacy was a pharmacy licensed by the Board, located at 3006 South Maryland Parkway #101, Las Vegas, Nevada. At all times pertinent to this matter, Ms. Moghadam was the owner and managing pharmacist of Sunrise Medical Pharmacy.

On or about May 2, 2003, Ms. D took her four-month old son, Baby B, to his pediatrician where he was diagnosed with a digestive problem and was given two prescriptions. One of the prescriptions was for Reglan 5 mg./5 ml. syrup with directions for use of, "1 cc QID AC 30 HS."

111.

Ms. Moghadam filled Baby B's Reglan prescription with the directions for use of,
"TAKE ONE TEASPOONFUL BY MOUTH FOUR TIMES DAILY 30 MINUTES BEFORE
MEALS AND BEDTIME." Thus, the directions for use created and approved by Ms.
Moghadam were five times as much Reglan as had been ordered.

IV.

Ms. D indicated that when she picked up the medication, the only counseling she received from Ms. Moghadam was to follow the directions on the label and to have a nice day. Ms. D began her son's medication therapy when they arrived home. Ms. D gave Baby B the medication as directed on the bottle three times.

V.

On May 3, 2003, Patient B began having seizures. Ms. D called 911 and Patient B was taken to the hospital. It was determined at that time that the Reglan prescription label was written with directions to take 1 teaspoonful four times a day, a five times overdose. The physicians who treated Baby B found that Baby B had a "positive toxic reaction secondary to Reglan." Baby B was given Benadryl and Cogentin and the seizures subsided over time. Baby B remained hospitalized for 24 hours for treatment

and observation and is now doing well.

FIRST CAUSE OF ACTION

VI.

In dispensing Baby B's prescription with incorrect directions for use, Ms.

Moghadam violated NRS 639.210(4) and Nevada Administrative Code (NAC) 639.945

(1)(d) and (i).

SECOND CAUSE OF ACTION

VII.

The dosage on the label of Baby B's Reglan plainly and clearly exceeded the maximum dose for Baby B. Ms. Moghadam knew that Baby B was a four-month old infant who would be the patient who would receive the Reglan.

VIII.

In failing to properly or adequately advise Ms. D during counseling that the dose of Reglan was excessive, Ms. Moghadam violated NRS 639.210(4) and NAC 639.707(1)(a) and (3)(a) and (e) and 639.945(1)(i).

THIRD CAUSE OF ACTION

IX.

On June 25, 1997, the Board approved and entered as an order a Stipulation and Order regarding Ms. Moghadam (Case No. 97-008A-RPH-S). The facts underlying the Stipulation and Order were that Ms. Moghadam had placed a Tylenol #3 with 30 mg. Codeine label (the label was correct) upon a stock bottle of Tylenol #4 with 60 mg. of Codeine and dispensed the incorrectly labeled prescription to a patient. The patient

became groggy, fell in her bathroom, and suffered a cut and some bruising. By the Stipulation and Order, Ms. Moghadam paid a fine of \$150.00 and an administrative fee of \$100.00.

X.

On October 22, 1997, the Board approved and entered as an order a Stipulation and Order regarding Ms. Moghadam (Case No. 97-027A-RPH-S). The facts underlying the Stipulation and Order were that Ms. Moghadam had filled a patient's Altace 5 mg. prescription with Hytrin 5 mg. and had dispensed the incorrectly filled prescription to a patient. The patient suffered temporary headache, weakness, tachycardia, disorientation, and anxiety, but suffered no permanent harm. By the Stipulation and Order, Ms. Moghadam paid a fine of \$300.00 and an administrative fee of \$100.00.

XI.

In being repeatedly negligent in the filling of prescriptions, as evidenced by Ms. Moghadan's actions in Case Nos. 97-008A-RPH-S, 97-027A-RPH-S, and the instant action, Ms. Moghadan violated NRS 639.210(4) and (16).

FOURTH CAUSE OF ACTION

XII.

In owning and operating the pharmacy in which Ms. Moghadam committed her violations, from which the incorrect prescription was dispensed, and from which Ms. D was improperly or inadequately counseled, Sunrise Medical Pharmacy violated NRS 639.210(4) and NAC 639.945 (1)(d) and (i) and (2).

WHEREFORE it is requested that the Nevada State Board of Pharmacy take appropriate disciplinary action with respect to the licenses or registrations of the Respondents.

Signed this 3 day of March, 2004.

Keith W. Mackenald Keith W. Macdonald, Executive Secretary Nevada State Board of Pharmacy

NOTICE TO RESPONDENT

You have the right to show the Nevada State Board of Pharmacy that your conduct, as alleged above, complies with all lawful requirements regarding your certificate of registration. To do so, you must mail to the Board within 15 days of your receipt of this Notice of Intended Action and Accusation a written statement showing your compliance.

NEVADA STATE BOARD OF PHARMACY,

Petitioner,

٧.

ANSWER AND
NOTICE OF DEFENSE

SIMA MOGHADAM, R.Ph. Certificate of Registration No. 11250

Case Number 03-049-RPH-S

Respondent.

Respondent above named, in answer to the Notice of Intended Action and Accusation filed in the above-entitled matter before the Nevada State Board of Pharmacy, declares:

- That a hearing on the Notice of Intended Action and Accusation (is) (is not)
 requested (delete inapplicable term).
- 2. That his objection to the Notice of Intended Action and Accusation as being incomplete or failing to state clearly the charges against him, is hereby interposed on the following grounds: (State specific objections or insert "none").

3. That, in answer to the Notice of Intended Action and Accusation, he admits, denies and alleges as follows:

7

I Would Like To Thank The Neadle STate Board of Pharmacy for giving me This right. The right To a hearing before The Newda State Board of Pharmacy. There fore I Can Personally appoing to the family of Patient and The Boar Personally appoing to the family of Patient and The Boar Members. My intention for Coming To This hearing is To Members. My intention for Coming To This hearing is To Members. My intention for Coming To This hearing is To Members. My intention for Coming to This hearing is To Members. To Know that Let the Patient and the Board members to Know that he mistake wasnot made intentionly, and I'm Sorry the mistake wasnot made intentionly, and I'm Sorry for all the incommice I caused for Parents of Cabriel Bassie S.

I hereby declare, under penalty of perjury, that the foregoing Answer and Notice of Defense, and all facts therein stated, are true and correct to the best of my knowledge.

DATED this B day of March, 2004.

Sima Moghadam, R.Ph.

NEVADA STATE BOARD OF PHARMACY,

Petitioner,

٧.

STATEMENT TO THE RESPONDENT NOTICE OF INTENDED ACTION AND ACCUSATION RIGHT TO HEARING

SIMA MOGHADAM, R.Ph.
Certificate of Registration No. 11250

Case Number 03-049-RPH-S

Respond	lent	t.

TO THE RESPONDENT ABOVE-NAMED: PLEASE TAKE NOTICE THAT:

1.

Pursuant to the authority and jurisdiction conferred upon the Nevada State Board of Pharmacy by NRS 639.241 to NRS 639.2576, inclusive, and NRS chapter 233B, a Notice of Intended Action and Accusation has been filed with the board by the Petitioner, Keith W. Macdonald, Executive Secretary for the board, alleging grounds for imposition of disciplinary action by the board against you, as is more fully explained and set forth in the Notice of Intended Action and Accusation served herewith and hereby incorporated reference herein.

11

You have the right to a hearing before the Nevada State Board of Pharmacy to answer the Notice of Intended Action and Accusation and present evidence and argument on all issues involved, either personally or through counsel. It is required that you complete two copies of the Answer and Notice of Defense documents served herewith and file said copies with the Nevada State Board of Pharmacy within fifteen

(15) days of receipt of this Statement and Notice, and of the Notice of Intended Action and Accusation served within.

The Board has reserved Wednesday, April 14, 2004 as the date for a hearing on this matter at the Las Vegas Chamber of Commerce, 3720 Howard Hughes Parkway, Las Vegas, Nevada. The hour of the hearing will be set by letter to follow.

IV.

Failure to complete and file your Notice of Defense with the board within the time allowed shall constitute a waiver of your right to a hearing in this matter and give cause for the entering of your default to the Notice of Intended Action and Accusation filed herein, unless the board, in its sole discretion, elects to grant or hold a hearing nonetheless.

DATED this 3 nd day of March, 2004.

Keith-W. Macdonald, Executive Secretary Nevada State Board of Pharmacy

NEVADA STATE BOARD OF PHARMACY.

Petitioner,

٧.

ANSWER AND NOTICE OF DEFENSE

SIMA MOGHADAM, R.Ph. Certificate of Registration No. 11250

Case Number 03-049-RPH-S

Respondent.

Respondent above named, in answer to the Notice of Intended Action and Accusation filed in the above-entitled matter before the Nevada State Board of Pharmacy, declares:

- That a hearing on the Notice of Intended Action and Accusation (is) (is not)
 requested (delete inapplicable term).
- 2. That his objection to the Notice of Intended Action and Accusation as being incomplete or failing to state clearly the charges against him, is hereby interposed on the following grounds: (State specific objections or insert "none").

3. That,	in answe	r to the N	otice of I	Intended /	Action and	d Accusa	ation, he a	admits, den	ies
	ges as fol								
i i i									
I hereby	declare,	under per	nalty of p	erjury, tha	at the fore	egoing Ar	nswer and	d Notice of	
Defense	, and all f	acts there	ein stated	d, are true	and corr	ect to the	e best of	my knowled	lge.
			DATED	this	day of		,,,	2004.	
			Sima Mo	oghadam,	R.Ph.				
				-2	5				



Financial Statements June 30, 2021

Nevada State Board of Pharmacy



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Independent Auditor's Report

To the Members of the Board Nevada State Board of Pharmacy Reno, Nevada

Report on the Financial Statements

We have audited the accompanying financial statements of the governmental activities and remaining fund information Nevada State Board of Pharmacy (Board) as of and for the year ended June 30, 2021, and the related notes to the financial statements, which collectively comprise the Board's basic financial statements as listed in the table of contents.

Management's Responsibility for the Financial Statements

Management is responsible for the preparation and fair presentation of these financial statements in accordance with accounting principles generally accepted in the United States of America; this includes the design, implementation, and maintenance of internal control relevant to the preparation and fair presentation of financial statements that are free from material misstatement, whether due to fraud or error.

Auditors' Responsibility

Our responsibility is to express opinions on these financial statements based on our audit. We conducted our audit in accordance with auditing standards generally accepted in the United States of America and the standards applicable to financial audits contained in *Government Auditing Standards*, issued by the Comptroller General of the United States. Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free from material misstatement.

An audit involves performing procedures to obtain audit evidence about the amounts and disclosures in the financial statements. The procedures selected depend on the auditor's judgment, including the assessment of the risks of material misstatement of the financial statements, whether due to fraud or error. In making those risk assessments, the auditor considers internal control relevant to the entity's preparation and fair presentation of the financial statements in order to design audit procedures that are appropriate in the circumstances, but not for the purpose of expressing an opinion on the effectiveness of the entity's internal control. Accordingly, we express no such opinion. An audit also includes evaluating the appropriateness of accounting policies used and the reasonableness of significant accounting estimates made by management, as well as evaluating the overall presentation of the financial statements.

We believe the audit evidence we have obtained is sufficient and appropriate to provide a basis for our opinion.

Opinions

In our opinion, the financial statements referred to above present fairly, in all material respects, the financial position of the governmental activities and remaining fund information of the Nevada State Board of Pharmacy, as of June 30, 2021, and the changes in financial position for the year then ended in accordance with accounting principles generally accepted in the United States of America.

Other Matters

Required Supplementary Information

Accounting principles generally accepted in the United States of America require that the management's discussion and analysis on pages 4 through 8, pension information - schedule of changes in net pension liability on page 26, pension information – schedule of contributions on page 27, other postemployment benefit information - schedule of changes in net other post-employment benefits liability on page 28, other post-employment benefit information – schedule of contributions on page 29, and the notes to the required supplementary information on page 30, be presented to supplement the basic financial statements. Such information, although not a part of the basic financial statements, is required by the Governmental Accounting Standards Board, who considers it to be an essential part of financial reporting for placing the basic financial statements in an appropriate operational, economic, or historical context. We have applied certain limited procedures to the required supplementary information in accordance with auditing standards generally accepted in the United States of America, which consisted of inquiries of management about the methods of preparing the information and comparing the information for consistency with management's responses to our inquiries, the basic financial statements, and other knowledge we obtained during our audit of the basic financial statements. We do not express an opinion or provide any assurance on the information because the limited procedures do not provide us with sufficient evidence to express an opinion or provide any assurance.

Other Information

Our audit was conducted for the purpose of forming an opinion on the financial statements that collectively comprise Nevada State Board of Pharmacy financial statements. The accompanying condensed schedules of net position and condensed schedules of activities are presented for purposes of additional analysis and are not a required part of the financial statements. The accompanying Schedule of Expenditures of Federal Awards is presented for purposes of additional analysis as required by the audit requirements of Title 2 U.S. Code of Federal Regulations (CFR) Part 200, Uniform, Administrative Requirements, Cost Principles and Audit Requirements for Federal Awards (Uniform Guidance) and is not a required part of the financial statements.

The condensed schedules of net position and condensed schedules of activities and the schedule of expenditures of federal awards are the responsibility of management and was derived from and relate directly to the underlying accounting and other records used to prepare the basic financial statements. Such information has been subjected to the auditing procedures applied in the audit of the basic financial statements and certain additional procedures, including comparing and reconciling such information directly to the underlying accounting and other records used to prepare the basic financial statements or to the basic financial statements themselves, and other additional procedures in accordance with auditing standards generally accepted in the United States of America. In our opinion, the condensed schedules of net position and condensed schedules of activities and the schedule of expenditures of federal awards are fairly stated, in all material respects, in relation to the basic financial statements as a whole.

Report on Summarized Comparative Information

We have previously audited, in accordance with accounting standards generally accepted in the United States of America, the basic financial statements of Nevada State Board of Pharmacy, as of and for the year ended June 30, 2020 and have issued our report dated November 19, 2020, which we expressed an unmodified opinion on the respective financial statements of the governmental activities and remaining fund information.

The condensed schedules of net position and condensed schedules of activities related to the 2020 financial statements are presented for purposes of additional analysis and were derived from and relate directly to the underlying accounting and other records used to prepare the 2020 financial statements. The information has been subjected to the auditing procedures applied in the audit of the 2020 basic financial statements and certain additional procedures, including comparing and reconciling such information directly to the underlying accounting and other records used to prepare those financial statements or to those financial statements themselves, and other additional procedures in accordance with auditing standards generally accepted in the United States of America. The condensed schedules of net position and condensed schedules of activities are consistent in relation to the basic financial statements from which they have been derived.

Other Reporting Required by Government Auditing Standards

In accordance with Government Auditing Standards, we have also issued our report dated November 2, 2021, on our consideration of the Nevada State Board of Pharmacy's internal control over financial reporting and on our tests of its compliance with certain provisions of laws, regulations, contracts, and grant agreements and other matters. The purpose of that report is solely to describe the scope of our testing of internal control over financial reporting and compliance and the results of that testing, and not to provide an opinion on the effectiveness of Nevada State Board of Pharmacy's internal control over financial reporting or on compliance. That report is an integral part of an audit performed in accordance with Government Auditing Standards in considering Nevada State Board of Pharmacy's internal control over financial reporting and compliance.

Ed Sailly LLP

Reno, Nevada November 2, 2021

The Board members' and management's discussion and analysis of the Nevada State Board of Pharmacy's (Board) financial condition and activities for the fiscal year ended June 30, 2021 is presented in conjunction with the audited financial statements.

Financial Highlights

- Program revenue for the fiscal year ended June 30, 2021 was approximately \$3,900,000 representing a \$1,100,000 increase from the fiscal year ended June 30, 2020.
- Then Nevada State Legislative Commission approved fee increases for three license types at their October 2019 meeting which will have a positive impact on the Board's financials in future years. The fee increases eliminate a structural deficit the Board experienced in the previous year.
- Fees were increased from \$80 to \$200 for controlled substances licenses, pharmacist license fees were increased from \$180 to \$200, and pharmacy tech license fees were increased from \$40 to \$50.
- The Board has enhanced the Prescription Drug Monitoring Program to include related services for healthcare professionals and reporting for State health officials. These additions were fully funded by grant funding and will cease upon loss of grant funds. These expansions do not represent a commitment of Board funds in future years.

Overview of Annual Financial Report

Management's Discussion and Analysis (MD&A) serves as an introduction to, and should be read in conjunction with, the basic audited financial statements and supplementary information. The MD&A represents the Board members' and management's examination and analysis of the Board's financial condition and performance. Summary financial statement data, key financial and operational indicators used in the Board's strategic plan, budget, and other management tools were used for this analysis.

The Board uses the modified accrual basis of accounting for internal financial statement reporting. The financial statements have been prepared in accordance with generally accepted accounting principles as they apply to governmental units. The financial statements include a balance sheet, a statement of revenues, expenditures, and changes in fund balance, and notes to the financial statements.

The Statement of Net Position and Governmental Fund Balance Sheet present the financial position of the Board on both the modified accrual basis under the general fund and the full accrual basis as net position. This statement provides information on the Board's assets, deferred outflows, liabilities and deferred inflows, with the difference reported as net position/fund balance. Over time, increases and decreases in net position/fund balance are one indicator of whether the financial position of the Board is improving or deteriorating.

The Statement of Net Position and Governmental Fund Balance Sheet provide information about the nature and amount of resources and obligations at year-end. The Statement of Activities and Governmental Fund Revenue, Expenditures and Changes in Fund Balance presents the results of the activities over the course of the fiscal year and information as to how the fund balance and net position changed during the year. The fund balance changes under the modified accrual method when revenue is received or the expenditure is made, while changes in net assets under the full accrual method are recorded as soon as the underlying event giving rise to the change occurs, regardless of the timing of the related cash flows. This statement also provides certain information about the Board's recovery of its costs.

The notes to financial statements provide required disclosures and other information that are essential to a full understanding of material data provided in the statements. The notes present information about the Board's accounting policies, significant account balances and activities, material risks, obligations, commitments, contingencies and subsequent events, if any.

The financial statements were prepared by the Board's staff from the detailed books and records of the Board. The financial statements were audited during the independent external audit process.

Financial Analysis

The basic financial statements, as well as the required supplementary information, serve as the key financial data for the Board members' and management's monitoring and planning.

Statement of Net Position

The Board's net position remains strong at year-end with adequate liquid assets to fulfill its responsibilities even though the net position is a deficit at year end. The Board members and management believe the current financial condition and staff capabilities are sufficient to meet anticipated operating expenses and operational objectives. During the year ended June 30, 2015, the Board implemented GASB 68 and 71, Accounting and Financial Reporting for Pensions and Pension Transitions for Contributions Made Subsequent to the Measurement Date., respectively. In March 2016, the GASB issued Statement No. 82, Pension Issues – An Amendment of GASB Statements No. 67, No. 68, and No. 73, effective for periods beginning after June 15, 2016, or June 15, 2017 when an employer's pension liability is measured on a date other than the employer's most recent fiscal year-end. The objective of this statement is to address certain issues that have been raised with respect to GASB Statements No. 67, No. 68, and No. 73 regarding (1) the presentation of payroll-related measures in required supplementary information, (2) the selection of assumptions and the treatment of deviations from the guidance in an Actuarial Standard of Practice for financial reporting purposes, and (3) the classification of payments made by employers to satisfy employee (plan member) contribution requirements. Management has implemented the statement during the year ended June 30, 2018.

During the year ended June 30, 2018, the Board implemented GASB Statement No. 75. Accounting and Financial Reporting for Postemployment Benefits other than Pensions (GASB 75) as required. The purpose of the statement is to improve accounting and financial reporting by state and local governments for postemployment benefits other than pensions (other postemployment benefits or OPEB). It also improves information provided by state and local governmental employers about financial support for OPEB that is provided by other entities. Total OPEB Liability (referred to as the Actuarial Accrued Liability under GASB 45) must be determined using the Entry Age Normal actuarial cost method as opposed to the Projected Unit Credit actuarial cost method used under GASB 45. This change in actuarial cost method resulted in a decrease in the Total OPEB Liability.

The impact of the implementation of these standards to the current year is to include certain deferred inflows and outflows of resources and reflect a net pension liability for the PERS retirement program and a net other post-employment liability as it relates to the Board. The financial impact resulted in the net position of the Board being a deficit of \$2,449,963 and \$2,793,082 at June 30, 2021 and 2020, respectively.

Statement of Activities

Revenue: The program revenue received by the Board is generated through the registration, renewal and licensure of pharmacies and pharmacists. Total revenue received by the Board for fiscal year ended June 30, 2021 was approximately \$5,600,000, representing a \$1,400,000 increase from the fiscal year ended June 30, 2020.

Expenses: Operating expenses for the fiscal year ended June 30, 2021 were approximately \$5,200,000, representing an increase over the fiscal year ended June 30, 2020 of approximately \$300,000. The increase primarily relates to an increase in retirement benefits, and professional contract services with offsetting decreases in salaries.

General Fund Budgetary Highlights

Total revenue received was more than the budgeted amount by approximately \$420,000. The categories of Credit Card Fees and Registrations were the primary variances. This was the first year that online renewals were accepted for all license types which resulted in credit cards being the most used form of payments. Credit card fees are equal to five percent of every transaction run as credit through the licensing system.

Total expenses were less than the budgeted amounts by approximately \$391,000. Savings were primarily due to salary savings related to vacant positions, and travel being restricted due to COVID-19. A planned increase in office space was also delayed due to COVID-19 concerns which resulted in substantial savings in the operating category of approximately \$190,000. All planned travel delayed due to COVID-19 concerns resulted in substantial savings in the travel category of approximately \$139,000.

Economic Factors and Next Year's Budget

The Board is charged with, and given statutory authority, to provide public protection through the licensure and regulation of pharmacists, pharmacies, and other businesses and their employees involved in the manufacture, distribution, and dispensation of drugs. The Board provides direction of staff actions toward its mission of public protection through licensure and disciplinary measures.

To this end, the Board has implemented a variety of changes that include continued software development to automate various job functions which provides cost savings in personnel services. Staff has been directed to continue seeking areas in which operating expenses can be reduced without jeopardizing the high level of customer service the licensees and public have come to know.

Through the Board members' and management's review of the annual budget and monthly income and expense statements, it is expected that these tools will continue to provide the Board with sufficient long and short-term planning information.

With COVID-19 restrictions lifting, it is anticipated that vacancies will be filled as they occur. It is also anticipated that travel will resume to pre-pandemic levels, and that expansion in office space and staff will occur as budgeted.

Following are the condensed statements of net position for the years ended June 30:

	2021 Actual Government- Wide	2020 Actual Government- Wide			
Assets Cash and cash equivalents	\$ 4,310,915	\$ 2,122,747			
Accounts and grants receivable	76,636	29,088			
Prepaid expenses and deposits	16,021	21,524			
Capital assets, net of accumulated depreciation	6,338	19,286			
Total assets	4,409,910	2,192,645			
Deferred Outflows of Resources	1,165,813	1,302,018			
Total assets and deferred outflows of resources	5,575,723	3,494,663			
Liabilities					
Accounts payable and accrued expenses	207,217	171,495			
Wholesaler license deposits	125,000	100,000			
Grants received in advance	33,622				
License fees received in advance	2,274,554	694,430			
Net other post-employment benefit liability	1,443,826	1,344,606			
Net pension liability	3,556,176	3,611,686			
Total liabilities	7,640,395	5,922,217			
Deferred Inflows of Resources	391,762	365,528			
Total liabilities and deferred inflows of resources	8,032,157	6,287,745			
Net Position					
Net position					
Invested in capital assets	6,338	19,286			
Unrestricted	(2,462,772)	(2,812,368)			
Total Net Position	\$ (2,456,434)	\$ (2,793,082)			

Following are the condensed statements of activities for the years ended June 30:

	2021 Actual Government- Wide	2020 Actual Government- Wide	
Expenses	4 1000	4 Village State	
Operations	\$ 1,778,933	\$ 1,735,954	
Personnel	3,411,467	3,073,565	
Travel	36,217	75,592	
Total expenses	5,226,617	4,885,111	
Program Revenue			
Fees, licensing, and permits (charges for services)	3,943,720	2,865,591	
General Revenue			
Grant revenue	1,344,052	1,208,099	
Investment income	16,783	32,412	
Other income	258,710	109,428	
Total general revenue	1,619,545	1,349,939	
Total revenue	5,563,265	4,215,530	
Change in Net Position	\$ 336,648	\$ (669,581)	

Nevada State Board of Pharmacy Statement of Net Position and Governmental Fund Balance Sheet June 30, 2021

		General Fund		djustments (Note 9)	Statement of Net Position	
Assets		A.m.A.				7777
Cash and investments	\$	4,310,915	\$	8	5	4,310,915
Prepaid expenses		16,021		-		16,021
Accounts receivable		76,636				76,636
Deposits						
Capital assets, net of accumulated depreciation	-		_	6,338	-	6,338
Total assets	_	4,403,572	_	6,338	-	4,409,910
Deferred Outflows of Resources						
Net other post-employment benefit liability related		17.		160,920		160,920
Net pension liability related	-		-	1,004,893	-	1,004,893
Total deferred inflows of resources		- 2	_	1,165,813		1,165,813
Total assets and deferred						
outflows of resources		4,403,572) <u>- </u>	1,172,151	_	5,575,723
Liabilities						
Accounts payable		24,992		(*)		24,992
Accrued compensated absences						2,6,7
Due within one year		14		64,000		64,000
Due in more than one year				118,225		118,225
Wholesaler license deposits		125,000		110,225		125,000
Grants received in advance		33,622				33,622
Licensing fees received in advance		2,274,554		0.430400		2,274,554
Net other post-employment benefit liability				1,443,826		1,443,826
Net pension liability	0		_	3,556,176	_	3,556,176
Total liabilities	_	2,458,168	_	5,182,227	_	7,640,395
Deferred Inflows of Resources				.00.260		-022.000
Net other post-employment benefit liability related		-		102,538		102,538
Net pension liability related	_		-	289,224	_	289,224
Total deferred inflows of resources	0		7	391,762	_	391,762
Total liabilities and deferred		Jan Tol				
inflows of resources	-	2,458,168	_	5,573,989		8,032,157
Fund Balance/Net Position Fund balance						
The state of the s						
Nonspendable		******		10.0 00.01		
Prepaid expenses and deposits Unassigned		16,021 1,929,383	-	(16,021) (1,929,383)		3
Total fund balances		1,945,404		(1,945,404)	_	
Total liabilities and fund balance	\$	4,403,572				
Net position						
Invested in capital assets				6,338		6,338
Unrestricted			_	(2,462,772)		(2,462,772
						(2,456,434

Nevada State Board of Pharmacy

Statement of Activities and Governmental Fund Revenue, Expenditures, and Changes in Fund Balance Year Ended June 30, 2021

	General Fund	Adjustments (Note 9)	Statement of Activities	
Expenditures/Expenses Board operations	\$ 4,960,624	\$ 265,993	\$ 5,226,617	
Program Revenue				
Charges for services, licensing revenue	3,943,720		3,943,720	
Net program revenue	(1,016,904)	(265,993)	(1,282,897)	
General Revenue				
Grant revenue	1,344,052		1,344,052	
Investment income	16,783		16,783	
Other income	258,710	-	258,710	
	1,619,545	-	1,619,545	
Excess (Deficiency) of Revenue over				
(under) Expenditures	602,641	(602,641)	12.	
Change in Net Position		336,648	336,648	
Fund Balance/Net Position				
Beginning of year	1,342,763	(4,135,845)	(2,793,082)	
End of Year	\$ 1,945,404	\$ (4,401,838)	\$ (2,456,434)	

Note 1 - Reporting Entity and Summary of Significant Accounting Policies

The Nevada State Board of Pharmacy (the Board) was created in 1901. The Board is regulated by the Nevada Revised Statutes, which also specify the authorized activities of the Board. The Board is the licensing and regulatory agency for pharmacists and pharmacies as well as fifteen other license types in the State of Nevada.

The financial statements of the Board have been prepared in accordance with generally accepted accounting principles as applied to governmental units. The Governmental Accounting Standards Board (GASB) is the accepted standard-setting body for establishing governmental accounting and financial reporting principles.

The following is a summary of the more significant accounting policies.

Reporting Entity

Effective July 1, 2001, Chapter 353 of the Nevada Revised Statutes (NRS) was amended to exempt certain professional and occupational boards from the state budget act and the provisions governing the administration of state funding. The provisions of Chapter 353 do not apply to boards created pursuant to chapters 623 to 625A, inclusive, 628, 630 to 640A inclusive, 641 to 644, inclusive, 654 and 656 of the NRS and the officers and employees thereof. Accordingly, the Board's budgeting and accounting practices and procedures have been removed from the oversight of the Department of Administration.

The Board's financial statements are not included in the financial statements of the State of Nevada since the State does not exercise financial or administrative control over the Board. This is in conformance with GASB codification Section 2100, Defining the Government Reporting Entity.

Basis of Presentation

The Board is defined as a single-program special-purpose entity under GASB Statement No. 14, paragraph 131 as amended by GASB Statement No. 39. This classification allows for the preparation of GASB 34 financial statements under an optional reporting method which combines the fund and government-wide statements into a single presentation. Under standard GASB 34 methodology, the government-wide statement of net position and statement of activities are presented independently from the respective fund balance sheet and statement of revenues, expenditures, and fund balance. A reconciliation of adjustments provided on the modified financial statements demonstrates the changes from the fund financial statements to the government-wide financial statements in order to assist the reader in evaluating these statements. The Board has utilized this optional method of presentation.

Basis of Accounting

The government-wide financial statements are reported using the economic resources measurement focus and the accrual basis of accounting. Revenues are recorded when earned and expenses are recorded when a liability is incurred, regardless of the timing of related cash flows.

Governmental fund financial statements are reported using the current financial resources measurement focus and the modified accrual basis of accounting. Revenue is recognized as soon as it is both measurable and available. "Measurable" means the amount of the transaction can be determined and "available" means collectable within the current period or soon enough thereafter to pay liabilities of the current period. For this purpose, the government considers revenues to be available if they are collected within 60 days of the end of the current fiscal period. Expenditures generally are recorded when a liability is incurred, as under accrual accounting.

Cash and Investments

Cash is maintained in two commercial banks in Reno, Nevada. The Board participates in the State of Nevada collateralization program to assure that funds deposited are protected.

Cash also consists of time certificates of deposit, which are stated at fair value. The net increase (decrease) in the fair value of the investments is the difference between the cost (if purchased during the fiscal year) or the fair value of the investments at the beginning of the year, and the fair value of the investments at the end of the year. Changes in fair value of the certificates are reflected, together with interest income, as investment income in the accompanying financial statements. The Board's certificates are held in its name and it participates in the State of Nevada collateralization program to assure that funds deposited are protected. By statutes, all cash must be deposited in entities that are located in the State of Nevada.

Capital Assets

Capital assets, which include furniture, fixtures, and equipment, are reported in the net position column in the government-wide financial statements. Capital assets are defined by the Board as assets with an initial, individual cost of \$500 and an estimated useful life of at least one year. Such assets are recorded at historical cost. Donated assets are recorded at estimated fair market value at the date of donation. The costs of normal maintenance and repairs that do not add to the value of the asset or materially extend asset lives are expensed as incurred. Capital assets are depreciated using the straight-line method over three to twenty years.

Under the modified accrual basis of accounting, acquisitions are considered expenditures in the year purchased.

Compensated Absences

Compensated absences are accounted for in accordance with GASB Statement 16, Accounting for Compensated Absences, which requires that a liability for compensated absences relating to services already rendered and that are not contingent on a specified event be accrued as an employee earns the rights to the benefits. Compensated absences relating to future services or that are contingent on a specified event will be accounted for in the period those services are rendered, or those events take place. The Board policy permits employees to accumulate earned but unused comp time, vacation and sick benefits subject to certain limitations on hours based on years of service. The sick time paid upon termination is limited to certain payout requirements and has hereby been reflected in the accompanying financial statements based upon these limitations. For the general fund, only the portion of the compensated absences paid from available resources, within 60 days following year-end, are reflected as a liability, if applicable. The full liability is reflected in the government-wide financial statements.

Wholesaler License Deposits

In accordance with statutes, non-publicly traded companies that are wholesalers of prescription drugs must provide a bond, cash deposit or other form of security. There are two companies that provided cash as security under this statute. The cash and liability are reflected in the accompanying financial statements.

Licensing and Licensing Fees Received in Advance

Licensing revenue includes fees for applications, registration and renewal, fines and penalties for late registration and disciplinary fines and charges for administrative duties performed by the Board.

The Board administers its licensing registration on biennial periods from November through October. Licensing fees received in advance represents revenue from the biennial renewals of licenses and the registration of new licenses and is recognized ratably over the license period.

Deferred Outflows and Inflows of Resources

In addition to assets, a separate section is reported for deferred outflows of resources. This separate financial statement element, deferred outflows of resources, represents a consumption of net position that applies to a future period and will not be recognized as an outflow of resources (expense/expenditure) until then. The differences between expected and actual experience, changes in assumptions, changes in proportion, and differences between employer contributions and proportionate share of contributions as well as contributions made after the measurement period for pensions qualify for reporting in this category.

In addition to liabilities, a separate section is reported for deferred inflows of resources. This separate financial statement element, deferred inflows of resources, represents an acquisition of net position that applies to a future period and will not be recognized as an inflow of resources (revenue) until that time. Differences between expected and actual experience and between projected and actual investment earnings on pension plan investments qualify for reporting in this category.

Fund Equity and Net Position

In the governmental fund financial statements, fund balances are classified as follows:

Nonspendable - represents amounts that are either not in a spendable form or are legally or contractually required to remain intact. The Board includes fund balances that have been prepaid for expenses and deposits on hand in this category.

Restricted – represents amounts which can be spent only for specific purposes because of state or federal laws, or externally imposed conditions. The Board has no restricted fund balances.

Committed – represents amounts which can be used only for specific purposes determined by the members of the governing Board's formal action through a resolution or action. The Board has no committed funds.

Assigned - represents amounts that are intended by the Board for specific purposes but do not require action by the governing Board. The Board has no assigned funds.

Unassigned - represents all amounts not included in spendable classifications.

The Board's policy is to first apply expenditures against restricted, committed, assigned fund balances and then unassigned balances. On an annual basis, assigned fund balances are determined based upon available resources.

In the government wide financial statements equity is classified as net position and displayed in the three following components, as applicable:

- Net invested in capital assets consists of capital assets, net of accumulated depreciation and any related debt.
- Restricted net position consists of net position with constraints placed on their use either by (1) external
 groups such as creditors, grantors, contributors, or laws and regulations of other governments; or (2) law
 through constitutional provisions or enabling legislation.
- Unrestricted net position net position that is neither classified as "invested in capital assets" nor as "restricted."

The Board's policy is to first apply expenditures against restricted net position and then unrestricted balances.

Pensions

For purposes of measuring the net pension liability, deferred outflows of resources, deferred inflows of resources and pension expense, information about the fiduciary net position of the Public Employees' Retirement System of Nevada (PERS) and additions to/deductions from PERS's fiduciary net position have been determined on the same basis as they are reported by PERS. For this purpose, benefit payments (including refunds of employee contributions) are recognized when due and payable in accordance with the benefit terms. Investments are reported at fair value.

Postemployment Benefits Other Than Pensions (OPEB)

For purposes of measuring the net OPEB liability, deferred outflows of resources and deferred inflows of resources related to OPEB, and OPEB expense, information about the fiduciary net position of the Self Insurance Trust Fund, Public Employees' Benefits Program (PEBP) and additions to/deductions from PEBP's fiduciary net position have been determined on the same basis as they are reported by PEBP. For this purpose, PEBP recognizes benefit payments when due and payable in accordance with the benefit terms. PEBP's cash and cash equivalents consist of short-term, highly liquid investments that are both (a) readily convertible to known amounts of cash and (b) so near to materiality that they present insignificant risk of changes in value due to charging interest rates.

Use of Estimates

The preparation of financial statements in conformity with generally accepted accounting principles requires management to make estimates and assumptions that affect certain reported amounts and disclosures. Accordingly, actual results could differ from those estimates.

Note 2 - Compliance with Nevada Revised Statutes and Nevada Administrative Code

The Board conformed to all significant statutory constraints on its financial administration during the fiscal year.

Note 3 - Deposits with Financial Institutions

The Board maintains its checking accounts and certificates of deposit in one commercial bank account and one brokerage account. The time certificates of deposit are held in the name of the Board. The accounts are insured by the Federal Deposit Insurance Corporation (FDIC) up to \$250,000 in the aggregate per bank for the checking accounts and \$250,000 for the time deposits.

The bank balances at June 30, 2021 is covered by the FDIC, and the amount not covered by the FDIC is collateralized with securities held by the Nevada Pooled Collateral program. By provisions of statutes, the Board is required to deposit all money in banks or savings and loan associations located in the State of Nevada.

Note 4 - Capital Assets

The Board has custodial responsibility to the State of Nevada for furniture, fixtures and equipment acquired with resources of the Board. The capital asset activity during the year is as follows:

	_Ju	ily 1, 2020		ncreases	Decr	eases	Jui	ne 30, 2021
Capital assets, being depreciated Office furniture and equipment Software Vehicle	\$	755,339 129,060 123,225	\$	1	\$	9.97	\$	755,339 129,060 123,225
	_	1,007,624	_					1,007,624
Less accumulated depreciation Office furniture and equipment Software Vehicle		(727,801) (142,028) (118,509)		(2,945) (6,860) (3,143)		1	_	(730,746) (148,888) (121,652)
	_	(988,338)		(12,948)			-	(1,001,286)
Net capital assets	\$	19,286	\$	(12,948)	\$	-	\$	6,338

Note 5 - Long-Term Obligations

Activity on long-term obligations as provided in the government-wide financial statements is as follows:

Licensing Face received	Ju	ly 1, 2020	Increases	Decreases	Ju	ne 30, 2021	_	Current Portion
Licensing fees received in advance Compensated absences	\$	694,430 135,329	\$ 4,575,930 110,486	\$ (2,995,806) (63,590)	\$	2,274,554 182,225	\$	2,274,554 64,000
	\$	829,759	\$ 4,686,416	\$ (3,059,396)	\$	2,456,779	\$	2,338,554

Note 6 - Operating Lease

The Board currently leases office space in Reno and Las Vegas, Nevada. The leases for Reno and Las Vegas expire on December 31, 2028 and March 31, 2022, respectively. The monthly rental payments range from \$2,845 to \$15,172. The Board also leases a mail machine with monthly payments of \$641 which expires in May of 2025.

The following is a schedule by years of future minimum rental payments:

Years Ending June 30,	
2022	\$ 166,479
2023	162,453
2024	167,098
2025	171,239
2026	169,114
Thereafter	 444,632
	\$ 1,281,015

Rent expense for the year ended June 30, 2021 was \$179,647.

Note 7 - Pensions

General Information About the Pension Plan

Plan Description

PERS (System) administers a cost-sharing, multiple-employer, defined benefit public employees' retirement system which includes both Regular and Police/Fire members. The System was established by the Nevada Legislature in 1947, effective July 1, 1948. The System is administered to provide a reasonable base income to qualified employees who have been employed by a public employer and whose earnings capacities have been removed or substantially impaired by age or disability.

Benefits Provided

Benefits, as required by the Nevada Revised Statutes (NRS or statute), are determined by the number of years of accredited service at time of retirement and the member's highest average compensation in any 36 consecutive months with special provisions for members entering the System on or after January 1, 2010 and for members entering the System on or after July 1, 2015. Benefit payments to which participants or their beneficiaries may be entitled under the plan include pension benefits, disability benefits, and survivor benefits.

Monthly benefit allowances for members are computed as 2.5% of average compensation for each accredited year of service prior to July 1, 2001. For service earned on and after July 1, 2001, this multiplier is 2.67% of average compensation. For members entering the System on or after January 1, 2010, there is a 2.5% service time factor and for regular members entering the System on or after July 1, 2015, there is a 2.25% multiplier. The System offers several alternatives to the unmodified service retirement allowance which, in general, allow the retired employee to accept a reduced service retirement allowance payable monthly during his or her lifetime and various optional monthly payments to a named beneficiary after his or her death.

Post-retirement increases are provided by authority of NRS 286.575 – 286.579.

Vesting

Regular members entering the System prior to January 1, 2010 are eligible for retirement at age 65 with five years of service, at age 60 with 10 years of service, or at any age with thirty years of service. Regular members entering the System on or after January 1, 2010, are eligible for retirement at age 65 with five years of service, or age 62 with 10 years of service, or any age with thirty years of service. Regular members who entered the System on or after July 1, 2015 are eligible for retirement at age 65 with 5 years of service, or at age 62 with 20 years of service or at age 55 with 30 years of service or at any age with 33 1/3 years of service.

The normal ceiling limitation on monthly benefits allowances is 75% of average compensation. However, a member who has an effective date of membership before July 1, 1985, is entitled to a benefit of up to 90% of average compensation. Both Regular and Police/Fire members become fully vested as to benefits upon completion of five years of service.

Contributions

The authority for establishing and amending the obligation to make contributions and member contribution rates is set by statute. New hires, in agencies which did not elect the Employer-Pay Contribution (EPC) plan prior to July 1, 1983 have the option of selecting one of two contribution plans. Contributions are shared equally by employer and employee. Employees can take a reduced salary and have contributions made by the employer (EPC) or can make contributions by a payroll deduction matched by the employer.

The System's basic funding policy provides for periodic contributions at a level pattern of cost as a percentage of salary throughout an employee's working lifetime in order to accumulate sufficient assets to pay benefits when due.

The System receives an actuarial valuation on an annual basis indicating the contribution rates required to fund the System on an actuarial reserve basis. Contributions actually made are in accordance with the required rates established by the Nevada Legislature. These statutory rates are increased/decreased pursuant to NRS 286.421 and 286.450.

The actuary funding method used is the Entry Age Actuarial Cost Method. It is intended to meet the funding objective and result in a relatively level long-term contributions requirement as a percentage of salary.

For the fiscal year ended June 30, 2020, the Statutory Employer/employee matching rate was 15.25% for Regular employees. The Employer-pay contribution (EPC) rate was 29.25%, for June 30, 2020 for Regular employees.

Pension Liabilities, Pension Expense, and Deferred Outflows of Resources and Deferred Inflows of Resources Related to Pensions

At June 30, 2021, the Board reported a liability of \$3,556,176 for its proportionate share of the net pension liability. The net pension liability was measured as of June 30, 2020, and the total pension liability used to calculate the net pension liability was determined by an actuarial valuation as of that date. The Board's proportion of the net pension liability was based on total contributions due on wages paid during the measurement period. Each employer's proportion of the net pension liability is based on their combined employer contributions relative to the total combined employer contributions for all employers for the period ended June 30, 2020. At June 30, 2020, the Board's proportion was .02553% percent, which was a decrease of .00096% from its proportion measured at June 30, 2019.

For the years ended June 30, 2021, the Board recognized pension expense of \$479,895. Amounts totaling \$304,004 resulting from Board contributions subsequent to the measurement date will be recognized as a reduction of the net pension liability in year ended June 30, 2021. For the year ended June 30, 2021, the Board contributed \$304,004 under the statutes requirements based on covered payroll of \$2,104,627 which equates to 14.44% overall to the plan. At June 30, 2021, the Board reported deferred outflows of resources and deferred inflows of resources related to pension from the following sources:

	0	Deferred utflows of esources	li	Deferred oflows of esources
Differences between expected and actual experience	\$	110,488	\$	45,919
Changes of assumptions		99,889		-
Net difference between projected and actual				
investment earnings on pension plan investments				134,337
Changes in proportion		490,512		108,968
Contributions subsequent to the measurement date	4	304,004		
	\$	1,004,893	\$	289,224

Amounts reported as deferred outflows of resources and deferred inflows of resources, without regard to the contributions subsequent to the measurement date, related to pensions will be recognized in pension expense as follows:

Years Ending June 30,		
2022	\$	110,291
2023 2024		168,038 116,199
2025 2026		34,253
Thereafter	_	(14,581) (2,535)
	\$	411,665

The net difference between projected and actual investment earnings on pension plan investments will be recognized over five years, all the other above deferred outflow and deferred inflows will be recognized over the average expected remaining services lives, which was 6.13 years for the measurement period ending June 30, 2020.

Reconciliation of the net pension liability at June 30, 2021 is as follows:

Beginning net pension liability Pension expense	\$ 3,611,686 479.895
Employer contributions	(266,075)
Current year net deferred (inflows) and outflows	(269,330)
Ending net pension liability	\$ 3,556,176

Actuarial Assumptions

The System's net pension liability was measured as of June 30, 2020, and the total pension liability used to calculate the net pension liability was determined by an actuarial valuation as of that date. The total pension liability was determined using the following actuarial assumptions, applied to all periods included in the measurement:

Inflation rate 2.75%

Payroll growth 5.00%, including inflation

Investment rate of return 7.50% Productivity pay increase 0.50%

Projected salary increases Regular: 4.25% to 9.15%, depending on service. Rates include

inflation and productivity increases.

Consumer price index 2.75%

Other assumptions Same as those used in the June 30, 2020 funding actuarial valuation

Actuarial assumptions used in the June 30, 2019 valuation were based on the results of the experience review completed in 2018.

The discount rate used to measure the total pension liability was 7.50% as of June 30, 2020. The projection of cash flows used to determine the discount rate assumed that employee and employer contributions will be made at the rate specified in statute. Based on that assumption, the pension plan's fiduciary net position at June 30, 2020, was projected to be available to make all projected future benefit payments of current active and inactive employees. Therefore, the long-term expected rate of return on pension plan investments was applied to all periods of projected benefit payments to determine the total pension liability as of June 30, 2020.

Investment Policy

The System's policies which determine the investment portfolio target asset allocation are established by the System. The asset allocation is reviewed annually and is designed to meet the future risk and return needs of the System. The following was the System's adopted policy target asset allocation as of June 30, 2020:

Asset Class	Target Allocation	Long-Term Geometric Expected Real Rate of Return *		
Domestic Equity	42%	5.50%		
International Equity	18%	5.50%		
Domestic Fixed Income	28%	0.75%		
Private markets	12%	6.65%		

^{*}As of June 30, 2020, PERS' long-term inflation assumption was 2.75%.

Discount Rate and Pension Liability Discount Rate Sensitivity

The following presents the net pension liability of the PERS as of June 30, 2020, calculated using the discount rate of 7.50%, as well as what the PERS net pension liability would be if it were calculated using a discount rate that is 1 percentage-point lower (6.50%) or 1 percentage-point higher (8.50%) than the current discount rate:

	1% Decrease			
	Discount Rate (6.50%)	Discount Rate (7.50%)	Discount Rate (8.50%)	
Net pension liability	\$ 5,545,837	\$ 3,556,176	\$ 1,901,423	

Pension Plan Fiduciary Net Position

Additional information supporting the Schedule of Employer Allocations and the Schedule of Pension Amounts by Employer is located in the PERS Comprehensive Annual Financial Report (CAFR) available on the PERS website at www.nvpers.org under Quick Links – Publications.

Note 8 - Other Post Employment Retirement Benefits (OPEB)

General Information About the OPEB Plan

Plan Description

Employees of the Board are provided with OPEB through the Self Insurance Trust Fund, Public Employees' Benefits Program (PEBP) – a cost-sharing multiple employer defined benefit OPEB plan administered by the Public Employees' Benefits Program Board (PEBP Board) which was created in 1983 by the Nevada Legislature to administer group health, life and disability insurance for covered employees, both active and retired, of the State, and certain other participating public employers within the State of Nevada. PEBP does not provide for refunds of employee contributions. The Self Insurance Trust Fund issues a publicly available financial report that can be obtained at https://pebp.state.nv.us/. The Board is reporting plan information consistently with the PEBP's accounting methods and assumptions as disclosed in the annual report. No information has come to our attention that indicates significant changes to the plan's disclosures.

Benefits Provided

Benefits other than pensions are provided to eligible retirees and their dependents through the payment of subsidies from the State Retirees' Health & Welfare Benefits Fund. The "base" subsidy rates are set by PEBP and approved by the Legislature and vary depending on the number of dependents and the medical plan selected. These subsidy rates are subtracted from the premium to arrive at the "participant premium". The "years of service" subsidy rates are then used to adjust the "participant premium" based on years of service. The current subsidy rates can be found on the PEBP website at www.pebp.state.nv.us. Benefits include health, prescription drug, dental and life insurance coverage. As required by statute, benefits are determined by the number of years

of service at the time of retirement and the individual's initial date of hire. Officers and employees hired after December 31, 2011 are not eligible to receive subsidies to reduce premiums. The following individuals and their dependents are eligible to receive subsidies from the Retirees' Fund:

Any PEBP covered retiree with State service whose last employer was the State or a participating local government entity and who:

- Was initially hired by the State prior to January 1, 2010 and has at least five years of public service; or
- Was initially hired by the State on or after January 1, 2010, but before January 1, 2012 and has at least fifteen years of public service; or
- Was initially hired by the State on or after January 1, 2010, but before January 1, 2012 and has at least five
 years of public service and has a disability; or
- Any PEBP covered retiree with State service whose last employer was not the State or a participating local government entity and who has been continuously covered under PEBP as a retiree since November 30, 2008.

State service is defined as employment with any Nevada State agency, the Nevada System of Higher Education and any State Board or Commission. Participating local government entity is defined as a county, school district, municipal corporation, political subdivision, public corporation or other local governmental agency that has an agreement in effect with PEBP to obtain group insurance.

Contributions

Per NRS 287 contribution requirements of the participating entities and covered employees are established and may be amended by the PEBP Board. The Boards' contractually required contribution for the year ended June 30, 2020 was \$42,134, actuarially determined as an amount that is expected to finance the costs of benefits earned by employees during the year. Employees are not required to contribute to the OPEB plan.

OPEB Liabilities, OPEB Expense, and Deferred Outflows of Resources and Deferred Inflows of Resources Related to OPEB

At June 30, 2021, the Board reported a liability of \$1,443,826 for its proportionate share of the net OPEB liability. The net OPEB liability was measured as of June 30, 2020, and the total OPEB liability used to calculate the net OPEB liability was determined by an actuarial valuation as of that date. The Board's proportion of the net OPEB liability was based on a projection of the Board's long-term share of contributions to the OPEB plan relative to the projected contributions of all participating state agencies, actuarially determined. At June 30, 2020, the Board's proportion was 0.0960%, which was a decrease of .0005% from its proportion measured at June 30, 2019.

For the year ended June 30, 2021, the Board recognized OPEB expense of \$82,360. At June 30, 2021, the Board reported deferred outflows of resources and deferred inflows of resources related to OPEB from the following sources:

	Outflows of Resources	Inflows of Resources
Differences between expected and actual experience Changes of assumptions Asset experience Fund contributions subsequent to the measurement date	\$ - 115,596 - 45,324	\$ 73,030 29,205 303
	\$ 160,920	\$ 102,538

Amounts reported as deferred outflows of resources and deferred inflows of resources related to OPEB will be recognized in OPEB expense as follows:

Years Ending June 30,	
2021	\$ (11,965)
2022	5,717
2023	11,226
2024	 8,080
	\$ 13,058

Actuarial Assumptions

The total OPEB liability in the June 30, 2020 actuarial valuation was determined using the following actuarial assumptions, applied to all periods included in the measurement, unless otherwise specified:

Inflation 2.50 percent

Salary increases 2.75 percent, average

Investment rate of return 3.51 percent

Healthcare cost trend rates 6.25 percent for 2020, see report for additional years

Mortality rates were based on Pub-2010 Public Retirement Plans General Mortality Table weighted by Headcount, projected by MP-2019.

The actuarial assumptions used in the June 30, 2020 valuation were based on the results of an actuarial valuation date of January 1, 2019, adjusted by using roll-forward procedures to determine the liability at the measurement date.

Discount Rate

The discount rate basis under GASB 75 is required to be consistent with a 20-Year Municipal Bond Index. The Bond Buyer General Obligation 20-Bond Municipal Bond Index is used for the determination of the discount rate.

The discount rates as of July 1, 2020 is 2.21%. Additional detail regarding the discount rates as of June 30, 2020, is provided in the "Actuarial Assumptions and Methods" section of the report proved by the PEBP Board.

Sensitivity of the Board's Proportionate Share of the Net OPEB Liability to Changes in the Discount Rate

The following presents the Board's proportionate share of the net OPEB liability, as well as what the Board's proportionate share of the net OPEB liability would be if it were calculated using a discount rate that is 1-percentage-point lower or 1-percentage-point higher than the current discount rate:

	1% Decrease 1,21%	Discount Rate 2.21%	1% Increase 3.21%
Net OPEB liability	\$ 1,615,714	\$ 1,443,826	\$ 1,298,593

Sensitivity of the Board's proportionate share of the net OPEB liability to changes in the healthcare cost trend rates.

The following presents the Board's proportionate share of the net OPEB liability, as well as what the Board's proportionate share of the net OPEB liability would be if it were calculated using healthcare cost trend rates that are 1-percentage-point lower or 1-percentage-point higher than the current healthcare cost trend rates:

	Heal	Ith Care Cost Trend F	Rates
	1% Decrease (5.25% Decreasing to 3.5%)	(6.25% Decreasing to 4.5%)	1% Increase (7.25% Decreasing to 5.5%)
Net OPEB liability	\$ 1,350,458	\$ 1,443,826	\$ 1,555,602

OPEB Plan Fiduciary Net Position

Detailed information about the OPEB plan's fiduciary net position is available in the separately issued PEBP financial report.

Note 9 - Conversion to Government-Wide Financial Statements

Adjustments on the face of the financial statements were made to the fund balance sheet and statement of revenue, expenditures, and changes in fund balance in order to reconcile the fund financial statements to the government-wide statements of net position and activities. These adjustments detail the effect of the capitalization of fixed assets of \$1,007,624, accumulated depreciation of \$1,001,286, depreciation expense of \$12,948, long-term accrued compensated absences of \$182,225, net deferred inflows and outflows of \$774,051 the net pension liability of \$3,556,176, and the net OPEB liability of \$1,443,826.

Nevada State Board of Pharmacy
Pension Information - Schedule of Changes in Net Pension Liability
Last Ten Fiscal Years

	2020	2019	2018	2017	2016	2015 2014	2014
Proportion of the net pension liability	0.02553%	0.02649%	0.02559%	0.02264%	0.01972%	0.01766%	0.01720%
Proportionate share of the net liability	\$3,556,176	\$3,611,686	\$3,490,261	\$3,010,553	\$2,654,412	\$2,024,299	\$1,793,062
Covered payroll	\$2,104,627	\$1,835,896	\$1,711,106	\$1,457,180	\$1,117,746	\$1,053,952	\$1,002,366
Proportionate share of the net pension liability as a percentage of covered payroll	168.97%	196.73%	203.98%	206.60%	237.48%	192.07%	178.88%
Plan fiduciary net position as a percentage of the total pension liability	77.04%	76.46%	75.24%	74.40%	72.20%	75.10%	76.30%

Note: Only seven years of information is available due to reporting changes with GASB 68 for Fiscal Year 2015.

Nevada State Board of Pharmacy Pension Information - Schedule of Contributions Last Ten Fiscal Years

	2021	2020	2019	2018	2017	2016	2015
Contractually required contributions	\$ 304,004	\$ 266,075	\$ 266,075 \$ 254,976 \$ 237,423 \$ 184,648 \$ 153,565 \$ 125,087	\$ 237,423	\$ 184,648	\$ 153,565	\$ 125,087
Contributions in relation to contractually required contributions	(304,004)	(266,075)	(254,976)	(237,423) (184,648)	(184,648)	(153,565)	(125,087)
Contribution deficiency (excess)	\$	*	\$	\$	\$	\$	\$
Covered payroll	\$2,104,627	\$1,845,447	\$1,835,986 \$1,711,106		\$1,457,180	\$1,117,745	\$1,053,952
Contributions as a percentage of covered payroll	14.44%	14.42%	13.89%	13.88%	12.67%	13.74%	11.87%

Note: Only seven years of information is available due to reporting changes with GASB 68 for Fiscal Year 2015.

Nevada State Board of Pharmacy

Other Post-Employment Benefit Information - Schedule of Changes in Net Other Post-Employment Benefits

Liability

Last Ten Fiscal Years

	2020	2019	2018	2017
Board's Proportion of the Net OPEB Liability	0.0960%	0.0965%	0.1000%	0.0846%
Board's Proportionate Share of the Net OPEB Liability	\$ 1,443,826	\$ 1,344,606	\$ 1,325,428	\$ 1,101,166
Board's Covered-Employee Payroll	\$ 2,104,627	\$ 1,845,447	\$ 1,711,106	\$ 1,407,868
Board's Proportionate Share of the Net OEPB Liability as a Percentage of its Covered-	C9 C00/	72.970/	77.460/	70 200/
Employee Payroll Plan Fiduciary Net Position as a	68.60%	72,87%	77.46%	78.30%
Percentage of the Total OPEB Liability	0.00%	0.00%	0.00%	0.00%

Note: Only four years of information is available due to reporting changes with GASB 75 for Fiscal Year 2017.

Nevada State Board of Pharmacy Other Post-Employment Benefit Information - Schedule of Contributions Last Ten Fiscal Years

		2021		2020		2019		2018
Contractually Required Contribution	ė	42,134	è	39,504	\$	39,654	\$	32,195
Contribution	À.	42,134	\$	35,304	Ą.	39,034	2	32,133
Contributions in Relation to the Contractually Required								
Contribution	_	(42,134)	_	(39,504)	-	(39,654)		(32,195)
Contribution Deficiency (Excess)	\$		\$	= ===	\$		\$	
Board's Covered-Employee Payroll	\$	2,104,627	\$	1,845,447	\$	1,711,106	\$	1,407,868
Contributions as A Percentage of Covered-Employee Payroll		2.00%	1	2.14%		2.32%		2.29%

Note: Only four years of information is available due to reporting changes with GASB 75 for Fiscal Year 2017.

Note 1 - Other Post-Employment Benefit (OPEB)

Changes of Benefit Terms

None.

Changes of Assumptions

The assumed discount rate used by the actuary to determine the post-employment benefits liability at June 30, 2021 was decreased to 2.21% from 3.51% at June 30, 2020. The effect of the change would result in an increase in the liability.



Supplementary Information June 30, 2021

Nevada State Board of Pharmacy



Nevada State Board of Pharmacy Condensed Schedules of Net Position Years Ended June 30, 2021 and 2020

	2021 Actual Government- Wide	2020 Actual Government- Wide
Assets Cash and cash equivalents	\$ 4,310,915	\$ 2,122,747
Accounts and grants receivable	76,636	29,088
Prepaid expenses and deposits	16,021	21,524
Capital assets, net of accumulated depreciation	6,338	19,286
Total assets	4,409,910	2,192,645
Deferred Outflows of Resources	1,165,813	1,302,018
Total assets and deferred outflows of resources	5,575,723	3,494,663
Liabilities		
Accounts payable and accrued expenses	207,217	171,495
Wholesaler license deposits	125,000	100,000
Grants received in advance	33,622	*******
License fees received in advance	2,274,554	694,430
Net other post-employment benefit liability	1,443,826	1,344,606
Net pension liability	3,556,176	3,611,686
Total liabilities	7,640,395	5,922,217
Deferred Inflows of Resources	391,762	365,528
Total liabilities and deferred inflows of resources	8,032,157	6,287,745
Net Position		
Net position	2222	85.5.5
Invested in capital assets	6,338	19,286
Unrestricted	(2,462,772)	(2,812,368)
Total Net Position	\$ (2,456,434)	\$ (2,793,082)

Nevada State Board of Pharmacy Condensed Schedules of Activities Years Ended June 30, 2021 and 2020

	2021 Actual Government- Wide	2020 Actual Government- Wide
Expenses	4 4 770 000	
Operations	\$ 1,778,933	\$ 1,735,954
Personnel	3,411,467	3,073,565
Travel	36,217	75,592
Total expenses	5,226,617	4,885,111
Program Revenue		
Fees, licensing, and permits (charges for services)	3,943,720	2,865,591
General Revenue		
Grant revenue	1,344,052	1,208,099
Investment income	16,783	32,412
Other income	258,710	109,428
Total general revenue	1,619,545	1,349,939
Total revenue	5,563,265	4,215,530
Change in Net Position	\$ 336,648	\$ (669,581)



Independent Auditors' Report on Internal Control over Financial Reporting and on Compliance and Other Matters Based on an Audit of Financial Statements Performed in Accordance with Government Auditing Standards

To the Members of the Board Nevada State Board of Pharmacy Reno, Nevada

We have audited, in accordance with auditing standards generally accepted in the United States of America and the standards applicable to financial audits contained in *Government Auditing Standards* issued by the Comptroller General of the United States, the financial statements of the governmental activities and the remaining fund information of the Nevada State Board of Pharmacy (Board), as of and for the year ended June 30, 2021, and the related notes to the financial statements which collectively comprise the Nevada State Board of Pharmacy's basic financial statements, and have issued our report thereon dated November 2, 2021.

Internal Control over Financial Reporting

In planning and performing our audit of the financial statements, we considered the Board's internal control over financial reporting (internal control) as a basis for designing audit procedures that are appropriate in the circumstances for the purpose of expressing our opinions on the financial statements, but not for the purpose of expressing an opinion on the effectiveness of the Board's internal control. Accordingly, we do not express an opinion on the effectiveness of the Board's internal control.

Our consideration of internal control over financial reporting was for the limited purpose described in the preceding paragraph and was not designed to identify all deficiencies in internal control over financial reporting that might be material weaknesses or significant deficiencies and therefore, material weaknesses or significant deficiencies may exist that have not been identified. However, as described in the accompanying schedule of findings and questioned costs, we identified certain deficiencies in internal control that we consider to be material weaknesses and significant deficiencies.

A deficiency in internal control exists when the design or operation of a control does not allow management or employees, in the normal course of performing their assigned functions, to prevent, or detect and correct, misstatements on a timely basis. A material weakness is a deficiency, or a combination of deficiencies, in internal control, such that there is a reasonable possibility that a material misstatement of the entity's financial statements will not be prevented, or detected and corrected on a timely basis. We consider the deficiency described in the accompanying findings and questioned costs as item 2021-001 to be a material weakness.

A significant deficiency is a deficiency, or a combination of deficiencies, in internal control that is less severe than a material weakness, yet important enough to merit attention by those charged with governance. We consider the deficiency described in the accompanying findings and questioned costs as item 2021-002 to be a significant deficiency.

Compliance and Other Matters

As part of obtaining reasonable assurance about whether the Board's financial statements are free from material misstatement, we performed tests of its compliance with certain provisions of laws, regulations, contracts, and grant agreements, noncompliance with which could have a direct and material effect on the financial statement. However, providing an opinion on compliance with those provisions was not an objective of our audit, and accordingly, we do not express such an opinion. The results of our tests disclosed no instances of noncompliance or other matters that are required to be reported under *Government Auditing Standards*.

Nevada State Board of Pharmacy's Response to Findings

Nevada State Board of Pharmacy's response to the findings identified in our audit are described in the accompanying schedule of findings and responses. Nevada State Board of Pharmacy's responses were not subjected to the auditing procedures applied in the audit of the financial statements and, accordingly, we express no opinion on the responses.

Purpose of this Report

The purpose of this report is solely to describe the scope of our testing of internal control and compliance and the results of that testing, and not to provide an opinion on the effectiveness of the entity's internal control or on compliance. This report is an integral part of an audit performed in accordance with *Government Auditing Standards* in considering the entity's internal control and compliance. Accordingly, this communication is not suitable for any other purpose.

Reno, Nevada

November 2, 2021

Esde Sailly LLP



Independent Auditors' Report on Compliance for the Major Program; Report on Internal Control over Compliance Required by the Uniform Guidance

To the Members of the Board Nevada State Board of Pharmacy Reno, Nevada

Report on Compliance for Each Major Federal Program

We have audited Nevada State Board of Pharmacy' compliance with the types of compliance requirements described in the *OMB Compliance Supplement* that could have a direct and material effect on Nevada State Board of Pharmacy's major federal program for the year ended June 30, 2021. Nevada State Board of Pharmacy's major federal program is identified in the summary of auditor's results section of the accompanying schedule of findings and questioned costs.

Management's Responsibility

Management is responsible for compliance with federal statues, regulations, and the terms and conditions of items federal awards applicable to its federal programs.

Auditor's Responsibility

Our responsibility is to express an opinion on compliance for Nevada State Board of Pharmacy's major federal program based on our audit of the types of compliance requirements referred to above. We conducted our audit of compliance in accordance with auditing standards generally accepted in the United States of America and our 2021 audit in accordance with the standards applicable to financial audits contained in *Government Auditing Standards*, issued by the Comptroller General of the United States; and the audit requirements of Title 2 *U.S. Code of Federal Regulations* Part 200, *Uniform Administrative Requirements, Cost Principles, and Audit Requirements for Federal Awards* (Uniform Guidance). Those standards and the Uniform Guidance require that we plan and perform the audit to obtain reasonable assurance about whether noncompliance with the types of compliance requirements referred to above that could have a direct and material effect on the major federal program occurred. An audit includes examining, on a test basis, evidence about Nevada State Board of Pharmacy's compliance with those requirements and performing such other procedures as we considered necessary in the circumstances.

We believe that our audit provides a reasonable basis for our opinion on compliance for the major federal program. However, our audit does not provide a legal determination of Nevada State Board of Pharmacy's compliance.

Opinion on the Major Federal Program

In our opinion, Nevada State Board of Pharmacy complied, in all material respects, with the types of compliance requirements referred to above that could have a direct and material effect on its major federal program for the year ended June 30, 2021.

Report on Internal Control over Compliance

Management of Nevada State Board of Pharmacy is responsible for establishing and maintaining effective internal control over compliance with the types of compliance requirements referred to above. In planning and performing our audit of compliance, we considered Nevada State Board of Pharmacy's internal control over compliance with the types of requirements that could have a direct and material effect on the major federal program to determine the auditing procedures that are appropriate in the circumstances for the purpose of expressing an opinion on compliance for the major federal program and to test and report on internal control over compliance in accordance with the Uniform Guidance, but not for the purpose of expressing an opinion on the effectiveness of internal control over compliance. Accordingly, we do not express an opinion on the effectiveness of the Nevada State Board of Pharmacy's internal control over compliance.

A deficiency in internal control over compliance exists when the design or operation of a control over compliance does not allow management or employees, in the normal course of performing their assigned functions, to prevent, or detect and correct, noncompliance with a type of compliance requirement of a federal program on a timely basis. A material weakness in internal control over compliance is a deficiency, or a combination of deficiencies, in internal control over compliance, such that there is a reasonable possibility that material noncompliance with a type of compliance requirement of a federal program will not be prevented, or detected and corrected, on a timely basis. A significant deficiency in internal control over compliance is a deficiency, or a combination of deficiencies, in internal control over compliance with a type of compliance requirement of a federal program that is less severe than a material weakness in internal control over compliance, yet important enough to merit attention by those charged with governance.

Our consideration of internal control over compliance was for the limited purpose described in the first paragraph of this section and was not designed to identify all deficiencies in internal control over compliance that might be material weaknesses or significant deficiencies. We did not identify any deficiencies in internal control over compliance that we consider to be material weaknesses. However, material weaknesses may exist that have not been identified.

The purpose of this report on internal control over compliance is solely to describe the scope of our testing of internal control over compliance and the results of that testing based on the requirements of the Uniform Guidance. Accordingly, this report is not suitable for any other purpose.

Reno, Nevada November 2, 2021

Ede Sailly LLP

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Federal Grantor/Pass-Through Grantor/ Program Title	Federal Financial Assistance Listing/Federal CFDA Number	Pass-through Entity Identifying Number	Expenditures
Department of Health and Human Services			
Pass through the State of Nevada Division of Public			
and Behavioral Health			
Injury Prevention and Control	6.5		
Research and State and Community Based Programs	93.136	NU17CE925001-02	\$ 988,321
Injury Prevention and Control	33.130	NO17CE323001-02	7 300,321
Research and State and Community Based			
Programs	93.136	NU17CE925001-01	29,987
			1,018,308
Block Grants for			
Prevention and Treatment of Substance			
Abuse	93.959	3B08TI0010039-19S2	75,000
Total Department of Health and Human Services			1,093,308
Department of Justice			
Bureau of Justice Assistance			
Harold Rogers Prescription Drug Monitoring			
Program	16.754	N/A	249,012
Total Federal Financial Assistance			\$ 1,342,320

Note 1 - Basis of Presentation

The accompanying schedule of expenditures of federal awards (schedule) includes the federal award activity of Nevada State Board of Pharmacy (Board) under programs of the federal government for the year ended June 30, 2021. The information in this schedule is presented in accordance with the requirements of Title 2 U.S. Code of Federal Regulations Part 200, *Uniform Administrative Requirements, Cost Principles, and Audit Requirements for Federal Awards* (Uniform Guidance). Because the Schedule presents only a selected portion of the operations of Nevada State Board of Pharmacy, it is not intended to and does not present the financial position or changes in net position of Nevada State Board of Pharmacy.

Note 2 - Summary of Significant Accounting Policies

Expenditures reported in the schedule are reported on the modified accrual basis of accounting. When applicable, such expenditures are recognized following the cost principles contained in the Uniform Guidance, wherein certain types of expenditures are not allowable or are limited as to reimbursement. No federal financial assistance has been provided to a subrecipient.

Note 3 - Indirect Cost Rate

The Board has elected to use the 10% de minimis indirect cost rate.

Section I - Summary of Auditor's Results

Financial Statements

Type of auditor's report issued:

Internal control over financial reporting:

Material weaknesses identified

Significant deficiencies identified not considered to be material

weaknesses

Noncompliance material to financial statements noted?

Unmodified

Yes

Yes

Federal Awards

Internal control over major program:

Material weaknesses identified

Significant deficiencies identified not considered to be material

weaknesses

None Reported

Type of auditor's report issued on compliance for major programs

Unmodified

Any audit findings disclosed that are required to be reported in

in accordance with Uniform Guidance 2 CFR 200.516

No

Identification of major programs:

Name of Federal Program	CFDA Number
Injury Prevention and Control Research and State and Community Based Programs	93.136
Dollar threshold used to distinguish between Type A and Type B programs:	\$750,000
Auditee qualified as low-risk auditee?	No

Section II - Financial Statement Findings

2021-001: Financial Close and Reporting - Material Weakness

Criteria: Management is responsible for establishing and maintaining an effective system of

internal control over financial reporting. One of the key components of an effective system of internal control over financial reporting is having the capability to prepare full disclosure financial statements in accordance with generally accepted accounting

principles (GAAP).

Condition: We proposed several audit adjustments for corrections to year-end adjustments. In

addition, year-end reconciliation for deferred revenue was not reviewed except for the prepare. The absence of controls over the reconciliations and year-end adjustments of the financial statements and related financial statement disclosures increases the possibility that a misstatement of the financial statements could occur and not be

prevented or detected and corrected in a timely manner.

Cause: Procedures have not been implemented to ensure final review procedures over the

financial statements once all year-end adjustments have been made, including all

reconciliations.

Effect: Financial information prepared by the Board may not comply with generally accepted

accounting principles.

Recommendation: We recommend the Board implement procedures to provide for internal controls over

year-end reconciliations and adjustments.

Views of Responsible

Officials: Nevada State Board of Pharmacy agrees with this finding.

2021-002: Review of the Actuary Reports - Significant Deficiency

Criteria: Management is responsible for establishing and maintaining an effective system of

internal control over financial reporting. Properly reviewing the actuary repots on the pension (PERS) and on the other post-employment benefit obligation (OPEB) is a key

component to effective internal control over financial reporting.

Condition: During our testing over PERS and OPEB related balances, we noted the actuary reports

were not adequately reviewed by Management to ensure consistency with current information and reasonableness over the assumptions used by the actuaries.

Cause: The Board did not have adequate controls to provide for the review of the PERS and

OPEB actuarial reports.

Effect: PERS and OPEB obligation balances at year-end may be misstated and contributions

paid into these plans are not appropriate.

Recommendation: We recommend the Board implement internal controls to provide for the review of the

actuarial reports and retain evidence of such a review.

Views of Responsible

Officials: Nevada State Board of Pharmacy agrees with this finding.

SFY22 MONTHLY BUDGET REPORT NEVADA STATE BOARD OF PHARMACY CURRENT MONTH: Nov 21

			HSOIR		CUBBENT MONTH	PRIOR MONTH(s)	PROJECTIONS THROITIGH	TOTAL BEVENIE/EXPENSE	
REVENUES	-1	APPROVED BUDGET	AMENDMENTS	REVISED BUDGET	REVENUE/EXPENSE	REVENUE/EXPENSE	6/30/2022	SFY22	DIFFERENCE
Beginning Balance	\$	4,267,562	\$ 125,940	\$ 4,393,502	\$	\$ 4,393,502		\$ 4,393,502	•
Renewal Fees	S	1,512,000		\$ 1,512,000	\$ 107,339	\$ 1,413,739	. \$	\$ 1,521,078	\$ 9,078
Registration Fees	S	668,834		\$ 668,834	\$ 102,792	\$ 579,042	\$ 87,000	\$ 768,834	\$ 100,000
Recovered Costs	S		\$ 15,000	\$ 15,000 \$	\$	5 18,177	\$ 25,448	\$ 43,625	\$ 28,625
CC Processing Fees	55		\$ 75,000	\$ 75,000	\$ 8,625	\$ 82,576	\$ 60,373	\$ 151,574	\$ 76,574
Change MGR RPh	\$		\$ 7,500	\$ 7,500	\$ 550	\$ 3,200	\$ 5,250	000'6 \$	\$ 1,500
Inspections	V		\$ 750	\$ 750	\$	\$ 150	\$ 210	\$ 360	\$ (390)
Interest Income	<i>y</i> ,	15,000		\$ 15,000	\$	\$ 1,792	\$ 2,508	\$ 4,300	\$ (10,700)
Indirect Grant Income	55	2,670	\$ (2,670) \$		\$				
Late Fees	5,	3 17,530		\$ 17,530	\$ 2,855	\$ 5,605	\$ 11,844	\$ 20,304	\$ 2,774
Unclaimed Property Refund	0,		\$ 28,050	\$ 28,050	\$	\$ 28,050	. \$	\$ 28,050	. +
Total Revenues	-	\$ 6,483,596	\$ 249,570 \$	\$ 6,733,166	\$ 222,161	\$ 6,525,833	\$ 192,633	\$ 6,940,627	\$ 207,461
EXPENSES	1								
Payroll	~	3,340,540		\$ 3,340,540	\$ 263,814	\$ 1,030,795	\$ 1,821,520	\$ 3,116,129	\$ (224,411)
Operating	55	825,000		\$ 825,000 \$	\$ 065'05 \$	\$ 194,526	\$ 426,694	\$ 671,810	\$ (153,190)
Equipment	35	30,000		\$ 30,000	5	\$ 6,063	\$ 23,937	\$ 30,000	
In-State Travel	55	000'011		\$ 110,000	\$ 617,61 \$	\$ 19,505	\$ 76,775	\$ 110,000	
Out-of-State Travel	57	000'59		\$ 65,000	\$		000'59 \$	\$ 65,000	
DAG Cost	57	12,000		\$ 12,000	\$ 1,467	\$ 2,103	\$ \$	\$ 13,467	\$ 1,467
Aid for Education	\$	2,000		\$ 2,000	\$		\$ 2,000	\$ 2,000	
Reserve	<i>y</i>	2,099,056	\$ 249,570	\$ 2,348,626	\$			\$ 2,932,221	\$ 583,595
Total Expenses	\$	\$ 6,483,596	\$ 249,570 \$	\$ 6,733,166	\$ 329,590 \$	\$ 1,252,993 \$	\$ 2,425,823	\$ 6,940,627	\$ 207,461
Balance									



11/30/2021

The Honorable Members of the Nevada Board of Pharmacy 985 Damonte Ranch Parkway, Suite 206 Reno, Nevada, 89521

RE: Notice of Intent to Act Upon a Regulation – specifically B. Amendment of Nevada Administrative Code 639

Dear Board of Pharmacy Member,

The Hemophilia Alliance is a national organization comprised of the over 120 Hemophilia Treatment Centers that are recognized by the Health Resources Services Administration (HRSA) as uniquely qualified to support the care and treatment of patients with hemophilia. We are sharing comments with you today regarding your October 22, 2021, Notice of Intent to Act Upon a Regulation – specifically B. Amendment of Nevada Administrative Code 639 and the BOP's intent to impose limitations of certain compounded drugs.

In July 2020, Ferring Pharmaceuticals, and its US distributor, CSL Behring, announced a voluntary recall of all batches of STIMATE® Nasal Spray (a solution of desmopressin acetate used to control bleeding in patients with mild hemophilia A and/or type 1 von Willebrand disease [VWD]). Since the July 2020 recall, many patients who relied on STIMATE® for bleeding control have experienced serious hardship. The only alternatives for bleeding events are to use injectable clotting factor medications, or intravenous DDAVP which is administered in the clinic or emergency room. They are both much more expensive and time-consuming treatment for patients, particularly for women of childbearing age, who must treat with injections of clotting factor for three days each month in the absence of desmopressin nasal spray.

With the prospect of a product shortage extending well into 2023, our organizations successfully petitioned the FDA to place STIMATE® on the FDA's 506E National Drug Shortage list. We encouraged STAQ Pharma, Inc., an FDA-registered and FDA-inspected 503B outsourcing facility located in Denver, CO, to produce the appropriate concentration of desmopressin for patient use via nasal spray route of administration. This regulation will effectively prevent the use of this significant effort to use FDA licensed and inspected 503B outsourcing facilities from providing needed patient access to essential therapeutics that, in this case would negatively impact the Nevada hemophilia patient. The complexity of safely compounding this product extemporaneously is significant because of the strength, stability and delivery system (imported from Germany). The FDA requirements for oversight and quality assurance for 503B outsourcing facilities assures patient access and safety that cannot be assured in a one-off compounding process without huge resource commitment from the compounder and the Nevada Board of Pharmacy.



Additionally, Nevada has a proportional share of its bleeding disorder population who live in rural areas and don't have quick and easy access to their Hemophilia Treatment Center (HTC). It is costly for them to take time away from work, costly to travel, and costly to pay their out-of-pocket expenses for more expensive clotting factor. This can and does result in patients forgoing treatment and suffering through the bleeding episodes which potentially could cause more serious long-term complications. Please consider the following:

- The use of a 503B outsourcing facility is the most expedient, safest, and least expensive way to alleviate the 18-month shortage of desmopressin (Stimate®) nasal spray;
- The process HA undertook will safely and cost effectively alleviate this shortage;
- The bleeding disorder community will immediately benefit from a safely and consistently compounded desmopressin nasal spray until such time a commercially available product is re-introduced to the market via a FDA approved 503A facility; and
- Failure to allow the use of this FDA approved process will have a negative impact to Nevada patients limiting their access to a needed, more effective, less costly, quality assured and more easily administered therapeutic agent than previously available to them.

Today, we can offer Nevada patients a safe and effective option for Mild Hemophilia A and VWD. To impose limitations on certain compounded drugs such as those supplied to HTC pharmacies by heavily regulated 503B compounding facilities could lead to a profoundly negative effect on NV bleeding disorder patients – just the opposite of your intentions. The Hemophilia Alliance respectfully asks that you reevaluate your position on imposing limitations on 503B outsourced therapeutic agents especially in such cases like desmopressin nasal spray. This is a safe and effective solution already is in place and effectively protects the bleeding disorders community of Nevada. The Hemophilia Alliance appreciates your consideration and is available to assist you if you wish.

Sincerely, Joseph N Pugliese

President and CEO

Hemophilia Alliance

Joseph Puglisse

Cc: George Oestreich, PharmD, MBA Mark Plencner, R. Ph. Attachments: Letter to Payers







July 27, 2021

Pharmacy Director State Medicaid Agency Office Address State and Zip Code

Re: Request for coverage of DDAVP nasal spray

Dear Director,

We write today to respectfully request coverage for Desmopressin Acetate Nasal Spray, on your Medicaid program. The Hemophilia Alliance (HA) is a not–for–profit national organization which advocates on behalf of patients to ensure they are not disadvantaged and to promote policies and procedures leading to successful patient outcomes. The National Hemophilia Foundation (NHF) and Hemophilia Federation of America (HFA) are national non-profit organizations that represent individuals with bleeding disorders across the United States. Our common mission is to ensure that individuals affected by hemophilia and other inherited bleeding disorders have timely access to necessary quality medical care, therapies, and services, regardless of financial circumstances or place of residence.

In July 2020, Ferring Pharmaceuticals, and its US distributor, CSL Behring, announced a voluntary recall of all batches of STIMATE® Nasal Spray (a solution of desmopressin acetate used to control bleeding in patients with mild hemophilia A and/or type 1 von Willebrand disease). The US recall was part of a global recall of the product, initiated when Ferring detected out-of-specification assay results in some vials of STIMATE® marketed outside the US. By letter dated February 3, 2021, Ferring noted that it had completed its investigation, was working to address the issues, and advised our organizations that it expects to restart manufacturing for first deliveries of STIMATE® to the market no earlier than the second half of 2023.

Since the July 2020 recall, many patients who relied on STIMATE® for bleeding control have experienced serious hardship. Desmopressin nasal spray has been an important therapeutic for these patients for management of bleeding complications and to facilitate surgical procedures and has been a first-line therapy for more than 25 years. Alternatives to STIMATE® include infusion products (which patients may not know how to self-administer). Pandemic circumstances, of course, heighten the difficulty for patients who may now need to visit emergency rooms or doctors' offices to obtain treatment for acute bleeding episodes. Clearly, a self-administered intranasal spray product is the preferred and least costly route of administration for the patient as compared to visiting an infusion center for an IV or an emergency room where costs are much greater.

With the prospect of a product shortage extending well into 2023, our organizations successfully petitioned the FDA to place STIMATE® on the FDA's 506E National Drug Shortage list. The Hemophilia Alliance then began searching for another source of a desmopressin nasal spray that could meet patients' needs. We encouraged STAQ Pharma, Inc., an FDA-registered and FDA-inspected 503B outsourcing facility located in Denver, CO, to produce the appropriate concentration of desmopressin for patient use via nasal spray route of administration. They have produced a product that has successfully passed the 75-day stability mark. We plan to start shipping August 1st. STAQ will continue to test the product at 30-day intervals with the goal of reaching 12-month dating.

We therefore ask that State Medicaid Agency cover this product as you did Stimate when it was available (using your analogous edits and policy). We are hopeful that this new treatment option will help patients who currently have no therapeutically equivalent options to treat their bleeding disorder in a timely and appropriate manner. It is also likely that the renewed availability of this therapeutic will provide lower cost and reduce the need for other services for the appropriate hemophilia patient. The product will be added to First DataBank™ and Medi-Span™. Please see Attachment A for additional information.

Thank you very much for your consideration. For more information, please contact: Joe Pugliese via cell: (215) 439-7173 or email: joe@hemoalliance.org or George L. Oestreich, PharmD., MPA at (573)230-7075 or george@hemoalliance.org.

Sincerely,

Hemophilia Federation of America National Hemophilia Foundation The Hemophilia Alliance

Exhibit A

Desmopressin Acetate Nasal Spray 1.5 mg/ml 0.6ml (6 sprays of 150mcg) vial* NDC: 73177-0114-15

J Code 3490

WAC price: \$400 per vial

Actual acquisition price (AAC): \$300 per vial

Terms: Net 30 days

DDAVP Dosages: 150 µg for body weight <50 kg (one spray into one nostril);

300 μg for body weight >50 kg (one spray into each nostril)

There are 6 actuations per vial which yields 3-6 doses depending on the size of the patient (see references for additional information).

References

https://reference.medscape.com/drug/ddavp-stimate-noctiva-desmopressin-342819

https://ashpublications.org/blood/article/90/7/2515/238641/Desmopressin-DDAVP-in-the-Treatment-of-Bleeding

The Hemophilia Alliance mailing address is:
Hemophilia Alliance
20 Vine Street, #1227
Lansdale, PA 19446



Joshua J. Hicks jhicks@mcdonaldcarano.com Reply to Reno

November 22, 2021

Via Email and U.S. Mail

Dave Wuest, Executive Secretary Nevada State Board of Pharmacy 985 Damonte Ranch Pkwy Suite 206 Reno Nevada 89521 dwuest@pharmacy.nv.gov

Re: Proposed Regulation LCB File No. R002-21

Dear Mr. Wuest.

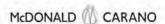
Our office represents Sincerus Pharmaceuticals, Inc. ("Sincerus"). Sincerus objects to the Board's continued attempts to adopt the premature and unnecessary regulation LCB File No. R002-21 (the "Regulation") without meaningfully engaging with any of the affected stakeholders: patients, hospitals, physicians and outsourcing facilities. The Regulation should be rejected—or, at a minimum, held in abeyance—for each of the independent reason listed below.

Pursuant to NRS 233B.0603(1), Sincerus, as an interested party, requests that the Board issue a meaningful statement of the need for and purpose of the Regulation, incorporating the Board's responses to each of the considerations below urged against the Regulation's adoption.

1. The Regulation Is More Stringent Than, and Is Preempted by, Section 503B of the Federal Food, Drug and Cosmetic Act

The Board states in its Notice of Intent—contrary to a position it had maintained previously—that the Regulation "is not required by federal law" and that the Board "is not aware of any similar federal regulation amendments of the same regulation activity in which the state regulation is more stringent." This is false in at least two ways.

First, the Board had previously maintained that the Regulation was required under Section 503B of the FDCA, 21 U.S.C. § 353(b)(a)(8), which is incorporated into Nevada law by NAC 639.6915. Sincerus, and others, have submitted extensive discussion of Section 503B to the Board in the past demonstrating that Section 503B permits products compounded by outsourcing facilities to be furnished to hospitals and physicians for further dispensing to patients—the very



conduct that the Regulation now seeks to prohibit. Without addressing any of these legal arguments, the Board now claims, for the first time, that the Regulation is not required by Section 503B after all.

More important, the Board's attempt to sidestep Section 503B does not resolve the core of the problem: the Regulation is more stringent than federal regulation on the very same issue, and is in fact preempted by federal law. Federal law permits the dispensing of compounded products prepared by outsourcing facilities by entities other than the outsourcing facilities themselves. Specifically, the statute provides, in relevant part, that certain exemptions applicable to outsourced products will apply if:

The drug will not be sold or transferred <u>by an entity other than</u> the outsourcing facility that compounded such drug. <u>This</u> paragraph <u>does not prohibit</u> administration of a drug in a health care setting or <u>dispensing</u> a drug pursuant to a prescription executed in accordance with [21 USC 353(b)(1)].

21 USC 353b(a)(8) (emphasis added).

It is clear from the text of the federal statute that Congress intended to prohibit a 503B outsourcing facility from selling compounded drug product to a *wholesaler*. In other words, Congress intended for there to be only a single transaction between the entity compounding the drug product and the entity dispensing or administering the product in a healthcare setting. This setup would allow for the traditional route for outsourced compounded products, whereby a healthcare provider purchases a compounded drug product and subsequently dispenses and/or administers the product to a patient.

The Regulation prohibits this very conduct and is thus more stringent that federal law. More important, according to federal preemption principles, a state law or requirement is preempted when it "stands as an obstacle to the accomplishment and execution of the full purposes and objectives of Congress." Thus, state requirements are preempted where they interfere with the delicate balance of policy objectives undertaken by Congress and/or a federal agency such as

¹ December 12, 2021 Letter to the Board from Hogan Lovells LLP on behalf of Sincerus Pharmaceuticals, Inc.; December 11, 2021 Letter from the Outsourcing Facilities Association. These letters are incorporated herein by reference in full.

² See Arizona v. United States, 132 S Ct. 2492, 2501 (2012) (quoting Hines v. Davidowitz, 312 US 341, 349 n.4 (2001)).



FDA.³ In addition, state requirements are preempted where they undermine a federal policy favoring a uniform national approach rather than patchwork state regulation.⁴

Here, the Regulation contradicts the clear language and intent of the statute and will impede the FDA's ability to implement the statute in a coherent and effective manner and ensure a consistent approach to regulated entities nationwide. In fact, the FDA has specifically identified state differences in licensing requirements for outsourcing facilities and differences among states' interpretations of the wholesaling prohibition that might cause as areas of concern. Federal law simply leaves no room for states to take a divergent view on these issues.

2. At Least One Other State Board of Pharmacy Has Taken a Diametrically Opposed Position to the Proposed Regulation

On August 27, 2021, the Ohio Board of Pharmacy published an updated Inspection Guide,⁵ which stated the following:

Personally Furnishing Compounded Drugs Obtained from an Outsourcing Facility

An outsourcing facility is permitted to provide non-patient specific compounded sterile drug products to healthcare professionals. These products are compounded under current good manufacturing practice (CGMP) requirements and the facilities are inspected by the FDA on a risk-based schedule. For more information on outsourcing facilities, including how to find those licensed by the Board of Pharmacy, visit: www.pharmacy.ohio.gov/outsourcing

The Board has confirmed with the FDA that non-patient specific drugs purchased directly from an outsourcing facility may be further prescribed and personally furnished to a patient. Please be advised that the 7-day supply limitation that applies to personally furnishing compounded drugs provided by a pharmacy (see 4729:7-2-05 (E)) does not apply to compounded drugs purchased from an outsourcing facility.

³ See Buckman Co. v. Plaintiffs' Legal Committee, 531 U.S. 341, 350 (2001).

⁴ Nash v. Florida Indus. Comm., 389 U.S. 235, 239 (1967) ("a state law cannot stand that 'either frustrates the purpose of the national legislation or impairs the efficiency of those agencies of the Federal government to discharge the duties, for the performance of which they were created."). ⁵ State of Ohio Board of Pharmacy, Inspection Guide, Terminal Distributor of Dangerous Drugs Veterinary Clinic, 4 (8/27/21), p.5 (emphasis added). Although this guide refers to veterinary clinics, the Ohio Board of Pharmacy was clarifying rules about outsourced products that apply to all drugs, human and veterinarian. Indeed, Section 503B only applies explicitly to human drugs.

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In the past, the Board has stated that there were no other states that addressed the issue of dispensing of outsourced drugs by hospitals or physicians, and thus, no apparent regulatory conflict would result from the Regulation. As is clear from the above official pronouncement by another Board of Pharmacy, that is no longer the case. The Ohio Board of Pharmacy's position alone should be enough to reject the proposed Regulation. At minimum, the Board should hold the Regulation in abeyance pending further discussions with the FDA and the Ohio Board of Pharmacy.

Critically, the FDA has highlighted the importance of alignment among the states and consistency regarding licensure and registration of compounders registered as outsourcing facilities under Section 503B of the FDCA. For example, as FDA noted in its preliminary recommendations for aligning federal and state regulation of 503B compounding outsourcing facilities, "FDA-State regulation alignment is particularly important in three specific areas: licensure and registration, filling of patient-specific prescriptions, and compliance with current good manufacturing practice (CGMP) requirements." FDA noted that consistency in the federal and state regulation of outsourcing facilities "is important to their success, and by extension, successful implementation of the Drug Quality and Security Act [section 503B]." FDA further highlighted that greater alignment between federal and state requirements would reduce conflicts, burdens, and confusion on the part of outsourcing facilities, which "in turn will help foster the success of the outsourcing facilities, which the [statute] created to help to address national concerns about oversight and quality of compounding practices." The Regulation will undermine all of the foregoing FDA's goals.

3. The Board Continues to Ignore Upcoming FDA Guidance That Will Address the Very Conduct that the Regulation Seeks to Prohibit

As noted in our letter of May 24, 2021, the FDA issued an agenda of all guidance planned for publication in 2021. That agenda included guidance on the "prohibition on Wholesaling Under Section 503B of the Federal Food, Drug and Cosmetic Act." The forthcoming guidance from the FDA is expected to address the exact type of circumstances in which an outsourcing facility such as Sincerus can provide compounded drugs to a health care facility or a practitioner for further dispensing.

⁶ Nash v. Florida Indus. Comm., 389 U.S. 235, 239 (1967) ("a state law cannot stand that 'either frustrates the purpose of the national legislation or impairs the efficiency of those agencies of the Federal government to discharge the duties, for the performance of which they were created."").

⁷ Id.

⁸ Id. at 3-4.



To date, the Board has refused to address the upcoming federal guidance, relying, instead, on the self-professed need to clarify an "ambiguity" in Section 503B by the Regulation. As an initial matter, and as discussed above, there is no ambiguity: the text of the federal statute is clear. In any event, any apparent ambiguity would counsel in favor of, not against, waiting for the FDA to issue formal guidance. Based on my client's discussions with the FDA, such guidance is expected within weeks after the appointment of a new FDA Commissioner. That is, now that President Biden has nominated a new permanent head of the FDA, it is very likely that mere weeks after the Board's December 2, 2021 public hearing on the Regulation, the FDA guidance will issue its guidance that entirely contradict the Regulation. We know this because of the position the FDA has taken with the Ohio Board of Pharmacy, see supra. Accordingly, as in our letter almost six months ago, our client again respectfully requests that the Board suspend proceedings on the Regulation until such time as FDA guidance is issued and any new rules can be crafted to be in harmony with federal law.

4. The Regulation Has Been Promulgated Without Any Meaningful Stakeholder Input

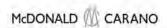
Notwithstanding the serious economic and clinical impact that the proposed Regulation will have, the record reflects *no* attempts by the Board to consult any affected stakeholders in connection with the passage of this Regulation.

First, the Board does not appear to have consulted the FDA (unlike the Ohio Board of Pharmacy). As is apparent from the conversations that the Ohio Board of Pharmacy had with the agency, the FDA intends to clarify (to the extent such clarification is necessary) that outsourced drugs may be further dispensed to patients by hospitals or physicians.

Second, the Board does not appear to have consulted other Boards of Pharmacy, for the same reasons noted above.

Third, the Board does not appear to have consulted hospitals or hospital associations. The proposed Regulation also prohibits hospital outpatient dispensing of outsourced drugs—including drugs covered by the 340B program—which is a widely used practice for non-patient-specific drugs—providing patient access to unavailable drugs by the safest means possible. For example, 503B-compounded drugs are often dispensed on an outpatient basis for home hospice use and for home dialysis. Outsourcing facilities have a played a critical role in addressing chronic shortages of a number of outpatient drugs, including peritoneal dialysis solutions, which have been exacerbated during the COVID-19 emergency. In addition, outsourcing facilities frequently

⁹ See Advancing American Kidney Health, Executive Order 13879, 84 FR 33817, 33818 (July 15, 2019) (creating payment incentives for greater use of home dialysis to "improve quality of life and care for patients who require dialysis.").



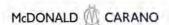
compound for outpatient dispensing certain drugs for parenteral antimicrobial therapy, where customized dosing and admixture are often necessary. ¹⁰ The Regulation entirely prohibits such outpatient dispensing and will have significant impact on patient care. We attach a list of 125 compounded products typically dispensed by hospitals and physicians that will be affected by the Regulation.

Fourth, the Board does not appear to have consulted any patient advocates. As discussed above, patients requiring outpatient compounded drugs—prepared in a cGMP environment in FDA-regulated 503B outsourcing facilities—will not be able to obtain them from the hospital or from their physician. This will result in diminished patient care, medication adherence and unnecessary delays.

Fifth, the Board does not appear to have consulted any outsourcing facilities. In particular, the Board has stated—without any analysis or justification—that the Regulation "should have no adverse economic impact . . . on the regulated entities or on the public." Not so. The Regulation will cause severe (or existential) harm to outsourcing facilities that prepare compounds for further dispensing by hospitals or physicians. The Regulation will also create undue regulatory burden by requiring outsourcing facilities to comply with an inconsistent hodge-podge of state regulatory requirements, including the Regulation's requirement to have policies to prevent dispensing by hospitals and physicians (addressed further below). Further, the Regulation effectively relegates outsourcing facilities to the level of 503A compounding pharmacies (which can only dispense pursuant to patient-specific prescriptions)—none of which have to expend the significant resources to register with the FDA and implement strict cGMP manufacturing requirements. Accordingly, the Regulation creates significant economic disincentives for outsourcing facilities to register as such with the FDA, undermining the federal scheme, and incentivizing potentially less safe compounded practices by other actors.

Sixth, the small business impact statement included in the Notice of Intent to Act Upon a Regulation dated October 22, 2021 ("Notice") is so generic as to be meaningless. The purported need for the Regulation is "for the protection, health and safety of the public." There are no examples provided of how the public's protection, health and safety has been in jeopardy. The Notice states that "there should be no adverse economic impact from this regulation amendment on the regulated entities or on the public." This is now the third letter submitted by my client pointing out adverse economic impacts. The Notice says the Regulation will "improve the delivery

¹⁰ See, e.g., 2018 Infectious Disease Society of America Clinical Practice Guideline for the Management of Outpatient Parenteral Antimicrobial Therapy, available at https://academic.oup.com/cid/article/68/1/e1/5175018#124458579. These drugs, which are compounded by outsourcing facilities and relied upon for outpatient dispensing, include cefazolin, ceftazidime, gentamicin, metronidazole, polymyxin B, vancomycin, and sodium thiosulfate.



of safe and reliable pharmaceutical care in Nevada." No examples or evidence for this statement are provided. The Notice is nothing more than a series of boilerplate self-serving conclusory statements that lack any kind of meaningful detail and omits adverse impacts on regulated entities. The Notice is therefore defective and fails to meet the requirements of Nevada law.

5. The Regulation Is Arbitrary and Will Not Advance Its Professed Justification

The sole justification offered for the Regulation is that the Regulation is "necessary for the protection, health and safety of the public" and that the Regulation will "ensur[e] the delivery of safe and reliable pharmaceutical care." On its face, the Regulation does not advance this goal, and is arbitrary.

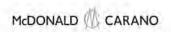
The premise of the Regulation appears to be that drugs compounded by outsourcing facilities are not "safe or reliable," undermine public "health and safety," and, as such, ought not be dispensed to patients. 11 Curiously, the Regulation nonetheless allows that these purportedly unsafe drugs be dispensed to patients by the outsourcing facilities themselves. The Board does not explain—nor could it—how a drug that is purportedly unsafe to be dispensed by a hospital or physician may nonetheless be dispensed by the outsourcing facility itself. This contradiction belies the very foundation of need for the Regulation, and underscores the arbitrariness of the Regulation.

The truth is that the FDA has repeatedly advocated for providers, including hospitals, to obtain their compounded drugs from outsourcing facilities precisely because they are manufactured in a safer, cGMP compliant environment. Most recently, the FDA issued a draft guidance on Hospital and Health System Compounding Under Section 503A of the Federal Food, Drug, and Cosmetic Act, where it repeatedly encouraged hospitals to procured medications from outsourcing facilities:

To the extent that hospitals and health systems have a need for compounded drug products, FDA encourages them to obtain such products from outsourcing facilities.

Outsourcing facilities, which are subject to CGMP requirements and other conditions that help to ensure drug quality and a clinical need for the drug products they compound from bulk drug substances, may compound and distribute drug products to healthcare facilities without first receiving prescriptions for identified individual patients.

¹¹ This premise is false because outsourcing facilities are subject to CGMP requirements and other conditions that help to ensure drug quality.



We encourage hospitals and health systems to look to outsourcing facilities, or to register their pharmacies as outsourcing facilities, when they wish to obtain non-patient-specific compounded drug products.¹²

The Board has articulated no reasoned, factual basis to support its claim that drugs compounded by outsourcing facilities—inspected by the FDA and state boards, and required to comply with strict cGMP requirements—are unsafe for the general public, particularly where the Board allows the very same drugs to be administered to patients by providers and *dispensed* by outsourcing facilities themselves.

6. The Regulation Is Duplicative and Unnecessary

Although not stated in the current Notice of Intent, the Board's prior justification for moving the Regulation forward was a supposed need to protect the "new drug application process," which some Board members felt may be undermined by outsourcing facilities manufacturing drugs in bulk without seeking a new drug application from the FDA. The Regulation is entirely unnecessary to achieve this goal, which, in any event, is the purview of the federal agency, not this Board.

Section 503B—which is incorporated into Nevada law by NAC 639.6915—already contains adequate mechanism to protect the "new drug application process." In particular, Section 503B prohibits outsourcing facilities from compounding drugs that are "essentially a copy of an approved drug." Both the FDA, and other boards of pharmacy, have taken enforcement actions when outsourcing facilities violated this provision. The Regulation does not additionally advance this already-addressed goal—instead, it creates undue burden, cost and impact on patients, doctors, hospitals and outsourcing facilities. Simply put, the Regulation is duplicative and unnecessary.

7. The Regulation Contains Impracticable Provisions

The Regulation purports to mandate that outsourcing facilities "establish and implement procedures to prohibit other persons and entities from dispensing . . . drugs compounded by the outsourcing facilities." However, it is entirely unclear (and the Board has provided no guidance) how an outsourcing facility could ever "implement" these unspecified procedures over third-party entities that it does not control. In effect, the Board intends to hold outsourcing facilities responsible for the actions of unrelated third parties. Unsurprisingly, no such regulations exist for any other drug manufacturers for the simple reason that they are impractical and imbue the Board

¹² FDA, Hospital and Health System Compounding Under Section 503A of the Federal Food, Drug, and Cosmetic Act, Guidance for Industry, October 2021.



with an unreasonable degree of enforcement discretion in potential violation of the parties' due process rights.

* *

Regulation R002-21 is premature, overly broad, and unnecessary. The Regulation should be put on hold or sent back to workshop for redrafting. Passing the Regulation at this point in time will create confusion and delay in the drug distribution supply chain, negatively impacting health care providers and their patients. Thank you for your consideration of these comments, and we ask that you make this letter part of the administrative record at the upcoming hearing on the Regulation currently scheduled for December 2, 2021.

Sincerely,

Joshua J. Hicks

Att.

JJH/ns

Cc: Brett Kandt, General Counsel to the Nevada State Board of Pharmacy (via email to bkandt@pharmacy.nv.gov)

Outsourcing facilities are required to provide FDA with a list of drugs they compounded during the previous six-month period upon initial registration and in June and December each year. The following represents common preparations that outsourcing facilities reported to the FDA for the reporting period ending December 2019 (2019-2). Injectable preparations have been excluded. It is common for prescribers to order compounded dosage forms including but not limited to creams, capsules, gels, ophthalmic drops, pastes, tablets, and troches to then dispense this compounded medication to patients.

Active Ingredients	Active Ingredients Info	Dosage	Package Description
.alphaEstradiol 3-Benzoate; Estriol	.0001 g/1 g; .0004 g/1 g	cream	30 G in 1 Jar
.alphaEstradiol 3-Benzoate; Estriol; Testosterone	.00045 g/l g; .00105 g/l g; .008 g/l g	cream	30 G in 1 Jar
7-Keto-Dehydroepiandrosterone	.05 g	capsule	1 Capsule in 1 Vial, Multi- Dose
Aluminum Hydroxide; Magnesium Hydroxide; Lidocaine Hydrochloride; Diphenhydramine Hydrochloride; Dimethicone	13.36 mg/1 mL; 13.36 mg/1 mL; 6.68 mg/1 mL; .835 mg/1 mL; 1.336 mg/1 mL	suspension	5 Ml in 1 Syringe
Anastrozole	.00025 g	capsule	1 Capsule in 1 Vial, Plastic
Benzocaine	95 mg/1 g	paste	480 G in 1 Jar
Benzocaine; Lidocaine Hydrochloride; Tetracaine Hydrochloride	.12 g/1 g; .04 g/1 g; .08 g/1 g	cream	240 G in 1 Jar
Benzocaine; Lidocaine; Tetracaine	.2 g/1 g; .06 g/1 g; .04 g/1 g	gel	100 G in 1 Jar
Butalbital	50 mg/50 mg	capsule	50 Mg in 1 Capsule

Calcium Gluconate; Propylene Glycol; Methylcellulose (1500 Mpa.s)	25 mg/1 g; 20 mg/1 g; 30 mg/1 g	gel	454 G in 1 Jar
Cantharidin	10 mg/1 mL	gel	10 Ml in 1 Vial, Multi- Dose (5446- 0572-03)
Chloramphenicol; Sulfamethoxazole; Amphotericin B	50 mg; 50 mg; 5 mg	capsule	I Capsule in 1 Not Applicable (5446-1157- 03)
Cholecalciferol	50000 [iU]/1 mL	suspension	10 Ml in 1 Vial
Chorionic Gonadotropin	600 [iU]	troche	1 Troche in 1 Vial, Plastic
Cyclosporine	20 mg/1 mL	suspension	10 Ml in 1 Bottle, Dropper
Diclofenac Sodium	16 mg/1 g	gel	24 Packet in 1 Box (71300- 6573-1) > 2.5 G in 1 Packet
Diltiazem Hydrochloride	20 mg/1 g	cream	30 G in 1 Jar (5446-1086- 03)

Diphenhydramine Hydrochloride; Magnesium Hydroxide; Aluminum Hydroxide; Dimethicone; Lidocaine Hydrochloride	.8 mg/1 mL; 13.4 mg/1 mL; 13.4 mg/1 mL; 1.34 mg/1 mL; 6.6 mg/1 mL	suspension	5 Ml in 1 Syringe, Plastic (62295-5023- 5)
Doxycycline	100 mg/1 mL	suspension	240 Ml in 1 Bottle
Doxycycline Anhydrous	50 mg/1 mL	suspension	240 Ml in 1 Bottle
Doxycycline Hyclate	300 mg	tablet	100 Tablet in 1 Bottle (71591-022- 01)
Doxycycline Hydrochloride	50 mg/50 mg	suspension	12000 Mg in 1 Bottle
Edetate Disodium	30 mg/1 mL	solution/ drops	10 MI in I Bottle, Dropper (70360-058- 39)
Enrofloxacin	204 mg	tablet	100 Tablet in 1 Bottle (71591-029- 01)
Estriol	2 mg	capsule	100 Capsule in I Vial, Plastic

			(26436-5343- 1)
Estriol; .alphaEstradiol 3-Benzoate; Testosterone	.001 g/1 g; .000257 g/1 g; .0005 g/1 g	cream	30 G in 1 Jar
Estriol; Estradiol	2 mg/1 g; .5 mg/1 g	cream	30 G in 1 Bottle, Pump (26436-5228- 3)
Estriol; Testosterone; Estradiol	1 mg/1 g; 1.5 mg/1 g; 1 mg/1 g	cream	35 G in 1 Bottle, Pump (26436-5364- 3)
Famotidine	10 mg/10 mg	suspension	4800 Mg in 1 Container
Fluconazole	300 mg	tablet	100 Tablet in 1 Bottle (71591-031- 01)
Fluocinonide; Hydroquinone; Tretinoin	.0001 g/1 g; .04 g/1 g; .0005 g/1 g	cream	30 G in 1 Jar
Fluoxetine Hydrochloride	5 mg/5 mg	capsule	10 Mg in 1 Capsule
Gabapentin	25 mg	tablet	100 Tablet in 1 Bottle (71591-035- 01)
Human Chorionic Gonadotropin	1000 [iU]	tablet, orally disintegrating	1 Tablet, Orally

			Disintegrating in 1 Vial, Plastic
Hydrocodone Bitartrate	10 mg/10 mg	capsule	10 Mg in 1 Capsule
Hydrocortisone; Chloramphenicol; Sulfamethoxazole; Amphotericin B	25 mg/1 g; 50 mg/1 g; 50 mg/1 g; 5 mg/1 g	capsule	1 G in 1 Not Applicable (5446-0457- 03)
Hyoscyamine Sulfate; Magnesium Hydroxide; Aluminum Hydroxide; Dimethicone; Lidocaine Hydrochloride	.007 mg/l mL; 17 g/l mL; 17 g/l mL; 1.7 g/l mL; 1.7 g/l mL; 5.7 mg/l mL	suspension	35 MI in 1 Bottle, Dispensing
Ibutamoren Mesylate	.025 g	capsule	1 Capsule in 1 Vial, Plastic
Ketoconazole; Amikacin Sulfate; Triamcinolone Acetonide	1 g/100 mL; .67 g/100 mL; .1 g/100 mL	ointment	3 Ml in I Syringe
Ketoconazole; Enrofloxacin; Triamcinolone Acetonide	1 g/100 mL; .23 g/100 mL; .1 g/100 mL	ointment	3 MI in 1 Syringe (71591-004- 06)
Ketoconazole; Gentamicin Sulfate; Triamcinolone Acetonide	1 g/100 mL; .67 g/100 mL; .1 g/100 mL	ointment	3 Ml in 1 Syringe
Ketoprofen	100 mg/1 g	cream	30 Packet in 1 Box (71300- 6442-1) > 1 G in 1 Packet

Levocetirizine Dihydrochloride; Clobetasol Propionate	20 g/100 mL; .05 g/100 mL	shampoo	120 MI in 1 Bottle, Plastic
Lidocaine	300 mg/1 g	ointment	114.12 G in 1 Bottle
Lidocaine Hydrochloride	40 mg/40 mg	solution/ drops	120 Mg in 1 Bottle, Dropper
Lidocaine Hydrochloride; Prilocaine Hydrochloride; Tetracaine Hydrochloride; Phenylephrine Hydrochloride	100 mg/1 g; 100 mg/1 g; 40 mg/1 g; 20 mg/1 g	gel	34 G in 1 Jar
Lidocaine Hydrochloride; Tetracaine	150 mg/150 mg; 150 mg/150 mg	cream	900 Mg in 1 Syringe, Plastic
Lidocaine Hydrochloride; Tetracaine Hydrochloride	.23 g/1 g; .07 g/1 g	cream	120 G in 1 Jar
Lidocaine Hydrochloride; Tetracaine Hydrochloride; Epinephrine Bitartrate	.04 g/1 g; 5 mg/1 g; .5 mg/1 g	gel	3 G in 1 Syringe, Plastic
Lidocaine Hydrochloride; Tetracaine Hydrochloride; Phenylephrine Hydrochloride	200 mg/1 g; 40 mg/1 g; 20 mg/1 g	gel	30 G in 1 Jar
Lidocaine Hydrochloride; Tetracaine Hydrochloride; Prilocaine Hydrochloride	100 mg/1 g; 40 mg/1 g; 100 mg/1 g	gel	100 G in 1 Jar
Lidocaine Hydrochloride; Tetracaine; Epinephrine	30 mg/30 mg; 30 mg/30 mg; 30 mg/30 mg	gel	270 Mg in 1 Syringe, Plastic

Lidocaine; Benzocaine; Tetracaine	2.4 g; 6 g; 2.4 g	cream	1 Cream in 1 Jar (73271- 2088-1)
Lidocaine; Dibucaine Hydrochloride; Phenylephrine Hydrochloride; Prilocaine	15 g/100 mL; .5 g/100 mL; 10 g/100 mL; 50 g/100 mL	ointment	120 Ml in 1 Bottle, Pump
Lidocaine; Prilocaine Hydrochloride; Tetracaine Hydrochloride	100 mg/1 mL; 100 mg/1 mL; 40 mg/1 mL	gel	30 Ml in 1 Jar (5446-0407- 10)
Lidocaine; Prilocaine Hydrochloride; Tetracaine Hydrochloride; Phenylephrine	100 mg/1 mL; 100 mg/1 mL; 40 mg/1 mL; 20 mg/1 mL	gel	30 Ml in 1 Jar (5446-1018- 10)
Lidocaine; Prilocaine; Tetracaine	1.5 g; 1.5 g; .6 g	gel	1 Gel in 1 Jar (73271-0552- 1)
Lidocaine; Racepinephrine; Tetracaine Hydrochloride	40 mg/1 mL; .5 mg/1 mL; 5 mg/1 mL	gel	5 Ml in 1 Syringe (5446-0607- 01)
Lidocaine; Tetracaine	230 mg/1 g; 70 mg/1 g	cream	100 G in 1 Bottle, Pump (72627-3102- 1)
Lidocaine; Tetracaine; Benzocaine	60 mg/1 g; 40 mg/1 g; 200 mg/1 g	ointment	90 G in I Bottle
Magnesium Hydroxide; Aluminum Hydroxide; Dimethicone; Lidocaine Hydrochloride	1.2 g/1 mL; 1.2 g/1 mL; .12 g/1 mL; 2 mg/1 mL	suspension	40 MI in 1 Bottle, Dispensing

Metformin Hydrochloride	500 mg	capsule	100 Capsule in 1 Vial, Plastic > 1 Capsule in 1 Capsule
Methimazole	25 mg/1 mL	cream	1 Ml in 1 Syringe
Metronidazole	50 mg/50 mg	suspension	24000 Mg in 1 Container
Metronidazole Benzoate	160 mg/1 mL	suspension	240 Ml in 1 Bottle
Mitomycin	.5 mg/l mL	irrigant	1 Syringe in 1 Package (71266-6412- 2) > 40 Ml in 1 Syringe (71266-6412- 1)
Mupirocin; Clotrimazole; Betamethasone Acetate	10 mg/1 g; 20 mg/1 g; .25 mg/1 g	ointment	15 G in 1 Jar (5446-0605- 01)
Mycophenolate Sodium	50 mg/50 mg	suspension	24000 Mg in 1 Container
Naltrexone	.00517 g	capsule	1 Capsule in 1 Vial, Plastic
Naltrexone Hydrochloride	4 mg/4 mg	capsule	4 Mg in 1 Capsule

Neomycin Sulfate; Polymyxin B Sulfate; Dexamethasone	3.5 mg/1 g; 10000 U/1 g; 1 mg/1 g	ointment	.1 G in 1 Syringe, Plastic
Omeprazole	10 mg/10 mg	suspension	4800 Mg in 1 Container
Ondansetron Hydrochloride	8 mg/1 mL	cream	1 Ml in 1 Syringe, Plastic
Oxandrolone	25 mg	capsule	30 Capsule in 1 Vial, Dispensing (26436-1115- 1)
Oxytocin; Tadalafil	50 U; 12 mg	capsule	1 Capsule in 1 Capsule (73271-9125- 1)
Papaverine; Sildenafil	.02 g; .15 g	troche	1 Troche in 1 Vial, Plastic
Pentoxifylline	400 mg	capsule	100 Capsule in 1 Vial, Plastic > 1 Capsule in 1 Capsule
Phenylephrine Hydrochloride; Tropicamide	25 mg/1 mL; 10 mg/1 mL	solution/ drops	15 Ml in 1 Bottle, Dropper

			(5446-0815- 01)
Phenylephrine Hydrochloride; Tropicamide; Ketorolac Tromethamine; Ciprofloxacin Hydrochloride	100 mg/1 mL; 10 mg/1 mL; 1.25 mg/1 mL; 3 mg/1 mL	solution/ drops	1 MI in I Syringe, Plastic (5446- 1270-01)
Phenylephrine; Lidocaine; Tetracaine	.0005 g/1 g; .02 g/1 g; .01 g/1 g	ointment	400 G in 1 Jar
Polyethylene Glycol 3350	1 g/1 g	powder	8.5 G in 1 Pouch
Polymyxin B Sulfate; Neomycin Sulfate	800 U/1 mL; .16 mg/1 mL	irrigant	500 Ml in 1 Bottle (5446- 0746-02)
Ponazuril	90 mg/1 mL	suspension	60 Ml in 1 Bottle (71591-043- 60)
Povidone-lodine	50 mg/1 mL	solution/ drops	.4 Ml in 1 Syringe, Plastic (61141-4538- 4)
Prasterone	.05 g	capsule	1 Capsule in 1 Vial, Plastic
Prasterone; Pregnenolone	.01 g; .025 g	capsule	1 Capsule in 1 Vial, Plastic
Prasterone; Sildenafil	.01 g/1 g; .05 g/1 g	cream	30 G in 1 Vial, Plastic

Praziquantel; Pyrantel Pamoate; Mebendazole	22,7 mg; 65.376 mg; 50 mg	tablet	100 Tablet in 1 Bottle, Plastic
Praziquantel; Pyrantel Pamoate; Oxantel Pamoate	22.7 mg; 65.376 mg; 252 mg	tablet	100 Tablet in 1 Bottle, Plastic
Prednisolone	5 mg/l mL	suspension	240 Ml in 1 Bottle (71591-047- 01)
Prednisolone Sodium Phosphate; Moxifloxacin Hydrochloride	7.5 mg/7.5 mg; 7.5 mg/7.5 mg	solution/ drops	75 Mg in 1 Bottle, Dropper
Pregnenolone; Progesterone	.01 g; .075 g	capsule	1 Capsule in 1 Vial, Plastic
Pregnenolone; Progesterone; Prasterone	.2 g; .02 g; .015 g	capsule	1 Capsule in 1 Vial, Plastic
Prilocaine Hydrochloride; Lidocaine Hydrochloride; Tetracaine	80 mg/80 mg; 80 mg/80 mg; 80 mg/80 mg	gel	720 Mg in 1 Syringe, Plastic
Progesterone	60 mg	capsule	30 Capsule in 1 Vial, Dispensing (26436-5247- 1)
Promethazine Hydrochloride	25 mg/1.2 mL	ointment	3 M1 in 1 Syringe,

		13	Plastic (5446- 1341-01)
Proparacaine Hydrochloride	5 mg/1 mL	solution/ drops	.4 Ml in 1 Syringe, Plastic (61141-4536- 4)
Racepinephrine Hydrochloride; Tetracaine Hydrochloride; Lidocaine Hydrochloride	1.5 mg/3 mL; 15 mg/3 mL; 120 mg/3 mL	gel	3 Ml in 1 Syringe
Salicylic Acid; Ciclopirox	20 g/100 mL; .77 g/100 mL	shampoo	400 Ml in 1 Cartridge
Salicylic Acid; Ciclopirox; Clobetasol Propionate	30 g/100 mL; .77 g/100 mL; .05 g/100 mL	shampoo	120 MI in 1 Bottle, Plastic
Salicylic Acid; Podophyllum Resin; Cantharidin	300 mg/1 mL; 50 mg/1 mL; 10 mg/1 mL	gel	10 Ml in 1 Vial, Multi- Dose (5446- 0970-03)
Secnidazole	140 mg/1 mL	suspension	30 Ml in 1 Bottle (71591-052- 01)
Sildenafil	.154 g	troche	1 Troche in 1 Vial, Plastic
Sildenafil Cítrate	50 mg	troche	1 Troche in 1 Package (73271-4005- 1)

:15 g; .02 g	troche	1 Troche in 1 Vial, Plastic
1000 mg	capsule	1 Capsule in 1 Capsule
.3 mg/1 mL	suspension	10 Ml in 1 Bottle, Dropper
10 mg	troche	1 Troche in 1 Package (73271-8010- 1)
100 mg/1 g	cream	30 G in 1 Jar (26436-0163- 5)
.01 g; .02 g	tablet	l Tablet in l Vial, Plastic
160 mg	capsule	1 Capsule in 1 Capsule (5446-1179- 03)
60 mg/1 g.	gel	10 G in 1 Syringe
106 mg/106 mg; 106 mg/106 mg; 106 mg/106 mg	cream	960 Mg in 1 Syringe
5 mg/1 mL; 1 mg/1 mL; 40 mg/1 mL	gel	3 Ml in 1 Syringe, Plastic
	1000 mg .3 mg/1 mL 10 mg 100 mg/1 g .01 g; .02 g 160 mg 106 mg/106 mg; 106 mg/106 mg; 106 mg/106 mg 5 mg/1 mL; 1 mg/1 mL; 40 mg/1	1000 mg capsule

Tetracaine; Benzocaine; Lidocaine	40 mg/1 mL; 200 mg/1 mL; 60 g/1 mL	cream	120 Ml in 1 Bottle, Pump (70713-103- 01)
Tetracaine; Lidocaine	.07 g/1 g; .23 g/1 g	ointment	400 G in 1 Jar
Tetracaine; Lidocaine; Benzocaine	40 mg/1 g; 60 mg/1 g; 200 mg/1 g	cream	100 G in 1 Jar
Tetracaine; Phenylephrine; Prilocaine; Lidocaine	1.2 g; .6 g; 3 g; 3 g	gel	1 Gel in 1 Jar (73271-1142- 1)
Tramadol Hydrochloride	50 mg/1 mL	suspension	60 Ml in 1 Bottle, Plastic
Tretinoin	.02 g/1 g	cream	30 G in 1 Vial, Plastic
Triamcinolone Acetonide	50 mg/1 mL	suspension	2 Ml in 1 Vial (62295-3317- 2)
Tropicamide; Phenylephrine Hydrochloride	10 mg/1 mL; 25 mg/1 mL	solution/ drops	1 Bottle, Dropper in 1 Bag (71449- 098-47) > 10 Ml in 1 Bottle, Dropper
Tropicamide; Phenylephrine Hydrochloride; Cyclopentolate Hydrochloride	10 mg/1 mL; 25 mg/1 mL; 10 mg/1 mL	solution/ drops	1 Bottle, Dropper in 1 Bag (71449- 093-46) > 5 Ml in 1

			Bottle, Dropper
Ursodiol	300 mg	tablet	100 Tablet in 1 Bottle
Vardenafil	.075 g	troche	1 Troche in 1 Vial, Plastic



November 23, 2021

Nevada Board of Pharmacy 985 Damonte Ranch Pkwy Ste 206 Reno, NV 89521

RE: Comment on Proposed Amendment to NAC 639; Dispensing Drugs Compounded by a Pharmacy, Practitioner or Outsourcing Facility

Dear Nevada Board of Pharmacy:

Epicur Pharma, a registered 503 outsourcing facility, thanks the Board for considering comments before making a final decision to amend the regulation at issue, NAC 639.

We hope the Board will reconsider the restriction on allowing an outsourcing facility product to be dispensed. Outsourcing facility products are safer and have higher assurances of quality than traditionally compounded preparations. By placing the same restrictions on cGMP outsourcing facility products as those placed on preparations compounded to meet USP standards, the Board is exposing Nevada residents to needless and avoidable risk.

FDA has Recently Spoken on this Issue and the DQSA Clearly Allows Dispensing

The FDA, which is tasked with primary regulation of outsourcing facilities, has recently spoken on the subject of further dispensing 503B products and has informed at least one other board of pharmacy (Ohio's) that cGMP outsourcing facility products may be further dispensed by practitioners (see attached guidance). The proposed rule the Nevada Board is considering directly contradicts the FDA's position in a sector of industry that is primarily regulated by the FDA. FDA has also given public notice that it will be releasing a guidance document this year on the issue of further dispensing. At a minimum the Nevada Board should wait to see what that guidance document conveys before adopting the proposed rule.



The Federal Government has stated clearly that there are two exceptions to the not for resale language in 503B's prohibition on wholesaling. Administration OR dispensing pursuant to a prescription.

21 U.S.C.S. § 353b

(8) Prohibition on wholesaling. The drug will not be sold or transferred by an entity other than the outsourcing facility that compounded such drug. This paragraph does not prohibit administration of a drug in a health care setting **Of dispensing a drug pursuant to a prescription** executed in accordance with section 503(b)(1) [21 USCS § 353(b)(1)].

In Nevada's proposed rule the Board is clearly acknowledging the first exception of administration, and clearly ignores the second. The reason for ignoring the second exception has not been addressed in any of its Board meetings discussing this issue.

Other States have Spoken on this Issue and Allow Dispensing

States are beginning to address practitioners dispensing 503B products. Recently, California, Georgia, Kansas, Minnesota, Missouri, Ohio, Pennsylvania, and Utah have clarified that federal law should be followed in their state regarding the dispensing of 503B products. As noted earlier, the DQSA is clear that dispensing is allowed and FDA has clarified to Ohio that outsourcing facility products may be further dispensed to clients by a practitioner per federal law.

The FDA's NDA Process is Already Protected by the FDA

The Nevada Board's previously stated goal of protecting the FDA's drug approval process by enacting this rule is unnecessary. The FDA protects its NDA process with its Essentially Copies guidance document (attached). Through this guidance FDA protects its NDA process and prevents outsourcing facilities from manufacturing copies of any marketed FDA approved drug. In addition, FDA requires outsourcing facilities to report every human use drug it manufactures to the FDA once every 6 months. By requiring these reports FDA is alerted to any copies of FDA approved drugs every 6 months.



CGMP Manufacturing Differs Significantly from USP Compounding

Traditionally compounded preparations are subject to USP standards. These standards were designed to safely supply drug preparations for one specific patient. In contrast, outsourcing facilities are required to follow cGMP standards. cGMP standards are designed to manufacture large batches to safely supply drug products for widespread distribution to patients. As the Board is aware, cGMP standards differ greatly from USP in terms of method, testing, documentation and even mindset. What the Board may not be aware of, is that outsourcing facilities are held to the same cGMP standards that traditional manufacturers are held to. There is no such thing as reduced cGMP or cGMP-lite. The FDA holds 503B outsourcing facilities to the same manufacturing standards that companies such as Pfizer or Johnson & Johnson are held to under cGMP.

Treating USP Compounds and CGMP Products the Same puts Nevada Residents at Risk

The higher assurances of quality that cGMP supplies, which allow for safe widespread distribution, come at a price. It is much more expensive to manufacture a cGMP product than it is to compound a preparation. By grouping all "compounding" together as the Board proposes to do in the proposed amendment at issue, practitioners will do the same. If a practitioner views all compounded products as equivalent, with the only difference being price, the practitioner will be incentivized to purchase the less expensive USP preparation. This unquestionably puts Nevada patients at higher risk of receiving a less safe product.

In addition, outsourcing facilities may be disincentivized from being licensed in Nevada, again depriving Nevada patients, human and animal, from having access to higher quality, safer, outsourcing facility products.



S. P. Zamanya Parway, Salata A. Marcillani, A. 1818. A 1883 S. 1937. Francisco Mar

Conclusion

We ask that the Board allow practitioners to administer and dispense outsourcing facility products as federal law unambiguously allows.

Respectfully submitted,

Michael Juliano

In-House Counsel Epicur Pharma 856-988-1889 Mjuliano@EpicurPharma.com



INSPECTION GUIDE

Terminal Distributor of Dangerous Drugs Veterinary Clinic

Updated 8/27/2021

To review updates, please see the <u>update history</u> section at the end of this document.

This document is reference material for licensees and applicants. The document does not bind the State of Ohio Board of Pharmacy, and does not confer any rights, privileges, benefits, or immunities for or on any person, applicant or licensee.

Positive Identification Guidance

"Positive identification" means a method of identifying a person that does not rely on the use of a private personal identifier such as a password, but must use a secure means of identification that includes any of the following:

- (1) A manual signature on a hard copy record;
- (2) A magnetic card reader;
- (3) A bar code reader;
- (4) A biometric method;
- (5) A proximity badge reader;
- (6) A board approved system of randomly generated personal questions;
- (7) A printout of every transaction that is verified and manually signed within a reasonable period of time by the individual who performed the action requiring positive identification. The printout must be maintained for three years and made readily retrievable; or
- (8) Other effective methods for identifying individuals that have been approved by the board.

NOTE: A method relying on a magnetic card reader, a bar code reader, a proximity badge reader, or randomly generated questions for identification must also include a private personal identifier, such as a password, for entry into a secure mechanical or electronic system.

REMINDER: Positive identification should be at the conclusion of a drug transaction. For electronic systems, positive identification required at log-in does not document the specific drug transaction and causes other security problems. For example, a nurse does not document the administration of a medication when they log in to an electronic drug record keeping system.

Personally Furnishing Compounded Drugs Obtained from an Outsourcing Facility

An outsourcing facility is permitted to provide non-patient specific compounded sterile drug products to healthcare professionals. These products are compounded under current good manufacturing practice (CGMP) requirements and the facilities are inspected by the FDA on a risk-based schedule. For more information on outsourcing facilities, including how to find those licensed by the Board of Pharmacy, visit: www.pharmacy.ohio.gov/outsourcing

The Board has confirmed with the FDA that non-patient specific drugs purchased directly from an outsourcing facility may be further prescribed and personally furnished to a patient. Please be advised that the 7-day supply limitation that applies to personally furnishing compounded drugs provided by a pharmacy (see 4729:7-2-05 (E)) does not apply to compounded drugs purchased from an outsourcing facility.

Compounded Drug Products That Are Essentially Copies of Approved Drug Products Under Section 503B of the Federal Food, Drug, and Cosmetic Act

Guidance for Industry

U.S. Department of Health and Human Services
Food and Drug Administration
Center for Drug Evaluation and Research (CDER)
Office of Compliance/OUDLC

January 2018
Compounding and Related Documents

Compounded Drug Products That Are Essentially Copies of Approved Drug Products Under Section 503B of the Federal Food, Drug, and Cosmetic Act

Guidance for Industry

Additional copies are available from:
Office of Communications
Division of Drug Information, WO51, Room 2201
Center for Drug Evaluation and Research
Food and Drug Administration
10903 New Hampshire Ave., Silver Spring, MD 20993
Phone: 301-796-3400; Fax: 301-847-8714
druginfo@fda.hhs.gov

http://www.fda.gov/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/default.htm

U.S. Department of Health and Human Services
Food and Drug Administration
Center for Drug Evaluation and Research (CDER)
Office of Compliance/OUDLC

January 2018
Compounding and Related Documents

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Guidance for Industry¹

Compounded Drug Products That Are Essentially Copies of Approved Drug Products Under Section 503B of the Federal Food, Drug, and Cosmetic Act

This guidance represents the current thinking of the Food and Drug Administration (FDA or the Agency) on this topic. It does not create any rights for any person and is not binding on FDA or the public. You can use an alternative approach if the approach satisfies the requirements of the applicable statutes and regulations. To discuss an alternative approach, contact the FDA staff responsible for this guidance as listed in the title page.

I. INTRODUCTION AND SCOPE

For a drug product compounded by an outsourcing facility to qualify for the exemptions under section 503B of the Federal Food, Drug, and Cosmetic Act (FD&C Act), it must not be "essentially a copy of one or more approved drug products," and must meet the other conditions in section 503B.³ This guidance sets forth FDA's policies concerning the *essentially a copy* provision of section 503B.⁴

In general, FDA's guidance documents do not establish legally enforceable responsibilities. Instead, guidances describe the Agency's current thinking on a topic and should be viewed only as recommendations, unless specific regulatory or statutory requirements are cited. The use of the word *should* in Agency guidances means that something is suggested or recommended, but not required.

All FDA guidances are available on the FDA guidance web page. FDA updates guidances regularly. To make sure you have the most recent version of a guidance, always consult the guidance web page at http://www.fda.gov/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/default.htm.

¹ This guidance was prepared by multiple offices in the Center for Drug Evaluation and Research, in consultation with the Office of Regulatory Affairs at the Food and Drug Administration.

² See section 503B(a)(5).

³ See section 503B(a)(11).

⁴ This guidance does not apply to drugs compounded for use in animals, to biological products subject to licensure in a biologics license application, or to repackaged drug products. For policies pertaining to mixing, diluting, and repackaging biological products, see FDA's guidance Mixing, Diluting, and Repackaging Biological Products Outside the Scope of an Approved Biologics License Application. For policies pertaining to repackaged drug products, see FDA's guidance Repackaging of Certain Human Drug Products by Pharmacies and Outsourcing Facilities.

II. BACKGROUND

A. Section 503B of the FD&C Act

In 2013, the Drug Quality and Security Act created a new section 503B of the FD&C Act, which describes a new category of compounders called *outsourcing facilities*. Section 503B of the FD&C Act describes the conditions that must be satisfied for human drug products compounded by, or under the direct supervision of, a licensed pharmacist in an outsourcing facility to qualify for exemptions from the following three sections of the FD&C Act:

- Section 502(f)(1) (concerning the labeling of drugs with adequate directions for use)
- Section 505 (concerning the approval of drugs under new drug applications (NDAs) or abbreviated new drug applications (ANDAs))
- Section 582 (concerning drug supply chain security requirements).

In contrast to drug products compounded under section 503A of the FD&C Act, drug products compounded by outsourcing facilities under section 503B cannot qualify for exemption from current good manufacturing practice (CGMP) requirements in section 501(a)(2)(B) of the FD&C Act. Outsourcing facilities are also subject to FDA inspections according to a risk-based schedule, specific adverse event reporting requirements, and other conditions that help to mitigate the risks of the drug products they compound.

One of the conditions that must be met for a compounded drug product to qualify for the exemptions under section 503B of the FD&C Act is that "the drug is not essentially a copy of one or more approved drugs." Section 503B(d)(2) defines essentially a copy of an approved drug as—

- A drug that is identical or nearly identical to an approved drug, or a marketed drug not subject to section 503(b) and not subject to approval in an application submitted under section 505, unless, in the case of an approved drug, the drug appears on the drug shortage list in effect under section 506E at the time of compounding, distribution, and dispensing (section 503B(d)(2)(A)); or
- A drug, a component of which is a bulk drug substance that is a component of an
 approved drug or a marketed drug that is not subject to section 503(b) and is not
 subject to approval in an application submitted under section 505, unless there is a
 change that produces for an individual patient a clinical difference, as determined

⁵ See Pub.L. No.113-54, § 102(a), 127 Stat. 587, 587-588 (2013). Under section 503B(b), a compounder can elect to register with FDA as an outsourcing facility. Section 503B(d)(4) defines an outsourcing facility as a facility at one geographic location or address that is engaged in the compounding of sterile drugs; has elected to register as an outsourcing facility; and complies with all of the requirements of section 503B. An outsourcing facility is not required to be a licensed pharmacy, although compounding must be done by, or under the direct supervision of, a licensed pharmacist. In addition, an outsourcing facility may or may not obtain prescriptions for identified individual patients.

⁶ See section 503B(a)(5).

by the prescribing practitioner, between the compounded drug and the comparable approved drug (section 503B(d)(2)(B)).

A compounded drug product only qualifies for the exemptions in section 503B if it is compounded by an outsourcing facility that compounds all of its drugs, both sterile and non-sterile, in accordance with all of the conditions of section 503B. A complete list of the conditions that must be met for a drug product to qualify for the exemptions in section 503B appears in the guidance For Entities Considering Whether to Register As Outsourcing Facilities Under Section 503B of the Federal Food, Drug, and Cosmetic Act.

B. Compounding, Generally

Compounded drug products serve an important role for patients whose clinical needs cannot be met by an FDA-approved drug product, such as for a patient who has an allergy and needs a medication to be made without a certain dye contained in an FDA-approved drug product, or an elderly patient or a child who cannot swallow a pill and needs a medicine in a liquid form that is not available in an approved product. Drug products for identified individual patients can be compounded by licensed pharmacists in State-licensed pharmacies and Federal facilities and by licensed physicians operating under section 503A of the FD&C Act. Drug products can also be compounded by outsourcing facilities for identified individual patients pursuant to prescriptions or for distribution to health care practitioners without receiving prescriptions. Sections 503A and 503B restrict the compounding of drug products that are essentially copies of commercially available (section 503A) or approved drug products (section 503B).

C. Compounded Drugs that are Essentially Copies of Approved Drug Products

Although compounded drugs can serve an important need, they can also pose a higher risk to patients than FDA-approved drugs. Drug products compounded by outsourcing facilities in accordance with the conditions of section 503B are exempt from FDA drug approval requirements and the requirement to be labeled with adequate directions for use. There are greater assurances of quality when drugs are compounded by outsourcing facilities that meet the conditions of section 503B and CGMP requirements than there are for drugs compounded by entities that are not required to comply with CGMP requirements and are not routinely overseen by FDA. However, as with all compounded drugs, drugs compounded by outsourcing facilities have not undergone FDA premarket review for safety, effectiveness, and quality, and lack a premarket inspection and finding of manufacturing quality that is part of the drug approval process. Because they are subject to a lower regulatory standard, compounded drugs should only be distributed to health care facilities or dispensed to patients to fulfill the needs of patients whose medical needs cannot be met by an FDA-approved drug.

⁸ Section 503A of the FD&C Act describes the conditions that must be met for a human drug product compounded by a licensed pharmacist in a State-licensed pharmacy or Federal facility, or by a licensed physician, to qualify for exemptions from sections 501(a)(2)(B), 502(f)(1), and 505 of the FD&C Act. The conditions applicable to compounders seeking to operate under section 503A are discussed in separate guidance documents applicable to these entities.

⁷ See sections 503B(a)(11) and 503B(d)(4)(A)(iii).

The restrictions on compounding drugs that are essentially copies of approved products ensure that outsourcing facilities do not compound drug products under the exemptions in section 503B for use in patients who could use an approved product. Compounding copies of these products would unnecessarily expose patients to drug products that have not been shown to be safe and effective.

In addition to these immediate public health risks, section 503B's prohibition on producing a drug product that is essentially a copy of an approved drug product protects the integrity and effectiveness of the new drug and abbreviated new drug approval processes. Sponsors would be less likely to invest in, and seek approval of, innovative, life-saving medications if an outsourcing facility could, after a drug is approved, compound "substitutes" that may be less expensive because they have not gone through the drug approval process.

Sponsors would also be less likely to seek approval of an ANDA for a generic drug if outsourcing facilities were permitted to compound drugs that are essentially copies of approved drugs without going through the ANDA process. An ANDA must include data to demonstrate that the drug has the same active ingredient and is bioequivalent to an approved drug. FDA also conducts a premarketing inspection of proposed manufacturing facilities before approving the application. Section 503B's restrictions on producing a drug product that is essentially a copy of an approved drug product protect the integrity of both the new drug and the abbreviated new drug approval processes.

D. Compounded Drugs that are Essentially Copies of Unapproved Non-Prescription Drug Products

The definition of essentially a copy of an approved drug in section 503B(d)(2) also refers to drug products that are not subject to section 503(b) (i.e., non-prescription drug products) and that are not subject to approval in an application submitted under section 505. Congress did not provide exemptions under section 503B for such drugs, which ensures that outsourcing facilities do not compound unapproved over-the-counter drug products under the exemptions in section 503B. Such products may only be produced under the requirements that apply generally to conventional drug manufacturers. Section 503B also protects FDA's over-the-counter (OTC) drug monograph process. FDA has an ongoing process to evaluate the safety and effectiveness of OTC medications, and if the Agency determines that an OTC drug meeting certain conditions is generally recognized as safe and effective, it will publish a final monograph specifying those conditions. Compounding copies of such drug products would undermine the OTC drug monograph process under which drug manufacturers must comply with the published monograph, which includes a set of specific regulatory requirements that limit the formulation of the drug product, and both the content and format of its labeling.

III. POLICY

Under section 503B(a)(5) of the FD&C Act, a compounded drug must not be essentially a copy of one or more approved drugs.9

⁹ FDA is considering the applicability of the policies described in this guidance to hospitals and health systems and

A. Definition of Essentially a Copy of an Approved Drug

The definition of essentially a copy of an approved drug has two components, specified in sections 503B(d)(2)(A) and 503B(d)(2)(B) of the Act. Section 503B(d)(2)(A) applies to a compounded drug that is "identical or nearly identical" to an approved drug or an unapproved non-prescription drug. All other compounded drugs are evaluated under section 503B(d)(2)(B). FDA applies these provisions as depicted in the diagrams in Appendices A and B.

The definition of essentially a copy of an approved drug in section 503B(d)(2) addresses both drug products approved under section 505 and marketed drug products that are not subject to section 503(b) and that are not subject to approval in an application submitted under section 505.

For purposes of this provision:

- Approved drug means a drug product that (1) is approved under section 505 of the FD&C
 Act, (2) does not appear on the list described in subsection 503B(a)(4) of drugs that have
 been withdrawn or removed from the market because such drugs or components of such
 drugs have been found to be unsafe or not effective.
- Marketed drug not subject to section 503(b) and not subject to approval in an application submitted under section 505 means any non-prescription drug product marketed without an approved application.¹⁰ We refer to these products as covered OTC drug products throughout the remainder of this guidance document.
- A drug appears on the drug shortage list in effect under section 506E if the drug is in "currently in shortage" status (and not in "resolved" status), as indicated in FDA's drug shortage database.¹¹

In addition, FDA does not intend to take action against an outsourcing facility for failing to compound in accordance with section 503B(a)(5) if it fills orders for a compounded drug that is essentially a copy of an approved drug that has been discontinued and is no longer marketed. 12

intends to address these is sues in separate guidance or rulemaking. FDA regards a health system as collection of hospitals that are owned and operated by the same entity and that share access to databases with drug order information for their patients. There is no definition of "health system" that applies to all sections of the FD&C Act. However, this is the definition of a "health system" used in section 506F of the Act concerning hospital repackaging of drugs in shortage.

¹⁰ This includes unapproved OTC drugs whether they are marketed under FDA's OTC Drug Monograph Review program or outside the monograph's ystem.

¹¹ See http://www.accessdata.fda.gov/scripts/drugshortages/default.cfm.

¹² FDA maintains a list of approved drug products that sponsors have indicated are not marketed in the discontinued section of the list of Approved Drug Products with Therapeutic Equivalence Evaluations (Orange Book). See http://www.accessdata.fda.gov/scripts/cder/ob/default.cfm. Specifically, the list includes approved drug products that have never been marketed, are for exportation, are for military use, have been discontinued from marketing and we have not determined that they were withdrawn for safety or effectiveness reasons, or have had their approvals withdrawn for reasons other than safety or effectiveness subsequent to being discontinued from marketing.

In the discussion that follows, in subsection 1, we explain how we intend to apply the definition of essentially a copy of an approved drug in section 503B(d)(2) when the compounded drug is compared to an approved drug, and then in subsection 2, we explain how we intend to apply this definition when the compounded drug is compared to a covered OTC drug product.

- 1. Application of the "Essentially a Copy" Definition in Section 503B(d)(2) When the Compounded Drug Is Compared to an Approved Drug (see Appendix A)
- a. Compounded drugs that are identical or nearly identical to an approved drug (section 503B(d)(2)(A))

Under section 503B(d)(2)(A), a compounded drug is essentially a copy of an approved drug if the compounded drug is identical or nearly identical to an approved drug unless the approved drug appears on the drug shortage list in effect under section 506E at the time of compounding, distribution, and dispensing.

i. Identical or nearly identical (Appendix A, box 1)

FDA intends to consider a compounded drug product to be identical or nearly identical to an approved drug if the compounded drug product and the FDA-approved drug have the same:

- active ingredient(s),
- route of administration, 13
- dosage form, 14
- · dosage strength, and
- excipients. 15

A compounded drug product that has all of these characteristics in common with an FDA-approved drug product is essentially a copy of an approved drug, unless the

https://www.fda.gov/Drugs/DevelopmentApprovalProcess/FormsSubmissionRequirements/ElectronicSubmissions/DataStandardsManualmonographs/ucm071667.htm. Additionally, if the approved drug can be used (regardless of how it is labeled) by the same route of administration prescribed for the compounded drug, we intend to treat the compounded drug as though it has the same route of administration for purposes of this analysis. For example, if the approved drug is an injectable drug sold in a vial that is labeled for intra-muscular use, but this drug can also be drawn from the vial by a smaller needle for subcutaneous administration, a compounded drug product sold in a similar vial and prescribed for sub-cutaneous use would be considered to have the same route of administration under this analysis.

¹³ See

¹⁴ See https://www.fda.gov/ForIndustry/DataStandards/StructuredProductLabeling/ucm162038.htm.

¹⁵ In some cases, information about the excipients contained in an approved drug is not publicly available and not known to the outsourcing facility. In such cases, FDA does not intend to consider whether the compounded drug has the same excipients that the approved drug is labeled to contain in determining whether a compounded drug is identical or nearly identical to an approved drug.

approved drug appears on FDA's drug shortage list at the time of compounding, distribution, and dispensing. If a compounded drug product is identical or nearly identical to an approved drug that is *not* on FDA's drug shortage list at the time of compounding, distribution, and dispensing, the compounded product is essentially a copy, and an outsourcing facility may not produce it under section 503B.

In establishing this policy, FDA considered the following. Under section 503B(d)(2)(A), the identical or nearly identical compounded product cannot be exempted from the copying restriction by a prescriber determination that there is a change to the compounded product that produces a clinical difference for an individual patient. Compounded products meeting the criteria outlined above are not expected to contain changes from an approved drug that would produce such a difference.

A compounded drug that is identical, or nearly identical, to an approved drug is not considered essentially a copy if the approved drug is in shortage at the time of compounding, distribution, and dispensing. ¹⁶ In such a case, the outsourcing facility can compound the drug provided that it complies with the other conditions of 503B. It is important to patients and prescribers that compounded drugs prepared to address a shortage closely resemble the drug in shortage, and for that reason, the statute seeks to allow compounders to compound drugs that are as close as possible to the drug in shortage. ¹⁷

A compounded drug product with the characteristics described in our policy would be the same as the approved drug in several important respects. The active ingredient is the substance in a drug product that is intended to furnish pharmacological activity or other direct effect in the diagnosis, cure, mitigation, treatment, or prevention of disease or to affect the structure or function of the body. Dosage form is the way of identifying the drug in its physical form, and route of administration describes the way a drug is administered to the body. Inactive ingredients (also known as "excipients") may include preservatives, dyes, and flavorings. The dosage strength of a drug product indicates the amount of the active ingredient that is present in each dosage.

If the outsourcing facility compounds a product that differs on one or more of these characteristics, we generally would not consider the product to be identical or nearly identical to the approved drug. As described below, if the compounded drug product is not considered identical or nearly identical under section 503B(d)(2)(A), it would then be evaluated under section 503B(d)(2)(B).

Outsourcing facilities seeking to compound drugs under this provision should also take note that other provisions of the FD&C Act contain requirements for drug product

¹⁶ For the purposes of this guidance, distribution means that a compounded human drug product has left the facility in which the drug was compounded. Distribution includes delivery or shipment to a physician's office, hospital, or other health care setting for administration and dispensing to an agent of a patient or to a patient for the patient's own use.

¹⁷ See footnote 11.

formulation and packaging that are important for patient safety. In particular, drug products compounded in accordance with section 503B remain subject to adulteration and misbranding provisions of the FD&C Act including, but not limited to, section 501(b) (concerning drug products that are recognized in an official compendium and whose strength differs from, or whose quality or purity falls below, the standards set forth in such compendium) and section 502(g) (concerning drug products that are recognized in an official compendium and that are not packaged and labeled as prescribed therein).

ii. Compounded drugs that are identical or nearly identical to an approved drug on FDA's drug shortage list after the shortage is resolved (Appendix A, box 2)

As explained above, under section 503B (d)(2)(A), a compounded drug is not essentially a copy of an approved drug if the approved drug appears on FDA's drug shortage list at the time of compounding, distribution, and dispensing. However, FDA recognizes that there may be circumstances in which a drug product is in shortage when the outsourcing facility compounds the drug, but the shortage is resolved before the outsourcing facility distributes it. FDA does not intend to take action against an outsourcing facility for filling orders that it received for a compounded drug that is identical, or nearly identical, to an approved drug that was on FDA's drug shortage list at the time that the outsourcing facility received the order, provided the drug also appeared on the FDA drug shortage list within 60 days of the outsourcing facility distributing or dispensing the drug. 18

b. Compounded drugs that contain a bulk drug substance that is a component of an approved drug (see Appendix A, boxes 3 and 4)

Under section 503B(d)(2)(B), a compounded drug product is essentially a copy of an approved drug if a component of the compounded drug product is a bulk drug substance ¹⁹ that is also a component of an approved drug, unless there is a change that produces, for an individual patient, a clinical difference, as determined by the prescribing practitioner, between the compounded drug and the comparable approved drug.

i. Using the same bulk drug substance (Appendix A, box 3)

¹⁸ An outsourcing facility may not be able to predict when a drug shortage will be resolved, and the facility may have orders for a compounded drug in-house that were in progress when the drug was removed from FDA's drug shortage list (e.g., the outsourcing facility may have compounded a drug while it was in shortage, but the shortage ended while the outsourcing facility awaited the results of sterility testing before release). This policy provides some regulatory flexibility when an outsourcing facility fills orders that it received for a compounded drug while the drug was in shortage. FDA may take regulatory action, however, if an outsourcing facility continues to fill new orders for the compounded drug after the approved drug is removed from FDA's drug shortage list, or if it continues to fill orders more than 60 days after the drug has been removed from FDA's drug shortage list.

¹⁹ Title 21, section 207.1 and 207.3 of the Code of Federal Regulations define the term *bulk drug substance* to mean "any substance that is intended for incorporation into a finished drug product and is intended to furnish pharmacological activity or other direct effect in the diagnosis, cure, mitigation, treatment, or prevention of disease, or to affect the structure or any function of the body. [It] does not include intermediates used in the synthesis of the substance."

If a component of the compounded drug is a bulk drug substance that is also a component of an approved drug, the compounded drug product is essentially a copy of an approved drug, and cannot be compounded under section 503B, unless there is a prescriber determination of clinical difference, as described below. ²⁰ This provision applies to a compounded drug whether it was compounded from bulk drug substances or from drugs in finished form.

ii. Prescriber determination of clinical difference (Appendix A, box 4)

If an outsourcing facility compounds a drug, the component of which is a bulk drug substance that is a component of an approved drug, there must be a change that produces a clinical difference for an individual patient as determined by the prescribing practitioner. If an outsourcing facility intends to rely on such a determination to establish that a compounded drug is not essentially a copy of an approved drug, the outsourcing facility should ensure that the determination is noted on the prescription or order (which may be a patient-specific prescription or a non-patient specific order) for the compounded drug.

FDA is aware that a health care practitioner who orders a compounded drug from an outsourcing facility for office stock will not know the identity of the individual patients who will receive the compounded drug at the time of the order. In that case, the outsourcing facility should obtain a statement from the practitioner that specifies the change between the compounded drug and the comparable approved drug and indicates that the compounded drug will be administered or dispensed only to a patient for whom the change produces a clinical difference, as determined by the prescribing practitioner for that patient. Such assurances should be provided by the health care practitioner or a person able to make the representation for the health care practitioner.

For example, a hospital may need an FDA-approved drug combined with a particular diluent in infusion bags to administer to patients during surgery, and this preparation may not be contemplated in the approved product labeling. The pharmacy manager for the hospital could order the compounded drug from an outsourcing facility and document on the order that the compounded drug will only be administered to patients for whom the prescriber determines that this formulation will produce a clinical difference from the comparable approved drug. Similarly, a physician who regularly treats patients with an allergy to an inactive ingredient in a particular approved injectable drug product could order a compounded version of the drug for office use from an outsourcing facility provided that he or she includes a statement on the order that removing the particular inactive ingredient produces a clinical difference for his or her individual patients and that he or she will provide the drug only to patients with that particular clinical need.

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²⁰ FDA expects that if a compounded drug has the same bulk drug substance as an approved drug, the two drugs have the same active ingredient.

Many outsourcing facilities compound non-sterile drugs in addition to sterile drugs.²¹ All drugs compounded by an outsourcing facility must be compounded in accordance with section 503B, including the prohibition on compounding drug products that are essentially copies of approved drug products in order for any of them to qualify for the exemptions provided in section 503B.²² For example, a hospice may need a compounded liquid formulation of a drug that is only approved in capsules to treat elderly patients who cannot swallow capsules. The pharmacy manager for the hospice could order the compounded drug from an outsourcing facility and document on the order that the liquid formulation produces a clinical difference for hospice patients who are unable to swallow capsules and that the compounded drug will be dispensed only to a patient whose prescribing practitioner determines that the liquid formulation will produce this clinical difference for the patient.

FDA does not believe that a particular format is needed, provided that an order for office stock (i.e., not patient-specific) clearly identifies the relevant change and the clinical difference that the change will produce for patient(s), as determined by the prescriber. For example, the following would be sufficient:

- "Liquid form, compounded drug will be prescribed to patients who can't swallow tablet" (if the comparable drug is a tablet)
- "Dilution for infusion solution to be administered to patients who need this
 formulation during surgery" (if the comparable drug is not available at that
 concentration, pre-mixed with the particular diluent in an infusion bag)
- "1 mg, pediatric patients need lower dose" (if the comparable drug is only available in 25 mg dose)

An order that only identifies the product formulation, without more information, would not be sufficient to establish that the determination described by section 503B(d)(2)(B) has been made.

Many outsourcing facilities also compound drug products based on prescriptions for identified individual patients. The following are examples of statements on a patient-specific prescription that could be used to document the prescriber's determination that a compounded drug has a change that produces a clinical difference for a particular patient:

- "No Dye X, patient allergy" (if the comparable drug contains the dye)
- "Liquid form, patient can't swallow tablet" (if the comparable drug is a tablet)

²¹ An entity that only compounds non-sterile drugs does not meet the statutory definition of an outsourcing facility in section 503B(d)(4) of the FD&C Act. The definition states, in part, that an outsourcing facility "is engaged in the compounding of sterile drugs" (section 503B(d)(4)(i)).

²² Under section 503B(a)(11), a compounded drug can qualify for the exemptions from section 503B only if all of the facility's compounded drugs are compounded in accordance with section 503B.

 "150 mg drug X in 120 ml cherry-flavored Syrup USP, patient needs alcohol-free preparation (if the comparable drug is only available in formulations that contain alcohol)

However, if a prescription identifies only a patient name and product formulation, this would not be sufficient to establish that the determination described by section 503B(d)(2)(B) has been made. Note also that the clinical difference identified on either a patient-specific prescription or order, or non-patient specific order, must be produced by the "change" between the outsourcing facility's product and the approved drug (i.e., a change in product formulation). Other factors such as a lower price are not sufficient to establish that the compounded product is not essentially a copy of the approved drug.

If a prescription or order does not make clear that the determination required by section 503B(d)(2)(B) has been made, the outsourcing facility may contact the prescriber or health care facility, and if the prescriber or health care facility contact confirms it, make a notation on the prescription or order that the prescriber has determined that the compounded product contains a change that produces a clinical difference for patient(s). The notations should be as specific as those described above, and should include the date of the conversation with the health care facility contact or prescriber and the name of the individual who provided the determination.²³

At this time, FDA generally does not intend to question the determinations of clinical difference that are documented in a prescription or order as described above. However, we do intend to consider whether a prescription or order relied upon by an outsourcing facility to establish that a drug is not essentially a copy documents that the determination was made.

iii. Essentially a copy of one or more approved drug products

Under section 503B(a)(5), a compounded drug product must not be essentially a copy of one or more (emphasis added) approved drug products. When applying section 503B(d)(2)(B), FDA intends to consider a compounded drug product that has bulk drug substances that are components of one or more approved drugs to be essentially a copy of an approved drug product, unless the prescribing practitioner determines that there is a change that produces a clinical difference for an individual patient between the compounded drug product and the comparable approved drug. For example, if there are two approved drug products that are tablets, one containing 5 mg of active ingredient A and the other containing 10 mg of active ingredient B and the outsourcing facility compounded a tablet that offered both active ingredients in the same dosage strengths, the compounded drug would be essentially a copy absent a prescriber determination of clinical difference.

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²³ See section IV of this guidance.

- 2. Application of the "Essentially a Copy" Definition in Section 503B(d)(2) When the Compounded Drug Is Compared to a Covered OTC Drug Product (Appendix B)
- a. Compounded drugs that are identical or nearly identical to a covered OTC drug product (section 503B(d)(2)(A)) (Appendix B, box 1)

Under section 503B(d)(2)(A), a compounded drug is not considered essentially a copy of an approved drug if it is identical or nearly identical to an approved drug that appears on FDA's drug shortage list at the time of compounding, distribution, and dispensing. The statute does not provide a similar exemption from the definition in section 503B(d)(2) if the compounded drug is identical or nearly identical to a covered OTC drug on FDA's drug shortage list. Therefore, FDA intends to apply the same policy described above in section III.A.1.a to OTC monograph drugs, with one exception.

If a compounded drug is identical or nearly identical to a covered OTC drug under section 503B(d)(2)(A), the compounded drug is essentially a copy of an approved drug, and the appearance of the covered OTC drug on FDA's shortage list does not change that result; the drug cannot be compounded under section 503B.²⁴ If the compounded drug is not identical or nearly identical to a comparable drug, it must be evaluated under section 503B(d)(2)(B), as described below.

b. Compounded drugs that contain a bulk drug substance that is a component of an covered OTC drug product (section 503B(d)(2)(B)) (Appendix B, box 2)

Under section 503B(d)(2)(B), a compounded drug product is essentially a copy and cannot be compounded under section 503B if a component of the compounded drug product is a bulk drug substance²⁵ that is also a component of a covered OTC drug, unless there is a change that produces for an individual patient a clinical difference, as determined by the prescribing practitioner, between the compounded drug and the comparable *approved* drug. A clinical difference between the compounded drug and an unapproved drug (such as a covered OTC drug) does not exempt the compounded drug from the definition in section 503B(d)(2)(B).

c. Essentially a copy of one or more approved drug products²⁶

Under section 503B(a)(5), a compounded drug product must not be essentially a copy of **one or more** approved drug products. When applying section 503B(d)(2)(B), FDA intends to consider a compounded drug product that has bulk drug substances that are components of one or more approved drugs to be essentially a copy of an approved drug product unless the prescribing practitioner determines that there is a change that produces a clinical difference for an individual patient between the compounded drug product and

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²⁴ The compounded drug would not be essentially a copy if it was also identical or nearly identical to an approved drug on FDA's drug shortage list, but this would be a very rare case.

²⁵ See footnote 19.

²⁶ This scenario is not depicted in the diagrams in the appendices.

the comparable approved drug. For example, if there are two approved drug products that are tablets, one containing active ingredient A and the other containing active ingredient B, and the outsourcing facility compounded a tablet that offered both active ingredients, the compounded drug containing active ingredients A and B would be essentially a copy absent a prescriber determination of clinical difference.

If a bulk drug substance is a component of a covered OTC drug and an approved drug, the bulk drug substance can be evaluated as a component of an approved drug, as described in section III.A.1 of this guidance.

B. Recordkeeping

Outsourcing facilities should maintain records to demonstrate compliance with the essentially a copy provision in section 503B(a)(5). For example, where an outsourcing facility has compounded a drug that is evaluated under 503B(d)(2)(B) and a component of the compounded drug is a bulk drug substance that is a component of an approved drug, the outsourcing facility should maintain prescription or order records of a prescriber's determination of clinical difference as described above in section III.A.1.b.ii.

In addition, if the outsourcing facility compounded a drug that is identical or nearly identical to an approved drug product that appeared on FDA's drug shortage list, the outsourcing facility should maintain documentation (e.g., a notation on the order for the compounded drug) regarding the status of the drug on FDA's drug shortage list at the time of compounding, distribution, and dispensing.²⁷

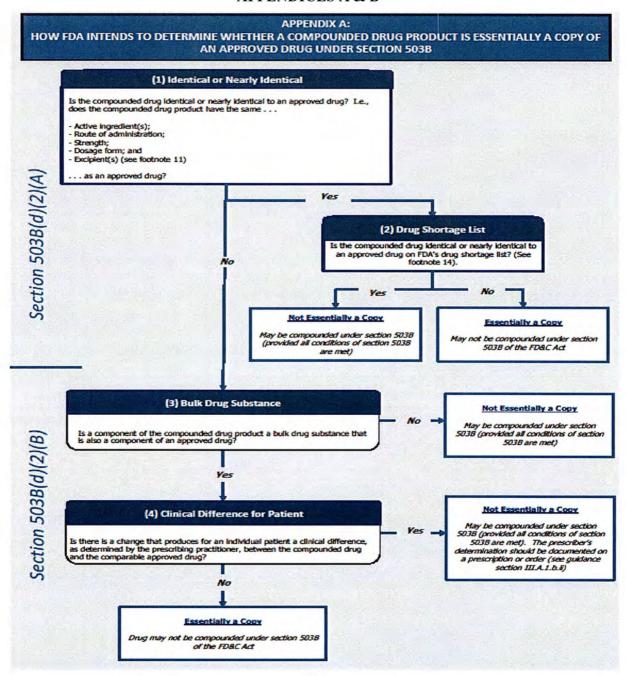
FDA recommends that outsourcing facilities maintain the records described above for a period of at least three years.

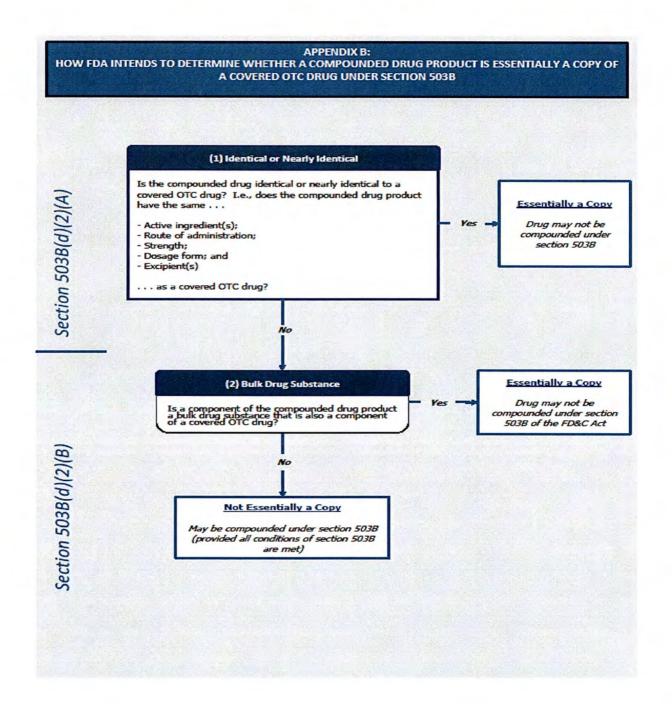
IV. PAPERWORK REDUCTION ACT

This guidance contains information collection provisions that are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501-3520). See footnotes 23 and 27. These provisions require review and are not in effect until they display a currently valid OMB control number. The information collection provisions in this guidance have been submitted to OMB for review as required by section 3507(d) of the Paperwork Reduction Act of 1995. FDA will publish a notice in the Federal Register announcing OMB's decision regarding the information collection provisions in this guidance.

²⁷ See section IV of this guidance.

APPENDICES A & B





DECLARATION OF MA LOURDES JAUREGUI, RPH

I, MA LOURDES JAUREGUI, declares under penalty of perjury that the following assertions are true and correct of my own personal knowledge unless stated upon information and belief:

- 1. I was the managing pharmacist of Trinity Pharmacy ("Trinity") located at 2797

 S. Maryland Parkway, Suite 28, Las Vegas, Nevada. I am writing this declaration because I am unavailable for the scheduled hearing for alleged violations against Stephen Dubin, MD ("Dubin") regarding the Nevada State Board of Pharmacy's ("Board") case number 20-020.
- 2. I have personal knowledge of all matters stated herein and, if called as a witness, could and would testify competently thereto, except as to those matters stated upon information and belief, and as to those matters, I believe them to be true.
- 3. On or about September 10, 2020, the Board's investigator, Dena McClish, requested certain prescription records from Trinity.
- 4. On or about September 10, 2020, I forwarded the requested prescription records, which included the following:
 - a) True and correct copies of Trinity's (redacted) prescription records for patient A.D. on December 18, 2018 are attached hereto as Exhibit "1".
 However, I was not the pharmacist who filled this particular prescription. It was filled by a floater pharmacist.
 - b) True and correct copies of Trinity's (redacted) prescription records for patient M.G. on December 19, 2018 are attached hereto as Exhibit "2".
 - c) True and correct copies of Trinity's (redacted) prescription records for patient M.H. on December 19, 2018 are attached hereto as Exhibit "3".

- d) True and correct copies of Trinity's (redacted) prescription records for patient D.B. on December 20, 2018 are attached hereto as Exhibit "4".
- 5. Exhibit "1" contains a prescription for patient A.D. which was telephoned by "Honey" on behalf of Dr. Dubin. "Honey" indicated that the prescription was authorized by Dr. Dubin. I did not speak with Dr. Dubin himself. The prescription for patient A.D. was taken by the floater pharmacist on December 18, 2018 for alprazolam 1 mg tablet. The alprazolam was dispensed on December 21, 2018 at 10:54 a.m. Dr. Dubin never signed this prescription.
- 6. Exhibit "2" contains a prescription for patient M.G. which was telephoned by "Speedy" at 4:00 p.m. on December 19, 2018 on behalf of Dr. Dubin. "Speedy" indicated that the prescription was authorized by Dr. Dubin. I did-not speak with Dr. Dubin himself. The prescription for patient M.G. was written in my handwriting for Percocet 10/325 mg, 60 tablets for a 15 day supply. The prescription was dispensed on December 21, 2018 at 10:55 a.m. On the prescription signature information, the patient or the guardian signed for receipt of the prescription. Dr. Dubin signed this prescription within 72 hours of December 19, 2018, but is undated.
- 7. Exhibit "3" contains a prescription for patient M.H. which was telephoned by "Speedy" at 1:00 p.m. on December 19, 2018. "Speedy" indicated that the prescription was authorized by Dr. Dubin. I did not speak with Dr. Dubin himself.

 The prescription for patient M.H. was for lorazepam 0.5 mg tablet, 45 tablets. The prescription was dispensed on December 21, 2018 at 10:55 a.m. On the prescription signature information, the patient or the guardian signed for receipt of the prescription. Dr. Dubin never signed this prescription.

- 8. Exhibit "4" contains a prescription for patient D.B. which was telephoned by "Honey" at 4:00 p.m. on December 20, 2018. "Honey" indicated that the prescription was authorized by Dr. Dubin. I did not speak with Dr. Dubin himself. The prescription for patient D.B. was for morphine sulfate OS 20mg/ml, quantity 30. The prescription was dispensed on December 20, 2018 at 5:58 p.m. On the prescription signature information, the patient or the guardian signed for receipt of the prescription. Dr. Dubin signed this prescription within 72 hours of December 20, 2018, but is undated.
- 9. Exhibit "4" also contains a prescription for patient D.B. which was telephoned by "Honey" at 4:00 p.m. on December 20, 2018. "Honey" indicated that the prescription was authorized by Dr. Dubin. I did not speak with Dr. Dubin himself. The prescription for patient D.B. was for lorazepam O/C, quantity 30. The prescription was dispensed on December 20, 2018 at 5:58 p.m. Dr. Dubin never signed this prescription.
- 10. My understanding was that the prescriptions for hospice patients from Jireh Healthcare were all considered emergency oral prescriptions. However, I did not notate on the telephoned or oral prescriptions, "Authorization for Emergency Dispensing" nor did I speak with Dr. Dubin directly. I understood that I did have to receive a signed prescription back from the practitioner within 72 hours after the telephoned prescription, which admittedly was not always accomplished.

Further, Your Declarant Sayeth Naught.

Dated: 11/18/202/

Lourdes Jauregui, RPH

JIREH HEALTHCARE SERVICES LLC (HOSPICE AGENCY)

PATIENT NAME: A

DOB:

START OF CARE: 09/07/2018 TO 12/31/2018

Claim Number	From Strvice D	ite. To Service Da	From Strying Date. In Service Date. Beneficially Nem Blaim Status	Total Charge	Type Or Bill Admir Date
219007004Z2207NVH	12/1/2018	12/91/2018	A P-Paid	87,790.97	813 - Hospice (9/7/2018
21833900538807NVR	11/1/2018	11/30/2018	And De P - Paid	\$8,724.43	B13 - Hospice (_9///2018
21830901126607NVR	10/1/2018	10/31/2018	A PuPaid	\$9,083,84	813Hospice (9/7/2018
21827601222707NVR	9/7/2018	9/30/2018	A D P-Paid	\$6,942.12	812 - Hospice (_ 9/7/2018
dain Number	From Service Date	Service Co.	To Service Date Temperary Nampham Status	Total Charge	Type of this Admit Date
21900700421907NVR	12/172018	12/31/2018	P-Papp	\$6,986.61	813-Ноѕрке (_10/10/2018
21833900542507NVR	11/1/2018	11/30/2018	P-Paid	\$8,197.48	813 - Hospice (_ 10/10/2018
21833900230804NVB	10/10/2018	10/31/2018	P-Paid	\$6,230.74	817-Hospice (10/10/2018
21830901121107NVR	10/10/2018	10/31/2018	A. P-Paid	\$5,630.74	812 - Hospice (± 10/10/2018
Claim Nomber	From Service Bate		To Sovice Date Beneficiary Non Elaim Status	Total Charge	Type OF BUIL Behalt Gare
219809811188507NVH	12/1/2018	12/26/2018	NE_P-Paid	\$6,626 68	814-Hospice (6/13/2018
21833900542007NVR	11/1/2018	11/30/2018	P-Paid	\$7,635.75	813 - Hospice (6/13/2018
21830901128607NVR	10/1/2018	10/31/2018	Padd Padd	\$8,065.97	813 - Hospice (_ 6/13/2018
21827601223807NVR	9/1/2018	9/30/2018	Pard - Pard	\$7,382.91	813 - Hospice (6/13/2018
21824701185107NVB	8/1/2018	8/31/2018	P-Paid	58,276.79	813 - Hospice (_6/13/2018
21821800581907NVR	7/1/2018	7/31/2018	P-Paid	\$9,216.50	813 - Ноѕрксе (_6/13/2018
21818500817807NVB	6/13/2018	6/30/2018:	P-Paid	\$5,126.72	812 Hospice (6/13/2018
Clarin Number	Hom Service Date	ite To Service Date	To Service Date Henefferary Nam Claim Status	क्टाबाधाया ।	Type Of Bill Admir Date
21900700421597NVR	12/20/2018	12/22/2018	D. P-Paid	3580.54	811 - Hospice (_ 12/20/2018

12/31/2018 Jireh Healthcar M0466	G30.1	Clark	ROUTINE	Medicare	'	178.2523	0
12/30/2018 Jireh Healthcar M0466	G30.1	Clark	ROUTINE	Medicare	Þ	178.2523	0
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12/27/2018 Jireh Healthcar M0466	G30.1	Clark	ROUTINE	Medicare		178.2523	0
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12/25/2018 Jireh Healthcar M0466	G30.1	Clark	ROUTINE	Medicare		178.2523	0
12/24/2018 Jireh Healthcar M0466	G30.1	Clark	ROUTINE	Medicare '		178.2523	0
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12/15/2018 Jireh Healthcar M0466	G30.1	Clark	ROUTINE	Medicare		178.2523	0
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12/7/2018 Jireh Healthcar M0466	G30.1	Clark	ROUTINE	Medicare		178.2523	0
12/6/2018 Jireh Healthcar M0466	G30.1	Clark	ROUTINE	Medicare		178.2523	0
12/5/2018 Jireh Healthcar M0466	G30.1	Clark	ROUTINE	Medicare		178.2523	0
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Jireh Healthcare Serrices LLC

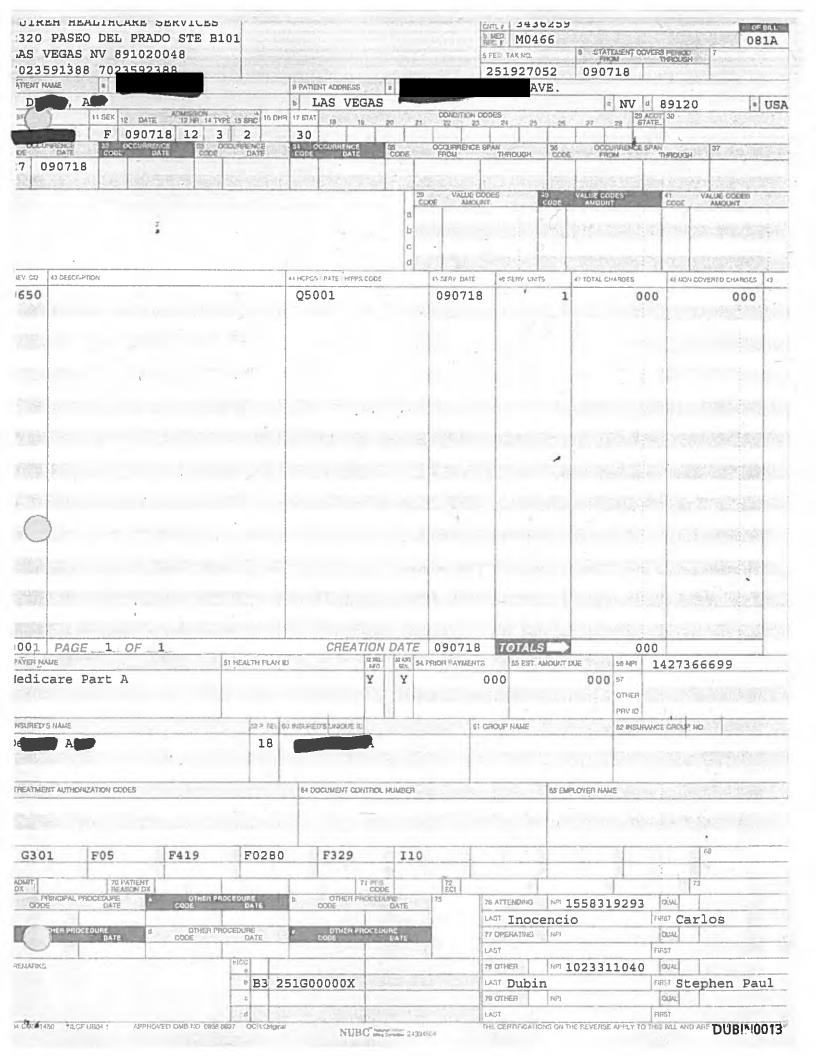


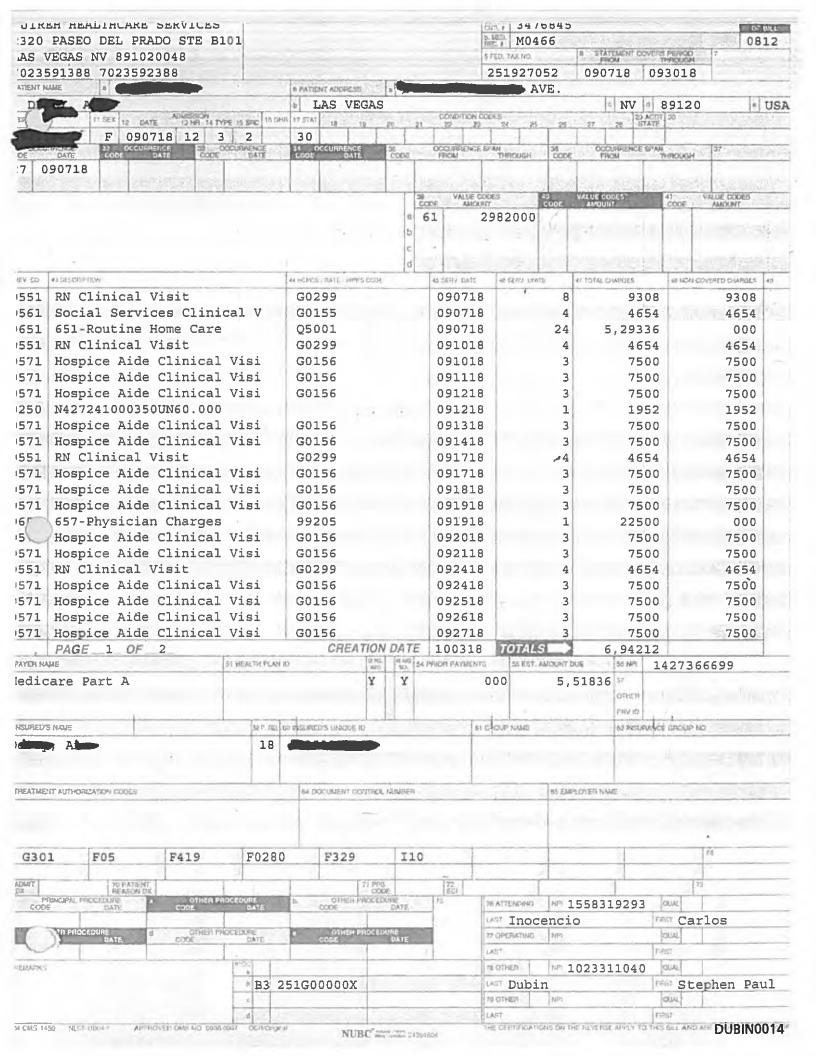
2320 Pased	Del Pra	do, Suite	B101, Las Veg	as, NV 89	9102 . Tel:	Fax:				¥				
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STARTING	CERT#	1												
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REFERRING CHOICE	3/ATTEN	DING/PA	ATIENT	CARLOS	INOCEN	CIO, Phone	9:(702	2) 733-0744,	rax:(702) 796-8	3262	2, NPI:155831929	13		
								-	· · · · · · · · · · · · · · · · ·				'	
Vendors														
PERMACY	/			MAIN:Tri	nity Pharm	nacy (Las V	/eas)						

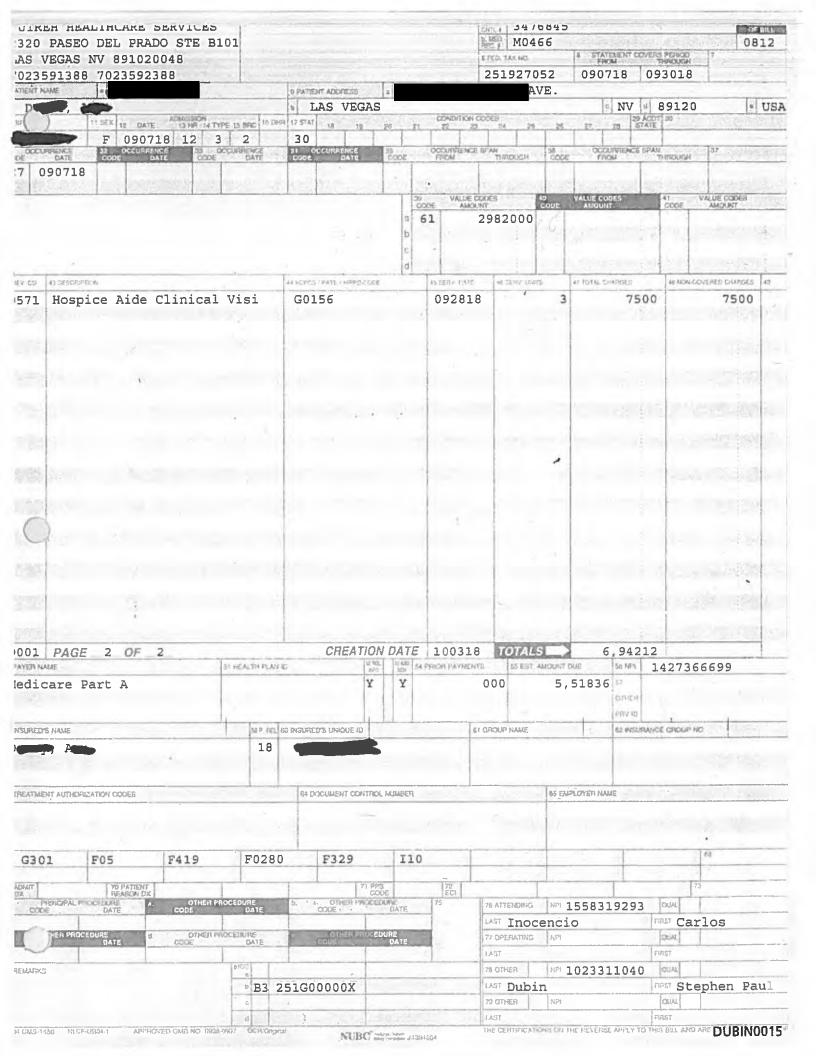
CONFIDENTIALITY NOTICE. The information contained in this facsimile is confidential and may contain privileged material or be otherwise protected by applicable law. It is intended only for the use of the individual (s) or entity named above. If the person receiving this facsimile, or any other reader of the facsimile, is not the intended recipient or the employee or agent responsible for delivering it to the intended recipient, then any use, dissemination, distribution, or copying of this communication is strictly prohibited and may be subject to civil and/or criminal liability. If you have received this communication in error, please destroy it and notify the sender immediately

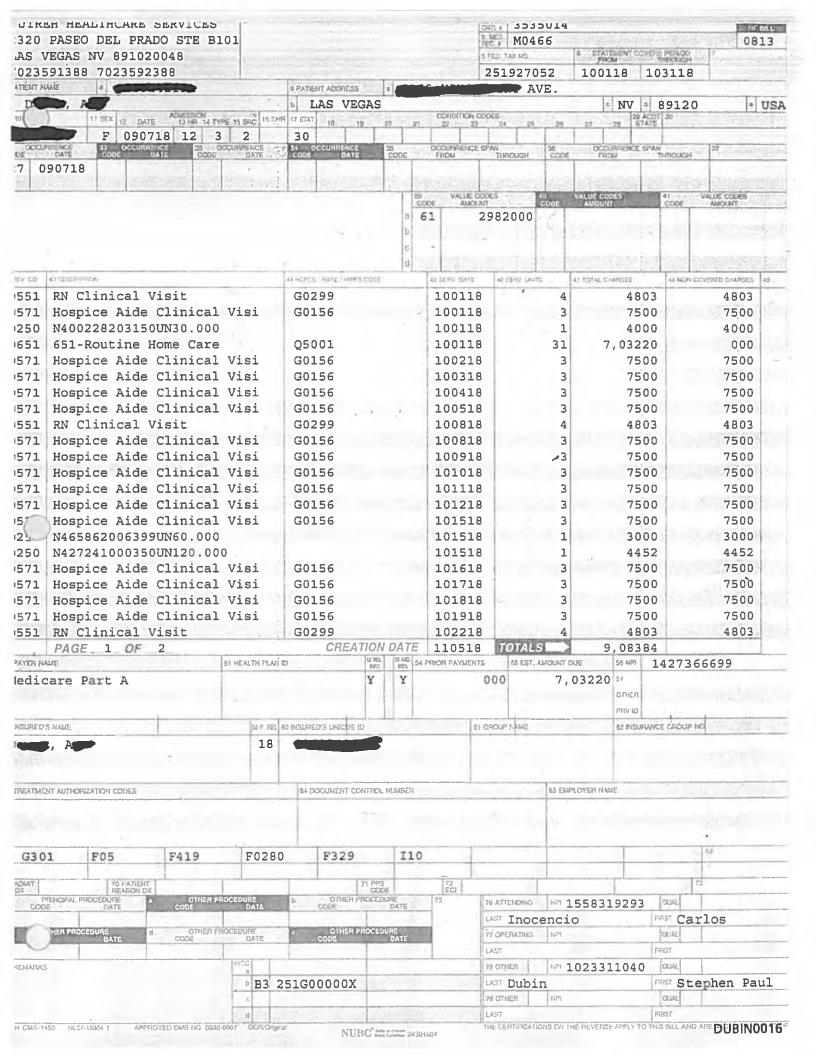
MAIN:Advantage Home Medical Services, Inc. (Las Vegas)



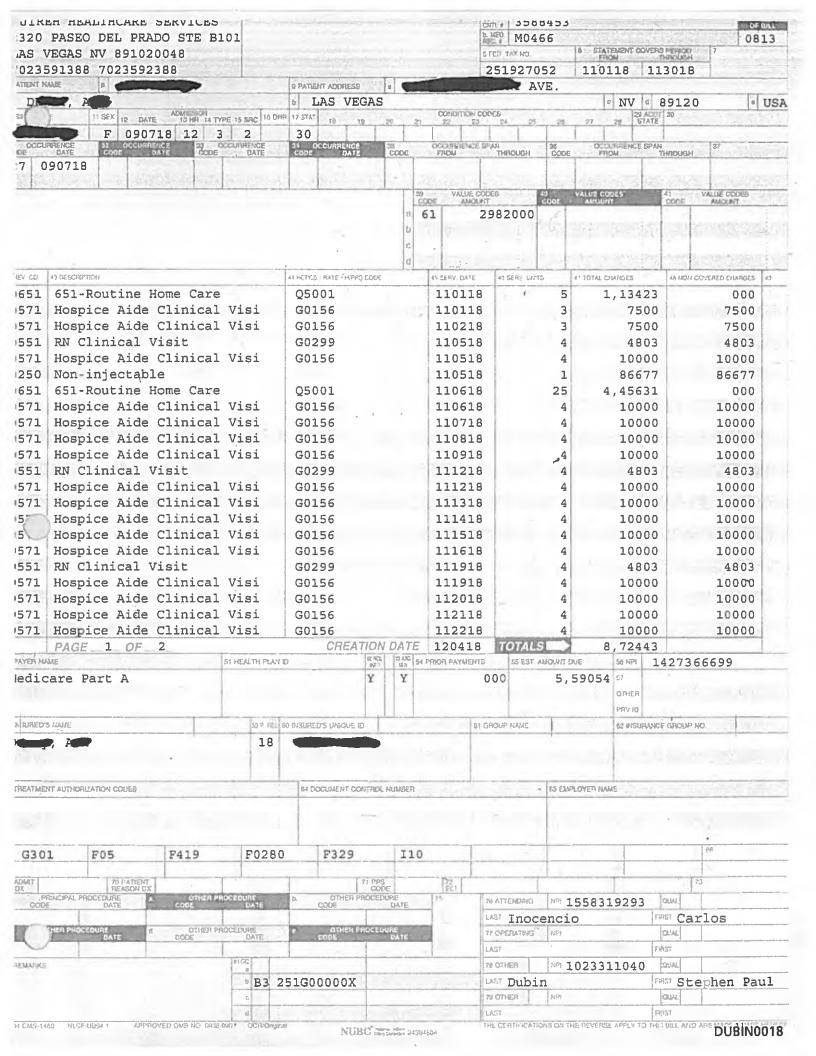


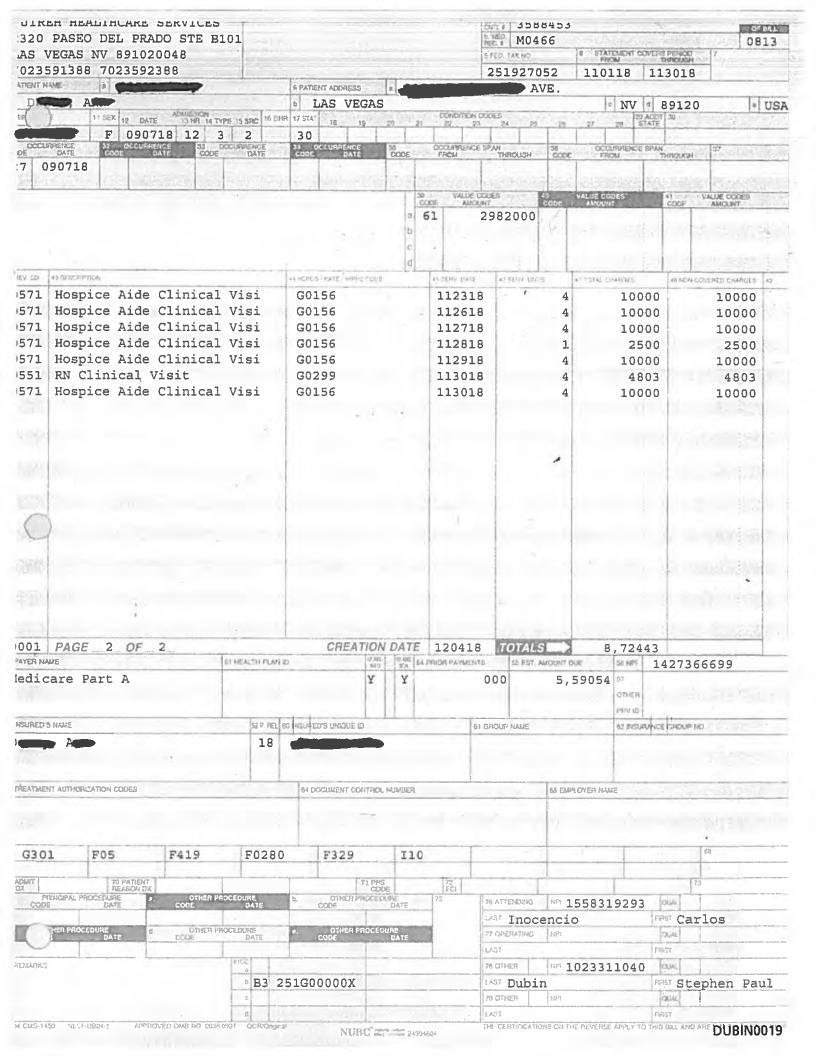


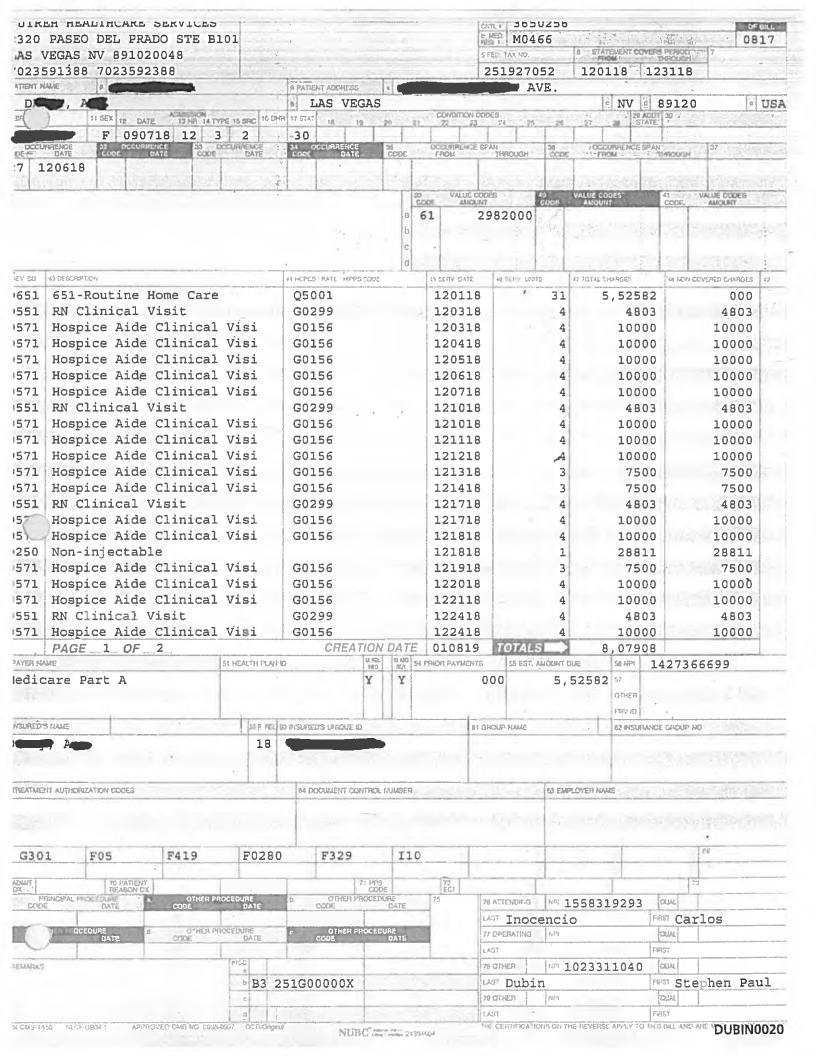




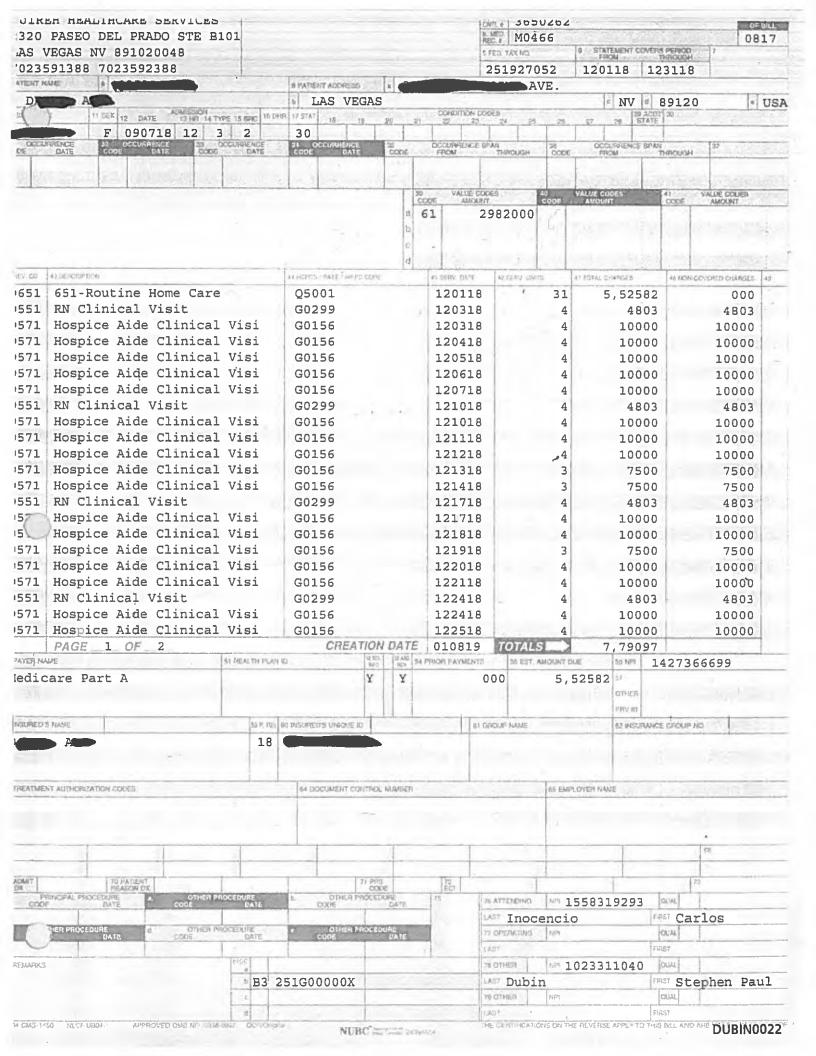
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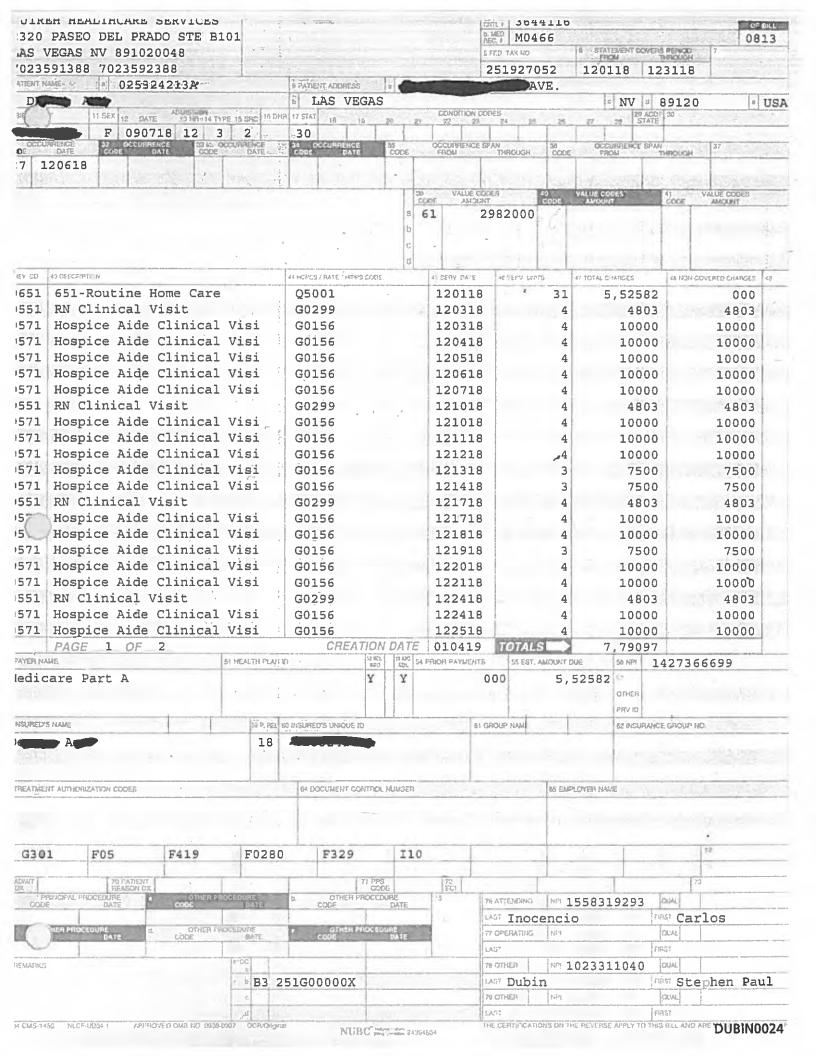




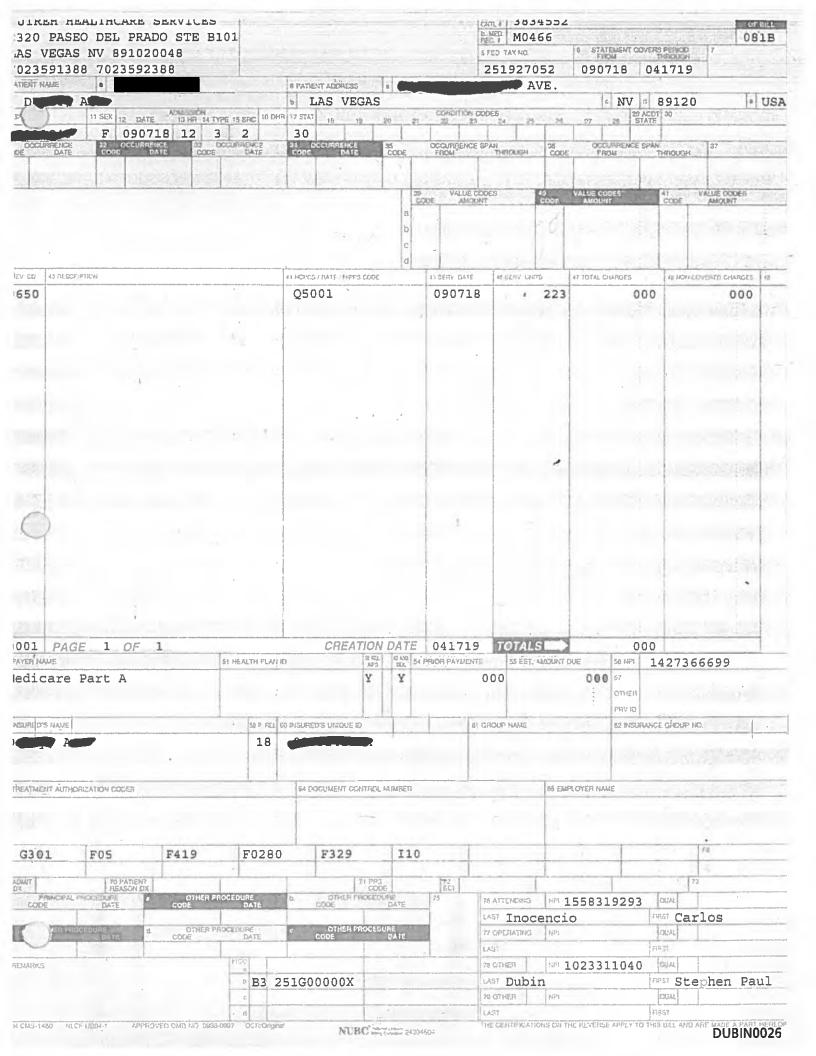
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Consolo Assignment Dated: 09/07/2018 for Date:

Patient Details for Deal, American ID 491783, MRN M0466)

Date Of Birth Sta

Status Pending - Discharged

Assignment Details

Effective Date: 09/07/2018, Medical Record No: M0466, Dme Provider: Advantage Home Medical Services, Inc., Pharmacy: Trinity Pharmacy, Staff (caregiver)

IDG Team

Discipline	Name	Primary?	Default for New Medications?	Visits	PRN Visits	Frequency Notes
Chaplain	CHAPLAIN_Murphy, Kathryn	Yes	No			Evaluation (frequency of visits to be determined post evaluation
Hospice Alde	CNA_Barba Magdalena	No	No	5 visit(s) Weekly		
MD Director	Dubin, Stephen Paul	Yes	No		1 PRN visit(s) every 30 day(s)	1x/mo PRN withn certification period for Emergency Response
Physician	Inocencio, Carlos	Yes	No		1 PRN visit(s) every 30 day(s)	Eval,1x/mo PRN withn certification period for Emergency Response
Skilled Nurse	RN_Cacanindin, Speedy	Yes	No	t visit(s) Weekly	2 PRN visit(s) every 7 day(s)	1x/wk &2x/wk PRN for fall & anxiety issues withn cert perd
Skilled Nurse	RN_Ruiz, Janyvill	No	No		2 PRN visit(s) every 7 day(s)	Eval &2x/wk PRN for fall & anxiety issues within cert perd

Consolo Assignment Dated: 09/10/2018 for Dam, A

Patient Details for Demand (Patient ID 491783, MRN M0466)

Date Of Birth

Status Pending - Discharged

Assignment Details

Effective Date: 09/10/2018, Medical Record No: M0466, Drne Provider: Advantage Home Medical Services, Inc. , Pharmacy: Trinity Pharmacy, Staff (caregiver)

IDG Team

			Default for			
			New			1
Discipline	Name	Primary?	Medications?	Visits	PRN Visits	Frequency Notes
Chaplain	CHAPLAIN_Murphy,	Yes	No			Evaluation (frequency of visits to
	Kathryn		50			be determined post evaluation
Hospice	CNA_Barba	Yes	No	5		
Aide	Magdalena			visit(s)		
				Weekly		
MD	Dubin, Stephen Paul	Yes	Na		1 PRN visit(s)	1x/mo PRN within certification
Director					every 30	period for Emergency Response
					day(s)	
Physician	Inocencio, Carlos	Yes	No		1 PRN visit(s)	Eval,1x/mo PRN withn certification
					every 30	period for Emergency Response
					day(s)	
Skilled	RN_Ruiz, Janyvill	Yes	No		2 PRN visit(s)	Eval 82x/wk PRN for fall 8 anxiety
Nurse					every 7 day(s)	issues within cert perd
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Skilled	RN_Cacanindin,	No	No	1	2 PRN visit(s)	1x/wk &2x/wk PRN for fall &
Nurse	Speedy			visit(s)	every 7 day(s)	anxiety issues within cert perd
				Weekly		

Consolo Assignment Dated: 09/12/2018 for Date, American Dated: 09/12/2018 for Dated: 09/

Patient Details for Despendent ID 491783, MRN M0466)

Date Of Birth

Status Pending - Discharged

Assignment Details

Effective Date: 09/12/2018, Medical Record No: M0466, Dme Provider: Advantage Home Medical Services, Inc. , Pharmacy: Trinity Pharmacy, Staff (caregiver)

IDG Team

Discipline	Name	Primary?	Default for New Medications?	Visits	PRN Visits	Frequency Notes
Chaplain	CHAPLAIN_Murphy, Kathryn	Yes	No	1 visit(s) Monthly	3 PRN visit(s) every 30 day(s)	Evaluation, 1x/mo, and 3x/mo PRN for spiritual issues within cer
Hospice Aide	CNA_Barba, Magdalena	Yes	No	5 visit(s) Weekly		
MD Director	Dubin, Stephen Paul	Yes	No .		1 PRN visit(s) every 30 day(s)	1x/mo PRN withn certilication period for Emergency Response
Physician	Inocencio, Carlos	Yes	No		1 PRN visit(s) every 30 day(s)	Eval,1x/mo PRN withn certification period for Emergency Response
Skilled Nurse	RN_Ruiz, Janyvill	No	No		2 PRN visit(s) every 7 day(s)	Eval &2x/wk PRN for fall & anxiety issues within cert perd
Skilled Nurse	RN_Cacanindin, Speedy	Yes	No	1 visil(s) Weekly	2 PRN visit(s) every 7 day(s)	1x/wk &2x/wk PRN for fall & anxiety issues withn cert perd

MD ORDERS, TREATMENT, MEDICATION & CTI

Xanax 1 MG Oral Tablet - Alprazu um Oral Tablet 1 MG Take 1 tablet (1 mg) by mouth every night as needed for anxiety

N/A -

Galantamine Hydrobromide ER 24 MG Oral Capsule Extended Release 24 Hour - Galantamine Hydrobromide Oral Capsule Extended Release 24 Hour 24 MG

Take 1 capsule (24 mg) by mouth daily in the morning with food

N/A

Twice to trace

Tihere is no allergy, history recorded for this patient

Encounter - 09/05/2011	8:		-5
SEEN BY		SEEN ON	
Anthony Denina		09/05/2018	
HEIGHT	WEIGHT	BMI	BLOOD PRESSURE
63:0:ln	18/1.4/lbs	32.1	153/82
TEMP	PULSE	RESP RATE	HEAD CIRC
97.9 F	64:0)bpm	17.0 rpm	- N/A
CC			

89120-Confirmed eval for Hospice St Anne Care Home

3:158.Kingspoint Ave 89:129 (Appt time: 7:00 AM) (Arrival time: 12:41 PM)

S

Pt was referred to ARdha Mobile Minfor a routine follow-up. Pt's pcg and grouphome owner stated that she spoke with pt's son Team who is pt's POA and is contemplating on placing pt under hospice care.

0

HPI: Pt is a 75 y/o female, abox1, able to recall some long term memory but has very poor short term memory, due to worsening alzheimers. She is ambulatory and is denying any discomfort at this time. Per pcg Susana, pt has episodes of sundowning and is getting very confused and agitated at night times with episodes of wandering and is an elopement risk. She is requesting for a hospice eval per her conversation with pt's son Todd: Explained that I can order for an eval or screening but the hospice medical director will have to take over pt's care once pt's condition is deemed appropriate for hospice.

Pt is homebound due to the following reasons:

- > Requires considerable taxing effort to safely leave home.
- > Unable to safely leave home unassisted
- > Dependent upon adaptive device-
- > Requires assistance on activities.

Advance Directives / Advance care planning.

Patient has already executed:an Advance Directive: Yes-

If no, was patient given an opportunity, to execute; an Advance Directive today? ... n/a

Physician statement: "Patient has the ability to prepare an Advance Directive.".. Yes ?Rio X.

Physician has completed a Physician Order of Life-Sustaining Treatment, or similar document of another name, reflecting the patient's wishes. Yes ?No.X.

Physician is willing to follow the patient's wishes. Yes

A:

P:E:

V.S. as recorded:

General appearance: Well-kept, A&O.x. 1, pleasant with conversation, answers questions appropriately HEENT: Head is normocephalic, temporal arteries soft; non-tender with no bruits. Eyes appears to be healthy/no redness/ tearing. Right & left ear appears to be clean, no exudates or discharges noted. No thyromegaly. Neck is supple/no jugular venous distension. Dentition appears unremarkable, denies dental/ oral discomfort: Nose is symmetrical. Oral mucosa is moist: Respiratory: Respiration's are unlabored. Symmetrical chest expansion noted. Lungs sounds are clear. No cyanosis or clubbing of the fingers noted. Cardiovascular: Rhythm and rate are regular. Normal S-1& S-2 heard, no murmurs noted Musculoskeletal: ambulatory with a steady gait; both upper and lower extremitles are normal in strength and mobility, No swelling or deformities noted. Gastrointestinal: Abdomen soft with normal bowel sounds, stated normal bowel movement: Genitourinary: continent; denies dysuria, denies frequency. Lymph nodes; No palpable cervical nodes. Integumentary: Skin is pink, and warm. No acne/ scarring or dyspigmentation noted. Neurological: A.A.O.X.1 with very poor short term memory. Psychlatric: Pt is cooperative: Answers to questions appropriately denies suicidal/homicidal thoughts.

P

> Continue all meds

> Refer to business for eval once accepted, transfer care to hospice's medical director as pt's pcp, pcg is agreeable.

Provided an in-depth discussion of disease process and potential compilications.

Safety, measures discussed; fall precautions discussed, evaluated home/environment for safety issues.



Response of Prado, Suite B101, Las Vegas, NV 89102
Phone: 702.359.1388 Fax: 702.359.2388

	PHYS	SICIAN'S ORDER		
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ORDER READ BACK	AND VERIFIED BY:			
Signature/Discipline				
Staff Receiving Orde	er: Qrikub	Date Received:	9/10/W	
Physician Signature:		Date Signed:	9/11/	



2320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102 Phone: 702.359.1388 Fax: 702.359.2388

PHYSICIAN'S ORDER

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C. A)	40466	NKOY	Dr. Studen ?	John M
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Signature/D		1		e151	
Staff Receivi	ng Order:	C. Kuf	Date Received:	9/07/2017	
Physician Sig	nature:	12	Date Signed:	9/07/2017	

Consolo Physician for DEWxA at Office Jireh Healthcare Services LLC

Patient Details for Desay, A (Patient ID 491783, MRN M0466)

Date Of Birth	Status	Active
Screened Allergies No Known Drug Allergy, No Known Drug Allergy	Unscreened Allergies	See Screened Allergies

Physician's Order Details

<u>User</u>

RN_Ruiz, Janyvill (jruiz)

Patient

Order Date 09/07/2018 Time of Event

0730

<u>Physician</u>

Dubin, Stephen Paul

Nurse

RN_Ruiz, Janyvill (jruiz)

Related Yes Oversight

Orders

Admit to Jireh Healthcare Services, Ltc. for the 1st 90 day benefit period.* Pharmacist may use generic drug equivalents or compound medications as appropriate. * Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to: hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction machine, etc.) * Assess and evaluate cardiac, respiratory, GI,GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs.* Instruction on care of terminally ill, medications, safety, nutritional needs, pain and symptom control based on assessment. * Assess and evaluate current medications.* May give nutritional supplements 1 can a day as tolerated for supplement. May crush medications that are crush-able. * Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders. * Hospice Medical Director may assist patient as needed to manage symptoms. * Hospice nurse may pronounce death. * Check oxygen saturation level printers are supplement. * Cityweek and 2x/week as needed for fall. & anxiety issues) to promote comfort and symptom management within certification period. * CHHA visits 5x/week to assist with personal care, home making and to promote comfort per RN assignment within certification period. * Chaplain visits. (Eval. & 1x/mo as needed for spiritual issues) *Social worker visits. (Eval. & 1x/mo as needed for psychosocial issues) Volunteer Services continue to be offered every week per Skilled Nurse/Staff, Admit Patient to Jireh Healthcare Services with the following orders verified with M.D. * Patient: DNR * Diet: As Tolerated.* Admit with the following medications: Please see related medications:

Read Back

Non Verbal Order

Yes

No

Medications

Medication	Instructions	Quantity	Refilis
Losartan Potassium 1 tab 1 times a day PO (100 MG Tablet)			
ALPRAZolam 1 tab 1 times a day PO (1 MG Tablet)	to be taken every night at bedtime.	,	
Metoproloi Tartrate 1 tab 2 times a day PO (50 MG Tablet)			
Galantamine Hydrobromide ER 1 tab 1 times a day PO (24 MG Capsule ER 24HR)			36
Escitalopram Oxalate 0.5 tab 1 times a day PO (10 MG Tablet)			
Namenda XR 1 tab 1 times a day PO (28 MG Capsule ER 24HR)	R		A.

Related Assignees

Skilled Nurse - RN_Ruiz, Janyvill 2 PRN visit(s) every 7 day(s) effective 09/07/2018

Skilled Nurse - RN_Cacanindin, Speedy 1 visit(s) Weekly 2 PRN visit(s) every 7 day(s) effective 09/07/2018

Hospice Aide - CNA_Barba, Magdalena 5 visit(s) Weekly effective 09/07/2018

Chaplain - CHAPLAIN_Murphy, Kathryn effective 09/07/2018

Physician - Inocencio, Carlos 1 PRN visit(s) every 30 day(s) effective 09/07/2018

MD Director - Dubin, Stephen Paul 1 PRN visit(s) every 30 day(s) effective 09/07/2018

Consolo Physician for DEWxA at Office Jireh Healthcare Services LLC

Signatures ·

- 1. Medical Director Dubln, Stephen Paul (Medical Director) signed on 09/07/2018. Recorded by jruiz on 09/10/2018 11:47:02.
- 2. Skilled Nurse RN_Ruiz, Janyvill (User) signed on 09/10/2018. Recorded by jruiz on 09/10/2018 11:46:50.

Consolo Physician Order (2397060) Dated: 12/03/2018 for Dubin, Stephen Paul for D

Patient Details for Date 7, April Patient ID 491783, MRN M0466)

Date Of Birth Status Pending - Discharged

Physician's Order Details

<u>User</u>

Dugay, Julia (jduga)

Patient

Order Date 12/03/2018 Time of Event

Physician

Dubin, Stephen Paul

Nurse Dugay, Julia (jduga)

) <u>(</u>

Overslaht No

Orders

Re-certify to Jireh Healthcare Services for the 2nd 90 day benefit period * Pharmacist may use generic drug equivalents or compound medications as appropriate. Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to; hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction machine, etc.) * Assess and evaluate cardiac, respiratory, GI, GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs * Instructions on care of terminally ill, medications, safety, nutritional needs, pain and symptom control based on assessment * Review and reconcile current medications * May give nutritional supplements 1 can a day as tolerated for supplement * May crush medications that are crush-able. * Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders * Hospice Medical Director may assist patient as needed to manage symptoms * Hospice nurse may pronounce death * Oxygen 3.0 lpm via NC continuously as needed for SOB * Check oxygen saturation level pm every SN visit * Bed-rails up 2x * Skilled Nurse visits 1x/week and 2x/week as needed for pain, SOB, fall & anxiety Issues to promote comfort and symptom management within certification period. * CHHA visits 5x/week to assist with personal care, home making and to promote comfort per RN assignment within certification period. * Chaptain visits -1x/mo, 1x/mo as needed for spiritual issues. * Social worker visits - 1x/mo as needed for psychosocial issues. * Volunteer Services continue to be offered every week per Skilled Nurse/Staff. * Admit patient to Jireh Healthcare Services with the following orders verified with M.D: Patient: DNR Diet: As Tolerated * Admit with the following medications: Please see related medications.

Read Back

Non Verbal Order

Yes

No

Related Assignees

Chaplain - CHAPLAIN_Murphy, Kathryn 1 visit(s) Monthly 3 PRN visit(s) every 30 day(s) effective 09/12/2018 **
Hospice Aide - CNA_Barba, Magdalena 5 visit(s) Weekly effective 09/12/2018
MD Director - Dubin, Stephen Paul 1 PRN visit(s) every 30 day(s) effective 09/12/2018
Physician - Inocencio, Carlos 1 PRN visit(s) every 30 day(s) effective 09/12/2018
Skilled Nurse - RN_Cacanindin, Speedy 1 visit(s) Weekly 2 PRN visit(s) every 7 day(s) effective 09/12/2018
Skilled Nurse - RN_Ruiz, Janyvill 2 PRN visit(s) every 7 day(s) effective 09/12/2018

Signatures

1. Medical Director Dubln, Stephen Paul (Medical Director) signed on 12/03/2018. Recorded by jduga on 12/03/2018 15:13:16.

Consolo Physician for DEWxA at Office Jireh Healthcare Services LLC

Patient Details for December (Patient ID 491783, MRN M0466)

Date Of Birth	Status	Active
Screened Allergies No Known Drug Allergy, No Known Drug Allergy	Unscreened Allergies	See Screened Allergies

Physician's Order Details

User

Dugay, Julia (jduga)

Patient

Order Date 12/03/2018 Time of Event

Physician

Dubin, Stephen Paul

<u>Nurse</u>

Dugay, Julia (jduga)

Oversight

No

Orders

Re-certify to Jireh Healthcare Services for the 2nd 90 day benefit period * Pharmacist may use generic drug equivalents or compound medications as appropriate. * Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to: hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction machine, etc.) * Assess and evaluate cardiac, respiratory, GI, GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs * Instructions on care of terminally ill, medications, safety, nutritional needs, pain and symptom control based on assessment * Review and reconcile current medications * May give nutritional supplements 1 can a day as tolerated for supplement * May crush medications that are crush-able. * Medications to be administered by licensed staff, patient and/or family member/caregiver_per medication profile and current orders * Hospice Medical Director may assist patient as needed to manage symptoms * Hospice nurse may pronounce death * Oxygen 3.0 fpm via NC continuously as needed for SOB * Check oxygen saturation level pm every SN visit * Bed-rails up 2x * Skilled Nurse visits 1x/week and 2x/week as needed for pain, SOB, fall & anxiety issues to promote comfort and symptom management within certification period. * CHHA visits 5x/week to assist with personal care, home making and to promote comfort per RN assignment within certification period. * Chaplain visits -1x/mo, 1x/mo as needed for spiritual issues. * Social worker visits - 1x/mo as needed for psychosocial issues. * Volunteer Services continue to be offered every week per Skilled Nurse/Staff. * Admit patient to Jireh Healthcare Services with the following orders verified with M.D: Patient: DNR Diet: As Tolerated * Admit with the following medications: Please see related medications.

Read Back

Non Verbal Order

Yes

No

Related Assignees

Chaplain - CHAPLAIN_Murphy, Kathryn 1 visit(s) Monthly 3 PRN visit(s) every 30 day(s) effective 09/12/2018

Hospice Aide - CNA_Barba, Magdalena 5 visit(s) Weekly effective 09/12/2018

MD Director - Dubin, Stephen Paul 1 PRN visit(s) every 30 day(s) effective 09/12/2018

Physician - Inocencio, Carlos 1 PRN visit(s) every 30 day(s) effective 09/12/2018

Skilled Nurse - RN_Cacanindin, Speedy 1 visit(s) Weekly 2 PRN visit(s) every 7 day(s) effective 09/12/2018

Skilled Nurse - RN_Ruiz, Janyvill 2 PRN visit(s) every 7 day(s) effective 09/12/2018

Signatures

1. Medical Director Dubin, Stephen Paul (Medical Director) signed on 12/03/2018. Recorded by jduga on 12/03/2018 15:13:16.

Consolo Physician Order (2253673) Dated: 09/12/2018 for Dubin, Stephen Paul for Dated: 09/12/2018 for Dated:

Physician's Order Details

User

RN_Esteban, Randy

Pa

Order Date 09/12/2018 Time of Event 1700

(Reste)

Physician Dubin, Stephen Paul Nurse RN_Esteban, Randy (Reste) Overslaht Yes

Orders

Discontinue previous chaptain visit frequency. Start new chaptain visit frequency: 1 time a month and 3 times a month as needed for spiritual issues.

Read Back Yes Non Verbal Order
No

4/17/19 7:14 PM EST

Consolo Physician Order for DEWxA at Office Jireh Healthcare Services LLC

Patient Details for Details for Details Application ID 491783, MRN M0466)						
Date Of Birth	Status Active					

Physician's Order Details

<u>User</u>

RN_Esteban, Randy

(Reste)

.

<u>Patient</u>

Order Date 09/12/2018 Time of Event 1700

Physician

Dubin, Stephen Paul

RN_Esteban, Randy

(Reste)

Oversight Yes

Orders

Discontinue previous chaplain visit frequency. Start new chaplain visit frequency: 1 time a month and 3 times a month as needed for spiritual issues.

Read Back Yes Non Verbal Order

No

Signature:

Date:

9/12/18



Patient Details for Date

Patient ID 491783, MRN M0466)

Date Of Birth

Status Pending - Discharged

Hospice Certification of Terminal Illness

Benefit Period

Benefit Period Start Date 09/07/2018

Benefit Period End Date 12/05/2018

Certification Start Date 09/07/2018

Cartification End Date 12/05/2018

Signed Date 09/07/2018 Medical Director or Certifying

Physician

Dubin, Stephen Paul

Physician Address

5741 S. Fort Apache Rd. Suite 100 Las Vegas, NV

Medical Director or Certifying

89148

Attending Physician

Inocencio, Carlos

Attending Physician Address 2320 Paseo Del Prado

Ste. B303 Las Vegas, NV 89102

Patient Hi Claim No 025324213A

Disaster Aculty

Medical services required within 72 hours

Provider Information

Provider Name

Jireh Healthcare Services LLC

Provider Number 291533

Provider Address

2320 Paseo Del Prado Ste B101 Las Vegas, NV 89102-0048

Admission

New Admission 09/07/2018 (ROUTINE)

Verbal Certification

Verbal Certification received from Medical Director: Dubin, Stephen Paul by RN_Ruiz, Janyvill (jruiz) on 09/06/2018

I certify that D Asis terminally ill with a life expectancy of six months or less if the terminal illness runs its normal course.

Skilled Nurse RN_Ruiz, Janyvill (User) signed on 09/06/2018. Recorded by jruiz on 09/10/2018 09:55:52.

Verbal Certification received from Physician: Inocencio, Carlos by RN_Ruiz, Janyvill (jruiz) on 09/06/2018

Affiles terminally ill with a life expectancy of six months or less if the terminal illness runs its normal course.

Skilled Nurse RN_Rulz, JanyvIII (User) signed on 09/06/2018. Recorded by Jruiz on 09/10/2018 09:55:59.

Brief Narrative Statement and Attestation

Review the patient's clinical circumstances and synthesize the medical information to provide clinical justification for admission and continuation to hospice services. If in third (3rd) or more benefit periods, clinical findings from the face to face encounter have been used to determine continued eligibility for hospice care.

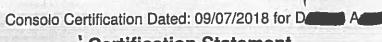
Dubin, Stephen Paul

Composing Physician Ms. Desired Strategies 75 year old, Caucasian Female, with very poor short term memory due to worsening alzheimer, requires considerable taxing effort to do her ADL's due to severe weakness and fatigue. Caregiver has to provide maximum to complete assist on patient as she is very high risk to fall and is very confused. As per caregiver Susan, patient refused to eat at times and would just stare on her plate. Todd the son of Ms. Dewey reported that her morn had lost 15 lbs in 2 months. Patient has imminence of death and if patient continues this course her condition is terminal and prognosis is six months or less.

Recorded By RN_Ruiz, Janyvill (jruiz)

Attestation: I confirm that I composed this narrative and that it is based on my review of the patient's medical record and/or examination of the patient.

Medical Director Dubln, Stephen Paul (Medical Director) signed on 09/07/2018. Recorded by jruiz on 09/10/2018 10:06:44.



, Certification Statement

I certify that this patient is under my care, and to the best of my medical knowledge given the data available, has a life expectancy of six (6) months or less if the terminal illness runs its normal course.

Medical Director or Certifying Medical Director Dubin, Stephen Paul (Medical Director) signed on 09/07/2018. Recorded by jruiz on 09/10/2018 10:06:56.

I certify that this patient is under my care, and to the best of my medical knowledge given the data available, has a life expectancy of six (6) months or less if the terminal illness runs its normal course.

Attending Physician: Physician Inocencio, Carlos (Physician) signed on 09/07/2018. Recorded by jrulz on 09/10/2018

Additional Signatures

1. Skilled Nurse RN_Rulz, Janyvill (User) signed on 09/10/2018. Recorded by jruiz on 09/10/2018 10:07:26.

Page 2 of 2

JIZEH HEALTHCARE SERVICES

2320 Paseo Del Prado Suite B101 Las Vegas Nevada 89102 PHONE: 702.359.1388 FAX: 702.359.2388 EMAIL ADDRESS: jirehhealthcare @gmail.com

PHYSICIAN'S CERTIFICATION FOR MEDICARE/MEDICAID HOSPICE BENEFIT

Physician's Certification of Terminal Illness for Hospice Benefit (Part -1)

MR# 140461 CERTIFICATION STATEMENT First 90-Day Period from 907 2018 to 12 05 2018 I (or WE) certify that is terminally ill and Has a life expectancy of six (6) months or fess if the terminal illness runs its normal course. I (WE) Reviewed the patient's clinical information and considered the primary terminal condition, related diagnoses, current subjective and objective medical findings, current medication and treatment orders and information about management of unrelated conditions in making this determination. Yerbal Order Date RN Signature 9/07/2012 Hospice Medical Director/Associate Hospice Doctor 9/07/2017 Date Attending Physician Signature RECERTIFICATION STATEMENT Second 90-Day Period from -i2-6-18 to 3-05-19I certify that I have reviewed the clinical record prior to recertification for the above noted patient and That patient is still considered to be terminally ill and has life expectancy of six (six) months or less, if The terminal illness runs its normal course. Hospice Medical Director/Associate Hospice Doctor Date RECERTIFICATION STATEMENT 60-Day Period from I certify that I have reviewed the clinical record prior to recertification for the above noted patient and That patient is still considered to be terminally ill and has life expectancy of six (6) months or less, if The terminal illness runs its normal course. Hospice Medical Director/Associate Hospice Doctor Date

B.

C.

Consolo Certification for DEWxA at Office Jireh Healthcare Services LLC

Patient Details for Description Apple (Patient ID 491783, MRN M0466)

Date Of Birth	-11	Status	Active
Screened Allergies	No Known Drug Allergy	Unscreened Allergies	See Screened Allergies
Start Of Care	09/07/2018	Gender	Female
Patient Address	3158 Kingspoint Ave. Las Vegas, NV	89120	1

Hospice Certification of Terminal Illness

Benefit Period

Benefit Period Start Date

Benefit Period End Date

Certification Start Date

Signed Date

12/05/2018

09/07/2018

Certification End Date 12/05/2018

09/07/2018

Medical Director or Certifying

Medical Director or Certifying Physician Address

09/07/2018

Physician

5741 S. Fort Apache Rd.

Dubin, Stephen Paul

Suite 100 Las Vegas, NV

89148

Attending Physician

Attending Physician Address

Patient Hi Claim No

Disaster Aculty

Inocencio, Carlos

2320 Paseo Del Prado Ste. B303 Las Vegas, NV . 025324213A

Medical services required within 72 hours

89102

Provider Information

Provider Name

Provider Number Jireh Healthcare Services

291533

Provider Address

2320 Paseo Del Prado Ste

B101 Las Vegas, NV

89102-0048

Admission

LLC

New Admission 09/07/2018 (ROUTINE)

Verbal Certification

Verbal Certification received from Medical Director: Dubin, Stephen Paul by RN_Ruiz, Janyvill (jruiz) on 09/06/2018

I certify that Demanded Amis terminally ill with a life expectancy of six months or less if the terminal illness runs its normal course.

Skilled Nurse RN_Ruiz, Janyvill (User) signed on 09/06/2018. Recorded by jruiz on 09/10/2018 09:55:52.

Verbal Certification received from Physician: Inocencio, Carlos by RN_Ruiz, Janyvill (fruiz) on 09/06/2018

I certify that D is terminally ill with a life expectancy of six months or less if the terminal illness runs its normal course.

Skilled Nurse RN_Ruiz, Janyvill (User) signed on 09/06/2018. Recorded by jruiz on 09/10/2018 09:55:59.

Consolo Certification for DEWxA at Office Jireh Healthcare Services LLC

Brief Narrative Statement and Attestation

Review the patient's clinical circumstances and synthesize the medical information to provide clinical justification for admission and continuation to hospice services. If in third (3rd) or more benefit periods, clinical findings from the face to face encounter have been used to determine continued eligibility for hospice care.

Composing Physician Dubin, Stephen Paul

Ms. Days is 75 year old, Caucasian Female, with very poor short term memory due to worsening alzheimer, requires considerable taxing effort to do her ADL's due to severe weakness and fatigue. Caregiver has to provide maximum to complete assist on patient as she is very high risk to fall and is very confused. As per caregiver Susan, patient refused to eat at times and would just stare on her plate. Todd the son of Ms. Dewey reported that her mom had lost 15 lbs in 2 months. Patient has imminence of death and If patient continues this course her condition is terminal and prognosis is six months or less.

Recorded By

RN Ruiz, Janyvill (jruiz)

Attestation: I confirm that I composed this narrative and that it is based on my review of the patient's medical record and/or examination of the patient.

Medical Director Dubin, Stephen Paul (Medical Director) signed on 09/07/2018. Recorded by jruiz on 09/10/2018 10:06:44.

Certification Statement

I certify that this patient is under my care, and to the best of my medical knowledge given the data available, has a life expectancy of one six (6) months or less if the terminal illness runs its normal course. Provider Marie

Medical Director or Certifying Physician: Medical Director Dubin, Stephen Paul (Medical Director) signed on 09/07/2018. Recorded by jruiz on 09/10/2018 10:06:56.

I certify that this patient is under my care, and to the best of my medical knowledge given the data available, has a life expectancy of six (6) months or less if the terminal illness runs its normal course. TENTON SHIPE!

Attending Physician: Physician Inocencio, Carlos (Physician) signed on 09/07/2018. Recorded by jruiz on 09/10/2018 10:07:05.

Vieted Cathigation have an from Madiso Time on Time and a cooker Fatt in Rt. Mula steep vill yours, or 08/06/2018 the first story of the statement of our or the expensions of our regular is took the density at the second

Million Murse RM Ruis Jam William Femedon St 193 2018 Recontro to long on 22 17279 174551

Additional Signatures

1. Skilled Nurse RN Ruiz, Janyvill (User) signed on 09/10/2018. Recorded by jruiz on 09/10/2018 10:07:26. Ann is herceastly के क्षेत्रि के तीव क्षत्रकार्यात्रामु को जोत तरानांकी प्रशासिक निर्माण विश्वास कर्माव d secon

ridach death

Screened Allergies No Known Drug Allergy, No Known Drug Allergy

(491783)

MRN M0466

Start Of Care 09/07/2018

Level Of Care ROUTINE More

Hospice Certification of Terminal Illness

Benefit Period

Benefit Period Start Date

12/06/2018

Signed Date 12/03/2018 **Benefit Period End Date** 03/05/2019

Medical Director or Certifying Physician

Dubin, Stephen Paul

Patient Hi Claim No

Certification Start Date 12/06/2018

Medical Director or Certifying Physician Address 5741 S. Fort Apache Rd. Suite 100 Las Vegas, NV 89148

Disaster Acuity Medical services required within 72 hours

Attending Physician Inocencio, Carlos

Certification End Date

03/05/2019

Attending Physician Address 2320 Paseo Del Prado Ste. B303 Las Vegas, NV 89102

025324213A

Provider Information

<u>Provider Name</u> Jireh Healthcare Services LLC <u>Provider Number</u> 291533

Provider Address 2320 Paseo Del Prado Ste B101 Las Vegas, NV 89102-0048

Admission

New Admission 09/07/2018 (ROUTINE)

Verbal Certification

Verbal Certification received from Medical Director: Dubin, Stephen Paul by Dugay, Julia (jduga) on 12/03/2018

I certify that D A is terminally ill with a life expectancy of six months or less if the terminal illness runs its normal course.

Skilled Nurse Dugay, Julia (User) signed on 12/03/2018. Recorded by Jduga on 12/03/2018 14:06:53

Brief Narrative Statement and Attestation

Review the patient's clinical circumstances and synthesize the medical information to provide clinical justification for admission and continuation to hospice services. If in third (3rd) or more benefit periods, clinical findings from the face to face encounter have been used to determine continued eligibility for hospice care.

Composing Physician Dubin, Stephen Paul

Jireh healthcare services for the 2nd 90-day benefit episode. She is a 75 y/o white Recertifying Ms ABD female with worsening Alzhelmer's Disease requiring maximum assist with her ADL/IADLs. She is oftentimes confused and eats 20-25% of her meals. If patient continues this course, her prognosis is 6 months or less.

Recorded By Dugay, Julia (jduga)

Attestation: I confirm that I composed this narrative and that it is based on my review of the patient's medical record and/or examination of the

Medical Director Dubin, Stephen Paul (Medical Director) signed on 12/03/2018. Recorded by Jduga on 12/03/2018 14:06:16

Certification Statement

ertify that this patient is under my care, and to the best of my medical knowledge given the data available, has a life expectancy of six (6) months is if the terminal illness runs its normal course.

Medical Director or Certifying Physician: Medical Director Dubin, Stephen Paul (Medical Director) signed on 12/03/2018. Recorded by Jduga on 12/03/2018

I recertify that this patient is under my care, and to the best of my medical knowledge given the data available, has a life expectancy of six (6) months or less if the terminal illness runs its normal course.

Attending Physician: Physician Inocencio, Carlos (Physician) signed on 12/03/2018. Recorded by jduga on 12/03/2018 14:05:58.

Additional Signatures

1. Skilled Nurse Dugay, Julia (User) signed on 12/03/2018. Recorded by Jduga on 12/03/2018 14:08:23.

Jireh Healthcare Services LLC

PHYSICIANS ORDER

2320 Pas@0 Del Prado, Suite B101, Las Vegas, NV 89102 . Tel: Fax:

F. ENT:

De A

ORDEREPBY: STEPHEN DUBIN (MD)

DOB:

ORDER DATE:

09/07/18

MR#: ORDER#: M0466

1598165

List of New/Refill Orders

Start Date Stop Date	Туре		Order	Strength	Quantity	Dosage	Route	Frequency	Indication	Payer
09/14/18	Medication	New	RISPERDAL (risperidone)	0.5 mg		1 Tab	ORAL	2 times a day		

art Date	Stop Date	Туре		Order	Strength	Quantity	Dosage	Route /	Frequency	Indication	Payer
09/07/18		Medication	Admit	Alprazolam (Alprazolam)	1 mg	1 3	1 Tab	ORAL	Once a day - At bedtime		117
	==>	Note: To be to	aken ever	night at bedtime.						Jan 1241 241	
09/07/18		Medication	Admit	Escitalopram (Escitalopram Oxalate)	10 mg		0.5 Tab	ORAL	Daily	111	
-	==>	Note: Take h	alf a tablet	once daily							
09/07/18		Medication	Admit	Galantamine (Galantamine)	24 mg	17	1 Cap ER	ORAL	Daily		
	==>	Note: Take or	ne tablet o	nce daily.							
09/07/18		Medication	Admit	Metoprolol Tartrate (Metoprolol tartrate)	50 mg		1 Tab	ORAL	2 times a day		

A TIONAL NOTE

Merging information from Consolo

PHONE ORDER / READ BACK ORDER BY MARY ROSE FRONDA (RN)

Electronically Signed: 9/7/2018, By STEPHEN DUBIN, MD

CONFIDENTIALITY NOTICE: The information contained in this facsimile is confidential and may contain privileged material or be otherwise protected by applicable law. It is intended only for the use of the individual (s) or entity named above. If the person receiving this facsimile, or any other reader of the facsimile, is not the intended recipient or the employee or agent responsible for delivering it to the Intended recipient, then any use, dissamination, distribution, or copying of this communication is strictly prohibited and may be subject to civil and/or criminal liability. If you have received this communication in error, please destroy it and notify the sender immediately.

Jireh Healthcare Services LLC

PHYSICIANS ORDER

2320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102 . Tel: Fax:

NT: DE A

ORDERED BY: STEPHEN DUBIN (MD)

DOB: ORDER DATE: 09/10/18

MR#: ORDER#: M0466 1598200

List of New/Refill Orders

Start Date	Stop Date	Туре		Order	Strength	Quantity	Dosage	Route	Frequency	Indication	Payer
09/10/18		Medication	New	Losartan (Losartan Potassium)	100 mg		1 Tab	ORAL	Daily		
	==>	Note: Take	one tablel	t by mouth once daily.				-			
09/10/18	-	Medication	New	Namenda XR (memantine hydrochloride)	28 mg		1 Cap ER	ORAL	Daily		-

==> Note: Take one tablet by mouth once daily.

List of 'ADMIT' Items - DO NOT FILL ITEMS BELOW (Patient has supply of item at admission)

Start Date Stop Date Type Order Strength Quantity Dosage Route Frequency Indication Payer

PHONE ORDER / READ BACK ORDER BY MARY ROSE FRONDA (RN)

Electronically Signed: 9/10/2018, By STEPHEN DUBIN, MD



Y PHARMACY

LAS VEGAS, NV 89109 TEL 702.776.8210

Name:	1	6	
Address	;		
riuui 600 Tala			

Date: 18/18/18

Time:

alp. Img

Tahs Anxist #

Ro



DR.: TEL:

NPI:

DEA:

Honey

FD 2419706 359-1388

DUBIN004

Trinity Pharmacy 2797 S. Maryland Pkwy 28 Las Vegas, NV 89109 Phone: 702-776-8210 Fax: 702-776-7195 **Telephoned Prescription** STEPHEN DUBIN 5380 S RAINBOW STE 306 DEA #: FD2419706 LAS VEGAS NV 89128 Lic#: 13772 Phone: 702-359-1388 : 702-362-9954 NPI#: 1023311040 Rx#: 168817 Rx Written: 11-30-2018 **Patient** Address AVE LAS VEGAS NV 89120 Phone # 7024 D.O.B. Gender : Female PRAZOLAM 1 MG TABLET Generic for XANAX 1MG TABLET uantity: 15 a Instructions: KE ONE TABLET BY MOUTH AT BEDTIME FOR ANXIETY/SLEEP Refills Authorized: 0 RPh: L. Diagnosis Codes: F41.9 Signature THIS PRESCRIPTION WILL BE FILLED GENERICALLY UNLESS PRESCRIBER WRITES "DAW" IN THE BOX BELOW

Dispense As Written

DUBINO050

Trinity Pharmacy 2797 S. Maryland Pkwy 28 Las Vegas, NV 89109 Phone: 702-776-8210 Fax: 702-776-7195 **Telephoned Prescription** STEPHEN DUBIN **5380 S RAINBOW STE 306** DEA#: FD2419706 LAS VEGAS NV 89128 Lic# : 13772 Phone: 702-359-1388 NPI#: 102331104 : 702-362-9954 Fax Rx#: 168116 Rx Written: 11-19-2018 Patient 1 Address AVE **LAS VEGAS NV 89120** Phone # 702-D.O.B. Gender **Female** ALPRAZOLAM 1 MG TABLET Generic for XANAX 1MG TABLET Quantity: 15 Sig Instructions: TAKE ONE TABLET BY MOUTH AT BEDTIME FOR ANXIETY/SLEEP Refills Authorized: 0 RPh: LJ Diagnosis Codes : F41.9 Signature THIS PRESCRIPTION WILL BE FILLED GENERICALLY UNLESS PRESCRIBER WRITES "DAW" IN THE BOX BELOW Dispense As Written

कुरियाः

Trinity Pharmacy

2797 S. Maryland Pkwy 28 Las Vegas, NV 89109

Phone: 702-776-8210

Fax: 702-776-7195

Telephoned Prescription

STEPHEN DUBIN

5380 S RAINBOW STE 306

DEA#: FD2419706 Lic#: 13772

LAS VEGAS NV 89128

NPI#: 1023311040

Rx Written: 10-29-2018

Rx#: 168757

Phone: 702-359-1388

Fax: 702-362-9954

Patient

: Daniel, A

Address

AVE

LAS VEGAS NV 89120

Phone #.

: 702-4

D.O.B.

Gender :

Female

ALPRAZOLAM 1 MG TABLET

Generic for XANAX 1MG TABLET

Quantity: 15

Sig Instructions:

TAKE ONE TABLET BY MOUTH AT BEDTIME FOR ANXIETY/SLEEP

Refills Authorized: 0

Diagnosis Codes: F41.9

MN

RPh: MLJ/C

@ L.41bw

Date

Signature

THIS PRESCRIPTION WILL BE FILLED GENERICALLY UNLESS PRESCRIBER WRITES "DAW"

AVE LAS VEGAS NV 89120

Dispense As Written

Trinity Pharmacy 2797 S. Manyland Pkwy 28 Las Vegas, Phone: 702-776-8210 Fax: 702 elephoned Prescription STEPHEN DUBIN 5380 S RAINBOW STE 308 LAS VEGAS NV 89128 Fax: 702-352-9554	NV 89109 : 2-776-7195 DEA # : FD2419708 Lic # : 13772 NPI# : 1023311040
Ref: 18-5873 Patient: Description AVE LAS VEGAS NV 89120 Phone #: 702- D.O.B. Gender: Female ALPRAZOLAM 1 MG TABLET Generic tor XANAX 1MG TABLET Quantity: 15 (fifteen) Big Instructions: AME ONE TABLET BY MOUTH AT BEDTIME FOR ANXIE	Ro Written: 10-15-2018 4 PM 5 WWW
Refills Authorized: 0 RPh: LJ Diagnosis Cores: F41.9 USignature THE PRESCRIPTION WILL BE FILED GENERICALLY UNLESS IN THE BOX BELOW	Date Date
Dispense As. Written Dispense As. Written 16507.	7-

THE HILL AS VEGAS HV MIZE R X O 1 6 5 8 7 3 a Principal Columbia

Trinity Pharmacy
2707 S. Maryland Provy 28 Las Vegas, NV 86109
Phone: 702-776-8210 Fax: 702-776-7195 elephoned Prescription STEPHEN DUBIN 5380 S RAINBOW STE 306 DEA#: FD2419706 Lig#: 13772 NPI#: 1023311040 Phone: 702-359-1388 Fax: 702-362-9954 LAS VEGAS NV 89128 Rud : 164958 Rx Written: 10-1-2018 Patient Address Phone # D.O.B. Gender ALPRAZOLAM 1 MG TABLET Generic for XANAX 1MG TABLET Quantity: 15 Sig Instructions: TAKE ONE TABLET BY MOUTH AT BEDŢĪME FOR ANXIETY/SLEEP PRESCRIPTION WILL BE FILLED GENERICALLY UNLESS PRESCRIBER WRITES "DAW"
IN THE BOX BELOW Dispense As Written

JIREH HEALTHCARE SERVICES LLC (HOSPICE AGENCY)

PATIENT NAME: M

C

DOB:

START OF CARE: 10/10/2018 TO 12/31/2018

EXHIBIT 7

Claim Number	From Service Date To Ser	To Service Date E	vice Date Beneficiary Nam Chaim Status	Total Charge	Type Of Bill Admit Date
21900700422207NVR	12/1/2018	12/31/2018	P - Paid	75.067,78	813 - Hospice (9/7/2018
21833900538807NVR	11/1/2018	11/30/2018	P - Paid	\$8,724,43	813 - Hospice (9/7/2018
21830901126607NVR	10/1/2018	10/31/2018	P - Paid	\$9,083.84	813 - Hospice (9/7/2018
21827601222707NVR	9/7/2018	9/30/2018	P - Paid	\$6,942.12	812-Hospice (_9/7/2018
Claim Number	From Service Date To Serv	To Service Date	म टिवाम Status	Total Charge	Type Of Ball Admit Bate
21900700421907NVR	12/1/2018	12/31/2018	P - Paid	\$6,986.61	813 - Hospice (10/10/2018
21833900542507NVR	11/1/2018	11/30/2018	P - Paid	\$8,197.48	813 - Hospice (10/10/2018
21833900230804NVR	10/10/2018	10/31/2018	P - Paid	\$6,230.74	817 - Hospice (10/10/2018
21830901121107NVR	10/10/2018	10/31/2018	b-Paid	\$5,630.74	812-Hospice (10/10/2018
Claim Number	From Service Bate To Service Date	To Service Date	am Glaim Status	Total Charge	Type Of Bill Admin Date
21900901108507NVR	12/1/2018	12/26/2018	I.,P-Paid	86,626,68	814 - Hospice (6/13/2018
21833900542007NVR	11/1/2018	11/30/2018	P - Paid	\$7,635.75	813 - Hospice (6/13/2018
21830901128607NVR	10/1/2018	10/31/2018	P. P. Paid	\$8,065.97	813, Hospice (6/13/2018
21827601223807NVR	9/1/2018	9/30/2018	P. Paid	\$7,382.91	813 - Hospice (6/13/2018
21824701185107NVR	8/1/2018	8/31/2018	F. P. Paid	\$8,276.79	813 - Hospice (6/13/2018
21821800581907NVR	7/1/2018	7/31/2018	F. P - Paid	\$9,216.50	813 - Hospice (_6/13/2018
21818600817807NVR	6/13/2018	6/30/2018	E P Paid	\$5,126.72	812 - Hospice (6/13/2018
Claim Number	From Service Bute To Service Bate	To Service Date	m Glaim Status	Tetal Charge	Type Of Bill Admit.Date
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Jireh Healthcare Services LLC



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MR#		10474	-0.041.21=		SSN				DOB		56 yrs)
LAST NAME			G	-	FIRST	NAME	М		MI		
ADDRESS			Dr.	ive			Marting State gastages		APT / SUITE #		
CITY	L	as Veg	as		STATE		NV		ZIP	89108	
PHONE	_(702			CELL	- 1			PRÍM LANGUAGE	• English	
RACE	E	llack			ETHNIC	CITY	Afro-American		MARITAL ST	Single	
GENDER	F	emale			RELIGI	ON	Baptist		DENOMINATION		
Insuranc	e						- 4	4			4.5
PAY SOURCE	INSURA	NCE	SHARE OF COST	POLICY	/ MBI #	GROUP#	BIN#	EFFECTIV DATE	E AUTH TYPE	R&B AUTH NO	R&B AUTH VALID TILI
rimary	Medicare		\$0.00					10/10/2	018		
Datas of	Camina	Dia	ovenie e	(<i>11 aum</i> i a.			y .				
REF DATE		, <i>Diu</i> j	gnosis & A	utergies	SOC		10/10/2018		RE-CERT	1/7/2019	
	-				EOC	F	1/7/2019		DISC CODE	Status Impro	ved
ADMIT TYPE New Admission STARTING CERT# 1 PRIMARY DX Unspecified sev		nission									
		ified severe pr	otein-calor	ie malnut	trition (E43)	a granden de de minimum promo	An .	DISEASE	FTT / Debility Disease)	(Non-Specific	
SECOND D	< ,	Adult fa	ilure to thrive (R62.7)	And the second s				DISEASE	-	
COMORBID	ITIES					*					
ALLERGIES											LAST UPDA
Place of	Service	(POS	5)								
ГҮРЕ		lome	7		FACILI	TY NAME					
EFFECTIVE	DATE			and the second	ROOM	NO			PCG		
ADDRESS			Dr	rive	CITY		Las Vegas		STATE-ZIP	NV 8910	8 -
PHONE		702)		Pri Terri	ALT-PH	HONE		jk-	FAX	And the second second	
,	ed Rep	(AR)	/Emergen	cy Con	tact (E	(C)					
Authoriza	Address: ,	, , Pho	one: , Alt Phon	e: , AR/EC	is:						
			tending M	D							
AR/EC ,,	Source	& At			N DUBIN	. Phone:, Fa	x:(702) 362-9930	, NPI:10233110)40		
ARIEC Referral		& At	,	STEPHE	10000111						
AR/EC ,,	RECTOR		de la				2) 362-9930, Fax.	(702) 362-9930	, NPI:1023311040		
Referral MEDICAL DI	RECTOR		de la				2) 362-9930, Fax. =	(702) 362-9930	, NPI:1023311040		

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Date Of Birth

Status Pending - Discharged

Assignment Details

Effective Date: 10/17/2018, Medical Record No: M0474, Dme Provider: Advantage Home Medical Services, Inc., Pharmacy: Trinity Pharmacy, Patient

IDG Team

			Default for New	1	PRN	4
Discipli	ne Name	Primary?	Medications?	Visits	Visits	Frequency Notes
Chaplai	n CHAPLAIN_Murphy, Kathryn	Yes	No	- =		Eval, 1x/mo, & 3x/mo PRN for spiritual distress w/in cert pd
Hospice Aide	CNA_Munive,	Yes	No	5 visit(s) Weekly		5X/W
Hospice Aide	Young, Grace Marie	No	No	5 visit(s) Weekly		
MD Director	Dubin, Stephen Paul	Yes	No			1x/mo PRN within certification period for Emergency Respons
Physicia	n Dubin Stephen Paul	Yes	No			1x/mo PRN withn certification period for Emergency Respons
Skilled Nurse	RN_Cacanindin, Speedy	Yes	No			1x/wk82x/wk PRN for fall & anxiety issues within cert perd
Skilled Nurse	RN_Domondon, Mary Alyn	No	No	1		Eval &2x/wk PRN for fall & anxiety issues within cort perd
Social Services	MSW_Minnlck, Joseph	Yes	No			Eval 1x/mo PRN for psychosocial issues within cert period



Patient Details for 1

G y, M (Patient ID 501364, MRN M0474)

Date Of Birth

Status Pending - Discharged

Assignment Details

Effective Date: 10/10/2018, Medical Record No. M0474, Drne Provider: Advantage Home Medical Services, Inc. , Pharmacy: Trinity Pharmacy

IDG Team

Discipline	Name	Primary?	Default for New Medications?	Visits	PRN Visits	Frequency Nates
Chaplain	CHAPLAIN_Murphy Kathryn	Yes	No	•		Eval. 1x/mo, & 3x/mo PRN for spiritual distress with cert pd
Hospice Aide	Young, Grace Marie	No	No	5 visit(s) Weekly		
MD Director	Dubin Stephen Paul	Yes	No		T -	1x/mo PRN withn certification period for Emergency Respons
Physician	Oubin, Stephen Paul	Yes	No			1x/mo PRN withn certification period for Emergency Respons
Skilled Nurse	RN_Domondon, Mary Alyn	No .	No			Eval &2x/wk PRN for fall & anxiety issues withn cert perd
Skilled Nurse	RN_Cacanindin Speedy	Yes	No			1x/wk82x/wk PRN for fall & anxiety issues within cert perd
Social Services	MSW_Minnick, Joseph	Yes	No	1		Eval, 1x/mo PRN for psychosocial issues within cert period

DUBIN0060

Consolo Discharge - No Longer Appropriate 01/07/2019 (ROUTINE) for Telephone

E

Patient Details for

G

y, M (Patient ID 501364, MRN M0474)

Date Of Birth

Status Pending - Discharged

Care Level Change Details

Discharge - No Longer Appropriate Effective Datetime 01/07/2019 13:27 PST Level Of Care ROUTINE Office

Jirch Healthcare Services

LLC

Status COMPL

Checklist

Completed 01/07/2019 - Call Primary Physician

Completed 01/07/2019 - Complete Advance Beneficiary Notice form

Completed 01/07/2019 - Complete Notice Of MEDICARE Provider Non-Coverage

Completed 01/07/2019 - Discharge Planning & Coordination of Care

Completed 01/07/2019 - Notify Ancillary Services (i.e. Massage, Music, Pet Therapy, etc)

Completed 01/07/2019 - Notify Bereavement Coordinator

Completed 01/07/2019 - Notify Care Facility

Completed 01/07/2019 - Notily Chaplain

Completed 01/07/2019 - Notify Clinical Director

Completed 01/07/2019 - Notify DME provider and request pickup

Completed 01/07/2019 - Natily Hospice Aide Supervisor

Completed 01/07/2019 - Notify Insurance Company/Caseworker

Completed 01/07/2019 - Notify Medical Director

Completed 01/07/2019 - Notify Medical Records

Completed 01/07/2019 - Notify Pharmacy

Completed 01/07/2019 - Notify Primary Nurse

Completed 01/07/2019 - Notily Senior Services/Caseworker

Completed 01/07/2019 - Notity Social Worker

Completed 01/07/2019 - Notify Volunteer

Completed 01/07/2019 - Notify Volunteer Coordinator

Completed 01/07/2019 - Print IDG report.

Completed 01/07/2019 - Verify HIS Data

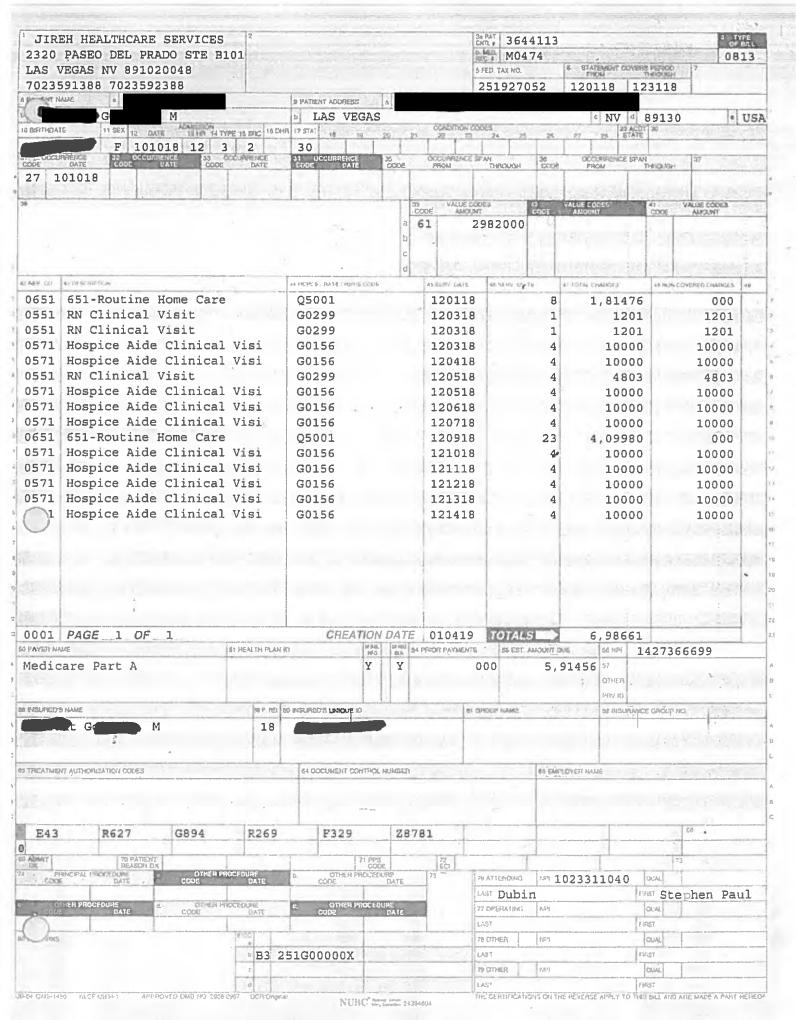
Discharge Details

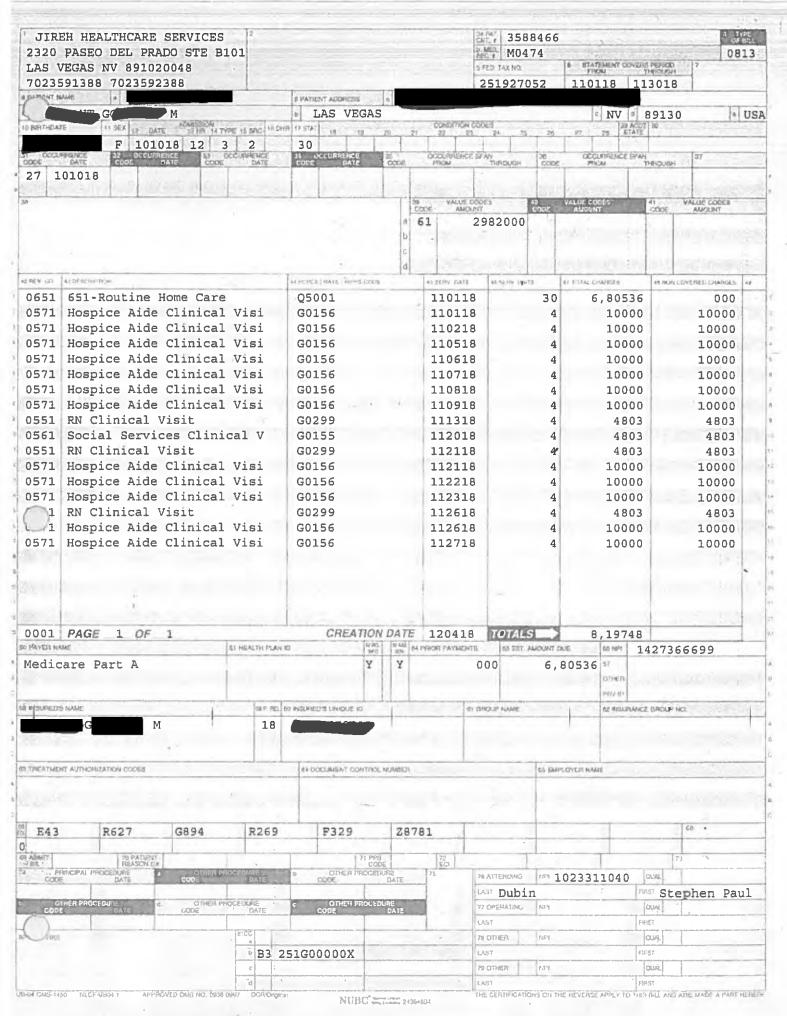
Notes

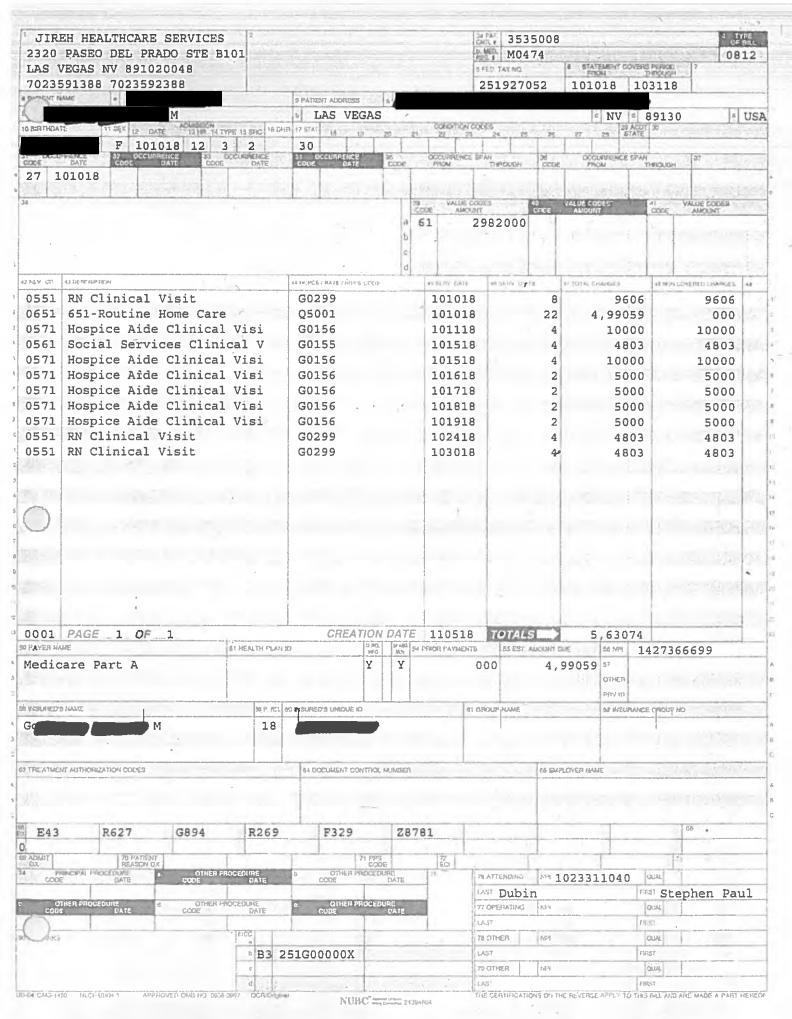
Patient is discharge to hospice care today 01/07/2019. Patient is no longer appropriate for hospice. She is also seeking for a physical therapy treatment. RN Speedy Cacanindin discussed to patient about safety measure, emergency plan, and to see primary physician in 2 weeks to discuss about medication reconciliation, and to manage patient overall health.

2320 LAS	EH HEALTHCARE SERVICE PASEO DEL PRADO STE VEGAS NV 891020048 591388 7023592388	The state of the s			366659 M0474 FFD TAX NO. 251927052	8 STATEMENT COVS	213 PETROD 7 7 THROUGH 010719	0814_
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T HAME	P G M		b LAS VEGAS			c NV	89130	• USA
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Jireh Healthcare Services LLC

PHYSICIANS ORDER

88 South Rainbow Blvd Suite A, Las Vegas, NV 89146, Tel. Fax: MR#: PATIENT: DOB: M0474 ORDER DATE: ORDERED BY: 10/08/18 ORDER#: 4672884 **ALLERGIES:** List of New/Refill Orders Start Date Stop Date Type Order Indication Strength Quantity Dosage Route Frequency Payer 10/08/18 Other Please send an RN to do hospice evaluation and treat List of 'ADMIT' Items - DO NOT FILL ITEMS BELOW (Patient has supply of item at admission) Order Payer Start Date Stop Date Type Strength Quantity Dosage Route Indication Frequency PHONE ORDER / READ BACK ORDER BY MARY ALYN DOMONDON (RN) Electronically Signed: 10/8/2018, By STEPHEN DUBIN, MD

CONFIDENTIALITY NOTICE. The information contained in this facsimile is confidential and may contain privileged material or be otherwise protected by applicable law. It is intended only for the use of the Individual (s) or entity named above. If the person receiving this facsimile, or any other reader of the facsimile, is not the intended recipient or the employee or agent responsible for delivering it to the intended recipient, then any use, dissemination, distribution, or copying of this communication is strictly prohibited and may be subject to civil and/or criminal liability. If you have received this communication in error, please destroy it and notify the sender immediately.

Consolo Physician for TARxM at Office Jireh Healthcare Services LLC

Patient Details for Games, M (Patient ID 501364, MRN M0474)

Date Of Birth	Status Pending - Discharged
Screened Allergies Not Assessed	Unscreened Allergies See Screened Allergies

Physician's Order Details

<u>User</u>

RN_Ruiz, Janyvill (jruiz)

Patient M

Order Date 10/10/2018 Time of Event

<u>Physician</u>

Dubin, Stephen Paul

Nurse

RN_Ruiz, Janyvill (jruiz)

Oversight No

Orders

Admit to Jireh Healthcare Services, Lfc. for the 1st 90 day benefit period * Pharmacist may use generic drug equivalents or compound medications as appropriate. * Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to: hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction machine, etc.) * Assess and evaluate cardiac, respiratory, GI,GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs * Instruction on care of terminally ill, medications, safety, nutritional needs, pain and symptom control based on assessment * Assess and evaluate current medications * May give nutritional supplements 1 can a day as tolerated for supplement * May crush medications that are crush-able, * Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders * Hospice Medical Director may assist patient as needed to manage symptoms * Hospice nurse may pronounce death * Check oxygen saturation level prn every SN visit * Bed-rails up 2x * Skilled Nurse visits - (1x/week and 2x/week as needed for respiratory & anxiety issues) to promote comfort and symptom management within certification period. * CHHA visits 5x/week to assist with personal care, home making and to promote comfort per RN assignment within certification period. * Chaplain visits - (1x/mo as needed for spiritual issues) *Social worker visits - (1x/mo as needed for psychosocial issues) Volunteer Services continue to be offered every week per Skilled Nurse/Staff, Admit Patient to Jireh Healthcare Services with the following orders verified with M.D. * Patient: DNR * Diet: As Tolerated * Admit with the following medications: Please see related medications

Read Back Yes Non Verbal Order

No

Medications

Medication	Instructions	Quantity	Refills
SUMAtriptan Succinate 1 Tablet 2 times a day PO (50 MG Tablet)	take 1 tab by mouth 2 times per day at least 2 hours between doses	1.	
Sertraline HCl 1 Tablet 1 (Imes a day PO (100 MG Tablet)			
Oxycodone-Acetaminophen 1 Tablet every 6 hours PO As Needed Pain (10-325 MG Tablet)			0
Oxybutynin Chloride ER 1 Tablet ER 24HR 1 times a day PO (5 MG Tablet ER 24HR)			
Omeprazole 1 Capsule DR 1 times a day PO As Needed Gas Relief (40 MG Capsule DR)			
Lidocaine 1 Patch every 12 hours EX (5 % Patch)			

Related Assignees

Social Services - MSW_Minnick, Joseph effective 10/17/2018

Chaplain - CHAPLAIN_Murphy, Kathryn effective 10/17/2018

Hospice Aide - Young, Grace Marie 5 visit(s) Weekly effective 10/17/2018

Skilled Nurse - RN_Cacanindin, Speedy effective 10/17/2018

Skilled Nurse - RN_Domondon, Mary Alyn effective 10/17/2018

MD Director - Dubin, Stephen Paul effective 10/17/2018

Physician - Dubln, Stephen Paul effective 10/17/2018

Consolo Physician for TARxM at Office Jireh Healthcare Services LLC

Signatures

- 1. Medical Director Dubin, Stephen Paul (Medical Director) signed on 10/18/2018. Recorded by jruiz on 10/17/2018 18:32:50.
- 2. Skilled Nurse RN_Ruiz, Janyvill (User) signed on 10/17/2018. Recorded by jruiz on 10/17/2018 18:32:40.

Patient Details for Garage, M (Patient ID 501364, MRN M0474)

Date Of Birth

Status Pending - Discharged

Indicator Dated 10/10/2018 for M

Medical Record No M0474

Effective Date 10/10/2018

Diagnoses

ICD10s

Primary ICD10 - E43 Unspecified severe protein-calorie malnutrition Secondary ICD10 - R62.7 Adult failure to thrive Tertlary ICD10 - G89.4 Chronic pain syndrome 4th ICD10 - R26.9 Unspecified abnormalities of gait and mobility

5th ICD10 - F32.9 Major depressive disorder, single episode, unspecified 6th ICD10 - Z87.81 Personal history of (healed) traumatic fracture

P G	-			NA
		4	-	IVI

Patient Details for G

M (Patient ID 501364, MRN M0474)

Date Of Birth

Status Pending - Discharged

Physician's Order Details

User

RN_Ruiz, Janyvill (jruiz)



Order Date 10/10/2018 Time of Event

Physician

Dubin, Stephen Paul

Nurse RN_Ruiz, Janyv I (jruiz) Overslaht

Orders

Admit to Jireh Healthcare Services, Uc. for the 1st 90 day benefit period. Pharmacist may use generic drug equivalents or compound medications as appropriate. Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheekchair, walker, suction machine, etc.) Assess and evaluate cardiact, respiratory, Gl,GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs. Instruction on care of terminally ill, medications, safety, nutritional needs, pain and symptom control based on assessment. Assess and evaluate current medications. May give nutritional supplements 1 can a day as tolerated for supplement. May crush medications that are crush-able. Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders. Hospice Medical Director may assist patient as needed to manage symptoms. Hospice nurse may pronounce death. Check oxygen saturation level prin every SN visit. Bed-rails up 2x. Skilled Nurse visits. (1x/week and 2x/week as needed for respiratory & anxiety issues) to promote comfort and symptom management within certification period. CHHA visits 5x/week to assist with personal care, home making and to promote comfort per RN assignment within certification period. Chaplain visits. (1x/mo as needed for psychosocial issues) Volunteer Services continue to be offered every week per Skilled Nurse/Staff. Admit Patient to Jireh Healthcare Services with the following orders verified with M.D. Patient DNR. Diet: As Tolerated. Admit with the following medications.

Read Back

Non Verbal Order

Yes

No

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Sertraline HCl 1 Tablet 1 times a day PO (100 MG Tablet)	10 th H		
Oxycodone-Acetaminophen 1 Tablet every 6 hours PO As Needed Pain (10-325 MG Tablet)	¥ 14 - 12		0
Oxybutynin Chloride ER 1 Tablet ER 24HR 1 times a day PO (5 MG Tablet ER 24HR)			À
Omeprazole 1 Capsule DR 1 times a day PO As Needed Gas Relief (40 MG Capsule DR)	5		
Lidocalne 1 Patch every 12 hours EX (5 % Patch)			

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Hospice Aide - Young, Grace Marie 5 visit(s) Weekly effective 10/17/2018

Skilled Nurse - RN_Cacanindin, Speedy effective 10/17/2018 Skilled Nurse - RN_Domandon, Mary Alyn effective 10/17/2018

MD Director - Dubin, Stephen Paul effective 10/17/2018 Physician - Dubin, Stephen Paul effective 10/17/2018

Signatures

- 1. Medical Director Dubln, Stephen Paul (Medical Director) signed on 10/18/2018. Recorded by Jruiz on 10/17/2018 18:32:50.
- 2. Skilled Nurse RN_Ruiz, Janyvill (User) signed on 10/17/2018. Recorded by jruiz on 10/17/2018 18:32:40.

Physician's Order Details

User

RN_Ruiz, Janyvill (jruiz)

Patient M

Order Date 10/10/2018 Time of Event

<u>Physician</u>

Dubin, Stephen Paul

Nurse RN_Ruiz, Janyvill (Jruiz) Oversight No

Orders

Admit to Jirch Healthcare Services, Ltc. for the 1st 90 day benefit period * Pharmacist may use generic drug equivalents or compound medications as appropriate. * Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction machine, etc.) * Assess and evaluate cardiac, respiratory, Gl.GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs * Instruction on care of terminally ill, medications, safety, nutritional needs, pain and symptom control based on assessment * Assess and evaluate current medications * May give nutritional supplements 1 can a day as tolerated for supplement * May crush medications that are crush-able. * Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders * Hospice Medical Director may assist patient as needed to manage symptoms * Hospice nurse may pronounce death * Check oxygen saturation level prin every SN visit * Bed-rails up 2x * Skilled Nurse visits - (1x/week and 2x/week as needed for respiratory & anxiety issues) to promote comfort and symptom management within certification period. * CHHA visits 5x/week to assist with personal care, home making and to promote comfort per RN assignment within certification period. * Chaptain visits - (1x/mo as needed for spiritual Issues) * Social worker visits - (1x/mo as needed for psychosocial issues) Volunteer Services continue to be offered every week per Skilled Nurse/Staff. Admit Patient to Jirch Healthcare Services with the following orders verified with M.D. * Patient: DNR * Diet: As Tolerated * Admit with the following medications.

Read Back

Non Verbal Order

Yes

No

Related Assignees

Social Services - MSW_Minnick, Joseph effective 10/17/2018
Chaplain - CHAPLAIN_Murphy, Kathryn effective 10/17/2018
Hospice Aide - Young, Grace Marie 5 visit(s) Weekly effective 10/17/2018
Skilled Nurse - RN_Cacanindin, Speedy effective 10/17/2018
Skilled Nurse - RN_Domondon, Mary Alyn effective 10/17/2018
MD Director - Dubin, Stephen Paul effective 10/17/2018
Physician - Dubin, Stephen Paul effective 10/17/2018

Signatures

1. Medical Director Dubin, Stephen Paul (Medical Director) signed on 10/18/2018. Recorded by jruiz on 10/17/2018 18:32:22.

Jireh Healthcare Services LLC

PHYSICIANS ORDER

2320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102 . Tel: Fax:

TIENT:

G, M

DOB:

MR#:

M0474

ORDERED BY: STEPHEN DUBIN (MD)

ORDER DATE:

10/10/18

ORDER#:

1606966

List of New/Refill Orders

Start Date Stop Date Type

Order

Strength

Quantity

Dosage

Route Frequency Indication

Payer

tart Date	Stop Date	Туре		Order	Strength	Quantity	Dosage	Route	Frequency	Indication	Paye
10/10/1B		Medication	Admit	Oxybutynin Chloride Extended Release (Oxybutynin Chloride)	5 mg	,	1 Tab ER	ORAL	Daily		
10/10/18		Medication	Admit	OXYCODONE AND ACETAMINOPHEN 10/ 325 mg (oxycodone hydrochloride and acetaminophen)	10-325 mg		1 Tab	ORAL	Every 6 hrs - PRN	Pain	-
10/10/18		Medication	Admit	Sertraline (Sertraline Hydrochloride)	100 mg		1 Tab	ORAL	Daily	-	
10/10/18		Medication	Admit	IMITREX (SUMATRIPTAN)	50 mg	4	1 Tab	ORAL	2 times a day		

ADDITIONAL NOTE

Merging of information from Consolo Services

ONE ORDER / READ BACK ORDER BY MARY ROSE FRONDA (RN)

Electronically Signed: 10/10/2018, By STEPHEN DUBIN, MD

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Jireh Healthcare Services LLC

PHYSICIANS ORDER

2320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102. Tel: Fax:

TIENT:

G M

DOB:

MR#:

M0474

ORDERED BY: STEPHEN DUBIN (MD)

EPHEN DUBIN (MD)

ORDER DATE: 10/17/18

ORDER#:

1606911

List of New/Refill Orders

Start Date	Stop Date			Order	Strength	Quantity	Dosage	Route	Frequency	Indication	Payer
10/17/18		Medication	New	Lidocaine	5% patch	1	1 Patch	TOPICAL	Every 12 Hours		
10/17/18		Medication	New	Omeprazole (omeprazole)	40 mg		1 Cap DR	ORAL	Daily - PRN	Gas Relief	

List of 'ADMIT' Items - DO NOT FILL ITEMS BELOW (Patient has supply of item at admission)								
Start Date Stop Date Type	Order	Strength	Quantity	Dosage	Route	Frequency		Payer

ADDITIONAL NOTE

Merging of information from Consolo Services

PHONE ORDER / READ BACK ORDER BY MARY ROSE FRONDA (RN)

Electronically Signed: 10/17/2018, By STEPHEN DUBIN, MD

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JIREH HEALTHCARE
2320 PASEO DEL PRADO SUITE B101, LAS VEGAS, NV 89102
TEL: 702-359-1388

14/9/18 4 PM 5 peedy

DATE OF BIRTH:

RX NUMBER:

Penevcet 18/325 mg # 60

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fil 15 days supply

1CD: M54.5



n-	HOSPICE PATIENT"
PHYSICIAN SIGNATURE:	DEA#: _FD2419706
HYSICIAN NAME: <u>STEPHEN DU</u>	BIN DATE:
'HYSICIAN ADDRESS: <u>5380 s r.</u>	AINBOW STE 306 LAS VEGAS NV 89128
	PLEASE FAX BACK TO 702-776-7195

TRINITY PHARMACY
2797 S. MARYLAND PKWY., #28, LAS VEGAS, NV 89109 | TEL: 702-776-8210 | FAX: 702-776-7195

			11/26/18		
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JIREH HEALTHCARE

2320 PASEO DEL PRADO SUITE B101, LAS VEGAS, NV 89102

TEL: 702-359-1388

RX NUMBER:

DATE OF BIRTH:



"HOSPICE PATIENT"

PHYSICIAN SIGNATURE: PHYSICIAN NAME: STEPHEN DUBIN

DEA#: FD2419706

DATE:

PHYSICIAN ADDRESS: __5380 \$ RAINBOW STE 306 LAS VEGAS NV 89128

PLEASE FAX BACK TO 702-776-7195

TRINITY PHARMACY

2797 S. MARYLAND PKWY., #28, LAS VEGAS, NV 89109 | TEL: 702-776-8210 | FAX: 702-776-7195

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PHYSICIAN ADDRESS: 5380 S	RAINBOW STE 306 LAS VEGAS NV 89128	
PHYSICIAIN ADDRESS:	PLEASE FAX BACK TO 702-776-7195	
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	TRINITY PHARMACY	. 702-776-7195
2797 S. MARYLAND PKWY.,	#28, LAS VEGAS, NV 89109 TEL: 702-776-8210 FA	. 102-110-1133

JIREH HEALTHCARE 2320 PASEO DEL PRADO SUITE B101, LAS VEGAS, NV 89102 TEL: 702-359-1388

PATIENT NAME: RX NUMBER:

DATE OF BIRTH:

Perescet 10/325 #60

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"HOSPICE PATIENT"

PHYSICIAN SIGNATURE:

DEA#: FD2419706

PHYSICIAN NAME: _STEPHEN DUBIN

DATE:

PHYSICIAN ADDRESS: 5380 5 RAINBOW STE 306 LAS VEGAS NV 89128

PLEASE FAX BACK TO 702-776-7195

TRINITY PHARMACY

2797 S. MARYLAND PKWY., #28, LAS VEGAS, NV 89109 | TEL: 702-776-8210 | FAX: 702-776-7195

Patient Details for M. (Patient ID 501364, MRN M0474)					
Date Of Birth	Status Pending - Discharged				

Physician's Order Details

User

RN_Ruiz, Janyvill (iruiz)

Patient G M

Order Date 10/10/2018 Time of Event

Physician

Dubin, Stephen Paul

Nurse RN_Ruiz, Janyvill (jruiz) Oversight No

Order

Admit to Jirch Healthcare Services, Llc. for the 1st 90 day benefit period. Pharmacist may use generic drug equivalents or compound medications as appropriate. Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to: hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction machine, etc.) Assess and evaluate cardiac, respiratory, Gl.,GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs. Instruction on care of terminally Ill, medications, safety, nutritional needs, pain and symptom control based on assessment. Assess and evaluate current medications. May give nutritional supplements 1 can a day as tolerated for supplement. May crush medications that are crush-able. Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders. Hospice Medical Director may assist patient as needed to manage symptoms. Hospice nurse may pronounce death. Check oxygen saturation level prin every SN visit. Bed-rails up 2x. Skilled Nurse visits. (1x/week and 2x/week as needed for respiratory.& anxiety issues) to promote comfort and symptom management within certification period. CHHA visits 5x/week to assist with personal care, home making and to promote comfort per RN assignment within certification period. Chaplain visits. (1x/mo as needed for psychosocial issues) Volunteer Services continue to be offered every week per Skilled Nurse/Staff, Admit Patient to Jurch Healthcare Services with the following orders verified with M.D. Patient, DNR. Diet: As Tolerated. Admit with the following medications: Please see related medications.

Read Back

Non Verbal Order

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Nn

Medications

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SUMAtriptan Succinate 1 Tablet 2 times a day PO (50 MG Tablet)	take 1 tab by mouth 2 times per day at least 2 hours between doses		
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Oxybutynin Chloride ER 1 Tablet ER 24HR 1 times a day PO (5 MG Tablet ER 24HR)			
Omeprazole 1 Capsule DR 1 times a day PO As Needed Gas Relief (40 MG Capsule DR)			
Lidocaine 1 Patch every 12 hours EX (5 % Patch)			

Related Assignees

Social Services - MSW_Minnick, Joseph effective 10/17/2018 Chaplain - CHAPLAIN_Murphy, Kathryn effective 10/17/2018

Hospice Alde - Young, Grace Marie 5 visit(s) Weekly effective 10/17/2018

Skilled Nurse - RN_Cacanindin, Speedy effective 10/17/2018
Skilled Nurse - RN_Domondon, Mary Alyn effective 10/17/2018
MD Director - Dubin, Stephen Paul effective 10/17/2018

Physician - Dubin, Stephen Paul effective 10/17/2018

Signatures

- 1. Medical Director Dubin, Stephen Paul (Medical Director) signed on 10/18/2018. Recorded by jruiz on 10/17/2018 18:32:50.
- 2. Skilled Nurse RN_Rulz, Janyvill (User) signed on 10/17/2018. Recorded by jruiz on 10/17/2018 18:32:40

, M (Patient ID 501364, MRN M0474)

Date Of Birth

Status Pending - Discharged

Physician's Order Details

User

RN_Ruiz, Janyvill (jruiz)

Physician

Dubin, Stephen Paul



Nurse RN_Ruiz Janyvi (jruiz) Order Date 10/10/2018

Oversight No Time of Event

Orders

Admit to Jirch Healthcare Services, Ltc. for the 1st 90 day benefit period. Pharmacist may use generic drug equivalents or compound medications as appropriate. Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction machine, etc.) Assess and evaluate cardiac, respiratory, Gl.GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs. Instruction on care of terminally ill, medications, safety, nutritional needs, pain and symptom control based on assessment. Assess and evaluate current medications. May give nutritional supplements 1 can a day as tolerated for supplement. May crush medications that are crush-able. Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders. Hospice Medical Director may assist patient as needed to manage symptoms. Hospice nurse may pronounce death. Check oxygen saturation level pm every SN visit. Bed-rails up 2x. Sk. ed Nurse visits. (1x/week and 2x/week as needed for respiratory & anxiety Issues) to promote comfort and symptom management within certification period. CHHA visits 5x/week to assist with personal care, home making and to promote comfort per RN assignment within certification period. Chaplain visits. (1x/mo as needed for spiritual issues) "Social worker visits. (1x/mo as needed for psychosocial issues) Volunteer Services continue to be offered every week per Skilled Nurse/Staff. Admit Patient to Jireh Healthcare Services with the following orders verified with M.D. Patient: DNR. Diet. As Tolerated. Admit with the following medications.

Read Back

Non Verbal Order

Related Assignees

Social Services - MSW_Minnick, Joseph effective 10/17/2018 Chaptain - CHAPLAIN_Murphy, Kathryn effective 10/17/2018

Hospice Aide - Young, Grace Marie 5 visit(s) Weekly effective 10/17/2018

No

Skilled Nurse - RN_Cacanindin, Speedy offactive 10/17/2018
Skilled Nurse - RN_Domondon, Mary Alyn effective 10/17/2018
MD Director - Dubin, Stephen Paul effective 10/17/2018
Physician - Dubin, Stephen Paul effective 10/17/2018

Signatures

1 Medical Director Dublin, Stephen Paul (Medical Director) signed on 10/18/2018 Recorded by jrulz on 10/17/2018 18:32:22.

, M (Patient ID 501364, MRN M0474)

Date Of Birth

Status Pending - Discharged

DOB

Screened

Unscreened

Current Indicator

11/06/1962

Allergies

Allergies

E43 - Unspecified severe protein-

Not Assessed

See Screened Allergies

Pending Medications

Current Medications

Lidocaine [5 %] Patch (ID: 6634907) - 1 Patch every 12 hours EX (5 % Patch)

Order Date	10/17/2018	Ordering MD	Dubin, Stephen Paul
Admin By	Patient	Coverage	Covered
Change	Updated	Updated Date	10/17/2018

Omeprazole [40 MG] Capsule DR (ID: 6634894) - 1 Capsule DR 1 times a day PO As Needed Gas Relief

(40 MG Capsule DR)

Order Date	10/17/2018	Ordering MD		Dubin, Stephen Paul	
Admin By	Patient	Coverage	1.	Covered	
Change	Updated	Updated Date		10/17/2018	
Reasons	Gas Relief				

Oxybutynin Chloride ER [5 MG] Tablet ER 24HR (ID: 6634895) - 1 Tablet ER 24HR 1 times a day PO (5

MG Tablet ER 24HR)

Order Date	10/10/2018	Ordering MD	Dubin, Stephen Paul	
Admin By	Patient	Coverage	Covered	
Change	Updated	Updated Date	10/17/2018	. 3

Oxycodone-Acetaminophen [10-325 MG] Tablet (ID: 6634899) - 1 Tablet every 6 hours PO As Needed

Pain (10-325 MG Tablet)

Reasons	Pain	-	
Change	Updated	Updated Date	10/17/2018
Admin By	Patient	Coverage	Covered
Order Date	10/10/2018	Ordering MD	Dubin, Stephen Paul
Order Date	10/10/2018	Ordering MD	Dubin Stanban Bard

Al. unsolo Medications for



Sertraline HCI [100 MG] Tablet (ID: 6634901) - 1 Tablet 1 times a day PO (100 MG Tablet)

Order Date	10/10/2018	Ordering MD	Dubin, Stephen Paul	
Admin By	Patient	Coverage	Covered	
Change	Updated	Updated Date	10/17/2018	

SUMAtriptan Succinate [50 MG] Tablet (ID: 6634906) - 1 Tablet 2 times a day PO (50 MG Tablet)

Order Date	10/10/2018	Ordering MD	Dubln, Stephen Paul	
Admin By	Patient	Coverage	Covered	
Change	Updated	Updated Date	10/17/2018	
Additional Notes	take 1 tab by mouth 2	times per day at least 2 hours bet	ween doses	

M (Patient ID 501364, MRN M0474)

12

Status Pending - Discharged

DOB

Screened

Unscreened

Current Indicator

11/06/1962

Allergies

Allergies

E43 - Unspecified severe proteincalorie malnutrition

Not Assessed

See Screened Allergies

Pending Medications

Current Medications

Lidocaine [5 %] Patch (ID: 6634907) - 1 Patch every 12 hours EX (5 % Patch)

 Order Date
 10/17/2018
 Ordering MD
 Dubin, Stephen Paul

 Admin By
 Patient
 Coverage
 Covered

 Change
 Updated
 Updated Qate
 10/17/2018

Omeprazole [40 MG] Capsule DR (ID: 6634894) - 1 Capsule DR 1 times a day PO As Needed Gas Rellef

(40 MG Capsule DR)

Order Date 10/17/2018 Ordering MD Dubin, Stephen Paul
Admin By Patient Coverage Covered

Change Updated Updated Date 10/17/2018

Roasons Gas Retief

Oxybutynin Chloride ER [5 MG] Tablet ER 24HR (ID: 6634895) - 1 Tablet ER 24HR 1 times a day PO (5

MG Tablet ER 24HR)

 Order Date
 10/10/2018
 Ordering MD
 Dubin, Stephen Paul

 Admin By
 Patient
 Coverage
 Covered

 Change
 Updated
 Updated Date
 10/17/2018

Oxycodone-Acetaminophen [10-325 MG] Tablet (ID: 6634899) - 1 Tablet every 6 hours PO As Needed

Pain (10-325 MG Tablet)

Order Date 10/10/2018 Ordering MD Dubin, Stephen Pauli
Admin By Patient Coverage Covered

Change Updated Updated Date 10/17/2018

Reasons Pain

All Consolo Medications for



Sertraline HCI [100 MG] Tablet (ID: 6634901) - 1 Tablet 1 times a day PO (100 MG Tablet)

Order Date	10/10/2018	Ordering MD	Dubin, Stephen Paul
Admin By	Patient	Coverage	Covered
Change	Updated	Updated Date	10/17/2018

SUMAtriptan Succinate [50 MG] Tablet (ID: 6634906) - 1 Tablet 2 times a day PO (50 MG Tablet)

Order Date	10/10/2018	1	Ordering MD	Dubin, Stephen Paul
Admin By	Patient		Coverage	Covered
Change	Updated		Updated Date	10/17/2018
Additional Notes	take 1 tab by m	outh 2 times p	er day at least 2 hours between riosi	ns f

2320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102 . Tel: Fax:

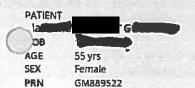
PHYSICIAN'S CERTIFICATION FOR HOSPICE BENEFIT

In order to determine if a patient is eligible for Hospice benefit we are required to have physician's authorization with a brief narrative supporting terminal illness

Cartification did.	CONTRACTOR OF THE PARTY OF THE	-	*	MR#:	M0474	
Certification #1:	Statement for First	90-day period				
I certify that	G M	has a life expectar	ncy of six months or less, if the to	erminal illness runs its normal cou	rse.	
Verbal Authorization Date	10/10/2018		Obtained by:	Mary Alyn Domondon/RN		
Effective Date of Certifica	tion: 10/10/2018 To	1/7/2019		Andrew Springer and Springer an		
Primary DX:	Unspecified severe pro	tein-calorie malnutrition (E	:43)	1 .		
Secondary/Related DX	Adult failure to thrive (F	(62.7)				-
Comorbidities	-					
Certification is based on:	✓ Medical history, reco	ord and patient status	✓ Team Assessment	Face to Face Encounter	Visit Date:	10/10/2018
ADL Assessment						
Ambulation	0	Transfer	0	Feeding	0	
Toileting	0	Dressing	0	Bathing	0	
Total ADL score	0	Total number of	factivities with complete depend	lence (0-6) 0 of 6 /	Activities	
Functional Status						
KPS		FAST				
PPS		NYHA		P		
LCD Determination	n Status:	-		1.00		
Physician Narrative:	Patient is at risk for falls an Ms. Of the is a 55-year-o fracture, sciatica, migraines and balance as well as sign	d lives in an apartment with d African American female major depressive disorde dificant weight loss. Patient	th brother. e with a terminal diagnosis of seer, adult failure to thrive. Patient thas moderate to severe generate.	nt with moderate to severé pain or vere protein calorie malnutrition. I will require maximum assistance alized pain; pain management will six months or less if terminal illnes	Patient with history o with ADL's and has continue. The patie	f traumatic a very poor gait
	care, and to the best of kno					e.
• 1 attest/confirm that I co	Attestation/Certific	ation	nedical record, team assessmen	t and/or examination of the patien		e.
● 1 attest/confirm that I co	Attestation/Certificomposed this narrative based of	ation on my review of patient's m	nedical record, leam assessmen Reviewed ar	nd electronically signed by STEPH		e. 10/10/2018
• 1 attest/confirm that I co	Attestation/Certific	ation on my review of patient's m	nedical record, leam assessmen Reviewed ar			e.
• 1 attest/confirm that I co	Attestation/Certificomposed this narrative based of	ation on my review of patient's m	nedical record, leam assessmen Reviewed ar	nd electronically signed by STEPH		e. 10/10/2018
• 1 attest/confirm that I co	Attestation/Certificomposed this narrative based of	ation on my review of patient's m	nedical record, leam assessmen Reviewed ar	nd electronically signed by STEPH		e. 10/10/2018
● 1 attest/confirm that I co	Attestation/Certificomposed this narrative based of	ation on my review of patient's m	nedical record, leam assessmen Reviewed ar	nd electronically signed by STEPH		e. 10/10/2018

CONFIDENTIALITY NOTICE: The information contained in this facsimile is confidential and may contain privileged material or be otherwise protected by applicable law. It is intended only for the use of the individual (s) or entity named above. If the person receiving this facsimile, or any other reader of the facsimile, is not the intended recipient or the employee or agent responsible for delivering it to the intended recipient, then any use, dissemination, distribution, or copying of this communication is strictly prohibited and may be subject to civil and/or criminal liability. If you have received this communication in error, please destroy it and notify the sender immediately.





FACILITY
RENATO RABARA Practice
T (702) 677-0561
9958 Corbridge st,
Las Vegas, NV 89178

ENCOUNTER

NOTE TYPE SOAP Note

SEEN BY RENATO RABARA

DATE 09/10/2018

AGE AT DOS 55 yrs

Electronically signed by RENATO

RABARA at 09/12/2018 07:08 am

Chief complaint

For Initial visit.

ADDRESS

Patient identifyl	ng details and den	-			
FIRST NAME	M	SEX DATE OF BIRTH	Female	ETHNICITY	Provider did not
MIDDLE NAME		DATE OF DEATH	•	PREF. LANGUAGE	
LAST NAME	G	PRN	GM88952Z	RACE	Provider did not
\$5N					ask
			2000	STATUS	Active patient
CONTACT INFORMA	TION		.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		4010011100114
ADDRESS LINE 1	MANAGELL	CONTACT BY	Mobile Phone		
		EMAIL.	-		
		HOME PHONE	•		
ADDRESS LINE 2	•	MOBILE PHONE	(702)		
CITY	Las Vegas	OFFICE PHONE			
STATE	NV	OFFICE	•		
ZIP CODE	89130	EXTENSION			مر
FAMILY INFORMATIO	ON				
NEXT OF KIN	•		PATIENT'S MOTHE	R'5 •	7
RELATION TO PATI	ENT -		MAIDEN NAME		
HONE				1	

PRIMARY PAYER			
PAYER	NV - Medicare B (MAC J1)	INSURED ID NUMBER	S41 76 7921-A
PRIORITY	Primary	GROUP NUMBER	
TYPE	Medicare	EMPLOYER NAME	
RELATIONSHIP TO INSURED	Self	INSURANCE PAYMENT TYPE	Copay
START DATE	09/10/2018	PAYMENT TYPE	Fixed
END DATE	-	COPAY AMOUNT	
		STATUS	Active

about:blank

DATE OF BIRTH

PAYMENT PREFERENCE
PATIENT'S RELATIONSHIP
TO GUARANTOR
SUARANTOR NAME
GUARANTOR ADDRESS

Primary Insurance

SEX SOCIAI PRIMA

SOCIAL SECURITY NUMBER PRIMARY PHONE NUMBER SECONDARY PHONE NUMBER

Vitals for this encounter	
	09/10/18 10:00 AM
Height	58 in
Welght	85 lb
Temperature == 7	97.50 ° ⊧ ′
Pulse	77 bpm
Respiratory rate	20 bpm
OZ Saturation	93 %
BMI	17.76
Blood pressure	106/82 mmHg

	Current Diagnoses	ACUITY	START	STOP
,	(Z87.81) Personal history of (healed) traumatic fracture	Chronic		
	(M84.4795) Pathological fracture, unspecified toe(s), sequela	Chronic		
	(M54,30) Sciatica, unspecified side	Chronic		
	(MS4.9) Dorsalgia, unspecified	Chronic	1	
	(R32) Unspecified urlnary incontinence	Chronic		1
_	(R25.9) Unspecified abnormalities of galt and mobility	Chronic		
	(G43.909) Migraine, unspecified, not intractable, without status migrainosus	Chronic		
	(G89.29) Other chronic paln	Chronic		
-	(F32.9) Major depressive disorder, single episode, unspecified	Chronic		
-	Historical Diagnoses No historical diagnoses	ACUITY	START	STOP
-				

Drug Allergies

DOR

PRN: 0M889622

ACTIVE ALLERGIES	SEVERITY/REACTIONS	ONSET
Patient has no known drug allergies		
Food Allergies		****
ACTIVE ALLERGIES	SEVERITY/REACTIONS	ONSET
No food allergies recorded		
		AND STANDS OF A CONTROL OF A CO
Environmental Allergies) = 141	26
ACTIVE ALLERGIES	SEVERITY/REACTIONS	ONSET
No environmental allergies recorded		

Active Medi	cations				مر
MEDICATION	100000000000000000000000000000000000000	SIG	START/STOP	ASSOCIATED DX	
Delayed Relea		take 1 capsule by mouth daily, PRN	•	•	May 4-12
exybutynin Ci	hloride (Oxybutynin MG Oral Tablet Extended	Take 1 tablet (5 mg) by mouth daily	ettisen rerit tradicionista.	and the second s	t belle folkerifter och sam serf. en e
	/ Acetaminophen cetaminophen) 10-325	tablet orally every 6 hours as needed	-	·	
Sertraline HCI	100 MG Oral Tablet	Take 1 tablet (100 mg) by mouth daily	-	•	
SUMAtriptan S Tablet	Succinate 50 MG Oral	Take 1 tablet (50 mg) by mouth 2 times per day at least 2 hours between doses as needed		•	Ε
	edications				
MEDICATION		SIG	START/STOP	ASSOCIATED DX	115-10
No historical n	nedications recorded				
mmunizatio	ns	***************************************	y 	A PROPERTY OF THE PROPERTY OF	والمراجع والمحاود والمراجع والم والمراجع والمراجع والمراجع والمراجع والمراجع والمراجع والمراج
DATE	VACCINE	SOURCE LOT	EXPIRES R	COMMENT	

ast medical history

about:blank

Page 3 of 6

MAJOR EVENTS

several surgeries due to MVA

FAMILY HEALTH HISTORY

DM

Hypertension

CA

Family health history

DIAGNOSIS

ONSET DATE

No Family health history recorded

Advance Directive

DIRECTIVE

ECORDED

No advance directives recorded for this patient.

Subjective

The patient is a 55 year old, female, African American who was seen for an Initial visit. The patient lives alone in an apartment. The patient stated that she is separated from her husband and has 2 independent living children. The patient was born in Spokane, Washington and ultimately moved to Las Vegas. The patient stated that she had several surgeries in the leg due to a MVA. The patient said that she became disabled due to the accident. The patient has an unsteady galt and uses a cane or cooter. The patient is also complaining of chronic pain and goes to a Pain Management Doctor for her pain issues. The patient is home bound and needs home visits because she is unable to leave home safely and needs assistive device due to abnormality of gait.

REVIEW OF SYSTEMS:.

Medications: Reviewed & Up-to-date

General: no fever, no fatigue, (+) reduction in weight, (+) change in appetite.

Head: No headaches, no dizziness

Eyes: no blurred vision, no eye pain.

Ears: No Otalgia, No change in hearing, no tinnitus, no discharge

Nose: no coryza, no stuffy nose, no discharge, no post nasal drip

Throat: No dental difficulties, no sore throat, no difficulty swallowing

Neck: No stiffness, no pain, no swollen glands

Breasts: No noted lumps, no tenderness, no swelling, no nipple discharge

Heart: No chest pains, no palpitations, no leg swelling.

Lung: No dyspnea, no wheezing, no cough

Abdomen: no abdominal pain, no nausea, no vomiting

Neurologic: No numbness/tingling, no weakness, no tremor, no seizures, no changes in memory, no ataxia

Extremitles: (+) pain in muscles or joints, (+) limitation of range of motion, no paresthesias or numbriess

Back! (+) back pain, no spasins,

Skin: no rashes, no lesions

GU: No frequency, no urgency, no dysuria, no hematuria.

Psychiatric: (+) depressive symptoms, no anxiety, no insomnia, no changes in thought content, no suicidal thoughts.

Objective

General: Normotensive, in no acute distress.

Head: Normocephalic, no lesions.

Eyes: PERRLA. EOM's full, conjunctivae clear,

Ears: EAC's clear, TM's normal.

Nose: Mucosa normal, no obstruction,

hroat: Clear, no exudates, no lesions.

Yeck: Supple, no masses, no thyromegaly, no bruits.

Chest: Lungs clear, no rales, no rhonchi, no wheezes,

about:bisnk

Page 4 of

Heart: RR, no murmurs, no rubs, no gallops.

Abdoment Soft, no tenderness, no masses, bowel sounds normal.

iU: Normal, no lesions, no discharge, no hemias noted.

Back: Normal curvature, no tenderness.

Extremities: (+) several scars, no edema, no enythema.

Neuro: Physiological, no localizing findings.

Skin: Normal, no rashes, no lesions noted.

Assessment

Diagnoses attached to this encounter:

Personal history of traumatic fracture [ICO-10: Z87.81], [ICD-9: V15.51], [SNOMED: 391095006]

Pathological fracture of toe [ICD-10: M84.479S], [SNOMED: 704168008]

Sciatica [ICD-10: M54.30], [ICD-9: 724.3], [SNOMED: 23056005]

Dorsalgla (ICD-10: M54.9], (ICD-9: 724.5), [SNOMED: 267981009]

Urine incontinence (ICO-10: R32), (ICD-9: 788.30), (SNOMED: 165232002)

Abnormality of galt (ICD-10: R26.9), [ICD-9: 781.2], [SNOMED: 22325002]

Migraine [ICD-10: G43.909], [ICD-9: 346.90], [SNOMED: 37796009]

Chronic pain [ICD-10: G89.29], [ICD-9: 338.29], [SNOMED: 82423001]

MDD [ICD-10: F32.9], [ICD-9: 296.20], [SNOMED: 370143000]

Plan

The visit was explained to the patient about the set up of the visiting provider. Primary care in the home by the nurse practitioner is being supervised by an MD. It was also explained to the patient thatch is electing King Rey to be her care provider and in an emergency situation the patient needs to call 911.

The patient verbalized understanding and consent was signed.

Labs will be ordered. (CBC, CMP, TSH with reflex free T4, HgbA1C, Lipid panel)

or HH evaluation, SN

atient will benefit from SN services for her overall health status, medications and disease process teachings.

SN to do regular check of blood pressure and log

Notify the provider if the BP is persistently <90/60 or >140/90

Continue current medications.

Low salt, Low fat dlet

Patient was advised on the risk of smoking.

Fall and safety precautions.

Follow up with her Pain Management Doctor for Pain Issues.

For follow up visit after 30 days...

Medications attached to this encounter:

Sertraline HCI 100 MG Oral Tablet 1 tablet (100 mg) orally daily

Oxycodone-Acetaminophen 10-325 MG Oral Tablet 1 tablet orally every 6 hours as needed

Oxybutynin Chloride ER 5 MG Oral Tablet Extended Release 24 Hour 1 tablet (5 mg) orally dally

Omephazole 40 MG Oral Capsule Delayed Release take 1 capsule by mouth daily, PRN

SUMAtriptan Succinate 50 MG Oral Tablet 1 tablet (50 mg) orally 2 times per day at least 2 hours between doses as needed

Orders

LAB ORDERS

No orders attached to this encounter.

IMAGING ORDERS

No orders attached to this encounter,

Screenings/interventions/assessments

lo screenings/interventions/assessments recorded.

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he practice rustors

JIREH HEALTHCARE SERVICES LLC (HOSPICE AGENCY)

PATIENT NAME: MENT H

DOB:

START OF CARE: 06/13/2018 TO 12/26/2018

Claim Number	From Service Date To	To Service Date	Beneficiary Nam Claim Status	To	Total Charge	Type Of Bill A	Admit Bate
21900700422207NVR	12/1/2018	12/31/2018	P-Paid	ST	79.067,78	813 - Hospice (9/7/2018	17/2018
21833900538807NVR	11/1/2018	11/30/2018	P-Paid	88	\$8,724,43	813 - Hospice (9/7/2018	/7/2018
21830901126607NVR	10/1/2018	10/31/2018	P - Paid	6\$	\$9,083,84	813 - Hospice (9/7/2018	17/2018
21827601222707NVR	9/7/2018	9/30/2018	P-Paid	Se	\$6,942.12	812-Hospice (9/7/2018	9102/17/
Claim Number	From Service-Date To Service Date	Fo Service Date	Beneficiary Nam Claim Status	Tota	Total Charge	Type Of Bill Adr	Admit Date
21900700421907NVR	12/1/2018	12/31/2018	P-Paid	86,9	56,986.61	813 - Hospice (10/10/2018	10/2018
21833900542507NVR		11/30/2018	P-Paid	58,1	\$8,197.48	813 - Hospice (10/10/2018	10/2018
21833900230804NVR	10/10/2018	10/31/2018	P - Paid	\$6,2	\$6,230.74	817 - Hospice (10/10/2018	10/2018
21830901121107NVR	10/10/2018	10/31/2018	P-Paid	\$5,6	\$5,630.74	812 - Hospice (10/10/2018	10/2018
Claim Number	From Service Date To	To Service Date	Service Date Beneficiary Nam Claim Status	Tota	Total Charge	Type Of Bill Adı	Admir Date
21900901108507NVR	12/1/2018	12/26/2018	Mark P-Paid	9,98	\$6,626.68	814 - Höspice (6/13/2018	3/2018
21833900542007NVR	11/1/2018	11/30/2018	Man Hammer P - Paid		\$7,635,75	813 - Hospice (6/13/2018	3/2018
21830901128607NVR	10/1/2018	10/31/2018	M. P. Paid	580	\$8,065.97	813 - Hospice (6/13/2018	3/2018
21827601223807NVR	9/1/2018	9/30/2018	M.P. Paid		87,382.9]	813 - Hospice (6/13/2018	3/2018
21824701185107NVR	8/1/2018	8/31/2018	M H P - Paid	\$8,	\$8,276.79	813 - Hospice (6/13/2018	3/2018
21821800581907NVR	7/1/2018	7/31/2018	MAN H. P. Paid	768	89,216.50	813 - Hospice (6/13/2018	3/2018
21818600817807NVR	6/13/2018	6/30/2018	Manney P - Paid	SS	\$5,126.72	812 - Hospice (6/13/2018	13/2018
Claim/Number	From Service Date	To Service Date	Beneficiary Nam Claim Status	Total	Total Charge	Type Of Bill Admi	Admit Date
	1000	otocicact	o d	\$680.54	54	811 - Hospice (12/20/2018	0/2018

										14-
	144.0	m1 1							*	
12/26/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
12/25/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
12/24/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
12/23/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare (178.2523				0
12/22/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
12/21/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
12/20/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
12/19/2018 Jireh Healthcare Servic M0456		Clark		ROUTINE	Medicare (178.2523				
	144.9									0
12/18/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare (178.2523				0
12/17/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		178.2523				0
12/16/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare (178.2523				0
12/15/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
12/14/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
12/13/2018 Jireh Healthcare Servic M0456	J44.9	Clark	1 1	ROUTINE	Medicare I	178.2523				0
12/12/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
12/11/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
12/10/2018 Jireh Healthcare Servic M0456	J44.9	Clark	- 6	ROUTINE	Medicare	178.2523				0
12/9/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare 4	178.2523				0
	144.9					178.2523				
12/8/2018 Jireh Healthcare Servic M0456		Clark		ROUTINE	Medicare					0
12/7/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
12/6/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		178.2523				0
12/5/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
12/4/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
12/3/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
12/2/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
12/1/2018 Jirch Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
11/30/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
11/29/2018 Jireh Healthcare Servic M04S6	144.9	Clark		ROUTINE	Medicare	178.2523				0
11/28/2018 Jireh Healthcare Servic M0456		Clark		ROUTINE						
	J44.9				Medicare	178.2523				0
11/27/2018 Jireh Healthcare Servic M0456	J44 9	Clark		ROUTINE	Medicare	178.2523				0
11/26/2018 Jireh Healthcare Servic M0456	J44 9	Clark		ROUTINE		178.2523				0
11/25/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
11/24/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare .	178.2523				0
11/23/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
11/22/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Meditare	178.2523				0
11/21/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare (178.2523				0
11/20/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare .	178.2523				0
11/19/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare *4	178.2523				0
11/18/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178 2523	. 4			0
11/17/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare					0
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11/16/2018 Jirch Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523	4 3			0
11/15/2018 Jireh Healthcare Servic M0456	144,9	Clark		ROUTINE	Medicare -	178.2523				0
11/14/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare (178,2523				0
11/13/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
11/12/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare (178.2523				0
11/11/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178 2523				0
11/10/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178,2523				0
11/9/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare (178.2523				0
11/8/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178,2523				0
11/7/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
11/6/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
11/5/2018 Jireh Healthcare Servic M0456	144.9									0
11/4/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
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11/3/2018 Jireh Healthcare Servic M0455	J44.9	Clark		ROUTINE	Medicare I	178.2523				0
11/2/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523		,		D
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10/31/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare (178.2523				0
10/30/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
10/29/2018 Jireh Healthcare Servic M04S6	144.9	Clark		ROUTINE	Medicare (178.2523				0
10/28/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
10/27/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
10/26/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178.2523				0
10/25/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
10/24/2018 Jireh Healthcare Servic M0456				ROUTINE						
	J44.9	Clark			Medicare (178.2523				0
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10/22/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
10/21/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178 2523				0
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10/18/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178 2523				0
10/17/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare (178-2523				0
10/16/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178-2523				0
10/15/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
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10/11/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE		178.2523				0
10/10/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	178 2523				0
10/9/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178 2523				0
10/8/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
10/7/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178 2523				0
10/6/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
10/5/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178 2523				0
10/4/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		178 2523				0
10/3/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	178.2523				0
10/2/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		178.2523				0
10/1/2018 Ilreh Healthcare Servic M0456	J44.9	Clark		ROUTINE		178.2523				0
9/30/2018 Jireh Healthcare Servic M0456	144.9	Clark			Medicare	173.2251				0
9/29/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		173 2251				0
9/28/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	-	173 2251				0
9/27/2018 Jireh Healthcare Servic M0456	J44.9	Clark	*	ROUTINE		173 2251 173 2251				0
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9/24/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		173 2251				0
9/23/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		173 2251				0
9/22/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		173 2251				0
9/21/2018 Jirch Healthcare Servic M0456	144.9	Clark		ROUTINE		173 2251				0
9/20/2018 Jireh Healthcare Servic M0456	J44.9	Clark			Medicare	173.2251				0
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9/18/2018 Jiréh Healthcare Servic M0456	144.9	Clark			Medicare	173 2251				0
9/17/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE		173 2251				0
9/16/2018 Jireh Healthcare Servic M0456	144.9	Clark			Medicare	173 2251				0
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9/14/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		173.2251				0
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9/12/2018 Jireh Healthcare Servic M04S6	144.9	Clark		ROUTINE	Medicare	173,2251				0
9/11/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	173.2251				0
9/10/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	Medicare	173.2251				0
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9/6/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE		173 2251				0
9/5/2018 Jireh Healthcare Servic M0456	144.9	Clark			Medicare	173 2251				0
9/4/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		173 2251				0
9/3/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE		173 2251				0
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8/29/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE		173 2251				0
8/28/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE		173 2251				0
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8/26/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		173 2251				0
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8/16/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		173 2251				0
8/15/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		173 2251				0
8/14/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE	Medicare	173 2251				0
8/13/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		173 2251				0
8/12/2018 Jireh Healthcare Servic M0456	J44.9	Clark		ROUTINE	_					0
8/11/2018 Jirch Healthcare Servic M0456	J44.9	Clark		ROUTINE		220.5569				0
8/10/2018 Jirch Healthcare Servic M0456	J44.9	Clark		ROUTINE		220.5569				0
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8/5/2018 Jireh Healthcare Servic M0456	144.9	Clark		ROUTINE		220.5569				0
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7/1/2018 Jireh Healthcare Servic M0456	144.9	Clark			ROUTINE	Medicare	220.5569	
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CITY	Las Ve	gas	, , , , , , , , , , , , , , , , , , , ,	STATE		NV ,	3	No. 1470 of all and address of goods above	ZIP	89121	
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Dates of Serv	ice, Dia	gnosis & A	llergies		- '	*					
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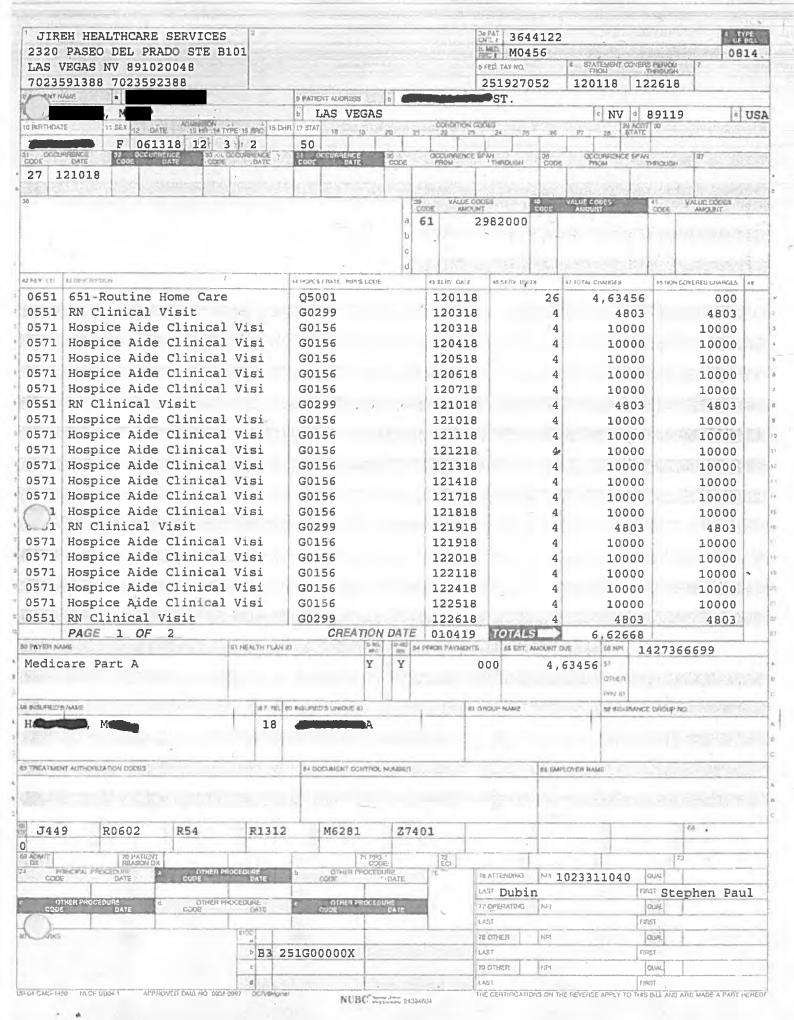




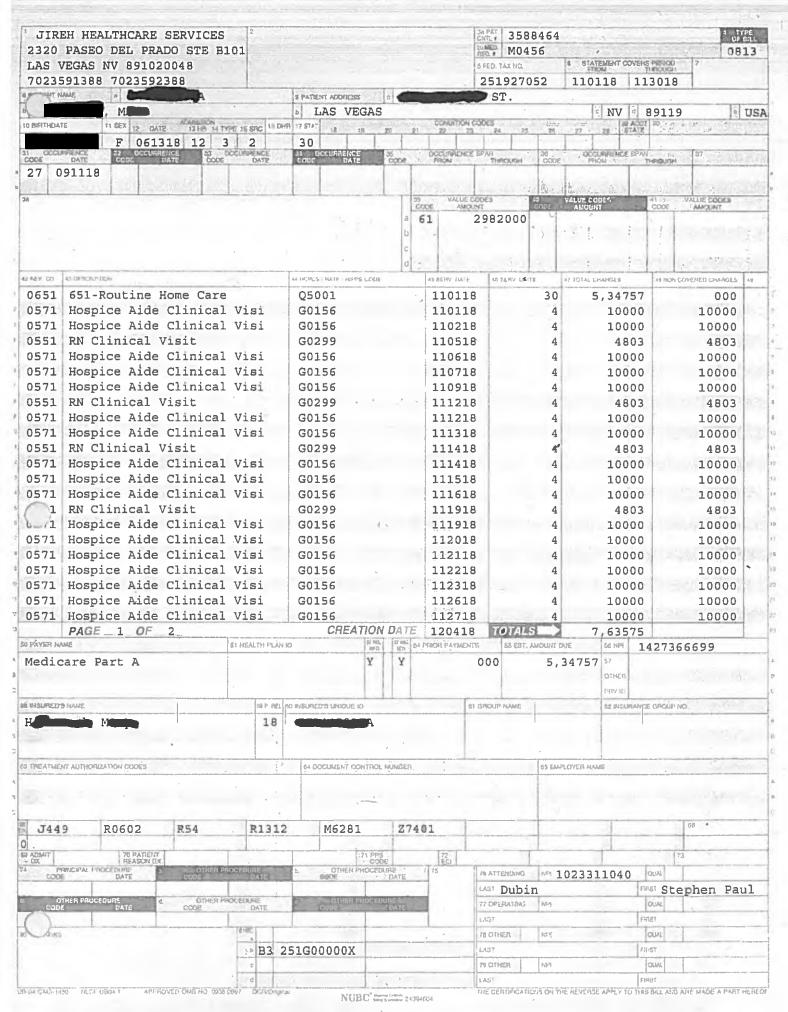
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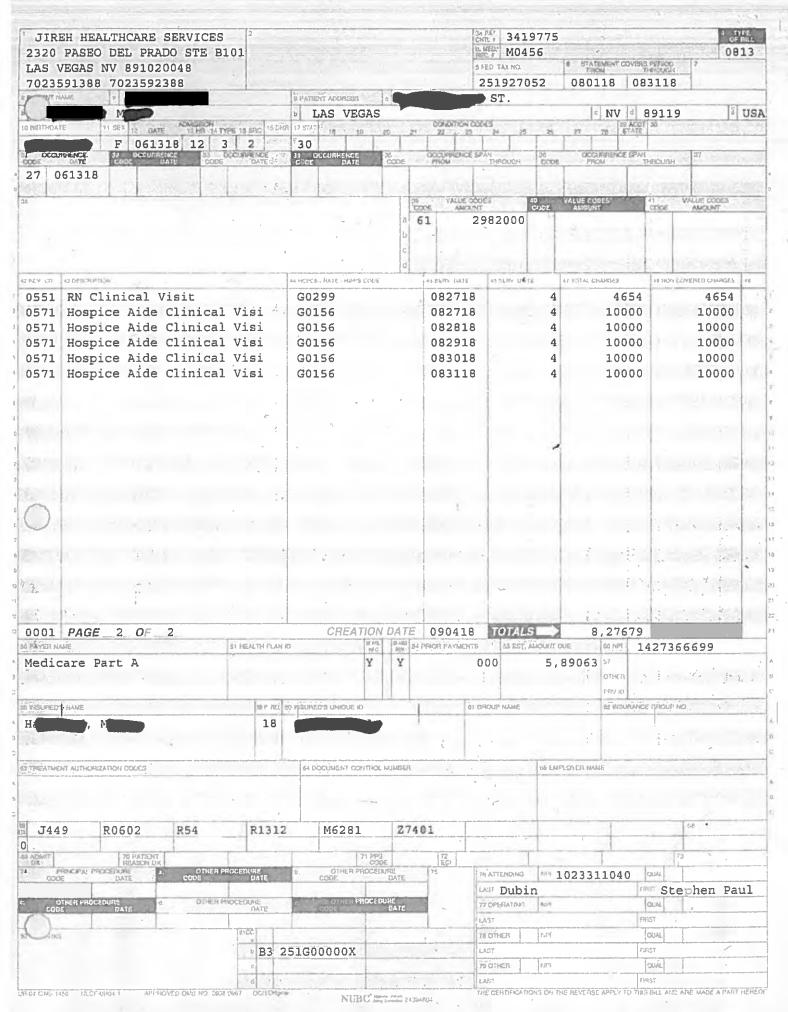


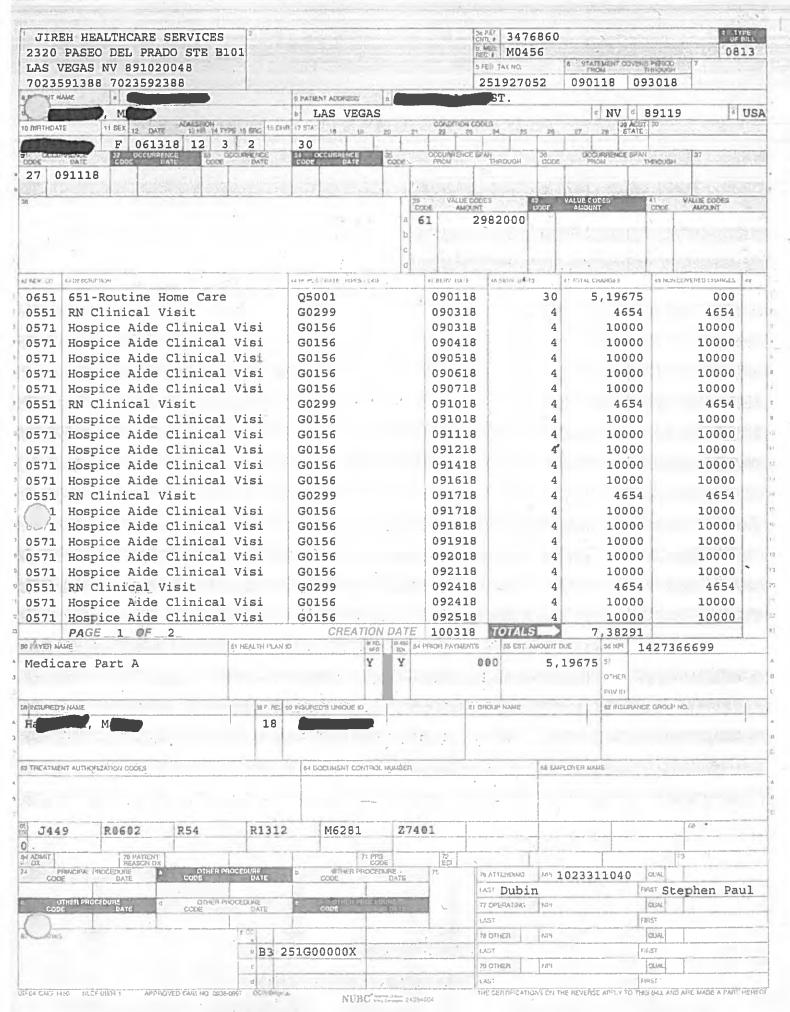
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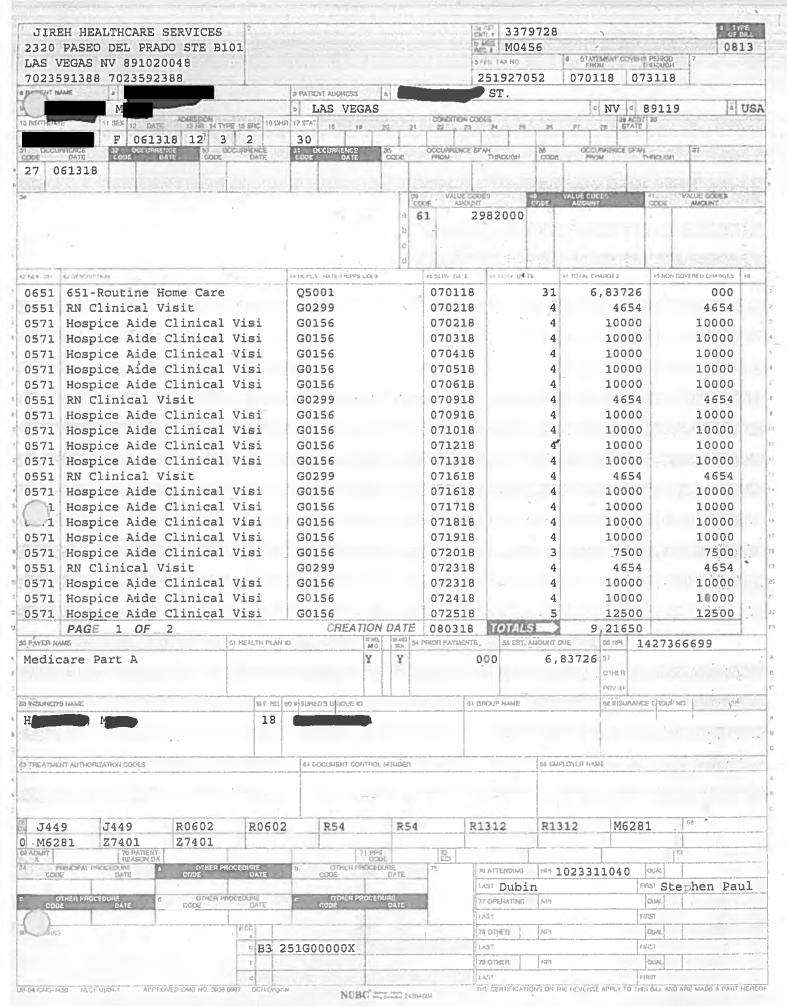
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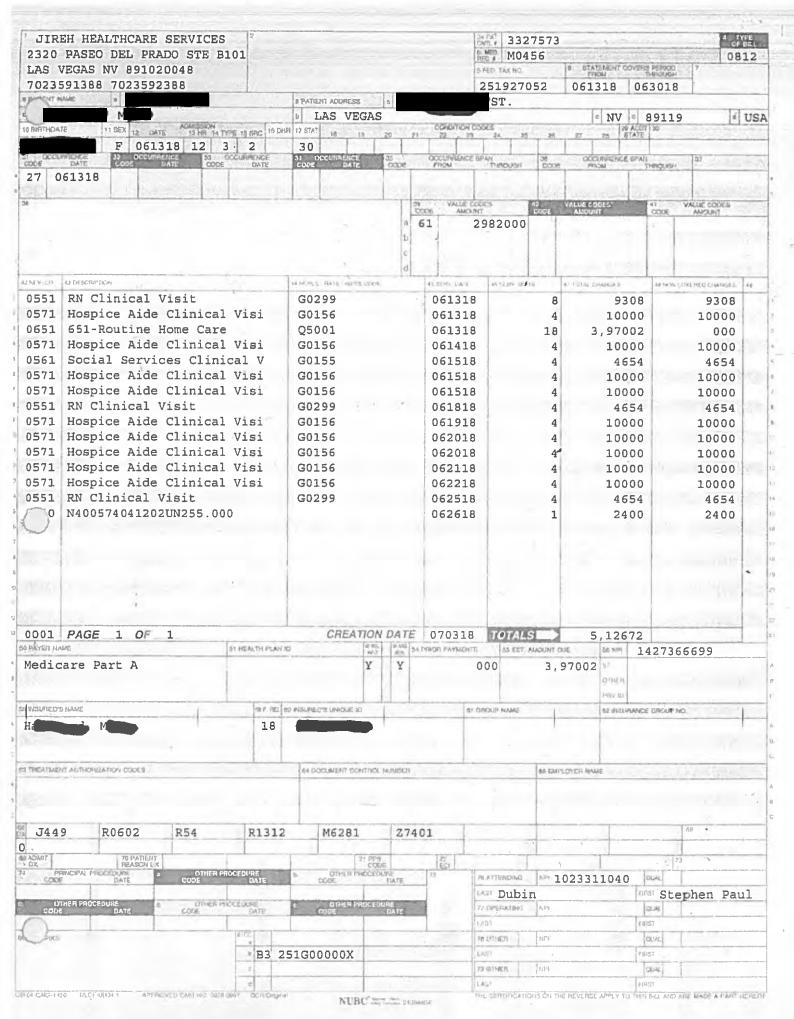




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2320 Paseo Del Prado, Suite B101, Las Vegas. NV 89102 . Tel: Fax:

Discharge Summary

PATIENT: H	Majo	in designation of the contract	apagetting gaing alleg a shape miles. Per some entertaintead despise filmati templologiste on plates a gift, entertain	MR#:	M0456
Këason For Discharge:	Transferred to Anothe	er Hospice			
EOC:	12/26/2018				
Based on the type of E extent to which goals	Discharge provide a b were and/or were not	rief summary related to; Dia met, and if applicable, sum	ignosis, health history, syn marize post-discharge con	nptom and p tinuity of ca	pain management, re orders.
Ms. Halstead transferred to	another hospice due to p	atient daughter decision to transi	er her mom to another hospice.	ratio for the Collection of Administrative Administration to	egilmiga ven versegarut versen, sp. tills till dillation hallt. Samman versegare som versegar i value som
Transfer forms checked	below sent to Receivin	g Agency	Attached Managery of	ter and de	elmen sprong, reported generalize spacks, mile hillineau sillenturalization proprieta elphonolo uniformità statuto della con-
POC Summary w/ List o	of Medications		_ ′		
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CONFIDENTIALITY NOTICE: The information contained in this fecsimile is confidential and may contain privileged material or be otherwise protected by applicable law. It is intended only for the use of the individual (s) or entity named above. If the person receiving this facsimile, or any other reader of the facsimile, is not the intended recipient or the employee or agent responsible for delivering it to the intended recipient, then any use, dissemination, distribution, or copying of this communication is strictly prohibited and may be subject to civil and/or criminal liability. If you have received this communication in error, please destroy it and notify the sender immediately.

PHYSICIANS ORDER

2320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102 . Tel. Fax:

IENT:

data Man

DOB:

ORDER DATE:

11/16/18

MR#:

M0456

ORDER#: 1

1600755

List of New/Refill Orders

ORDERED BY: STEPHEN DUBIN (MD)

Start Date	Stop Date	Туре	1	Order	Strength	Quantity	Dosage	Route	Frequency	Indication	Payer
11/16/18	described brown	Medication	New	Lorazepam (Ativan)	0.5 mg	4	1 tab	Oral	3x daily as needed	Anxiety	4
11/16/18		Medication	New	Furosemide (Furosemide)	20 mg		1 Tab	ORAL	once daily		
11/16/18		Medication	New	Klor-Con M (Potassium Chloride)	20 meq		Tab ER	ORAL	once daily		
11/16/18	amanan dina menangkan dinakan me	Medication	New	Ipratropium Bromide and Albuterol Sulfate 0.5/3 mg/3mL (Ipratropium Bromide and Albuterol Sulfate)	vial	1.0	1 Sol	RESPIRATORY (INHALATION)	3x daily as needed	SOB	

List of 'ADMIT' Items - DO NOT FILL ITEMS BELOW (Patient has supply of item at admission)

Start Date Stop Date Type Order Strength Quantity Dosage Route Frequency Indication Payer

ADDITIONAL NOTE

Ms Man Hamilla is an 85 year old, Caucasian female, with COPD on intermittent oxygen inhalation, hypertension, age related physical debility, polyneuropathy, has fracture on unspecified part of beck of left femur, chronic pain, dementia, muscle weakness. Daughter notified office of increasing anxiety, SOB and lower extremity edema.

SN performed vital signs and all body system assessments.

tient alert, pleasant, anxious; sitting comfortably in wheelchair. Complains of anxiety especially during the night, "could not sleep because I am scared".

- heart: regular heart rate
- respiratory: decreased breath sounds, no wheezing, no rales
- extremities: edema of LE, bilateral +3
- notified MD with orders given

PHONE ORDER / READ BACK ORDER BY JULIA DUGAY (RN)

Electronically Signed: 11/16/2018, By STEPHEN DUBIN, MD

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PHYSICIANS ORDER

2320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102, Tel: Fax:

IENT: M

DOB:

09/18/18

MR#:

M0456

1598477 ORDER#:

List of New/Refill Orders

ORDERED BY: STEPHEN DUBIN (MD)

Indication Payer Frequency Dosage Route Start Date Stop Date Type Order Strength Quantity ORAL. Daily Vitamin D 2000 units 2 Tab Medication New 09/18/18 ==> Note: Take one tablet by mouth once daily. 2 bottles Oral Daily - PRN Supplement Medication New Ensure 09/18/18

ORDER DATE:

List of 'ADMIT' Items - DO NOT FILL ITEMS BELOW (Patient has supply of item at admission)

Strength Frequency Indication Payer Order Quantity Dosage Route Stop Date Type Start Date

ADDITIONAL NOTE

Merging information from Consolo Services.

PHONE ORDER / READ BACK ORDER BY MARY ROSE FRONDA (RN)

Electronically Signed: 9/18/2018, By STEPHEN DUBIN, MD

PHYSICIANS ORDER

2320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102 . Tel: Fax:

IENT: HOUSE MAN

ORDERED BY: STEPHEN DUBIN (MD)

DOB:

ORDER DATE:

09/17/18

MR#:

M0456

ORDER#:

1598417

List of New/Refill Orders

Start Date	Stop Date	Туре		Order	Strength	Quantity	Dosage	Route	Frequency	Indication	Payer
09/17/18	3	Medication	New	Albuterol Sulfate (Albuterol Sulfate)	2.5 mg/3mL		1 Sol	RESPIRATORY (INHALATION)	2 times a day	SOB	
09/17/18	3	Medication	New	Sertraline Hydrochloride (Sertraline	50 mg		2 Tab	ORAL	Daily		-

==> Note: Take two tablets by mouth once daily

List of 'ADMIT' Items - DO NOT FIL	. ITEMS BELOW (Patient has supply of item at admission)
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Start Date Stop Date Type Order Strength Quantity Dosage Route Frequency Indication Payer

ADDITIONAL NOTE

Merging information from Consolo Services.

PHONE ORDER / READ BACK ORDER BY MARY ROSE FRONDA (RN)

Electronically Signed: 9/17/2018, By STEPHEN DUBIN, MD

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PHYSICIANS ORDER

320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102 . Tel: Fax.

PATIENT:

M

ORDERED BY: STEPHEN DUBIN

DOB:

ORDER DATE: 11/16/18

MR#:

M0456

ORDER#: 1600755

List of New/Refill Orders

Start Date	Stop Date	Туре		Order	Strength	Quantity	Dosage	Route	Frequency	Indication	Payer
11/16/18		Medication	New	Lorazepam (Ativan)	0.5 mg		1 tab	Oral	3x daily as needed	Anxiety_	
11/16/18		Medication	New	Furosemide (Furosemide)	20 mg		1 Tab	ORAL	once dally		
11/16/18		Medication	New	Klor-Con M (Potassium Chloride)	20 meq		Tab ER	ORAL	once dally		
11/16/18		Medication	New	Ipratropium Bromide and Albuterol Sulfate 0.5/3 mg/3ml. (Ipratropium Bromide and Albuterol Sulfate)	vial		1 Sol	RESPIRATORY (INHALATION)	3x daily as needed	SOB	

List of 'ADMIT' Items - DO NOT FILL ITEMS BELOW (Patient has supply of item at admission)

Start Date Stop Date Type Strength Quantity Dosage Route Frequency Indication Payer

ADDITIONAL NOTE

Ms Mary Halstead is an 85 year old, Caucasian female, with COPD on intermittent oxygen inhalation, hypertension, age related physical debility, polyneuropathy, has fracture on unspecified part of beck of left femur, chronic pain, dementia, muscle weakness. Daughter notified office of increasing anxiety, SOB and lower extremity edema.

- SN performed vital signs and all body system assessments.

- Patient alert, pleasant, anxious, sitting comfortably in wheelchair. Complains of anxiety especially during the night, "could not sleep because I am scared". - VS wnl:

- heart, regular heart rate

- respiratory; decreased breath sounds, no wheezing, no rales

- extremities: edema of LE, bilateral +3

- notified MD with orders given

PHONE ORDER / READ BACK ORDER BY JULIA DUGAY (RN)

MD Signature:

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PHYSICIANS ORDER

2320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102 . Tel Fax:

TIENT: Manager Manager DOB:

MR#:

M0456

ORDERED BY: STEPHEN DUBIN (MD)

ORDER DATE: 08/21/18 ORDER#:

1598450

List of New/Refill Orders

Indication Payer Frequency Dosage Route Strength Quantity Order Start Date Stop Date Type TOPICAL Daily Calmoseptine 20.6/ .44 1 Oint 08/21/18 Medication g00g (Zinc Oxide and menthol) ==> Note: Apply to affected area every diaper change SOB RESPIRATORY Every 6 hrs -**VENTOLINHFA HFA** 108 mcg 2 Aerosol Medication New 08/21/18 (INHALATION) PRN

==> Note: Take two puffs every 6 hours as needed for shortness of breath,

(albuterol sulfate)

List of 'ADMIT' Items - DO NOT FILL ITEMS BELOW (Patient has supply of item at admission)

Payer Indication Route Frequency Quantity Dosage Stop Date Order Strength Start Date

PHONE ORDER / READ BACK ORDER BY MARY ROSE FRONDA (RN)

Electronically Signed: 8/21/2018, By STEPHEN DUBIN, MD

PHYSICIANS ORDER

2320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102 . Tel. Fax:

CIENT:

Harmy Man

DOB:

ORDER DATE:

07/03/18

MR#:

M0456

ORDER#: 1

1598543

List of New/Refill Orders

ORDERED BY: STEPHEN DUBIN (MD)

Start Date Stop Date Type

Order

Strength Quantity

Dosage

Route

Frequency

Indication

Payer

Payer

07/03/18

Medication New

MiraLAX (Polyethylene Glycol 3350)

17 g7g

1 Powder

ORAL

L Daily - PRN

Constipation

,

==> Note, Mix 17 gram to 8 oz glass of water. Taken once daily by mouth as needed for constipation. To administer if no bowel movement for 3 days.

List of 'ADMIT' Items - DO NOT FILL ITEMS BELOW (Patient has supply of item at admission)

Start Date Stop Date Type Order Strength Quantity Desage Route Frequency Indication

ADDITIONAL NOTE

Merging of information from Consolo Services.

PHONE ORDER / READ BACK ORDER BY MARY ROSE FRONDA.(RN)

Electronically Signed: 7/3/2018, By STEPHEN DUBIN, MD

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Trinity Pharmacy
2797 S. Maryland Pkwy 28 Las Vegas, NV 89109
Phone: 702-776-8210 Fax: 702-776-7195 **Telephoned Prescription** STEPHEN DUBIN 5380 S RAINBOW STE 306 DEA#: FD2419708 Phone : 702-359-1388 Fax : 702-362-9954 Lig# : 13772 NPI# : 1023311040 LAS VEGAS NV 89128 Rx#: 170055 Rx Written: 12-19-2018 AVE (JIREH) LAS VEGAS NV 89121 702-4 Female LORAZEPAM 0,5MG TABLET Generic for ATIVAN TAB 0,5MG 100 Quantity: 45 Sig instructions: TAKE ONE TABLET BY MOUTH THREE TIMES DAILY AS NEEDED

Patient Address

Phone # D.O.B.

Gender

Refills Authorized : 0 0 Signature

THIS PRESCRIPTION WILL BE FILLED GENERICALLY UNLESS PRESCRIBER WRITES "DAW" IN THE BOX BELOW

AVE (AREA) LAS VEGAS NV 89121

Trinity Pharmacy
2797 S. Maryland Pkwy 28 Las Vegas, NV 89109

Phone: 702-776-8210 Fax: 702-776-7195

Telephoned Prescription



STEPHEN DUBIN

5380 S RAINBOW STE 306

LAS VEGAS NV 89128

NPI#: 1023311040

Rx Written: 11-16-2018

Lic# : 13772

DEA #: FD2419706

Rx#: 167999

Phone: 702-359-1388

: 702-362-9954

Patient Address

LAS VEGAS NV 89121

Phone #

D.O.B.

Gender

Female

LORAZEPAM 0.5MG TABLET

Generic for ATIVAN TAB 0.5MG 100

Quantity: 45

Sig Instructions:

TAKE ONE TABLET BY MOUTH THREE TIMES DAILY

Refills Authorized: 0

RPh: MLJ/CC

a a'. war

Signature

THIS PRESCRIPTION WILL BE FILLED GENERICALLY UNLESS PRESCRIBER WRITES "DAW" IN THE BOX BELOW

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Dispense As vynuen

Page _____ of __

Patient Name;

Hospice Medical Director or Designee Name (Printed):

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DUBIN0126

M **BENEFIT PERIOD** 60-Day Period from Face to Face Encounter (Note: Must be completed no more than Hospice Nurse Practitioner Attestation: I confirm that I had a face-to-face on /__/___(Date) and that the clinical findings of that encounter h continued eligibility for hospice care. NP Name (Printed): NP Signat Physician Attestation: I confirm that I had a face-to-face encounter with ... __/____(Date) and that I used the clinical findings of that enco Hospice Medical Director or Designee Name (Printed): Hospice M STEPHEN DUBIN, MD Recertification Statement: I recertify that I have reviewed the clinical is still considered to be terminally ill and has a life expectancy of six (6) mon Physician Brief Narrative Statement (Note: Must be completed no Review the patient's clinical circumstances and synthesize the medical infor Attestation: I confirm that I composed this narrative based on my review of Hospice Medical Director or Designee Name (Printed): Hospice N DUBIN, MD STEPHEN BENEFIT PERIOD 60-Day Period from Face to Face Encounter (Note: Must be completed no more than Hospice Nurse Practitioner Attestation: I confirm that I had a face-to-face __/____/ _____/ Date) and that the clinical findings of that encounter h continued eligibility for hospice care NP Name (Printing): NP Signal Physician Attestation: I confirm that I had a face-to-face encounter with (Data) and that I used the clinical findings of that enco Hospice Medical Director or Designee Name (Printed) Recertification Statement: I recertify that I have reviewed the clinical is still considered to be terminally ill and has a life expectancy of six (6) mor Physician Brief Narrative Statement (Note Must be completed no Review the patient's clinical circumstances and synthesize the medical infor

Attestation: I confirm that I composed this narrative based on my review of the patient's medical record and/or examination of the patient.

Hospice Medical Director or Designee Signature

JI 3H HEALTHCARE SERVIL 5

2320 Paseo Del Prado Suite B101 Las Vegas Nevada 89102 PHONE: 702,359,1388 FAX: 702,359,2388 EMAIL ADDRESS: jirehhealthcare @gmail.com

PHYSICIAN'S CERTIFICATION FOR MEDICARE/MEDICAID HOSPICE BENEFIT

Physician's Certification of Terminal Illness for Hospice Benefit
(Part - 1)

		MR# MO	156
CERTIFICATION STATEMENT First 90-Day Period from	6/13/2018	to 9/10/2	613
I (or WE) certify that Has a life expectancy of six (6) months or less if the terminal illnes Reviewed the patient's clinical information and considered the prin diagnoses, current subjective and objective medical findings, curren orders and information about management of unrelated conditions	ss runs its norm nary terminal o	is terminally all course. I (We condition, relation treatment	ill ar /E) ed
a. Pula		6/13/2014	
RN Signature	Ver	6 3 2017 bal Order Date	3
		6/13/2018 te 6/13/2018	
Hospice Medical Director/Associate Hospice Doctor	Da	te	
		6/13/2018	
Attending Physician Signature	Da	te	
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RECERTIFICATION STATEMENT Second 90-Day Period I certify that I have reviewed the clinical record prior to recertificat That patient is still considered to be terminally ill and has life expect The terminal illness runs its normal course.	tion for the abo	ve noted patier	กโลก
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JIREH HEATHCARE SERVICES

2320 Paseo Del Prado Sulte B101 Las Vegas, Nevada 89102 Tel 702.359.1388 Fax 702.359.2388

Patient Instructions, Narrative, Initial IDT Record and Initial Plan of Care

(Write a history of illness, how patient appears today, and reaction to hospice program)

THE FOLLOWING INFORMATION WAS REPORTED TO THE IDT, AND WAS CONSIDERED WHEN DEVELOPING THE PLAN OF CARE.

Pt. is an 85 year old, Caucasian female, with disabling dyspnea, hypoxemic on room air, COPD, hypertension, age related physical debility, polyneuropathy, has fracture on unspecified part of beck of left femur, chronic pain, dementia, GERD, muscle weakness, dysphagia, with other symbolic dysfunction. Pt. is currently bed bound, and has very poor appetite. Needs maximum to complete assistance on all her ADL's

MR#: M0456

Patient Name:

NARRATIVE:

They are also aware if pat original coverage. Medicate which includes information Advance Directives, and the changes in condition, problem.	i, verbalized by Fitient chooses to ions & Hospice in on palliative in the "Medicare Holems, or question port given, order	'atient & Family revoke Hospice 24 hour on call nedical care gui pspice Benefits'' s. Report given to admit to ho	They are awa, a revocation a service, review delines, planni, reviewed & le to Hospice Med	ed. Understanding of all hospice admission forms, & the re patient can choose not to continue Hospice at any time, statement must be signed & he/she will revert to his/her red, understanding verbalized. Hospice admissions packet, ng for death, feeding concerns, State Hotline no., DNR, eft with patient. They will call Hospice if they have any dical Director, admission approved by IDT, Telephone call olement palliative care protocols, & approval for hospice
Interdisciplinar	y Team Sign	atures	√	MEDICATIONS REVIEWED
Medical Director	Date:	Time:	1	TREATMENTS REVIEWED
Signature: Registered Nurse Signature:	- 6/13/18 Date:	Time:	Director, Patient concurs with Init	s, Medical Director, Medical Social Worker, Skilled Nurse, Chaplain Care Service, Volunteer were informed about Patient Admission and tial Plan of Care per telephone conversation. FREQUENCY FOR VISITS: Eval, 1x/week and 2x/week as needed for respiratory issues within certification period
Psychosocial Counselor	Date	Time:	MSW:	1x/month as needed for psychosocial issues within certification period
Signature: Spiritual Counseler	Daile:	SW 6	3)(PC:	Eval, */mo and */mo PRN for spiritual issues within certification period 5x/wk
Signature: RMM	rhy . Cl	naplain	6/10/18 Vol:	Refused
СННА	Date.	Time:	OTHER:	N/A
Signature:	4-13-18		RN Signature:	a. Rus
Other /	Date:	Time:	DATE:	0 6/13/18
Signature:				, - 1, -

TIME:

Consolo Physician Order for HALxM at Office Jireh Healthcare Services LLC

Patient Details for Hamman Man (Patient ID 47	70362, MRN M0456)
Date Of Birth	Status Active
	,

Physician's Order Details

User

RN_Esteban, Randy (Reste)

Physician Dubin, Stephen Paul Patient H

> <u>irse</u> RN_Esteban, Randy

Order Date 06/13/2018 Time of Event

Oversight No

Orders

Admit to Jirch Healthcare Services, LLC, for the 1st 90 day benefit period

(Reste)

- * Pharmacist may use generic drug equivalents or compound medications as appropriate
- * Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to: hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction machine, etc.)
- * Assess and evaluate cardiac, respiratory, GI, GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs
- * Instruction on care of terminally ill, medications, safety, nutritional needs, pain and symptom control based on assessment
- * Assess and evaluate current medications
- * Administer oxygen via nasal cannula at 2.5 lpm as needed for shortness of breath or until oxygen saturation is 95% and above
- * May give nutritional supplements 1 can a day as tolerated for supplement
- * May crush medications that are crush-able.
- * Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders
- * Hospice Medical Director may assist patient as needed to manage symptoms
- * Hospice nurse may pronounce death
- * Check oxygen saturation level every SN visit
- * Bed-rails up 2x
- * Skilled Nurse visits (Eval, 1x/week and 2x/week as needed for respiratory issues within certification period) to promote comfort and symptom management.
- * CHHA visits (5x/week within certification period) to assist with personal care, home making and to promote comfort per RN assignment.
- * MSW visits (1x/month as needed for psychosocial issues within certification period) to provide psychosocial support to patient and family.
- * Chaplain visits (1x/month as needed for spiritual issues within certification period) to provide spiritual

	· · · · · · · · · · · · · · · · · · ·
Read Back	Non Verbal Order
Yes	No

Signature G/13/18

			_
		Care	Co
L-III	nanv	Lair	UU

Chinomso Ezenwata, APRN DEA #: _

□ Chinyare Okeke, MD DEA #:

3110 S Durango Dr #200 Las Vegas, NV 89032 Tel: 702-415-9760 • Fax: 702-478-6211

Address

DOB . Date 7. (. 101(18.

Label

times PRN

Dispense Only as Written

SCRIPT# 1137

Order # 2439194-1

MD, APRN

Label

Address.

times PRN

Dispense Only as Written L

inany Care Corp

□ Chinyare Okeke, MD DEA #:____

OChinomso Ezenwate, APRN 3110 S Durango Dr #200 Las Vegas, NV 89032 Tel: 702-415-9780 • Fax: 702-478-6211

PX_2_NA_V

FligRx.com 800-307-7717 RxPads.com

MD, APRN

FloRx.com 630-3072717 RxPads.com MD, APRN Chinyere Okeke, MD DEA #: 3110-S Durango Dr #200 Las Vegas, NV 89032 Tel; 702-415-9760 • Fax: 702-478-6211 Chinany Care Corp times PRN Dispense Only as Written Lychinomeo Ezerwela, APRN DEA #:

Consolo Certification for HALxM at Office Jireh Healthcare Services LLC

Patient Details for Header, Many Patient ID 470362, MRN M0456)

Date Of Birth		Status	Active
Screened Allergies	Erythromycin	Unscreened Allergies	opiate/narcotics
Start Of Care	06/13/2018	Gender	Female
Patient Address	Ave Las Vegas, I	NV 89121	

Hospice Certification of Terminal Illness

Benefit Period

Benefit Period Start Date

06/13/2018

Benefit Period End Date

09/10/2018

Certification Start Date

06/13/2018

Certification End Date

09/10/2018

Signed Date

06/13/2018

Medical Director or Certifying

Physician

Dubin, Stephen Paul

Medical Director or Certifying

Physician Address

5741 S. Fort Apache Rd. Suite 100 Las Vegas, NV

89148

Attending Physician

Dubin, Stephen Paul

Attending Physician Address

5741 S. Fort Apache Rd. Sulte 100 Las Vegas, NV

89148

Patient Hi Claim No.

571445912A

Disaster Acuity

Medical services required

within 72 hours

Provider Information

Provider Name

Jireh Healthcare Services

LLC

Provider Number

291533

Provider Address

2320 Paseo Del Prado Ste B101 Las Vegas, NV 89102-0048

Admission

New Admission 06/13/2018 (ROUTINE)

Verbal Certification

Verbal Certification received from Medical Director: Dubin, Stephen Paul by RN_Rulz, Janyvill (Jruiz) on 06/13/2018

I certify that Halstead, Mary is terminally III with a life expectancy of six months or less if the terminal illness runs its normal course.

Skilled Nurse RN_Ruiz, Janyvill (User) signed on 06/13/2018. Recorded by jruiz on 06/14/2018 10:04:57.

Verbal Certification received from Physician: Dubin, Stephen Paul by RN_Ruiz, Janyvill (jruiz) on 06/13/2018

, Magnis terminally ill with a life expectancy of six months or less if the terminal Illness runs its normal course.

Skilled Nurse RN_Ruiz, JanyvIII (User) signed on 06/13/2018. Recorded by jruiz on 06/14/2018 10:05:12.

Consolo Certification for HALxM at Office Jireh Healthcare Services LLC

Brief Narrative Statement and Attestation

Review the patient's clinical circumstances and synthesize the medical information to provide clinical justification for admission and continuation to hospice services. If in third (3rd) or more benefit periods, clinical findings from the face to face encounter have been used to determine continued eligibility for hospice care.

Dubin, Stephen Paul

Composing Physician Pt. is an 85 year old, Caucasian female, with disabling dyspnea, hypoxemic on room air, COPD, hypertension, age related physical debility, polyneuropathy, has fracture on unspecified part of beck of left femur, chronic pain, dementia, GERD, muscle weakness, dysphagia, with other symbolic dysfunction. Pt. is currently bed bound, and has very poor appetite. Needs maximum to complete assistance on all her ADL's

Recorded By

RN_Ruiz, Janyvill (jruiz)

Attestation: I confirm that I composed this narrative and that it is based on my review of the patient's medical record and/or examination of the patient.

Medical Director Dubin, Stephen Paul (Medical Director) signed on 06/13/2018. Recorded by jruiz on 06/14/2018 10:06:24.

Certification Statement

I certify that this patient is under my care, and to the best of my medical knowledge given the data available, has a life expectancy of six (6) months or less if the terminal illness runs its normal course.

Medical Director or Certifying Physician: Medical Director Dublin, Stephen Paul (Medical Director) signed on 06/13/2018. Recorded by jruiz on 06/14/2018 10:06:32.

I certify that this patient is under my care, and to the best of my medical knowledge given the data available, has a life expectancy of six (6) months or less if the terminal illness runs its normal course.

Attending Physician: Medical Director Dubin, Stephen Paul (Medical Director) signed on 06/13/2018. Recorded by jruiz on 06/14/2018 10:06:32.

Additional Signatures

Skilled Nurse RN_Rulz, JanyvIII (User) signed on 06/14/2018. Recorded by jruiz on 06/14/2018 10:06:40.

Unscreened Allergies oplate/narcotics

Screened Allergies Erythromycin

Hammel, Man (470362)

MRN M0456

Start Of Care 06/13/2018

Level Of Care ROUTINE More

Hospice Certification of Terminal Illness

Benefit Period

Certification End Date

Benefit Period Start Date 12/10/2018

Signed Date - 12/03/2018 Benefit Period End Date 02/07/2019

Medical Director or Certifying Physician ' Dubin, Stephen Paul Certification Start Date 12/10/2018

Medical Director or Certifying Physician Address 5741 S. Fort Apache Rd. Suite 100 Las Vegas, NV 89148

Patient Hi Claim No

<u>Disaster Acuity</u> Medical services required within 72 hours

Provider Information

Provider Name
Jireh Healthcare Services LLC

Provider Number 291533 Provider Address 2320 Paseo Del Prado Ste B101 Las Vegas, NV 89102-0048

Admission

New Admission 06/13/2018 (ROUTINE)

eptional Face to Face Circumstances

Exceptional Face to Face Circumstances

Verbal Certification

Verbal Certification received from Medical Director: Dubin, Stephen Paul by Dugay, Julia (jduga) on 12/03/2018

I certify that Harman, Mannis terminally ill with a life expectancy of six months or less if the terminal illness runs its normal course.

Skilled Nurse Dugay, Julia (User) signed on 12/03/2018. Recorded by Jduga on 12/18/2018 14:29:49.

Brief Narrative Statement and Attestation

Review the patient's clinical circumstances and synthesize the medical information to provide clinical justification for admission and continuation to hospice services. If in third (3rd) or more benefit periods, clinical findings from the face to face encounter have been used to determine continued eligibility for hospice care.

Composing Physician Dubin, Stephen Paul Ms Mary Halstead is an 85 year old, white female with PMH of COPD (hypoxemic on room air), hypertension, age related physical debility, polyneuropathy, chronic pain, dementia, GERD, and anxiety disorder. Pt. is chair bound and is mostly in the wheelchair. She has very poor appetite and has generalized muscle weakness. She needs maximum to complete assistance on all her ADL's.

Recorded By Dugay, Julia (jduga)

Attestation: I confirm that I composed this narrative and that it is based on my review of the patient's medical record and/or examination of the patient

Medical Director Dubin, Stephen Paul (Medical Director) signed on 12/03/2018. Recorded by Jduga on 12/18/2018 14:29:12

Certification Statement

recertify that this patient is under my care, and to the best of my medical knowledge given the data available, has a life expectancy of six (6) months less if the terminal illness runs its normal course.

Medical Director or Certifying Physician: Medical Director Dubin, Stephen Paul (Medical Director) signed on 12/03/2018. Recorded by jduga on 12/18/2018 14:28:30.

Consolo Physician for HALxM at Office Jireh Healthcare Services LLC

Patient Details for Hammel, Man (Patient ID 470362, MRN M0456)

Date Of Birth	Status Active
Screened Allergles Erythromycin	Unscreened Allergles opiate/narcotics

Physician's Order Details

<u>User</u>

Dugay, Julia (jduga)

<u>Patlent</u>

Order Date 12/03/2018

Time of Event 1447

Physician Dubin, Stephen Paul **Nurse**

Dugay, Julia (jduga)

Oversight

Orders

Admit to Jireh Healthcare Services for the 1st 60 day benefit period * Pharmacist may use generic drug equivalents or compound medications as appropriate. * Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to: hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction machine, etc.) * Assess and evaluate cardiac, respiratory, GI,GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs * Instruction on care of terminally ill, medications, safety, nutritional needs, pain and symptom control based on assessment * Assess and evaluate current medications * May give nutritional supplements 1 can a day as tolerated for supplement * May crush medications that are crush-able. * Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders * Hospice Medical Director may assist patient as needed to manage symptoms * Hospice nurse may pronounce death * Oxygen 2.5 lpm via NC continuously or as needed for SOB * Check oxygen saturation level prn every SN visit * Bed-ralls up 2x * Skilled Nurse visits 1x/week and 2x/week as needed for fall, agitation, SOB, leg edema & change in level of consciousness issues to promote comfort and symptom management within certification period. * CHHA visits 5x/week to assist with personal care, home making and to promote comfort per RN assignment within certification period. * Chaplain visits - evaluation, 1x/mo, 1x/mo as needed for spiritual issues. * Social worker visits - evaluation, 1x/mo as needed for psychosocial issues. * Volunteer Services continue to be offered every week per Skilled Nurse/Staff. * Admit patient to Jireh Healthcare Services with the following orders verified with M.D: Patient: DNR Diet: As Tolerated * Admit with the following medications: Please see related medications.

Read Back

Non Verbal Order

Yec

Medications

Medication	Instructions	Quantity	Refills
Furosemide 1 Tablet 1 times a day PO (20 MG Tablet)	1	*	- 3
LORazepam 1 Tablet 3 times a day PO As Needed Agitation (0.5 MG Tablet)			
PoTASsium ChLORide 1 Tablet 1 times a day PO (20 mEq Tablet)			75

Related Assignees

Social Services - MSW_Minnick, Joseph 1 PRN visit(s) every 30 day(s) effective 09/03/2018

Skilled Nurse - RN_Cacanindin, Speedy 1 visit(s) Weekly 2 PRN visit(s) every 7 day(s) effective 09/03/2018

Physician - Dubin, Stephen Paul 1 PRN visit(s) every 30 day(s) effective 09/03/2018

MD Director - Dubin, Stephen Paul 1 PRN visit(s) every 30 day(s) effective 09/03/2018

Hospice Alde - Young, Grace Marie 5 visit(s) Weekly effective 09/03/2018

Chaplain - CHAPLAIN_Murphy, Kathryn 1 visit(s) Monthly 3 PRN visit(s) every 30 day(s) effective 09/03/2018

Signatures

1. Medical Director Dubin, Stephen Paul (Medical Director) signed on 12/03/2018. Recorded by jduga on 12/18/2018 14:57:45.

Consolo Physician for HALxM at Office Jireh Healthcare Services LLC

Patient Details for Haintan, Mann (Patient ID 470362, MRN M0456)

Date Of Birth	Status	Active
Screened Allergles Erythromycin	Unscreened Allergies	opiate/narcotics

Physician's Order Details

User Fronda, MaryRose

(mary1)

Patient

Order Date 09/03/2018 Time of Event 1000

Physician Dubin, Stephen Paul <u>Nurse</u> Fronda, MaryRose (mary1)

Related Yes

Oversight Nο

Orders

Re-certify to Jirch Healthcare Services, LLC, for the 2nd 90 day benefit period

- Pharmacist may use generic drug equivalents or compound medications as appropriate.
- * Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to: hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction machine, etc.)
- * Assess and evaluate cardiac, respiratory, GI, GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs
- Instruction on care of terminally III, medications, safety, nutritional needs, pain and symptom control based on assessment
- * Assess and evaluate current medications
- Administer oxygen via nasal cannula at 2.5 lpm as needed for shortness of breath or until oxygen saturation is 95% and above
- * May give nutritional supplements 1 can a day as tolerated for supplement
- * May crush medications that are crush-able.
- * Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders
- Hospice Medical Director may assist patient as needed to manage symptoms
- Hospice nurse may pronounce death
- Check oxygen saturation level every SN visit
- * Bed-rails up 2x
- * Skilled Nurse visits (Eval, 1x/week and 2x/week as needed for respiratory issues within certification period) to promote comfort and symptom management.
- CHHA visits (5x/week within certification period) to assist with personal care, home making and to promote comfort per RN assignment.
- * MSW visits (1x/month as needed for psychosocial issues within certification period) to provide psychosocial support to patient and
- Chaplain visits (1k/month as needed for spiritual issues within certification period) to provide spiritual

Read Back

Non Verbal Order

Yes

No

Related Assignees

Hospice Aide - Young, Grace Marie 5 visit(s) Weekly effective 06/20/2018

MD Director - Dubin, Stephen Paul effective 06/20/2018

Physician - Dubin, Stephen Paul effective 06/20/2018

Skilled Nurse - RN_Ruiz, Janyvill effective 06/20/2018

Skilled Nurse - RN_Cacanindin, Speedy effective 06/20/2018

Social Services - MSW_Minnick, Joseph effective 06/20/2018

Chaplain - CHAPLAIN_Murphy, Kathryn effective 06/20/2018

Signatures

1. Medical Director Dubin, Stephen Paul (Medical Director) signed on 09/03/2018. Recorded by mary1 on 09/16/2018 22:58:18.

Consolo Physician for HALxM at Office Jireh Healthcare Services LLC

Signature:		
Date:	£	

Patient Details for Hammel, Many (Patient ID 470362, MRN)

Date Of Birth Status Active

Physician's Order Details

<u>User</u>

RN_Ruiz, Janyvill (jruiz)

<u>P</u>

Time of Event

1629

Physician

Dubin, Stephen Paul

Nurse

RN_Ruiz, Janyvill (jruiz)

Oversight

Order Date

06/12/2018

No

<u>Orders</u>

Read Back Yes Non Verbal Order

No

Medications

Medication	Instructions	Quantity	Refills
Spiriva HandiHaler 2 cap(s) 1 times a day IN (18 MCG Capsule)			
Folic Acid 1 tab 1 times a day PO (1 MG Tablet)			
Levothyroxine 1 tab 1 times a day PO (25 mcg Tablet)	to be taken every morning		
Vitamin D 1 tab 1 times a day PO (2000 UNIT Capsule)	25 u S		
Vitamin C 1 tab 1 times a day PO (500 MG Tablet)		.~.	
vitamin B12 1 tab 1 times a day PO (1000 mcg Tablet)	12.0		
Sertraline HCl 1 tab 1 times a day PO (50 MG Tablet)			
Ferrous Sulfate 1 tab 2 times a day PO (325 (65 Fe) MG Tablet)	<u> </u>		11
Carvedilol 1 tab 2 times a day PO (3.125 MG Tablet)			
RisperiDONE 1 tab 1 times a day PO (0.5 MG Tablet)	take one tab orally daily at bedtime		
Oxygen Concentrator 2.5 L/M of 1 each Device continuously XX	I G		

- 1. Medical Director Dubin, Stephen Paul (Medical Director) signed on 06/16/2018. Recorded by jruiz on 06/16/2018 16:30:11.
- 2. Skilled Nurse RN_Ruiz, JanyvIII (User) signed on 06/16/2018. Recorded by jruiz on 06/16/2018 16:30:05.

Patient Details for Hallacett, Many (Patient ID 470362, MRN)

Date Of Birth Status Active

Physician's Order Details

<u>User</u> RN_Ruiz, Janyvill (jruiz)

Dubin, Stephen Paul

Patlent

Order Date 06/12/2018 Time of Event 1629

Physician //

Nurse

06/12/201

<u>rse</u> RN_Ruiz, Janyvill (jruiz) Oversight No

<u>Orders</u>

Read Back Yes

Non Verbal Order

No

Medications

Medication	Instructions	Quantity	Refills
Spiriva HandiHaler 2 cap(s) 1 times a day IN (18 MCG Capsule)		1	
Folic Acid 1 tab 1 times a day PO (1 MG Tablet)			
Levothyroxine 1 tab 1 times a day PO (25 mcg Tablet)	to be taken every morning		
Vitamin D 1 tab 1 times a day PO (2000 UNIT Capsule)			
Vitamin C 1 tab 1 times a day PO (500 MG Tablet)			
vitamin B12 1 tab 1 times a day PO (1000 mcg Tablet)			
Sertraline HCl 1 tab 1 times a day PO (50 MG Tablet)			
Ferrous Sulfate 1 tab 2 times a day PO (325 (65 Fe) MG Tablet)			
Carvedilol 1 tab 2 times a day PO (3.125 MG Tablet)			
RisperiDONE 1 tab 1 times a day PO (0.5 MG Tablet)	take one tab orally daily at bedtime		
Oxygen Concentrator 2.5 L/M of 1 each Device continuously XX			

- 1. Medical Director Dubin, Stephen Paul (Medical Director) signed on 06/16/2018. Recorded by jruiz on 06/16/2018 16:30:11.
- 2. Skilled Nurse RN_Ruiz, Janyvill (User) signed on 06/16/2018. Recorded by jruiz on 06/16/2018 16:30:05.

Consolo Indicator Dated: 09/03/2018 for Hammer, Ma

Patient Details for Hamman (Patient ID 470362, MRN M0456)

Date Of Birth Status Pending - Discharged

Indicator Dated 09/03/2018 for M Hand Medical Record No M0456

Effective Date 09/03/2018

Diagnoses

ICD10s

Primary ICD10 - J44.9 Chronic obstructive pulmonary disease, unspecified Secondary ICD10 - R06.02 Shortness of breath
Tertiary ICD10 - R54 Age-related physical debility
4th ICD10 - R13.12 Dysphagia, oropharyngeal phase
5th ICD10 - M62.81 Muscle weakness (generalized)
6th ICD10 - Z74.01 Bed confinement status

Patient Details for Hamman, Mark (Patient ID 470362, MRN M0456)

Date Of Birth

Status Pending - Discharged

Physician's Order Details,

User

Dugay, Julia (jduga)

Order Date 12/03/2018 Time of Event 1447

Physician

Dubin, Stephen Paul

Nurse Dugay, Julia (jduga) Oversight

No

Orders

Admit to Jireh Healthcare Services for the 1st 60 day benefit period * Pharmacist may use generic drug equivalents or compound medications as appropriate. * Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to: hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction machine, etc.) * Assess and evaluate cardiac, respiratory, GI,GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs * Instruction on care of terminally ill, medications, safety, nutritional needs, pain and symptom control based on assessment * Assess and evaluate current medications * May give nutritional supplements 1 can a day as tolerated for supplement * May crush medications that are crush-able. * Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders.* Hospice Medical Director may assist patient as needed to manage symptoms * Hospice nurse may pronounce death * Oxygen 2.5 lpm via NC continuously or as needed for SOB * Check oxygen saturation level prin every SN visit * Bed-rails up 2x * Skilled Nurse visits 1x/week and 2x/week as needed for fall, agilation, SOB, leg edema & change in level of consciousness issues to promote comfort and symptom management within certification period. * CHHA visits 5x/week to assist with personal care, frome making and to promote comfort per RN assignment within certification period. * Chaplain visits - evaluation, 1x/mo, 1x/mo as needed for spiritual issues. * Social worker visits - evaluation, 1x/mo as needed for psychosocial issues. * Volunteer Services continue to be offered every week per Skilled Nurse/Staff, * Admit patient to Jirch Healthcare Services with the following orders verified with M.D. Patient: DNR Diet: As Tolerated * Admit with the following medications: Please see related medications.

Read Back Yes

Non Verbal Order

Medications

Medication	Instructions	Quantity	Refills
Furosemide 1 Tablet 1 times a day PO (20 MG Tablet)		-	
LORazepam 1 Tablet 3 times a day PO As Needed Agitation (0.5 MG Tablet)	(3)		
PoTASsium ChŁORide 1 Tablet 1 times a day PO (20 mEq Tablet)	16: E		-

Related Assignees

Social Services - MSW_Minnick, Joseph 1 PRN visit(s) every 30 day(s) effective 09/03/2018

Skilled Nurse - RN_Cacanindin, Speedy 1 visit(s) Weekly 2 PRN visit(s) every 7 day(s) effective 09/03/2018

Physician - Dubin, Stephen Paul 1 PRN visit(s) every 30 day(s) effective 09/03/2018

MD Director - Dubin, Stephen Paul 1 PRN visit(s) every 30 day(s) effective 09/03/2018

Hospice Aide - Young, Grace Marie 5 visit(s) Weekly effective 09/03/2018

Chaplain - CHAPLAIN_Murphy, Kathryn 1 visit(s) Monthly 3 PRN visit(s) every 30 day(s) effective 09/03/2018

Signatures

1. Medical Director Dubln, Stephen Paul (Medical Director) signed on 12/03/2018. Recorded by jduga on 12/18/2018 14:57:45,

Consolo Physician Order (2253456) Dated: 09/17/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Order (2253456) Dated: 09/17/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Order (2253456) Dated: 09/17/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Order (2253456) Dated: 09/17/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Order (2253456) Dated: 09/17/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Order (2253456) Dated: 09/17/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Order (2253456) Dated: 09/17/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Consolo Physicia

Patient Details for Hamman Mine (Patient ID 470	0362, MRN M0456)
Date Of Birth	Status Pending - Discharged

Physician's Order Details

User

RN_Ruiz, Janyvill ([ruiz)

Order Date 09/17/2018 Time of Event 1946

Physician

Dubin, Stephen Paul

<u>Nurse</u> RN_Ruiz, Janyvill (jruiz) Oversight No

Orders

Ensure 2 Bottle 1 times a day PO As Needed Supplement (Liquid)

Read Back Yes

Non Verbal Order

No

Medications

Medication	Instructions	Quantity	Refills
Ensure 2 Bottle 1 times a day PO As Needed Supplement (Liquid)	310 M		

- 1. Medical Director Dubln, Stephen Paul (Medical Director) signed on 09/17/2018. Recorded by jruiz on 09/18/2018 19:49 21.
- 2. Skilled Nurse RN_Ruiz, JanyvIII (User) signed on 09/18/2018. Recorded by jruiz on 09/18/2018 19:49:04.

Consolo Physician Order (2253447) Dated: 09/18/2018 for Okeke, Chinyere for Ha

Patient Details for H

(Patient ID 470362, MRN M0456)

Date Of Birth

Status Pending - Discharged

Physician's Order Details

User

RN_Rulz, Janyvill (jruiz)

Order Date 09/18/2018 Time of Event 1944

Physician

Okeke, Chinyere

Nurse

RN_Ruiz, Janyvill (jrulz)

Oversight Nο

Orders

Discontinue Medication Ensure 2 Bottle 1 times a day PO (Liquid) on 09/17/2018

Read Back

Non Verbal Order No

Yes

Consolo Physician Order (2253444) Dated: 09/17/2018 for Dubin, Stephen Paul for Halander, Me

Patient Details for Hames, Mar (Patient ID 470362, MRN M0456)

Date Of Birth Status Pending - Discharged

Physician's Order Details

User

RN_Rulz, Janyvill (jruiz)

Patient Head, Man Order Date 09/17/2018 Time of Event 1942

Physician

Dubin, Stephen Paul

Nurse

RN_Ruiz, Janyvill (jruiz)

Oversight No

Orders

Albuterol Sulfate 1 vial 2 times a day IN ((2.5 MG/3ML) 0.083% Nebu Soln

Read Back Yes Non Verbal Order No

Medications

Medication Instructions Quantity Refills
Albuterol Sulfate 1 vial 2 times a day IN ((2.5 MG/3ML) 0.083% Nebu Soln)

- 1. Medical Director Dublin, Stephen Paul (Medical Director) signed on 09/17/2018. Recorded by Jruiz on 09/18/2018 19:43:41.
- 2. Skilled Nurse RN_Ruiz, Janyvill (User) signed on 09/18/2018. Recorded by jruiz on 09/18/2018 19:43:32.

Consolo Physician Order (2253392) Dated: 09/17/2018 for Dubin, Stephen Paul for Hammer, Management of the Consolo Physician Order (2253392) Dated: 09/17/2018 for Dubin, Stephen Paul for Hammer, Management of the Consolo Physician Order (2253392) Dated: 09/17/2018 for Dubin, Stephen Paul for Hammer, Management of the Consolo Physician Order (2253392) Dated: 09/17/2018 for Dubin, Stephen Paul for Hammer, Management of the Consolo Physician Order (2253392) Dated: 09/17/2018 for Dubin, Stephen Paul for Hammer, Management of the Consolo Physician Order (2253392) Dated: 09/17/2018 for Dubin, Stephen Paul for Hammer (2253392) Dated: 09/17/2018 for Dubin, Stephen Paul for Hammer (2253392) Dated: 09/17/2018 for Dubin, Stephen Paul for Hammer (2253392) Dated: 09/17/2018 for Dubin, Stephen Paul for Hammer (2253392) Dated: 09/17/2018 for Dubin, Stephen Paul for Hammer (2253392) Dated: 09/17/2018 for Dubin, Stephen Paul for Hammer (2253392) Dated: 09/17/2018 for Dubin, Stephen Paul for Hammer (2253392) Dated: 09/17/2018 for Dubin (225392) Dated: 09/17/2018 for

Patient Details for Hamilton Many (Patient ID 470362, MRN M0456)

Date Of Birth Status Pending - Discharged

Physician's Order Details

User

RN_Ruiz, Janyvill (jruiz)

Hammer, Market

Order Date 09/17/2018 Time of Event

Physician

Dubin, Stephen Paul

Nurse

RN_Ruiz, Janyvill (jruiz)

Oversight No

Orders

Sertraline HCl 2 tab 1 times a day PO (50 MG Tablet

Read Back Yes

Non Verbal Order No

- 1. Medical Director Dubin, Stephen Paul (Medical Director) signed on 09/17/2018. Recorded by jruiz on 09/18/2018 19:15:54.
- 2. Skilled Nurse RN_Rulz, Janyvill (User) signed on 09/18/2018. Recorded by jruiz on 09/18/2018 19:15:47.

Consolo Physician Order (2253401) Dated: 08/21/2018 for Dubin, Stephen Paul for Ham

Patient Details for H Patient ID 470362, MRN M0456) Date Of Birth Status Pending - Discharged

Physician's Order Details

RN_Rulz, Janyvill (jrulz)

Order Date 08/21/2018 Time of Event 1924

Physician Dubin, Stephen Paul Nurse AN_Ruiz, Janyvill (jruiz) Oversight No

Orders

Ventolin HFA 2 puff(s) every 6 hours IN As Needed SOB (108 (90 Base) MCG/ACT Aerosol Soln

Read Back Yes

Non Verbal Order

Medications

1	Medication	Instructions	Quantity	Refills
	Ventolin HFA 2 puff(s) every 6 hours IN As Needed SOB (108 (90 Base) MCG/ACT Aerosol Soln)	3		

- 1. Medical Director Dubln, Stephen Paul (Medical Director) signed on 08/21/2018. Recorded by jruiz on 09/18/2018 19:25:23.
- 2. Skilled Nurse RN_Rulz, JanyvIII (User) signed on 09/18/2018. Recorded by Jruiz on 09/18/2018 19:25:11.

Consolo Physician Order (2253414) Dated: 09/17/2018 for Dubin, Stephen Paul for He

Patient Details for Hamilton, Management ID 470362, MRN M0456)

Date Of Birth Status Pending - Discharged

Physician's Order Details

User

RN_Ruiz, Janyvill (jruiz)

H H

Order Date 09/17/2018 Time of Event 1930

Physician

Dubin, Stephen Paul

Nursa

RN_Ruiz, Janyvill (jruiz)

Oversight No

(Jruiz)

Orders

Sertraline HCl 2 (ab 1 times a day PO (50 MG Tablet)

Read Back

Non Verbal Order

Medications

1	Medication	Instructions **	Quantity	Refills
I	Sertraline HCl 2 tab 1 times a day PO (50 MG Tablet)			

- 1. Medical Director Dublin, Stephen Paul (Medical Director) signed on 09/17/2018. Recorded by jruiz on 09/18/2018 19:31:58.
- 2. Skilled Nurse RN_Rulz, JanyvIII (User) signed on 09/18/2018. Recorded by Iruiz on 09/18/2018 19:31:45.

Consolo Physician Order (2253386) Dated: 09/18/2018 for Dubin, Stephen Paul for H

Patient Details for Hamman (Patient ID 470362, MRN M0456)

Date Of Birth

Status Pending - Discharged

Physician's Order Details

User RN_Ruiz, Janyvill (jruiz)

Order Date 09/18/2018

Time of Event 1910

Physician

Dubin, Stephen Paul

Nurse

RN_Ruiz, Janyvill (Iruiz)

Oversight No

Orders

Discontinue Medication Sertraline HCl 1 tab 1 times a day PO (50 MG Tablet) on 09/10/2018

Read Back

Yes

Non Verbal Order

No

Patient Details for House Manua (Patient ID 470362, MRN M0456)

Status Pending - Discharged Date Of Birth

Physician's Order Details

User RN_Ruiz, Janyvill (jruiz)

Order Date 09/18/2018 Time of Event 1845

Physician Dubin, Stephen Paul Nurse

Oversight

RN_Ruiz, Janyvill ([ruiz)

No

Orders

Discontinue Medication Vitamin D 1 lab 1 times a day PO (2000 UNIT Capsule) on 09/15/2018

Read Back

Non Verbal Order No

Yes

Medications

Medication	Instructions	Quantity	Refills
Vitamin D 2 tab 1 times a day PO (2000 UNIT Tablet)	Take one tablet by mouth once dally.		

- 1. Medical Director Dubln, Stephen Paul (Medical Director) signed on 09/19/2018. Recorded by Juiz on 09/18/2018 18:49:29.
- 2. Skilled Nurse RN_Ruiz, JanyvIII (User) signed on 09/18/2018. Recorded by ruiz on 09/18/2018 18:49:21.

Consolo Physician Order (2253357) Dated: 09/18/2018 for Dubin, Stephen Paul for Hamman

Patient Details for Hamm , May (Patient ID 470362, MRN M0456) Date Of Birth Status Pending - Discharged

Physician's Order Details

User

RN_Ruiz, Janyvill (ruiz)

Order Date 09/18/2018

Time of Event 1845

Physician Dubin, Stephen Paul

<u>Nurse</u> RN_Ruiz, Janyvill (jruiz) Oversight No

Orders

Discontinue Medication Vitamin D 1 tab 1 times a day PO (2000 UNIT Capsule) on 09 16/2018

Read Back Yes

Non Verbal Order

Medications

Medication		Instructions	Quantity	Refills
Vitamin D t tab 1 times a day PO (20)	0 UNIT Capsule)	Take one tablet by mouth once dally.		

Consolo Physician Order (2249033) Dated: 09/03/2018 for Dubin, Stephen Paul for H

Patient Details for Hamman Manus Patient ID 470362, MRN M0456)

Date Of Birth

Status Pending - Discharged

Physician's Order Details

User

Fronda, MaryRose (mary1)

Order Date 09/03/2018 Time of Event 1000

Dubin, Stephen Paul

Nurse Fronda, MaryRose (mary1)

Related Yes

Oversight No

Orders

Physician

Re-certify to Jirch Healthcare Services, LLC, for the 2nd 90 day benefit period

- * Pharmacist may use generic drug equivalents or compound medications as appropriate.
- * Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction
- * Assess and evaluate cardiac, respiratory, Gl. GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs
- * Instruction on care of terminally III, medications, safety, nutritional needs, pain and symptom control based on assessment
- Assess and evaluate current medications
- Administer oxygen via nasal cannula at 2.5 lpm as needed for shortness of breath or until oxygen saturation is 95% and above
- * May give nutritional supplements 1 can a day as tolerated for supplement
- * May crush medications that are crush-able.
- * Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders
- * Hospice Medical Director may assist patient as needed to manage symptoms
- * Hospice nurse may pronounce death
- Check oxygen saturation level every SN visit
- * Bed-rails up 2x
- * Skilled Nurse visits (Eval, 1x/week and 2x/week as needed for respiratory issues within certification period) to promote comfort and symptom management
- * CHHA visits (5x/week within certification period) to assist with personal care, home making and to promote comfort per RN assignment.
- * MSW visits (1x/month as needed for psychosocial issues within certification period) to provide psychosocial support to patient and
- * Chaplain visits (1x/month as needed for spiritual issues within certification period) to provide spiritual

Read Back

Non Verbal Order

Related Assignees

Hospice Aide - Young, Grace Marie 5 visit(s) Weekly effective 06/20/2018

No

MD Director - Dubin, Stephen Paul 1 PRN visit(s) every 30 day(s) effective 06/20/2018

Physician - Dubin, Stephen Paul 1 PRN visit(s) every 30 day(s) effective 06/20/2018

Skilled Nurse - RN_Ruiz, Janyvill 2 PRN visit(s) every 7 day(s) effective 06/20/2018

Skilled Nurse - RN_Cacanindin, Speedy 1 visit(s) Weekly 2 PRN visit(s) every 7 day(s) effective 06/20/2018

Social Services - MSW_Minnick, Joseph 1 PRN visit(s) every 30 day(s) effective 06/20/2018

Chaplain - CHAPLAIN_Murphy, Kathryn 1 visit(s) Monthly 3 PRN visit(s) every 30 day(s) effective 06/20/2018

Signatures

1. Medical Director Dubin, Stephen Paul (Medical Director) signed on 09/03/2018. Recorded by mary1 on 09/16/2018 22:58:18.

Consolo Physician Order (2247289) Dated: 06/18/2018 for Dubin, Stephen Paul for Hamman

Patient Details for H Mem (Patient ID 470362, MRN M0456) Date Of Birth Status Pending - Discharged

Physician's Order Details

User

RN_Esteban, Randy (Reste)

Patient

Order Date 06/18/2018 Time of Event

1639

Physician

Dubin, Stephen Paul

Nurse RN_Esteban, Randy (Reste)

Oversight

No

Orders

Discontinue previous frequency order for chaplain. Start new chaplain frequency: 1/x mo and 3x/mo as needed for spiritual issues within certification period.

Read Back Yes

Non Verbal Order No

Consolo Physician Order (2211649) Dated: 08/24/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Order (2211649) Dated: 08/24/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Order (2211649) Dated: 08/24/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Order (2211649) Dated: 08/24/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Order (2211649) Dated: 08/24/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Order (2211649) Dated: 08/24/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Order (2211649) Dated: 08/24/2018 for Dubin, Stephen Paul for Hamiltonian Consolo Physician Consolo Physicia

Patient Details for Health Man, (Patient ID 470362, MRN M0456)

Date Of Birth 4 Status Pending - Discharged

Physician's Order Details

User RN_Ruiz, Janyvill (jruiz)

Patient Hand, M

Order Date 08/24/2018 Time of Event 1109

Physician Dubin, Stephen Paul Nurse RN_Ruiz, Janyvill (jruiz) Oversight No

Orders

Discontinue Medication Oxygen Concentrator 2.5 L/M of 1 each Device continuously XX on 07/19/2018

Read Back Yes Non Verbal Order

No

Medications

Medication	Instructions	Quantity	Refills
Oxygen Concentrator 2.5 L/M of 1 each Device continuously XX			

Consolo Physician Order (2204657) Dated: 07/19/2018 for Okeke, Chinyere for Hamman

Mem

Patient Details for House, Man (Patient ID 470362, MRN M0456)

Date Of Birth

Status Pending - Discharged

Physician's Order Details

User

RN_Esteban, Randy (Reste) Patient M

Order Date 07/19/2018 Time of Event

Physician Okeke Chinyere Nurse RN_Esteban, Randy (Reste) Oversight No

Orders

- 1. Discontinue Carvedilol 3.125 MG Tablet, 1 tablet by mouth twice daily
- 2. Discontinue RisperIDONE 0.5 MG Tablet, 1 tablet by mouth once daily
- 3. Start Spiriva 18 mcg cp handihaler, inhale once daily
- 4. Start Ensure, 2 can by mouth daily
- 5. Start Albuterol 2.5 mg/3 ml via nebulizer, inhale three times daily as needed for shortness of breath, weezing

Read Back No Non Verbal Order

Yes

Medications

Medication	Instructions	Quantity	Refilis
Spiriva HandiHaler 1 cap(s) 1 times a day IN (18 MCG Capsule)	Take one capsule once dally.		
Ensure 2 Bottle 1 times a day PO (Liquid)	Administer 2 bottles by mouth once daily.		
Albuterol Sulfate 1 vial 3 times a day IN As Needed SOB and Wheezing ((2.5 MG/3ML) 0.083% Nebu Soin)	Administer three times a day intranasally via nebulizer as needed for shortness of breath and wheezing.		

- 1. Physician Okeke, Chinyere (Physician) signed on 07/19/2018. Recorded by Reste on 08/21/2018 04:04:36.
- 2. Skilled Nurse RN_Esteban, Randy (User) signed on 08/21/2018. Recorded by Reste on 08/21/2018 04:00:50.

Consolo Physician Order (2204655) Dated: 07/19/2018 for Okeke, Chinyere for Ha

Patient Details for Halstead, Mary (Patient ID 470362, MRN M0456)

Date Of Birth

Status Pending - Discharged

Physician's Order Details

User

RN_Esteban, Randy (Reste)

Order Date 07/19/2018

Time of Event

Physician Okeke, Chinyere

<u>Oversight</u>

RN_Esteban, Randy (Reste)

Νo

Orders

- 1. Discontinue Carvedilol 3.125 MG Tablet, 1 tablet by mouth twice daily
- 2. Discontinue RisperiDONE 0.5 MG Tablet, 1 tablet by mouth once daily
- 3. Start Spiriva 18 mcg cp handihaler, inhale once daily
- 4. Start Ensure, 2 can by mouth daily
- 5. Start Albuterol 2.5 mg/3 ml via nebulizer, inhale three times daily as needed for shortness of breath, weezing

Read Back

Non Verbal Order

- 1. Skilled Nurse RN_Esteban, Randy (User) signed on 08/21/2018. Recorded by Reste on 08/21/2018 04:01:27.
- 2. Physician Okeke, Chinyere (Physician) signed on 07/19/2018. Recorded by Reste on 08/21/2018 04:00:39.

Physician's Order Details

User

RN_Esteban, Randy (Reste)

Order Date 07/19/2018

Time of Event

Physician Okeke, Chinyere

Nurse RN_Esteban, Randy (Reste) Oversight No

Orders

Calmoseptine Cream topical

Apply to affected area each diaper change

Read Back No Non Verbal Order

Yes

Medications

	Medication	Instructions	Quantity	Refills
Ca	almoseptine 1 application TP (0.44;20.625 %;% Lotion)	Apply to affected area every diaper change		

- 1. Physician Okeke, Chinyere (Physician) signed on 07/19/2018. Recorded by Reste on 08/21/2018 03 36 38.
- 2. Skilled Nurse RN_Esteban, Randy (User) signed on 08/21/2018. Recorded by Reste on 08/21/2018 03:36;32.

Patient Details for H Mana Patient ID 470362, MRN M0456) Date Of Birth

Status Pending - Discharged

Physician's Order Details

User RN_Esteban, Randy

Order Date 07/19/2018 Time of Event

Physician Okeke, Chinyere

(Reste)

Nurse RN_Esteban, Randy (Reste)

Oversight No

Oxygen 2 lpm via nasal cannula as needed for shortness of breath

Read Back

Non Verbal Order

No

Yes

Medications

Medication	Instructions	Quantity	Reflits
Oxygen Concentrator 2 LiterPerMin INS As	Administer oxygen via nasal canula intranasally at two		
Needed SOB (1 each Device)	liters/minute as needed for shortness of breath.		

- 1. Skilled Nurse RN_Esteban, Randy (User) signed on 08/21/2018. Recorded by Reste on 08/21/2018 03:42:09.
- 2. Physician Okeke, Chinyere (Physician) signed on 07/19/2018. Recorded by Reste on 08/21/2018 03:41:47.

Consolo Physician Order (2182304) Dated: 06/13/2018 for Dubin, Stephen Paul for H



Patient Details for Hamilton (Patient ID 470362, MRN M0456)			
Date Of Birth	Status Pending - Discharged		

Physician's Order Details

User RN_Esteban, Randy Patient M

Order Date 06/13/2018 Time of Event

(Resta)
Physician

Oubin, Stephen Paul

Nurse RN_Esteban, Randy (Reste) Oversight No

Orders

Admit to Jireh Healthcare Services, LLC, for the 1st 90 day benefit period

- * Pharmacist may use generic drug equivalents or compound medications as appropriate.
- * Interdisciplinary team to determine equipment and supply needs based on assessment for comfort and symptom control (may include but not limited to: hospital bed, special mattress, bedside commode, over-bed-table, shower chair, wheelchair, walker, suction machine, etc.)
- * Assess and evaluate cardiac, respiratory, GI, GU, pain, mobility, skin, nutrition/dehydration, comfort level and symptom management, disease progression, safety and family/caregiver needs
- * Instruction on care of terminally ill, medications, safety, nutritional needs, pain and symptom control based on assessment
- * Assess and evaluate current medications
- Administer oxygen via nasal cannula at 2.5 lpm as needed for shortness of breath or until oxygen saturation is 95% and above
- * May give nutritional supplements 1 can a day as tolerated for supplement
- * May crush medications that are crush-able.
- * Medications to be administered by licensed staff, patient and/or family member/caregiver per medication profile and current orders
- * Hospice Medical Director may assist patient as needed to manage symptoms
- * Hospice nurse may pronounce death
- * Check oxygen saturation level every SN visit
- * Bed-rails up 2x
- * Skilled Nurse visits (Eval, 1x/week and 2x/week as needed for respiratory issues within certification period) to promote comfort and symptom management.
- * CHHA visits (5x/week within certification period) to assist with personal care, home making and to promote comfort per RN assignment.
- * MSW visits (1x/month as needed for psychosocial issues within certification period) to provide psychosocial support to patient and family.
- * Chaplain visits (1x/month as needed for spiritual issues within certification period) to provide spiritual

Read Back

Non Verbal Order

Yes

No

- 1. Medical Director Dubin, Stephen Paul (Medical Director) signed on 06/13/2018. Recorded by mary1 on 09/16/2018 22:57:00.
- 2. Skilled Nurse RN_Esteban, Randy (User) signed on 08/07/2018. Recorded by Reste on 08/07/2018 13:08:37.

Physician's Order Details

User RN_Ruiz, Janyvill (jrulz)

Patient Head Man Order Date 06/26/2018 Time of Event 1703

Physician Dubin, Stephen Paul Nurse RN_Ruiz, Janyvill ([ruiz) Oversight No

Orders

Read Back Yes Non Verbal Order

Medications

Medication	Instructions	Quantity	Refills
Miralax 17 gm 1 times a day PO As	Mix 17 gm to 8oz glass of water. Take once daily by mouth as needed for		
Needed Constipation (17 gm Powder)	constipation. To administer if no bowel movement for three days.		

- 1. Medical Director Dubln, Stephen Paul (Medical Director) signed on 07/04/2018. Recorded by jruiz on 07/03/2018 17:03:30.
- 2. Skilled Nurse RN_Rulz, JanyvIII (User) signed on 07/03/2018. Recorded by jruiz on 07/03/2018 17:03:24.

Patient Details for H (Patient ID 470362, MRN)

Date Of Birth

Status Pending - Discharged

Physician's Order Details

User

RN_Ruiz, Janyvill (jruiz)

<u>Patient</u>

06/12/2018

Time of Event

Physician

Dubin, Stephen Paul

Nurse RN_Ruiz, Janyvill (jruiz) Oversight No

1629

Orders

Read Back Yes

Non Verbal Order No

Medications

Medication	Instructions	Quantity Refil:
Vitamin C 1 tab 1 times a day PO (500 MG Tablet)	Take one tablet by mouth once daily.	
Levothyroxine 1 tab 1 times a day PO (25 mcg Tablet)	Take one tablet by mouth once daily in the morning.	
Vitamin D 1 tab 1 times a day PO (2000 UNIT Capsule)	Take one tablet by mouth once daily.	
RisperiDONE 1 tab 1 times a day PO (0.5 MG Tablet)	take one tab orally daily at bedtlime	
Carvedilol 1 tab 2 times a day PO (3.125 MG Tablet)		
Spiriva HandiHaler 2 cap(s) 1 times a day IN (18 MCG Capsule)	Take two capsules once daily.	
Folic Acid 1 tab 1 times a day PO (1 MG Tablet)	Take one tablet by mouth once dally.	
vitamin B12 1 tab 1 times a day PO (1000 mcg Tablet)	Take one tablet by mouth once daily.	
Sertraline HCl 1 tab 1 times a day PO (50 MG Tablet)	Take one tablet by mouth once daily.	
Ferrous Sulfate 1 tab 2-times a day PO (325 (65 Fe) MG Tablet)	Take one tablet by mouth twice daily.	
Oxygen Concentrator 2.5 L/M of 1 each Device continuously XX		

- 1. Medical Director Dublin, Stephen Paul (Medical Director) signed on 06/16/2018. Recorded by jruiz on 06/16/2018 16:30:11.
- 2. Skilled Nurse RN_Ruiz, Janyvill (User) signed on 06/16/2018. Recorded by jruiz on 06/16/2018 16:30:05.

Consolo Indicator Dated: 06/13/2018 for Hamman

Patient Details for Hammer, March Patient ID 470362, MRN M0456)

Date Of Birth

Status Pending - Discharged

Medical

Indicator Dated 06/13/2018 for Men H

Record No M0456

Effective Date 06/13/2018

Diagnoses

ICD10s

Primary ICD10 - J44.9 Chronic obstructive pulmonary disease, unspecified Secondary ICD10 - R06.02 Shortness of breath
Tertiary ICD10 - R54 Age-related physical debility
4th ICD10 - R13.12 Dysphagla, oropharyngeal phase
5th ICD10 - M62.81 Muscle weakness (generalized)
6th ICD10 - Z74.01 Bed confinement status

Consolo Payer Group Details [Effective 06/13/2018 (506493)] for Hammer, Me

Patient Details for H

Maga (Patient ID 470362, MRN M0456) Date Of Birth

Status Pending - Discharged

Payer Group Details

Effective Date 06/13/2018

Hospica Item Sets

Medicare (traditional fee-for-service)

Payer 1: Medicare Part A

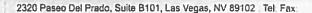
Payer Type Medicare

Payer Number 571445912A

Will Pay
Primary Hospice, Notice of Election, Pre Election

Suspend Claims?

Jireh Healthcare Services LLC





PHYSICIAN'S CERTIFICATION FOR HOSPICE BENEFIT

In order to determine if a patient is eligible for Hospice benefit we are required to have physician's authorization with a brief narrative supporting terminal illness.

Patient Name:

(First)

(MI)

M0456

MR#

Certification #3: Re-Certification Statement for Next 60-day period

• I certify that Hamilton, Many has a life expectancy of six months or less, if the terminal illness runs its normal course.

✓ Verbal Authorization (if needed) Date: 12/3/2018

Obtained by: Julia Dugay/ RN

Name / Title

Effective Date of Certification: 12/10/2018 To 2/7/2019

Primary DX:

Chronic obstructive pulmonary disease, unspecified (J44.9)

Secondary DX:

Shortness of breath (R06.02)

Comorbidities:

Certification is based on: Medical history, record and patient status

☑ Team Assessment

Face

0

to Face Encounter

Visit Date: 12/3/2018

(see MD F2F visit note)

RN Assessment

Narrative:

Patient is alert and oriented to person, place, and time. With +3 bilateral lower extremity edema, pedal pulses faint. Patient exhibits fatigue on exertion, fair coordination, and poor balance. Diet is as tolerated with significant worsening of appetite and nutritional intake. Patient with oxygen and breathing treatment at night for comfort. Patient requires complete assistance with activities of daily living (ADL's).

Physician Narrative: Ms. Halstead is an 85-year-old Caucasian female with diagnosis of COPD, hypoxemic on room air, hypertension, agerelated physical debility, polyneuropathy, chronic pain, dementia, GERD, and anxiety disorder. Patient is chair-bound and mostly on wheelchair. Patient with very poor appetite and has generalized muscle weakness. Patient requires maximum to complete assistance with ADL's. I recertify that this patient is under my care, and to the best of my medical knowledge given the data available, has a life expectancy of six months or less if the terminal illness runs its normal course.

Medical Director Attestation/Certification

· I attest/confirm that I composed this narrative based on my review of patient's medical records, team assessment and/or examination of the patient.

STEPHEN DUBIN (MD) / 1023311040

Signature on File/Verified by MARY ROSE FRONDA.

Date: 12/3/2018

Medical Director / Physician Designee / Referring MD (if attending)

Hospice Medical Director or Designee Name (Printed):

PHYSICIAN FACE-TO-FACE ENCOUNTER/ RECERTIFICATION OF TERMINAL ILLNESS

åbent Name:			ID# 10 10 10 10 10 10 10 10 10 10 10 10 10	Admission Date
M	n		M 0 456	6-13-18
BENEFIT PERIOR	D 60-Day Period from	12-10-18 to 00	2 - 07 - 2019	
ace to Face Encounter	(Note: Must be compl	leted no more than 30 days prior	to this benefit period)	
lospice Nurse Practitioner	r Attestation: I confirm that i	had a face-to-face encounter wit	th	(Patient's Na
n (Date)	and that the clinical findings	of that encounter have been pro-	vided to the certifying phys	sician for use in determining
ontinued eligibility for hospic	ce care.			
IP Name (Printed):		NP Signature:	•	Date
hysician Attestation: I cor	nfirm that I had a face-to-face	e encounter with	H	(Patient's Name)
		indings of that encounter in deter		for hospice care.
lospice Medical Director or		Hospice Medical Directo	or or Designee Signature:	
	MIN , MD			
still considered to be termi	lnally ill and has a life expect	reviewed the clinical record prior tancy of six (6) months or less, if	the terminal illness runs it	's normal course
leview the patient's clinical of		ist be completed no more than 15 ze the medical information to prov 86 ub white	ide clinical justification for	continued Hospice services
Chypoxedic o	in dement	HTN, Jace related	disorder of	t is chair bound
+ is mostly	in ser wheeld		nery poor a	pretite & has
generalized	mucle meats	ress; needs may	worth approx	House wifer her
ONDL'SO I	her condition	a continued to	rum its no	mad course,
all hors 6	no on less	to line		
Attestation: I confirm that I	composed this narrative bas	ed on my review of the patient's a	medical record and/or exa	mination of the patient.
lospice Medical Director or		Hospice Medical Directo	or or Designee Signature	Datou Britania
C TOULDAY 1	DUBIN, MD			
OVELLON !	Olive in the			The second secon
BENEFIT PERIO	pade Mandalysian to a	to .		
BENEFIT PERIO	D 60-Day Period from		to this benefit period)	
BENEFIT PERIO	D 60-Day Period from (Note: Must be comple	leted no more than 30 days prior		(Patents N
BENEFIT PERIOR ace to Face Encounter lospice Nurse Practitioner	60-Day Period from (Note: Must be compler Attestation: I confirm that	leted no more than 30 days prior I had a face-to-face encounter wi	th	
BENEFIT PERIOR ace to Face Encounter lospice Nurse Practitioner n(Date)	D 60-Day Period from (Note: Must be compler Attestation: I confirm that and that the clinical findings	leted no more than 30 days prior	th	
BENEFIT PERIOR ace to Face Encounter ospice Nurse Practitioner n/ (Date) ontinued eligibility for hospic	D 60-Day Period from (Note: Must be compler Attestation: I confirm that and that the clinical findings	leted no more than 30 days prior I had a face-to-face encounter wi s of that encounter have been pro	th	
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Hospice Medical Director or Designee Signature:

Date

DUBIN0165

THE HEALTHCARE SERVICES

PHONE: 702.359.1388 FAX: 702.359.2388
EMAIL ADDRESS: Jirehhealthcare @gmail.com

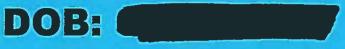
PHYSICIAN'S CERTIFICATION FOR MEDICARE/MEDICAID HOSPICE BENEFIT

Physician's Certification of Terminal Illness for Hospice Benefit
(Part - 1)

		MR# 10456
A.	CERTIFICATION STATEMENT First 90-Day Period from 6 13 201	8 to 9/10/2019
	Has a life expectancy of six (6) months or less if the terminal illness runs its not Reviewed the patient's clinical information and considered the primary terminal diagnoses, current subjective and objective medical findings, current medication orders and information about management of unrelated conditions in making the	l condition, related
	RN Signature	6/3/2017 Verbal Order Date
	Hospice Medical Director/Associate Hospice Doctor	6 13 2018 Date 6 13 2018
	Attending Physician Signature	6 13 14018 Date
В.	RECERTIFICATION STATEMENT Second 90-Day Period from Old I certify that I have reviewed the clinical record prior to recertification for the all That patient is still considered to be terminally ill and has life expectancy of six The terminal illness runs its normal course.	
		9-11-18 Date
	Hospice Medical Director/Associate Hospice Doctor	Date
c.	RECERTIFICATION STATEMENT 60-Day Period from 12 - 10 - 1 I certify that I have reviewed the clinical record prior to recertification for the at That patient is still considered to be terminally ill and has life expectancy of six The terminal illness runs its normal course.	
	Hospics Medical Division (A	
	Hospice Medical Director/Associate Hospice Doctor	Date

JIREH HEALTHCARE SERVICES LLC (HOSPICE AGENCY)

PATIENT NAME: DELEGE BELLE



START OF CARE: 12/20/2018 TO 12/22/2018

Claim Number	From Service Date, To S.		ervice Date Beneficiary Nam Glaim Status	Total Charge	e Type OffBill Admit Date
21 900700422207NVR	12/1/2018	12/31/2018	p - Paid	26'062'28	813 - Hospice (,, 9/7/2018
2183390053BB07NVR	11/1/2018	11/30/2018	P-Paid	\$8,724.43	813 - Hospice (9/7/2018
21830901126607NVR	10/1/2018	10/31/2018	P-Paid	\$9,083.84	813 - Hospice (9/7/2018
21827601222707NVR	9/7/2018	9/30/2018	P - Paid	\$6,942.12	812 - Hospice (9/7/2018
Claim Number	From Setvice Date To Service Date	To Service Date	Slaim Searus	Total Glarye	Type Off Bill Admit Bate
21900700421907NVR	12/1/2018 —	12/31/2018	p - Paid	\$6,986.61	813 - Hospice (10/10/2018
21833900542507NVR	11/1/2018	11/30/2018	b.Paid	\$8,197.48	813 - Hospice (10/10/2018
21833900230804NVR	10/10/2018	10/31/2018	P-Paid	\$6,230.74	817 - Hospice (10/10/2018
21830901121107NVR	10/10/2018	10/31/2018	P. Paid	\$5,630.74	812 - Hospice (10/10/2018
Claim Number	From Service Bate	To Service Date	Gaim Status	Total Grange	Typerof Bill Admyr Bate
21900901108507NVR	12/1/2018	12/26/2018	Paid	\$6,626.68	814-Hospice (6/13/2018
21833900542007NVR	11/1/2018	11/30/2018	- Paid	\$7,635.75	813 - Hospice (6/13/2018
21830901128607NVR	10/1/2018	10/31/2018	P - Paid	5 98,065.97	813 - Hospice (6/13/2018
21827601223807NVR	9/1/2018	9/30/2018	- Paid	** \$7,382.91	813 - Hospice (6/13/2018
21824701185107NVR	8/1/2018	8/31/2018	pine Paid	* \$8,276.79	813 - Hospice (6/13/2018
21821800581907NVR	7/1/2018	7/31/2018	P-Paid	\$9,216.50	813 - Hospice (_6/13/2018
21818600817807NVR	6/13/2018	6/30/2018	.p - Paid	\$5,126.72	812-Hospice (6/13/2018
Claim Number	From Service Date	From Service Date To Service Date	Slaim Status	Total Grange	Type Of Bill Admit.Date
91900700070001C	12/20/2018	12/22/2018	P - Paid	\$680.54	811 - Hospice (12/20/2018



PLEASF PRINT

Clark County Coroner/Medical Examiner Office

1704 Pinto Lane, Las Vegas, Nevada 89106 Phone number (702) 455-3210 Fax # (702) 455-3101 h

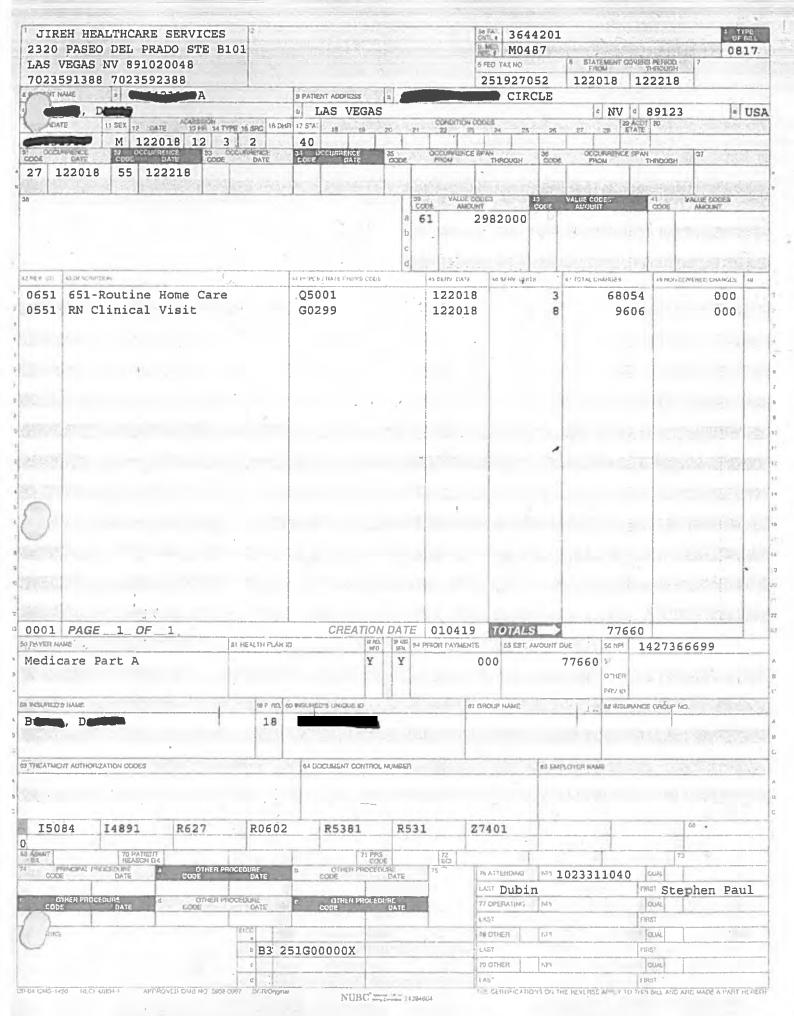
HOSPICE REPORT

All lines must be completed

1) Date: 12/22/2018 Time: 1430 Reported By: Roberto Alfajora PM
2) Agency: Jirch Healthcare Servius Phone # you are at Medical Record #: MOYY 6
3) Last Name: B First Name: O
4) Time pronounced: 1406 Date of Birth: Social Security #:
5) Race: Asian/Black/Caucasian/Hispanic/Indian/Multi-Cultured/Other (Circle One) Sex/Male// Female (Circle One)
6) Home address: (Arc)
City: Las Wigas Zip: 89123 Phone: 702
7) Death occurred at home(group home/)facility (Circle one)
8) Name of Group home or facility: Oliva Grouphord
9) Location of Death address: Circle City: Las Ugas Zip: 89123
10) Local Mortuary Used: Simple Genation Martial status: Married Single Widowed / Divorced (circle one)
11) Name of Next of Kin Notified: State United Relation: POA
12) NOK address:
City: Las Ugas Zip: 89184 Phone: 702
13) Who notified next of kin: Roberto Alfajora, Rr Method (Telephone) At Bedside (Circle One)
14) Death Certificate will be Signed By: Dr. Stephen Outin Phone: 702 359 1387
15) Cause of Death: End Stoge Heart foilure
16) Additional Information:
If the cause of death includes any traumatic injury, contact our office at 455-3210 immediately. If the cause of death is of natural causes, please fax this completed form to our office and the information will be entered into our database. Our FAX number is 455-3101

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Trinity Pharmacy
2797 S. Marytand Phony 28 Las Vegas, NV 89109
Phone: 702-776-8210 Fax: 702-776-7195 **Telephoned Prescription** STEPHEN DUBIN 5380 8 RAINBOW STE 308 LAS VEGAS NV 89128 DEA#: FD2419706 Lic#:: 13772 NPI#: 1023311040 Phone: 702-369-1388 Fax : 702-362-9954 Rod: 170144 Rx Written: 12-20-2018 Patient Address LAS VEGAS NV 89123 Phone # D.O.B. : 702-4 Gender LORAZEPAM O/C 2MG/ML Generio for LORAZEPAM O/8 2MG/A/L Quantity: 30 (thirty) instructions: E 0.25 M. EVERY 4 HOURS AS NEEDED FOR MILD ANXIETY, GIVE 0.5ML ERY 4 HOURS FOR MODERATE ANXIETY, GIVE 1.0 ML EVERY 4 HOURS FOR ERE ANXIETY ile Authorized: 0 RPh: MLJ/CC Signature Date THIS PRESCRIPTION WILL BE FILLED GENERICALLY UNLESS PRESCRIBER WRITES "DAW" IN THE BOX BELOW Dispense As Written

R

JIREH HEALTHCARE

2320 PASEO DEL PRADO SUITE B101 LAS VEGAS, NV 89102 TEL: 702-359-1388

	TEL: 702-3	e B101 Las 1 359-1388	/EGAS, N	V 89102
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TRINITY PHARMACY 2810 W CHARLESTON BLVD STE E44 LAS VEGAS, NV 89102 T: 702-776-8210 F: 702-776-7195

Jireh Healthcare Ser ices LLC

PHYSICIANS ORDER

2320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102 . Tel: Fax:

PATIENT:

Busine, D.

DOB:

ORDER DATE:

12/20/18

MR#:

M0487

ORDER#:

3310180

List of New/Refill Orders

ORDERED BY: STEPHEN DUBIN (MD)

Start Date	Stop Date	Туре		Order	Strength	Quantity	Dosage	Route		Frequency	Indication	Dave
12/20/18	I	Medication	New	Digoxin 0.125mg Tab	0.125 mg		Tab	Oral	6	1 tablet Daily	Afib	Paye
12/20/18	******	Medication	New	Diltiazem Hydrochloride 90mg Extended-Release Capsule	90 mg		Capsule	Oral	7	1 capsule two times daily	Hypertension/ Afib	1 _
12/20/18		Medication	New	Primidone 50mg Tab	50 mg		Tab	Oral		1 tablet by mouth daily	zeisure	1
12/20/18		Medication	New	Morphine Sulfate 20mg/1mL Solution	20 mg	,	Solution	Oral		Every 4 hours as needed for pain	Pain - Severe	Ĥ
		Give 10mg (0.5ml) sub	si) sublingual every 4 hours a lingual every 4 hours as nee igual every 4 hours as neede	ded for mod	ferate nain		-			:#1 #	
12/20/18	i	Medication	New	Lorazepam 2mg/ml Conc Soln	2 mg	-	Soln	Oral		every 4 hours as needed for anxiety/agitation	Agitation/restlessne	Н
	==>	Note: Give 0 Give 0.5ml e	.25ml ever	y 4 hours as needed for mild rs as needed for moderate a	l anxiety/ag	Itation			- 1	anxiety/agitation	55	

List of 'A	DMIT' Itei	ns - DO NOT FIL	L ITEMS BELOW (P	atient has	supply o	of item at	admission)			
Start Date	Stop Date	Туре	Order	Strength	Quantity	Dosage	Route	Frequency	Indication	Payer

PHONE ORDER / READ BACK ORDER BY MARY ALYN DOMONDON (RN)

Give 1ml every 4 hours as needed for moderate anxiety/agitation

Electronically Signed: 12/20/2018, By STEPHEN DUBIN, MD

Jireh Hearthcare Services LLC

2320 Paseo Del Prado, Suite B101, Las Vegas, NV 89102. Tel: Fax:

Patient Name: Burner, D.

MR#: M0487

HospiceMD

CURRENT TREATMENT / MEDICATION / DME LIST

All medications have been reviewed for effectiveness of drug therapy, drug side effects, actual or potential drug / food interactions, duplicate drug therapy and drug currently associated with laboratory monitoring.

				The second secon	THE PERSON NAMED IN COLUMN	1				
Order Date	Order Date Start Date Stop Date Type	Stop Date	Type	CEDS/DIJE	Strength Quantit	Quantity Dosage	Route	[Fraguenc)	Indication	Payer Administered By
12/20/18	12/20/18		Medication	Medication Digoxin 0.125mg Tab	0.125 mg	Tab	Oral	1 tablet Daily	Afib	3&4
12/20/18	12/20/18		Medication	Diltiazem Hydrochloride 90mg Extended-Release Capsule	60 шб	Capsule	Oral	1 capsule two times daily	Hypertension/ Afib I	3&4
12/20/18	12/20/18		Medication	Medication Lorazepam 2mg/ml Conc Soln	2 mg	Soln	Oral	every 4 hours as needed for anxiety/agitation	Agitation/restless H ness	384
		A II	Note: Give (Give 0.5ml (Give 1ml ev	==> Note: Give 0.25ml every 4 hours as needed for mild anxiety/agitation Give 0.5ml every 4 hours as needed for moderate anxiety/agitation Give 1ml every 4 hours as needed for moderate anxiety/agitation	mild anxiety/agitation ate anxiety/agitation e anxiety/agitation			,		
12/20/18	12/20/18		Medication	Medication Morphine Sulfate 20mg/1mL. Solution	20 mg	Solution	Oral	Every 4 hours as needed for pain	Pain - Severe H	3&4
		<u> </u>	Note: Give (Give 10mg (Give 20mg (==> Note: Give 5mg (0.25ml) sublingual every 4 hours as needed for mild pain Give 10mg (0.5ml) sublingual every 4 hours as needed for moderate pain Give 20mg (1ml) sublingual every 4 hours as needed for severe pain	urs as needed for mild pair needed for moderate pain eeded for severe pain	pain bain				
12/20/18	12/20/18		Medication	Medication Primidone 50mg Tab	50 mg	Тар	Oral	1 tablet by mouth daily	zeisure	384
12/20/18	12/20/18		Treatment	Andrewsky along market, and district the second of the sec	in the construction of the state of the stat				8	
			oxygen at 3	oxygen at 3L/min via nasal canula for comfort				1		1

(3&4) - Fam/PCG & (2&4) - Self & Hospice Nurse, (2&3) - Self & Fam/PCG, ADMINISTERED BY:(1) - SNF Nurse, (2) - Self, (3) - Fam/PCG, (4) - Hospice Nurse, (1&4) - SNF & Hospice Nurse, Hospice Nurse, (2,3&4) - Self & Fam/PCG & Hospice Nurse Pane 1 of 1

Date Of Birth

Hospice Certification of Terminal Illness

Benefit Period

Benefit Period Start Date

12/20/2018

Benefit Period End Date 03/19/2019

Certification Start Date 12/20/2018

Certification End Date

03/19/2019

Signed Date 12/20/2018

Medical Director or Certifying **Physician**

Dubin, Stephen Paul

Medical Director or Certifying Physiclan Address

5741 S. Fort Apache Rd. Suite 100 Las Vegas, NV

Medical services required

89148

Attending Physician

Dubin, Stephen Paul

Attending Physician Address

5741 S. Fort Apache Rd. Suite 100 Las Vegas, NV 89148

Patient Hi Claim No.

546421189A

Disaster Acuity

today

Provider Information

Provider Name

Jireh Healthcare Services

Provider Number 291533

Provider Address

2320 Paseo Del Prado Ste B101 Las Vegas, NV 89102-0048

Admission

New Admission 12/20/2018 (ROUTINE)

Verbal Certification

Verbal Certification received from Medical Director: Dubin, Stephen Paul by Fronda, MaryRose (mary1) on 12/20/2018

I certify that Burney Demis is terminally ill with a life expectancy of six months or less if the terminal illness runs its normal course

Admin Fronda, MaryRose (User) signed on 12/20/2018. Recorded by mary1 on 12/21/2018 17:06:18.

Verbal Certification received from Physician. Dubin, Stephen Paul by Fronda, MaryRose (mary1) on 12/20/2018

I certify that Beam. Demiss terminally ill with a life expectancy of six months or less if the terminal illness runs its normal course

Admin Fronda, MaryRose (User) signed on 12/20/2018. Recorded by mary1 on 12/21/2018 17:06:33.

Brief Narrative Statement and Attestation

Review the patient's clinical circumstances and synthesize the medical Information to provide clinical justification for admission and continuation to hospice services. If in third (3rd) or more benefit periods, clinical findings from the face to face encounter have been used to determine continued eligibility for hospice care.

Dubin, Stephen Paul

Composing Physician Mr. Bearis an 83 yr old Caucasian male, lives in a group home with 24/7 assistance, POA present at time of assessment. Patient's POA Stella Urbin signed consent as the Power of Attorney. Patient admitted to Jireh hospice under routine level of care, with terminal diagnosis of end stage Heart Failure. somnolerice, sleeps most of the visit, confused and anxious. Co-morbidities include atrial fibrillation/flutter, hypertension, BPH, severe weakness, latigue, severe SOB on non re-breather oxygen 15 l/min continuously. Patient has a worsening generalized and worsening adult failure to thrive with consequent loss of weight, MAC 20.7cm, POA claimed that patient to be over 180 lbs prior to this decline, dependent on ADLs, bed confined with complete assist, PPS 30 [KPS 30. Patient is incontinent of bladder and bowel, Patient is DNR, no agressive treatment nor hospitalization per POA request. Medications reviewed and POC discussed with POA, verbally agreed on plan. IDG notified.

Recorded By

Fronda, MaryRose

Attestation. I confirm that I composed this narrative and that it is based on my review of the patient's medical record and/or examination of the patient.

Medical Director Dublin, Stephen Paul (Medical Director) signed on 12/20/2018. Recorded by mary1 on 12/21/2018 17:11:45

Consolo Certification Dated: 12/20/2018 for Barry, Date



Certification Statement

I certify that this patient is under my care, and to the best of my medical knowledge given the data available, has a life expectancy of six (6) months or less if the terminal illness runs its normal course.

Medical Director or CertifyIng Medical Director Dubin, Stephen Paul (Medical Director) signed on 12/20/2018. Recorded by mary1 on Physician: 12/21/2018 17:11:53.

I certify that this patient is under my care, and to the best of my medical knowledge given the data available, has a life expectancy of six (6) months or less if the terminal illness runs its normal course.

Attending Physician: Medical Director Bubin, Stephen Paul (Medical Director) signed on 12/20/2018. Recorded by mary1 on 12/21/2018 17 11:53.

Additional Signatures

1. Admin Fronda, MaryRose (User) signed on 12/21/2018. Recorded by mary1 on 12/21/2018 17:12:03.

JIREH HEALTHCARE SERVICES

2320 Paseo Del Prado Suite B101 Las Vegas Nevada 89102 PHONE: 702,359,1388 FAX: 702,359,2388 EMAIL ADDRESS: jirehhealthcare @gmail.com

PHYSICIAN'S CERTIFICATION FOR MEDICARE/MEDICAID HOSPICE BENEFIT

Physician's Certification of Terminal Illness for Hospice Benefit
(Part - 1)

MR# F10436 CERTIFICATION STATEMENT First 90-Day Period from 12/20/2018 to 03/14/2019 Dan Barre is terminally ill and I (or WE) certify that ___ Has a life expectancy of six (6) months or less if the terminal illness runs its normal course. I (WE) Reviewed the patient's clinical information and considered the primary terminal condition, related diagnoses, current subjective and objective medical findings, current medication and treatment orders and information about management of unrelated conditions in making this determination. Verbal Order Date Hospice Medical Director/Associate Hospice Doctor Attending Physician Signature Second 90-Day Period from RECERTIFICATION STATEMENT B. I certify that I have reviewed the clinical record prior to recertification for the above noted patient and That patient is still considered to be terminally ill and has life expectancy of six (six) months or less, if The terminal illness runs its normal course. Date Hospice Medical Director/Associate Hospice Doctor 60-Day Period from RECERTIFICATION STATEMENT C. I certify that I have reviewed the clinical record prior to recertification for the above noted patient and That patient is still considered to be terminally ill and has life expectancy of six (6) months or less, if The terminal illness runs its normal course. Date Hospice Medical Director/Associate Hospice Doctor

DUBINO18

Jireh Healthcare Services LLC

PHYSICIANS ORDER

88 South Rainbow Blvd Suite A, Las Vegas, NV 89146 . Tel: Fax:

PATIENT: ORDERED BY: STEPHEN DUBIN (MD)

Bush, D/

DOB:

ORDER DATE:

12/19/18

MR#:

M0487

ORDER#: 4672846

ALLERGIES:

List of New/Refill Orders

Start Date Stop Date Type Order

Strength

Quantity Dosage

Route

Frequency

Indication

Payer

12/19/18

Other New

Please do hospice evaluation and treat

List of 'ADMIT' Items - DO NOT FILL ITEMS BELOW (Patient has supply of item at admission)

Start Date Stop Date Type

Order Strength Quantity Dosage

Frequency

Indication

Payer

PHONE ORDER / READ BACK ORDER BY MARY ALYN DOMONDON (RN)

Electronically Signed: 12/19/2018, By STEPHEN DUBIN, MD

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