NEVADA STATE BOARD OF PHARMACY

431 W Plumb Lane - Reno, NV 89509 - (775) 850-1440

APPLICATION FOR OUT-OF-STATE OUTSOURCING FACILITY LICENSE

\$500.00 Fee made payable to: Nevada State Board of Pharmacy (non-refundable and not transferable money order or cashier's check only)

Application must be printed legibly or typed

Any misrepresentation in the answer to any question on this application is grounds for refusal or denial of the application or subsequent revocation of the license issued and is a violation of the laws of the State of Nevada.

MagNew Outsourcing Facility □ Ownership Change (Provide current license number if making changes:) OUT □ 503a OR 🕱 503b Apply as retail pharmacy only.				
Check box below for type of ownership and complete all required forms for type of ownership that you have selected. If LLC use Non Publicly Corporation or Partnership □ Publicly Traded Corporation – Pages 1-3 & 4 □ Partnership - Pages 1-3 & 6 Non Publicly Traded Corporation – Pages 1-3 & 5 □ Sole Owner – Pages 1-3 & 7				
GENERAL INFORMATION to be completed by all types of ownership				
Facility Name: Cantrell Drug Company				
Physical Address: 7321 Cantrell Road				
City: Little Rock State: Arkansas Zip Code: 72207				
Telephone: 501-663-3642 Fax: 501-296-9936				
Toll Free Number: 877-666-5222 (Required per NAC 639.708)				
E-mail: kallen@cantrelldrug.com Website: www.cantrelldrug.com				
Supervising Pharmacist: Ashley D. Wagner Nevada License #: 19708				
SERVICES PROVIDED				
Yes/No				
🚨 🛘 Parenteral				
☑ □ Sterile Compounding				
☐ 🖾 Mail Service Sterile Compounding				
☐ 💆 Other Services:				
All boxes must be checked for the application to be complete				
An appearance will be required at a board meeting before the license will be issued.				
Roard Use Only Date Processed: Amount: \$ 500,00				

APPLICATION FOR OUT-OF STATE OUTSOURCING FACILITY - Page 3

I hereby certify that the answers given in this application and attached documentation are true and correct. I understand that any infraction of the laws of the State of Nevada regulating the operation of an authorized Outsourcing Facility may be grounds for the revocation of this permit.

I have read all questions, answers and statements and know the contents thereof. I hereby certify, under penalty of perjury, that the information furnished on this application are true, accurate and correct. I hereby authorize the Nevada State Board of Pharmacy, its agents, servants and employees, to conduct any investigation(s) of the business, professional, social and moral background, qualification and reputation, as it may deem necessary, proper or desirable. The facility must be registered with the FDA as an outsourcing facility (503B) to obtain an outsourcing facility from the Board of Pharmacy.

Federal and State law require a licensed pharmacist to supervise the compounding taking place in a registered outsourcing facility. This supervising pharmacist must be licensed by the Nevada Board of Pharmacy.

Does your outsourcing facility wholesale compounded medication for resale? Yes

No

No

The Law prohibits the resale of compounded medication. By signing this application you are attesting that your medications will be labeled with the statement "Not for Resale" and that the outsourcing facilities products will not be resold.

Original Signature of Person Authorized to Submit Application, no copies or stamps

JAMES L. MILLARLEY, DR CEO

Print Name of Authorized Person

Date

Nevada State Board of Pharmacy 431 W. Plumb Lane Reno, Nevada 89509

Ladies and Gentlemen:

In reference to question #2, Georgia Board of Pharmacy denied our application for a Manufacturer license. We will provide additional documentation to Georgia Board of Pharmacy. We are currently licensed in Georgia as a Wholesaler Pharmacy. See attached letter from Georgia Board of Pharmacy.

In reference to question #3, after an FDA Inspection of our Outsourcing Facility and our remediation of all the observations of the FDA, the Boards of Pharmacy in South Carolina, Alabama, and Illinois asked for additional information and suspended our right to ship into those Sates until they are satisfied. Upon receiving our newly acquired Verified-Accredited Wholesale Distributors certification, Illinois has reinstated our right to ship. The South Carolina and Alabama Boards of Pharmacy have now held hearings. Following the hearing in South Carolina, its Board concluded that Cantrell Drug Company will be placed on a two year probationary period. Following the hearing in Alabama, its Board concluded that there had been a deficiency in sterile compounding and imposed a fine. See attached final orders from Alabama Board of Pharmacy and South Carolina Board of Pharmacy.

The Boards of Pharmacy in Colorado, Florida, Indiana, Missouri, and Minnesota have investigated the same facts surrounding this FDA Inspection and allowed us to continue shipping. In Florida, we have voluntarily agreed to restrict our practice in the state until we have a new Florida-approved inspection of our facility.

Also in reference to question #3, James L. McCarley was disciplined by the Kentucky Board of Pharmacy due to a miscalculation in completing continuing education credits which has now been rectified.

In reference to question #4, in 2003, the Drug Enforcement Administration investigated Cantrell Drug Company for an alleged violation of Title 21 USC in regard to compounded intrathecal pump refills sent to the ordering physician for administration by the physician. This practice is standard in most compounding pharmacies dispensing intrathecal medication refills in the United States. A settlement was reached in 2004 upon the terms set forth in a written agreement, a copy of which is attached. Furthermore, Cantrell Drug Company complied with DEA request to register the pharmacy as a "manufacturer" with the agency.

Let me know if you need further information.

Dell McCarley, Pharm D

CEO

ALABAMA BOARD OF PHARMACY

SUSAN ALVERSON R.Ph. Executive Secretary

<u>Location</u>: 111 Village Street Birmingham, AL 35242

(205) 981-2280 (205) 981-2330 Fax www.albop.com



MEMBERS 2017

BUDDY BUNCH, R.Ph. President

DAVID DARBY, R.Ph. Vice President

DONNA YEATMAN, R.Ph. Treasurer

RALPH SORRELL, R.Ph.

Brenda Denson, PharmD.

May 4, 2017

CANTRELL DRUG COMPANY 7321 CANTRELL RD LITTLE ROCK AR 72207

RE:

Final order

Dear Cantrell Drug Company:

Enclosed you will find a FINAL ORDER resulting from your hearing before the Board. While the entire order is important, I particularly direct your attention to the portion of the Order setting forth discipline and specifically the mandatory obligation of your payment of a fine and costs. As you will see, those amounts are due within a specified period of time from the date of the Final Order and not the date of this letter.

If the referenced fine and costs are not received by the Board within the prescribed period of time, or special arrangements have not been made with the Secretary of the Board, the Board will file a lawsuit to enforce the Final Order which can result in the entry of a judgment against you and subsequent collection procedures.

Sincerely,

Wendy Passmore

Legal / Executive Assistant Alabama State Board of Pharmacy Phone 205-981-4764 Fax 205-803-6481 Email - wpassmore@albop.com

- The Respondent made no objection to the timeliness of the Notice of Hearing.
- 4. The Respondent committed and is guilty of the acts specified as violations in the Statement of Charges and Notice of Hearing dated December 28, 2016 as Amended on March 23, 2017.

Conclusions of Law

- 1. The Alabama State Board of Pharmacy has jurisdiction in this cause pursuant to Code of Alabama (1975). § 34-23-32. § 34-23-32.1, § 34-23-34. § 34-23-92 (11) and (12) and Code of Alabama (1975). § 41-22-12.
- 2. The Respondent was properly notified of the charges; the Respondent was represented at the administrative hearing by counsel.
 - 3. The Respondent made no objection to the timeliness of the Notice of Hearing.
- 4. The Respondent's permit as a manufacturer wholesaler/distributor in the State of Alabama is due to be have disciplinary sanctions imposed in that it is guilty of the acts specified in Count One of the Statement of Charges and Notice of Hearing dated December 28, 2016 and as Amended on March 23, 2017.

ORDER

In accordance with the foregoing Stipulation and Agreement. Findings of Fact and Conclusions of Law, it is hereby ORDERED as follows:

1. The Respondent is ORDERED to pay to the Board an administrative fine of Thirty Thousand (\$30,000.00) Dollars: said fine shall be paid in sixty (60) days from the date of this Final Order; and

SOUTH CAROLINA DEPARTMENT OF LABOR, LICENSING AND REGULATION BEFORE THE BEFORE THE STATE BOARD OF PHARMACY

IN THE MATTER OF:

PY.10776 & PY.16647

CANTRELL DRUG COMPANY
CANTRELL DRUG COMPANY INC
7321 Cantrell Rd.
Little Rock, AR 72207

OIE # 2016-149

FINAL ORDER (PUBLIC)

Respondent.

On March 15, 2017, the above licensing board ("Board"), with a quorum present, held a hearing on the Memorandum of Agreement and Stipulations ("MOA") in the above referenced matter entered into between the State and Respondent. The Board also heard Respondent's Petition to Resume Shipping Compounded Products. Patrick Hanks, Esquire, Chief Disciplinary Counsel, represented the State. Respondent was represented by Jon Wallace, Esquire. Dell McClary, CEO of Respondent; Dr. Eric Goode, Interim Chief of Compliance and Regulatory Affairs; and Ashley Wagener, Pharmacist in Charge, appeared on behalf of Respondent.

FINDINGS OF FACT

- 1. Respondent was properly served with a Notice of Hearing.
- 2. In the MOA, Respondent admitted to the following, which the Board adopts:
 - a. Respondent is an FDA Registered Outsourcing Facility under Section 503B of the Federal Food, Drug, and Cosmetic Act and is permitted in this state as a Nonresident Outsourcing facility, duly permitted by the State Board of Pharmacy (the "Board") in this State, and was so permitted at all times relevant to the matters asserted herein; thus, the Board has jurisdiction over this matter.
 - b. As a registered Outsourcing Facility under Section 503B of the Federal Food, Drug, and Cosmetic Act, Respondent must comply with cGMP requirements, be routinely inspected by FDA, and must meet certain other conditions, such as adverse event reporting, among other requirements.
 - c. FDA conducted an Outsourcing Facility Inspection of Respondent ending on October 14, 2016. As a result of the inspection, FDA issued Form 483 observations.
 - d. Respondent fully responded with a corrective action plan to the FDA Dallas District Office on November 4, 2016.

(2012, as amended) provides that upon determination by the Board that one or more grounds for disciplining a licensee or permittee exist, the Board may impose a fine of \$500 per violation, not to exceed a total of \$25,000 per action, plus the costs of the disciplinary action.

- 5. In this case, the Board concludes that Respondent may resume shipping compounded drugs into South Carolina subject to its compliance with certain conditions. First, Respondent's permit shall be placed on a probationary status for a minimum of two years. Prior to resuming shipping, Respondent must provide an inspection by the Arkansas Board of Pharmacy, which must be approved by the Board Administrator. Further, Respondent must provide an FDA End of Inspection ("EIR") Report indicating no further disciplinary action taken by the FDA. During Respondent's probationary period, Respondent must report any and all correspondence with the FDA to the Board Administrator. Respondent must reappear before the Board to have the probation lifted.
- 6. The sanctions and conditions imposed by this Order are within the scope of those permitted by S.C. Code Ann. §§ 40-1-120 and 40-43-150 (2011) and are designed not to punish the Respondent but to protect the life, health and welfare of the people at large.

IT IS THEREFORE ORDERED:

- 1. The Board accepts the MOA and finds that Respondent violated the Pharmacy Practice Act.
- 2. The Petition to resume shipping compounded products is granted, subject to Respondent's submission to, and approval by, the Board Administrator of a new Arkansas inspection report. Upon receipt and approval of the same, the license shall be immediately placed on a probationary status for a period of two years, subject to the following conditions of probation: 1) Respondent must provide the Board with an FDA EIR Report indicating no further violations; 2) Respondent shall submit all correspondence, documentation, etc. received by the FDA to the Board; and 3) Respondent must reappear before the Board to have its permit removed from probationary status.

AND IT IS SO ORDERED.

STATE BOARD OF PHARMACY

Carole Small Russell, R.Ph.

Board Chair

September 19, 2017



KENTUCKY BOARD OF PHARMACY

Matt Bevin Governor

> State Office Building Annex, Suite 300 125 Holmes Street Frankfort KY 40601 Phone (502) 584-7910 Fax (502) 696-3808 http://pharmacy.kv.goy

Board Members
Deborah L. Brewer, R.Ph.
Brian C. DeWire, DC, Consumer
Scott A. Greenwell, Pharm.D.
Cathy Hanna, Pharm.D.
Craig Martin, Pharm D,
Ron Poole, R.Ph.

Executive Director B. Steven Hart, R.Ph.

February 22, 2017

James McCarley Jr 7700 Northshore Place North Little Rock AR 72118

Re: Case No. 17-0202

Dear Pharmacist,

This letter follows a recent investigation by Board staff.

The purpose of this letter is to offer you an opportunity to informally resolve this matter through an Agreed Order prior to the filing of a formal Complaint. Find enclosed a proposed Agreed Order setting forth terms I believe the Board will accept.

Review the proposed Agreed Order carefully. Feel free to consult with legal counsel. If acceptable, sign and return the Agreed Order to the Board office by March 22, 2017. Upon receipt, the proposed Agreed Order will be signed by the Board President, and a copy will be sent to you.

If this proposed Agreed Order is unacceptable and you in good faith believe this matter can be resolved, please feel free to contact me during normal business hours.

Should you fail to respond by returning the proposed Agreed Order or contacting me by March 22, 2017, your case will be referred to the Office of the Attorney General to conduct an administrative hearing.

Sincerely.

Steve Hart, R.Ph. Executive Director

Enclosure



- (B) On or before March 22, 2017, Respondent shall submit to the Board office proof of no less than ten (10) continuing education hours, which programs shall not be used in any way to satisfy Respondent's continuing education requirements for renewal.
- (C) By entering into this Agreed Order, Respondent expressly acknowledges that the Respondent was fully and completely informed of Respondent's right to due process, that the Respondent fully understands those rights, and that the Respondent knowingly, voluntarily, and willingly agrees to waive those rights and to enter into this Agreed Order.
- (D) The above information shall be reported to the National Association of Boards of Pharmacy ("NABP") and is subject to disclosure under the Kentucky Open Records Act.

Date	
	Date

James McCarley Jr, Respondent

3-15-11 Date

UNITED STATES DISTRICT COURT EASTERN DISTRICT OF ARKANSAS WESTERN DIVISION

UNITED STATES OF AMERICA

V.

USAO: 2004V00173

CANTRELL DRUG STORE DELL MCCARLEY

SETTLEMENT AGREEMENT

This Settlement Agreement is made and entered into this 20 day of September, 2004, by and among the United States of America, acting through the United States Attorney for the Eastern District of Arkansas (hereinafter referred to as "USAO""), and Cantrell Drug Company.

PREAMBLE

WHEREAS, the United States contends that Cantrell Drug Company has violated 21 U.S.C. § 828(a), § 829(a) and § 842(a)(1), (a)(2) and (a)(5);

WHEREAS, the Cantrell Drug Company denies it has violated any provision of Title 21 U.S.C.

WHEREAS, the parties desire to reach an agreement that would settle, compromise and resolve the United States' claims under Title 21 U.S.C. in order to avoid the expense and uncertainty of litigation.

TERMS OF AGREEMENT

NOW, THEREFORE, in reliance on the representations contained herein and in consideration of the mutual promises, covenants, and obligations in this Agreement, and for good and valuable consideration, receipt of which is hereby acknowledged, the parties agree as

- 5. It is understood and agreed that this Settlement Agreement is in compromise of disputed claims and that it shall not be construed as an admission of or evidence of liability or wrongdomg on the part of any of the released entities.
- 6. This document contains the complete agreement between the parties with respect to the matters herein.
- 7. This Agreement may be executed in identical counterparts, each of which shall constitute an original and all of which shall constitute one and the same agreement.
- 8. This Agreement may be modified only by a written document signed by all of the parties. No waiver of this Agreement or of any of the promises, obligations, terms or conditions hereof shall be valid unless it is written and signed by the party against whom the waiver is to be enforced.
- 9. If any part or any provision of this Agreement shall be finally determined to be invalid or unenforceable under applicable law by a court of proper jurisdiction, that part shall be ineffective to the extent of such invalidity or unenforceability only, without in any way affecting the remaining part of said provision or the remaining provision of this Agreement.
- 10. Each person who signs this Agreement in a representative capacity represents that he or she is duty authorized to do so.
 - 11. This Agreement is effective upon the date of the signature of the last signatory.

IN WITNESS WHEREOF, we have hereunder set our hand as of the date first above written.

NEVADA STATE BOARD OF PHARMACY

431 W Plumb Lane - Reno, NV 89509 - (775) 850-1440

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 New Outsourcing Facility □ Ownership Change (Provide current license number if making changes:) OUT □ 503a OR □ 503b Apply as retail pharmacy only. 				
Check <u>box</u> below for type of ownership and complete all required forms for type of ownership that you have selected. If LLC use Non Publicly Corporation or Partnership ☐ Publicly Traded Corporation – Pages 1-3 & 4 ☐ Partnership - Pages 1-3 & 6 ☐ Non Publicly Traded Corporation – Pages 1-3 & 5 ☐ Sole Owner – Pages 1-3 & 7				
GENERAL INFORMATION to be completed by all types of ownership				
Facility Name:LEITERS HEALTH				
Physical Address: 13796 COMPARK BLVD				
City: ENGLEWOOD State: CO Zip Code: 80112 Telephone: (800) Fax: 408-288-8252				
Toll Free Number: 800-292-6772 (Required per NAC 639.708)				
E-mail: DenverRegulatoryAdmin@Leiters.com Website: www.Leiters.com				
Supervising Pharmacist: STACY ANN HAVER Nevada License #: 19777				
SERVICES PROVIDED				
Yes/No				
⊠ □ Parenteral				
☑ ☐ Sterile Compounding				
□ 💢 Non Sterile Compounding				
□ 🙀 Mail Service Sterile Compounding				
□ ☑ Other Services:				
All boxes must be checked for the application to be complete				
An appearance will be required at a board meeting before the license will be issued.				
Board Use Only Date Processed: Amount:				

APP	LICATION FOR OUT-OF STATE OUTSOURCING FACILITY	Page 2
FELI	Number (From FDA application): 3013438582	
Plea Di	se provide the name of the facility as registered with the FDA and the registr ENVER SOLUTIONS, LLC dba LEITERS HEALTH 3013438582	ration number:
Plea:	se provide a list of all DBA's used by outsourcing facility. A separate sheet i	s acceptable.
	se provide the name and Nevada license number of the supervising pharma e: STACY ANN HAVER Nevada License Number: 19	cist: 777
	vada business license is not required, however if the Outsourcing Facility hates license please provide the number: n/a	as a Nevada
	page must be submitted for all types of ownership.	
Withi	n the last five (5) years:	
1)	Has the corporation, any owner(s), shareholder(s) or partner(s) with any interest, ever been charged, or convicted of a felony or gross misdemeanor (including by way of a guilty plea or no contest plea)?	Yes □ No 🏻
2)	Has the corporation, any owner(s), shareholder(s) or partner(s) with any interest, ever been denied a license, permit or certificate of registration?	Yes □ No 🛚
3)	Has the corporation, any owner(s), shareholder(s) or partner(s) with any interest, ever been the subject of an administrative action, board citation, cite fine or proceeding relating to the pharmaceutical industry?	Yes □ No 🔀
/ 1)	Has the corporation and compar(s) at sociality ()	

4) Has the corporation, any owner(s), shareholder(s) or partner(s) with any interest, ever been found guilty, pled guilty or entered a plea of nolo contendere to any offense federal or state, related to controlled substances?

Yes □ No 🏻

Has the corporation, any owner(s), shareholder(s) or partner(s) with any interest, ever surrendered a license, permit or certificate of registration voluntarily or otherwise (other than upon voluntary close of a facility)?

Yes □ No Ⅸ

If the answer to question 1 through 5 is "yes", a signed statement of explanation must be attached. Copies of any documents that identify the circumstance or contain an order, agreement, or other disposition may be required.

APPLICATION FOR OUT-OF STATE OUTSOURCING FACILITY - Page 3

I hereby certify that the answers given in this application and attached documentation are true and correct. I understand that any infraction of the laws of the State of Nevada regulating the operation of an authorized Outsourcing Facility may be grounds for the revocation of this permit.

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Federal and State law require a licensed pharmacist to supervise the compounding taking place in a registered outsourcing facility. This supervising pharmacist must be licensed by the Nevada Board of Pharmacy.

Does your outsourcing facility wholesale compounded medication for resale? Yes □ No □

The Law prohibits the resale of compounded medication. By signing this application you are attesting that your medications will be labeled with the statement "Not for Resale" and that the outsourcing facilities products will not be resold.

Original Signature of Person Authorized to Submit Application, no copies or stamps

ROBIN SMITH HOKE

3/2

Print Name of Authorized Person

APPLICATION FOR OUT-OF-STATE OUTSOURCING FACILITY

Page 5

OWNERSHIP IS A NON PUBLICLY TRADED CORPORATION

State of Incorporation: Delaware	
Parent Company if any:See attached	
Address: See attached	
City: <u>See attached</u> State: <u>See attached</u> Zip: <u>see attached</u>	
Fax: (408) 288- 8252	
Contact Person: Robin S. Hoke, President & CEO	
or any corporation non publicly traded, disclose the following:	
) List top 4 persons to whom the shares were issued by the corporation?	
a) See attached Name Address	
Name Address	
b)	
Name Address	
c)	
Name Address	
d) Name Address	
Name Address	
Provide the number of shares issued by the corporationn/ 9	
) What was the price paid per share?	
) What date did the corporation actually receive the cash assets?h a	
Provide a copy of the corporation's stock register evidencing the above information.	l a

Include with the application for a non publicly traded corporation

<u>Certificate of Corporate Status</u> (also referred to as Certificate of Good Standing). The Certificate is obtained from the Secretary of State's office in the State where incorporated. The Certificate of Corporate status must be dated within the last 6 months.

List of officers and directors

NEVADA STATE BOARD OF PHARMACY

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GENERAL INFORMATION to be completed by all types of ownership Facility Name: Stokes Healthcare Inc.				
and a first control of the above the first control of the control				
Physical Address: 8000 Commerce Parkway, Suite 600				
City: Mt. Laurel State: NJ Zip Code: 08054				
Telephone: 800-754-5222 Fax: 856-505-5899				
Toll Free Number: 800-754-5222 (Required per NAC 639.708)				
E-mail: licensing@stokespharmacy.com Website: www.stokespharmacy.com				
Supervising Pharmacist: Emmett McVey Nevada License #: 19796				
SERVICES PROVIDED				
Yes/No				
□ ☑ Parenteral				
☑ □ Sterile Compounding				
☑ □ Non Sterile Compounding				
☑ □ Mail Service Sterile Compounding				
□ Other Services:				
All boxes must be checked for the application to be complete				
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Board Use Only Date Processed: Amount: #500-00				

FEIN	lumber (From FDA application):3002815949	
	e provide the name of the facility as registered with the FDA and the registr tokes Healthcare Inc. 3002815949	ation number:
	e provide a list of all DBA's used by outsourcing facility. A separate sheet is kes Pharmacy	s acceptable.
	e provide the name and Nevada license number of the supervising pharma e: Emmett McVey Nevada License Number:	
	vada business license is not required, however if the Outsourcing Facility hat ess license please provide the number:NA	s a Nevada
This p	page must be submitted for all types of ownership.	
Withir	the last five (5) years:	
1)	Has the corporation, any owner(s), shareholder(s) or partner(s) with any interest, ever been charged, or convicted of a felony or gross misdemeanor (including by way of a guilty plea or no contest plea)?	Yes □ No ☑
2)	Has the corporation, any owner(s), shareholder(s) or partner(s) with any interest, ever been denied a license, permit or certificate of registration?	Yes □ No ☑
3)	Has the corporation, any owner(s), shareholder(s) or partner(s) with any interest, ever been the subject of an administrative action, board citation, cite fine or proceeding relating to the pharmaceutical industry?	Yes ☑ No □
4)	Has the corporation, any owner(s), shareholder(s) or partner(s) with any interest, ever been found guilty, pled guilty or entered a plea of nolo contendere to any offense federal or state, related to controlled substances?	Yes □ No ☑
5)	Has the corporation, any owner(s), shareholder(s) or partner(s) with any interest, ever surrendered a license, permit or certificate of registration voluntarily or otherwise (other than upon voluntary close of a facility)?	Yes □ No ☑

Page 2

APPLICATION FOR OUT-OF STATE OUTSOURCING FACILITY

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APPLICATION FOR OUT-OF STATE OUTSOURCING FACILITY - Page 3

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MM	2				
Original Signature of Pe	erson Author	ized to Subr	mit Application	on, no copies or	stamps
Michael Tursi				1-30-18	
Print Name of Authorize	ed Person		Annual all and Discussion of the Control of the C	Date	

APPLICATION FOR OUT-OF-STATE OUTSOURCING FACILITY

Page 5

OWNERSHIP IS A NON PUBLICLY TRADED CORPORATION

State of Incorporation: New Je	ersey					
Parent Company if any:						
Address: 8000 Commerce Parkw		•				
City: Mt. Laurel	State: NJ	Zip: _	08054			
Telephone: _800-754-5222						
Contact Person: Michael Turs						
For any corporation non publicly 1) List top 4 persons to who			noration?			
	in the shares were issu	ied by the con	porauon <i>:</i>			
a) See attached.						
Name	Address					
b)						
b)Name	Address					
c)						
Name	Address					
d)						
Name	Address					
2) Provide the number of sh	Provide the number of shares issued by the corporation.					
3) What was the price paid	What was the price paid per share?					
4) What date did the corpor	What date did the corporation actually receive the cash assets?					
5) Provide a copy of the cor	Provide a copy of the corporation's stock register evidencing the above information					

Include with the application for a non publicly traded corporation

<u>Certificate of Corporate Status</u> (also referred to as Certificate of Good Standing). The Certificate is obtained from the Secretary of State's office in the State where incorporated. The Certificate of Corporate status must be dated within the last 6 months.

List of officers and directors



8000 Commerce Parkway, Suite 600, Mount Laurel, NJ 08054 p: 800-754-5222 f: 800-440-5899

Stokes Healthcare Inc. Corporate Officers are as follows:

Emmett McVey, RPh - 50%

T: 609-471-1326

E. Monterey Ave., #601 Wildwood Crest, NJ 08260 Vice President/Owner Pharmacist – In – Charge

Michael Tursi - 50%

T: 609-471-1295

Union Mill Road

Mt. Laurel, NJ 08054

President/Owner