Proposed Regulation of the Nevada State Board of Pharmacy

Workshop July 20th, 2017

Explanation – Language in blue italics is new; language in red text [omitted material] is language to be omitted, and language in green text indicates prior Board-approved amendments that are in the process of being codified.

AUTHORITY: §1, NRS 639.070

1. A REGULATION relating to the transmission of information to the Prescription Monitoring Program. Modification of Day Supply Reporting and Addition of ICD-10 Code (International Classification of Diseases Tenth Revision). Adding Schedule V drugs to the list of controlled substances that must be reported to the Prescription Monitoring Program.

NAC 639.926 Transmission of information regarding dispensing of controlled substances to certain persons. (NRS 639.070)

1. Each pharmacy and practitioner that uses a computerized system to record information concerning prescriptions and that dispenses a controlled substance that is listed in schedule II, III, or IV, or V to a person who is not an inpatient of a hospital, correctional institution or nursing facility shall transmit to the Board or its agent the following information, as applicable, set forth in the 2011 ASAP Version 4.2 Standard for Prescription Monitoring Programs published by the American Society for Automation in Pharmacy. The following Segments and the accompanying Data Elements of the Implementation Guide for the 2011 ASAP Version 4.2 Standard for Prescription Monitoring Programs are hereby adopted by reference:

(a) The Segment entitled “TH Transaction Header” and the following Data Elements:
(1) Version/Release Number;
(2) Transaction Control Number;
(3) Transaction Type;
(4) Response ID;
(5) Creation Date;
(6) Creation Time;
(7) File Type; and
(8) Segment Terminator Character;

(b) The Segment entitled “IS Information Source” and the following Data Elements:
(1) Unique Information Source ID;
(2) Information Source Entity Name; and
(3) Message;

(c) The Segment entitled “PHA Pharmacy Header” and the following Data Elements:
(1) National Provider Identifier (NPI);
(2) DEA Number;
(3) Pharmacy or Dispensing Prescriber Name;
(4) Phone Number;
(5) Contact Name; and
(6) Chain Site ID;

(d) The Segment entitled “PAT Patient Information” and the following Data Elements:
(1) Last Name;
(2) First Name;
(3) Address Information - 1;
(4) City Address;
(5) State Address;
(6) ZIP Code Address;
(7) Phone Number;
(8) Date of Birth; and
(9) Gender Code;

(e) The Segment entitled “DSP Dispensing Record” and the following Data Elements:
   (1) Reporting Status;
   (2) Prescription Number;
   (3) Date Written;
   (4) Refills Authorized;
   (5) Date Filled;
   (6) Refill Number;
   (7) Product ID Qualifier;
   (8) Product ID;
   (9) Quantity Dispensed;
   (10) Days Supply, the fewest number of days necessary to consume the quantity as determined by the practitioner;
   (11) Transmission Form of Rx Origin Code;
   (12) Classification Code for Payment Type; and
   (13) Date Sold;
   (14) ICD-10 Code (International Classification of Diseases Tenth Revision) for which the prescription was prescribed;

(f) The Segment entitled “PRE Prescriber Information” and the following Data Elements:
   (1) National Provider Identifier (NPI);
   (2) DEA Number;
   (3) DEA Number Suffix;
   (4) Last Name;
   (5) First Name; and
   (6) Phone Number;

(g) The Segment entitled “CDI Compound Drug Ingredient Detail” and the following Data Elements:
   (1) Compound Drug Ingredient Sequence Number;
   (2) Product ID Qualifier;
   (3) Product ID;
   (4) Component Ingredient Quantity; and
   (5) Compound Drug Dosage Units Code;

(h) The Segment entitled “TP Pharmacy Trailer” and the Data Element Detail Segment Count; and

(i) The Segment entitled “TT Transaction Trailer” and the following Data Elements:
   (1) Transaction Control Number; and
   (2) Segment Count.

2. A copy of the publication may be obtained from the American Society for Automation in Pharmacy at the Internet address http://www.asapnet.org, or by telephone at (610) 825-7783, for the price of $175 for members and $875 for nonmembers.

3. A pharmacy that dispenses a controlled substance and is required to transmit information to the Board or its agent pursuant to subsection 1 shall transmit the information not later than the end of the
next business day after dispensing the controlled substance. A pharmacy that does not dispense a controlled substance as specified in subsection 1 shall transmit to the Board or its agent a zero report stating that the pharmacy did not dispense such a controlled substance on the immediately preceding business day.

4. The information required pursuant to this section or a zero report must be transmitted by means of:
   (a) A secure file transfer protocol;
   (b) An upload from an Internet web portal; or
   (c) A manual entry.