

**WEST VIRGINIA  
SECRETARY OF STATE  
KEN HECHLER  
ADMINISTRATIVE LAW DIVISION**

Form #2

Do Not Mark In This Box

**FILED**  
JUL 3 8 53 AM '00

OFFICE OF WEST VIRGINIA  
SECRETARY OF STATE

**NOTICE OF A COMMENT PERIOD ON A PROPOSED RULE**

AGENCY: West Virginia Board of Pharmacy TITLE NUMBER: 15

RULE TYPE: Legislative CITE AUTHORITY: 30-5-1, 60A-1-101

AMENDMENT TO AN EXISTING RULE: YES  NO

IF YES, SERIES NUMBER OF RULE BEING AMENDED: 2

TITLE OF RULE BEING AMENDED: Rules and Regulations of the Board of Pharmacy for the Uniform  
Controlled Substances Act

IF NO, SERIES NUMBER OF RULE BEING PROPOSED: \_\_\_\_\_

TITLE OF RULE BEING PROPOSED: \_\_\_\_\_

IN LIEU OF A PUBLIC HEARING, A COMMENT PERIOD HAS BEEN ESTABLISHED DURING WHICH ANY INTERESTED PERSON MAY SEND COMMENTS CONCERNING THESE PROPOSED RULES. THIS COMMENT PERIOD WILL END ON August 4, 2000 AT 4:00 p.m. ONLY WRITTEN COMMENTS WILL BE ACCEPTED AND ARE TO BE MAILED TO THE FOLLOWING ADDRESS:

West Virginia Board of Pharmacy  
232 Capitol Street  
Charleston, WV 25301

THE ISSUES TO BE HEARD SHALL BE LIMITED TO THIS PROPOSED RULE.

*W. S. In.*

Authorized Signature

ATTACH A **BRIEF** SUMMARY OF YOUR PROPOSAL

**SUMMARY OF PROPOSED RULE**

**TITLE 15**

**LEGISLATIVE RULES**

**SERIES 2**

**RULES AND REGULATIONS OF THE BOARD OF PHARMACY FOR THE UNIFORM  
CONTROLLED SUBSTANCES ACT**

This rule repeals and replaces existing series 2 which has not been amended since 1982 and removes antiquated language and brings the rule into compliance with federal regulations. The rule clarifies the requirements for controlled substance permits, establishes security requirements regarding the storage and handling of controlled substances, establishes labeling and packaging requirements for controlled substances, establishes the requirement for maintenance of records and inventories, establishes manner of issuance of prescriptions, the form of prescriptions, and time limitations on controlled substance prescriptions.

**STATEMENT OF CIRCUMSTANCES**

**TITLE 15**

**LEGISLATIVE RULES**

**SERIES 2**

**RULES AND REGULATIONS OF THE BOARD OF PHARMACY FOR THE UNIFORM  
CONTROLLED SUBSTANCES ACT**

This rule repeals and replaces existing series 2 which has not been amended since 1982 and removes antiquated language and brings the rule into compliance with federal regulations. Many changes have occurred in the past 18 years in relation to the dispensing of controlled substances and it is necessary to address those changes by amending these rules at this time.

□  
APPENDIX B

FISCAL NOTE FOR PROPOSED RULES

Rule Title: Rules and Regulations of the Board of Pharmacy for the Uniform Controlled Subs

Type of Rule:  Legislative     Interpretive     Procedural

Agency: West Virginia Board of Pharmacy

Address: 232 Capitol Street  
Charleston, WV 25301

1. Effect of Proposed rule:

	ANNUAL FISCAL YEAR				
	INCREASE	DECREASE	CURRENT	NEXT	THEREAFTER
<b>ESTIMATED TOTAL COST</b>	N/A	N/A	N/A	N/A	N/A
<b>PERSONAL SERVICES</b>	N/A	N/A	N/A	N/A	N/A
<b>CURRENT EXPENSE</b>	N/A	N/A	N/A	N/A	N/A
<b>REPAIRS &amp; ALTERATIONS</b>	N/A	N/A	N/A	N/A	N/A
<b>EQUIPMENT</b>	N/A	N/A	N/A	N/A	N/A
<b>OTHER</b>	N/A	N/A	N/A	N/A	N/A

2. Explanation of Above Estimates:  
Rule would not have any fiscal impact upon state government or citizens of state.

3. Objectives of These Rules:  
update and amend controlled substance regulations.

Rule Title: Rules and Regulations of the Board of Pharmacy for the Uniform Controlled Sub

4. Explanation of Overall Economic Impact of Proposed Rule:

A. Economic Impact on State Government:  
N/A

B. Economic Impact on Political Subdivisions; Specific Industries; Specific Groups of Citizens: N/A

C. Economic Impact on Citizens/Public at Large.  
N/A

Date: 07-03-2000

Signature of Agency Head or Authorized Representative:

Will S. D... ..

FILED

JUL 3 8 53 AM '00

TITLE 15  
LEGISLATIVE RULE  
BOARD OF PHARMACY

OFFICE OF WEST VIRGINIA  
SECRETARY OF STATE

SERIES 2  
RULES AND REGULATIONS OF THE BOARD OF PHARMACY  
FOR THE UNIFORM CONTROLLED SUBSTANCES ACT

**§15-2-1. General.**

1.1. Scope. -- The West Virginia Code, section nineteen, article five, chapter thirty et seq. mandates that the Board of Pharmacy shall make such Rules and Regulations, not inconsistent with law, as necessary, to carry out the purposes and enforce the provisions of this article.

1.2. Authority. -- W. Va. Code §60A.

1.3. Filing Date. -- \_\_\_\_\_

1.4. Effective Date. -- \_\_\_\_\_

1.5. Repeal of Former Rule. -- This legislative rule repeals and replaces WV 15CSR2 "Rules and Regulations of the Board of Pharmacy for the Uniform Controlled Substances Act" filed December 28, 1982 and effective December 28, 1982.

**§15-2-2. Adoption of federal law.**

2.1. The requirements of the federal regulations, Drug Enforcement Administration, Department of Justice, 21 CFR Parts 1300-1316, are adopted by the West Virginia Board of Pharmacy and licensed pharmacies shall comply with them. Any changes to the federal Controlled Substances Act and applicable federal regulations regarding controlled substances shall be incorporated by reference within this rule.

**§15-2-3 Controlled Substance Permits**

3.1. Persons required to register.

3.1.1 Every person who manufactures, distributes or dispenses any controlled substance or who proposes to engage in the manufacture, distribution or dispensing of any controlled substance shall obtain annually a controlled substance permit unless exempted by law or pursuant to Section 3.2 of these rules. Only persons actually engaged in such activities are required to obtain a registration; related or affiliated persons who are not engaged in such activities are not required to be registered. (For example, a stockholder or parent corporation of a corporation manufacturing controlled substances is not required to obtain a registration). "Registrant" means a person who has obtained a controlled substance permit from the West Virginia Board of Pharmacy.

3.2. The West Virginia Board of Pharmacy shall exempt from payment of a fee for a controlled substance permit the following persons:

3.2.1. Any official or agency of the United States Army, Navy, Marine Corps, Air Force, Coast Guard, Veterans' Administration or Public Health Service who is authorized to procure or purchase controlled substances for official use; and

3.2.2. Any official, employee or other civil officer or agency of the United States, of any state or any political subdivision or agency thereof, who is authorized to purchase controlled substances, to obtain such substances from official stocks, to dispense or administer such substances, to conduct research, instructional activities, or chemical analysis with such substances, or any

combination thereof, in the course of his official duties or employment.

3.3. In order to claim exemption from payment of a fee, the applicant shall have completed the certification on the appropriate application form, wherein the registrant's superior certifies to the status and address of the registrant and to the authority of the registrant to acquire, possess or handle controlled substances.

3.4. Exemption from payment of a fee does not relieve the registrant of any other requirements or duties prescribed by law.

3.5. Each application shall include all information called for in the form, unless the item is not applicable, in which case this fact shall be indicated.

3.6. Each application, attachment, or other document filed as part of an application, shall be signed by the applicant, if an individual; by a partner of the applicant if a partnership; or by an officer of the applicant, if a corporation, corporate, corporate division, association, trust or other entity. Another person may be authorized to sign for the applicant, if proof of authority (e.g., general power of attorney) accompanies the application.

3.7. If the applicant is a pharmacy, , it shall be necessary for the pharmacist in charge of the pharmacy to sign the application; and where the owner of such pharmacy is a person, other than the practicing pharmacist, it shall be necessary for such other person, partnership or corporation, corporate division, association, trust or other entity, to sign the application form as provided in section 3.6 in addition to any other persons required to sign said application.

3.8 If the applicant is a rest home, nursing home, hospital, orphanage, clinic, home for the aged, governmental agency or institution or other place requiring the use of pharmacist consultants or coordinators of pharmaceutical services, it shall be necessary for such consultant or coordinator to sign the application in addition to any other

persons required to sign the application.

### 3.9. Filing of application; joint filings.

3.9.1 All applications for registration shall be submitted for filing to the office of the the Board of Pharmacy.

3.9.2 Any person required to obtain more than one (1) registration may submit all applications in one (1) package. Each application must be complete and should not refer to accompanying application for required information.

### 3.10. Acceptance for filing; defective applications.

3.10.1 Applications submitted for filing are dated upon receipt. If found to be complete, the application will be accepted for filing. Application failing to comply with the requirements of this Part will not generally be accepted for filing. In the case of minor defects as to completeness the West Virginia Board of Pharmacy may accept the application for filing with a request to the applicant for additional information. A defective application will be returned to the applicant within ten (10) days following its receipt with a statement of the reason for not accepting the application for filing. A defective application may be corrected and resubmitted for filing at any time; the West Virginia Board of Pharmacy shall accept for filing any application upon resubmission by the applicant, whether complete or not.

3.10.2 Accepting an application for filing does not preclude any subsequent request for additional information pursuant to Section 3.20 of these rules and has no bearing on whether the application will be granted.

### 3.11. Additional information.

3.11.1 The West Virginia Board of Pharmacy may require an applicant to submit such documents or written statements of fact relevant to the application as it deems necessary to

determine whether the application should be granted. The failure of the applicant to provide such documents or statements within a reasonable time after being requested to do so shall be deemed to be a waiver by the applicant of an opportunity to present such documents or facts for consideration by the West Virginia Board of Pharmacy in granting or denying the application.

3.12. Amendments to and withdrawal of applications.

3.12.1 An application may be amended or withdrawn without permission of the West Virginia Board of Pharmacy at any time before the date on which the applicant receives an order to show cause pursuant to Section 3.29 of these rules, or before the date on which a notice of hearing on the application is published pursuant to section three hundred five, article three, chapter sixty-a of the West Virginia Code, whichever is sooner. An application may be amended or withdrawn with permission of the West Virginia Board of Pharmacy at any time where good cause is shown by the applicant or where the amendment or withdrawal is in the public interest.

3.12.2 After an application has been accepted for filing the request by the applicant that it be returned or the failure of the applicant to respond to official correspondence regarding the application, when sent by registered or certified mail, return receipt requested, shall be deemed to be a withdrawal of the application.

3.13. Administrative review generally.

3.13.1 The West Virginia Board of Pharmacy may inspect, or cause to be inspected, the establishment of an applicant or registrant, pursuant to article five of the Uniform Controlled Substances Act (chapter sixty-a, West Virginia Code). The West Virginia Board of Pharmacy shall review the application for registration and other information gathered by the West Virginia Board of Pharmacy regarding an applicant in order to determine whether the applicable standards of section three hundred three of the

Uniform Controlled Substances Act (chapter sixty-a, West Virginia Code), have been met by the applicant.

3.14. Applications for research in Schedule I substances.

3.14.1 In the case of an application for registration to conduct research with controlled substances in Schedule I, the West Virginia Board of Pharmacy shall determine the qualifications and competency of the applicant as well as the merits of the research protocol. The Board, in determining the merits of a research protocol, shall confer as to effective procedures to safeguard adequately against diversion of such controlled substances from legitimate medical or scientific use. If the Board finds the applicant qualified and competent and the research protocol meritorious and adequately safeguarded, it shall register the applicant unless it finds registration should be denied for reasons set forth in section three hundred three, article three of the Uniform Controlled Substances Act (chapter sixty-a, West Virginia Code).

3.14.2 If the Board is unable to find the applicant qualified or the Board finds that grounds exist for the denial of the application, it shall issue an order to show cause and, if requested by the applicant, shall hold a hearing on the application.

3.15 The controlled substance permit shall contain the name, address and registration number of the registrant, the activity authorized by the registration, the schedules of the controlled substances which the registrant is authorized to handle, the amount of fee paid (or exemption) and the expiration date of the registration. The registrant shall prominently display the controlled substance permit at the registered location.

3.16 No registration or any authority conferred thereby shall be assigned or otherwise

transferred except upon such conditions as the West Virginia Board of Pharmacy may specifically designate and then only pursuant to its written consent.

#### §15-2-4. Security Requirements.

##### 4.1. Security requirements.

##### 4.2. Security requirements generally.

4.2.1 All applicants and registrants shall provide effective controls and procedures to guard against theft and diversion of controlled substances. In order to determine whether a registrant has provided effective controls against diversion, the West Virginia Board of Pharmacy shall evaluate the overall security system and needs of the applicant or registrant.

4.2.2 Physical security controls shall be commensurate with the schedules and quantity of controlled substances in the possession of the registrant in normal business operations. If a controlled substance is transferred to a different schedule or a noncontrolled substance is listed on any schedule, or the quantity of controlled substances in the possession of the registrant in normal business operations significantly increases, physical security controls shall be expanded and extended accordingly.

4.2.3 All registrants who receive or transfer substantial quantities of controlled substances in normal business operations shall employ security procedures to guard against in-transit losses.

4.3 Before distributing a controlled substance to any person who the registrant does not know to be registered to possess the controlled substance, the registrant shall make a good faith inquiry either with the Board or with the appropriate state controlled substances registration agency, if any, to determine that the person is registered to possess the controlled substance.

4.4. The registrant shall design and operate a

system to disclose to the registrant suspicious orders of controlled substances. The registrant shall inform the Office of the West Virginia Board of Pharmacy of suspicious orders when discovered by the registrant. Suspicious orders include orders of unusual size, orders deviating substantially from a normal pattern, and orders of unusual frequency.

4.5. The registrant shall notify the Office of the West Virginia Board of Pharmacy of any theft or significant loss of any controlled substances upon discovery of such theft or loss. The registrant shall also complete the form provided by the Secretary regarding such theft or loss.

##### 4.6.. Physical security controls

4.6.1. When a pharmacy is closed, controlled substances listed in Schedule II shall be stored in a securely locked narcotic cabinet made of 20 gauge metal or better or may be dispersed throughout the stock of noncontrolled substances in such a manner as to obstruct the theft or diversion of the controlled substance. Any other method of storage of controlled substances listed in Schedule II is not allowed unless specifically approved by the Board for that particular pharmacy. Only pharmacists practicing at the pharmacy and authorized by the pharmacist-in-charge may possess any keys or combinations to the narcotic cabinet. Controlled substances listed in Schedule III, IV, or V may be stored in the narcotic cabinet or may be dispersed throughout the stock of noncontrolled substances in such a manner as to obstruct the theft or diversion of the controlled substance. A secure automated distribution system, approved by the Board, may contain controlled substances within an institutional setting in lieu of a narcotic cabinet.

4.6.2. The registrant shall not employ as an agent or employee who has access to controlled substances, any person who has been convicted of a felony offense relating to controlled substances or who, at any time, had an application for registration denied, or has had his registration revoked.

**§15-2-5. Labeling And Packaging Requirements For Controlled Substances.**

5.1. Labeling and packaging requirements for controlled substances.

5.2. Scope of article.

Requirements governing the labeling and packaging of controlled substances pursuant to section three hundred five of the Uniform Controlled Substances Act (chapter sixty-a, West Virginia Code) are set forth generally by that section and specifically by the sections of this Part.

5.3. Definitions.

5.3.1. As used in this Part, the following terms shall have the meanings specified:

(a) The term "Commercial Container" means any bottle, jar, tube, ampule, or other receptacle in which a substance is held for distribution or dispensing to an ultimate user, and in addition, any box or package in which the receptacle is held for distribution or dispensing to an ultimate user. The term "Commercial Container" does not include any package liner, package insert or other material kept with or within a commercial container, nor any carton, crate, drug, or other package in which commercial containers are stored or are used for shipment of controlled substances.

(b) The term "Label" means any display of written, printed, or graphic matter placed upon the commercial container of any controlled substance by any manufacturer of such substance.

(c) The term "Labeling" means all labels and other written, printed or graphic matter (1) upon any controlled substance or any of its commercial containers or wrappers, or (2)

accompanying such controlled substance.

(d) The term "Manufacture" means the producing, preparation, propagation, compounding or processing of a drug or other substance or the packaging or repackaging of such substance, or the labeling or relabeling of the commercial container of such substance, but does not include the activities of a practitioner who, as an incident to his administration or dispensing such substance in the course of his professional practice, prepares, compounds, packages or labels such substance. The term "Manufacture" means a person who manufactures a drug or other substance, whether under a registration as a manufacturer or under authority of registration as a researcher or chemical analyst.

(e) Any term not defined in this section shall have the definition set forth in section one hundred one of the Uniform Controlled Substances Act, West Virginia Code Chapter sixty-a.

5.4. Symbol required; exceptions.

5.4.1. Each commercial container of a controlled substance except for a controlled substance accepted by the West Virginia Board of Pharmacy pursuant to section three hundred eight of this chapter, shall have printed on the label the symbol designating the schedule in which such controlled substance is listed. Each such commercial container, if it otherwise has no label, must bear a label complying with the requirement of this Part.

5.4.2 Each manufacturer shall print upon the labeling of each controlled substance distributed by him the symbol designating the schedule in which such controlled substance is listed.

5.4.3. The following symbols shall designate the schedule corresponding thereto:

Schedule I .....	CI or C-I.	Schedule II .....
	CII or C-II.	Schedule III .....
		CIII or C-III.
Schedule IV .....	CIV or C-IV.	Schedule V

..... CV or C-V.

The word "Schedule" need not be used. No distinction need be made between narcotic and nonnarcotic substances.

5.4.4 The symbol is not required on a carton or wrapper in which a commercial container is held if the symbol is easily legible through such carton or wrapper.

5.4.5. The symbol is not required on a commercial container too small or otherwise unable to accommodate a label, if the symbol is printed on the box or package from which the commercial container is removed upon dispensing to an ultimate user.

5.4.6. The symbol is not required on a commercial container containing, or on the labeling of, a controlled substance being utilized in clinical research involving blind and double blind studies.

5.4.7. The symbol is not required on a commercial container containing, or on the labeling, of a controlled substance intended for export from the United States.

#### 5.5. Location and size of symbol on label.

5.5.1 The symbol shall be prominently located on the right upper corner of the principal panel of the label of the commercial container and/or the panel of the commercial container normally displayed to dispensers of any controlled substance listed in Schedule I through V. The symbol must be at least two (2) times as large as the largest type otherwise printed on the label.

5.5.2. In lieu of locating the symbol in the corner of the label, as prescribed in Paragraph (a) of this section, the symbol may be overprinted on the label, in which case the symbol must be printed at least one half ( $\frac{1}{2}$ ) the height of the label and in a contrasting color providing clear visibility against the background color of the label.

5.5.3 In all cases, the symbol shall be clear and large enough to afford easy identification of the schedule of the controlled substance upon inspection without removal from the dispenser's shelf.

#### 5.6. Location and size of symbol on labeling.

5.6.1 The symbol shall be prominently located on all labeling other than labels covered by Section 5.5 of these rules. In all cases the symbol shall be clear and large enough to afford prompt identification of the controlled substance upon inspection of the labeling.

#### 5.7. Effective dates of labeling requirements.

5.7.1 All labels on commercial containers of, and all labeling of, a controlled substance which is listed in any schedule on June 15, 1971, and which is packaged after December 1, 1971, shall comply with the requirements of Section 5.4 of these rules.

5.7.2. All labels on commercial containers of, and all labeling of, a controlled substance which either is listed in any schedule on June 15, 1971, and thereafter transferred to another schedule or is added to any schedule after June 15, 1971, and which is packaged more than one hundred eighty (180) days following the date on which the transfer or addition becomes effective, shall comply with the requirements of Section 5.4 of these rules.

5.7.3. The West Virginia Board of Pharmacy may, in the case of any controlled substance, require compliance with the requirements of Section 5.4 of these rules, within a period of time shorter than required by this section if it finds that public health or safety necessitate an earlier effective date.

5.7.4. Until compliance is required under this section, the label on commercial container containing, and the labeling of, any controlled substance shall comply with any requirements under Federal Law as to labels of such containers and as to labeling of such

substances existing prior to the effective date prescribed in this section.

#### 5.8. Sealing of controlled substances.

5.8.1. On each bottle, multiple dose, vial or other commercial container of any controlled substance listed in Schedules I and/or II, and of any narcotic controlled substance listed in Schedule III or IV, there shall be securely affixed to the stopper, cap, lid, covering or wrapper or such container a seal to disclose upon inspection any tampering or opening of the container.

5.8.2. Any seal accepted for use under Federal Law prior to May 1, 1971, shall be deemed acceptable for use under this section.

### §15-2-6. Records And Reports Of Registrants.

6.1. All records required to be kept must be readily retrievable. "Readily Retrievable" means that certain records are kept by automatic data processing systems or other electronic or mechanized record keeping systems in such a manner that they can be separated out from all other records in a reasonable time and/or records are kept on which certain items are asterisked, red-lined or in some other manner visually identifiable apart from other items appearing on the records.

#### 6.2 Maintenance of records and inventories.

6.2.1. Every inventory and other record required to be kept under the Act shall be kept by the registrant and be available, for at least five (5) years from the date of such inventory or record, for inspecting and copying by authorized employees of the West Virginia Board of Pharmacy.

6.2.2. Each registered manufacturer, distributor, importer, and exporter shall maintain

inventories and records of controlled substances as follows:

(a) Inventories and records of controlled substances listed in Schedules I and II shall be maintained separately from all of the records of the registrant; and

(b) Inventories and records of controlled substances listed in Schedules III, IV and V shall be maintained either separately from all other records of the registrant or in such form that the information required is readily retrievable from the ordinary business records of the registrant.

6.2.3. Each registered individual practitioner required to keep records and institutional practitioner shall maintain inventories and records of controlled substances in the manner prescribed in Paragraph (b) of this section.

6.2.4 Each registered pharmacy shall maintain the inventories and records of controlled substances as follows:

(a) Inventories and records of all controlled substances listed in Schedules I and II shall be maintained separately from all other records of the pharmacy, and prescriptions for such substances shall be maintained in a separate prescription file; and

(b) Inventories and records of controlled substances listed in Schedules III, IV and V shall be maintained either separately from all other records of the pharmacy or in such form that the information required is readily retrievable from ordinary business records of the pharmacy, and prescriptions for such substances shall be maintained either in separate prescription file for controlled substances listed in Schedules III, IV and V only, or in such form that they are readily retrievable from the other prescription records of the pharmacy. Prescriptions will be deemed readily retrievable if, at the time they are initially filed, the face of the prescription is stamped in red ink in the lower right corner with the letter "C" no less than 1 inch high and filed either in the

prescription file for controlled substances listed in Schedules I and II or in the usual consecutively numbered prescription file for noncontrolled substances. However, if a pharmacy employs an automated data processing system or other electronic record-keeping system for prescriptions which permits identification by prescription number and retrieval of original documents by prescriber's name, patient's name, drug dispensed, and date filled, then the requirement to mark the hard copy prescription with a red "C" is waived.

### 6.3. General requirements for inventories.

6.3.1. Each inventory shall contain a complete and accurate record of all controlled substances on hand on the date the inventory is taken. Controlled substances shall be deemed to be "On Hand" if they are in the possession of or under the control of the registrant, including substances returned by a customer, substances ordered by a customer but not yet invoiced, substances stored in a warehouse on behalf of the registrant and substances in the possession of employees of the registrant and intended for distribution as complimentary samples.

6.3.2. A separate inventory shall be made by a registrant for each registered location. In the event controlled substances in the possession or under the control of the registrant at a location for which he is not registered, the substances shall be included in the inventory of the registered location to which they are subject to control or to which the person possessing the substance is responsible. Each inventory for a registered location shall be kept at the registered location.

6.3.3. A separate inventory shall be made by a registrant for each independent activity for which he is registered, except as provided in Section 7.14 of these rules.

6.3.4. A registrant may take an inventory either as of the opening of business or as of the close of business on the inventory date. The registrant shall indicate on the inventory

records whether the inventory is taken as of the opening or as of the close of business and the date the inventory is taken.

6.3.5. An inventory must be maintained in a written, typewritten or printed form. An inventory taken by use of an oral recording device must be promptly transcribed.

### 6.4. Initial inventory date.

6.4.1. Every person required to keep records shall take an inventory of all stocks of controlled substances on hand on the date he or she first engages in the manufacture, distribution or dispensing of controlled substances, in accordance with Sections 6.4 through 6.7 of these rules, as applicable. In the event a person commences business with no controlled substances on hand, he or she shall record this fact as the initial inventory.

### 6.5. Biennial inventory date.

6.5.1. After the initial inventory is taken, the registrant shall take a new inventory of all stocks of controlled substances on hand at least every two years. The biennial inventory may be taken on any date which is within two years of the previous biennial inventory date.

### 6.6. Inventory date for new controlled substances.

6.6.1 On the effective date of a rule or statutory change by the West Virginia Board of Pharmacy or the DEA adding a substance to any schedule of controlled substances, which substance was, immediately prior to that date, not listed on any such schedule, every registrant required to keep records who is manufacturing, distributing or dispensing that substance shall take an inventory of all stocks of the substance on hand. Thereafter such substance shall be included in each inventory made by the registrant pursuant to Section 6.4 of these rules.

### 6.7. Inventories of manufacturers.

6.7.1. Each registered manufacturer shall include the following information in his inventory:

(a) For each controlled substance in bulk form to be used in (or capable of use in) the manufacture of the same or other controlled or noncontrolled substances in finished form:

(1) The name of the substance; and

(2) The total quantity of the substance to the nearest metric unit weight consistent with unit size (except that for inventories made in 1971, avoirdupois weights may be utilized where metric weights are not readily available).

(b) For each controlled substance in the process of manufacture on the inventory date:

(1) The name of the substance;

(2) The quantity of the substance in each batch and/or stage of manufacture, identified by the batch number or other appropriate identifying number; and

(3) The physical form which the substance is to take upon completion of the manufacturing process (e.g. granulations, tablets, capsules or solutions), identified by the batch number or other appropriate identifying number, and if possible the finished form of the substance (e.g. ten (10) milligram tablet or ten (10) milligram concentration per fluid ounce of milliliter) and the number or volume thereof.

(c) For each controlled substance in finished form:

(1) The name of the substance;

(2) Each finished form of the substance (e.g., ten (10) milligram tablet or ten (10) milligram concentration per fluid ounce or milliliter);

(3) The number of units or volume of each

finished form in each commercial container (e.g., one hundred (100) tablet bottles or six (6) three (3) milliliter vials); and

(4) The total quantity of the substance in all forms to the nearest metric unit weight.

(d) For each controlled substance not included in Paragraphs (a), (b) or (c) of this section (e.g., damaged, defective or impure substances awaiting disposal, substances held for quality control purposes, or substances maintained for extemporaneous compoundings):

(1) The name of the substance;

(2) The total quantity of the substance to the nearest metric unit weight or the total number of units of finished form; and

(3) The reason for the substance being maintained by the registrant and whether such substance is capable of use in the manufacture of any controlled substance in finished form.

## 6.8. Inventories of distributors.

6.8.1. Each registered distributor shall include in his inventory the same information required of manufacturers pursuant to Section 6.8.1 (c) and (d) of these rules.

## 6.9. Inventories of dispensers and researchers.

6.9.1. Each person registered to dispense or conduct research with controlled substances and required to keep records pursuant to Section 6.4 of these rules, shall include in his inventory the same information required of manufacturers pursuant to Section 6.8.1. (c) and (d) of these rules. In determining the number of units of each finished form of a controlled substance in a commercial container which has been opened, the dispenser shall do as follows:

(a) If the substance is listed in Schedule I or II, he shall make an exact count or measure of the content; and

(b) If the substance is listed in Schedule III, IV or V, he shall make an estimated count or measure of the contents, unless the container holds more than one thousand (1,000) tablets or capsules in which case he must make an exact count of the contents.

#### 6.10.. Inventories of importers and exporters.

6.10.1. Each registered importer or exporter shall include in his inventory the same information required of manufacturers pursuant to Section 6.7.1 (a), (c) and (d) of these rules. Each registered importer and exporter who is also registered as a manufacturer or as a distributor shall include in his inventory as an importer or exporter only those stocks of controlled substances that are actually separated from his stocks as a manufacturer or as a distributor (e.g., in transit or in storage for shipment).

#### 6.11. Inventories for chemical analysts.

6.11.1 Each analytical laboratory registered to conduct chemical analysis with controlled substances shall include in its inventory the same information required of manufacturers pursuant to Section 6.8.1 (a), (c) and (d) of these rules, as to substances which have been manufactured, imported or received by the laboratory conducting the inventory, if less than one (1) kilogram of any controlled substance (other than a hallucinogenic controlled substance listed in Schedule I), or less than twenty (20) grams of a hallucinogenic substance listed in Schedule I, (other than lysergic acid diethylamide), or less than five tenths (0.5) gram of lysergic acid diethylamide, is on hand at the time of inventory, that substance need not be included in the inventory. Laboratories of the West Virginia Board of Pharmacy may possess up to one hundred fifty (150) grams of any hallucinogenic substance in Schedule I without regard to a need for an inventory of those substances.

6.12. General requirements for continuing records.

6.12.1 On and after June 15, 1971, every registrant required to keep records pursuant to Section 6.3 of these rules, shall maintain on a current basis a complete and accurate record of each such substance manufactured, imported, received, sold, delivered, exported or otherwise disposed of by him.

6.12.2. Separate records shall be maintained by a registrant for each registered location. In the event controlled substances are in the possession or under the control of a registrant at a location for which he is not registered, the substances shall be included in the records of the registered location to which they are subject to control or to which the person possessing the substance is responsible.

6.12.3. Separate records shall be maintained by a registrant for each independent activity for which he is registered.

6.12.4. In recording dates of receipt, importation, distribution, exportation or other transfer, the date on which the controlled substances are actually received, imported, distributed, exported or otherwise transferred shall be used as the date of receipt or distribution of any documents of transfer (e.g., invoices or packing slips).

#### 6.13. Records of manufacturers.

6.13.1. Each registered manufacturer shall maintain records with the following information:

(a) For each controlled substance in bulk form to be used, or capable of use in, or being used in, the manufacture of the same or other controlled or noncontrolled substances in finished form:

- (1) The name of the substance;
- (2) The quantity manufactured in bulk form by the registrant, including the date, quantity and batch or other identifying number of each batch manufactured;
- (3) The quantity received from other

persons, including the date and quantity of each receipt and the name, address and registration number of the other person from whom the substance was received;

(4) The quantity imported directly by the registrant (under a registration as an importer) for use in manufacture by him, including the date, quantity and import permit or declaration number for each importation;

(5) The quantity used to manufacture the same substance in finished form, including:

A) The date and batch or other identifying number of each manufacture;

B) The quantity used in the manufacture;

C) The finished form (e.g., ten (10) milligram tablets or ten (10) milligram concentration per fluid ounce or milliliter);

D) The number of units of finished form manufactured;

E) The quantity used in quality control;

F) The quantity lost during manufacturing and the causes therefore, if known;

G) The total quantity of the substance contained in the finished form;

H) The theoretical and actual yields; and

I) Such other information as is necessary to account for all controlled substances used in the manufacturing process.

(6) The quantity used to manufacture other controlled and noncontrolled substances, including the name of each substance manufactured and the information required in Subparagraph (5) of this paragraph;

(7) The quantity distributed in bulk

form to other persons, including the date and quantity of each distribution and the name, address and registration number of each person to whom a distribution was made;

(8) The quantity exported directly by the registrant (under a registration as an exporter), including the date, quantity and export permit or declaration number of each exportation;

(9) The quantity distributed or disposed of in any other manner by the registrant (e.g., by distribution of complimentary samples or by destruction), including the date and manner of distribution or disposal, the name, address and registration number of the person to whom distributed and the quantity distributed or disposed.

(b) For each controlled substance in finished form.

(1) The name of the substance;

(2) Each finished form (e.g., ten (10) milligram tablet or ten (10) milligram concentration per fluid ounce or milliliter) and the number of units or volume of finished form in each commercial container (e.g., one hundred (100) tablet bottle or three (3) milliliter vial);

(3) The number of containers of each such commercial finished form manufactured from bulk form by the registrant, including the information required pursuant to Subparagraph (5) of Paragraph (a) of this section;

(4) The number of units of finished forms and/or commercial containers received from other persons, including the date of and number of units and/or commercial containers in each receipt and the name, address and registration number of the person from whom the units were received;

(5) The number of units of finished forms and/or commercial containers imported directly by the registrant (under a registration as an importer), including the date of and the number of units and for commercial containers in each

importation;

(6) The number of units and/or commercial containers manufactured by the registrant from units in finished form received from others or imported, including:

A) The date and batch or other identifying number of each manufacture;

B) The operation performed (e.g., repackaging or relabeling);

C) The number of units of finished form used in the manufacture, the number manufactured and the number lost during manufacture, with the causes therefore, if known; and

D) Such other information as is necessary to account for all controlled substances used in the manufacturing process.

(7) The number of commercial containers distributed to other persons, including the date of and number of containers in each distribution, and the name, address and registration number of the person to whom the containers were distributed;

(8) The number of commercial containers exported directly by the registrant (under a registration as an exporter), including the date, number of containers and export permit or declaration number for each exportation; and

(9) The number of units of finished forms and/or commercial containers distributed or disposed of in any other manner by the registrant (e.g., by distribution of complimentary samples or by destruction), including the date and manner of distribution or disposal, the name, address and registration number of the person to whom distributed and the quantity in finished form distributed or disposed.

#### 6.14. Records for distributors.

6.14.1 Each registered distributor shall maintain records with the following information

for each controlled substance:

(a) The name of the substance;

(b) Each finished form (e.g., ten (10) milligram tablet or ten (10) milligram concentration per fluid ounce or milliliter) and the number of units or volume of finished form in each commercial container (e.g., one hundred (100) tablet bottle or three (3) milliliter vial);

(c) The number of commercial containers of each such finished form received from other persons, including the date of and number of containers in each receipt and the name, address and registration number of the person from whom the containers were received;

(d) The number of commercial containers of each such finished form imported directly by the registrant (under a registration as an importer), including the date of and the number of containers in each importation;

(e) The number of commercial containers of each such finished form distributed to other persons, including the date of and number of containers in each distribution and the name, address and registration number of the person to whom the containers were distributed;

(f) The number of commercial containers of such finished form exported directly by the registrant (under a registration as an exporter), including the date of and the number of containers in each exportation; and

(g) The number of units or volume of finished forms and/or commercial containers distributed or disposed of in any other manner by the registrant (e.g., by distribution as complimentary samples), including the date and manner of distribution or disposal, the name, address and registration number of the person to whom distributed and the quantity of the substance in finished form distributed or disposed.

#### 6.15. Records for dispensers and researchers.

6.15.1. Each person registered to dispense or conduct research with controlled substances and required to keep records pursuant to Section 6.3 of these rules, shall maintain records with the following information for each controlled substance:

- (a) The name of the substance;
- (b) Each finished form (e.g., ten (10) milligram tablet or ten (10) milligram concentration per fluid ounce or milliliter) and the number of units or volume of finished form in each commercial container (e.g., one hundred (100) bottle or three (3) milliliter vial);
- (c) The number of commercial containers of each such finished form received from other persons, including the date of and number of containers in each receipt and the name, address and registration number of the person from whom the containers were received;
- (d) The number of units or volume of such finished form dispensed, including the name and address of the person to whom it was dispensed, the date of dispensing, the number of units or volume dispensed, and the written or typewritten name or initials of the individual who dispensed or administered the substance on behalf of the dispenser; and
- (e) The number of units or volume of such finished forms and/or commercial containers disposed of in any other manner by the registrant, including the date and manner of disposal and the quantity of the substance in finished form disposed.

#### 6.16. Records for importers.

6.16.1 Each registered importer shall maintain records with the following information for each controlled substance:

- (a) The name of the substance;
- (b) The quantity (or number of units or volume in finished form) imported, including the

date, quantity (or number of units or volume) and import permit or declaration number for each importation;

- (c) The quantity (or number of units or volume in finished form) distributed to other persons, including the date and quantity (or number of units or volume) of each distribution and the name, address and registration number of each person to whom a distribution was made;

- (d) The quantity disposed of in any other manner by the registrant (except quantities used in manufacturing by an importer under a registration as a manufacturer, which quantities are to be recorded pursuant to Section 7.18 (a) (4) or (b) (5) of these rules, including the date and manner of disposal and the quantity disposed.

#### 6.17. Records for chemical analysis.

6.17.1. Each person registered to conduct chemical analysis with controlled substances shall maintain records with the following information (to the extent known and reasonably ascertainable by him) for each controlled substance:

- (a) The name of the substance;
- (b) The form or forms in which the substance is received, imported or manufactured by the registrant (e.g., powder, granulation, tablet, capsule or solution) and the concentration of the substance in such form (e.g., C.P., U.S.P., N.F., ten (10) milligram tablet or ten (10) milligram concentration per milliliter);
- (c) The total number of the forms received, imported or manufactured (e.g., one hundred (100) tablets, thirty (30) one (1) milliliter vial, or ten (10) grams of powder), including the date and quantity of each receipt, importation or manufacture and the name, address and registration number, if any, of the person from whom the substance was received;

- (d) The quantity distributed, exported or destroyed in any manner by the registrant (except quantities used in chemical analysis or

other laboratory work), including the date, the manner of distribution, exportation or destruction and the name, address and registration number, if any, of each person to whom the substance was distributed or exported.

6.17.2. Order forms, import and export permits, import invoices and export declarations relating to controlled substances shall be maintained separately from all other records of the registrant.

6.17.3. Records of controlled substances used in chemical analysis are not required.

6.17.4. Records relating to known or suspected controlled substances received as samples for analysis are not required under this section.

#### §15-2-7. Prescriptions.

7.1. Rules governing the issuance, filling and filing of prescriptions are set forth generally in West Virginia Code §60A-3-308.

##### 7.2.. Definitions.

7.2.1. As used in this Part, the following terms shall have the meanings specified:

(a) The term "Act" means the Uniform Controlled Substances Act (chapter sixty-a, West Virginia Code).

(b) The term "Individual Practitioner" means a physician, dentist, veterinarian or other individual licensed, registered or otherwise permitted, by the state jurisdiction in which he practices, to dispense a controlled substance in the course of professional practice, but does not include a pharmacist, a pharmacy or an institutional practitioner.

(c) The term "Institutional Practitioner" means a hospital or other person (other than an

individual) licensed, registered or otherwise permitted, by the state jurisdiction in which it practices, to dispense a controlled substance in the course of professional practice, but does not include a pharmacy.

(d) The term "Pharmacist" means any pharmacist licensed by a state to dispense controlled substances and shall include a pharmacist intern authorized by a state to dispense controlled substances under the supervision of a pharmacist licensed by such state.

(e) The term "Prescription" means an order for medication which is dispensed to or for an ultimate user but does not include the immediate administration to the ultimate user.

(f) Any term not defined in this section shall have the definition set forth in section one hundred one, article one of the Act (chapter sixty-a, West Virginia Code).

##### 7.3.. Persons entitled to issue prescriptions.

7.3.1. A prescription for a controlled substance may be issued only by an individual practitioner who is authorized to prescribe controlled substances in the jurisdiction in which he or she practices.

7.3.2. A prescription issued by an individual practitioner except for Schedule II controlled substance, may be communicated to a pharmacist by an employee or agent of the individual practitioner.

##### 7.4.. Purpose of issue of prescription.

7.4.1. A prescription for a controlled substance to be effective must be issued for a legitimate purpose by an individual practitioner acting in the usual course of his professional practice. The responsibility for the proper prescribing and dispensing of controlled substances is upon the prescribing practitioner, but a corresponding responsibility rests with the pharmacist who fills the prescription. An order purporting to be a prescription issued not in the

usual course of professional treatment or in legitimate and authorized research is not a prescription within the meaning and intent of section three hundred eight, article three of the Act and the person knowingly filling such a purported prescription, as well as the person issuing it, shall be subject to the penalties provided for violations of the provisions of law relating to controlled substances.

7.4.2. A prescription may not be issued in order for an individual practitioner to obtain controlled substances for supplying the individual practitioner for the purpose of general dispensing to patients; i.e. office use.

7.4.3. A prescription may not be issued for the dispensing of narcotic drugs listed in any schedule to a narcotic drug dependent person for the purpose of continuing his dependence upon such drugs, except in the course of conducting an authorized clinical investigation in the development of a narcotic addict rehabilitation program.

#### 7.5. Manner of issuance of prescriptions.

7.5.1. All prescriptions for controlled substances shall be dated as of, and signed on, the day when issued and shall bear the full name and address of the patient, and name, address and registration number of the practitioner. A practitioner may sign a prescription in the same manner as he would sign a check or legal document (e.g. J.H. Smith or John H. Smith). Where an oral order is not permitted, prescriptions shall be written with ink or indelible pencil or a typewriter and shall be manually signed by the practitioner. The prescriptions may be prepared by a secretary or agent for the signature of a practitioner, but the prescribing practitioner is responsible in case the prescription does not conform in all essential respects to the law and regulations. A corresponding liability rests upon the pharmacist who fills a prescription not prepared in the form prescribed in these regulations.

#### 7.6. Form of controlled substance

prescription.

7.6.1. Each controlled substance prescription shall be written on a separate blank and no non-controlled substance can be ordered on a blank with a controlled substance. No more than one controlled substance can be written per prescription blank. A controlled substance prescription issued by a practitioner located outside the state of West Virginia that does not comply with this section may be accepted by the pharmacist if it is issued pursuant to the laws in the state in which the practitioner resides.

7.6.2. If a pharmacist receives a prescription with more than one controlled substance on the blank or a non-controlled substance on a blank with a controlled substance, then the pharmacist shall refuse to fill the prescription; Provided, that if the pharmacist in his or her professional judgement determines the immediate necessity for the patient to receive their medication, then the prescriptions may be dispensed and the pharmacist must document in a log the prescription numbers and drugs dispensed. This log must be kept in the pharmacy and be available for inspection. The pharmacist shall contact the prescriber as soon as possible to inform them that the prescription was not written according to this rule. If the pharmacist continues to receive prescriptions from the same practitioner that do not comply with this rule, then the pharmacist shall inform the West Virginia Board of Pharmacy.

7.6.3. Every controlled substance prescription shall have the name of the practitioner stamped, typed, or printed legibly on the face of the prescription, as well as the signature of the practitioner. Institutional prescription blanks shall include the DEA number of the hospital or other institution and the special internal code number (suffix) assigned to him or her by the hospital or other institution, in lieu of the individual DEA number of the practitioner. If multiple practitioners are listed on a prescription blank, then the specific name of the prescriber must be clearly distinguished upon the prescription. If a pharmacist receives a

prescription that does not comply with this section, then the pharmacist shall refuse to fill the prescription; Provided, that if the pharmacist in his or her professional judgement determines the immediate necessity for the patient to receive their medication, then the prescriptions may be dispensed and the pharmacist must document in a log the date, patient name, practitioner name, prescription numbers, and drugs dispensed. This log must be kept in the pharmacy and be available for inspection. The pharmacist shall contact the prescriber as soon as possible to inform them that the prescription was not written according this rule. If the pharmacist continues to receive prescriptions from the same practitioner that do not comply with this rule, then the pharmacist shall inform the West Virginia Board of Pharmacy.

#### 7.7. Persons entitled to fill prescriptions.

7.7.1. A prescription for controlled substances may only be filled by a pharmacist acting in the usual course of his professional practice and either registered individually or employed in a registered pharmacy or registered institutional practitioner (e.g., a hospital, nursing home, home for the aged, clinic, orphanage, governmental agency or institution or other place of similar character which dispenses controlled substances).

7.8. Dispensing of narcotic drugs for maintenance purposes.

7.8.1 The administering or dispensing directly (but not prescribing) of narcotic drugs listed in any schedule to a narcotic drug dependent person for "detoxification treatment" or "maintenance treatment" shall be deemed to be within the meaning of the term "in the course of his professional practice or research.": Provided, that the practitioner is separately registered with the U.S. Attorney General as required by section 303(g) of the federal Controlled Substances Act (21 U.S.C. 823(g)) and then thereafter complies with the regulatory standards imposed relative to treatment qualification, security, records and unsupervised use of drugs pursuant to the Act.

7.8.2 Nothing in this section shall prohibit a physician who is not specifically registered to conduct a narcotic treatment program from administering (but not prescribing) narcotic drugs to a person for the purpose of relieving acute withdrawal symptoms when necessary while arrangements are being made for referral for treatment. Not more than one day's medication may be administered to the person or for the person's use at one time. Such emergency treatment may be carried out for not more than three days and may not be renewed or extended.

7.8.3 This section is not intended to impose any limitations on a physician or authorized hospital staff to administer or dispense narcotic drugs in a hospital to maintain or detoxify a person as an incidental adjunct to medical or surgical treatment of conditions other than addiction, or to administer or dispense narcotic drugs to persons with intractable pain in which no relief or cure is possible or none has been found after reasonable efforts.

7.9. Controlled substances listed in Schedule II.

#### 7.9.1.. Requirement of prescription.

(a) A pharmacist may dispense a controlled substance listed in Schedule II, which is a prescription drug as determined under the Federal Food, Drug and Cosmetic Act, only pursuant to a written prescription signed by the prescribing individual practitioner, except as provided in Rule 7.9.2 of this section. A prescription for a Schedule II controlled substance may be transmitted by the practitioner or the practitioner's agent to a pharmacy via facsimile equipment or electronic transmission, provided that the original written, signed prescription is presented to the pharmacist for review prior to the actual dispensing of the controlled substance, except as provided for CSR §15-1-21.1.3. A prescription for a Schedule II controlled substance is valid for thirty (30) days from the date issued; Provided that a pharmacist may fill the prescription after thirty (30) days if the prescriber

confirms to the pharmacist that he or she still wants the prescription filled and the pharmacist documents upon the prescription that such confirmation was obtained.

(b) An individual practitioner may administer or dispense a controlled substance listed in Schedule II in the course of his professional practice without a prescription, subject to Section 9.8 of these rules.

(c) An institutional practitioner may administer or dispense directly (but not prescribe) a controlled substance listed in Schedule II only pursuant to a written prescription signed by the prescribing individual practitioner or to an order for medication made by an individual practitioner which is dispensed for immediate administration to the ultimate user.

7.9.2. In the case of an emergency situation, a prescription for a Schedule II controlled substance may be communicated by the practitioner orally or by way of electronic transmission, provided that if the prescribing practitioner is not known to the pharmacist, he or she shall make a reasonable effort to determine that the oral authorization came from a registered practitioner, which may include a callback to the practitioner using the practitioner's phone number as listed in the telephone directory and other good faith efforts to insure his identity; and:

a. the quantity prescribed and dispensed is limited to the amount adequate to treat the patient during the emergency period (dispensing beyond the emergency period must be pursuant to a written prescription signed by the prescribing practitioner);

b. the orally communicated prescription is immediately reduced to writing by the pharmacist, or, if necessary, the prescription communicated by way of electronic transmission is immediately reduced to a hard copy;

c. within seven (7) days after authorizing an emergency oral prescription, the practitioner has a written prescription for the

emergency quantity prescribed delivered to the dispensing pharmacist. The prescription shall have written on its face "Authorization for Emergency Dispensing" and the date of the orally or electronically transmitted prescription. The written prescription may be delivered to the pharmacist in person or by mail, but if delivered by mail, it must be postmarked within the seven (7) day period. Upon receipt, the dispensing pharmacist shall attach this written prescription to the emergency oral prescription which had earlier been reduced to writing or to the hard copy of the electronically transmitted prescription. The pharmacist shall notify the nearest office of the U.S. Drug Enforcement Administration and the West Virginia Board of Pharmacy if the prescribing practitioner fails to deliver a written prescription.

#### 7.10.. Refilling Schedule II prescriptions.

7.10.1 The refilling of a prescription for a controlled substance listed in Schedule II is prohibited.

#### 7.11.. Partial filling of Schedule II prescriptions.

7.11.1. The partial filling of a prescription for a controlled substance listed in Schedule II is permissible, if the pharmacist is unable to supply the full quantity called for in a written or emergency oral prescription and he makes a notation of the quantity supplied on the face of the written prescription (or written record of the emergency oral prescription). The remaining portion of the prescription may be filled within seventy-two (72) hours of the first partial filling, however, if the remaining portion is not or cannot be filled within the seventy-two (72) hour period, the pharmacist shall so notify prescribing individual practitioner. No further quantity may be supplied beyond seventy-two (72) hours without a new prescription.

#### 7.12.. Labeling of Schedule II prescriptions.

7.12.1. The pharmacist filling a written or emergency oral prescription for a controlled

substance listed in Schedule II shall affix to the package a label showing date of filling, the pharmacy name and address, the serial number of the prescription, the name of the patient, the name of the prescribing practitioner and directions for use and cautionary statements, if any, contained in such prescription or required by law.

#### 7.13.. Filing of prescriptions.

7.13.1. All written prescriptions and written records of emergency oral prescriptions shall be kept in accordance with requirements of this chapter.

7.14.. Controlled substances listed in Schedules III , IV, and V.

#### 7.14.1.. Requirement of prescription.

(a) A pharmacist may dispense a controlled substance listed in Schedule III, IV, or V, which is a prescription drug as determined under the Federal Food, Drug and Cosmetic Act, only pursuant to either a written prescription signed by a prescribing individual practitioner and/or an oral prescription made by a prescribing individual practitioner and promptly reduced to writing by the pharmacist containing all information required in these rules, except for the signature of the prescribing individual practitioner.

(b) An individual practitioner may administer or dispense a controlled substance listed in Schedule III, IV, or V in the course of his professional practice without a prescription, subject to the provisions of Section 7.8 of these rules.

(c) An institutional practitioner, as described in Section 7.2 of these rules, may administer or dispense directly (but not prescribe) a controlled substance listed in Schedules III, IV, or V pursuant to a written prescription signed by a prescribing individual practitioner, or pursuant to an oral prescription made by a prescribing individual practitioner and promptly reduced to writing by the pharmacist (containing all

information required in Section 7.5 of these rules, except for the signature of the prescribing individual practitioner), or pursuant to an order for medication made by an individual practitioner which is dispensed for immediate administration to the ultimate user, subject to Section 7.8 of these rules.

7.15. Refilling of Schedule III or IV prescriptions.

7.15.1. No prescription for a controlled substance listed in Schedule III or IV shall be filled or refilled more than six (6) months after the date on which such prescription was issued and no such prescription authorized to be refilled may be refilled more than five (5) times. Each refilling of a prescription shall be entered on the back of the prescription (or on another uniformly maintained appropriate record, such as medication records, which indicate prescription refills), initialed, and dated by the pharmacist as of the date of dispensing, and shall state the amount dispensed. If the pharmacist merely initials and dates the back of the prescription, he shall be deemed to have dispensed a refill for the full face amount of the prescription. Additional quantities of controlled substances listed in Schedule III or IV may only be authorized by a prescribing practitioner through issuance of a new prescription as provided in Section 7.14 of these rules, which shall be a new and separate prescription. The number of partial fills may be more than five times as long as the total quantity prescribed is not exceeded.

7.16. Partial Filling of Schedule III, IV, or V prescriptions.

7.16.1. The partial filling of a prescription for a controlled substance listed in Schedule III, IV, or V is permissible provided that:

- (a) Each partial filling is recorded in the same manner as a refilling,
- (b) The total quantity dispensed in all partial fillings does not exceed the total quantity prescribed, and
- (c) No dispensing occurs after 6 months

after the date on which the prescription was issued.

7.17.. Labeling of Schedule III, IV, or V prescriptions.

7.17.1. The pharmacist filling a prescription for a controlled substance listed in Schedule III, IV, or V shall affix to the package a label showing the pharmacy name and address, the serial number and date of initial filling the name of the patient, the name of the practitioner issuing the prescription, and directions for use and cautionary statements, if any, contained in such prescription as required by law.

7.18.. Filing of Schedule III, IV, or V prescriptions.

7.18.1. All prescriptions for controlled substances listed in Schedules III, IV, or V shall be kept in accordance with Section 6.3.4 of this rule.

7.19.. Dispensing without prescription.

7.19.1. A controlled substance listed in Schedules II, III, IV, or V which is not a prescription drug as determined by the Federal Food, Drug, and Cosmetic Act, may be dispensed by a pharmacist without a prescription to a purchaser at retail: **Provided**, That

(a) Such dispensing is made only by a pharmacist and not by a nonpharmacist employee even if under the direct supervision of a pharmacist (although after the pharmacist has fulfilled his professional and legal responsibilities set forth in this section, the actual cash, credit transaction or delivery, may be completed by a nonpharmacist);

(b) Not more than 240 cc. (8 ounces) of any such controlled substance containing opium, nor more than 120 cc. (4 ounces) of any other such controlled substance nor more than 48 dosage units of any such controlled substance containing opium, nor more than 24 dosage units

of any other such controlled substance may be dispensed at retail to the same purchaser in any given forty eight (48) hour period;

(c) The purchaser is at least eighteen (18) years of age;

(d) The pharmacist requires every purchaser of a controlled substance under this section not known to him to furnish suitable identification (including proof of age where appropriate);

(e) A bound record book for distributions of controlled substances under this section (other than by prescription) is maintained by the pharmacist, which book shall contain the name and address of the purchaser, the name and quantity of controlled substance purchased, the date of each purchase and the name or initials of the pharmacist who dispensed the substance to the purchaser (the book shall be maintained in accordance with the record keeping requirement of Section 7.5 of these rules); and

(f) A prescription is not required for distribution or dispensing of the substance pursuant to any other federal, state or local law.

**§15-2-8. Miscellaneous.**

8.1. Distribution upon discontinuance or transfer of business.

8.1.1. Any registrant desiring to discontinue or transfer business activities altogether or with respect to controlled substances shall notify the Board of Pharmacy immediately and on such notification shall submit a complete and detailed closing inventory of all controlled substances in the registrant's possession.

8.2. Disposal of controlled substances.

8.2.1. Compliance with federal law and regulations shall be deemed in compliance with this section. A registrant shall document the

destruction or disposal of all controlled substances on the appropriate form approved by the Board. The disposal of excessive amounts of residual and wasted controlled substances accrued by extemporaneous compounding in an institutional setting may be completed by two (2) registered or licensed health care professionals with a record of the destruction indicating the two witnesses with their signatures.

### 8.3. Reporting theft of drugs.

8.3.1. In the event of any controlled substances being stolen, it shall be the duty of the registrant to immediately submit a report of drug theft with the Board of Pharmacy.

### 8.4 Ordering of Controlled Substances.

8.4.1. An order form (DEA Form 222) is required for each transfer of a Schedule II controlled substance to another registrant without a prescription.

8.4.2. A pharmacist shall verify the receipt within the pharmacy of all controlled substances listed in Schedule II-V by reviewing and countersigning the invoices or packing documents.