

## DEPARTMENT OF REGULATORY AGENCIES

### State Board of Pharmacy

#### 3 CCR 719-1

### STATE BOARD OF PHARMACY RULES

*[Editor's Notes follow the text of the rules at the end of this CCR Document.]*

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#### 1.00.00 RULES OF PROFESSIONAL CONDUCT.

- 1.00.11 A pharmacist shall at all times conduct his/her profession in conformity with all federal and state drug laws, rules and regulations; and shall uphold the legal standards of the current official compendia.
- 1.00.12 A pharmacist shall not be a party or accessory to nor engage in any fraudulent or deceitful practice or transaction in pharmacy, nor knowingly participate in any practice which detrimentally affects the patient, nor discredit his/her profession.
- 1.00.13 A pharmacist shall not enter into any agreement or arrangement with anyone for the compounding of secret formula or coded orders, except for investigational drugs.
- 1.00.15 A pharmacist shall not, directly or indirectly, be employed as a pharmacist to dispense drugs by a person authorized to prescribe drugs. For the purpose of this rule, the term person shall include any person or persons, partnership or business entity in which the person or persons authorized to prescribe drugs has an ownership interest individually or jointly greater than 10 percent.
- 1.00.16 Confidentiality.
- a. A pharmacist shall not exhibit, discuss, or reveal the contents of any order or prescription, the therapeutic effect thereof, the nature, extent, or degree of illness suffered by any patient or any medical information furnished by the practitioner with any person other than the patient or his authorized representative, the practitioner or another licensed practitioner then caring for the patient, another pharmacist or intern serving the patient, or a person duly authorized by law or by the patient to receive such information.
  - b. A pharmacist may disclose patient information to pharmacy technicians, authorized law enforcement personnel, another pharmacist acquiring and maintaining the records, third party entities responsible for payment and any other parties allowed by federal privacy regulations.
  - c. The pharmacist shall exercise his professional judgment in the release of patient information to a patient or his authorized agent.
- 1.00.17 A pharmacist or prescription drug outlet shall not pay or offer to pay or imply that payment might be made of any sum of money or other thing of value to a practitioner, health care facility, nursing care or assisted living facility, or any other health care provider or entity as consideration for any referral to, or promotion of, a prescription drug outlet.
- 1.00.18 Patient Counseling. When the patient seeks advice, or when, in the pharmacist's professional judgment, the best interest of the patient will be served, the pharmacist shall offer to advise the patient regarding the prescription.

- 1.00.21 Violation of Board Orders or Negotiated Stipulations or Diversion Program Contracts. It shall be considered unprofessional conduct for a Colorado-licensed pharmacist or intern to violate a lawful Board order or negotiated stipulation issued in result of a formal complaint against the licensee or to violate a peer health assistance diversion program contract.
- 1.00.22 A pharmacist has a professional responsibility to report to the Board in a timely manner any pattern of misconduct in the practice of pharmacy which constitutes a danger to the health, safety, or welfare of a patient or the public.
- 1.00.23 Severability Clause. If any word, clause, sentence, paragraph, or section of these Rules of Professional Conduct shall for any reason be adjudged by any court of competent jurisdiction to be unconstitutional or otherwise invalid, such judgment shall not affect, repeal, or invalidate the remainder thereof, but shall be confined in its operation to the word, clause, sentence, paragraph, section thereof so found to be unconstitutional or otherwise invalid.
- 1.00.24 A prescription drug outlet shall ensure that all prescription drugs and controlled substances are procured from another entity or person registered by the Board. Any drug designated as an Investigational New Drug from the Federal Food and Drug Administration is exempt from this requirement provided the research requirements for the receipt of the product are followed and it meets the requirements of CRS 12-42.5-128(2).

**2.00.00 ORDERS.**

**2.00.10 Receipt of Order.**

- a. Only a pharmacist or intern may receive and reduce to writing an oral order except for chart orders as provided in CRS 12-42.5-118(11).
- b. An electronically transmitted order (ETO) may be accepted in a PDO for dispensing.

**2.01.10 Information to Appear on Each Order.** The following information must appear on each written or oral order except as provided for chart orders for hospitalized patients (hospital chart orders):

- a. The date the order was compounded and dispensed;
- b. The assigned serial number (hospital chart orders are exempt from this requirement).
- c. The quantity dispensed if differs from the quantity ordered (LTCF chart orders are exempt from this requirement provided this information is recorded within another appropriate uniformly maintain and readily retrievable permanent record of the dispensing pharmacy); and.
- d. In the case of a controlled substance order, the patient address, prescriber address, and prescriber's Drug Enforcement Administration (DEA) registration and.
- e. Patient address, prescriber address, and prescriber DEA registration number need not appear on any type of order for a non-controlled substance prescription.

**2.01.20 Additional Information.** The following shall also appear on the prescription or LTCF chart order when appropriate:

- a. Any change in or clarification of an order shall be documented on the order and shall bear the initials of the responsible pharmacist or intern, the date contacted and the name of the individual conveying such change or clarification.

- b. When a substitution is made, the order shall indicate the following:
  - (1) The names of both the drug prescribed and the drug actually dispensed, as well as the date on which such substitution was initially made.
  - (2) The order shall also indicate the name of the distributor of the drug dispensed as it appears on the package or the national drug code number.
  - (3) On an order for a schedule II controlled substance, substitution shall not be deemed to be an alteration of the order.
  - (4) On subsequent refilling of any order, any change in the name of the distributor or the national drug code number as it appears on the package shall be recorded on the order unless the computer system used at that prescription drug outlet changes only the affected transaction(s) (any computer entry change must not alter previous transaction records).
- c. In the case of a chart order for a hospitalized patient (hospital chart order), the following information need not necessarily appear on the chart order, provided that such information is recorded on another appropriate, uniformly maintained and readily retrievable permanent record which reflects:
  - (1) The identity of the pharmacist making the initial interpretation;
  - (2) The identity of the pharmacist making the final evaluation each time a drug is dispensed, if different from the pharmacist making the initial interpretation;
  - (3) The quantity dispensed and
  - (4) The date of dispensing.
  - (5) Any record of a controlled substance dispensed pursuant to a chart order for an individual patient shall be visually identifiable from records of non-controlled substances.

2.01.30 Responsibility of a Pharmacist in Recording Refills. When a prescription order is refilled, the following information must be recorded on the back of the prescription order, or on the daily computer printout as specified in rule 11.00.00, and may be entered by a pharmacy technician if no interpretation is required: Date refilled and quantity, if different from the quantity shown on the face of the prescription order. If authority to refill is obtained, the name of the individual conveying such authority must be recorded. The entry shall also bear the name, initials, license number, or secure electronic identifier of the pharmacist making the final evaluation. This information shall be maintained and available for inspection for a period of two years from the date of any transaction relating to the order unless otherwise required by statute.

2.01.40 Prescription Order Copies. A pharmacist may issue a written copy conspicuously marked "COPY FOR REFERENCE ONLY" to the patient or patient's agent. A pharmacist who issues such a written copy of a prescription order shall place on the original prescription order his/her initials, the date, and an indication that a written copy has been issued. No information regarding authority to refill shall be issued in a written copy.

**2.01.50 Transfer of Prescription Orders Between Prescription Drug Outlets.**

- a. A prescription label or a written copy of a prescription order from another pharmacy may be used for informational purposes only and shall not be considered to be a valid prescription order. A pharmacist who receives such a label or prescription order copy shall either contact the prescribing practitioner for authorization to dispense the prescription, or, alternatively, shall comply with 2.01.52 through 2.01.59.
- b. A pharmacist may orally transfer prescription order information for non-controlled substances for the purpose of dispensing a prescription if the information is communicated by one pharmacist to another pharmacist or an intern, or by an intern under the direct supervision of a pharmacist to another pharmacist.
- c. A prescription drug outlet may transfer a prescription order electronically to another prescription drug outlet for the purpose of dispensing a prescription order.
  - (1) If the prescription order information is transmitted by facsimile, the transferring pharmacist shall comply with rule 2.01.52.
  - (2) Prescription order information may be transmitted electronically between two compatible computer systems that are capable of complying with the requirements of rules 2.01.52 and 2.01.53 (1)-(10). In the case of electronic transfers, the transferring and receiving pharmacist may be the same person.
  - (3) In the case of prescription drug outlets that access and share the same data storage device and that can electronically retrieve all necessary information, if the original prescription order information is not invalidated, each dispensing prescription drug outlet shall be capable of accessing a transaction record that indicates the following information: (a) date, (b) time, and (c) location from which the prescription was dispensed. If the prescription order is assigned a new prescription number at the receiving pharmacy, the prescription information at the originating pharmacy shall be invalidated.
- d. The one-time transfer of original prescription information for a controlled substance listed in schedules III, IV, or V for the purpose of dispensing is permissible between pharmacies. However, pharmacies electronically sharing a real-time, on-line database may transfer up to the maximum refills permitted by law and the prescriber's authorization. If the prescription order is assigned a new prescription number at the receiving pharmacy, the prescription may be transferred on a one-time basis only.
- e. A pharmacist may authorize pharmacy technician to electronically transfer an order, for the purpose of redispensing said order, provided that the electronic transfer is between two compatible computer systems and no changes are made. The pharmacist shall be identified on the transfer record as required by 2.01.52 and 2.01.53.

**2.01.52 The transferring pharmacist shall:**

- a. Write the word "void" across the face of the original prescription order to make the order invalid;
- b. Record on the reverse side of the invalidated prescription order:
  - (1) His/her name, license number, initials, or secure electronic identifier;

- (2) The name, license number, initials, or secure electronic identifier of the receiving pharmacist or intern;
  - (3) The name of the receiving prescription drug outlet;
  - (4) The address and telephone number of the receiving prescription drug outlet; and
  - (5) The date of the transfer.
  - (6) In the case of a controlled substance in schedule III through V, the Drug Enforcement Administration registration number of the receiving prescription drug outlet.
- c. A pharmacy utilizing a computer for storage and retrieval of information regarding prescription transactions shall be exempt from the requirements of paragraphs (a) and (b) of this rule if the computer is capable of invalidating the prescription order and retaining as part of the permanent record the information specified in paragraph (b) of this rule.

2.01.53 The pharmacist receiving the transferred prescription order information shall:

Reduce the transferred information to writing or print; write or print the word "transfer" on the face of the transferred prescription order; and provide all information required by law or rule to be on the prescription order, including:

- (1) The date of issue of the original prescription order;
- (2) The date of initial compounding and dispensing of the original prescription order;
- (3) The number of refills authorized and the original quantity prescribed or any limitations placed on the prescription;
- (4) The number of valid refills remaining;
- (5) The date of the last refill of the original prescription order;
- (6) The prescription order number from which the prescription order information was transferred;
- (7) The name, license number, initials, or secure electronic identifier of the transferring pharmacist or intern;
- (8) The name of the transferring prescription drug outlet;
- (9) The address and telephone number of the transferring prescription drug outlet;
- (10) In the case of a controlled substance in schedules III through V, the Drug Enforcement Administration number of the transferring prescription drug outlet, and the practitioner's Drug Enforcement Administration number.

2.01.54 The transferring prescription drug outlet shall retain the original prescription order as required by rule 11.04.10.

2.01.55 The receiving prescription drug outlet shall retain the transferred prescription order as required by rule 11.04.10.

- 2.01.56 The pharmacist at the receiving prescription drug outlet at the time of the dispensing of the transferred prescription, shall inform the patient that the prescription order is now invalid at the prescription drug outlet from which it was transferred.
- 2.01.58 Nothing in this rule shall be deemed to permit the transfer of a prescription order for a schedule II controlled substance.
- 2.01.59 A prescription order for a controlled substance in schedule III through V may be transferred only one time, that transfer being from the prescription drug outlet where the prescription was originally filled. It shall not be further transferred by, or to, any other prescription drug outlet.
- 2.01.60 A prescription order for a non-controlled prescription drug may be transferred from a prescription drug outlet to another prescription drug outlet as provided in 2.01.50 only so long as there are refills remaining and each prescription drug outlet can establish that a valid refill existed at the time of dispensing.
- 2.01.80 When a prescription drug outlet discontinues business and the prescription order files are moved to another prescription drug outlet, those orders shall be considered void and shall not be refilled. However, if the receiving pharmacist can establish that an authorized refill or authorized refills remain on any such order, such authorization may, at the sole discretion of the pharmacist, be used to establish a new order.
- a. If the record which reflects the authorized refill or refills is the original prescription order, the serial number of the original prescription order shall be recorded on the new order, and the serial number of the new prescription order shall be recorded on the original order.
  - b. If the record which reflects the authorized refill or refills is electronic, the pharmacist shall maintain in written or printed form a record which indicates both the serial number of the original prescription order and the serial number of the new prescription order. This record may be made part of the daily printout required by rule 11.04.20 if it is routinely recorded in such printout. The refill authorization(s) contained in the original electronic record must be invalidated to prevent further refilling.
  - c. The files from the prescription drug outlet that has discontinued business may be transferred to another prescription drug outlet under the following conditions:
    - (1) The computer or electronic database from the prescription drug outlet that discontinued business is located and will remain at the pharmacy to which it is transferred for at least two years.
    - (2) The computer or electronic database must be capable of complying with rule 2.01.52(c).

**3.00.00        DISPENSING.**

- 3.00.10 Limitations. Except as provided in CRS 12-42.5-120(2), no order shall be dispensed or refilled after one year from the date of issue by the practitioner.
- 3.00.20        Medical Need.
- (a) No licensee or registrant shall compound, dispense, deliver or distribute any drug to any person in such quantity or in any situation where the licensee or registrant knows or reasonably should know said drug has no recognized medical utility or application. Violation of this rule shall constitute prima facie proof of violation of CRS 12-42.5-123.

- (b) One additional bottle of a prescription eye drop may be dispensed to a patient if the following conditions are met:
    - 1. The corresponding patient's health benefit plan provides coverage for the prescription eye drops;
    - 2. The additional bottle is requested by the insured or the health care provider at the time the original prescription is dispensed;
    - 3. The original order states that one additional bottle is needed by the insured for use in a day care center, school, or adult day program;
    - 4. The additional bottle is limited to one additional bottle every three (3) months; and
    - 5. The total number of bottles dispensed does not exceed the total number of bottles prescribed as stated on the original order when accounting for authorized refills assigned to the original order by the prescriber, if applicable.
  - (c) A prescription eye drop may be refilled if the following conditions are met:
    - 1. The refill is requested by the insured at least twenty-one (21) days for a thirty (30) day supply of eye drops, forty-two (42) days for a sixty (60) day supply of eye drops, or sixty-three (63) days for a ninety (90) day supply of eye drops, from the later of the date that the original prescription was dispensed to the insured or the date that the last refill of the prescription was dispensed to the insured; and
    - 2. The original prescription order states that additional quantities of prescription eye drops are needed and the refill requested by the insured does not exceed the number of additional quantities needed.
  - (d) The pharmacist may not dispense a prescription drug or a controlled substance to a practitioner based on an order that does not list a specific patient. A prescription order for "office use" is not a valid order. Compounded prescription drugs distributed to veterinarians for "office stock" as defined in CRS 12-42.5-118.5(5)(b) must comply with the requirements of Rules 11.00.00 and 21.00.00.
- 3.00.21 A pharmacist shall make every reasonable effort to ensure that any order, regardless of the means of transmission, has been issued for a legitimate medical purpose by an authorized practitioner. A pharmacist shall not dispense a prescription drug if the pharmacist knows or should know that the order for such drug was issued without a valid preexisting patient-practitioner relationship. Such relationship need not involve an in-person encounter between the patient and practitioner if otherwise permissible under Colorado law. A pharmacist may, in good faith, dispense an opiate antagonist pursuant to an order that was issued without a valid preexisting patient-practitioner relationship under the following conditions:
- a. The opiate antagonist is not a controlled substance; and
  - b. The opiate antagonist is approved by the Federal Food and Drug Administration for the treatment of a drug overdose.
- 3.00.22 The dispensing of an opiate antagonist, as described in Rule 3.00.21, by a pharmacist shall not constitute unprofessional conduct pursuant to CRS 12-42.5-123 if he or she dispensed the opiate antagonist in good faith pursuant to an order or standing orders and protocols issued to or for the following:

- a. A person who is at increased risk of experiencing or likely to experience an opiate-related drug overdose event; or
- b. A family member, friend, or other person who is in a position to assist a person who is at increased risk of experiencing or likely to experience an opiate-related drug overdose event; or
- c. An employee or volunteer of a harm reduction organization; or
- d. A first responder.
- e. For the purpose of this Rule 3.00.22, the following definitions apply:
  - 1) "First responder" means a peace officer, firefighter, or volunteer firefighter.
  - 2) "Harm reduction organization" means an organization that provides services, including medical care, counseling, homeless services, or drug treatment, to individuals at risk of experiencing an opiate-related drug overdose event or to the friends and family members of an at-risk individual.
  - 3) "Opiate-related drug overdose event" means an acute condition, including but not limited to, a decreased level of consciousness or respiratory depression resulting from the consumption or use of a controlled substance, or another substance with which a controlled substance was combined, and that a layperson would reasonably believe to be an opiate related drug overdose event that requires medical attention.
  - 4) "Protocol" means a specific written plan, as maintained in a uniform and readily retrievable manner for the purpose of inspection at the prescription drug outlet for at least two (2) years from the date of the latest dispensing transaction related to protocol, for a course of medical treatment containing a written set of specific directions created by a physician, group of physicians, hospital medical committee, pharmacy and therapeutics committee, or other similar practitioners or groups of practitioners with expertise in the use of opiate antagonists.
  - 5) "Standing order" means a prescription order, as maintained in a readily retrievable manner for the purpose of inspection at the prescription drug outlet for at least two (2) years from the date of the latest dispensing transaction related to order, written by a practitioner that is not specific to and does not identify a particular patient.
- f. Each prescription drug outlet shall maintain, in a uniform and readily retrievable manner for at least two (2) years from the date of latest transaction related to a standing order, the following record detailing the dispensing of a non-controlled substance opioid antagonist pursuant to a standing order:
  - 1) The full name of the patient, person who is in a position to assist a person who is at increased risk of experiencing or likely to experience an opiate-related drug overdose event, first responder, or harm reduction organization receiving the drug;
  - 2) The full address of the first responder or harm reduction organization receiving the drug;
  - 3) The name, strength and dosage form of the drug dispensed;



- 4) The quantity of drug dispensed; and
- 5) The date of dispensing.

3.00.25 First Dose Dispensing. A pharmacist at a prescription drug outlet may dispense up to a seventy two (72) hour supply of a non-controlled substance prescription drug to an LTCF resident pursuant to a duplicate copy of an LTCF chart order provided by another prescription drug outlet for the purpose of providing immediate patient care, on a one time per order basis, if the following conditions are met:

- a. The receiving prescription drug outlet records on the prescription order the name and address of the originating prescription drug outlet and the date the order was received by the receiving prescription drug outlet;
- b. The receiving prescription drug outlet maintains the order as a prescription order and complies with all requirements for prescription orders specified in Board Rules 2.01.10 through 2.01.40, 3.00.10 through 3.00.51, and 11.04.10; and
- c. The originating prescription drug outlet records on the LTCF chart order the name and address of the receiving prescription drug outlet and the date the order was provided to the receiving prescription drug outlet.

3.00.27 Outlet to Outlet Drug Reconstitution. A pharmacist at a prescription drug outlet may reconstitute a prescription originally dispensed in an unreconstituted form pursuant to a patient-specific order at another prescription drug outlet or nonresident prescription drug outlet provided the following conditions are met:

- a. The prescription is delivered directly from the originating outlet to the receiving outlet;
- b. The prescription is at no time in the physical possession of the patient until after the prescription has been reconstituted;
- c. The prescription is reconstituted according to the corresponding manufacturer's directions;
- d. The prescription is not a controlled substance;
- e. The pharmacist at the receiving outlet does not alter the prescription or its original labeling in any way other than to reconstitute, re-label for re-dispensing for administration, and properly store the prescription; and
- f. The originating outlet is ultimately accountable to the Board for the accurate dispensing of the original prescription, and the receiving outlet is ultimately accountable for the accurate reconstitution and re-dispensing of the prescription.

3.00.30 Labeling. When a prescription drug is dispensed pursuant to an order, the name of the drug that appears on the container label shall correspond with the identity of the drug contained therein unless otherwise requested by the practitioner.

3.00.40 Expiration Dating. No drug or device shall be dispensed which will be outdated prior to utilization by the consumer, based on the practitioner's directions for use.

3.00.50 Initial Interpretation and Final Evaluation.

- a. Initial interpretation means the review of an order accompanied by order entry. The pharmacist(s) conducting the initial interpretation shall be held accountable for the accuracy of the electronic order entry/manual transcription and for drug regimen review.
- b. Final evaluation means the review of the final prescription to ensure that the ordered medication is properly prepared and placed in a suitable container with appropriate labeling. The pharmacist(s) conducting the final evaluation shall be held accountable for assuring that the identity of the drug that appears on the prescription label corresponds with identity of drug contained therein. When refills are dispensed, the pharmacist conducting the final evaluation shall be held accountable for the appropriate dispensing of refills including all drug utilization reviews as they pertain to refill dispensing.
- c. Drug regimen review includes but is not limited to the evaluation of order(s) and patient records(s) for:
  - 1) Known allergies;
  - 2) Rational therapy and contraindications;
  - 3) Reasonable dose, duration of use, and route of administration considering age, gender, and other patient factors;
  - 4) Reasonable directions for use;
  - 5) Potential or actual adverse drug reactions;
  - 6) Drug-drug interactions;
  - 7) Drug-food interactions;
  - 8) Drug-disease contraindications;
  - 9) Therapeutic duplication;
  - 10) Proper utilization (including over- or under-utilization) and optimum therapeutic outcomes; and
  - 11) Abuse/misuse.
- d. A pharmacist shall conduct an initial interpretation of each new order and a pharmacist shall conduct the final evaluation of each order dispensed. When refills are dispensed, the pharmacist making the final evaluation shall be held accountable for the appropriate dispensing of refills. The pharmacist manager shall be held accountable for the maintenance of all appropriate records.
- e. The pharmacist making the initial interpretation and final evaluation on prescription or LTCF chart orders shall be identified by either license number, initials, name, or secure electronic identifier on a uniformly maintained, readily retrievable document. The uniformly maintained, readily retrievable document shall bear the license number, initials, name, or secure electronic identifier of any additional pharmacists involved in the dispensing of the order. The pharmacist conducting the initial interpretation and final evaluation may be the same person.

- f. In the case where the computer software utilized is not password protected, the initial interpretation and final evaluation shall be maintained in a handwritten format bearing the license number, initials, or name of the responsible pharmacist. In addition, the identification of any other pharmacists involved in the dispensing shall be maintained in the same handwritten format.

**3.00.51 Records of Initial Interpretation and Final Evaluation.**

- a. Records detailing both the initial interpretation and final evaluation shall be retained at the prescription drug outlet for each prescription dispensed and for at least two years from the date of any transaction pertaining to the order. These records shall include at least the following:
  - 1. The license number, initials, name, or secure electronic identifier of the pharmacist conducting the initial interpretation for each new order;
  - 2. The license number, initials, name, or secure electronic identifier of the pharmacist conducting the final evaluation for each new and refill prescription; and
  - 3. The specific date on which each initial interpretation and final evaluation occurred. In the event the initial interpretation and final evaluation for a new order are conducted on separate dates, both dates shall be recorded to state specifically when both occurred.
- b. Each outlet shall maintain, in written format, a notice detailing how initial interpretations and final evaluations are documented in the outlet. Such notice shall include and comply with the following:
  - 1. The manner in which initial interpretations are recorded and maintained in the outlet for all new orders.
  - 2. The manner in which final evaluations are recorded in the outlet for all new and refill prescriptions.
  - 3. A statement that all pharmacy personnel involved in the dispensing of prescriptions have the ability to print, upon request, a record detailing the initial interpretation for each new prescription dispensed and final evaluation for each new and refill prescription dispensed.
  - 4. Such written notice shall be signed and dated by the pharmacist manager. In the event the pharmacist manager changes, the incoming pharmacist manager shall review, sign and date the notice within 72 hours of assuming the duties of pharmacist manager. In the event there is a lapse between the time one pharmacist manager ceases the duty and another assumes the duty, the previous method of recording initial interpretations and final evaluations shall remain in effect.
  - 5. If there are any changes to the outlet's method of documenting initial interpretations and final evaluations, a new written notice detailing the requirements of sections 1, 2, 3, and 4 above shall be executed. This notice shall detail the effective date of change.
  - 6. The outlet shall post these notices on a wall directly next to the outlet's most current Board registration.

7. These notices shall be retained at the outlet for a period of three years from the date last utilized.
8. In the event such notices are not posted, the pharmacist manager shall be held accountable for the failure to post the required notice and any dispensing errors. In the event such notices are not posted during the period of time between one pharmacist manager leaving the position and another assuming the position, the outlet shall be held accountable for the failure to post the required notice and any dispensing errors.

3.00.55 Prescription Flavoring. A flavor additive may be incorporated into a non-sterile prescription under the following conditions:

- a. The patient, patient's caregiver, or practitioner who authorized the original prescription shall authorize the flavoring of each new and, if applicable, refilled prescription;
- b. The flavor additive shall in no way compromise the stability, safety, or efficacy of the dispensed drug.
- c. No expired flavor additive shall be incorporated into a prescription. No flavor additive shall be incorporated which will expire prior to utilization by the patient, based on the practitioner's directions for use.
- d. For flavoring additives that do not have expiration dates assigned by the manufacturer or supplier, a pharmacist shall clearly and legibly label the container with the date of receipt and assign a conservative expiration date, not to exceed three (3) years after receipt, to the flavoring additive. In no event shall the labeled date of receipt or assigned expiration date be later altered after originally labeling the container.
- e. The following information shall be recorded and maintained in a suitable hard-copy or electronic dispensing record for a period of two years from the date of flavoring the corresponding new or refilled prescription. This record shall be made available, in printed form, for the Board or its representatives immediately upon the request of the Board or its representatives.
  - (1) Additive's flavor;
  - (2) Flavor additive's manufacturer
  - (3) Flavor additive's lot number (if available); and
  - (4) Flavor additive's expiration date.
- f. The pharmacist responsible for conducting the final evaluation of a new or refilled prescription shall also be responsible for the flavoring of the prescription as specified in subsections a., b., and c. of this Rule 3.00.55.
- g. The pharmacist manager shall be responsible for subsection d. of this Rule 3.00.55 and the maintenance of records as specified in subsection e. of this Rule 3.00.55.

3.00.60 When a substitution is made on a prescription order, a patient shall be given oral and written notice of this fact at the time such substitution initially occurs, except as provided in CRS 12-42.5-122. On subsequent refilling of a prescription order, such oral and written notices shall not be required unless, in the professional judgment of the pharmacist, the best interest of the patient will be served by giving such notices.

3.00.70 Responsibility for pharmacy technicians. A pharmacist shall be responsible for pharmacy technicians and shall at all times comply with CRS 12-42.5-116(5).

3.00.75 The placement of a prescription into another outer container and the labeling of the container with the patient's name or any other identifying information constitutes the "Practice of Pharmacy" as a function of preparation, packaging, labeling and delivery under CRS 12-42.5-102(31). Individuals who perform this function shall be included in the ratio of pharmacy technicians or interns a pharmacist is permitted to supervise pursuant to 12-42.5-119(1).

3.00.80 Return or Exchange of Drugs, Prescriptions, Medical Devices, and Medical Supplies for Dispensing or Donation.

3.00.81 Definitions.

For the purposes of this rule 3.00.00, the following definitions apply:

- a. "Automated cassette" is a container that is filled with a drug. This container may count the drug and may package the drug into a container suitable for dispensing, and may affix a label to the container. These cassettes may be used to dispense drugs in a traditional dispensing system or may be used to package unit-dose medication, or drugs in a unit of issue packaging system. An automated cassette shall not be used for schedule II controlled substances.
- b. "Correctional facility" means a facility under the supervision of the United States, the Department of Corrections, or a similar state agency or department in a state other than Colorado in which persons are or may be lawfully held in custody as a result of conviction of a crime; a jail or an adult detention center of a county, city, or city and county; and a private contract prison operated by a state, county, city or city and county.
- c. "Customized patient medication package" means a package which contains two or more drugs.
- d. "Licensed Facility" means any of the following facilities licensed by the Colorado Department of Public Health and Environment: community mental health center, acute treatment unit, hospital unit, inpatient hospice, nursing care facility, assisted living residence, or long-term care facility.
- e. "Medical Device" means an instrument, apparatus, implement, machine, contrivance, implant, or similar or related article that is required to be labeled pursuant to 21 CFR Part 801.
- f. "Medical Supply" means a consumable supply item that is disposable and not intended for reuse.
- g. "Nonprofit Entity" means a Board registered prescription drug outlet or other outlet which has nonprofit status, or an out-of-state entity with legal authority to both possess a prescription drug and receive a donated prescription drug distributed from a Board-registered outlet in the state of Colorado.
- h. "Originating Prescription Drug Outlet" means the prescription drug outlet which initially dispensed the prescription for a resident of a facility.

- i. "Package" means to prepare a drug in a container other than the original container. The packaging might include a unit dose dispensing system, single dose, automated cassette, or a container suitable for a traditional system. Unless otherwise specified, this includes preparing a drug in advance of the immediate need for dispensing (prior to the receipt of an order), or pursuant to an existing order.
- j. "Single dose package" means a package which contains a quantity of a drug intended for administration as a single dose.
- k. "Traditional dispensing system" means a drug package system in which individual doses are not packaged in unit dose packages or unit of issue packages.
- l. "Unique identifier" means an implicit or explicit unique identifier from which the originating prescription number can be determined.
- m. "Unit dose dispensing system" means a drug distribution system which is in a prescription drug outlet or hospital other outlet and uses unit dose packages or unit of issue packages that enable distribution of packaged doses in a manner that preserves the identity of the drug until the time of administration.
- n. "Unit dose package" means a package which contains one pharmaceutical unit.
- o. "Unit of issue package" means a package which provides multiple units of doses but separated in a medication card or other specifically designed container.

#### 3.00.82 General Provisions

- a. No prescription drug outlet shall accept returned or donated prescriptions, medical devices, or medical supplies for dispensing, or donation except in the following situations:
  - (1) A prescription drug outlet that complies with rules 3.00.82 through 3.00.89 may accept prescriptions, medical devices, and medical supplies for return, dispensing, and donation.
  - (2) A hospital prescription drug outlet may accept prescriptions and drugs for dispensing or reissue from all areas of the hospital, provided that the integrity of the product and package are maintained and the following requirements are met:
    - (a) An appropriate, uniformly maintained and readily retrievable record shall be maintained which indicates at least the total number of doses of the drug which were actually administered. This record may be combined with the record permitted by rule 2.01.20(c); or
    - (b) If the drug was distributed as floor stock in the facility, an appropriate, uniformly maintained and readily retrievable record of such return shall be made. This record shall state the following:
      - (I) The name of the drug;
      - (II) The strength of the drug;
      - (III) The dosage form of the drug if appropriate;
      - (IV) The quantity of the drug;

(V) The location within the facility to which the drug was originally distributed; and

(VI) The date of the return.

- b. No prescription drug returned for redispensing or donation from a facility or donated by a prescription drug outlet shall be redispensed if it expires prior to utilization by the consumer based on the prescribing practitioner's directions for use.
- c. Rules 3.00.80 through 3.00.89 do not apply to the Colorado Cancer Drug Repository.

**3.00.83 Entities Eligible to Donate or Return Prescriptions.**

The following may donate or return drugs:

- a. A correctional facility as defined in 3.00.81(b), a licensed facility as defined in 3.00.81(d), or any other facility that is required to be licensed pursuant to section 25-3-101, C.R.S. may return prescriptions to a prescription drug outlet.
- b. A correctional facility, a licensed facility, or any other facility that is required to be licensed pursuant to section 25-3-101, C.R.S. may donate prescriptions to a nonprofit entity as defined in 3.00.81(g) or to a practitioner authorized by law to dispense the prescription.
- c. A prescription drug outlet may donate a returned or donated prescription to a nonprofit entity as defined in 3.00.81(g) or to a practitioner authorized by law to dispense the prescription.

**3.00.84 Eligibility for Return or Donation.**

- a. For all prescriptions, medical devices, or medical supplies accepted for return or donation, the prescription drug outlet must ensure that the prescription, medical device, or medical supply was properly stored prior to return or donation. This includes storage at the facility, and shipment to and from the facility.
- b. Drugs which have been dispensed to a resident of a correctional facility, licensed facility, or any other facility that is required to be licensed pursuant to section 25-3-101, C.R.S. that are eligible for return or donation are as follows:
  - 1) Drugs which are liquid and the vial is still sealed and properly stored;
  - 2) Drugs that have been individually packaged and the packaging has not been damaged; and
  - 3) Drugs that are in the original, unopened, sealed, and tamper-evident unit dose package, unit of issue package, or unit dose dispensing system.
- c. Drugs which have been dispensed to a resident of a correctional facility, licensed facility, or any other facility that is required to be licensed pursuant to section 25-3-101, C.R.S. that are not eligible for Return or Donation are as follows:
  - 1) Any drug declared to be a controlled substance under any state or federal law or rule except as provided in 3.00.82(a)(2);
  - 2) Any drug dispensed in a traditional dispensing system;

- 3) Any drugs dispensed in a customized patient medication package;
- 4) Any drug packaged in a single dose package, a unit dose dispensing system, a unit dose package, or a unit of issue package that is not labeled in accordance with 3.01.20 and 3.01.21;
- 5) A compounded drug;
- 6) Drugs that are adulterated or misbranded as determined by the pharmacist;
- 7) Drugs that require refrigeration, freezing, or special storage;
- 8) Drugs that require special registration with the manufacturer;
- 9) Drugs that will expire prior to utilization by the consumer, based on the prescribing practitioner's directions for use;
- 10) Dispensed drugs that are received from facilities or pharmacies located outside of Colorado; and
- 11) Any drug that was not dispensed pursuant to an order.

**3.00.85 Records of Receipt of Returned or Donated Prescriptions, Medical Devices, and Medical Supplies.**

- a. The prescription drug outlet shall retain records for at least two years detailing receipt of donated or returned prescriptions that contain at least the following information:
  - (1) Name and address of facility or donating prescription drug outlet;
  - (2) Name and address of originating prescription drug outlet;
  - (3) Prescription number or unique identifier assigned at originating prescription drug outlet;
  - (4) Name and address of each prescription drug outlet having possession of the drug, device, or supply after the originating prescription drug outlet and the dates the product was in each prescription drug outlet's possession.
  - (5) Date of return or donation;
  - (6) Name, strength, and NDC number of drug received;
  - (7) Name of medical device or medical supply received; if applicable;
  - (8) Quantity received;
  - (9) Date received;
  - (10) Drug, medical device, or medical supply expiration date;
  - (11) Receipt record must state, "Returned or Donated Prescription, Device, or Supply"



- b. Records detailing the receipt of returned or donated prescriptions, devices, and supplies, as required by rule 3.00.84(a)(1) through (11) may be maintained electronically if the following requirements are met:
  - (1) The prescription drug outlet must ensure a daily (i.e., every twenty-four hours) back up is performed for use in restoring required information in case of a system failure;
  - (2) Have and maintain a complete on-line receipt file that is printable on the inspector's request;
  - (3) Have a "lock-out" feature that prevents editing of receipt information;
  - (4) The Board or its inspectors must be able to inspect and review all of the prescription drug receipt transactions of the outlet for the preceding two years. Therefore, immediately upon the oral or written request of the Board or its inspectors, the outlet shall either:
    - (a) Print a report of all prescription drug receipt transactions for a period of time as the Board or its inspector(s) may specify. The system must be capable of retrieving and printing such a report within a limited time not to exceed two hours; or
    - (b) Provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review prescription drug receipt transactions, and if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. If the outlet elects to comply with this subparagraph (2), the system must also be capable of printing the same reports described in subparagraph (1); or
  - (5) It is the responsibility of the pharmacist manager of the outlet to ensure that all outlet staff is aware of the requirements of subparagraphs (1) and (2). Any failure or refusal by the pharmacist manager and/or staff to comply with a request by the Board or its inspector(s) will be deemed to be a willful violation of these rules.

**3.00.86 Storage of Returned or Donated Prescription, Medical Devices/Supplies, and Establishment of Handling Fee.**

- a. Returned or donated prescriptions, medical devices, and medical supplies shall be stored in a separate area from other drug stocks belonging to the pharmacy. This area shall be conspicuously labeled with a sign indicating that such area contains only returned or donated prescriptions, medical devices, or medical supplies.
- b. An entity that receives a donated medication, medical device or medical supply may charge the end user a handling fee, which shall not exceed three (3) dollars for each complete prescription, medical device or medical supply dispensed to the end user and shall not resell the donated medication, medical device or medical supply for profit.

**3.00.87 Dispensing of Returned or Donated Prescriptions, Medical Devices, or Medical Supplies.**

- a. Special Conditions for Dispensing Returned or Donated Drugs:

- (1) Drug products in manufacturer's unit dose or unit of issue packages may be redispensed as often as necessary, provided that the integrity of the product and package are maintained.
- (2) Drug products which have been packaged into unit dose or unit of issue packages in the prescription drug outlet may be redispensed one time only, except as provided for in 3.00.82((a)(2), provided that the integrity of the product and the package are maintained.
- (3) Drug products which have been packaged into unit of issue packages in the prescription drug outlet may be redispensed one time only and then only in the package in which originally dispensed, except as provided in (5) below. Partially-used unit of issue packages may not be emptied and the drugs removed and packaged, nor may additional units of medication be added to partially-used unit of issue packages.
- (4) Drug products which have been packaged into single dose packages in the prescription drug outlet may be redispensed one time only and then only in the package in which originally dispensed, except as provided in (5) below. Single dose packages may not be emptied and the drugs removed and packaged.
- (5) Drug products which have been packaged into unit of issue packages or single dose packages may be removed from such packages and packaged for dispensing in a traditional dispensing system.
- (6) Prescriptions dispensed using returned or donated prescriptions shall be labeled according to CRS 12-42.5-121. Additionally, the label shall state, "Donated or Returned Drug."

**b. Records of Dispensing**

All records of dispensing shall be compliant with rules 2.00.00, 3.00.00, and 11.00.00. These records of dispensing, including prescription orders, shall be maintained separately from dispensing records of drugs that were not donated or returned.

**3.00.88 Donating Returned or Donated Prescriptions, Medical Devices, or Medical Supplies.**

**a. Prescription drug outlets may donate the returned or donated prescriptions, medical devices, or medical supplies. to any of the following:**

- (1) Nonprofit entity as defined in 3.00.81(g); or
- (2) A practitioner authorized by law to dispense the drug.

**b. Records of donation shall include the following:**

- (1) The name of the drug, medical device, or medical supply;
- (2) The strength of the drug;
- (3) The dosage form if appropriate;
- (4) The quantity of the drug, medical device, or medical supply;

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- (5) The manufacturer name and/or NDC number of the drug if labeled only with its generic name;
  - (6) The date of donation;
  - (7) The name and address of the donating prescription drug outlet;
  - (8) The name and address and registration number of the nonprofit entity receiving the drug, medical device, or medical supply, or the name, address, and license number of the practitioner receiving the drug, medical device, or medical supply.
  - (9) The name and address of the originating prescription drug outlet;
  - (10) The prescription number or unique identifier assigned to the prescription at the originating prescription drug outlet.
  - (11) The date the medication expires; and
  - (12) The name and address of each prescription drug outlet, other than the originating prescription drug outlet, having possession of the prescription and the dates the prescription was in that prescription drug outlet's possession.
- c. A copy of the donation record shall be maintained at the prescription drug outlet and a copy of the same record shall be furnished to the receiving individual or entity.
- d. Records detailing the donation of prescriptions, medical devices, and medical supplies, as required by rule 3.00.88(b)(1) through (12) may be maintained electronically if the following requirements are met:
- (1) The prescription drug outlet must ensure a daily (i.e., every twenty-four hours) back up is performed for use in restoring required information in case of a system failure;
  - (2) Have and maintain a complete on-line donation file that is printable on the inspector's request;
  - (3) Have a "lock-out" feature that prevents editing of donation information;
  - (4) The Board or its inspectors must be able to inspect and review all of the donation transactions of the outlet for the preceding two years. Therefore, immediately upon the oral or written request of the Board or its inspectors, the outlet shall either:
    - (a) Print a report of all donation transactions for a period of time as the Board or its inspector(s) may specify. The system must be capable of retrieving and printing such a report within a limited time not to exceed two hours; or
    - (b) Provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review donation transactions, and if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. If the outlet elects to comply with this subparagraph (2), the system must also be capable of printing the same reports described in subparagraph (1).
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- (5) It is the responsibility of the pharmacist manager of the outlet to ensure that all outlet staff are aware of the requirements of subparagraphs (1) and (2). Any failure or refusal by the pharmacist manager and/or staff to comply with a request by the Board or its inspector(s) will be deemed to be a willful violation of these rules.

**3.00.89 Record Retention**

- a. All records of receipt and dispensing shall be maintained for a period of two years from the date of receipt, or from the last dispensing transaction date. Such records shall be maintained separately from all other records of the prescription drug outlet.
- b. All records of donation shall be maintained for a period of three years from the date of donation. Such records shall be maintained separately from all other records of the prescription drug outlet.

**3.00.90 Prescriptions Dispensed but Not Delivered.** When a drug has been dispensed pursuant to a prescription or LTCF chart order but has not been delivered to the ultimate consumer, the drug may be returned to stock for subsequent redispensing provided that:

- a. It is stored in the container in which it was dispensed, with the original prescription label intact;
- b. A separate written record or a separate record printable upon request is maintained for prescriptions returned to stock. Such record shall indicate only prescriptions returned to stock and shall list at minimum the following:
  - (1) Prescription number;
  - (2) Drug name and strength;
  - (3) Quantity returned to stock;
  - (4) Date of return; and
  - (5) If centrally filled, the location where filled.
- c. The expiration date of the drug shall not be more than one year from the date it was dispensed. Unless it was dispensed in the manufacturer's original container and bears the manufacturer's original label and expiration date; and
- d. The drug remains under the same ownership from which it was originally dispensed or is dispensed from a pharmacy in which the pharmacy has a contractual affiliation for central fill processing;
- e. If the drug was delivered to another prescription drug outlet for delivery to the ultimate consumer, the following apply:
  - (1) The lot number and manufacturer's expiration date must be placed on the label of the drug container by the original dispensing prescription drug outlet; or
  - (2) The original dispensing prescription drug outlet can access and provide the expiration date and lot number upon request.
  - (3) No controlled substance prescriptions may be returned to stock.

- (4) No compounded or flavored prescriptions may be returned to stock.

3.00.91 Prescriptions dispensed by prescription drug outlets for delivery to consumers in other other outlet settings. When a drug has been dispensed pursuant to prescription order at a prescription drug outlet but has not been delivered to the ultimate consumer at an other outlet, the drug may be returned to stock only at the originating Prescription Drug Outlet, for subsequent redispensing provided that:

- a. The prescription drug outlet complies with Board Rules 3.00.90(a), (b), and (c);
- b. The storage conditions during the transport of the prescription to and from the other outlet do not in any way compromise the integrity or stability of the drug;
- c. No controlled substance prescriptions may be returned to stock; and
- d. No compounded or flavored prescription may be returned to stock.

3.01.00 Packaging.

3.01.10

- a. In a prescription drug outlet packaging shall only be done by a pharmacist, or by an intern or pharmacy technician under the supervision of a pharmacist. In an other outlet, packaging may be done by a person not licensed as a pharmacist pursuant to protocols approved by the Board.
- b. Such packaged drugs shall only be dispensed or distributed from the premises where packaged. Such packaged drugs shall only be distributed as provided in 3.01.10(d).
- c. Any container used for packaging shall meet compendia requirements.
- d. The following prescription drug outlets may distribute packaged medications without limitation to prescription drug outlets under common ownership:
  1. Prescription drug outlets owned and operated by a hospital that is accredited by the joint commission on accreditation of healthcare organizations or a successor organization pursuant to 12-42.5-118(15)(b), C.R.S.;
  2. Prescription drug outlets operated by a health maintenance organization as defined in section 10-16-102, C.R.S.; and
  3. The Colorado Department of Corrections.

3.01.20 Each packaged container, whether for use in a unit dose distribution system or a traditional dispensing system, shall be labeled in accordance with this rule. Any packaged unit dose, single dose or unit of issue container for which return for restocking and redispensing, pursuant to 3.00.80, is anticipated, shall be labeled in accordance with this rule. Additionally, any packaged container from which subsequent dispensing may occur, shall be labeled in accordance with this rule. Such labeling shall include at least the following:

- a. If a suitable internal record is maintained in the prescription drug outlet or other outlet, the requirements of (d), (e), (f), (g), and (h) of this rule may be omitted from the labeling and maintained in such record. An internal lot number shall be assigned and shall appear in the labeling. In a prescription drug outlet the record shall be signed by the pharmacist responsible for each lot packaged. In another outlet the record shall be signed by the person specified in the Board approved protocol. The record shall be retained for two years from the date of packaging unless otherwise required by law or rule.
- b. Name and strength of the medication, and, in the case of a single dose package, the total number of individual tablets or capsules per dose;
- c. A suitable expiration date, which shall be not later than the expiration date on the manufacturer's container, or one year from the date the drug is packaged, whichever is less. Sterile packaged product beyond-use dating shall comply with Rule 3.01.34(h)(3));
- d. The identity of the manufacturer or distributor;
- e. The manufacturer's or distributor's lot number;
- f. The manufacturer's or distributor's expiration date;
- g. The date the product was packaged;
- h. The identity of the pharmacist responsible for packaging in a prescription drug outlet, or, in the case of an other outlet, the identity of the non-pharmacist permitted to do so pursuant to protocols approved by the Board.
- i. The name and address of the packaging pharmacy if the drug is distributed by a prescription drug outlet owned and operated by a hospital that is accredited by the Joint Commission of Accreditation of Healthcare Organizations or a successor organization or by a prescription drug outlet operated by a health maintenance organization as defined in section 10-16-102, C.R.S. Such drugs may only be distributed to prescription drug outlets under common ownership.

3.01.21 If the unit dose package or unit of issue package is obtained from the manufacturer or distributor and complies with applicable federal requirements, such package may be dispensed without additional labeling as required in 3.01.20 above.

3.01.22 Filling of automated cassettes.

- a. If a multi-source drug, the outlet may not use drugs in the same cassette from multiple manufacturers or distributors;
- b. Schedule II controlled substances may not be packaged into automated cassettes.
- c. Automated cassettes, without electronic maintenance or records, shall be labeled with the following:

1. If a suitable internal record is maintained in the prescription drug outlet or other outlet, the requirements of 4, 5, 6, 7, and 8 of this rule may be omitted from the labeling and maintained in such record. An internal lot number shall be assigned and shall appear in the labeling. In a prescription drug outlet the record shall be signed by the pharmacist responsible for each lot packaged. In an other outlet the record shall be signed by the person specified in the Board approved protocol. The record shall be retained for two years from the date of packaging, unless otherwise required by law or rule.
  2. Name and strength of the medication;
  3. A suitable expiration date, which shall be not later than the expiration date on the manufacturer's container, or one year from the date the drug is packaged, whichever is sooner;
  4. The identity of the manufacturer or distributor;
  5. The manufacturer's or distributor's lot number(s);
  6. The manufacturer's or distributor's expiration date;
  7. The date the product was packaged;
  8. The identity of the pharmacist responsible for packaging in a prescription drug outlet, or, in the case of an other outlet, the identity of the non-pharmacist permitted to do so pursuant to protocols approved by the Board;
  9. All records detailing item 1-8 above, shall be retained at the pharmacy for at least two years.
- d. In the event that the automation associated with the cassettes deactivates the cassette when the suitable expiration date is reached, and the outlet either prints packaging printouts on a daily basis or is capable of electronically maintaining the packaging information, the cassette need only be labeled with the name and strength of the drug.
- e. In the event of a product recall, the pharmacist manager shall reasonably ensure that all recalled drug has been removed from the cassette.

**3.01.23 Maintenance of automated cassette records.**

A prescription drug outlet may utilize a computer (automated data processing system) for storage and retrieval of information regarding packaging in automated cassettes. The following requirements shall be met:

- a. All information required by rule 3.01.22 c (1-8) shall be entered into the system at the time of the transaction.

- b. Every 24 hours the system must produce a hard-copy document that, for the purposes of these rules, shall be known as the “packaging printout”. It shall consist of a single, uniform, complete document. The packaging printout shall list, separately, each packaging transaction for the previous 24 hours and shall contain all information required by this rule. Packaging printouts shall be retained in a chronological manner. If the printouts are bound, the sheets shall be separated into individual pages that are then placed in the same order as printed and bound uniformly. If the pages are bound in any other manner so that they are not uniform in placement or appearance, they shall be deemed not readily retrievable and available.

**3.01.24 Electronic Maintenance of Packaging Records.**

A prescription drug outlet which utilizes a computer (automated data processing system) for storage and retrieval of information regarding packaging transactions need not print the packaging printout required by rule 3.01.23 if the prescription drug outlet and the computer system utilized are capable of complying with the following requirements:

- a. The prescription drug outlet must be able to provide on-line retrieval of all information required by this rule for all packaging transactions during the two years preceding the request.
- b. The prescription drug outlet must ensure a daily (i.e., every twenty-four hours) back up is performed for use in restoring required information in case of a system failure.
- c. The prescription drug outlet must:
  - (1) Have and maintain a complete on-line transaction file that is printable on the inspector’s request,
  - or
  - (2) Have a “lock-out” feature that prevents editing of packaging information.
- d. The Board or its inspectors must be able to inspect and review the packaging transactions of the prescription drug outlet. Therefore, immediately upon the oral or written request of the Board or its inspectors, the prescription drug outlet shall either:
  - (1) Print a report of all packaging transactions for such period of time as the Board or its inspector(s) may specify. The system must be capable of retrieving and printing such a report within a limited time not to exceed two hours. Additionally within 72 hours, the system must be capable of retrieving and printing the information sorted according to the variables which include, but are not limited to, date packaged; drug name, strength and dosage form; lot number, manufacturer/distributor; or expiration date.
  - (2) Provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review packaging transactions, and, if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. If the prescription drug outlet elects to comply with this subparagraph (d), the system must also be capable of printing the same reports described in subparagraph (1).



- (3) It is the responsibility of the prescription drug outlet manager to ensure that all prescription drug outlet staff are aware of the requirements of subparagraphs (1) and (2). Any failure or refusal by the prescription drug outlet manager and/or a staff pharmacist to comply with a request by the Board or its inspector(s) will be deemed to be a willful violation of these rules.
- e. Whether the prescription drug outlet elects to comply with rule 3.01.24(d), the system and any reports printed on request shall contain, as a minimum, the following information for each transaction:
  - 1. Name and strength of the medication;
  - 2. A suitable expiration date, which shall be not later than the expiration date on the manufacturer's container, or one year from the date the drug is packaged, whichever is sooner;
  - 3. The identity of the manufacturer or distributor;
  - 4. The manufacturer's or distributor's lot number(s);
  - 5. The manufacturer's or distributor's expiration date;
  - 6. The date the product was packaged;
  - 7. The identity of the pharmacist responsible for packaging in a prescription drug outlet, or, in the case of an other outlet, the identity of the non-pharmacist permitted to do so pursuant to protocols approved by the Board;

**3.01.25 Maintenance and cleaning of automated cassettes**

- a. The outlet must maintain, on-site and available for inspection, the manufacturer's guidelines for maintenance and cleaning of the cassettes.
- b. The maintenance and cleaning schedule recommended by the manufacturer shall be adhered to and records of performed maintenance shall be available for inspection for a period of at least two years.
- c. If the outlet changes the drug used in a cassette, the cassette must be thoroughly cleaned per manufacturer's recommendations prior to using the cassette for a different drug.

**3.01.26 Responsibility for unit-dose medications packaged with automated cassettes is the responsibility of the pharmacist responsible for loading the cassette.**

**3.01.27 The pharmacist responsible for the final evaluation of any prescriptions dispensed using drugs packaged in automated cassettes shall be held accountable for the accuracy of the product.**

**3.01.30 Sterile Product Packaging.**

**3.01.32 "Sterile product packaging" means the transfer of a sterile drug, in part or in whole, from one container or device to another container or device in advance of immediate need for the purposes of future dispensing or distribution.**

3.01.34

- a. Each packaged container shall be labeled according to Rule 3.01.20(a) through (i).
- b. All sterile products shall be packaged under the environmental quality, controls and monitoring specified in Rules 21.20.60 through 21.60.90.
- c. Each outlet engaged in sterile product packaging shall maintain a policy and procedure manual that shall be reviewed, signed and dated by the pharmacist manager at least once annually, and within 30 days of a new pharmacist manager assuming that position. The manual shall at least address the following:
  - (1) Responsibilities of sterile product packaging personnel;
  - (2) Verification of packaging sterilization if not using packaging that is sterile
  - (3) Personnel training and evaluation in aseptic manipulation skills;
  - (4) Environmental quality and control;
  - (5) Aseptic processing;
  - (6) Labeling and recordkeeping;
  - (7) Finished preparation release check;
  - (8) Storage and beyond-use dating;
  - (9) Maintaining product quality and control during transportation and delivery after the packaged sterile product leaves the pharmacy;
  - (10) Patient or caregiver training;
  - (11) Adverse event reporting and recalls;
  - (12) Quality assurance program;
  - (13) Quality control procedures; and
  - (15) Manner by which storage excursions are handled and documented.
- d. All pharmacy personnel engaged in sterile product packaging shall receive suitable didactic and experiential training which, at minimum, includes aseptic processing, environmental testing, as well as the selection and use of containers, equipment and closures.
- e. Written procedures outlining equipment used in sterile product packaging, calibration, appropriate maintenance, monitoring for proper function, controlled procedures for use of the equipment, and specified time frames for these activities shall be established and followed. Results of equipment calibration and appropriate maintenance reports shall be maintained on file at the outlet for at least two years from the report date and shall be available for inspection.

- f. Accuracy assessments of automated sterile packaging devices shall be conducted daily for each day used. At least annually, the pharmacist manager, or his or her designee, shall review these assessments to avoid potentially significant errors. These assessments shall be documented and available for inspection at the outlet for at least two years.
- g. All sterile product packaging shall be individually inspected pursuant to written procedures. Immediately after packaging, and prior to dispensing or distribution, each packaged product shall be inspected for evidence of particulates or other foreign matter, container-closure integrity, precipitation, cloudiness, and any other apparent visual defect. Defective product shall be segregated from other product and shall not be dispensed or distributed.
- h. Storage and Beyond-Use Dating (BUD).
  - (1) All sterile packaged products shall be stored in accordance to the corresponding manufacturers' storage directions. The temperature of drug storage areas of sterile packaged products shall be monitored and recorded daily, either manually or electronically. Temperature records shall be maintained and available for inspection at the outlet for at least two years.
  - (2) Finished packaged sterile products that are not immediately dispensed, distributed, or administered shall be refrigerated or frozen unless their chemical and physical stability are known to be adversely affected by cold or freezing temperatures.
  - (3) In the absence of sterility testing compliant with the most current United States Pharmacopeia/National Formulary Chapter 71 [Sterility Tests], the BUD of all sterile product packaging shall not exceed the following:
    - (a) Low risk packaged sterile products with greater than 12-hour BUD:

Room temperature:	No more than 48 hours
Refrigerated temperature:	No more than 14 days
Frozen:	No more than 45 days
    - (b) Low risk packaged sterile products with 12-hour or less BUD:

Room temperature:	No more than 12 hours
Refrigerated temperature:	No more than 12 hours
Frozen:	Not applicable
- i. If, after tests or observations, a sterile packaged product is believed to be defective in any way, the outlet shall immediately recall any product dispensed or distributed. Any product remaining in the outlet shall be immediately quarantined and shall not be dispensed or distributed. Recall records shall be maintained and available for inspection at the outlet for at least two years and shall include at least the following:
  - (1) Product name, strength, dosage form;
  - (2) Reason for recall;
  - (3) Amount of product packaged;

- (4) Date packaged; and
- (5) Amount of product dispensed and/or distributed.

**3.03.00 Customized Patient Medication Packages (Med Paks).**

**3.03.10** When a unit dose, single dose, unit of issue or customized patient medication package is dispensed pursuant to an order, the prescription shall comply with all requirements of CRS12-42.5-121(2). Container requirements of a prescription for the purpose of unit dose systems may be broadened to include trays, bins, carts and locked cabinets or drawers. Additionally, a customized patient medication package shall comply with all the following requirements:

**a. Labeling**

The patient med pak shall bear a label stating

- (1) The name of the patient;
- (2) A serial number for the patient med pak itself and a separate identifying serial number for each of the prescription orders for each of the drug products contained therein;
- (3) The name, strength, and total quantity of each drug product contained therein;
- (4) The directions for use and cautionary statements, if any, contained in the prescription order for each drug product therein;
- (5) Any storage instructions or cautionary statements;
- (6) The name of the prescriber of each drug product therein;
- (7) The date of preparation of the patient med pak, the expiration date which may not exceed 60 days from the date of preparation;
- (8) The name, address, and telephone number of the dispenser.

**b. Record Keeping.**

- (1) Patient name and address;
- (2) The serial number of the prescription order for each drug in product contained therein;
- (3) Descriptive information sufficient to allow subsequent preparation of an identical patient med pak;
- (4) Date of preparation of the patient med pak and the expiration date assigned;
- (5) Any special labeling instructions; and
- (6) The identity of the pharmacist who prepared the patient med pak.

**c. Packaging**

- (1) Each container shall meet or exceed United States Pharmacopoeia standards.

- (2) Each container shall be either not reclosable or so designed as to show evidence of having been opened.

3.03.20 It shall not be considered redispensing for a prescription drug outlet to modify a customized medication package which it has previously dispensed if the following criteria are met:

- a. The med pak is modified for the same patient for which it was originally dispensed.
- b. The med pak is returned to the prescription drug outlet from which it was originally dispensed.
- c. Only discontinued medication may be removed from the med pak. Additional medications may not be added.
- d. The medications removed from the med pak are destroyed. They may not be redispensed.
- e. The med pak is assigned a new serial number.
- f. The labeling of the med pak is modified to comply with 3.03.10(a). The expiration date affixed to the label prior to modification must be retained.
- g. Records are maintained for the modified med pak which comply with rule 3.03.10(b).

3.04.00 Colorado Cancer Drug Repository Program.

3.04.10 A prescription drug outlet may accept donations of non-controlled cancer drugs and medical devices (prescription drugs and devices that are used to treat cancer or the side effects of cancer) from cancer patients or the cancer patient's family, provided the drugs or devices meet the following requirements:

- a. The drug or device is in the original, unopened, sealed, and tamper-evident unit-dose packaging, or if in a single unit dose package, the single unit dose package is unopened;
- b. The drug or device is not expired;
- c. The drug or device is not adulterated or misbranded as determined by the pharmacist;
- d. The drug or device does not require refrigeration, freezing, or other special temperature requirements beyond controlled room temperature; and
- e. The drug or device does not require patient registration with the drug manufacturer prior to dispensing.

3.04.20 The drugs and devices shall be stored in the compounding/dispensing area under manufacturer's recommended storage conditions. The drugs and devices shall be stored separately from the other drug stock.

3.04.30 Dispensing/distribution of repository drugs/devices.

3.04.31 The prescription drug outlet may distribute the donated drugs or devices to the following:

- a. A medical clinic, which is defined as a community health clinic required to be licensed or certified by the Colorado Department of Public Health and Environment. Such clinic must be registered with the Board as an other outlet; or

- b. A registered prescription drug outlet.

3.04.32 The prescription drug outlet may dispense the drugs or devices to eligible patients based on a valid order from a practitioner. The prescription drug outlet shall establish criteria for individuals to receive donated cancer drugs or devices. The pharmacy may only charge a handling fee for such dispensing. This fee shall be determined by the State Board of Health.

3.04.40 Recordkeeping.

3.04.41 The prescription drug outlet shall retain separate records detailing the receipt and distribution/dispensing of repository drugs and devices.

3.04.42 Records of receipt shall include at least the following:

- a. Name and address of person donating the drug or device;
- b. Drug or device name and strength;
- c. Manufacturer of drug or device;
- d. Manufacturer's lot number;
- e. Drug or device expiration date, if applicable;
- f. Date received; and
- g. Quantity received.

3.04.43 Records of distribution shall include at least the following:

- a. Name and address of medical clinic or prescription drug outlet;
- b. Drug or device name;
- c. Drug strength;
- d. Dosage form, if appropriate;
- e. Quantity distributed;
- f. Identity of manufacturer of drug or device;
- g. Manufacturer's lot number;
- h. Drug or device expiration date, if applicable;
- i. Date of distribution; and
- j. Name and address of distributing pharmacy.

3.04.44 Records of dispensing shall include at least the following:

- a. Patient name;
- b. Prescription number;

- c. Drug or device name and drug strength;
- d. Quantity dispensed;
- e. Practitioner's name;
- f. Date dispensed;
- g. Identity of drug or device manufacturer; and
- h. Drug or device lot number.

**4.00.00 LICENSING.**

**4.00.10 Definitions**

- a. "Academic examination" is the North American Pharmacist Licensure Examination.
- b. "Board-approved foreign pharmacy graduate certification" means the Foreign Pharmacy Graduate Equivalency Certification.
- c. "Board-approved jurisprudence examination" means the Colorado-specific Multistate Pharmacy Jurisprudence Examination.
- d. "Board-approved school or college of pharmacy" is a professional degree program of a school or college of pharmacy that has an accredited or preaccredited status from the Accreditation Council for Pharmacy Education ("ACPE").
- e. "Board-designated clearinghouse for license transfer" means the National Association of Boards of Pharmacy Clearinghouse operated by the National Association of Boards of Pharmacy.
- f. "Disenrollment" means the current status of a pharmacy student who no longer possesses the right or capacity to complete the curriculum in the allotted time as set forth in the policies of the corresponding Board-approved school or college of pharmacy.
- g. "Enrollment" means the current status of a pharmacy student who possesses the right or capacity to complete the curriculum in the allotted time as set forth in the policies of the corresponding Board-approved school or college of pharmacy.
- h. "Intern" means a person who is:
  - (1) Enrolled in a professional degree program of a Board-approved school or college of pharmacy, licensed by the Board to engage in the practice of pharmacy, and satisfactorily progressing toward meeting the requirements for licensure as a pharmacist;
  - (2) A graduate of a Board-approved school or college of pharmacy or a graduate who has established education equivalency by obtaining a Board-approved foreign pharmacy graduate certification and who is currently licensed by the Board for the purpose of obtaining practical experience as a requirement for licensure as a pharmacist; or
  - (3) A qualified pharmacist applicant awaiting examination for licensure as a pharmacist or meeting Board requirements for pharmacist licensure.

- i. "License transfer or endorsement" is the licensing of an individual who is licensed as a pharmacist by examination in another state and whose license in that state is in good standing.
- j. For the purposes of this rule 4.00.00, "manufacturer" means a manufacturer of prescription drugs which is registered by the Board.
- k. "Pharmacist" means an individual licensed by this state to engage in the practice of pharmacy.
- l. "Regulated individual" means any of the following individuals holding an active, unrestricted license, registration, or certification from the Colorado Department of Regulatory Agencies:
  - (1) Clinical Social Worker;
  - (2) Dentist;
  - (3) Occupational Therapist;
  - (4) Optometrist;
  - (5) Physical Therapist;
  - (6) Physician;
  - (7) Physician Assistant;
  - (8) Podiatrist;
  - (9) Psychologist;
  - (10) Registered Nurse or Advanced Practice Nurse;
  - (11) Respiratory Therapist; and
  - (12) Veterinarian.
- m. "Score transfer" is the transfer of the academic examination score to Colorado by participation in the NAPLEX Score Transfer Program operated by the National Association of Boards of Pharmacy.

**4.00.20 Requirements for Intern Licensure include the following;**

- a) Submission of a completed application form provided by the Division of Professions and Occupations with the appropriate fee.
- b) Submission of one of the following:
  - 1) Proof of enrollment in a Board-approved school or college of pharmacy. A person on suspension from a Board-approved school or college of pharmacy may not be licensed as an intern. A person in good standing with a Board-approved school or college of pharmacy may be licensed as an intern.



- 2) If a graduate of a foreign school or college of pharmacy, a Foreign Pharmacy Graduate Equivalency Certification;
- 3) Proof of graduation within the prior two years from a Board-approved school or college of pharmacy. If the applicant ceased to be enrolled in a Board-approved school or college of pharmacy more than two years prior to application, the applicant shall include an explanation of "good cause" for licensure which the Board or its designee shall review and act on in the normal course of business.
- 4) If a pharmacist in another state awaiting pharmacist licensure in Colorado, verification of an active, unrestricted license in another state.

4.00.25 Requirement for Intern Reporting. An actively licensed intern shall report to the Board, in writing, within 30 days of meeting the definition of "Disenrollment" as defined in Rule 4.00.10(f) from a Board-approved school or college of pharmacy.

4.00.30 Requirements for Pharmacist License by Exam or Score Transfer include the following:

- a. Submission of a completed application form provided by the Division of Professions and Occupation with the appropriate fee.
- b. Submission of a transcript and proof of graduation from a Board-approved school or college of pharmacy or a Foreign Pharmacy Graduate Equivalency Certification.
- c. Successful passage of the academic examination and Board-approved jurisprudence examination. The passing scores for these examinations are set by the examining entity. If an applicant passes only one of the required examinations, the applicant shall be required to repeat the failed examination. If, within the previous 24 months, the applicant has not passed both required examinations, he or she shall be required to also repeat the previously passed examination. Score transfer applicants shall complete licensure within one year from the date their scores are received by the Division of Professions and Occupations.
- d. Proof of completion of 1500 intern hours completed no more than five years after graduation from a Board-approved school or college of pharmacy. If a graduate of an unapproved school or college of pharmacy, receipt of the Foreign Pharmacy Graduate Equivalency Certification. Intern hours must be obtained under one or more of the following conditions:
  - (1) Engaged in the practice of pharmacy under the direct supervision of a pharmacist.
  - (2) Directly supervised by a manufacturer as part of the curriculum of an approved school or college of pharmacy.
  - (3) Directly supervised by a regulated individual as part of the curriculum of an approved school or college of pharmacy. The scope of practice of the regulated individual must overlap with that of a pharmacist for the course of the hours supervised.
  - (4) One year of practice of pharmacy as a licensed pharmacist in another state may be accepted by the Board in lieu of the 1500 hours if the applicant has completed this year of pharmacy practice prior to taking the examination.

- e. Education, training, or service gained in military services outlined in C.R.S. 24-34-102(8.5), to be accepted and applied towards receiving a license, must be substantially equivalent, as determined by the Board, to the qualifications otherwise applicable at the time of receipt of application. It is the applicant's responsibility to provide timely and complete evidence for review and consideration. Satisfactory evidence of such education, training, or service will be assessed on a case by case basis.

**4.00.40 Requirements for License Transfer or Endorsement are as follows:**

- a. Submission of a completed application and fee to the Board designated clearinghouse for license transfer.
- b. Submission of a completed application form provided by the Division of Professions and Occupations with the appropriate fee.
- c. Successful passage of the Board-approved jurisprudence examination. The passing score is set by the examining entity.
- d. Applicants for license transfer must have been licensed as a pharmacist for at least one year in another state or have served an Internship meeting the Colorado requirements at the time of original licensure.
- e. An applicant for license transfer shall apply for license transfer using a license issued by examination in another state. Such license shall be active, current, and in good standing. If the applicant holds pharmacist licenses in multiple states, all licenses must be in good standing. For the purposes of these rules, "good standing" means that the applicant is not currently subject to active disciplinary actions in any state.

**4.03.00 Reinstatement or Reactivation of Pharmacist License.**

- a. If the license has been inactive or expired for over 24 months, a person wishing to reinstate or reactivate such license shall do the following:
  - (1) Submit the appropriate application with the required fee;
  - (2) Submit 1 hour of continuing education for each month such license was inactive or expired. Twenty-four (24) of these hours shall have been completed in the 24 months prior to application for reinstatement or reactivation; and
  - (3) Take and pass the approved jurisprudence examination. The passing score shall be set by the examining entity.
- b. If the license has been expired or inactive for less than 24 months, a person wishing to reinstate or reactivate such license shall do the following:
  - (1) Submit the appropriate application with the required fee; and
  - (2) Submit 24 hours of continuing education completed within the 24 months prior to application.

**4.05.00 License Changes.**

- a. Name change. A licensee shall report a name change and provide appropriate supporting documentation within 30 days of such change. If the licensee wishes to obtain a new wall license with the new name, the licensee shall pay the requisite fee.

- b. Change of residential and electronic mail address. All pharmacists and interns shall notify the Board in writing within 30 days of any change of residential or electronic mail address.
- c. Change of manager. A pharmacist shall immediately notify the Board in writing of the date he ceases to be the pharmacist manager of a prescription drug outlet.

4.06.00 Identification of Licensee. A pharmacist, pharmacy intern, pharmacy technician, pharmacy clerk, store manager, or assistant store manager shall at all times while on duty within a prescription drug outlet wear a badge which is visible to the patient and which shall state at least the title accurately reflecting a person's role in the outlet such as Pharmacist, Pharmacy Intern, Pharmacy Technician, Pharmacy Clerk, Store Manager, or Assistant Store Manager.

#### **5.00.00 OUTLETS.**

5.00.01 Definitions. The following words and terms shall have the following meanings, unless the context clearly indicates otherwise.

- a. Compounding / Dispensing Area: means any area in a prescription drug outlet where "compounding / dispensing" is performed.
- b. In-State Prescription Drug Outlet: means any prescription drug outlet located within Colorado that is registered pursuant to CRS Title 12, Article 42.5, where prescriptions are compounded and dispensed.
- c. Non-Resident Prescription Drug Outlet: means any pharmacy outlet located outside this state that is registered pursuant to CRS Title 12, Article 42.5, which ships, mails, or delivers, in any manner, drugs or devices into this state pursuant to a prescription order.

5.00.10 Registration. The applicant for registration shall obtain the appropriate form as approved by the Board to register an outlet. In the case of an application for a new in-state or non-resident prescription drug outlet, for a transfer of ownership of an in-state or non-resident prescription drug outlet, or for the relocation of an in-state or non-resident prescription drug outlet, the applicant shall submit such additional documentation as the Board may require.

5.00.15 Registration for Nonresident Prescription Drug Outlet. An applicant for a new nonresident prescription drug outlet registration shall submit the following:

- a. The current application with required fee;
- b. A verification of the current pharmacy license or registration issued by the applicant's resident state board of pharmacy;
- c. A copy of the most recent report detailing an inspection of the nonresident prescription drug outlet by either its resident state board of pharmacy or the National Association of Boards of Pharmacy's Verified Pharmacy Program dated within the previous two (2) years of submission of the application; and
- d. An affidavit attesting that the nonresident prescription drug outlet shall not ship compounded or other prescription drugs into the State of Colorado without a prescription order for a specific patient, except as provided pursuant to Rule 21.00.20.

5.00.20 Applications. The Board, or its agent, may require any applicant or pharmacist manager of an outlet to meet with the Board, or its agent, before the Board takes action on any registration.

5.00.30 No two registered in-state or non-resident prescription drug outlets may occupy the same physical space. If there are two (or more) registrants co-located within the same building or at the same address, each must have its own area, separated by floor to ceiling walls, and separate entrances.

5.00.40 Transfer of Ownership. Application to transfer registration of an in-state or non-resident prescription drug outlet shall be submitted to the Board as provided in CRS 12-42.5-116, immediately upon the transfer of ownership. A transfer of ownership shall be deemed to have occurred:

- a. In the event the in-state or non-resident prescription drug outlet is owned by a corporation, upon sale or transfer of 20 percent or more of the shares of said corporation to a single individual or entity.
- b. In the event the in-state or non-resident prescription drug outlet is owned by a partnership, upon sale or transfer of 20 percent or more of any ownership interest.
- c. In the event the in-state or non-resident prescription drug outlet is owned by a limited liability company (LLC), upon sale or transfer of 20 percent or more of the membership interests.
- d. Upon incorporation of an existing in-state or non-resident prescription drug outlet.

5.00.50 Relocation.

- a. In the event of a relocation of an in-state or non-resident prescription drug outlet, the outlet shall submit an application provided by the board along with the prescribed fee at least 30 days prior to the effective date of relocation.
- b. The registration of a non-resident prescription drug outlet shall become void and shall be cancelled if the non-resident prescription drug outlet relocates to a state other than that which appears on its registration. In the event the non-resident prescription drug outlet wishes to continue shipping prescriptions into Colorado, it must apply for and receive a new Colorado registration prior to such shipment.

5.00.55 Reinstatement of an In-State or Non-Resident Prescription Drug Outlet Registration.

- a. In-state Prescription Drug Outlet. If a registration has expired, a facility seeking to reinstate such registration shall submit the following:
  - (1) The current reinstatement application with the required fee;
  - (2) If the owner of the in-state prescription drug outlet is a corporation, submit either a copy of the articles of incorporation as they were filed with the Colorado Secretary of State or a Certificate of Good Standing issued by the Colorado Secretary of State;
  - (3) A letter stating whether the corporation is public or private as follows:
    - (A) If the corporation is a public corporation, submit a list of all stockholders owning five percent or more of the stock; or
    - (B) If the corporation is a private corporation, submit a list of all stockholders;

- (4) An accurate drawn-to-scale floor plan of the prescription drug outlet's compounding / dispensing area detailing all counters, bays, sinks, refrigerators and, if applicable, sterile and non-sterile compounding hoods;
  - (5) A completed, dated and signed minimum equipment self-inspection form as provided with the reinstatement application; and
  - (6) A statement, signed by the pharmacist manager, stating whether or not greater than ten percent of the business is owned by a person or persons authorized by law to prescribe drugs.
- b. Non-resident Prescription Drug Outlet. If a registration has expired, a facility seeking to reinstate such registration shall submit the following:
  - (1) The current reinstatement application with the required fee;
  - (2) A verification of the current pharmacy license or registration issued by the resident state board of pharmacy;
  - (3) If the registration has expired for more than two years, a copy of the most recent report detailing an inspection of the non-resident prescription drug outlet by its resident state board of pharmacy dated within 5 years of submission of the reinstatement application.

**5.00.60 Closure.**

- a. Closure shall mean the permanent cessation of the practice of pharmacy in any in state or non-resident prescription drug outlet. For in-state prescription drug outlets, closure shall also be deemed to have occurred if the compounding/dispensing area is not open for business the minimum hours specified in 5.01.40(a).
- b. Upon the closure of any in-state or non-resident prescription drug outlet, it shall be the responsibility of the last pharmacist manager of record to remove the prescriptions and/or chart orders to another prescription drug outlet where patrons and/or practitioners are afforded reasonable access to a pharmacist's interpretation of such orders. Such relocation of records shall be made within 72 hours after closure. The pharmacist manager shall submit a notice, on a form and manner approved by the Board, detailing the closure of the prescription drug outlet or nonresident prescription drug outlet within 72 hours after closure. If the last pharmacist manager of record fails to relocate the records as required herein, the Board may direct the removal of the records to a suitable location. The last pharmacist manager of record shall make a reasonable effort to inform patrons of the prescription drug outlet of the location of the records.
- c. The Board on request shall provide the owner of any prescription drug outlet an instruction sheet applicable to the transaction prior to closure, or conducting bankruptcy proceedings, or transferring or selling the prescription drug inventory.

**5.00.70 Change in Pharmacist manager.**

- a. An in-state and non-resident prescription drug outlet shall be under the direct charge of a pharmacist manager. A proprietor who is not a pharmacist shall comply with this requirement and shall provide a manager who is a pharmacist.

- b. The registration of any in-state and non-resident prescription drug outlet shall become void if the pharmacist manager in whose name the registration was issued ceases to be engaged as the manager, and the owner shall close the outlet unless such owner has employed a pharmacist manager and, within thirty days after termination of the former manager's employment, has made application to transfer the registration to the new pharmacist manager and has paid the transfer fee therefor.

5.00.80 Disclosure. Any Board registered non-resident prescription drug outlet shall disclose to the Board, in writing, the location, names, and titles of all principal entity officers and all pharmacists who are dispensing drugs to residents of this state on an annual basis and within thirty days after any change of office, officer or pharmacist.

5.01.00 Prescription Drug Outlets (In-State).

5.01.10 Controlled Substance Inventory.

- a. Upon the change of pharmacist manager of a prescription drug outlet, an inventory of all controlled substances shall be taken within seventy-two hours, by the new pharmacist manager or the new pharmacist manager's designee. The inventory shall be taken either as of the opening or as of the close of business activity on the inventory date and such time and date taken shall be entered on the inventory record.
- b. Upon the transfer of ownership of a prescription drug outlet, an inventory of all controlled substances shall be taken by the pharmacist manager or the pharmacist manager's designee. The inventory shall be taken either as of the opening or as of the close of business activity on the inventory date and such time and date taken shall be entered on the inventory record.

5.01.20 Compounding/Dispensing Area (In-State)

5.01.21 In the event a transfer of ownership of a prescription drug outlet occurs, and the principal compounding/dispensing area or any satellite compounding/dispensing area does not meet the physical requirements of this rule, the transfer of the registration may be approved, provided that compliance with such requirements shall be accomplished within six months of the approval of the transfer of the registration or by the next prescription drug outlet registration renewal date, whichever time is greater.

5.01.31 Within every prescription drug outlet as defined in CRS 12-42.5-102(35), there shall be one area designated as the principal compounding/dispensing area. In addition to the principal compounding/dispensing area there may be satellite compounding/dispensing areas and drug storage areas ("satellites") which are located at the same location as the principal compounding/dispensing area. The principal compounding/dispensing area and any satellite shall comply with the following conditions:

- a. The principal compounding/dispensing area shall not be less than 225 continuous square feet, except that prescription drug outlets registered by the Board prior to the effective date of this regulation that do not meet this space requirement are hereby exempted from such requirement. However, any new prescription drug outlet shall comply with this requirement prior to the granting of the initial registration. Any existing prescription drug outlet which is being remodeled or is being moved from one location to another, whether or not there is a change of address, shall submit documentation required by the Board prior to remodeling or relocation.

- b. All compounding/dispensing satellites and any drug storage satellites in excess of the two permitted in subsection c below that are at the same location as the principal compounding/dispensing area must not be less than 100 continuous square feet and must be approved by the Board prior to use for compounding/dispensing.
- c. In addition to the satellite areas permitted in the previous paragraph, up to two satellites at the same location may be used solely for storage of prescription drugs and controlled substances. Such drug storage satellites must possess square footage commensurate for the safe storage and removal of drugs within the affected satellites and approved by the Board prior to use.
- d. Any room included within or adjacent to the principal compounding / dispensing area that is separated from the principal compounding / dispensing area by a door must meet the following:
  - (1) The prescription drug outlet shall submit documentation required by the board to remodel the principal compounding / dispensing area prior to the utilizing the room or rooms for the purposes of compounding and dispensing or for the storage of prescription drugs and controlled substance stocks;
  - (2) The door must have a conspicuously displayed sign attached to it, and facing the principal compounding / dispensing area, that states "This room is part of the Board-approved designated principal compounding / dispensing area";
  - (3) If a locked or otherwise secured door is used to separate parts of the compounding / dispensing area, it shall be unlocked immediately upon the request of the Board or of its inspectors and be available for inspection.
- e. All compounding/dispensing areas and satellites shall be well-lighted and well-ventilated with clean and sanitary surroundings devoted primarily to compounding/dispensing or drug storage. These areas shall provide necessary protection for drugs, chemicals and devices from deterioration due to light, heat or evaporation and shall be arranged to protect all prescription drugs and devices from pilferage or other unauthorized removal. No areas shall be subject to any condition likely to lead to errors.
- f. In every prescription drug outlet and in every satellite where compounding or dispensing is physically occurring, there shall be a minimum of 12 continuous square feet of free and clear counter space, and a minimum of 6 continuous square feet of free and clear counter space for each person engaged in compounding/dispensing as defined. These counters and surfaces shall be kept free and clear at all times for the purpose of compounding/dispensing. Any computer workstation or other equipment for the preparation of prescription labels and/or storage and retrieval of records shall be in addition to the minimum free compounding/dispensing area.
  - (1) The free floor space behind all compounding/dispensing counters or work surfaces shall be not less than 30 inches in width;
  - (2) The free floor space between shelving rows shall be not less than 24 inches; and
  - (3) There shall be sufficient shelf, drawer and/or cabinet space for proper storage of prescription drugs and devices.

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- g. In every satellite used for the sole purpose of storing prescription drugs or controlled substances, there shall be:
- (1) At least 24 inches of free floor space between shelving rows; and
  - (2) At least 30 inches of free floor space behind any counters, if counters are available.
- h. In the principal compounding/dispensing area there shall be a sink, equipped with running hot and cold water, which is attached to an approved drain, waste and vent system, or to a portable enclosed tank which is emptied as frequently as necessary. Each satellite area shall also be so equipped if appropriate to the compounding/dispensing activities which are or will be performed therein.
- i. The prescription drug outlet shall have all the technical equipment necessary for the appropriate compounding and dispensing it conducts.
- j. If refrigerated drugs are stored in the principal compounding/dispensing area or in any satellite, there shall be a refrigerator, dedicated to storing only drugs, meeting the compendia requirements and with an accurate thermometer in the refrigerator. The temperature of which shall be maintained between two and eight degrees Celsius (2 and 8 degrees C.) or thirty-six and forty-six degrees Fahrenheit (36 and 46 degrees F.). The temperature shall be electronically monitored each calendar day. Records detailing instances in which temperatures fall outside the aforementioned range requirement, for any period of time, shall be maintained at the prescription drug outlet and shall be made readily available for inspection upon request by the Board or its representatives for a period of at least two years preceding the request. Such records shall include the duration of time the temperature fell outside the aforementioned range requirement, based on the best available data, and measures taken by the outlet as a result of the temperature falling outside the aforementioned range requirement.
- k. If frozen drugs are stored in the principal compounding/dispensing area or in any satellite, there shall be a freezer, dedicated to storing only drugs, meeting the compendia requirements and with an accurate thermometer in the freezer. The temperature of which shall be maintained between twenty-five degrees below zero and ten degrees below zero Celsius (– 25 and – 10 degrees C.) or thirteen degrees below zero and fourteen degrees Fahrenheit (– 13 and 14 degrees F.). The temperature shall be electronically monitored each calendar day
- Records detailing instances in which temperatures fall outside the aforementioned range requirement, for any period of time, shall be maintained at the prescription drug outlet and shall be made readily available for inspection upon request by the Board or its representatives for a period of at least two years preceding the request. Such records shall include the duration of time the temperature fell outside the aforementioned range requirement, based on the best available data, and measures taken by the outlet as a result of the temperature falling outside the aforementioned range requirement.
- l. There shall be a professional reference library available in the prescription drug outlet. If an electronic library is provided, workstations must be provided in a compounding/dispensing area and must be readily available for use by staff, interns and Board personnel. This library shall contain current copies of the following:
- (1) A CRS Title 12, Article 42.5; the Pharmacists, Pharmacy Businesses, and Pharmaceuticals Act;
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- (2) CRS Title 18, Article 18, the Uniform Controlled Substances Act of 1992;
  - (3) Board rules;
  - (4) 21 Code of Federal Regulations ("CFR") Part 1300 to End containing Drug Enforcement Administration rules relating to controlled substances;
  - (5) If compounding sterile products, Guide to Parenteral Admixtures or Handbook on Injectable Drugs or other comparable references as determined by the pharmacist manager;
  - (6) If compounding cytotoxic products, Technical Manual Section VI: Chapter 2, Controlling Occupational Exposure to Hazardous Drugs or ASHP Technical Assistance Bulletin on Handling Cytotoxic and Hazardous Drugs; and
  - (7) Any other references that the pharmacist manager of the prescription drug outlet may deem necessary.
- m. If telephone prescription orders are accepted while the compounding/dispensing area is closed, a voice recording device shall be provided to receive them, and they shall be played back by the pharmacist or intern.
  - n. Written prescription orders and refill requests for prescription orders may be delivered to the prescription drug outlet while the compounding/dispensing areas are closed, provided a slot or drop box is provided for the prescription order or prescription order refill requests.
  - o. All prescription drug outlets shall maintain an adequate inventory of prescription drugs and shall offer adequate pharmaceutical service to the public they normally serve.
  - p. Every prescription drug outlet shall display in the principal compounding/dispensing area the report of the most recent inspection conducted by the Board or a photocopy of the most recent self-inspection performed by the pharmacist manager using the form provided by the Board, whichever is more recent, and have readily available documents sent or provided by the Board to clarify or assist in the legal operation of the prescription drug outlet.
  - q. No person other than a pharmacist or intern employed by the prescription drug outlet shall be permitted in the compounding/dispensing area without the consent of the pharmacist in charge of the compounding/dispensing area.
- 5.01.33 The use of any tobacco product in any compounding/dispensing area is hereby prohibited. However, this rule shall not apply to the compounding, dispensing or use of a drug which has been derived from a tobacco product and which is being used as an adjunct to a smoking cessation program.
- 5.01.34 Delivery and Temporary Storage of Prescriptions. Upon the request of a patient or an agent of the patient and with the approval of the pharmacist on duty a prescription may be delivered or temporarily stored outside the confines of a compounding/dispensing area. The pharmacist manager of the prescription drug outlet shall determine or approve procedures for the storage and security of, the access to, the confidentiality of, and the counseling regarding, prescriptions, including record keeping.

5.01.40 Minimum Hours of Operation.

- a. The principal compounding/dispensing area of a prescription drug outlet shall be open for normal business a minimum of two designated days per week (Monday through Sunday) and at least four continuous hours on each such designated day.
- b. In the event that the principal compounding/dispensing area is open less than 32 hours per week, the pharmacist manager shall submit to the Board a written statement of the designated days and hours when the principal compounding/dispensing area will be open for business, and this statement shall be submitted at least 30 days prior to the date on which the hours of operation will be less than 32 hours per week.

5.01.50 Security. In every prescription drug outlet, all compounding/dispensing areas shall comply with this regulation.

- a. When any compounding/dispensing area of a prescription drug outlet is occupied by any employee, a pharmacist must be physically present within the same building of the prescription drug outlet.
- b. In the event a pharmacist is within the building but absent from a compounding/dispensing area, it is the responsibility of the pharmacist to ensure the proper safeguard of all drugs.
- c. If a compounding/dispensing area is continually attended by a pharmacist when other people are in the building, the compounding/dispensing area need not be enclosed. However, if other people are in the building when there is not a pharmacist present, every compounding/dispensing area must be enclosed by a barrier as specified in paragraph e below.
- d. If more than one prescription drug outlet is located within the same building, a pharmacist shall not operate more than one outlet at the same time. If a pharmacist physically leaves one outlet for the purpose of entering into another outlet within the same building, any outlet not being physically attended to by a pharmacist shall be enclosed by a barrier as specified in paragraph e below and a non-pharmacist shall not remain inside the enclosed outlet during that time.
- e. A prescription drug outlet constituting part of a large establishment may be closed while the balance of the establishment is open for business, provided every compounding/dispensing area is enclosed with a secure floor-to-ceiling physical barrier, which shall be a divider or secure total enclosure, in which any openings shall not be large enough to permit removal of items from the compounding/dispensing area. The barrier must be of weight and strength sufficient to prevent it from being readily lifted, removed, penetrated or bent.
- f. All entrances to every compounding/dispensing area shall be secured from unauthorized entry when the pharmacist leaves the building. No one other than a pharmacist shall be permitted to enter any compounding /dispensing area except in extreme emergencies, which shall be defined as a threat to property, public disaster or other catastrophe whereby the public is better served by overlooking the security restrictions of drugs and devices. If any compounding/dispensing area is opened in the absence of a pharmacist or left unsecured from unauthorized entry when the pharmacist leaves the building, the pharmacist manager shall notify the Board in writing within ten days of the discovery of the occurrence. This written notice shall state:

- (1) The name of the person authorizing the opening of the compounding/dispensing area if known, or the name of the pharmacist responsible for securing the compounding/dispensing area from unauthorized entry;
  - (2) The name of the person opening the compounding/dispensing area if known; and
  - (3) A description of the situation requiring opening of the compounding/dispensing area including the date and time of the opening.
- g. While the compounding/dispensing area is closed and the rest of the establishment is open, a person on duty in the establishment shall be able to contact a pharmacist in case of emergency.
- h. The hours of business of the compounding/dispensing area shall be submitted to the Board in writing.
- i. No prescription drug outlet shall avail itself of the privileges of this rule until the barrier system and other requirements have been acknowledged, subject to final approval by the Board.
- j. This paragraph applies only to the compounding/dispensing areas of a hospital which operates a prescription drug outlet pursuant to a certificate of compliance; or which operates a registered prescription drug outlet on the premises of the hospital for the primary purpose of providing pharmaceutical services to the hospital's in-patients; or permits a registered prescription drug outlet to be operated on the premises of the hospital by another business entity for the primary purpose of providing pharmaceutical service to the hospital's in-patients.
  - (1) In an emergency situation and when a pharmacist is not on the premises of the hospital and administration of a drug to, or use of a device by or on, an in-patient is necessary pursuant to a chart order, and such drug or device is only available from a locked compounding/dispensing area, an authorized registered nurse may enter a locked compounding/dispensing area to obtain the drug or device. In the case of a drug, only pre-labeled packages, such as unit dose or unit-of-use packages, or a pre-labeled container, may be removed from the compounding/dispensing area.
  - (2) The following information regarding the removal of such drug or device shall be consistently recorded and maintained in a retrievable document: date; time; name, strength and dosage form of drug, and/or name, and size, if applicable, of device; total quantity of drug or device removed; name and location of patient for whose use the drug or device is necessary; name of the practitioner ordering the drug or device; and the initials or signature of the nurse obtaining the drug or device. This document shall be available for inspection by the Board for a period of 2 years. Additionally, the original, duplicate or electronic or mechanical facsimile of the chart order shall be left with the above document by the nurse at the time of obtaining the drug or device.
  - (3) Any unused portion of a drug or device so removed shall be returned to the compounding/dispensing area when a pharmacist is again on the premises. Additional quantities of the drug or device shall be supplied by a pharmacist and properly recorded as required by law and rule.

**6.00.00 PHARMACEUTICAL CARE, DRUG THERAPY MANAGEMENT AND PRACTICE BY PROTOCOL.**

**6.00.10 Definitions.**

- a. "Pharmaceutical care" means the provision of drug therapy and other pharmaceutical patient care services by a pharmacist intended to achieve outcomes related to the cure or prevention of a disease, elimination or reduction of a patient's symptoms, or arresting or slowing of a disease process. In addition to the preparation, dispensing, and distribution of medications, "pharmaceutical care" may include assessment and evaluation of the patient's medication related needs and development and communication of a therapeutic plan with defined outcomes in consultation with the patient and the patient's other health care professionals to attain the desired outcome. This function includes efforts to prevent, detect, and resolve medication related problems for individual patients. "Pharmaceutical care" does not include prescriptive authority.
- b. For the purpose of this Board Rule 6.00.00, a "prescriber" means a physician who is actively and unconditionally licensed by the Colorado Medical Board or an advanced practice registered nurse with prescriptive authority who is actively and unconditionally licensed by the Colorado State Board of Nursing.
- c. Drug therapy management means the review and evaluation of drug therapy regimens for patients undertaken by a pharmacist in order to provide drug therapy, monitor progress, and modify drug therapy. Drug therapy management may only be undertaken pursuant to an initial diagnosis made by a prescriber, a valid order for the therapy, and a written agreement, which delineates proper protocols, to be used and the type of interaction that must occur between the pharmacist and the prescriber. Therapeutic interchange programs in inpatient and group model integrated closed HMO settings that are approved by medical staff committees are not considered drug therapy management for purposes of these rules.
- d. Drug therapy management may include:
  - 1. Collecting and reviewing patient drug histories;
  - 2. Obtaining and checking vital signs;
  - 3. Ordering and evaluating the results of laboratory tests directly, related to management of the drug therapy when performed in compliance with the protocol ordered by the prescriber;
  - 4. Modifying drug therapy, when appropriate, in compliance with the protocol ordered by the prescriber; and
  - 5. Implementing the drug therapy plan agreed upon between the prescriber and the pharmacist, using protocols and managing the therapy according to those protocols.

- e. Protocol means a specific written plan for a course of medical treatment containing a written set of specific directions created by the prescriber, groups of prescribers, hospital medical committee, or pharmacy and therapeutics committee.
  - 1. Protocols must describe the nature and scope of drug therapy management appropriate for certain conditions or diagnoses, and include specific directions for the drug to be used, the specified dosage regimen, dosage forms or route of administration which are authorized. Protocols must include clear criteria and specific directions pharmacists are to follow when implementing and monitoring drug therapy. If the protocol includes ordering and evaluating laboratory tests, the protocol must provide precise instruction as to what tests are to be ordered, the criteria for ordering the tests, how the tests are to be interpreted, and what action the pharmacist is to take dependent upon the test results. If the protocol includes modifying drug therapy, the protocol must provide precise instruction as to the criteria dictating a change, and exactly how the therapy is to be changed.
  - 2. Protocols without specific directions regarding patient treatment or those that are nonspecific, vague, or rely on discretion without definition, are insufficient and may not be used in drug therapy management by the prescriber or the pharmacist.
  - 3. Protocols must also include specific instructions for responding to acute allergic or other adverse reactions. The protocols shall be signed and dated by the authorizing prescriber or chairperson of the authorizing group or committee.
  - 4. Evidence based protocols. Protocols used by prescribers and pharmacists engaging in drug therapy management must demonstrate a plan of treatment that constitutes evidence-based medicine. This means that the plan of treatment must be guided by or based on current, objective, supportive scientific evidence as published in scientific literature rather than anecdotal observations. Through the use of such protocols, drug therapy management must provide care that meets the standard of care in all applicable professions.
  - 5. The protocols shall be signed and dated by the authorizing prescriber or chairperson of the authorizing group or committee.
- f. Agreement means a written agreement between a Colorado licensed pharmacist and a Colorado licensed prescriber, or a group of Colorado licensed pharmacists and a group of Colorado licensed prescribers that sets forth the specific information required to assure the competent practice of pharmacy in an integrated health care fashion. Either party may withdraw from the agreement at any time.

**6.00.20 Drug therapy management requirements for all practice settings.**

- a. Drug therapy management may only be conducted by a pharmacist upon the presentation of a valid order for a specific, individual patient from that patient's prescriber. The order must specify the protocol to be used, and the protocol must either accompany the order, or otherwise be provided to the pharmacist in advance of starting drug therapy management.
- b. The pharmacist must ensure that the prescriber with whom the pharmacist is working is licensed in Colorado, in good standing, and the protocols used are within the scope of the prescriber's current practice.

- c. Prior to initiation of drug therapy management in any setting, the pharmacist or institution must inform the patient that he/she may refuse to participate in drug therapy management. Inpatient or group model integrated closed HMO settings may use the patient's signature on the institution's general consent to treat as the patient's indication to participate in drug therapy management.
- d. At a minimum the written agreement for carrying out drug therapy management between prescribers and pharmacists shall be reviewed annually, and revised, if necessary.
- e. Pharmacists may perform by protocol all aspects of drug therapy management referenced in 6.00.10 c and d, provided the protocol complies with 6.00.10e, and the pharmacists performing these functions are qualified as set forth in section 6.00.30 and are working pursuant to a written agreement with an appropriately qualified prescriber.
- f. Filing requirements.
  - 1. Pharmacists engaging in drug therapy management must maintain a current copy of the written agreement between the prescriber and the pharmacist at the location where drug therapy management is occurring. Pharmacists conducting such therapy in inpatient settings or group model integrated closed HMO's shall maintain a current copy of the general authorization plan as required by 6.00.40 at the location where drug therapy management is occurring. Upon request by the Board or its inspectors such written agreements and general authorization plans shall be submitted to the Board.
  - 2. Pharmacists practicing drug therapy management must also provide the Board documentation of their successful completion of all qualification requirements as set forth below in 6.00.30 upon request. Copies of pharmacy degrees are not required. Copies of completion of residency or other educational programs or certifications must be on file in the location of practice. Attestations from the supervising pharmacist or prescriber for clinical practice must be on file.
  - 3. Pharmacists practicing drug therapy management must have a copy of the pertinent protocols at the location at which they are practicing. Upon request by Board inspectors, pharmacists must produce the scientific literature upon which their protocols are derived.

#### 6.00.30 Pharmacist Qualifications.

Any pharmacist engaged in drug therapy management shall meet the following qualifications:

- a. Have and maintain an unrestricted license in good standing to practice pharmacy in Colorado; and
- b. Meet one of the following qualifications:
  - (1) Proof of completion of a pharmacy residency accredited by the American Society of Health Systems Pharmacists or the American Pharmacists Association in the specialty being practiced; or
  - (2) Proof of completion of one (1) year of practice experience in pharmacotherapy, and 40 hours of onsite supervised clinical practice and training in each area in which the pharmacist is choosing to practice; or

- (3) Completion of a certificate program accredited by the Accreditation Council for Pharmacy Education in each area of practice, and 40 hours of on-site supervised clinical practice and training in each area in which the pharmacist is choosing to practice; or
- (4) Completion of at least 40 hours of ACPE approved continuing education regarding clinical practice and 40 hours of onsite supervised clinical practice and training in the area in which the pharmacist is choosing to practice; or
- (5) Current Board specialty certification from the Board of Pharmaceutical Specialties, current certification from the National Institute for Standards in Pharmacist Credentialing, or current certification from the Commission for Certification in Geriatric Pharmacy. Such credentials must be in the area of pharmacy practice undertaken in the drug therapy management; or
- (6) In an inpatient or group model integrated closed HMO setting, all of the following criteria shall be met in order to practice drug therapy management:
  - (a) Forty (40) hours of onsite supervised clinical practice and training in the area(s) in which the pharmacist is choosing to practice;
  - (b) Protocols must be approved by the health-system's medical committee, or pharmacy and therapeutics committee; and
  - (c) Documented competency of each area of practice in which the pharmacist is choosing to practice shall be maintained on site.
- c. Licensed Colorado pharmacists practicing drug therapy management prior to August 1, 2005, must attest and certify that they were provided clinical training, experience, and oversight practicing in the disease state(s) that they work in, and the physician with whom they are currently practicing must attest that they are practicing to the standard of care required for management of the specific disease. Such attestations must be on file at the site of practice. Documentation of their employment dates must be on file as proof of practice prior to August 2, 2005.

**6.00.40 Drug Therapy Management in Inpatient and Group Model Integrated Closed HMO Settings.**

- a. Pharmacists engaging in drug therapy management in inpatient and group model integrated closed HMO settings must conduct activities pursuant to a valid order and must follow the protocols set forth by the hospital medical committee, or pharmacy and therapeutics committee. They must record all of the items required in subsection c. below for each patient, or the hospital may create a general authorization plan, identifying where such information will be located, and how it will be accessed throughout the facility by participating pharmacists and prescribers. The general authorization plan serves as the pharmacist/prescriber agreement in these settings. The general authorization plan must identify which prescribers and pharmacists are authorized and have agreed to participate in the facility to engage in drug therapy management. The hospital medical committee or pharmacy & therapeutics committee serves as the authorizing agent for the organization's medical staff, identifying which prescriber groups are authorized to participate, and may restrict authorization for certain protocols to specific prescriber groups or specialties. A pharmacist engaging in drug therapy management must read, sign and date the plan and the pertinent protocols that he/she agrees to use in the cases undertaken.

- b. The pharmacist manager shall ensure that the general authorization plans for drug therapy management are on file in the prescription drug outlet. Changes to the plan must be made as they occur, including the identification of persons participating. Protocols shall be onsite where the drug therapy management takes place and revised as medically necessary.
- c. Prior to initiation of drug therapy management, the pharmacist must review the following information:
  - 1. Patient's name, gender, date of birth, height, and weight;
  - 2. Patient diagnosis or diagnoses (from physician);
  - 3. Medication history;
  - 4. Prior lab values;
  - 5. Patient vital signs;
  - 6. Patient known allergies;
  - 7. Emergency contact number.
- d. Records of all activity by the pharmacist shall be documented in the patient's chart prior to administration.
- e. Pharmacists engaging in drug therapy management shall not delegate drug therapy management activities to any other staff.

**6.00.50 Drug Therapy Management in other settings.**

- a. Every pharmacist or group of pharmacists engaged in drug therapy management in an outpatient setting must have a valid order from the patient's prescriber for each specific patient for such therapy, and must operate according to a written agreement and protocol referenced in section 6.00.10.
- b. Written agreements shall contain the following information:
  - 1. Participating pharmacist name(s);
  - 2. Participating prescriber name(s);
  - 3. Diagnoses relevant to the drug therapy to be managed and other patient conditions relevant to maintenance of the patient's health during drug therapy management;
  - 4. Protocols to be employed;
  - 5. Functions and activities the pharmacist will perform, and restrictions or limitations on the pharmacist's management;
  - 6. Method, content and frequency of reports to the prescriber;
  - 7. Manner in which pharmacist's drug therapy management will be monitored by the prescriber, including method and frequency;



8. A specified time, not to exceed 24 hours, within which the pharmacist must notify the prescriber of any modifications of drug therapy;
  9. A provision that allows the prescriber to override any action taken by the pharmacist when the prescriber deems it to be necessary;
  10. An effective date of the agreement, and signatures of both parties.
  11. A provision addressing how drug therapy management will be handled when the patient has more than one prescriber involved in evaluating or treating the medical condition which is the subject of the agreement. All prescribers who are actively involved in the management of the relevant conditions shall be parties to the agreement.
- c. Prior to implementation of drug therapy management, pharmacists shall secure the following information:
1. Patient's name, gender, date of birth, height, and weight;
  2. Patient diagnosis or diagnoses (from prescriber);
  3. Medication history;
  4. Prior lab values;
  5. Patient vital signs;
  6. Patient known allergies;
  7. Emergency contact number.
- d. Pharmacists engaging in drug therapy management shall not delegate drug therapy management responsibilities to any other staff.

**6.00.60 Recordkeeping.**

- a. Pharmacists must document all actions taken in drug therapy management, including but not limited to any data required by the protocol. Records of each patient visit must be transmitted to the prescriber in the manner specified in the agreement. Records must indicate when and how the record was transmitted to the prescriber.
- b. Pharmacists must keep patient records that include:
1. Patient's name, gender, date of birth, height, and weight;
  2. Patient diagnosis or diagnoses (from physician);
  3. Medication history;
  4. Prior lab values;
  5. Patient vital signs;
  6. Patient known allergies;

7. Date and time the service was rendered;
8. Type of service rendered;
9. Results of interviews with the patient and any diagnostic tests or other pertinent information about the patient's disease;
10. When and how the record was transmitted to the prescriber; and
11. Emergency contact number.

**6.00.70 Retention of Records.**

- a. All records of drug therapy management shall be retained for a minimum of seven years from the last date of drug therapy management, or seven years from the patient's 18<sup>th</sup> birthday, whichever is later. Such records shall be available for inspection by the patient, the prescriber, the Board, or any other authorized local, state, or federal law enforcement or regulatory agency.
- b. Records may be maintained in an alternative data retention system, such as a data processing system or direct imaging system provided that:
  1. The records maintained in the alternative system contain all of the information required on the manual record;
  2. The data processing system is capable of producing a hard copy of the record upon the request of the Board, its representative, or of other authorized local, state, or federal law enforcement or regulatory agencies;
  3. A back-up is conducted of the data processing system every 24 hours; and
  4. The records are immediately available for the previous two years.

**6.00.90 Confidentiality.**

- a. The pharmacist shall provide adequate security to prevent indiscriminate or unauthorized access to confidential records. If confidential health information is transmitted through a data communication device, the confidential health information may not be accessed or maintained by the operator of the data communication device unless specifically authorized to do so by the patient.
- b. Patient information is confidential and may be released only as authorized by state and federal law. All protected health information obtained and maintained, including that obtained from the physician or other providers, must be strictly controlled in accordance with the requirements of Health Insurance Portability and Accountability Act of 1996 and any rules promulgated pursuant to the act and other federal and state laws and rules. Specifically, pharmacists can only release patient information to:
  1. The patient or the patient's agent;
  2. A practitioner or another pharmacist if, in the pharmacist's professional judgment, the release is necessary to protect the patient's health and well-being;
  3. The Board or to a person or another state or federal agency authorized by law to receive the confidential record;

4. A person employed by a state agency that licenses a practitioner, if the person is performing the person's official duties; and/or
5. An insurance carrier or other third party payer authorized by the patient to receive the information.

**6.01.10 Participation Not Mandatory.**

- a. No person or entity, as a condition of employment, participation on an insurance provider panel, or otherwise, shall require any prescriber to participate in or authorize drug therapy management.

**6.01.20 Board Review.**

- a. Board staff will review compliance with this rule and report to the Board regarding complaints and other relevant data associated with the rule every three years.

**7.00.00 PHARMACIST MANAGER RESPONSIBILITIES.**

**7.00.10 Reporting Violations.** The pharmacist manager of a prescription drug outlet shall report to the Board, in writing, within the timelines set forth below:

- a. Diversion, theft or significant unaccountable loss of prescription drugs or controlled substances from the pharmacy, hospital or health maintenance organization (as defined in Section 10-16-102, C.R.S.) within one business day of discovery. When a Drug Enforcement Administration (DEA) Form 106 is submitted to the DEA in instances involving controlled substances, a copy of the completed DEA Form 106 along with a detailed written explanation shall be submitted to the Board. When determining whether an unaccountable loss is significant, the pharmacist manager shall consider, among others factors, the following:
  1. The actual quantity of drug lost in relation to the type of business;
  2. The specific drug lost;
  3. Whether the loss of the drug can be associated with access to those drugs by specific individuals, or whether the loss can be attributed to unique activities that may take place involving the drug;
  4. A pattern of losses over a specific time period, whether the losses appear to be random, and the results of efforts taken to resolve the losses;
  5. Whether the specific drug is a likely candidate for diversion; and
  6. Local trends and other indicators of the diversion potential of the missing drug.
- b. Security breaches within the pharmacy or pharmacy area of the establishment within 10 days of discovery.
- d. Any pharmacist working in the pharmacy who is impaired due to the use of alcohol or drugs, or a pharmacist with a mental or physical impairment which affects his ability to perform his job competently. In such instance the report shall be submitted to the Board immediately upon discovery.

- e. Significant errors related to the practice of pharmacy, including those related to compounding, such as those that result in serious personal injury or death of a patient. In such instance the report shall be submitted to the Board immediately upon discovery.

**7.00.20 Administrative Reporting Responsibilities:**

- a. A pharmacist manager shall immediately notify the Board in writing of the date he ceases to be the pharmacist manager of a prescription drug outlet.
- b. Upon the change of pharmacist manager of a prescription drug outlet, the new pharmacist manager or the new pharmacist manager's designee shall take an inventory of all controlled substances within seventy-two hours. The inventory shall be taken either as of the opening or as of the close of business activity on the inventory date and such time and date taken shall be entered on the inventory record.
- c. Upon the transfer of ownership of a prescription drug outlet, the pharmacist manager or the pharmacist manager's designee shall take an inventory of all controlled substances. The inventory shall be taken either as of the opening or as of the close of business activity on the inventory date and such time and date taken shall be entered on the inventory record.
- d. The pharmacist manager shall determine or approve procedures for prescriptions delivered or temporarily stored outside the confines of a compounding/dispensing area at the request of a patient or an agent of the patient. This procedure shall include the storage of, security of, the access to, the confidentiality of, and the counseling regarding, prescriptions and necessary record keeping.
- e. Upon the closure of a prescription drug outlet it shall be the responsibility of the last pharmacist manager of record to remove the prescription and/or chart orders to another prescription drug outlet where patrons and/or practitioners are afforded reasonable access to a pharmacist's interpretation of such orders.
- f. The daily printout shall contain all information as required by rule. This applies to both prescription order and chart order dispensing.
- g. It is the responsibility of the pharmacist manager to ensure that all prescription drug outlet staff are aware that they must be able to print a report of all prescription order or chart order transactions for such period of time as the Board or its inspector(s) may specify, or to provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review dispensing transactions, and, if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. (If the prescription drug outlet elects to comply with the latter requirement of providing equipment and/or personnel, the system must also be capable of printing the reports previously described.) Any failure or refusal by the pharmacist manager and/or a staff pharmacist to comply with a request by the Board or its inspector(s) will be deemed to be a willful violation of these rules.
- h. It is the responsibility of the pharmacist manager to maintain records as required by rule 11.00.00.
- i. It is the responsibility of the pharmacist manager to maintain records of initial interpretation and final evaluation as required by rule 3.00.51(a) and (b).

- J. It is the responsibility of the pharmacist manager to maintain and to assure the outlet's compliance with a policy and procedure manual, where applicable, encompassing all aspects of non-sterile and sterile compounding as required by rules 21.10.10 and 21.20.30, respectively. The annual review of such manual or manuals shall be signed and dated by the pharmacist manager. In the event the pharmacist manager changes, the new manager shall review, sign, and date the manual within 30 days of becoming the pharmacist manager.

**7.00.30 Compliance of Outlet:**

- a. The manager of a prescription drug outlet is responsible for the operation of the outlet in compliance with all state and federal laws, rules, and rules.
- b. The manager shall be responsible for posting the following information for each pharmacy technician working in the compounding/dispensing area:
  - 1. Certificate indicating the technician is certified by a nationally recognized certification Board; or
  - 2. Diploma indicating the technician has graduated from an accredited pharmacy technician training program; or
  - 3. Documentation that the pharmacy technician has completed five hundred hours of experiential training at the pharmacy. This documentation must be certified by the pharmacist manager of the prescription drug outlet; or
  - 4. Documentation that the pharmacy technician does not have certification from a nationally recognized certification Board, has not graduated from an accredited pharmacy technician training program, and has not completed 500 hours of experiential training at the pharmacy.
- c. The pharmacist manager is responsible for ensuring that all prescription drugs and controlled substances are procured by the outlet from an entity or person registered by the Board. Any drug designated as an Investigational New Drug from the Federal Food and Drug Administration is exempt from this requirement provided the research requirements for the receipt of the product are followed and it meets the requirements of CRS 12-42.5-128(2).

**8.00.00 ADVERTISING.**

- 8.00.10 Labels. At least one address shall appear on a prescription label and that shall include the address of the prescription drug outlet from which the prescription was dispensed. In the case of a central fill prescription processing contract, the label shall contain at least the name and address of the originating and/or fulfillment pharmacy.
- 8.00.20 Prescription Order Forms. No prescription drug outlet shall provide any practitioner with prescription order forms that refer to a pharmacist or prescription drug outlet.
- 8.00.30 Multiple Names. A prescription drug outlet shall only use, operate or advertise under the name that appears on the current registration issued by the Board.
- 8.00.40 Truth in Advertising. No pharmacist or prescription drug outlet shall advertise or allow advertisement that is untrue or misleading in any manner regarding prescription drugs.

**9.00.00           LEGAL PROCEEDINGS.**

**9.00.10 Reporting.**

- a. A licensee or registrant shall notify The Board in writing within 72 hours of the licensee or registrant receiving service of process or knowledge by other means of any legal proceedings in Colorado or anywhere wherein it is alleged that the licensee or registrant has violated any law or rule pertaining to drugs or devices. This includes civil malpractice cases.
  - 1. The notice to the Board shall include the following information:
    - (a) The court;
    - (b) The jurisdiction;
    - (c) The case name;
    - (d) The case number; and
    - (e) A description of the matter and a copy of the indictment or charges.
  - 2. The licensee or registrant shall notify the Board in writing within 30 days of the disposition of such proceeding.
- b. All licensees or registrants shall notify the Board in writing within 30 days of any disciplinary action against them in another state. Such notification shall include the following:
  - 1. The state;
  - 2. The jurisdiction;
  - 3. The case name;
  - 4. The case number;
  - 5. A description of the matter and a copy of the indictment or charges;
  - 6. A copy of the discipline; and
  - 7. Proof of completion of any requirements set forth in the order, if applicable.
- c. All licensees or registrants shall notify the Board in writing of any criminal conviction or deferred judgment against them (including, but not limited to, "driving under the influence" and "driving while ability impaired"), and petty offenses within 30 days after such conviction or judgment.
  - 1. For purposes of this rule, a "conviction" includes:
    - (a) A guilty verdict;
    - (b) A plea of guilty accepted by the court;
    - (c) A plea of nolo contendere (no contest) accepted by the court; or

- (d) A deferred judgment or sentence.
- 2. The notice to the Board shall include the following information:
  - (a) The court;
  - (b) The jurisdiction;
  - (c) The case name;
  - (d) The case number;
  - (e) A description of the matter and a copy of the indictment or charges;
  - (f) A copy of the plea agreement or verdict; and
  - (g) Proof of completion of court ordered requirements, if applicable.
- d. The registrant or licensee notifying the Board may submit a written statement with any notice required under this rule to be included in the registrant or licensee records.
- e. Each insurance company licensed to do business in Colorado and engaged in the writing of malpractice insurance for licensed pharmacists and each pharmacy that self-insures shall send to the Board, in the form prescribed by the Board, information relating to each malpractice claim against a licensed pharmacist which is settled or in which judgment is rendered against the insured. Such information shall be provided to the Board within 30 days of the settlement or judgment.

**10.00.00 EMERGENCY KITS.**

**10.00.05 Definitions.**

- a. "Emergency kit" or "kit" means a tamper-evident sealed and secured container or secured electronic system containing drugs which are used for either immediate administration to patients of facilities delineated in 10.00.10 or in an emergency situation or as a starter dose.
- b. "Starter dose" means a dose of medication contained in an emergency kit for the purpose of starting the initial therapy for a patient residing in a facility delineated in 10.00.10.

**10.00.10** A prescription drug outlet or a hospital other outlet may provide an emergency kit to any of the following facilities that are licensed or certified by the Colorado Department of Public Health and Environment: Long-Term Care Facilities, Hospices, Acute Treatment Units and Home Health Agencies. Such kit is to provide an emergency supply of drugs, both controlled and non-controlled as provided below. The drugs maintained in the emergency drug supply shall remain the property of the prescription drug outlet or the hospital other outlet who supplied the drugs.

- a. Only one prescription drug outlet or hospital other outlet may provide a kit to any of the above facilities. Multiple pharmacies or hospital other outlets may not supply emergency kits to the same facility.

- b. The pharmacist manager of the prescription drug outlet supplying the kit or the consultant pharmacist of the hospital other outlet supplying the kit shall be responsible for the accurate stocking or restocking of the kit. He/she may delegate this function to non-pharmacist personnel, but the pharmacist manager or other outlet consultant pharmacist assumes responsibility for the accuracy of the contents of the kit.

**10.00.20 Categories and Limits**

- a. For Long-Term Care Facilities, Acute Treatment Units, and Inpatient Hospices, the medical director of the facility, or equivalent, and the consulting pharmacist shall determine the specific drugs to be kept in the kit. The number of drugs allowed in the kit shall be limited to sixty (60). Of the 60, twelve (12) may be controlled substances. The kit may contain no more than thirty (30) doses of any separate drug dosage form or strength for each drug. The container size for each drug shall be limited to unit dose or unit of issue packaging.
- b. In the case of a Certified Home Health Agency or an Outpatient Hospice, the director of nursing of the Certified Home Health Agency or of the Licensed Hospice, and a pharmacist employed and designated by the prescription drug outlet or hospital other outlet providing the kit shall determine the specific drugs to be kept in the kit. A Certified Home Health Agency or Outpatient Hospice may not have oral dosage forms or controlled substances in the kit. The container size for each injectable drug shall be limited to unit dose or unit of issue packaging. The number of drugs allowed in the kit shall be limited to sixty (60). The kit may contain only thirty (30) doses of any separate drug dosage form or strength for each drug.

**10.00.30** The kit shall be sealed with a tamper-evident seal or an electronic system which notifies the pharmacy when the kit has been accessed. Paper or tape seals are unacceptable. If an electronic system is utilized, the pharmacy and facility must maintain a written procedure for how the kit can be accessed in the event of downtime.

**10.00.40** The following information shall be readily retrievable and up-dated as required:

- a. Name, address and telephone number of the prescription drug outlet or hospital other outlet providing the contents of the kit;
- b. The date of sealing of the kit;
- c. A suitable expiration date which shall be the earliest expiration date of any drug in the kit, but in no event shall it be more than one year from the date of sealing; and
- d. In the case of a Long-Term Care Facility, Acute Treatment Unit or Inpatient Hospice, the name of the consulting pharmacist, or, in the case of a Certified Home Health Agency or an Outpatient Hospice, the name of the designated pharmacist.

**10.00.41** A copy of the kit contents shall also be attached to the kit.

**10.00.50** Access. Access to the contents of the kit shall be limited as follows:

- a. In the case of a Long-Term Care Facility, Acute Treatment Unit, or Inpatient Hospice, only a pharmacist employed by the prescription drug outlet or hospital other outlet which provides the kit or his/her designee, the consulting pharmacist, and any nurse employed at the facility shall have access.



- b. In the case of a Certified Home Health Agency or an Outpatient Hospice, only a pharmacist employed by the prescription drug outlet or hospital other outlet which provides the kit or a nurse employed by the Certified Home Health Agency or an Outpatient Hospice shall have access.

10.00.60        Inspection. A pharmacist employed by the prescription drug outlet or hospital other outlet providing the kit or that pharmacist's designee shall inspect and inventory the contents of the kit at least annually and within 72 hours after being notified that the kit has been accessed. Inspection shall be documented by that pharmacist, and such documentation shall be maintained and available for inspection at the prescription drug outlet or hospital other outlet for a period of two years.

10.00.70        A separate record of use for each drug placed in the kit, and for each kit provided, which shall state the following Records. The prescription drug outlet or hospital other outlet providing the kit shall maintain:

- a. The name and address of the Acute Treatment Unit, Long-Term Care Facility, Certified Home Health Agency, or Hospice;
- b. The name and strength of the drug; and
- c. The container size and the quantity initially placed in the kit.

10.00.71        When a drug is removed for administration the prescription drug outlet or hospital other outlet shall obtain a prescription order or LTCF chart order for the drug within 72 hours after being notified that the kit was opened and the drug was used. The order shall indicate the total number of doses administered. The order shall be assigned a serial number and the order shall be retained as required by rule 11.04.10. Additionally, the separate record required for each drug in the kit shall reflect the following information:

- a. Date and quantity administered;
- b. Names of both the patient and practitioner;
- c. Date the drug was replaced in the kit;
- d. The quantity of the drug replaced, which shall not exceed the quantity administered or removed for administration; and
- e. The prescription order number assigned.

10.00.80        Use. The drugs shall only be administered to patients of the Acute Treatment Unit, Long-Term Care Facility, Certified Home Health Care Agency, or Hospice pursuant to the order of a practitioner.

**11.00.00 RECORDS AND RECORDKEEPING.**

11.01.00 Records in General. All outlets registered and/or licensed by the Board shall maintain such records and inventories of prescription drugs as may be required by these rules or any other state or federal law or rule pertaining to such drugs. Such records shall be maintained on a current basis and shall be complete and accurate for all drugs which the outlet manufactures, receives, dispenses, distributes or otherwise disposes of in any other manner. Records and inventories of controlled substances shall be deemed to be "complete" only if each individual record and inventory contains all required information regarding each specific transaction, and if the set of records and inventories contains all information and documents requires to be kept by state and federal laws, and rules. A record or inventory shall be deemed to be "accurate" only if it is a complete, true and factual statement regarding or reflecting each specific transaction. A set of records or inventories shall be deemed to be "accurate" only if they are complete, and when considered as a whole, they demonstrate that the controlled substances and/or the records and inventories pertaining thereto have been handled in compliance with all applicable laws or rules and that all such controlled substances are properly accounted for.

All such records shall be retained for a period of at least two years after the date of any transaction relating to such record or inventory by any process providing an exact duplicate of the original order in a reproducible quality acceptable to the Board. Records shall be retained in a format that cannot be altered.

Records on an automated data processing system or subsequent storage of such records must be immediately retrievable (via monitor display or hard copy printout). Upon written Board approval, outlets capable of meeting the above standards may not be required to retain the original prescription order or LTCF chart order for non-controlled drugs.

11.02.00 Retrievability of Records. For the purposes of these rules, records and inventories shall be deemed "readily retrievable" if they meet the following requirements:

For all Registered Prescription Drug Outlets:

- a. The following records shall be maintained on the premises of the prescription drug outlet at all times, unless written authorization for off-site storage has been approved by the Board, and shall be made available for inspection by the Board or its inspectors immediately upon request:
  - (1) All DEA-222 forms executed during the two years preceding the request;
  - (2) All inventories of controlled substances required to be taken during the two years preceding the request;
  - (3) All prescription orders or LTCF chart orders dispensed during the two years preceding the request;
  - (4) All records of dispensing, receipts (invoices for drugs received and credited) of controlled substances, distribution, loss, surrender or disposal in manner of prescription drugs and controlled substances during the two years preceding the request;
  - (5) All lists as required by rules 11.08.00 and 11.09.00.
- b. The following records shall be made available within 48 hours or two business days, whichever is longer, on request by the Board or its inspectors:

- (1) All unexecuted DEA-222 forms.
- (2) Specific records requested by the inspector if the inspector determines the records are not maintained in a readily retrievable manner.
- (3) Records of receipt of non-controlled prescription drugs.

11.03.00 Inventories of Controlled Substances. Any inventory of controlled substances shall comply with the following:

- a. If the outlet is registered with the Drug Enforcement Administration as a "hospital/clinic", or is owned and operated by a health maintenance organization (as defined in section 10-16-102, C.R.S), the inventory shall include all drugs located throughout the facility, excluding any drug which has been dispensed pursuant to a lawful chart order but which has not yet been administered to the patient.
- b. Each inventory shall contain a complete and accurate records of all controlled substances (including outdated controlled substances) on hand on the date the inventory is taken. The inventory shall be maintained in written, typewritten or printed form at the prescription drug outlet. Schedule II drugs shall be separated from schedule III, IV, and V drugs. Controlled substances shall be deemed to be "on hand" if they are in the possession of or under the control of the registrant. However, the inventory shall exclude any drug that has been dispensed pursuant to a lawful prescription order but which has not yet been delivered.
- c. The inventory shall be taken either as of opening of business or as of the close of business on the inventory date and it shall be recorded on the inventory. In the event the prescription drug outlet is open 24 hours per day, the inventory shall specify the time the inventory was conducted.
- d. After the initial inventory is taken, the prescription drug outlet shall take new inventory of all stocks of controlled substances on hand at least every two years.
- e. On the effective date of a law or rule on which a previously non-scheduled drug is added to any schedule of controlled substances, every prescription drug outlet that possesses that drug shall take an inventory of all stocks of the drug on hand. Thereafter, that drug shall be included in each inventory made by the prescription drug outlet.
- f. The following information shall be recorded on the inventory.
  - (1) The name of the drug;
  - (2) Each finished form of the drug (strength and dosage form);
  - (3) The number of units or volume of each finished form.
  - (4) All outdated controlled substances.
- g. In determining the number of units of each finished form of a controlled substance in a commercial container which has been opened, the prescription drug outlet shall do as follows:
  - (1) If the drug is a schedule II drug, an exact count of the contents shall be made.

- (2) If the substance is listed in schedule III, IV, or V, and estimated count of the measure of the contents may be made, unless the container holds more than 1,000 tablets or capsules, in which case an exact count of the contents must be made.

- h. All controlled substance inventories shall be retained at the prescription drug outlet for at least two years from the date of such inventory.

11.04.00 Records pertaining to prescription orders and chart orders for long-term care facility patients (LTCF chart orders).

11.04.10 A hard copy of every prescription order shall be readily retrievable, legible, and available for inspection for a period of two years from the date of any transaction relating to such prescription order unless the prescription drug outlet has received written Board approval to not retain the original prescription order for non-controlled drugs and Schedule III, IV, and V Controlled Substances. Prescription orders will be deemed to be readily retrievable, legible, and available if they are filed according to the numerical sequence of the serial numbers assigned pursuant to 2.01.10, and are easily readable without the aid of any special device. In addition to being filed in numerical sequence, three different prescription files shall be maintained: one file shall consist only of schedule II controlled substance prescription orders; the second file shall consist only of schedule III, IV and V controlled substance prescription orders; and the third file shall consist of all non-controlled substance prescription orders. Filing of prescription orders in any manner other than by numerical sequence will result in such prescription orders being deemed not readily retrievable and available.

A hard copy of every chart order shall be readily retrievable and available for inspection for a period of two years from the date of any transaction relating to such order. The LTCF chart orders will be deemed to be readily retrievable and available if they are filed according to the date of dispensing. Chart orders for schedule III, IV, and V controlled substances shall be readily identifiable from non-controlled chart orders. Schedule II orders shall be retained separately from all other orders.

If a prescription drug outlet utilizes both prescription orders and chart orders, assigning serial numbers to both with the same computer system, the orders must be filed sequentially by serial number.

11.04.20 Computer Use With Prescription Order or LTCF Chart Order Transactions.

A prescription drug outlet may utilize a computer (automated data processing system) for storage and retrieval of information regarding prescription and LTCF chart order transactions. The following requirements shall be met:

- a. All new and refill prescription and LTCF chart order transactions shall be entered into the system at the time of the transaction, except as provided in rule 11.04.20(i).
- b. Every 24 hours, except as provided in rule 11.04.30, the system must produce a hard-copy document which, for the purposes of these rules, shall be known as the "daily printout". It shall consist of a single, uniform, complete document, except as otherwise permitted in this rule. The daily printout shall list, separately, each prescription fulfillment or LTCF chart order transaction for the previous 24 hours and shall contain all information required by this rule. Daily printouts shall be retained in a chronological manner. If the printouts are bound, the sheets shall be separated into individual pages which are then placed in the same order as printed and bound uniformly. If the pages are bound in any other manner so that they are not uniform in placement or appearance, they shall be deemed not readily retrievable and available.

- c. The daily printout shall contain all of the following information for each prescription fulfillment or LTCF chart order transaction and shall differentiate between new and refill transactions. As an alternative, new and refill transactions may be separated into two separate uniform and complete documents which contain the following:
  - (1) The serial number;
  - (2) The name of the patient;
  - (3) The name of the practitioner;
  - (4) For each controlled substance dispenses, the practitioner's Drug Enforcement Administration registration number;
  - (5) The date of issue by the practitioner, the date dispensed shall be presumed to be the date of issue.
  - (6) The total number of refills authorized;
  - (7) Date dispensed;
  - (8) The name and strength of the drug dispensed;
  - (9) The quantity of the drug dispensed;
  - (10) In the case of a refill, the total number of refills dispensed to date.
- d. Records of prescription or LTCF chart order transactions involving controlled substances shall be identifiable from those involving non-controlled substances. Alternatively a separate complete printout listing only controlled substance transactions may be produced.
- e. The daily printout shall be available for inspection by the Board within 72 hours from the most recent date recorded on the printout.
- f. Prescription or LTCF chart order refill transactions must be uniformly recorded on the original prescription or LTCF chart order or on the daily printout. In the event of a discrepancy between the entry on the order and the entry on the daily printout, the information recorded on the daily printout shall be deemed to be correct.

- g. Documentation of the fact that the refill information entered into the automated data processing system each time a pharmacist refills an original prescription or LTCF chart order for a schedule III, IV, or V controlled substance is correct must be provided by the pharmacist who makes the final evaluation. This documentation may be retained in the following manner:
  - (1) If such a system provides a hard-copy printout of each day's controlled substance prescription or LTCF chart order refill data, the controlled substance refill information shall be verified, dated, and signed by the pharmacist making the final evaluation. The pharmacist shall verify that the date indicated is correct and then sign this document in the same manner as he/she would sign a check or legal document. This document shall be maintained in a separate file at the prescription drug outlet for a period of two years from the dispensing date. The printout of the day's controlled substance prescription or LTCF chart order refills must be generated by the prescription drug outlet within 72 hours of the date on which the refill was dispensed. It must be verified and signed by each pharmacist who is involved in dispensing controlled substance refills.
  - or
  - (2) The prescription drug outlet shall maintain a bound log book, or separate file, in which each pharmacist involved in dispensing controlled substance order refills shall sign attesting to the fact that the refill information entered into the computer that day has been reviewed by him/her and is correct as shown. Such a book or file must be maintained at the prescription drug outlet for a period of two years after the date of dispensing the appropriately authorized refill.
- h. The daily printout shall contain all information as required by this rule except that the identity of the pharmacist who makes the final evaluation may appear either on the daily printout or on another separate, uniformly maintained and readily retrievable record. The outlet manager shall determine which of the two methods for identifying the responsible pharmacist is more appropriate for the outlet, and only that method for recording such information shall be used.
- i. Because of the potential for a system malfunction or failure, the prescription drug outlet must have a manual procedure for recording all dispensing transactions during the system failure or malfunction. All recoverable transaction data and all manually recorded transaction data shall be entered or restored into the system within a reasonable period of time not to exceed seven days following the restoration or operation of the system.
- j. Any automated data processing system used by an outlet shall maintain the confidentiality of records in accordance with applicable laws, rules and regulations.

11.04.30 Electronic Maintenance of Prescription or LTCF Chart Order Records.

A prescription drug outlet which utilizes a computer (automated data processing system) for storage and retrieval of information regarding prescription fulfillment or LTCF chart order transactions need not print the daily printout required by rule 11.04.20 if the prescription drug outlet and the computer system utilized are capable of complying with the following requirements:

- a. The prescription drug outlet must be able to provide on-line retrieval of all information required by this rule for all prescription order transactions during the two years preceding the request.

- b. The prescription drug outlet must ensure a daily (i.e., every twenty-four hours) back up is performed for use in restoring required information in case of a system failure.
- c. The prescription drug outlet must:
  - (1) Have and maintain a complete on-line transaction file that is printable on the inspector's request,  
  
or
  - (2) Have a "lock-out" feature that prevents editing of dispensing information.
- d. The Board or its inspectors must be able to inspect and review the prescription order transactions of the prescription drug outlet. Therefore, immediately upon the oral or written request of the Board or its inspectors, the prescription drug outlet shall either:
  - (1) Print a report of all prescription or LTCF chart order transactions for such period of time as the Board or its inspector(s) may specify. The system must be capable of retrieving and printing such a report within a limited time not to exceed two hours. Additionally within 72 hours, the system must be capable of retrieving and printing the information sorted according to the variables which include, but are not limited to, date dispensed; drug name, strength and dosage form; patient name, and practitioner name; or
  - (2) Provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review dispensing transactions, and, if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. If the prescription drug outlet elects to comply with this subparagraph (d), the system must also be capable of printing the same reports described in subparagraph (1).
  - (3) It is the responsibility of the prescription drug outlet manager to ensure that all prescription drug outlet staff are aware of the requirements of subparagraphs (1) and (2). Any failure or refusal by the prescription drug outlet manager and/or a staff pharmacist to comply with a request by the Board or its inspector(s) will be deemed to be a willful violation of these rules.
- e. Whether the prescription drug outlet elects to comply with rule 11.04.30 (d), the system and any reports printed on request shall contain, as a minimum, the following information for each transaction:
  - (1) The prescription order serial number;
  - (2) The name of the patient;
  - (3) The name of the practitioner;
  - (4) For each controlled substance dispensed, the practitioner's Drug Enforcement Administration registration number;
  - (5) The date of issue by the practitioner; if the date is omitted by the practitioner, the date dispensed shall be presumed to be the date of issue;
  - (6) The total number of refills authorized;

- (7) Date dispensed;
- (8) The name and strength of the drug dispensed;
- (9) The quantity of the drug dispensed;
- (10) In the case of a refill, the total number of refills dispensed to date;
- (11) Whether the prescription order is a new or refill transaction;
- (12) In the case of a controlled substance, a means of visually identifying orders for such substances and differentiating them from non-controlled substances.

**11.05.00** Records Pertaining to Hospital Chart Orders.

**11.05.10** A chart order or the other appropriate, uniformly maintained records permitted by rule 2.01.20 (c) shall be readily retrievable and available for inspection for a period of two years from the date of any transaction relating to such order or record. However, if the records permitted by rule 2.01.20 (c) and 11.05.20 are retained and are complete, the prescription drug outlet copy of a chart order need not be retained. Prescription drug outlet copies of chart orders or the other appropriate, uniformly maintained records permitted by 2.01.20 (c) and 11.05.20 will be deemed to be readily retrievable and available if they are filed:

- a. In chronological order according to the date of discharge of the patient; or
- b. Alphabetically by patient surname by month of discharge; or
- c. By date of dispensing transaction.

Filing of chart order/dispensing transaction record in any other manner will result in such orders or records being deemed not readily retrievable and available.

**11.05.20** Records Pertaining to Chart Orders Shall Contain the Following Information:

- a. The identity of the pharmacist making the initial interpretation;
- b. The identity of the pharmacist making the final evaluation each time a drug is dispensed, if different from the pharmacist making the initial interpretation;
- c. The quantity dispensed and
- d. The date of dispensing;
- e. Any record of a controlled substance dispensed pursuant to a chart order for and individual patient shall be visually identifiable from records of non controlled substances.

**11.05.21** It is permissible to store different types of chart order dispensing records separately. For the purpose of this rule, different types of chart order dispensing records include fill lists, records of compounded injectable products, records of the initial dispensing of a chart order, and the records of redispensing of chart orders. If the prescription drug outlet chooses to maintain different types of dispensing records separately, they must be maintained as required by rule 11.05.10.



- 11.05.30 Computer Use with Hospital Chart Order Transactions. A prescription drug outlet may utilize a computer (automated data processing system) for storage and retrieval of information regarding chart order transactions if the following requirements are met:
- a. All new chart orders shall be entered into the system, except as provided in rule 11.05.30 (e). For the purpose of this rule, "dispensing transaction" is defined as delivery of a drug or device pursuant to a chart order.
  - b. All records produced by this computer system must comply with rule 11.05.20. These records shall be printed a minimum of every 24 hours unless the prescription drug outlet complies with rule 11.05.40. This documentation shall be retained for at least two years from the date of dispensing. This documentation shall be retained in a chronological manner. If printouts are bound, the sheets shall be separated into individual pages, which are then placed in the same order as printed and bound uniformly. If the pages are bound in any other manner so that they are not uniform in placement or appearance, they shall be deemed not readily retrievable. This documentation shall be available for inspection by the Board or its inspectors within 72 hours from the most recent date recorded on the documentation.
  - c. Any computer system utilized shall have the capability of producing a single-document printout, which shows for any controlled substance a complete history of all dispensing transactions during the previous two years for each patient admission. This printout shall be available within 72 hours of a request by the Board.
  - d. Because of the potential for a system malfunction or failure, the prescription drug outlet must have a manual procedure for recording all dispensing transactions during the system failure or malfunction. All recoverable transaction data and all manually entered transaction data shall be entered or restored into the system within a reasonable period of time not to exceed seven days following the restoration of operation of the system.
  - e. Any automated data processing system used by an outlet shall maintain the confidentiality of records in accordance with applicable laws, rules and rules.
- 11.05.40 Electronic Maintenance of Hospital Chart Order Records. A prescription drug outlet which utilizes a computer (automated data processing system) for storage and retrieval of information regarding chart order transactions need not print the records of chart order dispensing required by rule 11.05.20, if the prescription drug outlet and the computer system utilized are capable of complying with the following requirements.
- a. The prescription drug outlet must be able to provide on-line retrieval of all information required by this rule for all chart order transactions during the two years preceding the request.
  - b. The prescription drug outlet must ensure a daily (i.e., every twenty-four hours) back up is performed for use in restoring required information in case of a system failure.
  - c. The prescription drug outlet must:
    - (1) Have and maintain a complete on-line transaction file that is printable on the inspector's request,
    - or
    - (2) Have a "lock-out" feature that prevents editing of dispensing information.

- d. The Board or its inspectors must be able to inspect and review the chart order transactions of the prescription drug outlet. Therefore, immediately upon the oral or written request of the Board or its inspectors, the prescription drug outlet shall either:
- (1) Print a report of all chart order transactions for such period of time as the Board or its inspector(s) may specify. The system must be capable of retrieving and printing such a report within a limited time not to exceed two hours. Additionally, within 72 hours, the system must be capable of retrieving and printing the information sorted according to variables which include, but are not limited to, date dispensed; drug name, strength and dosage form; patient name; or
  - (2) Provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review dispensing transactions, and, if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. If the prescription drug outlet elects to comply with this subparagraph (b), the system must also be capable of printing the same reports described in subparagraph (1)
  - (3) It is the responsibility of the prescription drug outlet manager to ensure (1) and (2). Any failure or refusal by the prescription drug outlet manager and/or a staff pharmacist to comply with a request by the Board or its inspector(s) will that all prescription drug outlet staff are aware of the requirements of subparagraphs be deemed to be a willful violation of these rules.
- e. Whether the prescription drug outlet elects to comply with rules 11.05.40 (d) (1) or (2), the system and any reports printed on request shall contain, as a minimum, the following information for each transaction:
- (1) The name and/or other identifying factor of the patient;
  - (2) The identity of the pharmacist making the initial interpretation;
  - (3) The quantity dispensed;
  - (4) The date of dispensing;
  - (5) Any record of a controlled substance dispensed pursuant to a chart order for an individual patient shall be distinguishable from records of non-controlled substances. Alternatively, a separate complete printout listing on controlled substance transactions may be produced.
- f. The daily printout shall contain all information as required by this rule except that the identity of the pharmacist who makes the final evaluation may appear either on the daily printout or on another separate, uniformly maintained and readily retrievable record. The outlet manager shall determine which of the two methods for identifying the responsible pharmacist is more appropriate for the outlet, and only that method for recording such information shall be used.

11.06.00 Receipts

- 11.06.05 All prescription drugs and controlled substances received by a prescription drug outlet shall only be procured from another entity or person registered by the Board.

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- 11.06.10 Records of receipts of prescription drugs and controlled substances shall contain the following information for each such substance received:
- a. Name of the drug;
  - b. Strength of the drug;
  - c. Dosage form if appropriate;
  - d. Quantity received;
  - e. Date received;
  - f. Name of the labeler of the drug and/or NDC number of the drug if it is labeled only with its generic name;
  - g. Name and address of the distributor;
  - h. Name and address of the receiving outlet;
  - i. DEA number of distributor and receiver if a controlled substance; and
  - j. If a schedule II controlled substance, The DEA form 222 or a copy of the DEA form 222 and the corresponding invoice shall be attached to each other.
- 11.06.20 All records of receipt (including credit invoices) of prescription drugs and controlled substances shall be maintained for a period of time not less than two years from the date the drugs were received. Hard copy records of receipt (including credit invoices) of non-controlled prescription drugs need not be maintained on the premises provided that they shall be made available within 48 hours or two business days, whichever is longer, on request by the Board or its inspectors.
- 11.06.30 All records of receipt of schedule II controlled substances shall be maintained separately from all other records and be readily available for inspection in hard copy form at the outlet for a period of time not less than two years from the date the drugs were received.
- 11.06.40 All records of receipt of schedule III, IV, and V controlled substances shall be maintained separately from all other records and shall at all times be maintained and readily available for inspection in hard copy form at the outlet for a period of time not less than two years from the date the drugs were received.
- 11.06.50 Records detailing the receipt of prescription drugs, as required by rule 11.06.10(a) through (h), may be maintained electronically if the following requirements are met:
- a. The prescription drug outlet must ensure a daily (i.e., every twenty-four hours) back up is performed for use in restoring required information in case of a system failure;
  - b. Have and maintain a complete on-line receipt file that is printable on the inspector's request; or
  - c. Have a "lock-out" feature that prevents editing of receipt information;
  - d. The Board or its inspectors must be able to inspect and review all of the prescription drug receipt transactions of the outlet for the preceding two years. Therefore, immediately upon the oral or written request of the Board or its inspectors, the outlet shall either:

- (1) Print a report of all prescription drug receipt transactions for a period of time as the Board or its inspector(s) may specify. The system must be capable of retrieving and printing such a report within a limited time not to exceed two hours;
  - Or
  - (2) Provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review prescription drug receipt transactions, and if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. If the outlet elects to comply with this subparagraph (2), the system must also be capable of printing the same reports described in subparagraph (1).
  - (3) It is the responsibility of the pharmacist manager of the outlet to ensure that all outlet staff are aware of the requirements of subparagraphs (1) and (2). Any failure or refusal by the pharmacist manager and/or staff to comply with a request by the Board or its inspector(s) will be deemed to be a willful violation of these rules.
- e. If the outlet chooses to maintain records detailing the receipt of prescription drugs electronically, any reports printed upon the request shall contain, at a minimum, the following information for each receipt transaction:
  - (1) Name of the drug;
  - (2) Strength of the drug;
  - (3) Dosage form if appropriate;
  - (4) Quantity of the drug received;
  - (5) Date received;
  - (6) Name of the labeler of the drug and/or NDC number of the drug if it is labeled only with its generic name;
  - (7) Name and address of the distributor; and
  - (8) Name and address of the receiving outlet.

**11.07.00 Distribution**

**11.07.10** Records of distribution of controlled substances and prescription drugs within hospitals and facilities owned and operated by health maintenance organizations (as defined in section 10-16-102, C.R.S.). Records of distribution of controlled substances and prescription drugs shall comply with the following:

- a. In a hospital or a facility owned and operated by health maintenance organization which operates a registered prescription drug outlet, a controlled substance or prescription drug may be distributed for floor stock to appropriate areas within the hospital or facility. A record of any such distribution shall be made and retained by the Prescription Drug Outlet for a period of time not less than two years and shall include the following information:
  - (1) The location receiving the drug;

- (2) The name of the drug;
  - (3) The strength of the drug;
  - (4) The quantity of the drug;
  - (5) The dosage form if appropriate;
  - (6) The date the drug was supplied;
  - (7) The identity of the person in the prescription drug outlet who issued the drug;
  - (8) The identity of the person who placed the drug into floor stock.
- b. These records of distribution may be retained electronically provided the following requirements are met:
- (1) The prescription drug outlet must ensure a daily (i.e., every twenty-four hours) back up is performed for use in restoring required information in case of a system failure.
  - (2) The prescription drug outlet must:
    - (a) Have and maintain a complete on-line distribution file that is printable on the inspector's request,
    - or
    - (b) Have a "lock-out" feature that prevents editing of distribution information.
  - (3) The Board and its inspectors must be able to inspect and review the distribution transactions of the prescription drug outlet. Therefore, immediately upon the oral or written request of the Board or its inspectors, the prescription drug outlet shall either:
    - (a) Print a report of all distribution transactions for such period of time as the Board or its inspector(s) may specify. The system must be capable of retrieving and printing such a report within a limited time not to exceed two hours. Additionally, the system must be capable of retrieving and printing the information sorted according to variables which include, but are not limited to, date distributed, drug name, strength and dosage form, or
    - (b) Provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review distribution transactions, and, if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. If the prescription drug outlet elects to comply with this subparagraph (2), the system must also be capable of printing the same reports described in subparagraph (1)

- (c) It is the responsibility of the prescription drug outlet manager to ensure that all prescription drug outlet staff are aware of the requirements of subparagraphs (1) and (2). Any failure or refusal by the prescription drug outlet manager and/or a staff pharmacist to comply with a request by the Board or its inspector(s) will be deemed to be a willful violation of these rules.
- (4) If the prescription drug outlet chooses to maintain records of distribution electronically, any reports printed upon request shall contain, as a minimum, the following information for each transaction:
  - (a) The location receiving the drug;
  - (b) The name of the drug;
  - (c) The strength of the drug;
  - (d) The quantity of the drug;
  - (e) The dosage form if appropriate;
  - (f) The date the drug was supplied;
  - (g) The identity of the person in the prescription drug outlet who issued the drug;
  - (h) The identity of the person who placed the drug into floor stock.

11.07.20       Records of Distribution/Casual Sale of Controlled Substances and Prescription Drugs. A prescription drug outlet which distributes prescription drugs and/or controlled substances shall record the following:

- a. The name of the drug;
- b. The strength of the drug;
- c. The dosage form if appropriate;
- d. The quantity of the drug;
- e. The name of the manufacturer or the NDC number of the drug if labeled only with its generic name. In the case of a compounded product, the name of the pharmacy shall be recorded;
- f. If a compounded product, the batch or lot number;
- g. The date of distribution;
- h. The name and address of the distributing outlet;
- i. The name and address of the receiver;
- j. If a controlled substance is distributed, the record shall also indicate the Drug Enforcement registration number of the distributing outlet and the receiver.

- k. A schedule II controlled substance shall only be distributed pursuant to receipt of a properly executed DEA-222 form.
  - l. The internal lot number assigned if the drug is packaged and distributed by a prescription drug outlet owned and operated by a hospital that is accredited by the Joint Commission of Accreditation of Healthcare Organizations or a successor organization or by a prescription drug outlet operated by a health maintenance organization as defined in section 10-16-102, C.R.S. Such drugs may only be distributed to prescription drug outlets under common ownership.
- 11.07.21 These records of distribution and casual sale required by 11.07.20 shall be retained for a period of time not less than two years from the date of the distribution.
- 11.07.22 Records of distribution and casual sale required by rule 11.07.20 may be maintained electronically if the following requirements are met:
- a. The prescription drug outlet must ensure a daily (i.e., every twenty-four hours) back up is performed for use in restoring required information in case of a system failure.
  - b. Have and maintain a complete on-line distribution and casual sale file that is printable on the inspector's request, or
  - c. Have a "lock-out" feature that prevents editing of distribution and casual sale information.
  - d. The Board or its inspectors must be able to inspect and review the distribution transactions of the prescription drug outlet. Therefore, immediately upon the oral or written request of the Board or its inspectors, the prescription drug outlet shall either:
    - (1) Print a report of all distribution and casual sale transactions for a period of time as the Board or its inspector(s) may specify. The system must be capable of retrieving and printing such a report within a limited time not to exceed two hours. Additionally, the system must be capable of retrieving and printing the information sorted according to variables which include, but are not limited to; date of distribution, drug name, strength and dosage form, and licensee receiving the distribution;
    - or
    - (2) Provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review distribution transactions, and, if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. If the prescription drug outlet elects to comply with this subparagraph (2), the system must also be capable of printing the same reports described in subparagraph (1).
    - (3) It is the responsibility of the prescription drug outlet manager to ensure that all prescription drug outlet staff are aware of the requirements of subparagraphs (1) and (2). Any failure or refusal by the prescription drug outlet manager and/or a staff pharmacist to comply with a request by the Board or its inspector(s) will be deemed to be a willful violation of these rules.
  - e. If the prescription drug outlet chooses to maintain records of casual sales and distribution electronically, any reports printed upon request shall contain, as a minimum, the following information for each transaction:

- (1) The name of the drug;
- (2) The strength of the drug;
- (3) The dosage form if appropriate;
- (4) The quantity of the drug;
- (5) The manufacturer name and/or NDC number of the drug if labeled only with its generic name;
- (6) The date of distribution;
- (7) The name and address of the distributing outlet;
- (8) The name and address of the receiver;
- (9) If a controlled substance is distributed, the record shall also indicate the Drug Enforcement registration number of the distributing outlet and the receiver.

11.08.00 List of Employees. Each prescription drug outlet shall keep and maintain on a current basis a list of every licensed pharmacist and intern who has practiced pharmacy in that outlet at any time during the previous two years, including all part-time or relief personnel. This list shall show, for each such person, the following information:

- a. The printed name of the person;
- b. The person's license number;
- c. A sample of his/her initials and signature and any other identifying mark as affixed to any record required by law or rule; and
- d. The date upon which such person began practicing pharmacy in the prescription drug outlet.

11.09.00 Symbols and Codes. Symbols and codes may be used to identify any manufacturer, distributor or repackager. If such symbols and codes appear in the records of a prescription drug outlet, the prescription drug outlet shall keep a current, complete printed or typed list, which shows both the symbol and code and its complete definition. This list shall be readily retrievable and available for examination by the Board for at least two years.

## **12.00.00 NUCLEAR PHARMACY.**

12.00.10 Authorized handling.. It is .unlawful for any person to provide radiopharmaceutical services unless he or she is a nuclear pharmacist acting in accordance with CRS Title 12, Article 42.5, and the rules of the Board and rules of the Colorado Department of Health, with the exception of an authorized practitioner for administration to his patients. No person may receive, acquire, possess, use, transfer or dispose of any radioactive material except in accordance with the conditions of any radioactive material license required by the Colorado Department of Health pursuant to CRS 25-11-101 *et seq.* The requirements of this rule are in addition to, and not in substitution for, other applicable provisions of rules of the Board and the State Radiation Control Agency.



- 12.00.20 Definitions.
- 12.00.21 “Nuclear prescription drug outlet” means a prescription drug outlet which deals with the preparation and delivery of radioactive material as defined in CRS 25-11-101.
- 12.00.22 “Nuclear pharmacist” means a pharmacist who holds an active pharmacist license with the Board and has met the standards of training and experience for “Authorized User Status” in handling radioactive materials in accordance with either the State Radiation Control Agency or the U.S. Nuclear Regulatory Commission.
- 12.00.23 “Radiopharmaceutical service” shall mean, but shall not be limited to, the compounding, dispensing, labeling and delivery of radiopharmaceuticals; the participation in radiopharmaceutical selection and radiopharmaceutical utilization reviews; the proper and safe storage and distribution of radiopharmaceuticals; the maintenance of radiopharmaceutical quality assurance; the responsibility for advising, where necessary or where regulated, of therapeutic values, hazards, and use of radiopharmaceuticals, and the offering or performing of those acts, services, operations or transactions necessary in the conduct, operation, management and control of radiopharmaceuticals.
- 12.00.24 “Radiopharmaceutical” is any drug which exhibits spontaneous disintegration of unstable nuclei with the emission of nuclear particles or protons and includes any such drug which is intended to be made radioactive. This definition includes non-radioactive reagent kits and nuclide generators which are intended to be used in the preparation of any such substance but does not include drugs such as carbon-containing compounds or potassium-containing salts which contain trace quantities of naturally occurring radionuclides.
- 12.00.25 “Radiopharmaceutical quality assurance” means, but is not limited to, the performance of appropriate chemical, biological and physical tests on radiopharmaceuticals and the interpretation of the resulting data to determine their suitability for use in humans and animals, including internal test assessment, authentication of product history and the keeping of proper records.
- 12.00.26 “Internal test assessment” means, but is not limited to, conducting those tests of quality assurance necessary to ensure the integrity of the test.
- 12.00.27 “Authentication of product history” means, but is not limited to, identifying the purchasing source, the ultimate fate, and intermediate handling of any component of a radiopharmaceutical.
- 12.00.28 “Authorized practitioner” means a practitioner authorized by law to possess, use and administer radiopharmaceuticals, acting within the scope of such authority.
- 12.00.30 Requirements For Nuclear Prescription Drug Outlets. A nuclear prescription drug outlet shall only be managed by a nuclear pharmacist. All personnel performing tasks in the preparation and distribution of radiopharmaceuticals shall be under the direct supervision of a nuclear pharmacist. A nuclear pharmacist shall be in attendance at all times that the nuclear prescription drug outlet is open for business and shall be responsible for all operations of the registered area.

- 12.00.31 All nuclear prescription drug outlets shall have adequate space, commensurate with the scope of services required and provided. The nuclear prescription drug outlet area shall be separate from the areas for non-radioactive drugs and shall provide a radioactive storage and product decay area separate from and exclusive of the radioactive laboratory, compounding, dispensing, quality assurance and administrative area. Prior to registration, a nuclear prescription drug outlet that wishes to compound both radiopharmaceuticals and non-radiopharmaceuticals not directly pertaining to nuclear studies shall meet the space requirements set forth in rule 5.01.31. If a nuclear prescription drug outlet wishes to compound only radiopharmaceuticals, it shall not be required to meet the space requirements set forth in rule 5.01.31. All nuclear prescription drug outlets shall submit detailing drawing-to-scale floor plans to the Board that have been approved by the state radiation control agency before approval of the registration.
- 12.00.32 There shall be a professional reference library available in the nuclear prescription drug outlet. If an electronic library is provided, workstations must be provided in a compounding/dispensing area and must be readily available for use by staff, interns and Board personnel. This library shall contain current copies of the following:
- (1) A CRS Title 12, Article 42.5; the Pharmacists, Pharmacy Businesses, and Pharmaceuticals Act;
  - (2) CRS Title 18, Article 18, the Uniform Controlled Substances Act of 1992;
  - (3) Board rules;
  - (4) 21 Code of Federal Regulations ("CFR") Part 1300 to End containing Drug Enforcement Administration rules relating to controlled substances;
  - (5) If compounding sterile products, Guide to Parenteral Admixtures or Handbook on Injectable Drugs or other comparable references as determined by the pharmacist manager;
  - (6) If compounding cytotoxic products, Technical Manual Section VI: Chapter 2, Controlling Occupational Exposure to Hazardous Drugs or ASHP Technical Assistance Bulletin on Handling Cytotoxic and Hazardous Drugs;
  - (7) Any other references that the pharmacist manager of the prescription drug outlet may deem necessary; and
  - (8) The current rules of the State Radiation Control Agency and U.S. Nuclear Commission.
- 12.00.33 A nuclear prescription drug outlet shall comply with all applicable laws and regulations of federal and state agencies, including those laws and rules governing non-radioactive drugs.
- 12.00.34 A nuclear prescription drug outlet shall have adequate equipment commensurate with the scope of radiopharmaceutical services to be provided.
- 12.00.35 Nuclear prescription drug outlets which compound and dispense only radiopharmaceuticals shall be exempt from the security requirements of rule 5.01.50 provided the following conditions are met:
- a. Only individuals identified as having "Authorized User" status by the State Radiation Control Agency or the U.S. Nuclear Regulatory Commission may enter the compounding/dispensing area in the absence of a pharmacist and only for the purpose of equipment maintenance.

- b. The nuclear prescription drug outlet maintains a written record documenting such entry detailing the following information:
  - 1) Date and time of entry;
  - 2) Authorized users name;
  - 3) Reason for entry; and
  - 4) Signature of pharmacist manager.

Such record shall be maintained on the premises and available for inspection for at least two years from the date of entry.

**12.00.40 General Requirements for Nuclear Pharmacists. A nuclear pharmacist shall:**

- a. Be a pharmacist licensed to practice in Colorado;
- b. Meet the standards of training and experience for “authorized user status” in handling of radioactive materials in accordance with either the State Radiation Control Agency or the U.S. Nuclear Regulatory Commission; and
- c. Be specifically identified, by name, as an “authorized nuclear pharmacist” on either a radioactive materials license issued by the Colorado Department of Public Health and Environment or on a U.S. Nuclear Regulatory Commission master license.

**12.00.45 Nuclear prescription drug outlets shall post, in a conspicuous area of the compounding / dispensing area of the outlet, and shall have readily available for inspection, the follow:**

- a. The original copy of the current registration with the pharmacy board;
- b. The original copy, or a reference to its specific location in the outlet, of the most current radioactive materials license issued by the Colorado Department of Public Health and Environment;
- c. A copy, or a reference to its specific location in the outlet, of the most current U.S. Nuclear Regulatory Commission master license which details a listing of its authorized nuclear pharmacists if the current radioactive license issued by the Colorado Department of Public Health and Environment references the outlet’s U.S. Nuclear Regulatory Commission master license rather than detailing a listing of the outlet’s authorized nuclear pharmacists itself; and
- d. The outlet’s current list of employees that complies with rule 11.08.00.

**12.00.64 Nuclear Compounding.**

If a nuclear pharmacist compounds a preparation according to the manufacturer’s labeling instructions, then further documentation is not required. All other compounded preparations require further documentation.

- a. No expired components may be used in compounding. No component may be used which will expire prior to the beyond-use date of the final compounded product.
- b. The compounding of sterile radiopharmaceuticals shall comply with rule 21.00.00, including all recordkeeping requirements.

- 12.00.70      Dispensing.
- a.      A radiopharmaceutical shall only be dispensed pursuant to a valid, patient-specific prescription order that is issued by an authorized practitioner.
  - b.      A nuclear prescription drug outlet shall only dispense radiopharmaceuticals which comply with acceptable standards of radiopharmaceutical assurance.
  - c.      In addition to any labeling requirement of the Board for nonradiopharmaceuticals, the immediate outer container of the radiopharmaceutical to be dispensed shall also be labeled with:
    - (1)      The standard radiation symbol;
    - (2)      The words "Caution – Radioactive Material";
    - (3)      The name of the radiopharmaceutical;
    - (3)      The amount of radioactive materials contained, in millicuries or microcuries;
    - (4)      If a liquid, the volume in milliliters;
    - (5)      The requested calibration time for the amount of radioactivity contained; and
    - (6)      Expiration data, if applicable.
  - d.      The immediate inner container shall be labeled with:
    - (1)      The standard radiation symbol;
    - (2)      The words "caution – radioactive material";
    - (3)      The assigned serial number of the corresponding prescription order; and
    - (4)      The name of the radiopharmaceutical.
  - e.      The amount of radioactivity shall be determined by radiometric methods for each individual preparation immediately prior to dispensing.
- 12.00.71      Records of Dispensing.
- a.      In addition to any requirement of the board for non-radiopharmaceutical prescription orders, the prescription order shall include the following:
    - (1)      Address of the authorized practitioner and/or the address where the prescription is to be administered;
    - (2)      The name of radiopharmaceutical;
    - (3)      The amount of radioactive materials contained, in millicuries or microcuries; and
    - (4)      Calibration time for the amount of radioactivity contained.

For the purposes of this rule, the prescription drug outlets may record the address on the order or maintain it in a readily retrievable format.

- b. A hard copy of every prescription order shall be readily retrievable and available for inspection for a period of two years from the date of any transaction relating to such order. If a nuclear prescription drug outlet dispenses only radiopharmaceuticals, prescription orders will be deemed to be readily retrievable and available if they are filed according to the date of dispensing. If a nuclear prescription drug outlet dispenses both radiopharmaceuticals and non-radiopharmaceuticals not directly pertaining to nuclear studies, all prescription orders will be deemed to be readily retrievable and available if they are filed according to the numerical sequence of the serial numbers assigned pursuant to rule 2.01.10.

**12.00.72 Distribution.**

- a. A nuclear prescription drug outlet may distribute a compounded radiopharmaceutical to a practitioner authorized by law to prescribe the drug for the purposes of administration. Such distributions shall be limited to up to 10 percent of the total number of drug dosage units dispensed and distributed on an annual basis by such outlet.
- b. A nuclear prescription drug outlet may redistribute NDA approved radiopharmaceuticals if the outlet does not process the radiopharmaceuticals in any manner or violate the product packaging.
- c. The immediate outer container of the radiopharmaceutical to be distributed shall be labeled with:
  - (1) The standard radiation symbol;
  - (2) The words "Caution – Radioactive Material";
  - (3) "RX Only";
  - (4) The name of Radiopharmaceutical;
  - (5) The amount of radioactive materials contained, in millicuries or microcuries;
  - (6) If a liquid, the volume in milliliters;
  - (7) The requested calibration time for the amount of radioactivity contained;
  - (8) Expiration data, if applicable;
  - (9) The assigned batch (lot) number;
  - (10) Specific route of administration;
  - (11) Storage directions; and
  - (12) The name and address of the prescription drug outlet.
- d. The immediate inner container shall be labeled with:
  - (1) The standard radiation symbol;
  - (2) The words "Caution – Radioactive Material";
  - (3) The assigned batch (lot) number; and

- (4) The name of the radiopharmaceutical.

**12.00.73** Records of Distribution.

- a. A nuclear prescription drug outlet shall maintain records of acquisition and distribution of all radiopharmaceuticals in accordance with CRS Title 12, and CRS Title 25.
- b. A nuclear prescription drug outlet must retain verification of each practitioner's license from the jurisdiction in which licensed on a current basis for each practitioner to whom it distributes compounded radiopharmaceuticals.
- c. A nuclear prescription drug outlet that distributes radiopharmaceuticals shall record the following:
  - (1) The name of the radiopharmaceutical;
  - (2) The amount of radioactive materials contained, in millicuries or microcuries;
  - (3) If a liquid, the volume in milliliters;
  - (4) The requested calibration time for the amount of radioactivity contained;
  - (5) The date of distribution;
  - (6) The name and address of the authorized practitioner and the address where the preparation is to be administered; and
  - (7) The name and address of the distributing outlet.
- d. Records of distribution shall be retained at the outlet for a period of not less than two years from the date of distribution.

**13.00.00** **DECLARATORY ORDERS.**

- 13.00.10** Requests. Any person may petition the Board for a declaratory order to terminate controversies or to remove uncertainties as to the applicability to the petitioner of any statutory provision or of any rule or order of the Board.

Refer to existing definition of "person" in APA, rules or statute, if any.

- 13.00.11** The Board will determine, in its discretion and without notice to petitioner, whether to rule upon any such petition. If the Board determines that it will not rule upon such a petition, the Board shall promptly notify the petitioner of its action and state the reasons for such action.

- 13.00.12** In determining whether to rule upon a petition filed pursuant to this rule, the Board will consider the following matters, among others:

- a. Whether a ruling on the petition will terminate a controversy or remove uncertainties as to the applicability to petitioner of any statutory provision or rule or order of the Board.
- b. Whether the petition involves any subject, question, or issue which is the subject of a formal or informal matter or investigation currently pending before the Board or a court but not involving any petitioner. Whether the petition seeks a ruling on a moot or hypothetical question or will result in an advisory ruling or opinion.

- c. Whether the petitioner has some other adequate legal remedy, other than an action for declaratory relief pursuant to Rule 57 Colorado Rules of Civil Procedure, which will terminate the controversy or remove any uncertainty as to the applicability to the petitioner of the statute, rule or order in question.
- 13.00.13 Any petition filed pursuant to this rule shall set forth the following:
- a. The name and address of the petitioner and whether the petitioner is licensed pursuant to the provisions of CRS 12-42.5-101, et seq., as amended, and the statute, rule, or order to which the petition relates.
  - b. A concise statement of all of the facts necessary to show the nature of the controversy or uncertainty and the manner in which the statute, rule or order in question applies or potentially applies to the petitioner.
- 13.00.20 Ruling. If the Board determines that it will rule on the petition, the following procedures apply:
- a. The Board may rule upon the petition based solely upon the facts presented in the petition. In such a case: Any ruling of the Board will apply only to the extent of the facts presented in the petition and any amended to the petition.
  - b. The Board may order the petitioner to file a written brief, memorandum or statement of position. The Board may set the petition, upon due notice to petitioner, for a non-evidentiary hearing.
  - c. The Board may dispose of the petition on the sole basis of the matters set forth in the petition.
  - d. The Board may request the petitioner to submit additional facts, in writing. In such event, such additional facts will be considered as an amendment to the petition.
  - e. The Board may take administrative notice of facts pursuant to the Administrative Procedure Act (CRS 24-4-105(8)) and may utilize its experience, technical competence and specialized knowledge in the disposition of the petition.
  - f. If the Board rules upon the petition without a hearing, it shall promptly notify the petitioner of its decision.
  - g. The Board may, in its discretion, set the petition for hearing, upon due notice to petitioner, for the purpose of obtaining additional facts or information or to determine the truth of any facts set forth in the petition or to hear oral argument on the petition. The notice to the petitioner setting such hearing shall set forth, to the extent known, the factual or other matters into which the Board intends to inquire. For the purpose of such a hearing, to the extent necessary, the petitioner shall have the burden of proving all of the facts stated in the petition, all of the facts necessary to show the nature of the controversy or uncertainty and the manner in which the statute, rule or order in question applies or potentially applies to the petitioner and any other facts the petitioner desires the Board to consider.
- 13.00.30 Parties. The parties to any proceeding pursuant to this rule shall be the Board and the petitioner. Any other person may seek leave of the Board to intervene in such a proceeding, and leave to intervene will be granted at the sole discretion of the Board. A petition to intervene shall set forth the same matters as required by section 13.00.13 of this rule. Any reference to a "petitioner" in this rule also refers to any person who has been granted leave to intervene by the Board.

13.00.40 Review. Any declaratory order or other order disposing of a petition pursuant to this rule shall constitute agency action subject to judicial review pursuant to CRS 24-4-106.

**14.00.00 OTHER OUTLETS.**

14.00.05 Eligibility for registration. The following facilities may register as other outlets provided all requirements are met:

- a. Hospitals that do not operate registered prescription drug outlets. For such hospitals, dispensing shall be limited as provided in CRS 12-42.5-118(10);
- b. Federal Qualified Health Centers, as defined in section 1861(aa)(4) of the federal "Social Security Act", 42 U.S.C. sec. 1395x(aa)(4);
- c. Family Planning Clinics;
- d. Colleges, universities and schools (grades kindergarten through twelve) which operate a school-based clinic for students and faculty of that school. Schools must submit any contractual affiliations to the Board prior to registration;
- e. Jails. A jail which obtains prescription drugs solely on the basis of individual prescription orders which have been compounded in and dispensed from a registered prescription drug outlet do not need registration;
- f. County or district public health agencies;
- g. Community and Rural Health Clinics, registered, certified, or licensed as such as by the Colorado Department of Public Health and Environment;
- h. Ambulatory Surgical Centers licensed pursuant to Part 1 of Article 3 of Title 25, C.R.S. that engage in the compounding, dispensing, and delivery of drugs or devices for administration to patients while being treated in the facility;
- i. Medical Clinics operated by a hospital that engage in the compounding, dispensing, and delivery of drugs or devices for administration to patients while being treated in the facility; and
- j. Hospices licensed pursuant to Part 1 of Article 3 of Title 25, C.R.S. that engage in the compounding, dispensing, and delivery of drugs or devices for administration to patients while being treated in the facility.
- K. Acute treatment units, registered, certified, or licensed as such by the Colorado Department of Public Health and Environment.
- L. Telepharmacies as defined pursuant to 12-42.5-102(25).

14.00.10 General Criteria. Unless otherwise exempted, the general criteria, which shall be met by other outlets herein enumerated, which are seeking to be registered by the Board pursuant to CRS 12-42.5-117(1)(d) are stated below.

- a. For the purpose of this section, the consultant pharmacist is the pharmacist responsible for the other outlet registration and the overall operation pertaining to drug receipt and distribution.



- b. Except as provided in rule 14.07.00, all prescription drugs utilized by the outlet shall be obtained from an entity or individual registered with the Board or a state or local health agency.
  - c. For the purposes of this rule, "dispensing unit" means a container or containers of a drug, either packaged pursuant to rule 3.01.00 or the manufacturer's original container(s), containing a quantity suitable for the prescribed treatment or condition.
- 14.00.20      Protocols. Written protocols shall be developed by the consultant pharmacist and submitted to the Board for approval. These protocols shall be submitted on form(s) provided by the Board and shall establish:
- a. A system of recordkeeping to document the procurement, administration, compounding, dispensing, and/or distribution, including the return to the original source, of all prescription drugs and devices, including recalled items.
  - b. A system to ensure that no drug or device shall be dispensed which will be outdated prior to utilization by the consumer, based on the practitioner's directions for use.
  - c. A system by which drugs are dispensed complying with the labeling, drug identification and container requirements imposed by law.
  - d. The duties of the consulting pharmacist.
- 14.00.30      Revisions to other outlet protocols. Revisions to other outlet protocols shall be submitted as a complete set in duplicate for approval by the Board. Prior to becoming effective, the protocol changes must be approved by the Board or its designee.
- 14.00.40      Application Procedure.
- a. Original application. Original application for registration as an other outlet shall be made on a form provided by the Board. This application shall be accompanied by the appropriate fee and two copies of the protocols.
  - b. Other outlet relocation.
    - (1) When an other outlet changes location, the outlet shall submit an application on a form provided by the Board prior to outlet relocation.
    - (2) The consultant pharmacist for the other outlet shall submit two copies of revised protocols to the Board within 30 days of relocation.
  - c. Change of ownerships of other outlet. Application to transfer registration of an other outlet shall be submitted on a form provided by the Board. This application shall be accompanied by the appropriate fee and two copies of protocols. Transfer of ownership shall be deemed to have occurred:
    - (1) In the event the other outlet is owned by a corporation, upon sale or transfer of 20 percent or more of the shares of said corporation to a single individual or entity.
    - (2) In the event the other outlet is owned by a partnership, upon sale or transfer of 20 percent or more of any ownership interest.

- (3) In the event the other outlet is owned by a limited liability company (LLC), upon sale or transfer of 20 percent or more of the membership interests.
    - (4) Upon incorporation of an existing other outlet.
  - d. Change of name of other outlet. Changes in the name of an other outlet shall be submitted to the Board on a form provided by the Board. Two copies of protocols shall be submitted to the Board within 30 days of the other outlet changing its name.
  - e. Change of consultant pharmacist.
    - (1) A new application shall be submitted to the Board within 30 days after the former consultant pharmacist ceases to be the consultant pharmacist.
    - (2) If an application is not submitted within 30 days, the other outlet registration shall become void and the Board shall be informed in writing by the person responsible for the overall operation of the other outlet of the disposition of all drug stock possessed by the other outlet.
    - (3) The other outlet registration shall be issued in the name of the consultant pharmacist. At such time as the consultant pharmacist ceases to be engaged in said position, he/she shall immediately upon knowledge thereof, notify the Board in writing. The person responsible for the overall operation of the other outlet shall immediately notify the Board in writing when the consultant pharmacist ceases to function as such.
    - (4) A pharmacist assuming the duties as a consultant pharmacist for an other outlet shall notify the Board in writing within seven days of assuming said position.
    - (5) A pharmacist assuming duties as a consultant pharmacist for an other outlet shall review the current protocols and document the review within 30 days of assuming said position. Documentation shall include the date of review and the consultant pharmacist's signature. Said documentation shall be retained with the consultant pharmacist's record of inspection or the current Board approved protocols.
  - f. Change of Registration.
    - (1) Any other outlet located in a community health clinic, rural health clinic, college, or university which dispenses more than 25,000 dispensing units in a calendar year shall register with the Board as a prescription drug outlet.
    - (2) Any other outlet located in a hospital which has greater than 25 beds as stated on its license with the Colorado Department of Public Health and Environment shall register as a prescription drug outlet.
  - g. Reinstatement of Registration. If an Other Outlet registration has expired, a registrant wishing to reinstate such registration shall submit the following:
    - (1) The current reinstatement application with the required fee; and
    - (2) Two complete and duplicate copies of written protocols, on forms provided by the Board, which are signed and dated by the individual who is the consultant pharmacist at the time the reinstatement application is submitted to the Board.

- 14.00.50 Board request that protocols be submitted. When the Board requests that protocols be submitted, the consultant pharmacist shall comply within 30 days of said request.
- 14.00.60 Registration posting. Every other outlet shall display in the primary drug storage area, or other readily accessible area, all licenses and registrations applicable to the possession and distribution of prescription drugs and controlled substances. Furthermore, every other outlet shall display in the primary drug storage area, or other readily accessible area, the report of the last inspection conducted by the Board and have readily available Board approved protocols, consultant pharmacist reports of inspections and any other documents sent by the Board to clarify or assist in the legal operation of the other outlet.
- 14.00.70 Other required registrations. The other outlet shall obtain such state and/or federal registrations as may be required.
- 14.00.80 Consultant pharmacist.
- a. A consultant pharmacist shall either:
    - (1) Initially interpret all prescription orders dispensed from the other outlet, or (2) Provide written protocols for dispensing by unlicensed persons.
  - b. A consultant pharmacist shall be available for professional consultation.
  - c. A consultant pharmacist shall annually review the protocols for compliance with this rule 14.00.00. The review shall be documented in writing, signed, and dated by the consultant pharmacist. The consultant pharmacist shall record on the protocols at least annually the number of dispensing units dispensed in a calendar year for the following facility types: community clinics, rural health clinics, colleges, and universities. A calendar year is considered to run from January 1 through December 31.
  - d. The consultant pharmacist shall develop an inspection form to document the visit and the results thereof. Such form shall be dated and signed by the consultant pharmacist and shall be maintained and available for inspection at the other outlet by the Board for a period of two years.
  - e. The consultant pharmacist shall inspect and document the inspection in writing as detailed in 14.00.80(d) the following other outlets at the following frequencies:
    - (1) Quarterly inspections and visits shall be conducted for the following:
      - (a) Jails;
      - (b) County health departments;
      - (c) Schools, grade kindergarten through twelve;
      - (d) Hospitals;
      - (e) Family planning clinics;
      - (f) Hospices;
      - (g) Medical clinics operated by hospitals; and
      - (h) Ambulatory Surgical Centers.

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- (2) Community clinics, federally qualified health centers, rural health clinics, colleges, acute treatment units, telepharmacies, and universities shall be inspected and visited as follows:
    - (a) Monthly if 2,500 or less dispensing units are dispensed in a calendar year. A calendar year is considered to run from January 1 through December 31.
    - (b) Every other week if more than 2,500 but less than 7,501 dispensing units are dispensed in a calendar year; A calendar year is considered to run from January 1 through December 31.
    - (c) Each week if 7,501 dispensing units but less than 12,501 dispensing units are dispensed in a calendar year. A calendar year is considered to run from January 1 through December 31.
    - (d) Twice each week if 12,501 dispensing units but less than 25,001 dispensing units are dispensed in a calendar year. A calendar year is considered to run from January 1 through December 31.
  - f. The consultant pharmacist shall be responsible for the accuracy of records pertaining to drug stock returned to the original supplier, the manufacturer, or via a reverse distributor. The record of any returned drug stock shall indicate, as a minimum, the name and address of the original supplier, manufacturer or reverse distributor, the date of return, and the name, strength, and quantity of the drug returned. This record shall be signed by the consultant pharmacist, and shall be maintained on the premises for a minimum of two years.
  - g. The consultant pharmacist for a licensed hospital other outlet shall be notified of any casual sale or loan of a drug made by the licensed hospital other outlet to a practitioner authorized by law to prescribe the same prior to the transaction. The consultant pharmacist for a licensed hospital other outlet shall be notified within 72 hours of any casual sale or loan of a drug to a registered other outlet, a prescription drug outlet, or a mobile emergency care unit.
  - h. The consultant pharmacist is responsible for ensuring all prescription drugs obtained by the other outlet are procured from an individual or entity registered by the Board or a state or local health agency.
  - i. The consultant pharmacist shall be responsible for ensuring any significant errors related to the practice of pharmacy, such as those that result in significant harm to a patient or the death of a patient, are immediately reported to the Board upon discovery.
  - j. The consultant pharmacist shall be responsible for assuring that the other outlet complies with all applicable provisions of Board Rule 21.00.00 when compounding non-sterile and sterile products.
  - k. The consultant pharmacist shall be responsible for reporting diversion, theft or significant unaccountable loss of prescription drugs or controlled substances from the other outlet, hospital or health maintenance organization (as defined in Section 10-16-102, C.R.S.) within one business day of discovery. When a Drug Enforcement Administration (DEA) Form 106 is submitted to the DEA in instances involving controlled substances, a copy of the completed DEA Form 106 along with a detailed written explanation shall be submitted to the Board. When determining whether an unaccountable loss is significant, the consultant pharmacist shall consider, among others factors, the following:

1. The actual quantity of drug lost in relation to the type of business;
2. The specific drug lost;
3. Whether the loss of the drug can be associated with access to those drugs by specific individuals, or whether the loss can be attributed to unique activities that may take place involving the drug;
4. A pattern of losses over a specific time period, whether the losses appear to be random, and the results of efforts taken to resolve the losses;
5. Whether the specific drug is a likely candidate for diversion; and
6. Local trends and other indicators of the diversion potential of the missing drug.

14.01.00 Interim designated consultant pharmacist. In the event the consultant pharmacist in whose name the other outlet registration is issued is unable to perform the duties of a consultant pharmacist, the consultant pharmacist shall designate an individual pharmacist to assume the consultant pharmacist's duties for no more than 90 consecutive days. The consultant pharmacist in whose name the other outlet registration is issued shall notify the Board in writing within ten days of designating an individual pharmacist to assume said consultant pharmacist's duties. Said written notification shall include, as a minimum, the name and license number of the individual pharmacist, the beginning and ending dates for which said individual pharmacist assumes the consultant pharmacist's duties, and the reason for which said individual pharmacist is designated to assume the consultant pharmacist's duties. In the event the consultant pharmacist in whose name the other outlet registration is issued is unable to perform the duties of a consultant pharmacist for a period exceeding 90 days, an application identifying a new consultant pharmacist shall be submitted to the Board no later than 30 days following the end of the original 90 day period.

14.02.00 Records and recordkeeping in other outlets.

14.02.10 Records in general. All other outlets registered and/or licensed by the Board shall maintain such records and inventories of prescription drugs as may be required by these rules or any other state or federal law or rule pertaining to such drugs. Such records shall be maintained on a current basis and shall be complete and accurate for all drugs which the outlet manufactures, receives, dispenses, distributes or otherwise disposes of in any other manner. Records and inventories of controlled substances shall be deemed to be "complete" only if each individual record and inventory contains all required information regarding each specific transaction, and if the set of records and inventories contains all information and documents required to be kept by state and federal laws, rules, and rules. A record or inventory shall be deemed to be "accurate" only if it is a complete, true and factual statement regarding or reflecting each specific transaction. A set of records or inventories shall be deemed to be "accurate" only if they are complete, and, when considered as a whole, they demonstrate that the controlled substances and/or the records and inventories pertaining thereto have been handled in compliance with all applicable laws or rules and that all such controlled substances are properly accounted for.

14.02.20 Retrievability of records. For the purposes of these rules, records and inventories shall be deemed "readily retrievable" if they meet the following requirements:

- a. For all other outlets:

- (1) The following records shall be maintained on the premises of the other outlet at all times and shall be made available for inspection by the Board or its inspectors immediately upon request.
  - (a) All DEA-222 forms executed during the two years preceding the request;
  - (b) All inventories of controlled substances required to be taken during the two years preceding the request;
  - (c) All records of dispensing, receipt (invoices for drugs received and drugs credited), distribution, loss, surrender or disposal in any other manner of prescription drugs and controlled substances during the two years preceding the request;
- (2) The following records shall be made available within 48 hours or two business days, whichever is longer, on request by the Board or its inspectors:
  - (a) All unexecuted DEA-222 forms.
- b. In the case of a request by the inspector for specific records:
  - (1) Records shall be maintained in such a manner as to permit the inspector to retrieve specific records immediately.
  - (2) If the inspector determines the records are not maintained in the manner specified in (1) above, the inspector may give the consultant pharmacist or outlet staff a list of the items to be retrieved. The requested records shall be made available to the inspector within 48 hours of the request.

14.02.30 Inventories of controlled substances. Any inventory of controlled substances shall comply with the following:

- a. If the outlet is registered with the Drug Enforcement Administration as a "hospital/clinic", the inventory shall include all drugs located throughout the facility, excluding any drug which has been dispensed pursuant to a lawful chart order but which has not yet been administered to the patient.
- b. Each inventory shall contain a complete and accurate record of all controlled substances on hand on the date the inventory is taken. The inventory shall be maintained in written, typewritten or printed form at the other outlet. Schedule II drugs shall be separated from schedule III, IV, and V drugs. Controlled substances shall be deemed to be "on hand" if they are in the possession of or under the control of the outlet. However, the inventory shall exclude any drug that has been dispensed pursuant to a lawful order but which has not yet been delivered.
- c. The inventory shall be taken either as of opening of business or as of the close of business on the inventory date and this shall be recorded on the inventory. In the event the other outlet is open 24-hours per day, the inventory shall specify the time the inventory was conducted.
- d. After the initial inventory is taken, the other outlet shall take a new inventory of all stocks of controlled substances on hand at least every two years.

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- e. On the effective date of a law or rule on which a previously non-scheduled drug is added to any schedule of controlled substances, every other outlet that possesses that drug shall take an inventory of all stocks of the drug on hand. Thereafter, that drug shall be included in each inventory made by the other outlet.
  - f. The following information shall be recorded on the inventory.
    - (1) The name of the drug;
    - (2) Each finished form of the drug (strength and dosage form);
    - (3) The number of units or volume of each finished form;
    - (4) All outdated controlled substances.
  - g. In determining the number of units of each finished form of a controlled substance in a commercial container which has been opened, the other outlet shall do as follows:
    - (1) If the drug is a schedule II drug, an exact count of the contents shall be made.
    - (2) If the substance is listed in schedule III, IV, or V, an estimated count of the measure of the contents may be made, unless the container holds more than 1000 tablets or capsules, in which case an exact count of the contents must be made.
  - h. All controlled substance inventories shall be retained at the other outlet for at least two years from the date of such inventory.
- 14.03.00 Dispensing records.
- a. At minimum, dispensing records must include the following information for every transaction:
    - (1) Unique serial number;
    - (2) Patient name;
    - (3) Prescriber;
    - (4) Date dispensed;
    - (5) Name and strength of drug dispensed;
    - (6) Quantity dispensed;
    - (7) Whether the transaction is a new or refill transaction;
    - (8) If refill transaction, the date of the initial order;
    - (9) Number of refills authorized;
    - (10) Number of refills dispensed to date;
    - (11) Identification of individual responsible for dispensing;
-

- (12) If a controlled substance, the Drug Enforcement Administration registration number of the prescriber;

Records must be current and show all dispensing transactions, new and refill.

14.03.10 Computer use for dispensing transactions. An other outlet may utilize a computer (automated data processing system) for storage and retrieval of information regarding dispensing transactions. The following requirements shall be met:

- a. All new and refill transactions shall be entered into the system at the time of the transaction, except as provided in rule 14.03.10 i..
- b. Every 24 hours, except as provided in rule 14.03.20, the system must produce a hard-copy document which, for the purposes of these rules, shall be known as the "daily printout". It shall consist of a single, uniform, complete document, except as otherwise permitted by this rule. The daily printout shall list, separately, each prescription order transaction for the previous 24 hours and shall contain all information required by this rule. Daily printouts shall be retained in a chronological manner. If the printouts are bound, the sheets shall be separated into individual pages which are then placed in the same order as printed and bound uniformly. If the pages are bound in any other manner so that they are not uniform in placement or appearance, they shall be deemed not readily retrievable and available.
- c. The daily printout shall contain all of the following information for each dispensing transaction and shall differentiate between new and refill transactions. As an alternative, new and refill transactions may be separated into two separate uniform and complete documents which contain the following:
  - (1) The serial number;
  - (2) The name of the patient;
  - (3) The name of the practitioner;
  - (4) For each controlled substance dispensed, the practitioner's Drug Enforcement Administration registration number;
  - (5) The date of issue by the practitioner. If the date is omitted by the practitioner, the date dispensed shall be presumed to be the date of issue;
  - (6) The total number of refills authorized;
  - (7) The date dispensed;
  - (8) The initials, name, or secure electronic identifier of the individual making the final evaluation;
  - (9) The name and strength of the drug dispensed;
  - (10) The quantity of the drug dispensed;
  - (11) In the case of a refill, the total number of refills dispensed to date.



- d. Records of dispensing transactions involving controlled substances shall be identifiable from those involving non-controlled substances. Alternatively, a separate complete printout listing only controlled substance transactions may be produced.
  - e. The daily printout shall be available for inspection by the Board within 72 hours from the most recent date recorded on the printout.
  - f. Documentation of the fact that the refill information entered into the automated data processing system each time a person refills an original prescription order for a schedule III, IV, or V controlled substance is correct must be provided by the individual who makes the final evaluation. This documentation may be retained in the following manner:
    - (1) If such a system provides a hard-copy printout of each day's controlled substance prescription order refill data, the controlled substance refill information shall be verified, dated, and signed by the person making the final evaluation. This individual shall verify that the date indicated is correct and then sign this document in the same manner as he/she should sign a check or legal document. This document shall be maintained in a separate file at the other outlet for a period of two years from the dispensing date. The printout of the day's controlled substance dispensing transaction must be generated by the other outlet within 72 hours of the date on which the refill was dispensed. It must be verified and signed by each person who is involved in dispensing controlled substance refills.
    - or
    - (2) The other outlet shall maintain a bound log book, or separate file, in which each person involved in dispensing controlled substance refills shall sign attesting to the fact that the refill information entered into the computer that day has been reviewed by him/her and is correct as shown. Such a book or file must be maintained at the other outlet for a period of two years after the date of dispensing the appropriately authorized refill.
  - g. The daily printout shall contain all information as required by this rule except that the identity of the person who makes the final evaluation may appear either on the daily printout or on another separate, uniformly maintained and readily retrievable record. The consultant pharmacist shall determine which of the two methods for identifying the responsible person is more appropriate for the outlet, and only that method for recording such information shall be used.
  - h. Because of the potential for a system malfunction or failure, the other outlet must have a manual procedure for recording all dispensing transactions during the system failure or malfunction. All recoverable transaction data and all manually recorded transaction data shall be entered or restored into the system within a reasonable period of time not to exceed seven days following the restoration of operation of the system.
  - i. Any automated data processing system used by an outlet shall maintain the confidentiality of records in accordance with applicable laws, rules and regulations.
- 14.03.20 Electronic maintenance of dispensing records. An other outlet which utilizes a computer (automated data processing system) for storage and retrieval of information regarding dispensing transactions need not print the daily printout required by rule 14.03.10 if the other outlet and the computer system utilized are capable of complying with the following requirements:
- a. The other outlet must be able to provide on-line retrieval of all information required by this rule for all dispensing transactions during the two years preceding the request.

- b. The other outlet must ensure a daily (i.e., every twenty-four hours) back up is performed for use in restoring required information in case of a system failure.
- c. The other outlet must:
  - (1) Have and maintain a complete on-line transaction file that is printable on the inspector's request,  
  
or
  - (2) Have a "lock-out" feature that prevents editing of dispensing information.
- d. The Board or its inspectors must be able to inspect and review the dispensing transactions of the other outlet. Therefore, immediately upon the oral or written request of the Board or its inspectors, the other outlet shall either:
  - (1) Print a report of all dispensing transactions for such period of time as the Board or its inspector(s) may specify. The system must be capable of retrieving and printing such a report within a limited time not to exceed two hours. Additionally, the system must be capable of retrieving and printing the information sorted according to variables which include, but are not limited to, date dispensed; drug name, strength and dosage form; patient name, and practitioner name;  
  
or
  - (2) Provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review dispensing transactions, and, if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. If the other outlet elects to comply with this subparagraph (d), the system must also be capable of printing the same reports described in subparagraph (1).
  - (3) It is the responsibility of the consultant pharmacist to ensure that all outlet staff are aware of the requirements of subparagraphs (1) and (2). Any failure or refusal by the consultant pharmacist and/or outlet staff to comply with a request by the Board or its inspector(s) will be deemed to be a willful violation of these rules.
- e. Whether the other outlet elects to comply with rule 14.03.20(d), the system and any reports printed on request shall contain, as a minimum, the following information for each transaction:
  - (1) The prescription order serial number;
  - (2) The name of the patient;
  - (3) The name of the practitioner;
  - (4) For each controlled substance dispensed, the practitioner's Drug Enforcement Administration registration number;
  - (5) The date of issue by the practitioner; if the date is omitted by the practitioner, the date dispensed shall be presumed to be the date of issue;
  - (6) The total number of refills authorized;

- (7) Date dispensed;
- (8) The initials or other means of identification of the individual dispensing the order;
- (9) The name and strength of the drug dispensed;
- (10) The quantity of the drug dispensed;
- (11) In the case of a refill, the total number of refills dispensed to date;
- (12) Whether the prescription order is a new or refill transaction;
- (13) In the case of a controlled substance, a means of visually identifying orders for such substances and differentiating them from non-controlled substances.

**14.04.00 Receipts.**

**14.04.10** Records of receipts of prescription drugs and controlled substances shall contain the following information for each such substance received:

- a. Name of the drug;
- b. Strength of the drug;
- c. Dosage form if appropriate;
- d. Quantity received;
- e. Date received if a controlled substance;
- f. Name of the labeler of the drug if it is labeled only with its generic name;
- g. Name of the distributor;
- h. Drug Enforcement Administration number of distributor if a controlled substance.
- i. The DEA form 222 or a copy of the DEA form 222 and the corresponding invoice shall be attached to each other.

**14.04.20** All records of receipt of prescription drugs and controlled substances shall be maintained at the other outlet for a period of time not less than two years from the date the drugs were received.

**14.04.30** All credit invoices of prescription drugs and controlled substances shall be maintained at the other outlet for a period of time not less than two years from the date of the invoice.

**14.04.40** All records of receipt of schedule II controlled substances shall be maintained separately from all other records.

**14.04.50** Records of receipt of schedule III, IV, and V controlled substances may be maintained with other records of receipt. However, the record shall be readily identifiable from the records of receipt of non-controlled drugs.

14.05.00        Distribution.

14.05.10        Records of distribution of controlled substances and prescription drugs within hospital other outlets. Records of distribution of controlled substances and prescription drugs shall comply with the following:

- a.        In a hospital which operates a registered hospital other outlet, a controlled substance or prescription drug may be distributed for floor stock to appropriate areas of the facility. A record of any such distribution shall be made and retained for a period of time not less than two years and shall include the following information:
  - (1)        The location receiving the drug;
  - (2)        The name of the drug;
  - (3)        The strength of the drug;
  - (4)        The quantity of the drug;
  - (5)        The dosage form if appropriate;
  - (6)        The date the drug was supplied;
  - (7)        The identity of the person in the prescription drug outlet who issued the drug;
  - (8)        The identity of the person who received the drug into floor stock.
- b.        These records of distribution may be retained electronically provided the following requirements are met:
  - (1)        The other outlet must ensure a daily (i.e., every twenty-four hours) back up is performed for use in restoring required information in case of a system failure.
  - (2)        The other outlet must:
    - (a)        Have and maintain a complete on-line distribution file that is printable on the inspector's request,
    - or
    - (b)        Have a "lock-out" feature that prevents editing of distribution information.
  - (3)        The Board or its inspectors must be able to inspect and review the distribution transactions of the other outlet. Therefore, immediately upon the oral or written request of the Board or its inspectors, the prescription drug outlet shall either:
    - (a)        Print a report of all distribution transactions for such period of time as the Board or its inspector(s) may specify. The system must be capable of retrieving and printing such a report within a limited time not to exceed two hours. Additionally, the system must be capable of retrieving and printing the information sorted according to variables which include, but are not limited to, date distributed, drug name, strength and dosage form;
    - or

- (b) Provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review distribution transactions, and, if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. If the other outlet elects to comply with this subparagraph (2), the system must also be capable of printing the same reports described in subparagraph (1).
- (c) It is the responsibility of the consultant pharmacist to ensure that all other outlet staff are aware of the requirements of subparagraphs (1) and (2). Any failure or refusal by the consultant pharmacist and/or outlet staff to comply with a request by the Board or its inspector(s) will be deemed to be a willful violation of these rules.
- (4) If the other outlet chooses to maintain records of distribution electronically, any reports printed upon request shall contain, as a minimum, the following information for each transaction:
  - (a) The location receiving the drug;
  - (b) The name of the drug;
  - (c) The strength of the drug;
  - (d) The quantity of the drug;
  - (e) The dosage form if appropriate;
  - (f) The date the drug was supplied;
  - (g) The identity of the person in the prescription drug outlet who issued the drug;
  - (h) The identity of the person who received the drug into floor stock.

14.05.11 A county health department registered as an other outlet may distribute prescription drugs to another registered other outlet owned or operated by that county health department. The drug shall be distributed in the original sealed container in which it was received from the wholesaler.

14.05.20 Records of distribution (casual sales) of controlled substances and prescription drugs. A hospital or county health department other outlet which distributes prescription drugs and/or controlled substances shall record the following:

- a. The name of the drug;
- b. The strength of the drug;
- c. The dosage form if appropriate;
- d. The quantity of the drug;
- e. The manufacturer name or NDC number of the labeler of the drug if labeled only with its generic name;

- f. The date of distribution;
  - g. The name, and address of the distributing outlet;
  - h. The name, and address of the receiving practitioner or registered outlet.
  - i. If a controlled substance is distributed, the record shall also indicate the Drug Enforcement registration number of the distributing outlet and the receiving practitioner or registered outlet.
  - j. A schedule II controlled substance shall only be distributed pursuant to receipt of a properly executed DEA-222 form.
- 14.05.21 These records of distribution (casual sales) required by 14.05.20 shall be retained for a period of time not less than two years from the date of the distribution.
- 14.05.22 Records of distribution (casual sales) required by rule 14.04.20 may be maintained electronically if the following requirements are met:
- a. The other outlet must ensure a daily (i.e., every twenty-four hours) back up is performed for use in restoring required information in case of a system failure.
  - b. Have and maintain a complete on-line distribution file that is printable on the inspector's request,  
  
or
  - c. Have a "lock-out" feature that prevents editing of distribution information.
  - d. The Board or its inspectors must be able to inspect and review the distribution transactions of the other outlet. Therefore, immediately upon the oral or written request of the Board or its inspectors, the other outlet shall either:
    - (1) Print a report of all distribution transactions for such period of time as the Board or its inspector(s) may specify. The system must be capable of retrieving and printing such a report within a limited time not to exceed two hours. Additionally, the system must be capable of retrieving and printing the information sorted according to variables which include, but are not limited to, date of distribution; drug name, strength and dosage form; and licensee receiving the distribution;  
  
or
    - (2) Provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review distribution transactions, and, if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. If the other outlet elects to comply with this subparagraph (2), the system must also be capable of printing the same reports described in subparagraph (1).
    - (3) It is the responsibility of the consultant pharmacist to ensure that all outlet staff are aware of the requirements of subparagraphs (1) and (2). Any failure or refusal by the consultant pharmacist and/or outlet staff to comply with a request by the Board or its inspector(s) will be deemed to be a willful violation of these rules.

- e. If the other outlet chooses to maintain records of distribution electronically, any reports printed upon request shall contain, as a minimum, the following information for each transaction:
  - (1) The name of the drug;
  - (2) The strength of the drug;
  - (3) The dosage form if appropriate;
  - (4) The quantity of the drug;
  - (5) The manufacturer name or NDC number of the labeler of the drug if labeled only with its generic name;
  - (6) The date of distribution;
  - (7) The name, and address of the distributing outlet;
  - (8) The name, and address of the receiving practitioner or registered outlet;
  - (9) If a controlled substance is distributed, the record shall also indicate the Drug Enforcement registration number of the distributing outlet and the receiving practitioner or registered outlet.

**14.05.24 Advertising.**

- a. Only one address shall appear on a prescription label and that shall be the address of the other outlet from which the prescription was dispensed.
- b. An other outlet shall only use, operate or advertise under the name that appears on the current registration issued by the Board.
- c. An other outlet may not advertise, either orally or in writing, that it is a prescription drug outlet (pharmacy).

**14.06.00 Petition for a Reduced /Inspection Schedule.**

- a. The consultant pharmacist of an other outlet may petition the Board for a reduced inspection schedule by submitting a written request to the Board detailing the procedures or technology the other outlet has in place which eliminate the need for the required frequency of inspection. The Board will review these requests in the ordinary course of business. No other outlet may change its inspection schedule without receiving written notification from the Board approving the outlet's alternative inspection schedule. Such written notification shall be maintained in the other outlet posted next to the other outlet registration.

**14.07.00 Emergency Redistribution of Prescription Drugs**

- a. In the event of a shortage of medication or state or national emergency as dictated by either the Centers for Disease Control and Prevention (CDC) or the Colorado Department of Public Health and Environment (CDPHE), an other outlet located in a county health department or public health agency as defined in CRS 25-1-502 may obtain medications from facilities, physicians, and other entities in possession of the drugs, and redistribute the medication as directed by the CDC or CDPHE. The other outlet shall not be required to become licensed as a wholesaler to conduct distribution of drugs for the limited purpose set forth in this rule. The other outlet shall maintain written records of the distributions detailing the following:
  - a. The name of the drug;
  - b. The strength of the drug;
  - c. The dosage form if appropriate;
  - d. The quantity of the drug;
  - e. Lot number of the drug;
  - f. Expiration date of the drug;
  - g. The name of the manufacturer or the NDC number of the drug if labeled only with its generic name.
  - h. The date of distribution;
  - i. The name and address of the distributing outlet;
  - j. The name and address of the receiver;
  - k. If a controlled substance is distributed, the record shall also indicate the drug enforcement administration registration number of the distributing outlet and the receiver.
  - l. A schedule II controlled substance shall only be distributed pursuant to receipt of a properly executed DEA-222 form.

**15.00.00 WHOLESALERS.**

**15.01.00 Wholesale Drugs Distributor Registration Requirement.**

- a. A wholesaler means any person engaged in the wholesale distribution of prescription drugs, including, but not limited to repackagers, own-label distributors, private-label distributors, jobbers, brokers, warehouses, including manufacturers' and distributors' warehouses; manufacturers' exclusive distributors; authorized distributors of record; drug wholesalers or distributors; independent wholesale drug traders; specialty wholesaler distributors; pharmacy buying cooperative warehouses; retail pharmacies that conduct wholesale distribution; and chain pharmacy warehouses that conduct wholesale distribution.



- b. Every wholesaler must be registered with the Board if it resides in Colorado and distributes drugs or is located in another state or territory of the United States and ships prescription drugs into Colorado.
- 15.01.10 Requirements for Licensure.
- 15.01.11 Minimum required information for registration.
- a. The following minimum information shall be required from each wholesaler as part of the registration:
    - (1) The name, full business address, and telephone number of the applicant;
    - (2) All trade or business names used by the applicant;
    - (3) Addresses, telephone numbers, and the names of contact persons for all facilities used by the applicant for the storage, handling and distribution of prescription drugs;
    - (4) The type of ownership or operation (i.e., partnership, corporation, sole proprietorship, limited liability company, or government entity); and
    - (5) The name(s) of the owner and operator of the applicant including:
      - (a) If a person, the name of the person;
      - (b) If a partnership, the name of each partner, the name of the partnership, and the federal employer identification number (FEIN);
      - (c) If a corporation, the name and title of each corporate officer and director, the name of the parent company, the corporate names, the federal employer identification number of the business, and the name of the state of incorporation; and
      - (d) Name of the business entity. If a sole proprietorship, the full name of the sole proprietor, and the name and federal employer identification number of the business entity.
      - (e) If a government entity, identify the name of director and the name of the governmental agency he/she represents.
    - (6) If a limited liability company, the name and title of each member, federal employer identification number (FEIN) of the business, and name of parent company, if any.
    - (7) A list of the licenses and permits issued to the applicant by any other state that authorizes the applicant to purchase or possess prescription drugs.
    - (8) The name of the applicant's designated representative, who must meet the following requirements:
      - (a) Be at least twenty-one years of age;

- (b) Have at least three years of full-time employment history with a pharmacy or a wholesaler in a capacity related to the dispensing and distribution of and the recordkeeping related to prescription drugs;
  - (c) Be employed by the applicant in a full-time managerial position;
  - (d) Be actively involved in and aware of the actual daily operation of the wholesaler;
  - (e) Be physically present at the facility of the applicant during regular business hours, except when the absence of the designated representative is authorized, including, but not limited to, sick leave and vacation leave;
  - (f) Serve in the capacity of a designated representative for only one applicant or wholesaler at a time, except where more than one licensed wholesaler is co-located in the same facility and the wholesalers are members of an affiliated group as defined by section 1504 of the federal "Internal Revenue code of 1986."
  - (g) Not have any convictions under federal, state, or local law relating to wholesale or retail prescription drug distribution or controlled substances;
  - (h) Not have an felony convictions pursuant to federal, state, or local law; and
  - (i) Undergo a background check as required by CRS 12-42.5-304.
- (9) Wholesalers that distribute animal drugs exclusively must have a designated representative. However, the requirements of 15.01.11a(8)(g) through-(i) are not required.
- b. Changes in any information in section 15.01.11 shall be submitted to the Board within thirty calendar days thereof.
- c. Any registered wholesale drug distributor that is accredited by a Board approved accreditation body shall inform the Board, in writing, within 72 hours if its accreditation is:
  - (1) Expired;
  - (2) Suspended;
  - (3) Revoked; or
  - (4) Withdrawn.
- d. An out-of-state wholesaler's Colorado registration shall be deemed void and shall be cancelled if the wholesaler relocates to a state other than that which is listed on its Colorado registration. In the event the wholesaler wishes to continue distributing prescription drugs into and within Colorado, it must apply for and receive a new Colorado registration indicating its current state of residence.
- e. A wholesaler's Colorado registration shall be deemed void and shall be cancelled if it was registered in Colorado using an inspection from a board-approved accreditation body and the accreditation issued by that accreditation body is revoked or withdrawn.

15.01.12 Minimum Qualifications.

- a. The Board shall consider, at a minimum, the following factors in reviewing the qualifications of persons of businesses described in 15.01.11 above who engage in the wholesale distribution of prescription drugs within the state:
  - (1) Any conviction of the applicant under any federal, state, or local laws relating to drug samples, wholesale or retail drug distribution, or distribution of controlled substances;
  - (2) Any criminal or civil convictions of the applicant under federal or state laws;
  - (3) The applicant's past experience in the manufacture or distribution of prescription drugs, including controlled substances;
  - (4) The furnishing by the applicant of false or fraudulent material in any application made in connection with drug manufacturing or distribution;
  - (5) Disciplinary proceedings by any federal, state, or local government of any registration currently or previously held by the applicant for the manufacture, distribution or dispensing of any drugs, including controlled substances;
  - (6) Compliance with registration requirements under a previously granted registration, if any;
  - (7) Compliance with requirements to maintain and/or make available to the Colorado Board of Pharmacy or other governmental agency those records required under this section; and
  - (8) Any other factors or qualifications the Board considers relevant to and consistent with the public health and safety.
- b. The Board shall have the right to deny a registration to an applicant if it determines that the granting of such a registration would not be in the public interest.
- c. All applicants shall be inspected within the previous five years prior to registration. If the applicant is located in Colorado, inspectors from the Board shall conduct the inspection. If the wholesaler is located outside of Colorado, the board of pharmacy of the state in which the wholesaler resides shall conduct an inspection of the facility or the out of state wholesaler may be inspected by a Board-approved accreditation body.
- d. The Board may suspend, revoke, refuse to renew, or otherwise discipline the registration of any wholesale drug distributor if its Board approved accreditation has been suspended, revoked, or withdrawn.

15.01.13 A wholesaler must be located at a commercial location. It may not be located in a personal dwelling or residence.

15.01.14 Change of name, location, or ownership, or designated representative.

- a. Any change in the name or location of the wholesaler shall be reported to the Board on an application provided by the Board prior to such change.

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- b. Any change in ownership shall be reported on an application provided by the Board within fifteen calendar days of the change and the new owner(s) shall apply for a new registration from the Board and pay the appropriate fee. A change of ownership shall be deemed to have occurred:
    - (1) In the event the owner is a corporation, upon sale or transfer of 20 percent or more of the shares of the corporation to a single individual or entity;
    - (2) In the event the outlet is owned by a partnership, upon sale or transfer of 20 percent or more of any ownership interest.
    - (3) In the event the outlet is owned by a limited liability company (LLC), upon sale or transfer of 20 percent or more of the membership interests.
    - (4) Upon incorporation of an existing wholesaler.
  - c. Any change in the designated representative of a wholesaler shall be reported to the Board on a form supplied by the Board within thirty calendar days of such change. The incoming designated representative must undergo the required background check.
- 15.01.17 When a wholesaler changes location, the outlet shall submit an application on a form provided by the Board prior to outlet relocation.
- 15.01.18 Reinstatement of an Expired In-State or Out-of-State Prescription Drug Wholesaler Registration.
- a. In-State Prescription Drug Wholesaler. If a registration has expired, a registrant wishing to reinstate such registration shall submit the following:
    - (1) The current reinstatement application with the required fee;
    - (2) A newly completed designated representative affidavit, on a form provided by the Board, that is signed and dated by the designated representative; and
    - (3) If a different designated representative has been established for the applicant since the expiration of the registration, the applicant shall submit the new designated representative's fingerprints to the Colorado Bureau of Investigation for both a state and federal background check at the time of submission of the reinstatement application, unless otherwise statutorily exempt or previously waived by the Board.
  - b. Out-of-State Prescription Drug Wholesaler. If a registration has expired, a registrant wishing to reinstate such registration shall submit the following:
    - (1) The current reinstatement application with the required fee;
    - (2) The applicant shall submit the designated representative's fingerprints to the Colorado Bureau of Investigation for both a state and federal background check at the time of submission of the reinstatement application, unless otherwise statutorily exempt or previously waived by the Board.
    - (3) A verification of the current prescription drug wholesaler license or registration issued by the resident state board of pharmacy;
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- (4) A newly completed designated representative affidavit, on a form provided by the Board, that is signed and dated by the designated representative; and
- (5) If the registration has expired for over 2 years, a registrant shall submit one of the following:
  - (A) A copy of a report detailing an inspection of the out-of-state prescription drug wholesaler by its resident state board of pharmacy dated within 2 years of submission of the reinstatement application; or
  - (B) A current copy of the wholesaler's accreditation by a board-approved accreditation body; or
  - (C) Proof of the wholesaler's current registration with the Federal Food and Drug Administration (FDA).

15.02.00 Personnel.

15.02.10 Designated Representative. A single person shall be designated by name and title who has complete and overall responsibility for the operation of the facility in compliance with all applicable laws rules pertaining to drugs and devices. This person's name, social security number, and title shall be reported to the Board in writing.

15.02.11 Wholesalers shall certify that all staff, employees, and personnel have suitable education or experience for the position such staff and employees hold and the job functions they are assigned. The wholesaler shall affirm that such staff has disclosed any past criminal convictions or violations of state and federal law.

15.02.12 The Designated Representative shall have overall responsibility for the operation and compliance of the facility and shall have a minimum of three years verifiable full-time experience in a pharmacy or wholesaler.

15.03.00 Sanitation.

15.03.10 Adequate sanitary and plumbing facilities shall be installed. These facilities shall be maintained in good repair and shall be regularly cleaned.

15.03.11 All areas of the facility shall be regularly and routinely cleaned. The walls, ceilings, windows and floors of the premises shall be clean and maintained in good repair and order.

15.03.12 The premises shall be free from noxious odors.

15.03.13 There shall be adequate pest control.

15.03.14 All personnel shall keep themselves and their attire as clean as possible. Facilities for storage of additional clothing and changing shall be provided as necessary and appropriate.

15.04.00 Storage.

15.04.10 All drugs shall be stored at appropriate temperatures and under appropriate conditions in accordance with requirements, if any, in the labeling of such drugs, or with requirements in the current edition of an official compendium such as the USP/NF.

- a. If no storage requirements are established for a drug, the drug may be held at “controlled” room temperature, as defined in an official compendium such as USP/NF, to help ensure that its identity, strength, quality, and purity are not adversely affected.
  - b. Appropriate manual, electromechanical, or electronic temperature and humidity equipment, and/or logs shall be utilized to document proper storage of drugs. Refrigerator and freezer units shall be monitored each business day. If done manually, the temperature shall be recorded each business day. All electromechanical or electronic temperature equipment utilized shall alert the outlet if the temperature falls out of the acceptable range.
  - c. Packaging of the drugs should be in accordance with an official compendium such as USP/NF and identify any compromise in the integrity of the drugs due to tampering or adverse storage conditions.
  - d. Controlled substance drugs should be isolated from non-controlled substance drugs and stored in a secure area in accordance with Drug Enforcement Administration security requirements and standards.
  - e. All areas of the outlet shall be well lighted and ventilated.
- 15.04.11 There shall be adequate storage space. Products that are not stored on shelving or under special conditions, such as refrigeration, shall not be stored directly on the floor.
- 15.04.12 Drugs and devices shall be placed under proper storage conditions as soon as possible after receipt.
- 15.04.13 The wholesaler shall be responsible that proper storage requirements are met for all drugs during shipment to a purchaser or other person entitled to receive such products.
- 15.05.00 Security.
- 15.05.10
- a. All facilities used for wholesale drug distribution shall be secure from unauthorized entry:
    - (1) Access from outside the premises shall be kept to a minimum and be well-controlled;
    - (2) The outside perimeter of the premises shall be well-lighted; and
    - (3) Entry into areas where drugs are held shall be limited to authorized personnel.
  - b. All facilities shall be equipped with an alarm system to detect unauthorized entry. Such alarm systems shall be both external and centrally monitored with a dedicated line and systems back up. The systems and the back up shall be regularly inspected and tested.
  - c. All facilities shall be equipped with a security system that will provide suitable protection against theft and diversion. When appropriate, the security system shall provide protection against theft or diversion that is facilitated or hidden by tampering with computers or electronic records.
  - d. All facilities shall be equipped with inventory management and control systems that detect, protect against, and document any instances of theft, diversion, or counterfeiting.

- e. All facilities shall be equipped with security systems to protect the integrity and confidentiality of data and documents and make such data and documents readily available to the Board and other state and federal law enforcement officials.
- 15.05.11 One person shall be designated by name or title, in writing, to have ultimate responsibility for security of all keys or other methods of entry into the facility itself and into all limited access areas within the facility. There shall be a list that identifies all persons who are authorized to have access to controlled substances. This information shall be made available to the Board upon request.
- 15.05.12 Storage areas shall be constructed in such a manner as to reduce the possibility of illegal entry. The wholesaler shall take adequate precautions to ensure the security of controlled substances during shipment to a purchaser or other person entitled to receive and possess controlled substances.
- 15.05.13 Any theft, suspicious loss, or recurring loss of prescription drugs or any loss of controlled substances shall be reported to the Board within thirty calendar days of the loss, along with a description of the loss, cause of the loss and any other appropriate information. Any loss of controlled substances shall also be reported to the appropriate law enforcement agency.
- 15.05.14 Any computer system used by the wholesaler shall be protected from unauthorized use.
- 15.06.00 Drug receipt, handling, and shipment.
- 15.06.10 Drugs and devices shall be placed under proper storage conditions as soon as possible after receipt.
- 15.06.11 The wholesaler shall be responsible that proper storage requirements are met for all drugs during shipment to a purchaser or other person entitled to receive such products.
- 15.06.12 Upon receipt, each shipping container shall be visually examined for identity and to determine if it may contain contaminated, contraband, counterfeit, suspected of being counterfeit or damaged drugs or drugs that are otherwise unfit for distribution. This examination shall be adequate to reveal container damage that would suggest possible contamination, adulteration, misbranding, counterfeiting, suspected of being counterfeit, or other damage to the contents.
- 15.06.13 The drugs found to be unacceptable under section 15.06.12 shall be quarantined from the rest of stock until the examination and determination that the drugs are not outdated, damaged, deteriorated, misbranded, counterfeited, or adulterated and determined to be fit for human use.
- 15.06.14 Each outgoing shipment shall be carefully inspected for identity of the drugs and to ensure that the drugs for shipment have not been damaged in storage or held under improper conditions.
- 15.06.15 Upon receipt, a wholesale distributor must review records for the acquisition of drugs for accuracy and completeness, noting the wholesale distributors involved.
- 15.06.16 The recordkeeping requirement in 15.09.00 shall be followed for all incoming and outgoing drugs and devices.

- 15.07.00        Returned drugs.
- 15.07.10        A drug which has been returned to the wholesaler shall be segregated from other stock until it can be determined if the item is salable and suitable for placement into inventory or if it is unsalable.
- 15.07.11        Any drug or device returned to a manufacturer or wholesale distributor shall be kept under proper conditions for storage, handling, transport, shipment, and documentation showing that proper conditions were maintained prior to its return is provided to the manufacturer or wholesale distributor to which the drugs are returned.
- 15.07.12        If the conditions under which a drug or device has been returned cast doubt on the drug's or device's safety, identity, strength, quality, or purity, then the drug or device shall be destroyed, or returned to the supplier, unless examination, testing or other investigation proves that the drug or device meets appropriate standards of safety, identity, strength, quality, and purity.
- 15.07.13        Any returned drug which is deemed unsalable shall be handled in accordance with the procedures delineated in rule 15.08.00.
- 15.08.00        Unsalable drugs (outdated, damaged, adulterated, misbranded, counterfeit, or suspected of being counterfeit).
- 15.08.10        Counterfeit drugs are those in which the container, shipping container, seal, or labeling, without authorization, bears the trademark, trade name, or other identifying mark, imprint, device, or any likeness thereof, of a manufacturer, processor, packer, or distributor other than the person or persons who in fact manufactured, processed, packed, or distributed such drug and which thereby falsely purports or is represented to be the product of, or to have been packed or distributed by such other manufacturer, processor, packer, or distributor.
- 15.08.11        A drug or device shall be deemed to be adulterated if:
- a.        It consists in whole or in part of any filthy, putrid, or decomposed substance; or
  - b.        It has been produced, prepared, packed, or held under unsanitary conditions whereby it may have been contaminated with filth, or whereby it may have been rendered injurious to health; or
  - c.        If the methods used in, or the facilities or controls used for, its manufacture, processing, packing, or holding do not conform to or are not operated or administered in conformity with current good manufacturing practice to assure that the drug or device meets the requirements of this part as to safety and has the identity and strength, and meets the quality and purity characteristics which it purports or is represented to possess; or
  - d.        If its container is composed, in whole or in part, of any poisonous or deleterious substance which may render the contents injurious to health; or
  - e.        If it bears or contains, for purposes of coloring only, a color additive which is unsafe within the meaning of the federal food, drug and cosmetic act.
- (1)        It is a color additive, the intended use of which is for purposes of coloring only, and is unsafe within the meaning of the federal food, drug, and cosmetic act;



- (2) If it purports to be or is represented as a drug, the name of which is recognized in an official compendium, and its strength differs from, or its quality or purity falls below, the standard set forth in the compendium. Such determination as to strength, quality, or purity shall be made in accordance with the tests or methods of assay set forth in the compendium, or in the absence of or inadequacy of these tests or methods of assay, those prescribed under the authority of the federal food, drug, and cosmetic act. No drug defined in an official compendium shall be deemed to be adulterated under this paragraph because it differs from the standard of strength, quality, or purity therefore set forth in the compendium, if its difference in strength, quality, or purity from that standard is plainly stated on its label. Whenever a drug is recognized in both the United States Pharmacopoeia and the Homeopathic Pharmacopoeia of the United States it shall be subject to the requirements of the United States Pharmacopoeia unless it is labeled and offered for sale as a homeopathic drug, in which case it shall be subject to the Homeopathic Pharmacopoeia of the United States and not those of the United States Pharmacopoeia;
- (3) If it is not subject to paragraph (2) and its strength differs from, or its purity or quality falls below, that which it purports or is represented to possess;
- (4) If it is a drug and any substance has been (a) mixed or packed therewith so as to reduce its quality or strength; or (b) substituted wholly or partially into it.

15.08.12 A drug or device shall be deemed to be misbranded if the label is false or misleading in any particular; or the label does not bear the name and address of the manufacturer, packer, or distributor, and does not have an accurate statement of the quantities of the active ingredients in case of a drug; or if the label does not show an accurate monograph for legend drugs.

15.08.13 Any unsalable drug shall be segregated in a specific area away from salable stock.

15.08.14 Any drug or device whose immediate or sealed outer or secondary containers or labeling is adulterated, misbranded, counterfeited, or suspect of being counterfeit shall be quarantined and physically separated from other drugs or devices until it is returned to either the manufacturer or wholesale distributor from which it was acquired or destroyed. When the immediate or sealed outer or secondary containers or labeling of any drug or device is adulterated, misbranded, counterfeited, or suspect of being counterfeit, notice of the adulteration, misbranding, counterfeiting, or suspected counterfeiting shall be provided to the Board, FDA, and manufacturer and wholesale distributor from which it was acquired within three (3) business days.

15.08.15 Any drug or device that has been opened or used, but is not adulterated, misbranded, counterfeited, or suspect of being counterfeit, shall be identified as such, and shall be quarantined and physically separated from other drugs or devices until they are returned to the manufacturer or wholesale distributor from which acquired or destroyed.

15.08.16 Contraband, counterfeit, or suspected to be counterfeit drugs and devices, other evidence of criminal activity, and accompanying documentation shall be retained and not destroyed until its disposition is authorized by the Board and FDA.

15.08.17 The shipping container, immediate or sealed outer or secondary container or labeling, and accompanying documentation, suspected of or determined to be counterfeit or fraudulent shall not be destroyed until its disposition is authorized by the Board and FDA.

15.08.18 An unsalable controlled substance shall be disposed of in compliance with the requirements of the drug enforcement administration and appropriate records shall be kept.

15.08.19 In the case of a drug or a device which is unsalable, records shall be kept which contain the following:

- a. The name of the drug;
- b. The strength of the drug;
- c. The dosage form if appropriate;
- d. The quantity of the drug;
- e. The name and/or NDC number of the labeler of the drug if labeled only with its generic name;
- f. If just an NDC number is used the wholesaler or manufacturer shall maintain a current, complete printed or typed list, which shows both the symbol and code and its complete definition. This list shall be readily retrievable for examination by the Board for at least three years;
- g. Method of disposition of item;
- h. Date of disposition; and
- i. Method of destruction, if applicable; and
- j. Signature of individual destroying, if applicable, and signature of individual witnessing destruction.

15.09.00 Recordkeeping.

15.09.10 All records of receipt, distribution or other disposal of prescription drugs and/or controlled substances shall be available to the Board on request for inspection, copying, verifying or other proper use. If authorization has been granted to maintain certain records centrally at another location, these records shall be made available within two business days (48 hours maximum.) Records kept at an inspection site or other site than can be immediately retrieved by computer or other electronic means shall be readily available for authorized inspection during the retention period. If recap records are available, the Board may, at its option, utilize them, but the original records must also be produced if requested and shall be considered the document of record in any case.

15.09.11 Records in general. All wholesalers registered by the Board shall maintain such records and inventories of prescription drugs as may be required by these rules or any other state or federal law or regulation pertaining to such drugs. Such records shall be maintained on a current basis and shall be complete and accurate for all drugs which the outlet manufactures, receives, distributes or otherwise disposes of in any other manner. Records, including pedigrees, and inventories of controlled substances shall be deemed to be "complete" only if each individual record and inventory contains all required information regarding each specific transaction, and if the set of records and inventories contains all information and documents required to be kept by state and federal laws, rules, and regulations. A record or inventory shall be deemed to be "accurate" only if it is a complete, true and factual statement regarding or reflecting each specific transaction. A set of records or inventories shall be deemed to be "accurate" only if they are complete, and when considered as a whole, they demonstrate that the controlled substances and/or the records and inventories pertaining thereto have been handled in compliance with all applicable laws or rules and that all such controlled substances are properly accounted for.

- a. All such records, including pedigrees, shall be retained for a period of at least three years after the date of any transaction relating to such record or inventory by any process providing an exact duplicate of the original order in a reproducible quality acceptable to the Board. Records shall be retained in a format that cannot be altered.
- b. A wholesaler in the possession of a pedigree (a document or electronic file containing information that records each distribution of any given prescription drug that leaves the normal distribution channel) for a prescription drug shall verify that each transaction on the pedigree has occurred prior to distributing the prescription drug.
- c. The pedigree shall include all necessary identifying information concerning each sale in the chain of distribution of the product from the manufacturer or first authorized distributor of record through the acquisition and sale by a wholesaler until final sale to a pharmacy or other person dispensing or administering the prescription drug. When a wholesaler distributes a product to another wholesaler, both the distributing and receiving wholesaler shall maintain a copy of the pedigree. The pedigree shall include at least the following:
  - (1) The name, address, telephone number, and, if available, the e-mail address of each owner of the prescription drug and each wholesaler of the drug;
  - (2) The name and address of each location from which the prescription drug was shipped, if different from that of the owner;
  - (3) The transaction dates;
  - (4) Certification that each recipient has authenticated the pedigree;
  - (5) The name of the prescription drug;
  - (6) The dosage form and strength of the prescription drug;
  - (7) The size and number of containers;
  - (8) The lot number of the prescription drug; and
  - (9) The name of the manufacturer of the finished dosage form.
- d. Wholesalers that distribute animal drugs exclusively are exempt from the requirements of pedigrees.

Records on an automated data processing system or subsequent storage of such records must be immediately retrievable (via monitor display or hard copy printout).

15.09.12      Retrievability of records. For the purposes of these rules, records and inventories shall be deemed "readily retrievable" if they meet the following requirements:

- a. The following records shall be maintained on the premises of the registrant at all times and shall be made available for inspection by the Board or its inspectors immediately upon request:
  - (1) All DEA-222 forms executed during the three years preceding the request;
  - (2) All inventories of controlled substances required to be taken during the three years preceding the request;

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- (3) All records of receipt (invoices for drugs received and credited) of controlled substances, distribution, loss, surrender or disposal in manner of prescription drugs and controlled substances during the three years preceding the request;
      - (4) List(s) of symbols and codes, if applicable. Symbols and codes may be used to identify any manufacturer, distributor, or repackager. If such symbols and codes appear in the records of the registrant, the registrant shall keep a current, complete printed or typed list, which shows both the symbol and code and its complete definition. This list shall be readily retrievable and available for examination by the Board for at least three years.
    - b. The following records shall be made available within 48 hours or two business days, whichever is longer, on request by the Board or its inspectors:
      - (1) All unexecuted DEA-222 forms.
      - (2) Specific records requested by the inspector if the inspector determines the records are not maintained in a readily retrievable manner.
      - (3) Records of receipt of non-controlled prescription drugs.
    - c. Pedigrees shall be made available to the board or its inspectors within five business days of request.
  - 15.09.13 Inventories of controlled substances. Any inventory of controlled substances shall comply with the following:
    - a. Each inventory shall contain a complete and accurate record of all controlled substances (including outdated controlled substances, returns from customers, and items ordered but not yet invoiced) on hand on the date the inventory is taken. The inventory shall be maintained in written, typewritten or printed form at the outlet. Schedule II drugs shall be separated from schedule III, IV, and V drugs. Controlled substances shall be deemed to be "on hand" if they are in the possession of or under the control of the registrant.
    - b. The inventory shall be taken either as of opening of business or as of the close of business on the inventory date and it shall be recorded on the inventory. In the event the prescription drug outlet is open 24 hours per day, the inventory shall specify the time the inventory was conducted.
    - c. After the initial inventory is taken, the outlet shall take new inventory of all stocks of controlled substances on hand at least every two years.
    - d. On the effective date of a law or rule on which a previously non-scheduled drug is added to any schedule of controlled substances, every prescription drug outlet that possesses that drug shall take an inventory of all stocks of the drug on hand. Thereafter, that drug shall be included in each inventory made by the outlet.
    - e. The following information shall be recorded on the inventory:
      - (1) The name of the drug;
      - (2) Each finished form of the drug (strength and dosage form);
      - (3) The number of units or volume of each finished form; and

- (4) The number of commercial containers of each finished form.
- g. All controlled substance inventories shall be retained at the prescription drug outlet for at least three years from the date of such inventory.
- 15.09.14 Receipts.
  - a. In-state prescription drug wholesalers shall only receive prescription drugs and controlled substances from an entity that is registered by the Board. This section shall not apply to intracompany or reverse distribution transactions.
- 15.09.15 Records of receipt of prescription drugs and controlled substances shall contain the following information for each such substance received:
  - a. Name of the drug;
  - b. Strength of the drug;
  - c. Dosage form if appropriate;
  - d. Quantity received;
  - e. Date received if a controlled substance;
  - f. Name of the labeler of the drug if it is labeled only with its generic name;
  - g. Name of the receiver;
  - h. Address of the receiver;
  - i. Name of the distributor that physically distributed the drug directly to the receiver;
  - j. Address of the distributor where the drug was directly distributed from;
  - k. Drug Enforcement Administration registration number of the distributor if a controlled substance;
  - l. Drug Enforcement Administration registration number of the receiver if a controlled substance;
  - m. The DEA form 222 or an electronic order form shall be completed for each schedule ii controlled substance received.
- 15.09.16 All records of receipt (including credit invoices) of prescription drugs and controlled substances shall be maintained for a period of time not less than two years from the date the drugs were received. Records of receipt (including credit invoices) of non-controlled prescription drugs need not be maintained on the premises provided that they shall be made available within 48 hours or two business days, whichever is longer, on request by the Board or its inspectors.
- 15.09.17 All records of receipt of schedule II controlled substances shall be maintained separately from all other records.
- 15.09.18 Records of receipt of schedule III, IV, and V controlled substances may be maintained with other records of receipt. However, the record shall be readily identifiable from the records of receipt of non-controlled drugs.

- 15.09.19      Distribution.
- a.      A manufacturer or wholesaler as defined in rule 15.01.00 shall furnish prescription drugs only to a person or entity licensed by the appropriate regulatory board. Before furnishing prescription drugs to a person not known to the wholesaler, the wholesaler shall affirmatively verify that the person or entity is legally authorized to receive the prescription drugs by contacting the appropriate regulatory board.
  - b.      Prescription drugs furnished by a manufacturer or wholesaler shall be delivered only to a practitioner authorized by law to prescribe the drug or to an entity licensed or registered by the Board. In the case of such entities registered or licensed by the Board, drugs shall be distributed only to the registered or licensed address. The manufacturer or wholesaler may furnish prescription drugs to an authorized person or agent of the person listed on the license if the identity and authorization of the recipient is properly established and the method of receipt is employed only to meet the immediate needs of a particular patient of the authorized person or agent.
  - c.      Prescription drugs may be furnished to a hospital pharmacy receiving area provided that a pharmacist or authorized receiving agent signs, at the time of delivery, a receipt showing the type and quantity of the prescription drug received. Any discrepancy between the receipt and the type and quantity of the prescription drug actually received shall be reported to the delivering manufacturer or wholesaler by the next business day after the delivery to the pharmacy receiving area.
- 15.09.20      Records of distribution of controlled substances and prescription drugs. An outlet which distributes prescription drugs and/or controlled substances shall record the following:
- a.      The name of the drug;
  - b.      The strength of the drug;
  - c.      The dosage form if appropriate;
  - d.      The quantity of the drug;
  - e.      The name of the manufacturer or the NDC number of the drug if labeled only with its generic name;
  - f.      The date of distribution;
  - g.      If just an NDC number is used the wholesaler or manufacturer shall maintain a current, complete printed or typed list, which shows both the symbol and code and its complete definition. This list shall be readily retrievable for examination by the Board for at least three years;
  - h.      The name and address of the distributing wholesaler;
  - i.      The name and address of the receiver;
  - j.      If a controlled substance is distributed, the record shall also indicate the Drug Enforcement Administration registration number of the distributing outlet and the receiver; and
  - k.      A schedule II controlled substance shall only be distributed pursuant to receipt of a properly executed DEA-222 form or an electronic order.

- 15.09.21 These records of distribution shall be retained for a period of time not less than two years from the date of the distribution.
- 15.09.22 Records of distribution may be maintained electronically if the following requirements are met:
- a. The wholesaler must ensure a daily (i.e., every twenty-four hours) back up is performed for use in restoring required information in case of a system failure.
  - b. Have and maintain a complete on-line distribution file that is printable on the inspector's request, or
  - c. Have a "lock-out" feature that prevents editing of distribution information.
  - d. The Board or its inspectors must be able to inspect and review the distribution transactions of the wholesaler. Therefore, immediately upon the oral or written request of the Board or its inspectors, the outlet shall either:
    - (1) Print a report of all distribution transactions for a period of time as the Board or its inspector(s) may specify. The system must be capable of retrieving and printing such a report within a limited time not to exceed two hours. Additionally, the system must be capable of retrieving and printing the information sorted according to variables which include, but are not limited to; date of distribution, drug name, strength and dosage form, and registrants receiving the distribution;
    - or
    - (2) Provide a computer terminal and monitor for the sole use of the Board or its inspector(s) to inspect and review distribution transactions, and, if necessary, provide a person to assist the Board or its inspector(s) for a period of time not to exceed two hours in operating the system. If the outlet elects to comply with this subparagraph (2), the system must also be capable of printing the same reports described in subparagraph (1)
    - (3) It is the responsibility of the manager to ensure that all wholesale staff is aware of the requirements of subparagraphs (1) and (2). Any failure or refusal by the outlet manager and/or staff to comply with a request by the Board or its inspector(s) will be deemed to be a willful violation of these rules.
  - e. If the outlet chooses to maintain records of distribution electronically, any reports printed upon request shall contain, as a minimum, the following information for each transaction:
    - (1) The name of the drug;
    - (2) The strength of the drug;
    - (3) The dosage form if appropriate;
    - (4) The quantity of the drug;
    - (5) The manufacturer name and/or NDC number of the drug if labeled only with its generic name;
    - (6) The date of distribution;

- (7) The name and address of the distributing outlet;
- (8) The name and address of the receiver; and
- (9) When a controlled substance is distributed, the record shall also indicate the drug enforcement registration number of the distributing outlet and the receiver.

15.09.23 Wholesalers shall maintain a system for the mandatory reporting of any theft, suspected theft, diversion or other loss of any prescription drug or controlled substance to the Board or as required by the Drug Enforcement Administration or other state and/or federal agencies for prescription drugs and controlled substances.

15.09.24 Records detailing losses of prescription drugs and controlled substances shall be maintained on the premises of the registrant and shall be made readily available for inspection by the Board or its inspectors immediately upon request.

15.10.00 Policies and procedures.

15.10.10 Wholesale drug distributors shall establish, maintain and adhere to written policies and procedures, which shall be followed for the receipt, security, storage, inventory, and distribution of prescription drugs, including controlled substances, and including policies and procedure for identifying, recording, and reporting destruction, losses or thefts, and for correcting all errors and inaccuracies in inventories. Wholesale drug distributors shall include the following in their written policies and procedures:

- a. A procedure whereby the oldest approved stock of a prescription drug product is distributed first. The procedure may permit deviation from this requirement, if such deviation is temporary and is itself, an approved deviation procedure.
- b. The registrant shall have a procedure to assure that any outdated stock, or any stock with an expiration date that does not allow sufficient time for dispensing by the prescription drug outlet shall be segregated from other stock and shall be returned to the manufacturer or otherwise destroyed, and documented.
- c. A procedure to be followed for handling recalls and withdrawals of prescription drugs. Such procedure shall be adequate to deal with recalls and withdrawals due to:
  - (1) Any legal action initiated at the request of the food and drug administration or other government agency with jurisdiction:
  - (2) Voluntary action by the manufacturer to remove defective or potentially defective drugs from the market:or
  - (3) Any action undertaken to promote public health and safety by the replacing of existing merchandise with an improved product or new package design.
- d. A procedure to ensure that wholesale drug distributors prepare for, protect against, and handle any crisis that affects security of operation of any facility in the event of strike, fire, flood, or other natural disaster, or other situations of local, state, or national emergency.



- e. A procedure to ensure that any outdated, misbranded, counterfeit, adulterated or unsalable prescription drugs shall be segregated from other drugs and either returned to the manufacturer or destroyed. This procedure shall provide for written documentation which shall be maintained for three years after disposition of the outdated drugs.
  - f. Policies and procedures to cover the examination of materials to include the visual inspection of shipping containers for prescription drugs unfit for distribution and prescription drugs which have been damaged in storage or held under improper conditions.
  - g. Procedures which assure employees possess the necessary education or experience for the position they hold and the job functions they are assigned.
  - h. Procedures which assure that all prescription drugs and controlled substances are only received from entities that are registered by the Board. This section shall not apply to intracompany or reverse distribution transactions.
  - i. A procedure to ensure that drugs are distributed only to individuals or entities with authorization to possess them.
  - j. A procedure to ensure that drugs are only distributed to practitioners authorized by law to prescribe the drug or to an entity licensed or registered by the Board. In the case of such entities registered or licensed by the Board, drugs shall be distributed only to the registered or licensed address. In the event the license does not show the address, a written confirmation from the regulatory board licensing or registering the individual or entity shall be obtained.
  - k. A procedure to ensure verification of all transactions on a pedigree prior to distribution of the drug.
  - l. A procedure to ensure a pedigree is furnished when distribution occurs outside of the normal distribution channel.
  - m. A procedure to ensure that staff has disclosed any past criminal convictions or violations of state and federal law.
- 15.10.11 The policies and procedures shall contain a provision for review at least annually, at which time they shall be up-dated as necessary. A record documenting this review shall be kept with the policies and procedures and shall indicate the date of completion of the review and the signature of the responsible person as defined in rule 15.02.10.
- 15.10.12 These policies and procedures and the documentation of the annual review shall be available to the Board on request for review or other proper use.
- 15.10.13 Additional requirements for wholesalers which distribute veterinary drugs directly to a person responsible for control of an animal.

- 15.10.14 A wholesaler may sell or deliver to a person responsible for the control of an animal a drug intended for veterinary use provided the following conditions are met:
- a. A licensed veterinarian has issued, prior to such sale or delivery, either a written or oral prescription order for the drug in the course of an existing, valid veterinarian-client-patient relationship. If the order is for a Schedule III, IV or V controlled substance and it is transmitted orally, it must be immediately transcribed to writing and the practitioner's written prescription order shall be transmitted to the wholesaler within three business days of the oral order;
  - b. If the order was transmitted orally, the practitioner's written prescription order shall be attached to the oral order and retained as the original order;
  - c. The drugs, prior to distribution, may not be packaged or dispensed by the registrant;
  - d. The drugs, once distributed, may not be returned to the registrant for resale or redistribution;
  - e. The prescription order issued by the veterinarian becomes void after one year if for a non-controlled drug or a schedule II controlled substance, unless the veterinarian specifies a shorter expiration date. The registrant may not distribute larger quantities than the order authorizes.
  - f. If a schedule III, IV, or V controlled substance, the prescription order becomes void after six months from date of issue, unless the veterinarian specifies a shorter expiration date. The registrant may not distribute larger quantities than the order authorizes.
  - g. The original order must be retained on the premises of the registrant filed by client name. The invoices for each distribution authorized by the order must be attached to the order.
  - h. A drug distribution log must be retained on the premises of the registrant. It shall include the following information:
    - (1) Date sold/delivered;
    - (2) Client and patient name;
    - (3) Veterinarian name;
    - (4) Veterinarian's Drug Enforcement Administration registration if a controlled substance;
    - (5) Drug sold/delivered;
    - (6) Quantity drug;
    - (7) Date of issue of order;
    - (8) Expiration of order; and
    - (9) Invoice number.

**16.00.00 LIMITED LICENSE.**

16.00.10 General Criteria. The Board may issue a limited license to the following facilities ("outlets") to purchase, possess, store and administer drugs enumerated in this Rule 16.00.00 in a manner appropriate to the outlet as authorized by law.

- A. For the purpose of the capture, sedation or immobilization of animals prior to, and including, euthanasia of injured, sick, homeless, or unwanted pets and animals:
  - 1. a humane society which is duly registered with the Secretary of State and has been in existence and in business for at least five years in this state as a nonprofit corporation; or
  - 2. an animal control agency which is operated by a unit of government.
- B. Where the employees, agents or contractors of Colorado Division of Wildlife locations are authorized by the agency to capture or immobilize wildlife for animal control, management or research purposes, those locations are considered "animal control agencies" for purposes of 12-42.5-118(17) and this rule 16.00.00.
- C. All drugs purchased, possessed, stored and administered by the Outlet shall be obtained from an individual or entity registered by the Board.

16.00.20 Application Procedure.

- a. Original Application.

Original application for registration as a limited license Outlet shall be made on a form provided by the Board.

- b. Limited License Outlet Relocation

When a limited license Outlet changes location, the Outlet shall submit an application on a form provided by the Board prior to relocation.

- c. Change of Name of Limited License Outlet.

Changes in the name of a limited license Outlet shall be submitted to the Board on a form provided by the Board.

- d. Reinstatement of Limited License.

If a registration has expired, a registrant wishing to reinstate such registration shall submit the following:

- (1) The reinstatement application that is current at the time submitted with the required fee; and
- (2) A copy of the applicant's current registration with the Drug Enforcement Administration (DEA).

- 16.00.30        Security. Outlets shall maintain limited access to controlled substances and other drug. All drugs shall be stored in locked cabinets, a safe bolted to the floor, or an equivalent secure location. Drugs shall be stored at the address registered with the Drug Enforcement Administration, or when being transported for use in the field, drug shall be secured and in the immediate possession of the employees, agents or contractors of the Outlet who are authorized by the Agency to capture or immobilize wildlife.
- 16.00.40        Training. Staff shall receive adequate training to properly administer all drugs referenced in this section.
- 16.00.50        Records in General. All outlets registered and/or licensed by the Board shall maintain such records and inventories of prescription drugs as may be required by these rules or any other state or federal law or rule pertaining to such drugs. Such records shall be maintained on a current basis and shall be complete and accurate for all drugs which the outlet manufactures, receives, dispenses, distributes or otherwise disposes of in any other manner. Records and inventories of controlled substances shall be deemed to be "complete" only if each individual record and inventory contains all required information regarding each specific transaction, and if the set of records and inventories contains all information and documents required to be kept by state and federal laws, rules, and regulations. A record or inventory shall be deemed to be "accurate" only if it is a complete, true and factual statement regarding or reflecting each specific transaction. A set of records or inventories shall be deemed to be "accurate" only if they are complete, and, when considered as a whole, they demonstrate that the controlled substances and/or the records and inventories pertaining thereto have been handled in compliance with all applicable laws or rules and that all such controlled substances are properly accounted for.
- 16.00.60        Retrievability of records. For the purposes of these rules, records and inventories shall be deemed "readily retrievable" if they meet the following requirements:
- a.        For all limited licenses:
    - (1)        The following records shall be maintained on the premises of the limited license at all times and shall be made available for inspection by the Board or its inspectors immediately upon request:
      - (a)        All official DEA 222 forms executed during the two years preceding the request;
      - (b)        All inventories of controlled substances required to be taken during the two years preceding the request;
      - (c)        All records of administration, receipt (invoices for drugs received and drugs credited), distribution, loss, surrender or disposal in any other manner of prescription drugs and controlled substances during the two years preceding the request;
    - (2)        The following records shall be made available within 48 hours or two business days, whichever is longer, on request by the Board or its inspectors:
      - (a)        All unexecuted DEA-222 forms.
  - b.        In the case of a request by the inspector for specific records:
    - (1)        Records shall be maintained in such a manner as to permit the inspector to retrieve specific records immediately.

- (2) If the inspector determines the records are not maintained in the manner specified in (1) above, the inspector may give the OUTLET a list of the items to be retrieved. The requested records shall be made available to the inspector within 48 hours of the request.

16.00.70 Inventories of controlled substances. Any inventory of controlled substances shall comply with the following:

- a. Each inventory shall contain a complete and accurate record of all controlled substances on hand on the date the inventory is taken. The inventory shall be maintained in written, typewritten or printed form at the other outlet. Schedule II drugs shall be separated from schedule III, IV, and V drugs. Controlled substances shall be deemed to be "on hand" if they are in the possession of or under the control of the outlet.
- b. The inventory shall be taken either as of opening of business or as of the close of business on the inventory date and this shall be recorded on the inventory.
- c. After the initial inventory is taken, the Outlet shall take a new inventory of all stocks of controlled substances on hand at least every two years.
- d. The following information shall be recorded on the inventory.
  - (1) The name of the drug;
  - (2) Each finished form of the drug (strength and dosage form);
  - (3) The number of units or volume of each finished form;
  - (4) All outdated controlled substances.
- e. In determining the number of units of each finished form of a controlled substance in a commercial container which has been opened, the other outlet shall do as follows:
  - (1) If the drug is a schedule II drug, an exact count of the contents shall be made.
  - (2) If the substance is listed in schedule III, IV, or V, an estimated count of the measure of the contents may be made, unless the container holds more than 1000 tablets or capsules, in which case an exact count of the contents must be made.
- f. All controlled substance inventories shall be retained at the Outlet for at least two years from the date of such inventory.

16.00.80 Records of use. Records of use of sodium pentobarbital, sodium pentobarbital in combination with other prescription drugs, or drugs used for the purposes of chemical capture or immobilization of animals or wildlife shall contain the following information:

- a. Animal or wildlife number, if available, or general description.
- b. Animal or wildlife weight, if available, or estimate.
- c. Amount of drug administered, and method if drug was administered for the purposes of chemical capture or control.
- d. Identification of individual administering drug.

- e. Amount of drug wasted (if applicable).
- f. Date administered.

Records of use shall be maintained for a period of at least two years from the date of administration.

16.01.00 Receipts.

16.01.10 Records of receipts of prescription drugs and controlled substances shall contain the following information for each such substance received:

- a. Name of the drug;
- b. Strength of the drug;
- c. Dosage form if appropriate;
- d. Quantity received;
- e. Date received if a controlled substance;
- f. Name of the labeler of the drug if it is labeled only with its generic name;
- g. Name of the distributor;
- h. Drug Enforcement Administration number of distributor if a controlled substance;
- i. The DEA form 222 or a copy of the DEA form 222 and the corresponding invoice shall be attached to each other.

16.01.20 All records of receipt of prescription drugs and controlled substances shall be maintained at the Outlet for a period of time not less than two years from the date the drugs were received.

16.01.30 All credit invoices of prescription drugs and controlled substances shall be maintained at the Outlet for a period of time not less than two years from the date of the invoice.

16.01.40 All records of receipt of schedule II controlled substances shall be maintained separately from all other records.

16.01.50 All records of receipt of schedule III, IV, and V controlled substances may be maintained with other records of receipt. However, the record shall be readily identifiable from the records of receipt of non-controlled drugs.

16.02.00 Chemical capture and sedation of animals or wildlife for euthanasia or immobilization.

16.02.01. Outlets are authorized to purchase, possess and administer drugs commonly used for the chemical capture of animals or wildlife for control, management or research purposes or to sedate or immobilize pet animals prior to euthanasia in a manner appropriate to the Outlet as authorized by law. The drugs acceptable for this use are:

- a. Acepromazine.
- b. Ketamine.

- c. Xylazine.
  - d. Tiletamine and Zolazepam.
  - e. Sodium Pentobarbital.
  - f. Butorphanol.
  - g. Azaperone.
  - h. Medetomidine.
  - i. Midazolam.
  - j. Haloperidol.
  - k. Nalbuphine.
  - l. Atipamezole.
  - m. Tolazoline.
  - n. Naltrexone.
  - o. Doxapram.
  - p. Yohimbine.
  - q. Diphenhydramine.
- 16.02.02. Outlets must maintain records of the receipt, distribution, loss, surrender and/or disposal of these drugs in the manner specified in rules 16.00.50 – 16.01.50.
- 16.02.03. Outlets must demonstrate that staff are trained and capable of using the drugs as intended. Staff must demonstrate training as follows:
- a. Certification of successful completion of the chemical immobilization workshop provided by the Law Enforcement Training Institute of the University of Missouri at Columbia, Missouri; or
  - b. Certification of successful completion of the Chemical Immobilization Workshop (the level I or III workshop) provided by the National Animal Control Association; or
  - c. Certification of successful completion of other training programs that provide at least 6 hours of didactic classroom instruction which covers animal behavior, drug delivery equipment, drug delivery, drugs for immobilization, calculating drug dosages, dosage guidelines, post immobilization procedures, emergencies, records, and laws and safety. In addition, the course must provide a minimum of two hours of field training on the use of instruments used for chemical immobilization. Credentials of instructors at these courses must demonstrate their knowledge, experience and expertise in the field of chemical immobilization of animals; and

- d. In the case of euthanasia training, the consultant or staff veterinarian must certify that staff has received thorough and adequate training on the proper administration of the medications that comply with the dosage and routes guidelines of the American Veterinary Medical Association. Furthermore, the veterinarian must certify that he/she has provided direct supervision of staff administration of such drugs for at least 3 hours prior to staff administration without supervision.

**17.00.00 [Repealed eff. 01/01/2013]**

**18.00.00 PHARMACY PEER HEALTH ASSISTANCE PROGRAM.**

**18.00.10 Definitions.**

- a. "Board" means the Colorado State Board of Pharmacy.
- b. "Board-ordered" means a Licensee or Program Participant who has been ordered by the Board to enter or complete a Program Contract pursuant to:
  - 1) a Stipulation and Final Agency Order,
  - 2) a Final Agency Order issued subsequent to an Initial Decision entered by an Administrative Law Judge following a disciplinary hearing pursuant to CRS 24-4-105, or
  - 3) a Board order entered pursuant to CRS 12-42.5-204(3) directing a Licensee to be evaluated and/or participate in the Program.
- c. "Licensee" means a person who is a pharmacist or pharmacist intern, who possess an active license issued by the Board, or has applied for licensure and paid all required fees.
- d. "PHAO" means a Peer Health Assistance Organization under contract with the Board which provides a formal, structured program that meets the requirements specified in Title 12, Article 42.5, Part2 of the Colorado Revised Statutes. Such program shall be administered by appropriate professionals for the purpose of assisting Licensees and Program Participants experiencing impaired practice to obtain evaluation, treatment, short-term counseling, monitoring of progress, and ongoing support for the purpose of arresting and treating the Licensee's or Program Participant's psychiatric, psychological, or emotional conditions or excessive alcohol or drug use or addiction.
- e. "PHAO Contract" means the contract between a Peer Health Assistance Organization ("PHAO") and the Department of Regulatory Agencies, as awarded in accordance with State law, for operation of the Pharmacy Peer Health Assistance Program.
- f. "Pharmacy Peer Health Assistance Program" means, and refers to, the Pharmacy Peek Health Assistance Diversion Program under 12-42.5-201, ET SEQ., C.R.S..
- g. "Program" means the treatment program and all associated services provided by the PHAO to the Program participant.
- h. "Program contract" means a contract between the PHAO and a program participant to detail a treatment/recovery plan or other kind of support services plan as determined necessary by the PHAO, and to provide other program services as outlined in the contract.



- i. "Program Participant" means a Licensee who is enrolled in the Program and has a Program Contract with the PHAO.
- j. "Program Participants with active cases" means those Licensees who are currently Board-ordered to receive an evaluation, treatment referral and/or monitoring with the PHAO, and those Licensees whose cases have been referred for discipline or a confidential agreement.

18.01.00 Peer Health Assistance Organizations (PHAO).

18.01.10 General Responsibilities.

Each PHAO which enters a PHAO Contract to provide Program services for the Board shall be responsible for the following:

- a. Performing assessments and evaluations of licensees who self-refer or are referred to the PHAO by the Board, and such additional evaluations and assessments as are deemed necessary by the PHAO or requested by the Board.
- b. Entering into a Program Contract with Licensees admitted into the program ("Program Participants").
- c. Informing each Program Participant of his/her rights and responsibilities under the Program Contract and the possible consequences of non-compliance.
- d. Corresponding with Program Participants regarding Board actions relevant to the Program Participants.
- e. Notifying a Program Participant and the Board of instances of noncompliance by the Program Participant or of the termination of the Program Participant from the program.
- f. Destruction of all material maintained by the PHAO three years after a Program Participant's successful completion of or termination from the program.
- g. Other duties as set forth in the PHAO Contract.

18.01.11 Quarterly Reports to the Board by PHAO's.

The PHAO shall submit compliance reports for the previous quarter during the months of April, July, October and January for participants ordered to participate in the Program. Compliance reports may include summaries of, but are not be limited to:

- a. Records of attendance by Program Participants at all prescribed therapeutic activities including, but not limited to, counseling sessions, group meetings, and drug urine screens.
- b. Records of attendance and performance from the Program Participants' supervisors/employers.
- c. Records of monitored Antabuse or other relevant prescribed medications/agents.
- d. Reports by treatment provider(s).
- e. Evaluations and assessments.

- f. Self-status reports.
  - g. Reports as required by the Program Participants' Program Contracts.
  - h. Other details as required in the PHAO Contract.
- 18.01.12 Confidentiality.
  - a. Any compliance report submitted by a PHAO to the Board regarding the progress of a Program Participant in the Program shall be reported to the Board by case number only, except as specified in paragraphs b through d below.
  - b. Whenever any Program Participant tests positive for alcohol or drugs, or otherwise chronically and/or substantially fails to comply with his/her Program Contract, the PHAO shall report the Program Participant by name to the Board.
  - c. When the PHAO reports a participant's failure to comply with the Program contract the Program Participant's treatment records and reports will no longer be kept confidential from the Board. Such reports and records shall remain confidential and be subject to protection from further disclosure pursuant to CRS 24-72-204(3)(a)(I).
  - d. The PHAO shall maintain and keep confidential a Program Participant's Program records for three years after completion of or termination from the program and then destroy them.
- 18.03.00 Program - Eligibility, Participation, Program Completion or Termination of Individual Licensees.
- 18.03.10 Program Participation.
  - a. Voluntary Participation. To be eligible for voluntary participation in the Program, a Licensee shall:
    - 1) Be a pharmacist or intern who possesses a currently active license in this state.
    - 2) Have a psychiatric, psychological or emotional condition or abuse alcohol and/or drugs in a manner which may affect the Licensee's ability to practice with reasonable skill and safety.
    - 3) Voluntarily request admission into the program.
    - 4) Agree to undergo reasonable evaluation and examination necessary for the determination of need and ability to participate in the program.
    - 5) Bear the cost of the program.
    - 6) Cooperate by providing such evaluation and treatment information, disclosure authorizations and releases of liability as may be requested by the PHAO.
    - 7) Sign a written Program Contract with the PHAO including a treatment/recovery plan in which the Licensee agrees to comply with all elements of the Program.
  - b. Mandatory Participation. A Licensee is eligible for Program participation and services if the Licensee:

- 1) Enters into a Stipulation and Final Agency Order wherein the Licensee agrees to participate in the Program as a term of disciplinary probation; or
  - 2) Is ordered into the Program for treatment pursuant to a Final Agency Order following a disciplinary hearing; or
  - 3) Is Board-ordered to enter into the Program for treatment pursuant to CRS 12-42.5-204(3).
  - 4) Has a psychiatric, psychological or emotional condition or abuses alcohol and/or drugs in a manner which may affect the Licensee's ability to practice with reasonable skill and safety.
  - 5) Bears the cost of the program.
  - 6) Cooperates by providing such evaluation and treatment information, disclosure authorizations and releases of liability as may be requested by the PHAO.
  - 7) Signs a written Program Contract with the PHAO including a treatment/recovery plan in which the Licensee agrees to comply with all elements of the Program.
- c. In the event that a previously voluntary Program Participant is subsequently Board-ordered to participate in and/or complete the Program, the Program Participant shall enter into a new Program Contract with the PHAO in which it is indicated that the Program Participant's participation in the Program was Board-ordered.

**18.03.11 Admission Procedures.**

- a. Each Licensee requesting admission into the Program shall submit an application to the PHAO.
- b. Each Program Participant will be assigned a case number by the PHAO for the purpose of confidential identification during the Program Participant's participation in the program in a manner consistent with Rule 18.01.13, below.
- c. The Program Participant shall enter into a Program Contract with the PHAO signed by Program Participant and an authorized representative of the PHAO. The Program Contract is to be kept in the confidential files of the PHAO with a copy provided to the Program Participant and any other lawfully authorized parties.
- d. The term of any Program Contract between the Program Participant and the PHAO shall be determined by the PHAO unless superseded by Board order. The term of the Program Contract may be extended and/or retroactive credit may be given at the discretion of the PHAO unless superseded by Board order.
- e. In any case where the Program Participant has been Board-ordered into the Program, the PHAO shall submit a copy of the Program Contract to the Board for inclusion in the Board's files.

**18.03.12 Reports to the Board for Non-Compliance.**

Notwithstanding any other provision of these Rules, if the PHAO determines that any applicant, Licensee, or Program Participant is unable to practice with reasonable skill and safety, the applicant, Licensee, or Program Participant shall be reported by name with supporting written documentation to the Board by the next business day.

18.03.13 Successful Discharge of a Program Participant from the Program.

A Program Participant shall be considered to have completed the Program when the Program Participant has complied with all of the terms and conditions of the Program Contract, has completed the contractual treatment program, and the PHAO has determined that the Program Participant can safely practice pharmacy without further treatment or monitoring.

18.03.14 Termination of a Program Participant from the Program.

A Program Participant may be terminated from his/her Program Contract with the PHAO for failure to comply with the treatment/recovery plan or any terms of the Program Contract with the PHAO.

**19.00.00 ADMINISTRATION.**

19.01.00 Vaccines and Immunizations.

19.01.10 Qualifications.

- a. A pharmacist certified in immunization, or pharmacy intern under the supervision of a pharmacist certified in immunization, may administer vaccines and immunizations per authorization of a physician. A copy of the authorization shall be maintained at the prescription drug outlet. Routine childhood immunizations, as defined by the Colorado State Board of Health, shall comply with CDC guidelines.
- b. Licensees shall be considered "trained" to administer vaccines and immunizations to a person only if:
  - (1) The pharmacist or pharmacy intern has completed a pharmacy-based immunization delivery course of at least 20 hours of training, including didactic and live hands-on training that is either accredited by the Accreditation Council for Pharmacy Education or provided by an ACPE accredited school or college of pharmacy as part of obtaining a pharmacy degree. Proof of completion of this training shall be posted at the pharmacist's or pharmacy intern's main practice location(s).
  - (2) The pharmacist or pharmacy intern holds a current basic cardiopulmonary resuscitation (CPR) certification issued by the American Heart Association or the American Red Cross or a basic cardiac life support certification. If the CPR certification has no expiration date, current means the certification must have been issued within the last two years. Proof of certification shall be available at pharmacist's main practice location.
  - (3) The vaccines are administered in accordance with CDC guidelines.
  - (4) The prescription drug outlet shall have a current version available, either in hard copy or electronically available, of the CDC reference "Epidemiology and Prevention of Vaccine-Preventable Diseases".

19.01.20 A trained pharmacist may delegate the administration of vaccines and immunizations only to a trained pharmacy intern.

19.01.30 Policies and Procedures

- a. Prior to administering vaccines or immunizations, pharmacists and pharmacy interns must be trained in a pharmacy-based immunization delivery course accredited as detailed in rule 19.01.10(b).
- b. The prescription drug outlet must maintain and follow written policies and procedures for handling and disposal of used and contaminated equipment and supplies. The prescription drug outlet must obtain a physician protocol for addressing allergic reactions to immunizations.
- c. The prescription drug outlet must give the appropriate "Vaccine Information Statement" (VIS) to the patient or legal representative with each dose of vaccine covered by these forms. The pharmacist must ensure that the patient or legal representative has received and signed the informed consent form and has had their questions answered prior to the administration of the vaccine.
- d. The prescription drug outlet must report adverse events as required by the Vaccine Adverse Events Reporting System (VAERS) and to the primary care provider as identified by the patient.

19.01.40 Recordkeeping.

- a. The following information must be maintained by the prescription drug outlet for three years for each dose of vaccine or immunization administered:
  - (1) The name, address, and date of birth of the patient;
  - (2) Patient responses to screening questions for indications/contraindications to the immunization or vaccine being administered;
  - (3) The date of the administration and site of injection of the immunization or vaccine;
  - (4) The name, dose, manufacturer, lot number, and expiration date of the vaccine or immunization;
  - (5) The name and address of the patient's primary health care provider as identified by the patient;
  - (6) The name or identifiable initials of the administering pharmacist. If the administration is by a pharmacy intern, the initials of both the intern and supervising pharmacist;
  - (7) The signed informed consent document for each administration;
  - (8) Which vaccine information statement (VIS) was provided;
  - (9) The date the VIS was provided; and
  - (10) The name and address of the facility at which the vaccine or immunization was administered, if administered off-site.
- b. The above records shall be maintained separately from other records of the prescription drug outlet.

- c. All records required to be maintained pursuant to this Rule 19.00.00 may be maintained electronically so long as such records are maintained in a uniform and readily retrievable manner, are printable upon request of the Board or its inspectors, and can be reviewed at a viewable rate that may customarily be reviewed when otherwise in hard-copy form.

**19.01.50 Off-Site Administration of Immunizations and Vaccines**

- a. A prescription drug outlet may allow a licensed pharmacist to remove immunizations and vaccines from the prescription drug outlet, provided the following requirements are met:
  - (1) The prescription drug outlet maintains records which detail the removal of the immunizations and vaccines with at least the following information:
    - (a) Name, strength, dosage form, and NDC number of the immunization or vaccine removed;
    - (b) Quantity removed;
    - (c) Date removed;
    - (d) Name and license number of pharmacist removing the immunization or vaccine.
  - (2) The immunizations and vaccines are properly stored at compendial temperatures during transport and storage at the off-site location.
  - (3) The vaccines and immunizations shall be secured during transport and storage at the off-site location so as to allow only licensed pharmacists and interns affiliated with the prescription drug outlet to have access to them.
  - (4) The remaining vaccines and immunizations shall be returned to the prescription drug outlet the day they were removed.
  - (5) The prescription drug outlet shall maintain records detailing the vaccines and immunizations returned with at least the following information:
    - (a) Name, strength, dosage form, and NDC number of the immunizations or vaccines returned;
    - (b) Quantity returned;
    - (c) Date returned; and
    - (d) Name and license number of pharmacist returning the immunization or vaccine.
- b. All required records shall be maintained in a manner that is uniformly maintained, readily retrievable, and available for inspection for a period of three years from the date of removal off immunizations or vaccines for off-site administration.

**20.00.00 CENTRAL PRESCRIPTION PROCESSING.**

- 20.00.10 “Central prescription processing” means the dispensing of an order when more than one registered prescription drug outlet (pharmacy) is involved in the transaction. It is the processing by one pharmacy of a request from another pharmacy to fill or refill an order or to perform one or more dispensing functions, such as preparation, mixing, labeling, initial interpretation, and refill authorizations.
- 20.00.11 “Central prescription processing contract” means a written contract which outlines the services to be provided and the responsibilities and accountabilities of each pharmacy that is a party to the central prescription processing contract in compliance with federal and state laws and rules.
- 20.00.12 “Contract pharmacy” means a pharmacy that is a party to the same central prescription processing contract as another pharmacy performing a portion of the fulfillment of a given prescription in a shared pharmacy services arrangement.
- 20.00.20 “Initial interpretation” means the review of an order accompanied by order entry. The pharmacist(s) conducting the initial interpretation shall be held accountable for the accuracy of the electronic order entry/ manual transcription and for appropriateness of therapy (e.g. known allergies, reasonable dose, duration of use, and route of administration, considering age, gender, and other patient factors; reasonable directions for use; potential or actual adverse drug reactions; drug-drug interactions; drug-food interactions; drug-disease contraindications; therapeutic duplication; proper utilization (including over- or under-utilization) and optimum therapeutic outcomes; and abuse/misuse.)
- 20.00.30 “Fulfillment” means the preparation, mixing, and placement of the ordered medication in a suitable container with appropriate labeling.
- 20.00.31 “Fulfillment pharmacy” means the pharmacy where fulfillment is completed.
- 20.00.40 “Originating pharmacy” means the pharmacy or hospital where the order is initially presented.
- 20.00.41 “Network pharmacies” means pharmacies that are under common ownership, or are parties to a central prescription processing contract, which pharmacies may perform one or more parts of the fulfillment of a given prescription.
- 20.00.43 “Shared pharmacy services” means a system that allows a common ownership or contract pharmacy to request another common ownership or contract pharmacy to conduct the initial interpretation of a prescription order or chart order.

Pharmacies participating in shared pharmacy services shall comply with all provision of this Board Rule 20.00.00 unless otherwise specifically stated in the rule.

- 20.00.50 The dispensing, delivery, and return of prescriptions by one pharmacy for another pursuant to this section shall not be construed as the filling of a transferred prescription or as a wholesale distribution.
- 20.00.60 Operational Standards.
- a. A pharmacy may outsource one or more portions of the dispensing of an order to other pharmacies provided the pharmacies:

1. Have the same owner or have entered into a central prescription processing contract; and
  2. Share a common electronic file or have appropriate technology/interface to allow access to information required to process the order; and
  3. Are registered with the Board as either prescription drug outlets or non-resident prescription drug outlets, depending on the pharmacy's location, except that a nonresident pharmacy that does not physically ship, mail or deliver dispensed prescriptions directly into this state from the nonresident pharmacy location shall be exempt from the requirement of obtaining a nonresident prescription drug outlet registration pursuant to 12-42.5-130(2). All pharmacies participating in the central prescription processing contract, or who are engaged in shared pharmacy services, must be located within the United States regardless of the requirement of a Colorado registration.
- b. The pharmacist manager of the fulfillment pharmacy shall assure that:
1. The pharmacy maintains and uses adequate storage or shipment containers and shipping processes to ensure drug stability and potency. Such shipping processes shall include the use of appropriate packaging material and/or devices to ensure that the drug is maintained at an appropriate temperature range to maintain the integrity of the medication throughout the delivery process; and
  2. The filled prescriptions are shipped in containers, which are sealed in a manner as to show evidence of opening or tampering.

**20.00.70 Notification to Patients.**

- a. Prior to the outsourcing of any portion of the dispensing process to another pharmacy that is a contract pharmacy or pharmacy under common ownership, the pharmacy shall:
1. Notify the patient that the prescription may be outsourced to the other pharmacy; and
  2. Give the name of the contract pharmacy or common ownership pharmacy. If the pharmacy is part of a network of pharmacies that may participate in dispensing the prescription, the patient shall be notified of this fact. Such notification may be provided through a one-time written notice to the patient or through use of a sign in the pharmacy.
- b. Prescription drug outlets in hospitals are exempt from this requirement.

**20.00.80 Prescription Labeling.**

- a. Prescriptions shall be labeled with all information required by CRS 12-42.5-121. In addition, the following shall be included on the label of any prescription dispensed via central processing:
1. The name and address of the originating and/or fulfillment pharmacy involved in the dispensing; and
  2. The telephone number of the pharmacy that the patient or caregiver should contact regarding refills or questions about the prescription.



20.00.90 Responsibilities of Originating Pharmacy.

- a. The originating pharmacy, when transmitting a controlled substance order to a contract or common ownership pharmacy, shall write "Central Fill" on the face of the original order and record the following:
  1. The name, and address of the pharmacy to whom the order is transmitted;
  2. The Drug Enforcement Administration registration of the pharmacy if a controlled substance order;
  3. Name of pharmacist transmitting the order; and
  4. The date of transmission.
  5. Dispensing transactions in the Shared Pharmacy Services process are exempt from the requirement of writing "Central Fill" on the face of the original order.
- b. The originating pharmacy, when transmitting a non-controlled substance order to a contract or common ownership pharmacy, shall maintain records of the following:
  1. The name, and address of the pharmacy to whom the order is transmitted;
  2. Name of pharmacist transmitting the order; and
  3. The date of transmission.
- c. Upon receipt of the prescription from the fulfillment pharmacy, the originating pharmacy shall record the following:
  1. Date of receipt;
  2. Method of delivery (private, common, or contract carrier); and
  3. Name of pharmacy employee accepting delivery.
- d. The above records shall be retained for a period not less than two years.
- e. The originating pharmacy is responsible for the maintenance of the original order in accordance with rule 11.00.00.

20.01.00 Responsibilities of Fulfillment Pharmacy.

- a. The fulfillment pharmacy shall:
  1. Retain an electronic record of all information transmitted by the originating pharmacy, including the name, address, and Drug Enforcement Administration registration (for controlled substances only) of originating pharmacy.
  2. Retain a record detailing the following:
    - i) Date the transmitted order was received;
    - ii) Identity of the pharmacist responsible for the final evaluation;

- iii) Date the order was fulfilled;
- iv) Date prescription delivered to the originating pharmacy; and
- v) The method of delivery.

20.01.10 Records.

- a. Each pharmacy shall comply with all the laws and rules relating to the maintenance of records as required by rule 11.00.00 and be able to produce an audit trail showing all prescriptions dispensed by the pharmacy.
- b. The originating pharmacy is responsible for retaining the order in the manner specified in rule 11.00.00.
- c. All involved pharmacies shall maintain appropriate records which identify the identity, date, and location of each individual performing any processing function for an order.

20.01.20 Policies and Procedures.

- a. A policy and procedure manual as it relates to central prescription processing or shared pharmacy services shall be maintained and complied with by all pharmacies involved in the dispensing of the prescriptions. This policy and procedure manual shall be readily available for inspection. The manual shall:
  - 1. Outline the responsibilities of each involved pharmacy;
  - 2. Include a list of the names, addresses, telephone numbers, and all license/registration numbers (including Drug Enforcement Administration registrations) of involved pharmacies;
  - 3. Delineate which pharmacy name and address appears on the prescription label.
  - 4. Include policies and procedures for:
    - i) Notifying patients that their prescription may be outsourced to another pharmacy for centralized prescription processing and the name of that pharmacy or pharmacies;
    - ii) Protecting the confidentiality and integrity of patient information;
    - iii) Dispensing prescriptions when the filled prescription has not been received from the fulfillment pharmacy;
    - iv) Maintaining appropriate records to identify the location and pharmacist responsible for all aspects of dispensing of any order;
    - v). Complying with federal and state laws and rules;
    - vi) Identifying the pharmacist responsible for each aspect of prescription preparation including, but not limited to, the drug regimen review, the initial electronic entry, any changes or modifications of the prescription record or patient profile, and the final evaluation of the completed prescription;

- vii) Reviewing the policy and procedure at least annually. Such review shall be done by the pharmacist manager and documented as to the date of the review accompanied by the signature of the pharmacist manager.

**21.00.00 COMPOUNDING.**

The purpose of this rule is to codify the compounding of preparations to assure that they are of acceptable strength, quality and purity.

If the pharmacist compounds a preparation according to the manufacturer's labeling instructions, then further documentation is not required. All other compounded preparations require further documentation as set forth in this rule.

Compounding of investigational products may be exempt from sections of rule 21.00.00 when compounding is restricted to utilizing ingredients that are regulated by the Federal Food and Drug Administration through an Investigational Review Board (IRB) and when the IRB- approved protocol requires deviation from this rule.

**21.00.10 Limitations and Record-Keeping.**

- a. No non-controlled substance preparation shall be compounded in advance in such quantity as may exceed a 90-day supply or is necessary to accurately compound the preparation. A 90-day supply shall be determined by the average number of dosage units dispensed or distributed of said preparation during the previous 6 month period. All prescription drug outlets shall comply with all applicable federal laws and rules pertaining to the compounding and dispensing of controlled substance preparations, including any federal laws or rules pertaining to compounding controlled substance preparations in anticipation of immediate need.
- b. No expired components may be used in compounding. No component may be used which will expire prior to the beyond-use date of the final compounded product. No expired final compounded product shall be dispensed or distributed.
- c. All records required to be maintained pursuant to this Rule 21.00.00 may be maintained electronically so long as such records are maintained in a uniform and readily retrievable manner, are printable upon request of the Board or its inspectors, and can be reviewed at a viewable rate that may customarily be reviewed when otherwise in hard-copy form.

**21.00.20 Casual Sales/Distribution of Compounded Products.**

- a. An in-state prescription drug outlet shall only distribute a compounded product to:
  - (1) Practitioners licensed and located in Colorado and authorized by law to prescribe the drug;
  - (2) Colorado licensed/registered acupuncturists, direct-entry midwives, or naturopathic doctors who are located in Colorado and authorized by law to obtain the drug;
  - (3) Hospital prescription drug outlets registered and located in Colorado; or
  - (4) Hospital other outlets registered and located in Colorado.

Except as provided by Rule 21.00.20(d), distribution of the compounded product pursuant to this rule shall be for the sole purpose of drug administration. In-state Prescription Drug Outlets shall not distribute compounded products outside of the state. In-state Prescription Drug Outlets shall dispense compounded products and ship them out of the state only pursuant to patient-specific prescription orders.

- b. Unless otherwise allowed by state and federal law, nonresident prescription drug outlets shall not distribute compounded products into Colorado pursuant to 21 U.S.C. secs. 331(a), 353(b) and 355(a).
- c. Unless otherwise allowed by state and federal law, nonresident prescription drug outlets registered in Colorado may dispense compounded products and ship them into Colorado only pursuant to valid, patient-specific prescription orders.
- d. A nonresident prescription drug outlet may distribute a compounded product to a Colorado-licensed veterinarian who is located in Colorado and authorized by law to prescribe the drug only if:
  - i) The nonresident prescription drug outlet provides the Board with a copy of the outlet's most recent report detailing an inspection by the National Association of Boards of Pharmacy Verified Pharmacy Program, for which third-party inspection the nonresident prescription drug outlet shall obtain and pay for on an annual basis, and the Board approves the inspection report as satisfactorily demonstrating proof of compliance with the Board's own inspection procedures and standards; and
  - ii) The nonresident prescription drug outlet provides the Board, on an annual basis, with a copy of the outlet's current manufacturer registration obtained from the Drug Enforcement Administration.
- e. Distribution of a compounded product to a Colorado-licensed veterinarian may be for the purpose of dispensing by the receiving veterinarian only if:
  - i) The compounded product is necessary for the treatment of a companion animal's emergency medical condition; and
  - ii) As determined by the veterinarian, the veterinarian cannot access, in a timely manner, the compounded product from a prescription drug outlet or nonresident prescription drug outlet.
- f. Except as provided under CRS 12-42.5-118(15)(a), (b)(I) and (b)(II), the amount of compounded drug product a prescription drug outlet compounds and distributes shall be no more than ten (10) percent of the total number of drug dosage units the prescription drug outlet dispenses and distributes on an annual basis, and no more than ten (10) percent of the total number of drug dosage units the nonresident prescription drug outlet dispenses and distributes into Colorado on an annual basis pursuant to Board Rules 21.00.20(d) and (e). An in-state compounding prescription drug outlet registered pursuant to CRS 12-42.5-117(9) may distribute compounded product pursuant to CRS 12-42.5-118(15)(a), (b)(I) and (II). All prescription drug outlets shall comply with all applicable federal laws and rules pertaining to the distribution of controlled substance preparations.
- g. The distributing prescription drug outlet or compounding prescription drug outlet must retain the following information on a current basis for each practitioner, hospital prescription drug outlet or hospital other outlet or, when allowable, each prescription drug outlet, to whom it distributes compounded products:

- (1) Verification of practitioner's license, or hospital prescription drug outlet's or hospital other outlet's registration; and
    - (2) Verification of practitioner's or hospital prescription drug outlet's or hospital other outlet's current Drug Enforcement Administration registration, if controlled substances are distributed;
  - h. Labeling of compounded products which are distributed shall comply with rule 21.11.10(c) or (d) or 21.21.70(c) or (d), whichever is applicable.
  - i. Records of distribution shall comply with rule 11.07.10 or 11.07.20, whichever is applicable.
- 21.00.30 Definitions. When used in this Rule 21.00.00, the following words and terms shall have the following meanings, unless the context clearly indicates otherwise.
- a. Active Pharmaceutical Ingredient (API): Chemicals, substances or other components of preparations intended for use in the diagnosis, cure, mitigation, treatment, or prevention of diseases in human or other animals or for use as dietary supplements.
  - b. Batch (Lot): Multiple units of the same compounded preparation in a single discrete process, by the same individuals, carried out during one limited time period.
  - c. Beyond-Use Date (BUD): A date after which a compounded preparation should not be stored, used or transferred and is determined from the date the preparation is compounded.
  - d. Companion animal: An animal, other than a food animal, as defined by the Colorado Board of Veterinary Medicine.
  - e. Component (ingredient): Any substance which is contained in a compounded preparation.
  - r. Compounding:
    - (1) The preparation, mixing, or assembling, of one or more active ingredients with one or more other substances, or the assembling of a finished device:
      - (a) Formulated for use on or for the patient as the result of a practitioner's prescription drug order, chart order, or initiative, based on the relationship between the practitioner, patient, and pharmacist in the course of professional practice; or
      - (b) For the purpose of, or as an incident to, research, teaching, or chemical analysis and not for sale or dispensing; or
      - (c) In anticipation of prescription orders based on routine, regularly-observed prescribing patterns.

- (2) Compounding does not include the preparation of copies of commercially available drug products. Compounded preparations that produce, for the patient, a significant difference between the compounded drug and the comparable commercially available drug product as determined, by the prescriber, as necessary for the medical best interest of the patient are not copies of commercially available products. "Significant differences" may include, but are not limited to, the removal of a dye for medical reasons (such as allergic reaction), changes in strength, and changes in dosage form or delivery mechanism. Price differences are not a "significant" difference to justify compounding.
  - g. Preparation or Product: A compounded drug dosage form, a compounded dietary supplement, or a finished device.
  - h. Quality Assurance (QA): Set of activities used to ensure that the processes used in the preparation of non-sterile or sterile drug products lead to products that meet predetermined standards of quality.
  - i. Quality Control (QC): Set of testing activities used to determine that the ingredients, components and final non-sterile or sterile drug products prepared meet pre-determined requirements with respect to strength, identity, quality, and purity.
  - j. Repackaging: The subdivision or transfer of a product from one container or device to a different container or device. Repackaging does not constitute compounding, whether or not the product being repackaged was previously compounded.
  - k. SOPS: Standard operating procedures.
  - l. Stability: Extent to which a preparation retains, within specified limits, and throughout its period of storage and use, the same properties and characteristics that it possessed at the time of compounding.
  - m. USP/NF: The current edition of the United States Pharmacopeia/National Formulary.
  - n. Validation: Documented evidence providing a high degree of assurance that specific processes will consistently produce a product meeting predetermined specifications and quality attributes.
  - o. Vehicle: A component for internal or external use that is used as a carrier or diluent in which liquids, semisolids, or solids are dissolved or suspended. Examples include, but are not limited to, water, syrups, elixirs, oleaginous liquids, solid and semisolid carriers, and proprietary products.
- 21.10.00 Compounding of Non-Sterile Products.
- 21.10.10 Policy and Procedure Manual.
- a. A manual, outlining policies and procedures encompassing all aspects of non-sterile compounding shall be available for inspection at the pharmacy. The manual shall be complied with and shall be reviewed on an annual basis. Such review shall be signed and dated by the pharmacist manager. In the event the pharmacist manager changes, the new manager shall review, sign, and date the manual within 30 days of becoming pharmacist manager. The pharmacist manager shall ensure compliance with the manual.

b. The policy and procedure manual shall address at least the following:

- (1) Responsibility of compounding personnel;
- (2) Verification of compounding accuracy;
- (3) Personnel training and evaluation in compounding skills;
- (4) Environmental quality and control;
- (5) Labeling and recordkeeping;
- (6) Finished preparation release check;
- (7) Quality control procedures, as appropriate;
- (8) Storage and beyond-use dating;
- (9) Adverse event reporting and recalls; and
- (10) Quality assurance program.

**21.10.20 Personnel Education, Training and Evaluation.**

- a. All pharmacy personnel preparing non-sterile compounded products must receive suitable training.
- b. Documentation of training of personnel shall be retained at the pharmacy and be available for inspection.

**21.10.30 Environmental Quality and Controls.**

- a. The area used for compounding shall have adequate space for the orderly placement of equipment and materials to prevent mix-ups between ingredients, containers, labels, in-process materials, and finished preparations.
- b. The compounding area shall be designed, arranged, used, and maintained to prevent adventitious cross-contamination.
- c. Non-sterile compounding areas shall be separate and distinct from any sterile compounding area.
- d. The entire compounding area is to be well-lighted. Heating, ventilation, and air conditioning systems are to be controlled to avoid decomposition of chemicals.
- e. Storage areas shall provide an environment suitably controlled to ensure quality and stability of bulk chemicals and finished preparations.
- f. All components, non-freestanding equipment, and containers shall be stored off of the floor and in a manner to prevent contamination and permit inspection and cleaning of the compounding / dispensing area.

- g. Compounding areas shall be maintained in a clean and sanitary condition. Adequate washing facilities are to be provided, including hot and cold running water, soap or detergent, and air driers or single-service towels. The plumbing system shall be free of defects that could contribute to contamination of any compounded preparation.
- h. Purified water shall be used for compounding nonsterile preparations when formulations indicate the inclusion of water. Purified water shall also be used for rinsing equipment and utensils used in compounding.
- i. Sewage, trash, and other refuse in the compounding area are to be disposed of in a safe, sanitary, and timely manner.
- j. Special precautions shall be taken to clean equipment and compounding areas meticulously after compounding preparations that contain allergenic ingredients.

**21.10.40 Equipment.**

- a. Equipment shall be of appropriate design and capacity, and be operated within designed operational limits.
- b. Equipment shall be of suitable composition so that surfaces that contact components, in-process material, or drug products shall not be reactive, additive, or absorptive so as to alter the safety, identity, strength, quality, or purity of the drug product beyond the desired result.
- c. Appropriate cleaning processes shall be in place to insure cleanliness of equipment.
- d. Written procedures outlining required equipment, calibration, appropriate maintenance, monitoring for proper function, controlled procedures for use of the equipment and specified time frames for these activities shall be established and followed. Results of equipment calibration and appropriate maintenance reports shall be kept on file at the outlet for at least two years from the report date. These results shall be available for inspection.

**21.10.60 Components.**

- a. Compounding personnel shall ascertain that ingredients for compounded products are in compliance with rule 21.00.10(b) and are of the correct identity and appropriate quality using the following information: vendors' labels, labeling, certificates of analysis, direct chemical analysis, and knowledge of compounding facility storage conditions. No expired components may be used in compounding. No component may be used which will expire prior to the beyond-use date of the preparation.
- b. Ingredients used in a compounded preparation shall either originate from FDA-approved sources, when available, or be USP/NF grade substances, when such sources are not available and identified on the FDA drug shortage list.
- c. If neither USP/NF grade substances nor FDA-approved substances are available, or when food, cosmetics, or other substances are, or must be used, the substance shall be of a chemical grade in one of the following categories:
  - (1) Chemically Pure (CP);
  - (2) Analytical Reagent (AR); or



- (3) American Chemical Society (ACS); or
- (4) Food Chemical Codex.
- d. For all ingredients, unless FDA-approved, the pharmacist shall establish purity and stability by obtaining a certificate of analysis from the supplier. The certificate of analysis, when applicable, shall be maintained at the prescription drug outlet for at least two years from the date of preparation.
- e. For components that do not have expiration dates assigned by the manufacturer or supplier, a pharmacist shall clearly and legibly label the container with the date of receipt and assign a conservative expiration date, not to exceed three (3) years after receipt, to the component based on the nature of the component and its degradation mechanism, the container in which it is packaged, and the storage conditions. A pharmacist shall clearly and legibly label the container with the assigned expiration date. In no event shall the labeled date of receipt or assigned expiration date be later altered after originally labeling the container.
- f. A manufactured drug product may be a source of active ingredient. Only manufactured drugs from containers labeled with a lot number and an expiration date are acceptable as a potential source of active ingredients. When compounding with manufactured drug products, the compounder must consider all ingredients present in the drug product relative to the intended use of a compounded preparation.
- g. Drug preparations that have been withdrawn or removed from the market for safety reasons shall not be compounded. Such preparations may be compounded exclusively for veterinary use provided no documentation exists which indicates that the preparation is unsafe for such use.
- h. Any ingredient regulated by the FDA through an Investigational Review Board (IRB) is exempt from rule 21.10.60 provided the research requirements for the receipt of the ingredient is followed and meets the requirements of CRS 12-42.5-128(2).

**21.10.65 Packaging and Drug Preparation Containers**

- a. Pharmacy personnel shall ensure that the containers and container closures used in the packaging of compounded preparations meet all applicable USP requirements and, when available, compounding monographs.
- b. The containers and closures shall be made of suitable clean material in order not to alter the quality, strength, or purity of the compounded preparation in any way.
- c. The containers and closures shall be stored off of the floor, handled and stored to prevent contamination, and rotated so that the oldest stock is used first. The containers and closures shall be stored in such a way as to permit inspection and cleaning of the compounding / dispensing area.

21.10.70 Finished Preparation Release Checks.

a. Physical Inspection

- (1) Written procedures for physical inspection of compounded preparations shall be followed. Immediately after compounding, and prior to dispensing or distribution, each product shall be inspected for evidence of particulates or other foreign matter, container-closure integrity, and any other apparent visual defect. Defective product shall be segregated from other product and shall not be dispensed or distributed.

b. Compounding Accuracy Checks

- (1) Written procedures for double-checking compounding accuracy shall be followed for every compounded product during preparation and immediately prior to release. Outlets which compound shall have at least the following written procedures for verifying the correct identity and quality of compounded products prior to dispensing or distribution:
  - (a) Verification of label for accuracy; and
  - (b) Correct identities, purities, and amounts of ingredients have been used by comparing the original written order to the written compounding record for the compounded product.

21.10.80 Storage and Beyond-Use Dating.

- a. Completed compounded preparations that are not immediately dispensed or distributed shall be stored according to the guidelines in the formulation record.

- b. In the absence of stability information that is applicable to the lowest and highest dose or concentration of a specific preparation compounded at the outlet, the following maximum beyond-use dates are to be used for non-sterile compounded preparations that are packaged in tight, light-resistant containers and stored at controlled room temperature unless otherwise indicated.

- (1) For non-aqueous liquids and solid formulations
  - (a) Where the manufactured drug product is the source of the active ingredient, the beyond-use date shall not exceed 25% of the time remaining until the product's expiration date or 6 months, whichever is earlier;
  - (b) Where a USP/NF substance is the source of active ingredient, the beyond-use date shall not be greater than 6 months;
- (2) For water-containing oral formulations prepared from ingredients in solid form, regardless of whether an ingredient contains water or water by itself is an ingredient, the beyond-use date shall not be greater than 14 days when stored at cold temperatures;
- (3) For intranasal formulations, the beyond-use date shall not be greater than 30 days;

- (4) For all other formulations, including topical, dermal, mucosal, liquids and semi-solid formulations, the beyond-use date shall not be greater than the intended duration of therapy or 90 days;
- (5) The beyond-use date limits may be exceeded when there is supporting valid scientific stability information that is directly applicable to the specific preparation. This information shall be retained on-site at the outlet and be available for inspection.

**21.10.90 Formulation Record.**

- a. For each compounded preparation, a uniform, readily retrievable formulation record shall be maintained and available for inspection for two years from the date last utilized, documenting:
  - (1) The official or assigned name, strength, dosage form, and route of administration of the compounded preparation;
  - (2) Calculations needed to determine and verify quantities or concentrations of components and doses of APIs;
  - (3) All ingredients and their quantities;
  - (4) Compatibility and stability information, including references when available;
  - (5) The equipment used to compound the preparation;
  - (6) Mixing instructions that shall include:
    - (a) order of mixing;
    - (b) mixing temperatures or other environmental controls;
    - (c) duration of mixing; and
    - (d) other factors pertinent to the replication of the preparation as compounded;
  - (7) Sample labeling information which shall include, in addition to other required information;
    - (a) generic name and quantity or concentration of each API;
    - (b) assigned BUD;
    - (c) storage conditions; and
    - (d) assigned prescription or control number, whichever is applicable;
  - (8) The assigned BUD;
  - (9) The containers used in dispensing;
  - (10) Packaging and storage requirements;

- (11) Physical description of final product; and
- (12) Procedures for quality control, if applicable.

**21.11.00 Compounding Record.**

- a. For each compounded product prepared, a record shall be maintained and available for inspection for two years on the original order, or on a separate, uniform, and readily retrievable record documenting the following:

- (1) The official or assigned name and strength of the compounded preparation;
- (2) Formulation record reference for the preparation;
- (3) Names and corresponding quantities of all components used in the preparation;
- (4) Sources, lot numbers, and expiration dates of each component;
- (5) Total number of dosage units compounded;
- (6) Name of the person who compounded the preparation;
- (7) Name of the pharmacist who approved the preparation;
- (8) Batch (lot) number assigned, if multiple units compounded;
- (9) Date prepared;
- (10) Assigned BUD;
- (11) Assigned prescription number(s) or control number(s), whichever is applicable;
- (12) Storage conditions;
- (13) Physical description of the final product;
- (14) Results of quality control procedures, if applicable; and
- (15) Documentation of any quality control issues and any adverse reactions or preparation problems reported by the patient or caregiver.

**21.11.10 Labeling of Non-Sterile Compounded Preparations.**

- a. Labeling of non-sterile compounded products dispensed pursuant to a prescription order or LTCF chart order shall include at least the following:

- (1) All requirements of CRS 12-42.5-121;
- (2) Batch (lot) number, if appropriate;
- (3) Assigned BUD;
- (4) Storage directions when appropriate; and

- (5) A clear statement that this product was compounded by the pharmacy, except for radiopharmaceuticals prepared from FDA-approved, commercially available kits and/or drug products.
- b. Labeling of non-sterile compounded products dispensed pursuant to a hospital chart order shall include at least the following:
  - (1) All requirements of CRS 12-42.5-121;
  - (2) Batch (lot) number, if appropriate;
  - (3) Assigned BUD; and
  - (4) Storage directions, when appropriate.
- c. Labeling of non-sterile compounded products distributed to practitioners, other prescription drug outlets, or other outlets allowed by law or made in anticipation of orders shall include at least the following:
  - (1) Name and address of the outlet;
  - (2) Name and strength of the drug(s) / active ingredient(s) in the final product;
  - (3) Total quantity in package;
  - (4) Assigned BUD;
  - (5) Batch (lot) number;
  - (6) Specific route of administration;
  - (7) Storage directions, when appropriate;
  - (8) "Rx only"; and
  - (9) "This product was compounded by the pharmacy", except for radiopharmaceuticals prepared from FDA-approved, commercially available kits and/or drug products.
- d. Labeling of non-sterile compounded products distributed within hospitals as floor stock shall include at least the following:
  - (1) Name of the outlet;
  - (2) Name and strength of the drug(s);
  - (3) Total quantity in package;
  - (4) Quantity of active ingredient in each dosage unit;
  - (5) Assigned BUD;
  - (6) Batch (lot) number;
  - (7) Specific route of administration; and

- (8) Storage directions, if appropriate.

**21.11.20 Patient Monitoring, Adverse Events Reporting, and Product Recall.**

- a. Outlets which compound shall provide patients and other recipients of compounded preparations with a way to address their questions and report any concerns that they may have with these preparations.
- b. The outlet shall have written policies describing specific instructions for receiving, acknowledging; and for recording, or filing, and evaluating reports of adverse events and of the quality of preparation claimed to be associated with compounded preparations.
- c. The pharmacist manager shall report to the Board in writing significant errors related to compounded preparations such as those that result in serious personal injury or death of a patient.
- d. If a compounded preparation is believed to be defective in any way, or if the Board or Federal Food and Drug Administration makes a written request for a recall of a specific preparation, the outlet shall immediately recall any product dispensed or distributed. Any product remaining in the outlet shall be immediately quarantined and shall not be dispensed or distributed. Recall records shall include at least the following:
  - (1) Product name, strength, dosage form;
  - (2) Reason for recall;
  - (3) Amount of product made;
  - (4) Date made; and
  - (5) Amount of product dispensed or distributed.
- e. The outlet shall conduct tests, as appropriate, on the recalled product to identify reason product was defective. Results of these tests shall be retained at the outlet.
- f. Adverse event reports and product recall records shall be retained and available for inspection at the outlet for at least two years.

**21.20.00 Compounding of Sterile Products (CSPs).**

**21.20.10 Definitions.** In addition to the definitions set forth above in rule 21.00.30, when used in these rules 21.20.00 et seq., 21.21.00 et seq. and 21.22.00 et seq., the following words and terms shall have the following meanings, unless the context clearly indicates otherwise.

- a. **Anteroom:** An ISO Class 8 (Class 100,000) or better area where personnel perform hand hygiene and garbing procedures, staging of components, order entry, labeling, and other activities which generate particulates. It is a transition area that provides assurance that air flows from clean to dirty areas.
- b. **Aseptic Processing:** A mode of processing pharmaceutical and medical products that involves the separate sterilization of the product and of the packaging and the transfer of the product into the container and its closure under at least ISO Class 5 conditions.

- c. Biological Safety Cabinet (BSC): A ventilated containment unit for personnel, product, and environmental protections having an open front with inward airflow for personnel protection, downward HEPA filtered laminar airflow for product protections, and HEPA filtered exhausted air for environmental protections.
- d. Buffer Area: An ISO Class 7 (Class 10,000), or an ISO Class 8 for the preparation of radiopharmaceuticals, area where the primary engineering control is physically located. Activities conducted in this area include the preparation and staging of components and supplies when compounding sterile products. This area may also be referred to as a buffer or core room, buffer or cleanroom areas, buffer room area, buffer or clean area.
- e. Class 100 Environment (ISO Class 5): An atmospheric environment which contains less than one hundred (100) particles 0.5 microns in diameter per cubic foot of air, according to federal standards.
- f. Class 10,000 Environment (ISO Class 7): An atmospheric environment which contains less than ten thousand (10,000) particles 0.5 microns in diameter per cubic foot of air, according to federal standards.
- g. Class 100,000 Environment (ISO Class 8): An atmospheric environment which contains less than one hundred thousand (100,000) particles 0.5 microns in diameter per cubic foot of air according to federal standards.
- h. Clean Room: A room in which the concentration of airborne particles is controlled to meet a specified airborne particulate cleanliness class. Microorganisms in the environment are monitored so that a microbial level for air, surface, and personnel is not exceeded for a specified cleanliness class.
- i. Compounding Aseptic Containment Isolator (CACI): A compounding aseptic isolator (CAI) designed to provide worker protection from exposure to undesirable levels of airborne drug throughout the compounding and material transfer process and to provide an aseptic environment for compounding sterile preparations. Air exchange with the surrounding environment should not occur unless the air is first passed through a microbial retentive filter (HEPA minimum) system capable of containing airborne concentrations of the physical size and state of the drug being compounded. Where volatile drugs are prepared, the exhaust air from the isolator should be appropriately removed by properly designed building ventilation.
- j. Compounding Aseptic Isolator (CAI): A closed system made up of solid walls, an air-handling system, and transfer and interaction devices. The walls are constructed so as to provide surfaces that are cleanable with covering between wall junctures. The air-handling system provides HEPA filtration of inlet air. Transfer of materials is accomplished through air locks, glove rings, or ports. Transfers are designed to minimize the entry of contamination. Manipulations can take place through either glove ports or half suits. A barrier isolator is designed for compounding sterile products. It is designed to maintain an aseptic compounding environment within the isolator throughout the compounding and material transfer process. Air exchange into the isolator from the surrounding environment should not occur unless it has first passed through a HEPA filter.
- k. Compounded Sterile Products (CSPs): A sterile drug or nutrient compounded in a registered prescription drug outlet or other outlet. Such products may include, but are not limited to, implants, injectables, parenteral nutrition solutions, irrigation solutions, inhalation solutions, intravenous solutions and ophthalmic preparations.

- l. Critical Area: An ISO Class 5 environment.
- m. Critical Sites: Include sterile ingredients of CSPs and locations on devices and components used to prepare, package, and transfer CSPs that provide opportunity for contamination.
- n. Cytotoxic Drugs: A pharmaceutical product that has the capability of direct toxic action on living tissue that can result in severe leucopenia and thrombocytopenia, depression of the immune system and the alteration of a host's inflammatory response system.
- o. Disinfectant: An agent that frees from infections. It is usually a chemical agent but sometimes a physical one. It destroys disease-causing pathogens or other harmful microorganisms but may or may not kill bacterial spores. It refers to substances applied to inanimate objects.
- p. High-Efficiency Particulate Air (HEPA) filter: A filter composed of pleats of filter medium separated by rigid sheets of corrugated paper or aluminum foil that direct the flow of air forced through the filter in a uniform parallel flow. HEPA filters remove 99.97% of all particles three-tenths (0.3) microns or larger. When HEPA filters are used as a component of a horizontal or vertical-laminar-airflow workbench, an environment can be created consistent with standards for a class 100 clean room.
- q. Media-Fill Test: A test which is used to qualify aseptic technique of compounding personnel or processes and to ensure that the processes used are able to produce sterile product without microbial contamination. A microbiological growth medium such as soybean-casein digest medium (SCDM) is substituted for the actual drug product to simulate admixture compounding.
- r. Multiple-Dose Container: A multiple-unit container for articles or preparations intended for parenteral administration only. These containers usually contain antimicrobial preservatives. The beyond-use date (BUD) for an opened or entered multi-dose container with antimicrobial preservatives is 28 days, unless otherwise specified by the manufacturer.
- s. Parenteral: A sterile preparation of drugs for injection through one or more layers of skin.
- t. Pharmacy Bulk Package: A container of a sterile preparation for parenteral use that contains multiple single doses. The contents of the package are intended for use in a pharmacy admixture program and are restricted to the preparation of admixtures for infusion or, through a sterile transfer device, for the filling of empty sterile syringes. The closure shall be penetrated only one time after constitution with a suitable sterile transfer device or dispensing set, which allows measured dispensing of the contents. The pharmacy bulk package is to be used only in a suitable work area such as a laminar flow hood or an equivalent clean air compounding area. Such container shall be labeled with the following:
  - (1) The name, strength and quantity of drug or base solution;
  - (2) The statement "Pharmacy Bulk Package—Not For Direct Infusion;"
  - (3) Information on the proper technique to assure safe use of the product; and
  - (4) A statement limiting the time frame in which the container may be used once it has been entered, provided it is held under the labeled storage conditions.



- u. Primary Engineering Control (PEC): A device or room that provides an ISO Class 5 environment for the exposure of critical sites when compounding CSPs. Such devices include, but are not limited to, laminar airflow workbenches (LAFWs), biological safety cabinets (BSCs) and compounding aseptic isolators (CAIs), and compounding aseptic containment isolators (CACIs).
- v. Process Validation or Simulation: Microbiological simulation of an aseptic process with growth medium processed in a manner similar to the processing of the product and with the same container or closure system.
- w. Segregated Compounding Area: A part of the designated compounding / dispensing area that is a specifically designated space, either a demarcated area or room, and that is restricted to preparing low-risk level CSPs with a 12-hour or less BUD. Such area shall contain a device that provides unidirectional airflow of ISO Class 5 air quality for preparation of CSPs and shall be void of activities and materials that are extraneous to sterile compounding.
- x. Single-Dose Container: A single-unit container for articles or preparations intended for parenteral administration only. It is intended for single use and is labeled as such. Examples include, but are not limited to, prefilled syringes, cartridges, fusion-sealed containers, and closure-sealed containers when so labeled.
- y. Sterile Pharmaceutical: A dosage form free from living microorganisms.
- z. Sterilization: A validated process used to render a product free of viable organisms.
- aa. Sterilizing Grade Filter Membranes: Filter membranes that are documented to retain 100% of a culture of 10<sup>7</sup> microorganisms of a strain of *Brevundimonas* (*Pseudomonas*) *diminuta* per square centimeter of membrane surface under a pressure of not less than 30 psi (2.0 bar). Such filter membranes are nominally at 0.22 or 0.2 micrometer porosity, depending on the manufacturer's practice.
- bb. Sterilization by Filtration: Passage of a fluid or solution through a sterilizing grade filter to produce a sterile effluent.
- cc. Terminal Sterilization: The application of a lethal process (e.g. steam under pressure or autoclaving) to sealed containers for the purpose of achieving a predetermined sterile assurance level of usually less than 10<sup>-6</sup>, or a probability of less than one in one million of a non-sterile unit.
- dd. Temperatures:
  - 1. Frozen means temperatures between twenty five degrees below zero and ten degrees below zero Celsius (-25 and -10 degrees C.) or thirteen degrees below zero and fourteen degrees Fahrenheit (-13 and 14 degrees F.).
  - 2. Refrigerated means temperatures between two and eight degrees Celsius (2 and 8 degrees C.) or thirty-six and forty-six degrees Fahrenheit (36 and 46 degrees F.).
  - 3. Room temperatures mean room temperatures between fifteen and thirty degrees Celsius (15 and 30 degrees C.) or fifty-nine and eighty-six degrees Fahrenheit (59 and 86 degrees F.).

- ee. Unidirectional Flow: An airflow moving in a single direction, in a robust and uniform manner, and at sufficient speed to reproducibly sweep particles away from the critical processing or testing area.

**21.20.16 Allergen Extracts as CSPs**

- a. Allergen extracts as CSPs are single dose and multiple dose intradermal or subcutaneous injections and are not subject to the environmental and storage requirements for CSP Risk Levels provided they are compounded in accordance with the most recent USP <797> guidelines.
- b. The compounding process involves simple transfer of commercial sterile allergen products and shall contain appropriate substances in effective concentrations to prevent the growth of microorganisms.
- c. The label of each multiple dose container of allergen extracts as CSPs shall list the name of one specific patient and a BUD and storage temperature range based on manufacturers' recommendations or peer-review publications available at the outlet for inspection.
- d. Single dose allergen extracts as CSPs shall not be stored for subsequent use.

**21.20.20 Definitions of Sterile Compounded Products by Risk Level.**

- a. Immediate Use CSPS:
  - (1) Immediate use CSPS are intended only for emergency or immediate patient administration of a CSP, and are exempt from the requirements for low-risk CSPS if:
    - (a) The compounding process involves a transfer of not more than three (3) commercially manufactured sterile nonhazardous products from the manufacturers' original containers and not more than two (2) entries into any one (1) container;
    - (b) The compounding process takes less than one (1) hour;
    - (c) Aseptic technique is followed when compounding occurs outside of class 5 air quality;
    - (d) Product administration begins no later than one (1) hour after product preparation; and
    - (e) The product is labeled with a one (1) hour BUD.
- b. Low Risk CSPs;
  - (1) All low risk CSPs shall be compounded with aseptic manipulations within ISO Class 5 or better air quality. All PECs (CAI, CACI, LAFW, and BSC) shall be certified as required and shall maintain ISO Class 5 air quality.
  - (2) Low risk CSPs with greater than 12-hour BUD: Applies to compounding sterile products that exhibit characteristics (a) and (b) stated below. The products shall be prepared with sterile equipment, sterile ingredients and solutions and sterile contact surfaces for the final product. Low risk includes the following:

- (a) The compounding involves only transfer, measuring, and mixing manipulations using no more than three commercially manufactured sterile products and no more than two entries into any one container or package of sterile product to make the CSP; and
    - (b) Manipulations are limited to aseptically opening ampules, penetrating sterile stoppers on vials with sterile needles and syringes, and transferring sterile liquids in sterile syringes to sterile administration devices, package containers of other sterile products, and containers for storage and dispensing.
  - (3) Low risk CSPs with 12-hour or less BUD: The LAFW or BSC shall not be located within an ISO Class 7 area. Applies to low-risk, nonhazardous or radiopharmaceutical CSPs dispensed pursuant to a patient-specific order which are prepared and administered within 12 hours of the preparation or as stated in the corresponding manufacturer's package insert (whichever is less) and the following conditions are met:
    - (a) PECs shall be in a segregated compounding area restricted to sterile compounding activities that minimize the risk of CSP contamination;
    - (b) The segregated compounding area shall not be in a location that has unsealed windows or doors that connect to the outdoors or high traffic flow, or that is adjacent to construction sites, warehouses, or food preparation or any area that could cause contamination. The segregated area shall not be located next to a sink; and
    - (c) The specifications in cleaning and disinfecting the sterile compounding area, personnel training and competency evaluation of garbing, aseptic work practices and cleaning/disinfection procedures, and viable and non-viable environmental sampling testing shall be followed.
    - (d) This shall not apply to chemotherapeutic preparations subject to USP/NF Chapter 800.
- c. Medium Risk CSPs: Sterile products exhibit characteristics (1), (2), or (3) stated below. When CSPs are compounded aseptically under low risk conditions, and one or more of the following conditions exists, such CSPs are at a medium risk level of contamination:
  - (1) Multiple individual or small doses of sterile products are combined or pooled to prepare a CSP that will be administered either to multiple patients or to one patient on multiple occasions; or
  - (2) The compounding process includes complex aseptic manipulations other than the single volume transfer; or
  - (3) The compounding process requires unusually long duration, such as that required to complete dissolution or homogeneous mixing.
- d. High Risk CSPs: CSPs compounded under any of the following conditions are either contaminated or at high risk to become contaminated with infectious microorganisms:
  - (1) Products compounded from non-sterile ingredients or compounded with non-sterile components, containers or equipment before terminal sterilization; or

- (2) Sterile contents of commercially manufactured products, CSPs that lack effective antimicrobial preservatives, and sterile surfaces of devices and containers for the preparation, transfer, sterilization, and packaging of CSPs are exposed to air quality worse than ISO Class 5 for more than 1 hour; or
- (3) Before sterilization, non-sterile procedures such as weighing and mixing are conducted in air quality worse than ISO Class 7, compounding personnel are improperly garbed and gloved; or water-containing preparations are stored for more than 6 hours; or
- (4) It is assumed, and not verified by examination of labeling and documentation from suppliers or by direct determination, that the chemical purity and content strength of ingredients meet their original or compendial specifications in unopened or in opened packages of bulk ingredients.

**21.20.23 Single-Dose and Multiple-Dose Containers.**

- a. Opened or needle-punctured single-dose containers shall be used within 1 hour if opened in worse than ISO Class 5 air quality. Single-dose containers exposed to ISO Class 5 air quality or cleaner air may be used up to 6 hours after initial puncture.
- b. If multiple-dose containers include antimicrobial preservatives, the BUD shall not exceed 28 days from the initial date of entering or opening, unless otherwise specified by the manufacturer.

**21.20.25 Radiopharmaceuticals as CSPs.**

- a. Production of radiopharmaceuticals for positron emission tomography (PET) shall comply with the most current Chapter 823 of the USP/NF < Radiopharmaceuticals for Positron Emission >.
- b. Radiopharmaceuticals shall be compounded in conformity with rules 21.20.25(b)(1) through (4) below, rule 12.00.00, and all other applicable sections of rule 21.00.00.
  - (1) Radiopharmaceuticals compounded from FDA-approved, commercially sterile components in closed sterile containers and with a volume of 100 ml or less for a single-dose injection or not more than 30 ml taken from a multiple-dose container shall be designated as, and conform to, the standards for low risk CSPs.
  - (2) Radiopharmaceuticals shall be compounded using appropriately shielded vials and syringes in a properly functioning and certified ISO Class 5 PEC located in an ISO Class 8 or cleaner air environment to permit compliance with special handling, shielding, and negative air flow requirements.
  - (3) Radiopharmaceutical vials designated for multiple use, compounded with technetium-99m, exposed to an ISO Class 5 environment, and punctured by needles with no direct contact contamination may be used up to the time indicated by the manufacturer's recommendations.

- (4) Technetium-99m/molybdenum-99 generator systems shall be stored and operated under conditions recommended by the manufacturer and applicable state and federal rules. Such generator systems shall be operated in an ISO Class 8 or cleaner air environment to permit special handling, shielding, and air flow requirements. To limit acute and chronic radiation exposure of inspecting personnel to a level that is as low as reasonably achievable (ALARA), direct visual inspection of radiopharmaceutical CSPs containing high concentrations of doses of radioactivity shall be conducted in accordance with ALARA.

**21.20.30 Policy and Procedure Manual.**

- a. A manual, outlining policies and procedures encompassing all aspects of compounding low, medium or high risk products, shall be available for inspection at the pharmacy. This manual shall be complied with and shall be reviewed on an annual basis. Such review shall be signed and dated by the pharmacist manager. In the event the pharmacist manager changes, the new manager shall review, sign, and date the manual within 30 days of becoming pharmacist manager. The pharmacist manager shall ensure compliance with the manual.
- b. The policy and procedure manual shall address at least the following:
  - (1) Responsibility of compounding personnel;
  - (2) Verification of compounding accuracy and sterilization;
  - (3) Personnel training and evaluation in aseptic manipulation skills;
  - (4) Environmental quality and control;
  - (5) Aseptic processing;
  - (6) Labeling and recordkeeping;
  - (7) Finished preparation release check;
  - (8) Storage and beyond-use dating;
  - (9) Maintaining product quality and control during transportation and delivery after the CSP leaves the pharmacy;
  - (10) Patient or caregiver training;
  - (11) Adverse event reporting and recalls;
  - (12) Quality assurance program;
  - (13) Quality control procedures, as appropriate; and
  - (14) Verification of work area cleaning effectiveness.

**21.20.40 Personnel Education and Training.**

- a. Low risk: All pharmacy personnel preparing sterile products must receive suitable didactic and experiential training.

- b. Medium risk: In addition to low risk requirements, personnel training includes assessment of competency in all medium risk procedures.
- c. High risk: In addition to low and medium risk requirements, operators have specific education, training and experience to prepare high risk products. The pharmacist knows principles of good compounding practice for risk level products, including:
  - (1) Aseptic processing;
  - (2) Quality assurance of environmental, component, and end-product testing;
  - (3) Sterilization; and
  - (4) Selection and use of containers, equipment, and closures.

**21.20.50 Personnel Evaluation in Aseptic Manipulation Skills.**

- a. Personnel who prepare CSPs shall be provided appropriate training before they begin preparing CSPs.
- b. Compounding personnel shall perform didactic review and pass written and media-fill testing of aseptic manipulative skills initially; at least annually thereafter for low and medium risk products; and every six months, thereafter, for high risk products.
- c. Personnel who fail written tests, or whose media-fill test vials result in gross microbial colonization, must be immediately reinstructed and re-evaluated by expert compounding personnel to ensure correction of all aseptic practice deficiencies.
- d. Results of these tests shall be retained and be available for inspection at the outlet for at least two years.

**21.20.60 Environmental Quality and Controls.**

- a. All CSPs shall be compounded in air quality of a Class 100 (ISO Class 5) environment or better.
- b. For the compounding of non-radiopharmaceuticals, all primary engineering controls shall be placed in a buffer area that is of air quality Class 10,000 (ISO Class 7) or better. If the PEC is a CAI or a CACI that provides isolation from the room and maintains ISO Class 5 conditions during dynamic operating conditions, including transferring ingredients, components, and devices into and out of the isolator and during preparation of CSPs, then it is not required to be placed in an ISO Class 7 buffer area. For the compounding of radiopharmaceuticals, all primary engineering controls shall be placed in a buffer area that is of air quality Class 100,000 (ISO Class 8) or better.
- c. The surfaces of the ceiling, walls, floor, fixtures, shelving, counters, and cabinets in the buffer area or clean room shall be smooth, impervious, free from cracks and crevices and non-shedding. Junctures of ceilings to walls shall be coved or caulked. There shall be no sink or floor drains in the buffer area or clean room.
- d. An anteroom shall be physically isolated from the buffer area or clean room. In this area, supplies are uncartoned and disinfected. Hand sanitizing and gowning occurs in this area. A demarcation line or barrier identifies the separation of the buffer area from the anteroom area. The air quality of the anteroom shall be Class 100,000 (ISO Class 8) or better.

- e. Presterilization procedures for high-risk level CSPs, such as weighing and mixing, shall be completed in no worse than an ISO Class 8 environment.

**21.20.70 Environmental Monitoring.**

- a. Class 100 or better clean rooms and/or primary engineering controls shall be certified by qualified operators at least every six months and whenever the device or room is relocated or major service to the facility is performed. Certification records shall be maintained and be available for inspection at the outlet for at least two years from the certification date.
- b. Certification that each ISO classified area is within established guidelines shall be performed no less than every six months and whenever the primary engineering control is relocated or the physical structure of the buffer area or anteroom has been altered. The testing shall be performed by qualified operators using state-of-the-art electronic equipment with the following results:
  - (1) Not more than 3,520 particles 0.5 micrometer size and larger per cubic meter of air for any primary engineering control (ISO Class 5).
  - (2) Not more than 352,000 particles of 0.5 micrometer size and larger per cubic meter of air (ISO Class 7) for any buffer room; and
  - (3) Not more than 3,520,000 particles of 0.5 micrometer size and larger per cubic meter of air (ISO Class 8) for any anteroom/area.
- c. Certification records shall be maintained and be available for inspection at the outlet for at least two years from the certification date.
- d. Tests shall be done for airborne microorganisms. Electronic air samplers are the preferred method. The instructions in the manufacturer's user manual for verification and use of the electronic air sample that actively collects volumes of air for evaluation must be followed. The sampling is performed at locations judged by compounding personnel to be the most prone to contamination. These tests shall be done at least every six months. The outlet shall have written policies to reevaluate cleaning procedures, operational procedures, and air filtration efficiency if the number of colony forming units increases over the normal baseline level. Records of these tests shall be maintained and be available for inspection at the outlet for at least two years from the testing date.
- e. Glove fingertip sampling shall be conducted at least annually for all compounding personnel if compounding low and medium risk CSPs and semi-annually if compounding high risk CSPs. When a finger plate result for personnel monitoring after proper incubation exceeds the action limit, a review of hand hygiene and garbing procedures as well as glove and surface disinfection procedures and work practices shall occur.
- f. A pressure gauge shall be installed to monitor the pressure differential between the ISO Class 7 cleanroom and the ISO Class 8 anteroom and the anteroom and the general pharmacy area. The results shall be reviewed and documented on a daily basis. The pressure between the cleanroom and general pharmacy area shall not be less than 5 Pa (0.02-inch water column, w.c.). The pressure differential between the cleanroom and the anteroom shall be greater than the pressure differential between the anteroom and the general pharmacy area, except for the preparation of radiopharmaceuticals where there is no pressure differential.

21.20.80 Cleaning and Disinfecting the Workspaces.

- a. The cleaning and sanitizing of the workspaces shall be done pursuant to written procedures and shall be the responsibility of trained operators, using appropriate disinfecting agents.
- b. The direct and contiguous compounding area (DCCA), including ISO Class 5 areas, shall be cleaned and disinfected prior to the beginning of each shift. All items shall be removed from the DCCA and all surfaces shall be cleaned of loose material and residue from spills prior to cleaning.
- c. Work surfaces in the ISO Class 7 buffer areas and ISO Class 8 anteroom/areas are cleaned and disinfected at least daily.
- d. Dust and debris shall be removed as necessary from the storage areas for compounding ingredients and supplies.
- e. Storage shelving shall be disinfected at least monthly. All items shall be removed from the shelving prior to cleaning.
- f. The walls and ceilings in the buffer and anteroom areas shall be cleaned and disinfected at least monthly.
- g. Floors in the buffer and anteroom areas shall be mopped daily when no aseptic operations are in progress.
- h. All cleaning tools, such as wipers, sponges, and mops shall be non-shedding and dedicated to use in the buffer or clean area. Floor mops may be used in both the buffer or clean area and anteroom area, but only in that order. Most wipers shall be discarded after one use. If cleaning tools are reused, their cleanliness shall be maintained by thorough rinsing and disinfection after use and by storing in a clean environment between uses. Trash shall be collected in suitable plastic bags and removed with minimal agitation.

21.20.90 Personnel Cleansing and Garbing.

- a. Prior to entering the controlled (buffer) area, operators shall remove personal outer garments (such as lab jackets), makeup, and jewelry.
- b. After donning dedicated appropriate garbing that includes shoes or shoe covers, head and facial hair coverings, and face masks, hands and arms shall be thoroughly scrubbed up to the elbow. After drying hands and arms, operators shall properly don non-shedding gowns that fit snugly around the wrists and enclosed at the neck.
- c. Once inside the clean area, hands shall be cleansed with an antiseptic hand cleanser. Sterile, powder-free gloves shall then be donned, except for the preparation of radiopharmaceuticals where the prescription drug outlet can demonstrate statistically significant equivalence or superior sterility by using another method.
- d. During protracted compounding activities, personnel shall intermittently resanitize their gloves.
- e. For low and medium risk compounding: If personnel leave the buffer area, they shall don new hair covers, masks, shoe covers, and gloves prior to reentry. Gowns may be reused during the same compounding session if hung in the anteroom.



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- f. For high risk: If personnel leave the buffer area, they must don new hair covers, masks, shoe covers, gowns and gloves prior to reentry.
- 21.21.10 Components.
- a. Compounding personnel shall ascertain that ingredients for CSPs are in compliance with rule 21.00.10(b) and are of the correct identity and appropriate quality using the following information: vendors' labels, labeling, certificates of analysis, direct chemical analysis, and knowledge of compounding facility storage conditions. No expired components may be used in compounding. No component may be used which will expire prior to the beyond-use date of the finished CSP.
- b. Ingredients used in a compounded preparation shall either originate from FDA-approved sources, when available, or be USP/NF grade substances when such sources are not available and identified on the FDA drug shortage list.
- c. If neither USP/NF grade substances nor FDA-approved substances are available, or when food, cosmetics, or other substances are, or must be used, the substance shall be of a chemical grade in one of the following categories:
- (1) Chemically Pure (CP);
  - (2) Analytical Reagent (AR); or
  - (3) American Chemical Society (ACS); or
  - (4) Food Chemical Codex.
- d. For all ingredients, unless FDA-approved, the pharmacist shall establish purity and stability by obtaining a certificate of analysis from the supplier. The certificate of analysis, when applicable, shall be maintained at the prescription drug outlet for at least two years from the date of preparation.
- e. A manufactured drug product may be a source of active ingredient. Only manufactured drugs from containers labeled with a lot number and an expiration date are acceptable as a potential source of active ingredients. When compounding with manufactured drug products, the compounder must consider all ingredients present in the drug product relative to the intended use of a compounded preparation.
- f. Drug preparations that have been withdrawn or removed from the market for safety reasons shall not be compounded. Such preparations may be compounded exclusively for veterinary use provided no documentation exists which indicates that the preparation is unsafe for such use.
- g. Sterile ingredients and components:
- (1) A written procedure for physical inspection of ingredients and components prior to compounding shall be followed.
- h. Non-sterile ingredients and components:
- (1) If any non-sterile components or ingredients, including containers, devices, and ingredients, are utilized to make the CSP, the product shall be compounded at high risk.
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- (2) If non-USP or non-NF active ingredients, added substances, or excipients are utilized, a certificate of analysis from the supplier of the ingredient shall be maintained at the prescription drug outlet for at least two from the date of preparation.
- (3) When non-sterile ingredients and components are received at the outlet, their container shall be marked, in indelible pencil or ink, with the date of receipt. In the absence of a supplier's expiration date on the product, the expiration date of the ingredient shall be one-year from the date of receipt, unless either appropriate inspection or testing indicates that the ingredient has retained its purity and quality for use in CSPs.
- (4) Prior to compounding with non-sterile ingredients and components, the ingredients shall be visually inspected for evidence of deterioration, other types of unacceptable quality and wrong identification.
- i. Any ingredient regulated by the FDA through an Investigational Review Board (IRB) is exempt from rule 21.21.10 provided the research requirements for the receipt of the ingredient is followed and meets the requirements of CRS 12-42.5-128(2).

**21.21.20 Equipment.**

- a. Written procedures outlining required equipment, calibration, appropriate maintenance, monitoring for proper function, controlled procedures for use of the equipment and specified time frames for these activities shall be established and followed. Results of equipment calibration and appropriate maintenance reports shall be kept on file at the outlet for at least two years from the report date and shall be available for inspection.
- b. Accuracy assessments of automated compounding devices (ACD) shall be conducted daily for each day used. At routine intervals, the pharmacist manager, or his or her designee, shall review these assessments to avoid potentially clinically significant cumulative errors over time. These assessments shall be documented and be maintained and available for inspection at the outlet for at least two years.

**21.21.30 Finished Preparation Release Checks and Tests.**

- a. Physical Inspection
  - (1) Finished CSPs shall be individually inspected after compounding pursuant to written procedures. Immediately after compounding, and prior to dispensing or distribution, each product shall be inspected for evidence of particulates or other foreign matter, container-closure integrity, precipitation, cloudiness, and any other apparent visual defect. Defective product shall be segregated from other product and shall not be dispensed or distributed.
- b. Compounding Accuracy Checks.
  - (1) Written procedures for double-checking compounding accuracy shall be followed for every CSP during preparation and immediately prior to release. Outlets which compound CSPs shall have at least the following written procedures for verifying the correct identity and quality of CSPs prior to dispensing or distribution:
    - (a) Verification of label for accuracy;

- (b) Correct identities, purities, and amounts of ingredients have been used; and
  - (c) Correct fill volumes in CSPs and correct quantities of filled units of the CSPs were obtained.
- c. Sterility Testing.
  - (1) Sterility testing shall be done on the following high risk CSPs:
    - (a) Batches larger than 25 identical individual single-dose packages (ampules, bags, syringes, vials, etc);
    - (b) Multiple dose vials for administration to multiple patients;
    - (c) Product is exposed longer than 12 hours at refrigerator temperatures prior to sterilization; or
    - (d) Product is exposed longer than 6 hours to temperatures warmer than refrigerator temperature prior to sterilization.
  - (2) The sterility test shall be compliant with the most current USP/NF Chapter 71 < Sterility Tests >. A method not described in the USP/NF may be used if verification results demonstrate that the alternative is at least as effective and reliable as the USP/NF methods.
  - (3) When a high risk CSP is dispensed or distributed before receiving the results of the sterility test, there shall be a written procedure requiring daily observation of the incubating test specimens and requiring an immediate recall if there is any evidence of microbial growth. In addition, the patient and the practitioner of the patient to whom a potentially contaminated CSP was administered shall be notified of the potential risk. Positive sterility results shall prompt a rapid and systematic investigation of aseptic technique, environmental and other sterility assurance controls to identify sources of contamination and correct problems in the methods or processes.
- d. Bacterial Endotoxin (Pyrogen) Testing.
  - (1) Endotoxin testing shall be done on the following high risk CSPs that are to be administered parenterally:
    - (a) Batches larger than 25 identical individual single-dose packages (ampules, bags, syringes, vials, etc.);
    - (b) Multiple dose vials for administration to multiple patients;
    - (c) Product is exposed longer than 12 hours at refrigerator temperatures prior to sterilization; or
    - (d) Product is exposed longer than 6 hours to temperatures warmer than refrigerator temperature prior to sterilization.

- (2) The endotoxin test shall be compliant with the most current USP/NF Chapter 85 < Bacterial Endotoxins Test >. In the absence of a bacterial endotoxins limit in the official monograph or other CSP formula source, the CSP must not exceed the amount of USP/NF Endotoxin Units (EU per hour per kg of body weight) specified for the route of administration.

**21.21.40 Storage and Beyond-Use Dating.**

- a. The temperature of drug storage areas of CSPs shall be monitored and recorded daily, either manually or electronically. Temperature records shall be maintained and be available for inspection for at least two years.
- b. Finished CSPs that are not immediately dispensed or administered shall be refrigerated or frozen unless their chemical and physical stability are known to be adversely affected by cold or freezing temperatures.
- c. In the absence of sterility testing for each compounded batch compliant with the most current USP/NF Chapter 71 <Sterility Tests>, the beyond-use date (before administration) shall not exceed the following:

- (1) Low risk CSPs with greater than 12-hour BUD:

Room temperature: No more than 48 hours

Refrigerated temperature: No more than 14 days

Frozen: No more than 45 days

- (2) Low risk CSPs with 12-hour or less BUD:

Room temperature: No more than 12 hours

Refrigerated temperature: No more than 12 hours

Frozen: Not applicable

- (3) Medium risk CSPs:

Room temperature: No more than 30 hours

Refrigerated temperature: No more than 9 days

Frozen: No more than 45 days

- (4) High risk CSPs:

Room temperature: No more than 24 hours

Refrigerated temperature: No more than 3 days

Frozen: No more than 45 days

- d. For high risk products, there must be a reliable method for establishing all expiration dates, including sterility. There must be a reliable method for establishing all beyond-use dating. Products maintaining beyond-use dating of greater than thirty (30) days shall have lab testing of product stability and potency.
- e. Each outlet shall adhere to manufacturers' instructions for handling and storing of Add-Vantage®, Mini Bag Plus®, Add A Vial®, Add-Ease® products, and any similar products.

**21.21.50 Formulation Record.**

- a. For each CSP, a uniform, readily retrievable formulation record shall be maintained and available for inspection for two years from the date last utilized, documenting:
  - (1) The name, strength, dosage form, and route of administration of the compounded preparation;
  - (2) All ingredients and their quantities;
  - (3) The equipment used to compound the preparation, when appropriate, and mixing instructions;
  - (4) The beyond use date;
  - (5) The containers used in dispensing;
  - (6) Storage requirements; and
  - (7) Procedures for quality control, if applicable.

**21.21.60 Compounding Record.**

- a. For each CSP prepared, a record shall be maintained and available for inspection for two years on the original order, or on a separate, uniform, readily retrievable record documenting the following:
  - (1) Name and strength of the compounded preparation;
  - (2) Formulation record reference for the preparation;
  - (3) Sources and lot number of each ingredient;
  - (4) Manufacturer's expiration date of each ingredient, when applicable;
  - (5) Total number of dosage units compounded;
  - (6) Name of the person who compounded the preparation;
  - (7) Name of the pharmacist who approved the preparation;
  - (8) Batch (lot) number assigned, if multiple units compounded;
  - (9) Date of preparation;
  - (10) Beyond use date;

- (11) Prescription number(s), if appropriate;
- (12) Results of quality control procedures; and
- (13) If a high risk product, the record shall also include comparisons of actual with anticipated yields, sterilization methods, and quarantine specifications.

**21.21.70 Labeling of CSPs.**

- a. Labeling of CSPs dispensed pursuant to a prescription order or LTCF chart order shall include at least the following:
  - (1) All requirements of CRS12-42.5-121;
  - (2) Batch (lot) number, if appropriate;
  - (3) Beyond-use date;
  - (4) If for parenteral administration, the following shall be included:
    - (a) Name of base solution; and
    - (b) name and amounts of drugs added.
  - (5) Storage directions; and
  - (6) A clear statement that this product was compounded by the pharmacy, except for radiopharmaceuticals prepared from FDA-approved, commercially available kits and/or drug products.
- b. Labeling of CSPs dispensed pursuant to a hospital chart order shall include at least the following:
  - (1) All requirements of CRS 12-42.5-121;
  - (2) Batch (lot) number, if appropriate;
  - (3) Beyond-use date;
  - (4) If for parenteral administration, the following shall be included;
    - (a) Name of base solution; and
    - (b) Name and amounts of drugs added; and
  - (5) Storage directions.
- c. Labeling of CSPs distributed to practitioners, other prescription drug outlets, or other outlets allowed by law shall include at least the following:
  - (1) Name of the outlet;
  - (2) Name and strength of the drug(s);
  - (3) Total quantity in package;

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- (4) Quantity of active ingredient in each dosage unit;
    - (5) Beyond-use date;
    - (6) Batch (lot) number;
    - (7) Specific route of administration;
    - (8) Storage directions;
    - (9) "Rx only"; and
    - (10) A clear statement that this product was compounded by the pharmacy, except for radiopharmaceuticals prepared from FDA-approved, commercially available kits and/or drug products.
  - d. Labeling of CSPs distributed within hospitals as floor stock shall include at least the following:
    - (1) Name of the outlet;
    - (2) Name and strength of the drug(s);
    - (3) Total quantity in package;
    - (4) Quantity of active ingredient in each dosage unit;
    - (5) Beyond-use date;
    - (6) Batch (lot) number;
    - (7) Specific route of administration; and
    - (8) Storage directions.
- 21.21.80** Maintaining Product Quality and Control After the CSP Leaves the Outlet or Hospital Location.
- a. The outlet shall have written policies and procedures that are adhered to which shall ensure the CSP is packaged properly for transit, stored properly during transit, and stored properly at site of administration. Such policies and procedures shall also discuss patient or caregiver training.
- 21.21.90** **Patient Monitoring, Adverse Events Reporting, and Product Recall.**
- a. Outlets which compound CSPs shall provide patients and other recipients of CSPs with a way to address their questions and report any concerns that they may have with CSPs and their administration devices.
  - b. The outlet shall have written policies describing specific instructions for receiving, acknowledging, and dating receipts; and for recording, or filing, and evaluating reports of adverse events and of the quality of preparation claimed to be associated with CSPs.
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- c. The pharmacist manager shall report to the Board in writing significant errors related to compounded CSPs such as those that result in serious personal injury or death of a patient.
- d. If a CSP is believed to be defective in any way, or if the Board or Federal Food and Drug Administration makes a written request for a recall of a specific preparation, the outlet shall immediately recall any product dispensed or distributed. Any product remaining in the outlet shall be immediately quarantined and shall not be dispensed or distributed. Recall records shall include at least the following:
  - (1) Product name, strength, dosage form;
  - (2) Reason for recall;
  - (3) Amount of product made;
  - (4) Date made; and
  - (5) Amount of product dispensed or distributed.
- e. The outlet shall conduct tests, as appropriate, on the recalled product to identify the reason the product was defective. Results of these tests shall be maintained at the outlet for at least two years.
- f. Adverse event reports and product recall records shall be retained and be available for inspection at the outlet for at least two years.

**21.22.00 Quality Assurance Program.**

- a. Outlets that make CSPs shall have a formal written quality assurance (QA) program which shall provide a mechanism for monitoring, evaluating, correcting, and improving the activities and processes regarding the compounding of sterile products.
- b. At a minimum, the written QA program shall include the following:
  - (1) Consideration of all aspects of the preparation, dispensing, and distribution of products, including environmental testing, work area cleaning effectiveness, validation results, etc;
  - (2) Describe specific monitoring and evaluation activities;
  - (3) Specification of how results are to be reported and evaluated;
  - (4) Identification of appropriate follow-up mechanisms when action limits or thresholds are exceeded; and
  - (5) Delineation of the individuals responsible for each aspect of the QA program.

**21.22.10 Cytotoxic Drug Preparation.**

- a. Cytotoxic drugs shall be compounded in a vertical flow, Class II biological safety cabinet (BSC) or CACI. Such BSC or CACI shall be placed in an ISO Class 7 area that is physically separated from other preparation areas and is negative pressure to adjacent positive pressure anteroom. If used for other products, the cabinet must be thoroughly cleaned;



- b. Appropriate personnel protective equipment (PPE) shall be worn when compounding in a BSC or CACI and when using closed-system vial transfer devices (CSTDs). PPE should include gowns, face masks, eye protection, hair covers, shoe covers or dedicated shoes, double gloving with sterile chemo-type gloves, and compliance with manufacturers' recommendations when using a CACI;
- c. Appropriate safety and containment techniques for compounding cytotoxic drugs shall be used in conjunction with the aseptic techniques required for preparing sterile products;
- d. Appropriate disposal containers for used needles, syringes, and if applicable, cytotoxic waste from the preparation of chemotherapy agents and infectious waste from patients' homes. Disposal of cytotoxic waste shall comply with all applicable local, state and federal requirements;
- e. Written procedures for handling major and minor spills and generated waste of cytotoxic agents must be developed and must be included in the policy and procedure manual; and
- f. Prepared doses of cytotoxic drugs must be labeled with proper precautions inside and outside, and shipped in a manner to minimize the risk of accidental rupture of the primary container.

**21.22.20 Exemption for Sterile Compounding of Products in Closed or Sealed System.**

- a. Pharmacists and pharmacies or other outlets where sterile compounding is provided may be exempt from this rule when compounding is restricted to utilizing compounds or products that are contained only in a closed or sealed system and can be transferred or compounded within this self-contained system or topical products that require further transfer or combination in order to achieve a finished product without further modification of the product.

**22.00.00 [Repealed eff. 07/15/2010]**

**23.00.00 ELECTRONIC PRESCRIPTION MONITORING PROGRAM.**

**23.00.10 Definitions:**

- a. "Bona fide investigation," for purposes of an investigation of an individual prescriber under investigation by a state regulatory board, means:
  - 1. Any investigation conducted by any state regulatory board within the Colorado Division of Professions and Occupations, or the Director of the Colorado Division of Professions and Occupations and
  - 2. Investigations pertaining to matters which are the subject of a complaint or notice of charges pending in the Office of Administrative Courts so long as the information obtained from the PDMP is made available by the state regulatory board to the respondent in the pending case.

- b. "Bona fide research or education" means research conducted by qualified entities whose recognized primary purpose is scientific inquiry; the results of which would likely contribute to the basic knowledge of prescribing practitioners, dispensing pharmacists, or entities for the purpose of curtailing substance abuse of consumers. The Board shall determine in its discretion on a case-by-case basis whether an individual or entity seeking access to the PDMP pursuant to CRS 12-42.5-404(5) constitutes "bona fide research or education" conducted by qualified personnel for purposes of satisfying the statutory limitations therein.
- c. "Clinical patient care services" means pharmaceutical care provided in a clinical setting. The pharmacist providing clinical patient care services must be working closely with the physician/prescriber responsible for the patient's care. "Clinical patient care services" do not include monitoring previously dispensed prescriptions for any purpose in the absence of a current assessment of a patient whether in a clinical setting or not.
- d. "Law Enforcement Official" means any of the following:
  - 1. Sheriff;
  - 2. Undersheriff;
  - 3. Certified deputy sheriff;
  - 4. Coroner;
  - 5. Police Officer;
  - 6. Southern Ute Police Officer;
  - 7. Ute Mountain Ute police officer;
  - 8. Town marshal;
  - 9. CBI director and agents;
  - 10. Colorado state patrol officer;
  - 11. Colorado attorney general and any entity designated as "peace officers" by the Attorney General or acting on behalf of a state agency;
  - 12. Attorney general criminal investigator;
  - 13. District attorney and all assistants, deputies, etc. statutorily defined as "peace officers;"
  - 14. District Attorney chief investigator and investigators;
  - 15. Police administrator and police officers employed by the Colorado State Hospital in Pueblo; and
  - 16. Federal special agents.

- e. “Legitimate program to monitor a patient’s controlled substance abuse” means a program in which prescribers actively monitor a patient’s controlled substance use. Such programs shall only involve patients in pain management or other controlled substance management programs. Such programs shall actively monitor the patient’s controlled substance usage by means of urine or other drug screens in addition to the use of the PDMP. The patient must be informed in writing that his/her controlled substance usage is being actively screened by various methods, including review of the PDMP.
- f. “PDMP” means the Electronic Prescription Drug Monitoring Program.
- g. “Prescriber” or “practitioner” means a licensed health care professional with authority to prescribe a controlled substance.
- h. “Prescription Drug Outlet” or “Dispenser” means any resident or nonresident pharmacy registered with the Board.
- i. “Qualified personnel” means persons who are appropriately trained to collect and analyze data for the purpose of conducting bona fide research or education.
- j. “Valid photographic identification” means any of the following forms of identification which include an identifying photograph:
  - 1. A valid driver’s license, or identification issued by any United States state;
  - 2. An official passport issued by any nation; or
  - 3. A United States armed forces identification card issued to active duty, reserve, and retired personnel and the personnel’s dependents.

**23.00.30 Data Submission Timeline.**

Every prescription drug outlet must ensure that all controlled substance dispensing transactions are reported to the PDMP on a daily basis by no later than the outlet’s next regular business day.

**23.00.40 Data Submission Format.**

Prescription drug outlets shall submit to the PDMP the following data requirements:

- a. Identifier (Transmission type identifier), if applicable;
- b. Bin (Bank Identification Number);
- c. Version Number (a number to identify the format of the transaction sent or received);
- d. Transaction Code;
- e. NABP or Drug Enforcement Administration number assigned to pharmacy;
- f. Customer ID (number to identify the patient receiving the RX);
- g. Zip Code (3 digit US Postal Code identifying the State Code), if applicable;
- h. Customer’s Birth Date;
- i. Sex Code;

- j. Date Filled;
  - k. Prescription Number;
  - l. New/Refill Number;
  - m. Metric Quantity;
  - n. Days Supply;
  - o. Compound Code;
  - p. NDC Number of the drug dispensed;
  - q. Prescriber's Drug Enforcement Administration registration;
  - r. Drug Enforcement Administration suffix, if applicable;
  - s. Date RX Written;
  - t. Number of Refills Authorized;
  - u. RX Origin Code;
  - v. Customer Location;
  - w. Diagnosis Code, if available;
  - x. Alternate Prescriber #, if applicable;
  - y. Patient Last Name (if an animal, the owner's last name);
  - z. Patient First Name (if an animal, the animal's first name);
  - aa. Patient Street Address;
  - bb. Patient's state of residence;
  - cc. Patient's zip code;
  - dd. Triplicate Serial Number, if appropriate; and
  - ee. Filler Field to be populated with Payment Type as designated by PDMP vendor.
- 23.00.50 Data Correction.
- a. Any errors identified by the PDMP shall be corrected and resubmitted by the prescription drug outlet within 30 calendar days of original dispensing date of the affected prescription(s).
  - b. If errors cannot be corrected, the pharmacy must retain a record in written format detailing the following information for each uncorrected error:
    - 1. Detail of Error Notification highlighting uncorrected error(s); and

2. Detailed reason of why error cannot be corrected.

**23.00.60 PATIENT NOTIFICATION**

Prescription Drug Outlets shall disclose to patients receiving controlled substance prescriptions that their prescription information is being submitted to the PDMP, and that this prescription information may be queried by specific individuals for a limited number of purposes as authorized by statute.

- 23.00.65 Unsolicited Reporting.** In conjunction with other Colorado Boards who regulate prescribing practitioners and applicable stakeholders, the Board shall develop criteria for indicators of potential misuse, abuse and diversion of controlled substances and, based on those criteria, provide unsolicited reports of dispensed controlled substance prescriptions to the responsible prescribing practitioners and dispensing pharmacies of controlled substance(s) dispensed to the patient for purposes of education and intervention to prevent and reduce occurrences of controlled substance misuse, abuse, and diversion.

**23.00.70 PDMP Access**

The PDMP shall be available for query only to the following persons or groups of persons:

- a. Board staff responsible for administering the PDMP;
- b. Any licensed practitioner, or up to three (3) trained individuals designated by the practitioner by way of registered PDMP sub-accounts of the prescriber to act on the prescriber's behalf in accordance with 12-42.5-403(1.5)(b), (c) and (d), C.R.S., with the statutory authority to prescribe controlled substances to the extent the query relates to a current patient of the practitioner to whom the practitioner is prescribing or considering prescribing a controlled substance;
- c. Licensed pharmacists, or up to three (3) trained individuals designated by the pharmacist by way of registered PDMP sub-accounts of the pharmacist to act on the pharmacist's behalf in accordance with 12-42.5-403(1.5)(b), (c) and (d), C.R.S., or a pharmacist licensed in another state, with statutory authority to dispense controlled substances to the extent the information requested relates specifically to a current patient to whom the pharmacist is dispensing or considering dispensing a controlled substance or to whom the pharmacist is providing clinical patient care services;
- d. Practitioners engaged in a legitimate program to monitor a patient's controlled substance abuse;
- e. Law enforcement officials so long as the information released is specific to an individual patient, prescriber, or prescription drug outlet and part of a bona fide investigation and the request for information is accompanied by an official court order or subpoena. Such official court orders or subpoenas shall be submitted with the Board-provided form;
- f. The individual who is the recipient of a controlled substance prescription so long as the information released is specific to such individual. The procedure for individuals to obtain such information is as follows:
  1. The individual shall submit a written, signed request to the Board on the Board-provided form;
  2. The individual shall provide valid photographic identification prior to obtaining the PDMP information;

3. An individual submitting a request on behalf of another individual who is the recipient of a controlled substance prescription may only obtain PDMP information if the following documents are provided:
  - A. The original document establishing medical durable power of attorney of the individual submitting the request as power of attorney for the individual who is the recipient of the controlled substance prescription, and
  - B. Valid photographic identification of the individual submitting the request.
- g. State regulatory boards within the Colorado Division of Professions and Occupations and the Director of the Colorado Division of Professions and Occupations so long as the information released is specific to an individual prescriber and is part of a bona fide investigation and the request for information is accompanied by an official court order or subpoena. Such official court orders or subpoenas shall be submitted with the Board-provided form; and
- h. A resident physician with an active physician training license issued by the Colorado medical board pursuant to section 12-36-122 and under the supervision of a licensed physician to the extent the query relates to a current patient of the resident physician to whom the resident physician is prescribing or considering prescribing a controlled substance.
- i. The Department of Public Health and Environment for purposes of population-level analysis, but any use of the program data by the department is subject to the federal "Health Insurance Portability and Accountability Act of 1996 (HIPAA) and any rules promulgated pursuant to HIPAA, including the requirement to remove any identifying data unless exempted from the requirement.
- j. A person authorized to access the PDMP may knowingly release PDMP information specific to an individual or to the individual's treating providers in accordance with HIPAA, Pub.L. 104-191, as amended, and any rules promulgated pursuant to HIPAA without violating Part 4 of Title 12, Article 42.5.

**23.00.80 Research or Education Agreements**

The Board may enter into a written agreement to provide data to qualified personnel of a public or private entity for the purpose of bona fide research or education, so long as such information does not identify a recipient, prescriber, or dispenser of a prescription drug. Any public or private entity wishing to enter into or extend such an agreement shall submit a written request to the Board detailing the information it is seeking and the public benefit of such research or education. The Board will act on such request in the normal course of business.

**23.00.90 Exemptions**

- a. The following individuals or entities are exempt from reporting controlled substance dispensing transactions to the Prescription Drug Monitoring Program:
  1. Hospitals licensed or certified pursuant to CRS 25-1.5-103;
  2. A prescription drug outlet located within a hospital licensed or certified pursuant to CRS 25-1.5-103 that dispenses controlled substances only pursuant to chart orders or dispenses no more than a 24-hour supply of a controlled substance to an outpatient;

- 3. Emergency medical services personnel certified pursuant to CRS 25-3.5-203; and
  - 4. A prescription drug outlet which has applied to the Board and received a waiver from the Board. Waivers will only be considered if the pharmacy has no electronic automation. Such requests must be submitted in writing to the Board and will be considered in the normal course of business.
- b. Controlled substance dispensing transactions that occur solely for Institutional Review Board (IRB) approved interventional research trials using investigational drug products that are regulated by the Federal Food and Drug Administration shall be exempt from the data submission requirements of the PDMP.

**24.00.00 CONFIDENTIAL AGREEMENTS**

24.00.10 No later than thirty (30) days from the date a physical or mental illness or condition impacts a pharmacist's or pharmacy intern's ability to practice pharmacy with reasonable skill and safety, the pharmacist or pharmacy intern shall provide the Board, in writing, the following information:

- a. The diagnosis and a description of the illness or condition;
- b. The date that the illness or condition was first diagnosed;
- c. The name of the current treatment provider and documentation from the current treatment provider confirming the diagnosis, date of onset, and treatment plan;
- d. A description of the pharmacist's or pharmacy intern's practice and any modifications, limitations or restrictions to that practice that have been made as a result of the illness or condition; and
- e. Whether the pharmacist or pharmacy intern has been evaluated by, or is currently receiving services from the Board's authorized Peer Health Assistance Diversion Program related to the illness or condition and, if so, the date of initial contact and whether services are ongoing.

24.00.20 The pharmacist or pharmacy intern shall further notify the Board of any significant change in the illness or condition ("change of condition") that impacts the pharmacist's or pharmacy intern's ability to practice pharmacy with reasonable skill and safety. The pharmacist or pharmacy intern must notify the Board of a positive or negative change of condition. Such notification shall occur within thirty (30) days of the change of condition. The pharmacist or pharmacy intern shall provide the Board, in writing, the following information:

- a. The date of the change of condition;
- b. The name of the current treatment provider and documentation from the current treatment provider confirming the change of condition, the date that the condition changed, the nature of the change of condition, and the current treatment plan; and
- c. A description of the licensee's practice and any modifications, limitations or restrictions to that practice that have been made as a result of the change of condition.

24.00.30 Compliance with this rule is a prerequisite for eligibility to enter into a Confidential Agreement with the Board pursuant to Section 12-42.5-134, C.R.S. However, mere compliance with this rule does not require the Board to enter into a Confidential Agreement. Rather, the Board will evaluate all facts and circumstances to determine if a Confidential Agreement is appropriate.

24.00.40 If the Board discovers that a pharmacist or pharmacy intern has a mental or physical illness or condition that impacts the pharmacist's or pharmacy intern's ability to practice pharmacy with reasonable skill and safety and the pharmacist or pharmacy intern has not timely notified the Board of such illness or condition, the pharmacist or pharmacy intern shall not be eligible for a Confidential Agreement and may be subject to disciplinary action pursuant to Section 12-42.5-123(1)(r), C.R.S.

24.00.50 A pharmacist or pharmacy intern who is addicted to, dependent on, or engages in the habitual or excessive use or abuse of intoxicating liquors, a habit-forming drug, or a controlled substance as defined in Section 12-42.5-123(1)(e), C.R.S., shall seek assistance from the Diversion Program as governed by Section 12-42.5-204, C.R.S. Such pharmacists or pharmacy interns are not eligible to enter into a confidential agreement with the Board pursuant to Section 12-42.5-134, C.R.S.

**25.00.00 SPECIALIZED PRESCRIPTION DRUG OUTLETS.**

25.00.10 Definitions.

- a. "Automated device" or "AD" means a mechanical system that performs operations or activities relative to the storage, packaging, counting, labeling, and dispensing of medications, and which collects, controls, and maintains appropriate transaction information.
- b. "Long term care facility" or "LTCF" means a nursing facility as defined in section 25.5-4-103(14) C.R.S. that is licensed pursuant to 25-1.5-103 C.R.S. An LTCF is a nursing home, skilled nursing facility or a nursing care facility that provides supportive, therapeutic, or compensating services with the availability of a licensed nurse for observation and treatment on a twenty-four hour basis.
- c. "Managing prescription drug outlet" means the prescription drug outlet located within the State of Colorado which is responsible for ownership and operation of a specialized prescription drug outlet located at an LTCF within Colorado. The managing prescription drug outlet is responsible for the application for the specialized prescription drug outlet on behalf of the LTCF. The managing prescription drug outlet shall own and operate the SPDO and maintain ownership of the drugs.
- d. "Specialized prescription drug outlet" or "SPDO" means an outlet located at an LTCF which is owned and operated by a managing prescription drug outlet located within the State of Colorado. The managing prescription drug outlet engages in the compounding, dispensing, and delivery of drugs and devices, or the provision of pharmaceutical care, residents of the LTCF. The managing prescription drug outlet may use automated devices in the SPDO to provide drugs, as well as other Board-approved nontraditional methods, to provide pharmaceutical care to the residents of the LTCF
- e. "Stock drugs" mean non-patient specific prescription drugs or controlled substances that are distributed from a managing prescription drug outlet to a SPDO by means other than a patient-specific prescription order or LTCF chart order.



- 25.00.12 Requirements for Registration. Eligibility requirements for an SPDO include the following:
- a. A current Board-issued registration of the managing prescription drug outlet that engages in the compounding, dispensing, and delivery of drugs, or provision of pharmaceutical care to residents of an LTCF;
  - b. The submission of a separate application by the managing prescription drug outlet on behalf of the SPDO for a SPDO registration, on a form provided by the Division of Professions and Occupations. The managing prescription drug outlet shall submit an application for each individual SPDO to which the managing prescription drug outlet will provide stock drugs;
  - c. A Drug Enforcement Administration registration specifically assigned to the SPDO if the managing prescription drug outlet provides stock controlled substances to the SPDO;
  - d. Successful completion of a pre-registration inspection of the SPDO by the Board or its inspectors;
  - e. A pharmacist manager who, in addition to being responsible for the operations of the managing prescription drug outlet in compliance with all state and federal laws and rules, is responsible for the operations of the SPDO in compliance with all provisions of rule 25.00.00; and
  - f. A secure AD that prevents the diversion of drugs and that limits the access to drugs within the AD only to those persons whom have been given permission to access the AD.
- 25.00.14 Scope of Practice.
- a. An SPDO shall maintain and operate an AD for the purpose of storing drug stocks.
  - b. The managing prescription drug outlet shall ensure that all medications stocked in the AD are either in unit dose form, single dose packages, or packaged as such or in customized medication packs prior to release from the AD for administration to a patient. All records of packaging shall be maintained at the managing prescription drug outlet.
  - c. An SPDO shall only utilize stock prescription drugs or controlled substances it receives from the managing prescription drug outlet for the purpose of drug administration, and not for the purpose of further dispensing.
- 25.00.16 Records and Recordkeeping.
- a. The managing prescription drug outlet shall be exempt from any casual sale limitations specified in 12-42.5-102(6) C.R.S. only to the extent of distributing drug stocks to an SPDO.
  - b. Records of drug distribution from the managing prescription drug outlet to the SPDO shall be retained at the managing prescription drug outlet and shall be readily available for inspection by the Board or its inspectors for at least two years from the date of distribution. These records shall be maintained separately from all other records of distribution to Board-registered entities which are not SPDOs or individual practitioners authorized by law prescribe the drugs. The record of distribution shall include the following:
    - 1. The name of the drug;

2. The strength of the drug;
  3. The dosage form if appropriate;
  4. The quantity of the drug;
  5. The name of the manufacturer or the NDC number of the drug if labeled only with its generic name;
  6. The date of distribution;
  7. The name and address of the distributing prescription drug outlet;
  8. The name and address of the receiving SPDO;
  9. If a controlled substance is distributed, the record shall also indicate the DEA registration number of the distributing prescription drug outlet and the receiving SPDO;
  10. A schedule II controlled substance shall only be distributed pursuant to receipt of a properly executed DEA-222 form;
  11. The identity of the person in the prescription drug outlet who issued the drug; and
  12. The identity of the person who placed the drug into the SPDO's AD.
- c. A duplicate copy of the record of distribution outlined in Rule 25.00.16(b) shall be maintained at the SPDO in a readily retrievable manner for at least two years from the date of receipt. This record shall serve as the SPDO's record of receipt.
- d. Records of use from the AD shall be retained at the managing prescription drug outlet and shall be readily available for inspection by the Board or its inspectors for at least two years from the date of latest use transaction. The record of use shall include the following:
1. The name of the patient;
  2. The name of the practitioner;
  3. Date removed;
  4. The name, strength and dosage form of the drug removed;
  5. The quantity of the drug removed; and
  6. The identity of the person at the SPDO that removed the drug.

- e. Biennial Controlled Substance Inventory. A complete and exact inventory of all stocks of controlled substances shall be conducted at each SPDO at least once every two years. The inventory shall be recorded on a uniform and readily retrievable record, and this record shall be signed by the pharmacist manager of the managing prescription drug outlet or another pharmacist as delegated by the pharmacist manager. The inventory shall include the date and time of day the inventory was conducted. A copy of this recorded inventory shall be maintained and readily available for inspection at both the managing prescription drug outlet and SPDO for at least two years from the date the inventory was conducted. This inventory record shall be maintained separately from all other recorded inventories of the managing prescription drug outlet.
- 25.00.18 Policy and Procedure Manual.
- a. Each managing prescription drug outlet and corresponding SPDO shall maintain a policy and procedure manual which is approved by the Board or its designee prior to SPDO operation. This policy and procedure manual shall be reviewed, signed, and dated by both the pharmacist manager of the managing prescription drug outlet and the nursing home administrator or other accountable individual of the SPDO at least once annually. The pharmacist manager shall be responsible for assuring that the nursing home administrator or other accountable individual of the SPDO signs the policy and procedure manual.
  - b. If a change in the pharmacist manager, or nursing home administrator or other accountable individual at the SPDO occurs, the new pharmacist manager and/or nursing home administrator or other accountable individual shall review, sign, and date the policy and procedure manual within 30 days of assuming the respective positions. The pharmacist manager shall be responsible for assuring that the new nursing home administrator or other accountable individual of the SPDO signs the policy and procedure manual.
  - c. The policy and procedure manual shall, at minimum, address the accessibility to, the stocking of, the accountability and recordkeeping of, and the security of, the AD.
- 25.00.21 Relocation.
- a. In the event of a relocation of a SPDO, the managing prescription drug outlet shall submit an application form provided by the Division of Professions and Occupations along with the prescribed fee at least 30 days prior to the effective date of relocation.
- 25.00.22 Reinstatement of a SPDO Registration. If a registration of a SPDO has expired, the managing prescription drug outlet seeking to reinstate such SPDO registration shall submit the an application on a form provided by the Division of Professions and Occupations along with the required fee.
- 25.00.24 Closure.
- a. Upon the closure of the SPDO it shall be the responsibility of the managing prescription drug outlet's pharmacist manager to remove all drug stocks from the LTCF within 72 hours after closure.
  - b. The pharmacist manager of the managing prescription drug outlet shall notify the Board of the closure of the SPDO location within seven business days of the closure.

**26.00.00 REMOTE PHARMACY PRACTICE.**

**26.00.10 Definitions**

- a. "Drug regimen review" includes but is not limited to the evaluation of order(s) and patient records(s) for:
  - 1. Known allergies;
  - 2. Rational therapy and contraindications;
  - 3. Reasonable dose, duration of use, and route of administration considering age, gender, and other patient factors;
  - 4. Reasonable directions for use;
  - 5. Potential or actual adverse drug reactions;
  - 6. Drug-drug interactions;
  - 7. Drug-food interactions;
  - 8. Drug-disease contraindications;
  - 9. Therapeutic duplication;
  - 10. Proper utilization (including over- or under-utilization) and optimum therapeutic outcomes; and
  - 11. Abuse/misuse.
- b. "Final Evaluation" means the review of the final prescription to ensure that the ordered medication is properly prepared and placed in a suitable container with appropriate labeling. The pharmacist(s) conducting the final evaluation shall be held accountable for assuring that the identity of the drug that appears on the prescription label corresponds with identity of drug contained therein. When refills are dispensed, the pharmacist conducting the final evaluation shall be held accountable for the appropriate dispensing of refills including all drug utilization reviews as they pertain to refill dispensing.
- c. "Initial Interpretation" means the review of an order accompanied by order entry. The pharmacist(s) conducting the initial interpretation shall be held accountable for the accuracy of the electronic order entry/manual transcription and/or drug regimen review.
- d. "Remote Pharmacy Practice" means initial interpretation and/or final evaluation of orders which is conducted at a location other than a registered prescription drug outlet or other outlet registered with the Board.

**26.00.20 Requirements for Remote Pharmacy Practice.**

- a. A registered prescription drug outlet or other outlet may employ or contract with one or more pharmacists for the purpose of conducting Remote Pharmacy Practice provided that all requirements, including those of confidentiality, privacy, and security are met, as required in this rule.

- b. All pharmacists employed or contracted with to conduct Remote Pharmacy Practice must be Colorado licensed pharmacists, holding active, unrestricted licenses with the Board;
- c. No drug inventory shall be stocked or maintained at any Remote Pharmacy Practice location;
- d. Pharmacists engaged in Remote Pharmacy Practice are not considered in the computation of the technician to pharmacist ratio;
- e. All records of Remote Pharmacy Practice must be maintained at the prescription drug outlet or other outlet in accordance with applicable recordkeeping rules;
- f. Remote Pharmacy Practice shall have adequate security and be conducted in a setting sufficient to maintain patient privacy and confidentiality;
- g. Prior to engaging in Remote Pharmacy Practice, the prescription drug outlet or other outlet shall submit a Policy and Procedure Manual and sample Written Agreement to the Board for review. Such Policy and Procedure Manual and Written Agreement must be approved by the Board or its designee prior to the prescription drug outlet or other outlet engaging in Remote Pharmacy Practice. Any revisions to the Policy and Procedure Manual and Written Agreement must be approved by the Board or its designee prior to implementation;
- h. Each prescription drug outlet or other outlet which employs or contracts with pharmacists engaged in Remote Pharmacy Practice must maintain a written Policy and Procedure Manual. Such Policy and Procedure Manual shall be reviewed, updated, and revised as necessary to remain current, but at least annually. Written documentation of such review shall be maintained at the prescription drug outlet or other outlet. This policy and procedure manual shall be available for inspection upon request of the Board or its inspectors. This Policy and Procedure Manual shall include the following:
  - 1. Operation of the Remote Pharmacy Practice;
  - 2. Maintenance of security for Remote Pharmacy Practice;
  - 3. Procedure to ensure that Remote Pharmacy Practice shall be conducted in a manner in which patient privacy and confidentiality is maintained, including a provision that patient information may not be printed; and
  - 4. A detailed list of all pharmacists engaged in Remote Pharmacy Practice and contact information for each pharmacist.
- i. The prescription drug outlet or other outlet shall enter into Written Agreements with each pharmacist engaged in Remote Pharmacy Practice, detailing all conditions, and policies and procedures governing Remote Pharmacy Practice. Such agreement shall be reviewed, updated and revised no less than annually. Such agreement shall be maintained at the prescription drug outlet or other outlet and be available for inspection upon request of the Board or its inspectors; and
- j. A pharmacist conducting Remote Pharmacy Practice is responsible for ensuring that his or her Remote Pharmacy Practice is conducted in accordance with the Policy and Procedure Manual and Written Agreement.

- 26.00.40. Equipment. All equipment, including computer equipment, utilized for Remote Pharmacy Practice shall meet at least the following requirements:
- a. Computer equipment must be able to establish a secure internet connection;
  - b. Equipment must be configured so patient information may not be stored at the site at which Remote Pharmacy Practice is occurring;
  - c. Access to the site for Remote Pharmacy Practice must be locked or shut down when the pharmacist ceases to be engaged in Remote Pharmacy Practice; and
  - d. Security parameters must prevent unauthorized storage or transfer of patient information.
- 26.00.50 Requirements for Conducting Final Evaluation. If Remote Pharmacy Practice includes conducting the final evaluation of prescriptions for a prescription drug outlet or other outlet, the following requirements apply:
- a. The pharmacist must have a visual connection with the prescription drug outlet or other outlet for the pharmacist to review the finished product prior to delivery to the patient; and
  - b. The prescription drug outlet or other outlet shall maintain records of final evaluation in accordance with all other applicable recordkeeping requirements.

**27.00.00 HOSPITAL SATELLITE PHARMACY.**

27.00.10 Definitions.

- a. "Hospital satellite pharmacy (HSP)" is a pharmacy located in a facility under the same management or control as the building or site where the hospital's Primary pharmacy is located, has a different address than the primary pharmacy, and is housed in a building with a main entrance that is no more than one (1) mile from the main entrance to the building which houses the primary pharmacy. Hospital satellite pharmacies may stock drugs at areas of the building where the hospital pharmacy is located, provided the areas are under the same management or control as the building or site where the hospital's primary pharmacy is located.
- b. "Primary pharmacy" is a registered prescription drug outlet in the hospital where the principal compounding/dispensing area is located.

27.00.20 Registration requirements.

- a. Hospitals which own or operate a pharmacy shall register all HSPs.
- b. The primary pharmacy shall submit an application on a form provided by the Division of Professions and Occupations on behalf of the HSP and for any drug storage satellites at the same location as the HSP.
- c. HSPs and any drug storage satellites placed at the same location as the HSP must pass a pre-registration inspection by the Board or its inspectors prior to registration.
- d. Any existing HSP or drug storage satellite at the same location as the HSP which is being remodeled or is being moved from one area of the location of the HSP to another shall submit documentation required by the Board prior to remodeling or moving.

- e. The compounding/dispensing area of an HSP shall not be less than 100 continuous square feet and must be approved by the Board prior to use for the practice of pharmacy.
- f. Any room included within or adjacent to the compounding/dispensing area that is separated from the compounding/dispensing area by a door must meet the following:
  - 1. The HSP shall submit documentation required by the Board to remodel the compounding/dispensing area prior to the utilizing the room for the purposes of compounding and dispensing or for the storage of prescription drugs and controlled substance stocks;
  - 2. The door must have a conspicuously displayed sign attached to it, and facing the compounding/dispensing area, that states "This room is part of the approved designated compounding/dispensing area";
  - 3. Unless the door is used to secure a room dedicated to storing controlled substances, it shall not have the ability to be locked or otherwise secured. The Board or its representatives shall have readily available and unimpeded access to this room at all times during normal business hours; and
  - 4. If a locked or otherwise secured door is used to secure a room dedicated to storing controlled substances, it shall be unlocked immediately upon the request of the Board or its representatives.
- g. Up to two satellites at the same location as the HSP may be used solely for storage of prescription drugs and controlled substances. Such drug storage satellites must possess square footage commensurate for the safe storage and removal of drugs within the affected satellites and approved by the Board prior to use.
- h. All HSPs and all satellites shall be well-lighted and well-ventilated with clean and sanitary surroundings devoted primarily to compounding/dispensing or drug storage. These areas shall provide necessary protection for drugs, chemicals and devices from deterioration due to light, heat or evaporation and shall be arranged to protect all prescription drugs and devices from pilferage or other unauthorized removal. No areas shall be subject to any condition likely to lead to errors.
- i. If the HSP engages in compounding/dispensing, there shall be a minimum of 12 continuous square feet of compounding/dispensing area, and a minimum of 6 continuous square feet of compounding/dispensing area for each person engaged in compounding/dispensing. These counters and surfaces shall be kept free and clear at all times for the purpose of compounding/dispensing. Any computer workstation or other equipment for the preparation of prescription labels and/or storage and retrieval of records shall be in addition to the minimum free compounding/dispensing area.
- j. The free floor space behind any compounding/dispensing counters or work surfaces shall be not less than 30 inches in width;
- k. The free floor space between rows of shelving shall be not less than 24 inches;
- l. If the HSP engages in compounding/dispensing, there shall be sufficient shelf, drawer and/or cabinet space for proper storage of prescription drugs and devices.
- m. If the HSP engages in compounding/dispensing, there shall be a sink, equipped with running hot and cold water, which is attached to an approved drain, waste and vent system, or to a portable enclosed tank which is emptied as frequently as necessary.

- n. Any other professional and technical equipment appropriate and adequate for the type of practices the HSP engages in shall at all times be located within the compounding/dispensing area.
- o. If refrigerated drugs are stored at the HSP or drug storage satellite, there shall be a refrigerator, dedicated to storing only drugs, meeting the compendia requirements and with an accurate thermometer in the refrigerator. The temperature shall be maintained between two and eight degrees Celsius (2 and 8 degrees C.) or thirty-six and forty-six degrees Fahrenheit (36 and 46 degrees F.). The temperature shall be electronically monitored each calendar day. Records detailing instances in which temperatures fall outside the aforementioned range requirement, for any period of time, shall be maintained at the HSP and shall be made readily available for inspection upon request by the Board or its representatives for a period of at least two years preceding the request. Such records shall include the duration of time the temperature fell outside the aforementioned range requirement, based on the best available data, and measures taken by the outlet as a result of the temperature falling outside the aforementioned range requirement.
- p. If frozen drugs are stored at the HSP or drug storage satellite, there shall be a freezer, dedicated to storing only drugs, meeting the compendia requirements and with an accurate thermometer in the freezer. The temperature shall be maintained between twenty-five degrees below zero and ten degrees below zero Celsius (– 25 and – 10 degrees C.) or thirteen degrees below zero and fourteen degrees Fahrenheit (– 13 and 14 degrees F.). The temperature shall be electronically monitored each calendar day. Records detailing instances in which temperatures fall outside the aforementioned range requirement, for any period of time, shall be maintained at the HSP and shall be made readily available for inspection upon request by the Board or its representatives for a period of at least two years preceding the request. Such records shall include the duration of time the temperature fell outside the aforementioned range requirement, based on the best available data, and measures taken by the outlet as a result of the temperature falling outside the aforementioned range requirement.
- q. There shall be a professional reference library available in the HSP. If an electronic library is provided, workstations must be provided in the compounding/dispensing area and must be readily available for use by staff, interns and Board personnel. This library shall contain current copies of the following:
  - 1. CRS Title 12, Article 42.5; the Pharmacists, Pharmacy Businesses, and Pharmaceuticals Act.
  - 2. CRS Title 18, Article 18, the Uniform Controlled Substances Act of 1992;
  - 3. Board of Pharmacy Rules;
  - 4. 21 Code of Federal Regulations (“CFR”) Part 1300 to End containing Drug Enforcement Administration rules relating to controlled substances;
  - 5. If compounding sterile products, the Guide to Parenteral Admixtures or Handbook on Injectable Drugs or other comparable references as determined by the pharmacist manager;
  - 6. If compounding cytotoxic products, Technical Manual Section VI: Chapter 2, Controlling Occupational Exposure to Hazardous Drugs or ASHP Technical Assistance Bulletin on Handling Cytotoxic and Hazardous Drugs; and



7. Any other references that the pharmacist manager of the primary pharmacy may deem necessary.

**27.00.30 Requirements for operation of an HSP**

- a. The pharmacist manager of the primary pharmacy shall have responsibility for the operation and control of the HSP;
- b. For the purpose of recordkeeping, drug stocks of the HSP shall be included in the inventory of the primary pharmacy;
- c. All records from the HSP shall be maintained at the primary pharmacy in accordance with Rule 11.00.00;
- d. Pharmacist staffing at the HSP cannot be considered in the computation of the pharmacists to pharmacy technician ratio in the primary pharmacy;
- e. Pharmacist staffing at the primary pharmacy cannot be considered in the computation of the pharmacists to pharmacy technician ratio in the HSP;
- f. The primary pharmacy may distribute drugs to the HSP in the same manner it would to other units of the hospital, and records shall be maintained in accordance with Rule 11.07.10;
- g. Every HSP shall display in the HSP compounding/dispensing area the report of the most recent inspection conducted by the Board or a photocopy of the most recent self-inspection performed by the pharmacist manager using the form provided by the Board, whichever is more recent; and
- h. No person other than a pharmacist or intern employed by the HSP shall be permitted in the compounding/dispensing area without the consent of the pharmacist in charge of the compounding/dispensing area.

**27.00.40 Minimum Hours of Operation**

- a. The principal compounding/dispensing area of an HSP shall be open for normal business a minimum of two designated days per week (Monday through Sunday) and at least four continuous hours on each such designated day.
- b. In the event that the principal compounding/dispensing area is open less than 32 hours per week, the pharmacist manager shall submit to the Board a written statement of the designated days and hours when the principal compounding/dispensing area will be open for business, and this statement shall be submitted at least 30 days prior to the date on which the hours of operation will be less than 32 hours per week.

**27.00.50 Security. All HSPs and additional satellitesshall comply with this rule.**

- a. When any compounding/dispensing area of an HSP is occupied by any employee, a pharmacist must be physically present within the same building of the HSP.
- b. In the event a pharmacist is within the building but absent from a compounding/dispensing area, it is the responsibility of the pharmacist to ensure the proper safeguard of all drugs.

- c. If a compounding/dispensing area is continually attended by a pharmacist when other people are in the building, the compounding/dispensing area need not be enclosed. However, if other people are in the building when there is not a pharmacist present, every compounding/dispensing area must be enclosed by a barrier as specified in paragraph (e) below.
- d. If more than one HSP is located within the same building, a pharmacist shall not operate more than one outlet at the same time. If a pharmacist physically leaves one outlet for the purpose of entering into another outlet within the same building, any outlet not being physically attended to by a pharmacist shall be enclosed by a barrier as specified in paragraph (e) below and a non-pharmacist shall not remain inside the enclosed outlet during that time.
- e. An HSP constituting part of a large establishment may be closed while the balance of the establishment is open for business, provided every compounding/dispensing area is enclosed with a secure floor-to-ceiling physical barrier, which shall be a divider or secure total enclosure, in which any openings shall not be large enough to permit removal of items from the compounding/dispensing area. The barrier must be of weight and strength sufficient to prevent it from being readily lifted, removed, penetrated or bent.
- f. All entrances to every compounding/dispensing area shall be secured from unauthorized entry when the pharmacist leaves the building where the HSP is located. No one other than a pharmacist shall be permitted to enter any compounding /dispensing area except in extreme emergencies, which shall be defined as a threat to property, public disaster or other catastrophe whereby the public is better served by overlooking the security restrictions of drugs and devices. If any compounding/dispensing area is opened in the absence of a pharmacist or left unsecured from unauthorized entry when the pharmacist leaves the building, the pharmacist manager shall notify the Board in writing within ten days of the discovery of the occurrence. This written notice shall state:
  - 1. The name of the person authorizing the opening of the compounding/dispensing area if known, or the name of the pharmacist responsible for securing the compounding/dispensing area from unauthorized entry;
  - 2. The name of the person opening the compounding/dispensing area if known; and
  - 3. A description of the situation requiring opening of the compounding/dispensing area including the date and time of the opening.
- g. While the compounding/dispensing area is closed and the rest of the building where the HSP is located is open, a person on duty in the building shall be able to contact a pharmacist in case of emergency.
- h. No HSP shall avail itself of the privileges of this rule until the barrier system and other requirements have been acknowledged, subject to final approval by the Board.

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- i. Procedures to follow in an emergency situation when a pharmacist is not in the building where the HSP is located are as follows:
    - 1. In an emergency situation and when a pharmacist is not in the building where the HSP is and administration of a drug to, or use of a device by or on, a patient is necessary pursuant to a chart order, and such drug or device is only available from a locked compounding/dispensing area, an authorized registered nurse may enter a locked compounding/dispensing area to obtain the drug or device. In the case of a drug, only pre-labeled packages, such as unit dose or unit-of-use packages, or a pre-labeled containers, may be removed from the compounding/dispensing area.
    - 2. The following information regarding the removal of such drug or device shall be consistently recorded and maintained in a retrievable document: date; time; name, strength and dosage form of drug, and/or name, and size, if applicable, of device; total quantity of drug or device removed; name and location of patient for whose use the drug or device is necessary; name of the practitioner ordering the drug or device; and the initials or signature of the nurse obtaining the drug or device. This document shall be available for inspection by the Board for a period of two years. Additionally, the original, duplicate or electronic or mechanical facsimile of the chart order shall be left with the above document by the nurse at the time of obtaining the drug or device.
    - 3. Any unused portion of a drug or device so removed shall be returned to the compounding/dispensing area when a pharmacist returns to the building. Additional quantities of the drug or device shall be supplied by a pharmacist and properly recorded as required by law and rule.
- 27.00.70 Relocation. In the event of a relocation of an HSP, the primary pharmacy shall submit an application on a form provided by the Division of Professions and Occupations along with the required fee at least thirty days prior to the effective date of relocation.
- 27.00.80 Reinstatement of an HSP registration. If a registration of an HSP has expired, the primary pharmacy shall submit a reinstatement application on a form provided by the Division of Professions and Occupations along with the required fee.
- 27.00.90 Closure.
- a. Closure shall mean the permanent cessation of the practice of pharmacy in any HSP. Closure shall also be deemed to have occurred if the compounding/dispensing area is not open for business the minimum hours specified in 27.00.40.
  - b. Upon the closure of the HSP, it shall be the responsibility of the pharmacist manager of the primary pharmacy to relocate the chart orders and drugs to the primary pharmacy. Such relocation of records shall be made within 72 hours after closure of the HSP. The pharmacist manager shall notify the Board on a form provided by the Board, detailing the closure of the HSP within 72 hours after closure. If the pharmacist manager fails to relocate the drugs and records as required herein, the Board may direct the removal of the drugs and records to a suitable location.
- 28.00.00 VETERINARY PHARMACEUTICAL ADVISORY COMMITTEE.**
- 28.00.10 Definitions.
- a. "Board" means the Colorado State Board of Pharmacy.
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- b. “Veterinary Pharmaceutical” means a prescription drug that is any of the following:
  - (1) Intended solely for animal use;
  - (2) Distributed for animal use;
  - (3) Dispensed for animal use;
  - (4) Administered to an animal.
- c. “Veterinary Pharmaceutical Advisory Committee (Advisory Committee)” is a committee comprised of three (3) members, each appointed by the state veterinarian, which reviews matters concerning veterinary pharmaceuticals, as specified by this Board Rule 28.00.00, referred to it by the Board and which makes recommendations on how the Board should proceed on the matters.

28.00.20 Matters Referred by the Board to the Advisory Committee Specified. Unless a matter presented to the Board constitutes an emergency requiring prompt resolution, the Board shall refer the following matters that directly concern veterinary pharmaceuticals to the Advisory Committee for recommendation on how the Board should proceed on the matter:

- a. Whether and to what extent action, if any, should be taken on an investigation into or complaint of an alleged violation of Article 42.5 of Title 12, C.R.S. as it directly pertains to the distribution, dispensation or administration of a veterinary pharmaceutical to an animal, including whether to:
  - (1) Suspend or revoke a license or registration;
  - (2) Impose a fine against a registrant, whether the violation is egregious, and the amount of any fine recommended;
  - (3) Seek a cease and desist order or injunction in district court against an entity or person; or
  - (4) Pursue other disciplinary action against a licensee or registrant.
- b. Review of license and registration applications and renewal, reactivation, and reinstatement applications when there is evidence the applicant directly engages in the distribution, dispensation, or administration of veterinary pharmaceuticals solely to animals; and,
- c. Promulgation of rules as they pertain to the distribution, dispensation, or administration of veterinary pharmaceuticals solely to animals.

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### **Editor’s Notes**

#### **History**

Rules 2.01.10; 2.01.30; 3.00.50; 3.00.70, 6.00.20; 6.00.30; 6.00.40; 8.00.10; 11.04.20; 14.03.10 eff. 07/30/2007.

Rules 8.00.10; 11.04.10; 20.00.00 eff. 09/30/2007.

Rule 4.00.00 eff. 11/30/2007.

Rules 3.01.20, 10.00.00 eff. 03/01/2008.

Rules 5.01.31; 15.01.11; 15.01.12; 15.09.11; 15.09.14; 22.00.00 eff. 05/30/2008.

Rules 4.02.00 (c), 21.00.00, 23.00.00 eff. 06/30/2008.

Rules 1.00.00, 2.00.00, 3.00.00, 5.00.00, 7.00.00, 11.00.00, 12.00.00, 14.00.00 eff. 11/30/2008.

Rule 15.09.11 eff. 01/31/2009.

Rules 6.00.30, 11.06.00, 22.00.00 eff. 03/02/2009.

Rule 9.00.00 eff. 04/30/2009.

Rules 5.00.55, 5.01.31(a), 6.00.20(f), 14.00.40, 15.01.17, 15.01.18, 15.08.19(f), 15.09.11(d), 15.09.15, 15.09.19, 15.09.20(g-h), 15.09.23, 15.09.24, 15.10.10, 16.00.20(d), 19.01.10(b), 19.01.30(a) eff. 12/30/2009.

Rules 4.00, 18.00 eff. 03/17/2010.

Rules 3.00.80 – 3.00.90; 5.00.55; 15.01.12; 19.00.00 – 19.01.50. Rule 22.00.00 repealed eff. 07/15/2010.

Rules 1.00.21, 5.01.31(e), 5.01.50 eff. 08/30/2010.

Rules 5.00.55, 21.11.10 (a), 21.21.70 (a) eff. 11/14/2010.

Rules 1.00.18, 2.01.50 – 2.01.53, 3.00.50 – 3.00.51, 5.00.50, 5.00.60, 5.01.31.a, 11.04.10, 15.01.11, 15.09.11.e eff. 06/14/2011.

Rules 3.01.24, 4.00.00, 11.04.20, 11.04.30, 21.00.00 - 21.11.20, 23.00.00 eff. 04/14/2012.

Rule 14.00.10 eff. 05/15/2012.

Entire rule eff. 01/01/2013. Rule 17.00.00 repealed eff. 01/01/2013.

Rules 3.00.21 – 3.00.22, 3.00.55, 3.00.90.e.(4), 3.01.20.c, 3.01.30, 3.01.32, 3.01.34, 4.00.10.f, 4.00.20, 5.01.31.a.(1)(C), 15.10.14.a, 23.00.90 eff. 09/14/2013.

Rules 2.01.10, 3.00.25, 3.00.91, 5.00.15, 6.00.30, 10.00.00, 11.03.00, 11.07.10, 14.00.05.k-l, 14.00.80.e.(2), 14.00.80.j, 16.00.00, 18.00.00, 20.00.00, 21.00.20, 21.10.80, 21.11.00.a.(12), 21.11.10.c, 21.20.20, 21.20.30.b(14), 21.21.40.c, 21.21.70.c, 21.22.00.b(1), 23.00.30, 23.00.50, 23.00.65, 23.00.70, eff. 10/15/2014.

Rules 3.00.22, 3.00.81.l-o, 3.00.82-3.00.84, 3.00.85.a(3), 3.00.86, 3.00.88.a(2), 3.00.88.b(10), 4.06.00, 6.00.10-6.00.20, 6.00.40.a, 6.00.50, 6.00.60.a, 6.00.60.b.10, 6.00.70.a, 6.00.90.b, 6.01.10.a, 19.01.40.c, 21.00.10, 21.00.20.b, 21.10.60.b, 21.10.80.b(4), 21.11.10.a(5), 21.11.10.c(9), 21.20.10.d, 21.20.20.b(2)(a), 21.20.25.b, 21.20.70.f, 21.20.90.b-c, 21.21.10.b, 21.21.70.a(6), 21.21.70.c(10), 23.00.40.y-z, 23.00.70.h-j eff. 09/14/2015.

Rules 3.00.21, 3.00.27, 19.01.10(1), 21.00.20, 21.11.20.d, 21.20.16, 21.20.20.b.(2), 21.20.60.b, 21.20.60.e, 21.21.90.d eff. 03/16/2016.

Rules 3.00.20, 3.00.22 e, 3.00.81 g, 3.00.84, 3.01.10 d, 4.00.10, 4.00.25, 4.05.00, 5.00.15 d, 5.01.31, 6.00.20 e, 7.00.10, 8.00.10, 14.00.80 i-k, 19.01.10 b.(2), 20.00.80 a.1, 21.00.20, 21.00.30, 21.20.20 b, 27.00.00, 28.00.00 eff. 11/14/2016. Rule 10.00.51 repealed eff. 11/14/2016.