

475:10-1-4. Separate registration

(a) Every person who engages in, or who proposes to engage in, more than one group of independent activities shall obtain a separate registration for each group of activities, except as provided by this subsection. Any person, when registered to engage in the group of activities described in each paragraph of this subsection, shall be authorized to engage in the coincident activities described in that subparagraph without obtaining a registration to engage in such coincident activities; provided that, unless specifically exempted, he/she complies with all requirements and duties prescribed by law for persons registered to engage in such coincident activities.

(1) A person registered to manufacture any controlled dangerous substance or basic class of controlled dangerous substances shall be authorized to distribute that substance or class, but is not authorized to distribute any substance or class which he/she is not registered to manufacture.

(2) A person registered to manufacture any controlled dangerous substance listed in Schedules II-I through V shall be authorized to conduct chemical analysis and preclinical research (including quality control analysis) with narcotic and non-narcotic controlled dangerous substances listed in those schedules which he/she is authorized to manufacture.

(3) A person registered to conduct scientific research with a basic class of controlled dangerous substances listed in Schedule I shall be authorized to manufacture such substances if and to the extent that such manufacture is set forth in the research protocol and to distribute such substances to other persons registered or authorized to conduct scientific research with such substances or registered or authorized to conduct chemical analysis for scientific purposes with controlled dangerous substances provided such distribution is made in conformance with state law.

(4) A person registered or authorized to conduct analysis for scientific purposes with controlled dangerous substances shall be authorized to manufacture such substances for analytical or instructional purposes, to distribute such substances to other persons registered or authorized to conduct scientific analysis for scientific purposes or instructional activities or scientific research with such substances and to persons exempted from registration provided such distribution is made in conformance with state law.

(5) A person registered or authorized to conduct scientific research with controlled dangerous substances listed in Schedules II through V shall be authorized to conduct chemical analysis for scientific purposes with controlled dangerous substances listed in those schedules in which he/she is authorized to conduct scientific research, to manufacture such substances if and to the extent that such manufacture is set forth in the protocol filed with the application for registration, to distribute such substances to other persons registered or authorized to conduct chemical analysis for scientific purposes, instructional activities, or scientific research with such substances and to persons exempted from registration provided such distribution is made in conformance with state law, and to conduct instructional activities with controlled dangerous substances.

(6) Physicians, dentists, podiatrists, veterinarians, optometrists and other qualified persons who are authorized to carry on their respective activities under the laws of the State of Oklahoma and who are registered with the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control to dispense, prescribe or administer controlled dangerous substances shall be authorized to conduct instructional activities with those substances.

(7) Trainers or handlers of a canine controlled dangerous substance detector who, in the ordinary course of their profession, desire to possess any controlled dangerous substance for training said canine.

(8) A single registration to engage in any group of independent activities may include one or more controlled dangerous substances listed in the schedules authorized in that group of independent activities. A person registered to conduct scientific research with controlled dangerous substances listed in Schedule I may conduct scientific research with any substance listed in Schedule I for which he/she has filed and had approved a scientific research protocol.

(b) The following locations shall not be deemed to be principal places where controlled dangerous substances are manufactured, distributed, dispensed or prescribed:

(1) A warehouse where controlled dangerous substances are stored by or on behalf of a registered person, unless such substances are distributed directly from such warehouse to registered locations other than the registered location from which the substances were delivered or to persons not required to register.

(2) An office used by agents of a registrant where sales of controlled dangerous substances are solicited, made or supervised but which neither contain such substances (other than substances for display purposes or lawful distribution as samples only) nor serves as a distribution point for filling sales orders.

(3) An office used by a practitioner (who is registered at another location) where controlled dangerous substances are prescribed but neither administered nor otherwise dispensed as a regular part of the professional practice of the practitioner at such office, and where no supplies of controlled dangerous substances are maintained.

475:10-1-18. Certificate of registration form

The Certificate of Registration shall contain the name, business address, and registration number of the registrant, the schedules of controlled dangerous substances which the registrant is authorized to handle, any limitation or condition placed on the registration, and the expiration date of the registration. The registrant shall maintain the Certificate of Registration at the registered location and shall permit inspection of the Certificate by a peace officer or agency official in the enforcement of laws relating to controlled dangerous substances.

475:20-1-3. Physical security controls for nonpractitioners; storage areas

Physical security controls for nonpractitioners and storage areas shall comply with Title 21 Code of Federal Regulations §1301.72-, except physical security controls for medical marijuana retailers shall, at a minimum, meet the following requirements for each retail storage area:

(1) Each registered premises shall have a Security Alarm System which upon unauthorized entry shall transmit a signal directly to a central station protection agency or a local state or state police agency, each having a legal duty to respond, or to a 24-hour control station operated by the registrant, or to such other source of protection as the Director may approve.

(2) All retail storage areas shall be equipped with self-closing, self-locking doors constructed of substantial material commensurate with the type of building construction, provided, however, a door which is kept closed and locked at all times when not in use and when in use is kept under direct observation of a responsible employee or agent of the registrant is permitted in lieu of a self-closing, self-locking door. Doors may be sliding or hinged. If doors hinges are mounted on the outside, such hinges shall be sealed, welded or otherwise constructed to inhibit removal. Locking devices for such doors shall be either of the multiple-position combination, keyless entry, or key lock type and:

(A) In the case of key locks, shall require key control which limits access to a limited number of employees, or;

(B) In the case of multiple-position combination or keyless entry systems, the system shall be limited to a minimum number of employees and can be changed upon termination of employment of an employee having knowledge of the combination.

(3) The retail storage areas shall be accessible only to an absolute minimum number of authorized employees. When it is necessary for employee maintenance personnel, nonemployee maintenance personnel, business guests, or visitors to be present in or pass through controlled substances storage areas, the registrant shall provide for adequate observation of the area by an employee specifically authorized in writing.

475:20-1-4. Physical security controls for nonpractitioners; manufacturing areas

Physical security controls for nonpractitioners and manufacturing areas shall be in compliance with Title 21 Code of Federal Regulations §1301.73, except physical security controls for medical marijuana commercial growers, processors, packagers, and manufacturers shall, at a minimum, meet the following requirements:

(1) All in-process medical marijuana shall be returned to the storage area at the termination of the process. If the process is not terminated at the end of a workday (except where a continuous process or other normal manufacturing operation should not be interrupted), the processing area or tanks, vessels, bins or bulk containers containing medical marijuana shall be securely locked, with adequate security for the area or building.

(2) Each building shall require an alarm, that upon unauthorized entry, shall transmit a signal directly to a central station protection company, or local or state police agency that has a legal duty to respond, or a 24-hour control station operated by the registrant, or to such other source of protection as the Director may approve.

(3) Each building shall be equipped with self-closing, self-locking doors constructed of substantial material commensurate with the type of building construction, provided, however, a door which is kept closed and locked at all times when not in use and when in use is kept under direct observation of a responsible employee or agent of the registrant is permitted in lieu of a self-closing, self-locking door. Doors may be sliding or hinged. If doors hinges are mounted on the outside, such hinges shall be sealed, welded or otherwise constructed to inhibit removal. Locking devices for such doors shall be either of the multiple-position combination, keyless entry, or key lock type and:

(A) In the case of key locks, shall require key control which limits access to a limited number of employees, or;

(B) In the case of multiple-position combination or keyless entry systems, the system shall be limited to a minimum number of employees and can be changed upon termination of employment of an employee having knowledge of the combination.

(4) Any outdoor or greenhouse facilities shall provide adequate security measures for the area or building including the following:

(A) The entire outdoor or greenhouse facility shall be surrounded by a fence and entry gates. The fence shall measure at least eight (8) feet from the ground to the top of the fence and shall be constructed of at least six (6) gauge or higher metal chain link fence or another similarly secure material or wood. All support posts shall be steel and securely anchored.

(B) All entry gates shall measure at least eight (8) feet from the ground to the top of the entry gate and shall be constructed of six (6) gauge or higher metal chain link fence or a similarly secure material or wood.

(C) The fence shall be in good repair and obscure the Limited Access Area so that it is not easily viewed from outside the fence.

(5) The medical marijuana commercial growing, processing, packaging, and manufacturing areas shall be accessible only to an absolute minimum number of authorized employees. When it is necessary for employee maintenance personnel, nonemployee maintenance personnel, business guests, or visitors to be present in or pass through controlled substances storage areas, the registrant shall provide for adequate observation of the area by an employee specifically authorized in writing.

475:20-1-5. Other security controls for nonpractitioner registrants

- (a) Before distributing a controlled dangerous substance to any person whom the registrant does not know to be registered to possess the controlled dangerous substance, the registrant shall make a good-faith inquiry either with the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control or with the Drug Enforcement Administration, or when applicable, the Oklahoma Medical Marijuana Authority, to determine that the person is registered to possess the controlled substance.
- (b) The registrant shall design and operate a system to disclose to the registrant suspicious orders of controlled dangerous substances. The registrant shall inform the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control of suspicious orders when discovered by the registrant. Suspicious orders include orders of unusual size, orders deviating substantially from a normal pattern, and orders of unusual frequency.
- (c) All registrants shall notify the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control of any theft or significant loss of any controlled dangerous substances upon discovery of such theft or loss. Notification shall be made in writing and shall contain a list of the substances stolen or diverted by their trade name, quantities, descriptions, amount lost or stolen, and any cost code marks utilized. Thefts must be reported whether or not the controlled dangerous substances are subsequently recovered and/or the responsible parties are identified and action taken against them.
- (d) No person acting as an agent of a registered controlled dangerous substances manufacturer or distributor (i.e., detailman, salesman, etc.), or a medical marijuana commercial grower, processor, or manufacturer, shall distribute samples of controlled dangerous substances to a practitioner without first having been registered (no fee required) with the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control.
- (1) To register with the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control to distribute samples of controlled dangerous substances a form must be filled out and submitted to the Registration Department. Such forms may be obtained through the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control website or by calling the Registration Department.
 - (2) A new form shall be completed and submitted to the registration department each time the list of items to be distributed changes.
 - (3) A copy of the form submitted to the Oklahoma State Bureau of Narcotics shall be retained by the distributor.
 - (4) The practitioner receiving the samples shall keep a record each time he/she receives or distributes samples of controlled dangerous substances.
- (e) When shipping controlled dangerous substances, a registrant is responsible for selecting common or contract carriers which provide adequate security to guard against in-transit losses. When storing controlled dangerous substances in a public warehouse, a registrant is responsible for selecting a warehouseman which will provide adequate security to guard against storage losses; wherever possible, the registrant shall store controlled dangerous substances in a public warehouse which complies with the requirements set forth in this Chapter. In addition, the registrant shall employ precautions (e.g., assuring that shipping containers do not indicate that contents are controlled dangerous substances except in the case of medical marijuana) to guard against storage or in-transit losses and comply with all current Federal regulations, except medical marijuana transit shall comply with rules set forth in OAC 310:681-3. Reporting the loss of in-transit shipments is the responsibility of the registrant shipping the controlled dangerous substances.
- (f) When distributing controlled dangerous substances through agents (e.g., detailmen), a registrant is responsible for providing and requiring adequate security to guard against theft and diversion while the controlled dangerous substances are being stored or handled by the agent(s).
- (g) No registrant shall knowingly employ as an agent or employee any person who will have access to controlled dangerous substances if such person has been convicted, pled guilty or nolo contendere or otherwise ordered to complete a period of probation or supervision for a misdemeanor or felony relating to any controlled dangerous substances as defined by the Uniform Controlled Dangerous Substances Act

in this state, any other state, or the United States, or any person convicted, pled guilty or nolo contendere or otherwise ordered to complete a period of probation or supervision for any felony of this state, any other state, or the United States, unless, after full review of the circumstances, the Director waives this requirement in writing with respect to each person on a case-by-case basis. However, Schedule I medical marijuana registrants, employees, and agents shall be subject to the criminal history requirements pursuant to Title 63 Okl.St. Ann. §420A et seq., unless, after full review of the circumstances, the Director waives this requirement in writing with respect to each person on a case-by-case basis.

(h) The registrant shall immediately notify OBN and seek authorization to employ any individual as specified above.

475:25-1-2. General information

Registrants shall be required to maintain records, reports and inventory in accordance with this Chapter and pursuant to Title 21 Code of Federal Regulations, and Title 63 Okl.St. Ann. §2-307, except Schedule I medical marijuana registrants shall be required to maintain readily-retrievable inventory tracking, records, and reports in the format set forth in OAC 310:681-5-6.

475:25-1-4. Maintenance of records and inventories

(a) Every inventory and other record required to be kept by the Uniform Controlled Dangerous Substances Act and this Chapter shall be kept by the registrant ~~and be available~~ for at least two (2) years from the date of such inventory or record. Schedule I medical marijuana inventory and records shall be kept for at least seven (7) years from the date of such inventory or record. Every inventory and other record required to be kept shall be available for inspecting and copying by authorized peace officers or officers of agencies specifically directed to enforce the State of Oklahoma or the United States controlled dangerous substances laws, pursuant to and in the manner prescribed by Title 63 Okl.St. Ann. § 2-502, Title 21 Code of Federal Regulations § 1304.04, and this Chapter.

(b) Each registered manufacturer and distributor shall maintain inventories and records of controlled substances as follows:

(1) Inventories and records of controlled dangerous substances listed in Schedules I and II shall be maintained separately from all other records of the registrant.

(2) Inventories and records of controlled dangerous substances listed in Schedules III, IV and V shall be maintained separately from all other records of the registrant as of November 1, 1990.

(c) Each registered individual practitioner required to keep records and institutional practitioners required to keep records shall maintain inventories and records of controlled dangerous substances in the manner prescribed in (b) of this Section.

(d) Each registered pharmacy shall maintain the inventories and records of controlled dangerous substances as follows:

(1) Inventories, records, invoices and purchase records of all controlled dangerous substances listed in Schedules I and II shall be maintained separately from all other records of the pharmacy, and prescriptions for such substances shall be maintained in a separate prescription file and be readily retrievable.

(2) Inventories, records, invoices and purchase records of controlled dangerous substances listed in Schedules III, IV and V shall be maintained separately from all other records of the pharmacy and be readily retrievable. Prescriptions for such substances shall be maintained in separate prescription files for controlled dangerous substances listed in Schedules III, IV and V and shall be readily retrievable from the other prescription records of the pharmacy.

475:25-1-9. Inventories of manufacturers

Except for Schedule I medical marijuana registrants, Inventoriesinventories of manufacturers of controlled dangerous substances shall conform to Title 21 Code of Federal Regulations, §1304.11.

475:25-1-10. Inventories of distributors

Except for Schedule I medical marijuana registrants, Eacheach person registered or otherwise authorized to distribute controlled dangerous substances shall include in his/her inventory the same information required of a manufacturer pursuant to Title 21 Code of Federal Regulations, §1304.11.

475:25-1-14. Records for manufacturers

Except for Schedule I medical marijuana registrants, Recordsrecords for manufacturers of controlled dangerous substances shall conform to Title 21 Code of Federal Regulations, §1304.22.

475:25-1-15. Records for distributors

Each person registered or otherwise authorized to distribute controlled dangerous substances, except for Schedule I medical marijuana registrants, shall maintain records with the following information for each controlled dangerous substance:

- (1) The name of the substance.
- (2) Each finished form (e.g., 10-milligram tablet or 10-milligram concentration per fluid ounce or milliliter) and the number of units or volume of finished form in each commercial container (e.g., 100-tablet bottle or 3-milliliter vial).
- (3) The number of commercial containers of each such finished form received from other persons, including the date and number of containers in each receipt and the name, address and Federal Drug Enforcement Administration registration number of the person from whom the containers were received.
- (4) The number of commercial containers of each such finished form imported directly by the person, including the date of, the number of commercial containers in, and the import permit or declaration number for each importation.
- (5) The number of commercial containers of each such finished form distributed to other persons, including the date and number of containers in each distribution and the name, address and Federal Drug Enforcement Administration registration number of the person to whom the containers were distributed.
- (6) The number of commercial containers of each such finished form exported directly by the person, including the date of, the number of commercial containers in, and the export permit or declaration number for each exportation.
- (7) The number of units or volume of finished forms and/or commercial containers distributed or disposed of in any other manner by the person (e.g., by distribution as complimentary samples or by destruction), including the date and manner of distribution or disposal, the name, address and the Federal Drug Enforcement Administration registration number of the person to whom distributed, and the quantity of the substance in finished form distributed or disposed.

475:25-1-16. Records of scientific researchers

Each person registered or otherwise authorized to conduct scientific research with controlled dangerous substances and required to keep records shall maintain records with the following information for each controlled dangerous substance:

- (1) The name of the substance.
- (2) Each finished form (e.g., 10-milligram tablet or 10-milligram concentration per fluid ounce or milliliter) and the number of units or volume of finished form in each commercial container (e.g., 100-tablet bottle or 3-milliliter vial).
- (3) The number of commercial containers of each such finished form received from other persons, including the date of and number of containers in each receipt and the name, address and, if applicable, Federal Drug Enforcement Administration registration number of the person from whom the containers were received.
- (4) The number of units or volume of such finished forms and/or commercial containers disposed of in any other manner by the registrant, including the date and manner of disposal and the quantity of the substance in the finished form disposed.

475:25-1-17. Records of scientific analyst

(a) Each person registered or otherwise authorized to conduct scientific analysis with controlled dangerous substances shall maintain records with the following information to the extent known and reasonably ascertainable by him/her for each controlled dangerous substance:

- (1) The name of the substance.
- (2) The form or forms in which the substance is received, imported or manufactured by the registrant (e.g., powder, granulation, tablet, capsule or solution) and the concentration of the substance in such form (e.g., C.P., U.S.P., N.D., 10-milligram tablet or 10-milligram concentration per milli-liter).
- (3) The total number of the forms received, imported or manufactured (e.g., 100 tablets, 30 1-milliliter vials, or 10 grams of powder), including the date and quantity of each receipt, importation or manufacture and the name, address and Federal Drug Enforcement Administration registration number, if any, of the person from whom the substance was received.
- (4) The quantity distributed or destroyed in any manner by the registrant (except quantities used in chemical analysis or other laboratory work), including the date and manner of distribution or destruction and the name, address and Federal Drug Enforcement Administration registration number, if any, of each person to whom the substance was distributed.

~~(b) — Records of controlled dangerous substances used in chemical analysis or other laboratory work are not required.~~

~~(e)~~(b) Records relating to known or suspected controlled dangerous substances received as evidentiary material for analysis are not required under (a) of this Section.

~~(d)~~(c) Each person registered as a scientific analyst conducting scientific analysis of ~~anonymous~~ samples of suspected controlled dangerous substances shall maintain records containing the following information (to the extent known and reasonably ascertainable by him/her):

- (1) Laboratory identification number.
- (2) Date the sample received.
- (3) Purported contents and actual identification.

- (4) Quantity received.
- (5) Form of sample (i.e., powder, liquid, tablets, etc.).
- (6) Description of sample.
- (7) Quantity utilized in analysis.
- (8) Disposition of sample.
- (9) Street price, if known.
- (10) Method shipment is received.
- (11) Each laboratory shall submit to the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control a quarterly report containing at least the following information:
 - (A) Actual content of drug analyzed.
 - (B) Alleged content of drug analyzed.
 - (C) Description of sample.
 - (D) Origin of sample.
 - (E) Street price, if known.

~~(e)(d)~~ Qualitative and Quantitative analysis may be conducted of ~~anonymous~~ samples. ~~However, to prevent the possibility of illegal drug traffickers utilizing these laboratories as quality control, only qualitative results may be given to the donor. Analysis should be sufficient to determine if the strength is so great that use would be harmful to the user. In these cases, the submitter can only be told what the drug was and that use would be dangerous.~~

- (1) Security of standards and samples, including Schedule I medical marijuana, shall be in accordance with 475:20-1-6 and 475:20-1-7, with the exception that all standards and samples must be treated as Schedules I and II.
- (2) Any unused portion of a submitted ~~anonymous~~ sample shall be disposed of in accordance with 475:35-1-4.
- (3) All controlled dangerous substances distributed to canine handler registrants and scientific research registrants shall be analyzed quantitatively, and a record of such analysis shall be maintained prior to distribution. Oklahoma State Bureau of Investigation has discretion to refuse to distribute any controlled dangerous substances. Each such registrant shall receive a copy of the quantitative analysis.

475:30-1-1. Purpose

The rules of this Chapter describe the procedures to be followed for issuance of a valid prescription, and the information required to be placed on labels for controlled dangerous substances. Labeling for Schedule I medical marijuana shall be in accordance with OAC 310:681-7-1.

475:35-1-3. Distribution upon discontinuance or transfer of business

(a) Any registrant desiring to discontinue business activities altogether or with respect to controlled dangerous substances (without transferring such business activities to another person) shall return for cancellation of his/her Certificate of Registration. Any controlled dangerous substances in his/her possession may be disposed of in accordance with Title 21 Code of Federal Regulations, part 1317. Schedule I medical marijuana shall be disposed pursuant to standards set forth in the Uniform Controlled Dangerous Substances Act, 63 Okla.St. Ann. §2-101 et seq., and OAC 252:205.

(b) Any registrant desiring to discontinue business activities altogether or with respect to controlled dangerous substances (by transferring such business activities to another person) shall

submit in person or by registered or certified mail, return receipt requested, to the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control at least fourteen (14) days in advance of the date of the proposed transfer (unless the Director waives this time limitation in individual instances), the following information:

- (1) The name, address, registration number and authorized business activity of the registrant discontinuing the business (registrant-transferor).
 - (2) The name, address, registration number and authorized business activity of the person acquiring the business (registrant-transferee).
 - (3) Whether the business activities will be continued at the location registered by the person discontinuing the business or moved to another location (if the latter, the address of the new location should be listed).
 - (4) Whether the registrant-transferor has a quota to manufacture or procure any controlled dangerous substance listed in Schedule I or II (if so, the basic class or classes of the substance should be indicated).
 - (5) The date on which the transfer of controlled dangerous substances will occur.
- (c) Unless the registrant-transferor is informed by the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control, before the date on which the transfer was stated to occur, that the transfer may not occur, the registrant-transferor may distribute (without being registered to distribute) controlled dangerous substances in his/her possession to the registrant-transferee in accordance with the following:

- (1) On the date of transfer of the controlled dangerous substances, a complete inventory of all controlled dangerous substances being transferred shall be taken in accordance with 475:25-1-5 through 475:25-1-12. This inventory shall serve as the final inventory of the registrant-transferor and the initial inventory of the registrant-transferee, and a copy of the inventory shall be included in the records of each person. It shall ~~not~~ be necessary to file a copy of the inventory with the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control unless ~~requested~~ waived by the Director. Except for Schedule I medical marijuana, Transfer/transfers of any substances listed in Schedule I or II requires the use of order forms in accordance with Title 21 Code of Federal Regulations, § 1305.
- (2) On the date of transfer of the controlled dangerous substances, all records required to be kept by the registrant-transferor with reference to the controlled dangerous substances being transferred, pursuant to this Chapter and Title 21 Code of Federal Regulations, § 1304, or OAC 310:681-5-6, shall be transferred to the registrant-transferee. Responsibility for the accuracy of records prior to the date of transfer remains with the transferor, but responsibility for custody and maintenance shall be upon the transferee.

475:35-1-4. Procedure for disposing of controlled dangerous substances

Any registrant in possession of any controlled dangerous substances and desiring or required to dispose of such substances may do so according to the provisions of Title 63 Okl.St. Ann. §2-315 and Title 21 of the Code of Federal Regulations, part 1317-, except Schedule I medical marijuana shall be disposed pursuant to standards set forth in the Uniform Controlled Dangerous Substances Act, 63 Okla.St. Ann. §2-101 et seq., and OAC 252:205.

475:40-1-2. Authority to make inspections

Administrative inspections of OBN registrants shall include, but not be limited to, the following:

- (1) Inspecting, copying and verifying the correctness of records, reports or other documents required to be kept or made, including, but not limited to, inventory and other records required to be kept pursuant to the Uniform Controlled Dangerous Substances Act, this Title, ~~and~~ the Code of Federal Regulations governing controlled dangerous substances, or OAC 310:681-5-6; order form records required to be kept pursuant to Title 63 Okl.St. Ann. § 2-308 and other applicable state statutes and rules; prescriptions and distribution records required to be kept pursuant to Title 63 Okl.St. Ann. § 2-307 and other applicable state statutes and rules; shipping records identifying the name of each carrier used; and the date and quantity of each storage.
- (2) Inspecting within reasonable limits and in a reasonable manner all pertinent equipment, finished and unfinished controlled dangerous substances and other substances or materials, containers, and labeling found at the controlled premises relating to the Uniform Controlled Dangerous Substances Act and this Title.
- (3) Making a physical inventory of all controlled dangerous substances on hand at the premises.
- (4) Collecting samples of controlled dangerous substances or precursors (in the event any samples are collected during an inspection, the peace officer or officer so authorized shall issue a receipt for such samples to the owner, operator or agent in charge of the premises).