(b) The holder of a permit to operate a pharmacy department and the registered pharmacist-in-charge of the department shall comply with all requirements in this chapter and shall also be subject to the following additional requirements:

1. The pharmacy department shall be constructed so as to enable the closing off and securing of the department from the main store area. The department shall be separated from the main store area by a secured barrier or partition extending from the floor or fixed counter to the ceiling of either the department or main store and attached thereto. Any entrance to the pharmacy department shall be capable of being locked and connected to a security device or other Board approved security system.

2. The registered pharmacist on duty shall be responsible for keeping the pharmacy department secure and locked and the alarm system turned on at all times when he or she is not present within the department, except as provided in N.J.A.C. 13:39-6.4, and shall be responsible for the security of the keys to the department.

3. All medications requiring supervision of a pharmacist, including dispensed medication, shall remain within the confines of the department when the pharmacist is not in the pharmacy department.

4. The hours that the department is open and the name of the registered pharmacist-in-charge shall be posted in plain view at the entrance to the department and at the public entrance to the enterprise containing the department.

5. When the enterprise in which the department is located maintains different store hours from the pharmacy department, all advertising, announcements, signs or statements indicating store hours and the presence of the pharmacy department shall clearly and distinctly indicate the hours that the department is open.

6. The pharmacy department shall have a published telephone number different from that of the establishment in which the department is located. No extensions of this phone shall be located outside the department.

7. The telephone number of the registered pharmacist-in-charge shall be available in the office of the manager of the establishment.

8. There shall be provided a secure area for the receiving of prescription drugs from suppliers. No prescription drug shall be accepted from any supplier during the hours the pharmacy department is closed unless adequate security for the storage of department shipments has been provided.

9. If a drop-off device is utilized for prescriptions it shall be of a one-way, irretrievable design.

Amended by R.1994 d.351, effective July 18, 1994.
See: 26 N.J.R. 1596(a), 26 N.J.R. 2905(b).
Amended by R.1999 d.214, effective July 19, 1999.
See: 31 N.J.R. 1151(a), 31 N.J.R. 1932(a).
In (b), substituted references to registered pharmacists-in-charge for references to pharmacists-in-charge in 2 and 8.
Recodified from N.J.A.C. 13:39-4.15 and amended by R.2005 d.25, effective January 18, 2005.
See: 36 N.J.R. 3345(a), 37 N.J.R. 295(a).
Rewrote the section. Former N.J.A.C. 13:39-4.14, Contract pharmaceutical services, recodified to N.J.A.C. 13:39-9.4.

13:39-4.15 Permits; specialized permits

(a) The Board may issue a special permit, wherein the type of service is of a limited nature. The permit so issued, being based on special conditions of use imposed by the Board, may necessitate the waiver of certain rule requirements.

(b) Specialized permits shall pertain to pharmacies providing specific services as may be necessary and proper to efficiently meet a limited public need for pharmaceutical services. An applicant for any specialized pharmacy permit shall provide the Board with an application and a policy and procedure manual which sets forth a detailed description of the type of specialized pharmacy services to be provided within the pharmacy practice. The policy and procedure manual shall also contain detailed provisions which ensure the protection of the public welfare as determined by the Board.

Recodified from N.J.A.C. 13:39-4.16 by R.2005 d.25, effective January 18, 2005.
See: 36 N.J.R. 3345(a), 37 N.J.R. 295(a).
Former N.J.A.C. 13:39-4.15, Retail permit; prescription department or pharmacy department, recodified to N.J.A.C. 13:39-4.14.

13:39-4.16 Steering prohibited

It shall be unlawful for a pharmacy permit holder to enter into an arrangement with a health care practitioner who is licensed to issue prescriptions for the purpose of directing or diverting patients to or from a specified pharmacy or restraining in any way a patient's freedom of choice to select a pharmacy.

Recodified from N.J.A.C. 13:39-4.17 and amended by R.2005 d.25, effective January 18, 2005.
See: 36 N.J.R. 3345(a), 37 N.J.R. 295(a).
Rewrote the section. Former N.J.A.C. 13:39-4.16, Permits; specialized permits, recodified to N.J.A.C. 13:39-4.15.

13:39-4.17 Responsibilities of permit holders

(a) All permit holders shall be responsible for compliance with all the rules, regulations and laws governing the practice of pharmacy.

(b) Any permit holder may be held liable for violations of the Pharmacy Act, N.J.S.A. 45:14-1 et seq., and the rules in this chapter and may be subject to disciplinary action.

Recodified from N.J.A.C. 13:39-4.18 and amended by R.2005 d.25, effective January 18, 2005.
See: 36 N.J.R. 3345(a), 37 N.J.R. 295(a).
Rewrote the section. Former N.J.A.C. 13:39-4.17, Steering prohibited, recodified to N.J.A.C. 13:39-4.16.

13:39-4.18 Procedures for centralized prescription handling

(a) The four component functions of handling a prescription are intake, processing, fulfillment and dispensing.

(b) Central prescription handling entails two or more licensed pharmacies sharing responsibility for performing the four component functions of handling a prescription.

(c) The following pharmacies may engage in central prescription handling: an intake or originating pharmacy; a central processing pharmacy; a central fill pharmacy; and a dispensing pharmacy. The four component functions of handling a prescription shall be performed by the following pharmacies:

1. An intake or originating pharmacy, which is a licensed pharmacy that received the patient's or prescribing practitioner's request to fill or refill a prescription. A central processing pharmacy or a central fill pharmacy, as delineated in (c)2 and 3 below, may be considered the intake or originating pharmacy if the prescription was transmitted by the prescribing practitioner directly to the centralized pharmacy as provided in N.J.A.C. 13:39-5.8A and 5.8B or if the patient requested the refill from that pharmacy;

2. A central processing pharmacy, which is a licensed pharmacy that engages in prescription review by performing functions that may include, but are not limited to, data entry, prospective drug review, refill authorizations, interventions, patient counseling, claims submission, claims resolution and adjudication;

3. A central fill pharmacy, which is a licensed pharmacy engaging in central prescription handling by filling and/or refilling prescriptions which includes the preparation and packaging of the medication; and

4. A dispensing pharmacy, which is a licensed pharmacy that receives the processed prescription and/or the filled or refilled prescription for dispensing to the patient or to the patient's authorized representative.

(d) Two or more licensed pharmacies delineated in (c) above may engage in central prescription handling provided:

1. Any or all of the pharmacies participating in central prescription handling have a contractual agreement to provide such services or have the same owner;

2. Prior to engaging in central prescription handling, all pharmacies that are parties to the central prescription handling obtain Board approval. If a participating pharmacy is located outside the State of New Jersey, the pharmacy shall have registered with the Board pursuant to N.J.A.C. 13:39-4.19. The pharmacies shall make a single application to the Board, delineating the scope of practice of each pharmacy and the specific rules in this chapter with which each pharmacy shall comply;

3. An audit trail is maintained that records and documents the name(s) or other personal identifier(s) of the pharmacist(s) or pharmacy technician(s) and the component function(s) performed by each, at the time the functions are performed, for each step of prescription handling. The audit trail shall be maintained for not less than five years from the date the prescription is filled or refilled. The oldest four years of information shall be maintained in such a manner so as to be retrievable and readable within two weeks. The most recent one year of information shall be retrievable and readable within one business day;

4. The dispensed prescription for any product bears a permanently affixed label with at least the following information:

i. The brand name or generic name, and if generic, the name of the manufacturer;

ii. The strength of medication, where applicable;

iii. The quantity dispensed;

iv. The date upon which prescription medication is dispensed;

v. A CDS cautionary label, where applicable and when permitted by law;

vi. The patient name;

vii. The prescriber name;

viii. The prescription number;

ix. Directions for use;

x. The phrase "use by" followed by the product's use by date, if dispensed in any packaging other than the manufacturer's original packaging. For purposes of this paragraph, "use by date" means the earlier of one year from the date of dispensing or the expiration date on the manufacturer's container;

xi. All auxiliary labeling as recommended by the manufacturer and/or as deemed appropriate in the professional judgment of the dispensing pharmacist; and

xii. The name, address and telephone number of any or all of the following:

(1) The intake pharmacy;

(2) The central processing pharmacy;

(3) The central fill pharmacy; and/or

(4) The dispensing pharmacy;

5. The patient name, the brand or generic name of the medication, and the directions for use appear in larger type, in a different color type, or in bolded type, in comparison to the other information required to appear on the label of the dispensed container pursuant to (d)4 above;

6. The patient is provided with written information, either on the prescription label or with the prescription container, that indicates which pharmacy to contact if the patient has any questions about the prescription or the medication. The written information provided to the patient shall be in bold print, easily read, and shall include the hours a pharmacist is available and a telephone number where a pharmacist may be reached. The telephone service shall be available at no cost to the pharmacy's primary patient population;

7. All pharmacies that are to engage in central prescription handling maintain a common policies and procedures manual which designates who shall be responsible for each of the component functions of handling the prescription and for ensuring compliance with the Board rules set forth in this chapter. The policies and procedures manual shall also include maintenance of the audit trail required by (d)3 above. The policies and procedures manual shall be made available to the Board upon request;

8. All pharmacies that are to engage in central prescription handling share a common electronic file; and

9. All pharmacies that are to engage in central prescription handling are responsible for ensuring that the prescription has been properly filled.

(e) A prescription for a controlled substance may be filled or refilled by pharmacies engaging in central prescription handling when permitted by law, consistent with Federal requirements set forth at 21 C.F.R. §§ 1300 et seq.

New Rule, R.2004 d.380, effective October 4, 2004.
See: 36 N.J.R. 11(a), 36 N.J.R. 4480(a).
Recodified from N.J.A.C. 13:39-5.10 by R.2005 d.25, effective January 18, 2005.
See: 36 N.J.R. 3345(a), 37 N.J.R. 295(a).
Former N.J.A.C. 13:39-4.18, Responsibilities of pharmacists and permit holders, recodified to N.J.A.C. 13:39-4.17.
Amended by R.2007 d.351, effective November 19, 2007.
See: 38 N.J.R. 4630(a), 39 N.J.R. 4935(a).
In (d)2, inserted the second sentence.

13:39-4.19 Out-of-State pharmacy registration

(a) Any pharmacy located in a state other than New Jersey (hereinafter "out-of-State pharmacy") that ships, mails, distributes or delivers in any manner, legend drugs or devices or controlled dangerous substances pursuant to a prescription into the State, or which participates in a central prescription handling arrangement pursuant to N.J.A.C. 13:39-4.18, shall be registered with the Board pursuant to this section.

(b) It shall be unlawful for any out-of-State pharmacy not registered with the Board pursuant to this section to ship, mail, distribute or deliver in any manner, legend drugs or devices or controlled dangerous substances pursuant to a prescription into the State of New Jersey. Such conduct shall be deemed a violation of N.J.S.A. 45:14-73 and this section.

(c) An out-of-State pharmacy seeking to register with the Board shall submit a completed application for registration to the Board which shall include the following:

1. The name under which the pharmacy is to be operated, the type of practice in which the pharmacy will be engaging, the weekly hours of operation for the pharmacy, and a copy of the prescription label to be used by the pharmacy;

2. The location, names and titles of all principal corporate officers, if the applicant is a corporation, or the location, names and titles of any individuals in whom ownership is or will be vested, if the applicant is not a corporation;

3. The name of the pharmacist-in-charge and his or her license number in the state in which the pharmacy is located, and his or her weekly hours of employment;

4. A dated copy of the most recent inspection report resulting from an inspection of the out-of-State pharmacy conducted by the regulatory or licensing agency in the state in which the pharmacy is located;

5. A letter of good standing from the state licensing authority in the state in which the out-of-State pharmacy is licensed, permitted or registered; and

6. The application fee specified in N.J.A.C. 13:39-1.3(a)4.

(d) An out-of-State pharmacy registered with the Board shall maintain, at all times, a valid unexpired license, permit, or registration to conduct the pharmacy in compliance with the laws and regulations of the state in which it is located. The pharmacy shall notify the Board immediately upon the permanent closing of the pharmacy or upon the commencement of any action by the licensing authority in the state in which it is located concerning its license, permit or registration to conduct the pharmacy. Suspension or revocation of a pharmacy's license, permit or registration in the state in which it is located shall result in the immediate commencement of proceedings by the Board to suspend or revoke the out-of-State pharmacy's registration in New Jersey.

(e) An out-of-State pharmacy registered with the Board shall submit on an annual basis, prior to the expiration of the registration, a renewal application which shall contain the information set forth in (c)1 through 5 above, and the renewal fee set forth in N.J.A.C. 13:39-1.3(a)4. A registered out-of-State pharmacy that fails to submit the renewal application within 30 days after the registration expiration shall submit the late renewal fee set forth in N.J.A.C. 13:39-1.3(a)4 in addition to the renewal fee. An out-of-State pharmacy that continues to ship, mail, distribute or deliver legend drugs or devices or controlled dangerous substances into the State, or continues to participate in a central prescription handling arrangement pursuant to N.J.A.C. 13:39-4.18, with an expired registration shall be deemed to be engaging in the un-

authorized practice of pharmacy and shall be subject to the penalties set forth in N.J.S.A. 45:1-25 et seq.

(f) An out-of-State pharmacy registered with the Board shall submit the information set forth in (c)1 though 5 above and the fee set forth in N.J.A.C. 13:39-1.3(a)4, if applicable, within 30 days of the following:

1. Any change in ownership of the individual equity holder(s) or business entity holding the license, permit or registration to operate the pharmacy;

2. A change of registered agents or officers or a change of stock ownership involving 10 percent or more of the outstanding stock of a publicly traded corporation holding the license, permit or registration to operate the pharmacy;

3. A change in the location of the licensed, permitted or registered pharmacy;

4. A change in the name of the licensed, permitted or registered pharmacy; or

5. A change in the registered pharmacist-in-charge.

(g) An out-of-State pharmacy may obtain a replacement registration upon payment of the fee specified in N.J.A.C. 13:39-1.3(a)4 and upon submission of an affidavit describing the loss or destruction of the registration originally issued, or upon return of the damaged permit.

(h) An out-of-State pharmacy registered with the Board shall:

1. Inform the Board, upon request, of the results of any inspections or investigations conducted by the regulatory or licensing agency of the state in which the pharmacy is licensed, permitted or registered or by a Federal agency, including the filing of any action against the pharmacy by the agency;

2. Inform the Board, upon request, of any directions to, and requests for information from, the pharmacy issued by the regulatory or licensing agency of the state in which the pharmacy is licensed, permitted or registered or by a Federal agency; and

3. Comply with directions concerning compliance with this section and any requests for information issued by the Board.

(i) An out-of-State pharmacy registered with the Board shall maintain its record of prescriptions for patients in the State of New Jersey for a period of not less than five years. The oldest four years of record information shall be maintained in such a manner so as to be retrievable and readable within two weeks. The most recent one year of record information shall be retrievable and readable within one business day.

(j) An out-of-State pharmacy registered with the Board shall, during its regular hours of operation, but not less than five days per week, and for a minimum of 40 hours per week, provide a toll-free telephone service to facilitate communication between patients in the State of New Jersey and a pharmacist who has access to the patients' records. This toll-free number shall be disclosed on a label affixed to each container of drugs dispensed to patients in the State of New Jersey or the out-of-State pharmacy shall meet the requirements set forth in N.J.A.C. 13:39-4.18(d)6.

(k) The Board may forward a complaint against any out-of-State pharmacy registered with the Board for alleged violations of any New Jersey or Federal law or regulation, or any information concerning alleged violations of New Jersey or Federal law by the pharmacy, to the regulatory or licensing agency in the state in which the pharmacy is located, or the Board may institute disciplinary proceedings in New Jersey pursuant to N.J.S.A. 45:1-21 et seq., to resolve the complaint or alleged violation.

New Rule, R.2007 d.351, effective November 19, 2007.
See: 38 N.J.R. 4630(a), 39 N.J.R. 4935(a).

SUBCHAPTER 5. RETAIL FACILITY REQUIREMENTS

13:39-5.1 Purpose and scope

The rules in this subchapter shall apply to all retail pharmacies and retail pharmacy departments in the State. For purposes of this subchapter, "pharmacy" means a retail pharmacy or a retail pharmacy department.

New Rule, R.2005 d.25, effective January 18, 2005.
See: 36 N.J.R. 3345(a), 37 N.J.R. 295(a).
Former N.J.A.C. 13:39-5.1, Imprinted prescription blanks, repealed.

13:39-5.2 Pharmacy access and egress

Pharmacies shall maintain entrances which are easily and safely accessible to the general public. Access to and egress from the pharmacy shall not be such that the public must traverse or traffic through any area in which prescriptions are prepared.

Recodified from N.J.A.C. 13:39-7.1 and amended by R.2005 d.25, effective January 18, 2005.
See: 36 N.J.R. 3345(a), 37 N.J.R. 295(a).
Rewrote the section. Former N.J.A.C. 13:39-5.2, Lack of directions on original prescription, recodified to N.J.A.C. 13:39-7.2.

13:39-5.3 Pharmacy signs

(a) Pharmacies shall post a sign on the exterior of the building or a sign which is otherwise visible from a public roadway, conspicuously identifying the existence of a pharmacy on the premises, unless prohibited by lease agreement or municipal ordinance. In such case, a copy of the lease or ordinance shall be furnished to the Board.

Amended by R.1994 d.351, effective July 18, 1994.
See: 26 N.J.R. 1596(a), 26 N.J.R. 2905(b).
Amended by R.1998 d.167, effective April 6, 1998.
See: 29 N.J.R. 4740(b), 30 N.J.R. 1298(a).
In (d), deleted language regarding practical experience hours in an approved training pharmacy.
Amended by R.1999 d.214, effective July 19, 1999.
See: 31 N.J.R. 1151(a), 31 N.J.R. 1932(a).
In (a)2, substituted a reference to certified preceptors for a reference to preceptors; in (c), substituted "20 hours" for "35 hours" following "less than"; and in (d), inserted a reference to faculty preceptors in the first sentence, and substituted "40 hours" for "45 hours" in the last sentence.
Amended by R.2005 d.25, effective January 18, 2005.
See: 36 N.J.R. 3345(a), 37 N.J.R. 295(a).
In (b); inserted "American Council of Pharmaceutical Education" preceding "accredited college of pharmacy" and deleted the last sentence; in (c), inserted "American Council of Pharmaceutical Education" preceding "accredited college of pharmacy"; in (d), deleted "Board-approved" preceding "college of pharmacy externship program" and deleted "Board-accredited" following "Not more than 45 hours of"; in (f), deleted the last sentence.

13:39–8.5 Change in intern status

(a) A pharmacy intern applying for registration as a pharmacist in the State of New Jersey shall notify the Board within 10 days of any change in:

1. Beginning of a term of internship;
2. Termination of an internship;
3. Number of hours of employment;
4. Scheduled hours of employment;
5. Certified preceptor; and/or
6. Employing pharmacy.

Amended by R.1994 d.351, effective July 18, 1994.
See: 26 N.J.R. 1596(a), 26 N.J.R. 2905(b).
Amended by R.1999 d.214, effective July 19, 1999.
See: 31 N.J.R. 1151(a), 31 N.J.R. 1932(a).
In (a)5, substituted a reference to certified preceptors for a reference to preceptors.

13:39–8.6 (Reserved)

Amended by R.1999 d.214, effective July 19, 1999.
See: 31 N.J.R. 1151(a), 31 N.J.R. 1932(a).
In (a)4, substituted a reference to certified preceptors for a reference to preceptors; deleted a former (c); and recodified former (d) as (c), and deleted a former first sentence.
Repealed by R.2005 d.25, effective January 18, 2005.
See: 36 N.J.R. 3345(a), 37 N.J.R. 295(a).
Section was "Committee on Pharmacy Internship and Externship".

13:39–8.7 (Reserved)

Amended by R.1994 d.351, effective July 18, 1994.
See: 26 N.J.R. 1596(a), 26 N.J.R. 2905(b).
Amended by R.1998 d.167, effective April 6, 1998.
See: 29 N.J.R. 4740(b), 30 N.J.R. 1298(a).
In (a)2iii, substituted "Three examples" for "One example".
Amended by R.1999 d.214, effective July 19, 1999.
See: 31 N.J.R. 1151(a), 31 N.J.R. 1932(a).
In (a)2, inserted a reference to medication orders in i, substituted a reference to certified preceptors for a reference to preceptors in iv, and added v through ix.
Repealed by R.2005 d.25, effective January 18, 2005.
See: 36 N.J.R. 3345(a), 37 N.J.R. 295(a).
Section was "Pharmacist intern log".

SUBCHAPTER 9. PHARMACEUTICAL SERVICES FOR HEALTH CARE FACILITIES

13:39–9.1 Purpose and scope

(a) The rules in this subchapter shall apply to all retail pharmacies which contract to provide pharmaceutical services for healthcare facilities and to all institutional pharmacies which provide pharmaceutical services for their own health care system.

(b) An institutional pharmacy filling prescriptions for outpatient use shall comply with all retail pharmacy requirements of this chapter.

New Rule, R.2005 d.25, effective January 18, 2005.
See: 36 N.J.R. 3345(a), 37 N.J.R. 295(a).
Former N.J.A.C. 13:39–9.1, Definitions, recodified to N.J.A.C. 13:39–9.2.

13:39–9.2 Definitions

The following words and terms, as used in this subchapter, shall have the following meanings, unless the context clearly indicates otherwise.

"Drug administration" means a procedure in which a prescribed drug is given to a patient by an authorized person in accordance with all laws and rules governing such procedures.

"Formulary" means a continually revised compilation of pharmaceuticals available in the pharmacy for use in the facility developed by the Pharmacy and Therapeutics Committee.

"Health care facility" means a facility or institution licensed by the Department of Health and Senior Services pursuant to N.J.S.A. 26:2H–1 et seq.

"Health care system" means one or more health care facilities which are owned or controlled by the same legal entity.

"Institutional pharmacy" means the area in a health care facility or a health care system licensed by the Board as a pharmacy that maintains an institutional permit. "Institutional pharmacy" includes any areas of the health care facility or the health care system where pharmaceuticals are stored, compounded or dispensed.

"Medication order" means a written request for medication originated by an authorized prescriber and intended for patient use in the health care facility, and not for use of the institution's employees or their dependents or outpatients of the facility's clinics. A valid medication order contains the date ordered, the patient's name and location within the facility, the name, dose, route, and frequency of administration of the medication, and any additional instructions. Computer-generated medication orders within an institutional setting, utilizing the prescriber's electronic signature or password will meet legal requirements for a prescriber's original handwritten signature on medication orders. Computerized signatures or passwords will be accepted provided that the facility has adequate safeguards which assure the confidentiality of each electronic signature or password and which prohibit their improper or unauthorized use.

"Pharmacy and Therapeutics Committee" means the active standing committee of the institution or health care facility which is the organizational line of communication and liaison between the medical service and pharmacists and which acts to review and promote rational drug therapy and utilization in the facility.

"Unit dose drug distribution system" means a system of dispensing drugs to be administered to patients of the facility whereby the medications are delivered daily (or more frequently) by the pharmacy to the patient care units in amounts equal to a 24–hour supply or less and are prepared, whenever possible, in single unit use packaging.

"Unit use packaging" means a single unit use medication provided in sealed packaging which contains the following information for each unit in the package:

1. Product name;

2. Strength and/or quantity and/or volume, where appropriate;

3. Lot number;

4. Use by date;

5. Manufacturer or repackager; and

6. If there is more than one product in the single unit, a physical description of each medication in the single unit.

Amended by R.1994 d.351, effective July 18, 1994.
See: 26 N.J.R. 1596(a), 26 N.J.R. 2905(b).
Amended by R.1998 d.167, effective April 6, 1998.
See: 29 N.J.R. 4740(b), 30 N.J.R. 1298(a).
Amended N.J.A.C. reference in "Pharmacy and Therapeutics Committee" definition.
Amended by R.1999 d.214, effective July 19, 1999.
See: 31 N.J.R. 1151(a), 31 N.J.R. 1932(a).
Deleted "Authorized prescriber" and "Direct supervision"; in "Health care facility", substituted a reference to patients and residents for a reference to the sick and injured; in "Unit dose drug distribution system", deleted a reference to biologicals; and added "Unit use packaging".
Amended by R.2000 d.457, effective November 20, 2000.
See: 31 N.J.R. 3044(a), 32 N.J.R. 4123(a).
Inserted "Health care system"; in "Institutional pharmacy", inserted references to health care systems throughout, substituted "any" for "other" following "includes", and deleted "manufactured," following "stored,".
Recodified from N.J.A.C. 13:39–9.1 and amended by R.2005 d.25, effective January 18, 2005.
See: 36 N.J.R. 3345(a), 37 N.J.R. 295(a).
Rewrote "Health care facility"; in "Pharmacy and Therapeutics Committee", substituted "medical service and pharmacists" for "medical and pharmacy staff" and deleted the last sentence; rewrote "Unit use packaging". Former N.J.A.C. 13:39–9.2, Licensure of institutional pharmacies, recodified to N.J.A.C. 13:39–9.3.

13:39–9.3 Licensure of institutional pharmacies

(a) Any institutional pharmacy as defined under N.J.A.C. 13:39–9.2 shall be registered with and possess an institutional permit issued by the Board. The permit shall be conspicuously displayed in the facility's pharmacy. The institutional pharmacy is subject to and shall be conducted in accordance with all existing State and Federal rules and regulations.