



NON-RESIDENT PHARMACY PERMIT APPLICATION

This permit authorizes facilities outside of this state engaged in the business of filling mail order prescriptions to engage in the sale, distribution, or dispensing of legend drugs or devices in this State.

The pharmacist-in-charge for the applicant must attend an Application Review Committee meeting at the Board's office. Applicant will be notified by email of the date and time of the meeting for which its application review hearing is scheduled. All requested information and an emailed confirmation are required prior to the meeting date. Using false, fraudulent, forged statement or document, or committing a fraudulent, deceitful or dishonest act or omitting a material fact in obtaining licensure is grounds for discipline or licensure denial. A South Carolina Non-Resident Pharmacy Application has a one-year expiration.

Failure to complete all required fields and/or provide necessary supplemental documentation will delay the application process. **If an item is not applicable, please indicate N/A. In order to avoid delay, please do not provide the items below in a binder, folder or use dividers. Also, provide items in the order as listed below. Please write legibly.**

Include with your application:

- Check or money order in the amount of **\$420** made payable to SC Board of Pharmacy. (Application fee is non-refundable. A returned check fee of up to \$30, or an amount specified by law, may be assessed on all returned funds.)
- Copy of resident state pharmacy permit
- Copy of recent operational inspection report. Inspection must have been conducted within the last 2 years. You must also supplement your application by providing the Board's staff with an inspection report received between the date of the application and the date of the application review hearing.
- Copy of current DEA registration
- Copy of state controlled substance registration
- Copy of policy and procedure for shipping refrigerated products
- Copy of dispensed label
- Letter describing, in detail, the nature of your business
- Provide a list of all pharmacy permits/licenses held in other states
- Photographs
 - Exterior of pharmacy building to include identifiable parts of adjacent buildings
 - Front end of pharmacy to include consulting area and drop-off/pick-up
 - Compounding area
 - Work area
- Accreditation certificates
- Non-sterile Compounding requirements from list on page 8 of application
- Sterile Compounding requirements from list on page 9 of application
- Compounding – requirements even if you do not plan to ship sterile/non-sterile Compounds into SC, you must still submit all compounding information.
- Include organizational chart. If a change of ownership, include charts of before and after the change. The chart must show the legal business entities from the ultimate parent company down to and including the applicant and must include the legal business name, trade name, and type of ownership for each entity on the chart. Chart must include owner's name with a ten percent or greater ownership interest in a non-publicly traded company.



South Carolina Department of Labor, Licensing and Regulation
South Carolina Board of Pharmacy
 110 Centerview Dr. • Columbia • SC • 29210
 P.O. Box 11927 • Columbia • SC 29211-1927
 Phone: 803-896-4700 • Contact.pharmacy@llr.sc.gov • Fax: 803-896-4596
 llr.sc.gov/bop

NON-RESIDENT PHARMACY PERMIT APPLICATION

FOR BOARD USE ONLY	
Date Paid	
Amount Paid	
Check No.	

- New Permit
 Change to Existing Permit (Permit No.: _____)
 Change of Ownership **(include organizational chart before and after change)**
 Change of Name
 Change of Location

Federal Employee Identification No. (FEIN): _____

NABP e-Profile ID No.: _____

Legal Name of Pharmacy: _____

dba Name: _____

Street address of physical location of pharmacy: _____

City: _____ State: _____ Zip: _____

Toll-free number for patients: _____

Business Phone: _____ Resident State Permit/License No.: _____

Is application based on a change in ownership? Yes No

If Yes: _____
Previous Owner/Name of Pharmacy SC Permit/License Number

Mailing Address where all correspondence regarding licensure will be sent if other than facility physical address above:

Contact Person: _____ Email: _____

Facility Name: _____

Address: _____ City: _____ State: ___ Zip: _____

EMPLOYEE INFORMATION

Pharmacist-In-Charge: _____ License No.: _____

Email: _____ State of Issuance: _____

Pharmacist Full-Time (Use separate sheet if necessary)	License No.	State Issuance

Pharmacist Part-Time (Use separate sheet if necessary)	License No.	State Issuance

Pharmacy Technicians (Use separate sheet if necessary)	License No.	State Issuance

What is the daily working ratio of pharmacists to pharmacy technicians? _____

PHARMACY

Pharmacy website address: _____

Hours of Operation: _____ Hours a pharmacist is available: _____

Do you fill prescriptions via the internet? Yes No

If Yes, is there a policy and procedure to validate patient-practitioner relationship? Yes No

Provide all website addresses used: _____

When was your last Board of Pharmacy operational inspection? _____
(Attach a copy of the inspection report)

Date your pharmacy began dispensing to South Carolina patients: _____

Approximate number of South Carolina patients served annually: _____

Has the **applicant or any entity** under common control with applicant ever applied for a pharmacy permit in South Carolina? Yes No

If Yes, state business name on application: _____

Month and year submitted: _____ Permit No. Issued: _____

CONTROLLED SUBSTANCES

Non-resident pharmacies permitted by the SC Board of Pharmacy that dispense controlled substances are required to obtain a South Carolina Controlled Substances Registration from the SCDHEC-Bureau of Drug Control. Access the application via the website at www.dhec.sc.gov/Health/FHPF/DrugControlRegisterVerify/NewRegistrations/. All SC-permitted non-resident pharmacies must register with the SCDHEC-Bureau of Drug Control Prescription Monitoring Program. If you do not dispense controlled substances, you must request an exemption from reporting requirements. Contact PMP at scripts@dhec.sc.gov or (803) 896-0688.

Does your pharmacy dispense controlled substances? Yes No

COMPOUNDING

Does your pharmacy do compounding? Yes No (If No, skip compounding questions.)

Sterile compounding? Yes No

Non-sterile compounding? Yes No

COMPOUNDING QUESTIONS Additional documentation required (See attached lists)

Do you have PCAB accreditation? Yes No

If Yes, expiration date of accreditation: _____

Is the pharmacist-in-charge trained and knowledgeable in the preparation, labeling and dispensing of compounded drugs? Yes No

If No, the pharmacist-in-charge designates the following pharmacist employee as knowledgeable in the preparation, labeling and dispensing of compounded drug preparations:

Pharmacist Name: _____ License No.: _____

Do you use an independent lab or qualified individual for end product testing? Yes No

If Yes, provide a name and address: _____

Do you use an independent lab or qualified individual for routine environmental testing of hood, ante room and clean room? Yes No

If Yes, provide a name and address: _____

Do you provide patient specific compounded products? Yes No

Do you provide non-patient specific compounded products? Yes No

OWNERSHIP (Check appropriate box and provide complete information)

Do any of the owners of this facility own any portion of, control, or have any beneficial interest in any other pharmacy/wholesaler/3PL (whether permitted in SC or not) other than a publicly-traded corporation? Yes No

If Yes, provide a list with names and addresses.

Sole Proprietorship Name of Business Entity: _____

Name	City, State	Birth Year

General Partnership **LLP** Name of Partnership: _____

Partner Name	City, State	Birth Year	% of Ownership

Corporation LLC Name of Corporation/LLC: _____

Name of Parent Company: _____ State of Incorporation: _____

(Only include individuals with 10% or greater ownership.)

Name of Individual Owners and Principal Officers	Title	City, State	Birth Year	% of Ownership
1.				
2.				
3.				

DISCIPLINARY HISTORY

If you answer “Yes” to any part of this section, provide a detailed explanation on a separate sheet and attach copies of applicable court documentation. Include the city and state where the offense(s) occurred.

HAS THE APPLICANT: to include the entity, the undersigned pharmacist-in-charge, undersigned permit holder, any person or entity identified in the ownership management section above, or any entity under common control with the applicant **EVER:**

1. Had a permit or professional license disciplined, denied, refused or revoked for violation of any federal or state pharmacy laws or drug laws? Yes No
If No, is there any pending disciplinary action? Yes No

2. Been convicted, fined or entered in a plea of guilty or nolo contendere in any criminal prosecution, felony or misdemeanor in South Carolina or any other state, or in a United States court for:
 - a. any offense relating to drugs, narcotics, controlled substances or alcohol, whether or not a sentence was imposed? Yes No
 - b. any offense involving the practice of pharmacy, or relating to acts committed within a pharmacy or drug distributor setting or incident to pharmacy practice, whether or not a sentence was imposed? Yes No
 - c. any offense involving fraud, dishonesty or moral turpitude whether or not a sentence was imposed? Yes No

3. Had an application for a pharmacy; pharmacist license, permit or certificate denied, refused, or not issued in South Carolina or any other state or country? Yes No
If Yes, please explain: _____

4. Operated, or allowed the facility to operate without a valid permit? Yes No

5. Violated the drug laws, rules, statutes and/or regulations of South Carolina or any other state or country? Yes No

ATTESTATION

I declare that I have read and approve the foregoing and the statements are true and correct to the best of my knowledge and belief; I will comply with the South Carolina Pharmacy Practice Act and I understand I am responsible for any violation(s) occurring during my tenure.

Pharmacist-in-Charge Signature

Print Name of Pharmacist-in-Charge

Email Address of Pharmacist-in-Charge

Date

I declare that the foregoing statements are true and correct to the best of my knowledge and belief; the permit applied for is to cover only the pharmacy indicated above and at the location specified; and that I will comply with the South Carolina Pharmacy Practice Act.

Permit Holder Signature

Print Name

Email Address of Permit Holder

Title

Date

Information from this application may be shared.

Mail completed application to:

South Carolina Board of Pharmacy
110 Centerview Dr.
Columbia SC 29210

Compounded medications cannot be re-sold, therefore the compounded medications must be sent and billed to the patient or sent to a physician/facility to be administered on site.

SECTION 40-43-86 (CC)(2) *Pharmacists may not offer compounded medications to other pharmacies for resale; however, pharmacists may compound products based on an order from a practitioner for use by practitioners for patient use in institutional or office settings. Compounding pharmacies/pharmacists may advertise or otherwise promote the fact that they provide prescription compounding services, e.g., chemicals, devices, and information, when requested; however, they may not solicit business by promoting to compound specific drug products, e.g., like a manufacturer;*

The pharmacy must keep all records and a pharmacist must be present during all compounding functions.

SECTION 40-43-89 (N)(1) *A facility located outside this State, whose primary business is mail order prescription service, shall have a permit issued by the board to ship, mail, or deliver a controlled substance or dangerous drug or device into this State pursuant to a prescription of a licensed practitioner. The facility shall report to the board:*

- (b) that it complies with the applicable laws for operation in the state in which it is located and with the provisions of this section. The facility shall have a valid unexpired license, permit, or registration in compliance with the laws of the state in which it is located and must be constantly under the personal and immediate supervision of a licensed pharmacist. The facility shall submit to the board with its initial application and with each renewal application a copy of its most recent inspection report resulting from an inspection conducted by the regulatory or licensing agency of the state in which it is located. These inspections are deemed to meet all inspection requirements contained in this chapter;*
- (c) that it maintains its records of controlled substances or dangerous drugs or devices dispensed to patients in this State so that the records are readily retrievable.*

If the state in which the pharmacy is located does not have a pharmacist to auxiliary personnel ratio, a pharmacist is not allowed to supervise more than two pharmacy technicians at one time.

SECTION 40-43-89 (N)(3) *If the state in which the facility is located does not establish, by statute or regulation, a ratio describing the number of auxiliary personnel that a pharmacist may supervise, or otherwise define the role of the pharmacist in the compounding and dispensing of prescription drugs, then that facility may not allow a pharmacist to supervise more than two pharmacy technicians at any time in the compounding and dispensing of prescription drugs.*

The pharmacy label must contain a toll-free number and have a pharmacist available during its regular house of operation, but not less than 6 days or 40 hours a week.

SECTION 40-43-89 (N)(4) *A pharmacy, as described in this section, during its regular hours of operation but not less than six days or forty hours a week, shall provide a toll-free telephone service to facilitate communication between patients in this State and a pharmacist at the pharmacy who has access to their records. This telephone number must be printed on a label affixed to the container for the substance, drug, or device.*

NON-RESIDENT PHARMACY
NON-STERILE COMPOUNDING REQUIREMENTS

- A.** Continuing Education: Documentation of CE in the science and art of compounding for pharmacists and technicians involved in compounding. Six (6) hours initially and four (4) hours annually. Does not have to be ACPE-approved.
- B.** Diagram and photographs of compounding area.
- C.** Refrigerator temperature log: Copy of one page of the most current month to include time, date, temperature, initials.
- D.** Room temperature and humidity log: Copy of one page of the most current month to include time, date, temperature, humidity, and initials.
- E.** Cleaning logs: Copy of one month of logs to include, at a minimum,
 - a. Daily cleaning log – countertops, hoods, equipment, utensils, floors swept, trash discarded
 - b. Weekly cleaning log – floors mopped
 - c. Monthly cleaning log – shelves, refrigerator/freezer, cabinet exteriors (all sanitized)
- F.** Documentation that equipment is routinely inspected, calibrated and cleaned.
- G.** Copies of completed logs/completed product formula worksheets for top 5 compounded products with a copy of the actual prescription. Also provide a reprint/duplicate of the final compounded product label. Include assigned BUD and reasoning for BUD assigned.
- H.** Copies of procedures (choose any 3) done within the last 6 months to monitor the output of compounded prescriptions such as potency, capsule size and weight.
- I.** A printed batch (stock) label, if applicable.
- J.** Standard operating policies and procedures for:
 - a. General compounding procedures
 - b. Maintenance and cleaning of area and equipment

* Do NOT send entire SOP library.

NON-RESIDENT PHARMACY
STERILE COMPOUNDING REQUIREMENTS

- A.** Documentation of training and/or continuing education in the science and art of compounding of sterile products for all pharmacists and technicians involved in compounding.
- B.** Diagram and photographs of Sterile Compounding Area.
- C.** Refrigerator temperature log: Copy of one page of the most current month to include time, date, temperature, initials.
- D.** One page of Logs monitoring:
 - pressure differential
 - room temperature/humidity in compounding area
- E.** Logs for one full month to include:
 - cleaning of all areas used in sterile compounding process
- F.** Copy of last inspection, by qualified individual, of hoods, buffer, clean and ante areas including ISO classification, particle counts and microbiology.
- G.** Copies of completed logs/completed product formula worksheets for top 5 sterile compounded products with a copy of the actual prescription. Also provide a reprint/duplicate of the final compounded product label. Include assigned BUD and reasoning for BUD assigned.
- H.** Reprint/duplicate of final product label
 - minibag
 - large volume
 - TPN
 - syringe
 - vial
- I.** Compounding Policies and Procedures, specific to your facility, as applicable for the following:
 - (1) quality control
 - (2) sterile compounding technique
 - (3) cleaning/maintenance of compounding area and equipment

* Do not send the entire SOP library.