DIVISION OF MEDICAL QUALITY ASSURANCE BOARD OF PHARMACY 4052 BALD CYPRESS WAY, BIN #C-04 TALLAHASSEE, FLORIDA 32399-3254 (850) 245-4474



INSTITUTIONAL PHARMACY PERMIT APPLICATION AND INFORMATION

January 2018



Dear Florida Pharmacy Permit Applicant,

Thank you for applying for a pharmacy permit in the State of Florida. The information in this packet has been designed to provide the essential information required to process your application in a timely manner. Your assistance in providing all required information will enable the Florida Board of Pharmacy (the Board) staff to process your application as soon as possible. You are encouraged to apply as early as possible, to avoid delays due to a large volume of applicants.

Florida Statutes require a completed application and fees before your application can be reviewed. Please read these instructions carefully and fully before submitting the application. You should keep a copy of the completed application and all other materials sent to the board office for your records. When you mail the completed application and fees, use the address noted in the instructions and on the application form.

When your application arrives, your fees will be deposited and verified before the staff review can begin. You will receive a letter acknowledging receipt of your application. The staff will notify you within 30 days if any materials are incomplete.

If you need to communicate with the board staff, you are encouraged to email the board staff at <u>info@floridaspharmacy.gov</u> or you may call us at (850) 245-4474. Phone calls are returned within 24 hours and emails are responded to within 48 hours during normal business hours. Our staff is committed to providing prompt and reliable information to our customers. Many procedures have been streamlined to expedite the processing of applications; we certainly welcome your comments on how our services may be improved.

Sincerely,

The Board of Pharmacy

INSTITUTIONAL PHARMACY PERMIT APPLICATION INFORMATION

Whether opening a new establishment, changing locations, or changing owners, a pharmacy permit is required prior to operating in the State of Florida. The permit application must be completed and returned to the Florida Board of Pharmacy with the required fee of \$255.00. The application MUST have the original signatures of the owner or officer of the establishment and the Consultant Pharmacist of Record (COR). If compounding sterile preparations, submit an additional application on Form DH-MQA 1270, "Special Sterile Compounding Permit" and pay the additional permitting fee.

There are three types of Institutional Pharmacy Permit applicants. Please read the description below. Check which permit type you are applying for on the application.

1. <u>Institutional Class I Pharmacy</u> – An Institutional Class I pharmacy is an institutional pharmacy in which all medicinal drugs are administered from individual prescription containers to the individual patient and in which medicinal drugs are not dispensed on the premises. No medicinal drugs may be dispensed in a Class I Institutional pharmacy. A Special- Closed System Pharmacy Permit, Special Parenteral and Enteral Pharmacy Permit, or Community Pharmacy Permit allow dispensing of individual patient prescriptions.

2. <u>Institutional Class II Pharmacy Permit</u> – An Institutional Class II pharmacy is an institutional pharmacy that employs the services of a registered pharmacist or pharmacists who, in practicing institutional pharmacy, provide dispensing and consulting services on the premises to patients of that institution, for use on the premises of that institution. An Institutional Class II pharmacy is required to be open sufficient hours to meet the needs of the hospital facility.

3. <u>Modified Institutional Class II Pharmacy Permits</u> - Modified Institutional Class II pharmacies are those institutional pharmacies in short-term, primary care treatment centers that meet all the requirements for a Class II permit, except space and equipment requirements. Modified Class II Institutional pharmacies are designated as Type "A", Type "B" and Type "C" according to the type of specialized pharmaceutical delivery system utilized. Please review Rule 64B16-28.702, Florida Administrative Code for specific requirements.

Section 465.022(4), Florida Statutes, also provides that an application for a pharmacy permit must include the applicant's written policies and procedures for preventing controlled substance dispensing based on fraudulent representations or invalid practitioner-patient relationships. The policy and procedure manual shall contain the procedures implemented to minimize the dispensing of controlled substances based on fraudulent representations as follows:

- 1. Provisions to identify and guard against invalid practitioner-patient relationships.
- 2. Provisions to guard against filling fraudulent prescriptions for controlled substances.
- 3. Provisions to identify prescriptions that are communicated or transmitted legally.
- 4. Provisions to identify the characteristics of a forged or altered prescription.

Application Processing

Please read all application instructions before completing your application.

1) Mail Application.

Please mail the application and the \$255.00 application fee (cashier's check or money order made payable to the FLORIDA DEPARTMENT OF HEALTH) to the following address:

Application & Fees:

Department of Health Board of Pharmacy P.O. Box 6320 Tallahassee, Florida 32314-6320

Express Mail ONLY

Department of Health Board of Pharmacy 4052 Bald Cypress Way, Bin C-04 Tallahassee, FL 32399-3254

Within 30 days of receipt of your application and fees, the Board office will notify you of the receipt of your application, any required documents, and your status. If the application is complete, you will be notified that an inspector will contact you to setup an inspection appointment. If your application is incomplete, you will be notified in writing of what is required to make your application complete.

2) Submit fingerprint results

Failure to submit fingerprints will delay your application. All owners, officers, and Consultant Pharmacists of Record (CORs) are required to submit a set of fingerprints unless the corporation is exempt under Section 465.022, Florida Statutes, for corporations having more than \$100 million of business taxable assets in this state. These corporations are only required to have the COR to submit fingerprints.

Electronic fingerprint information ("EFI") that has been submitted to the Florida Agency for Health Care Administration may be accessible by the Florida Department of Health for a period of sixty (60) months. If the Department is able to access EFI from AHCA, applicants will not be required to resubmit EFI for additional or new applications submitted during this time period. After sixty (60) months, new electronic fingerprint information must be submitted as part of all applications. <u>Note: If your officer, owner, or Consultant</u> <u>Pharmacist of Record has already been fingerprinted at the time you are completing this Institutional Pharmacy permit application, please ensure to provide the Transaction Control Number (TCN), if known, with the requested information in the application.</u>

Applicants may use any Livescan vendor that has been approved by the Florida Department of Law Enforcement to submit their fingerprints to the department. Please ensure that the Originating Agency Identification (ORI) number is provided to the vendor when you submit your fingerprints. If you do not provide an ORI number or if you provide an incorrect ORI number to the vendor, the Board of Pharmacy <u>will not</u> receive your fingerprint results. The applicant is fully responsible for selecting the vendor and ensuring submission of the prints to the Department.

How do I find a Livescan vendor in order to submit my fingerprints to the Department? The Department of Health accepts electronic fingerprinting service offered by Livescan device vendors that are approved by the Florida Department of Law Enforcement and listed at their site. You can view the vendor options and contact information at:

http://www.floridahealth.gov/licensing-and-regulation/background-screening/livescanservice-providers.html What information must I provide to the Livescan vendor I choose?

- If you are an applicant seeking a license for any profession regulated by the Department of Health, which requires a criminal background search as a condition of licensure, you must provide accurate demographic information at the time your fingerprints are taken, *including your Social Security number*. The Department will not be able to process a submission that does not include your Social Security number.
- You must provide the correct ORI number.

Where do I get the ORI number to submit to the vendor?

The ORI number for the pharmacy profession is **EDOH4680Z**.

Attestation for Business Taxable Assets

If the applicant has more than \$100 million dollars of business taxable assets in this state, please submit a formal opinion letter from a Certified Public Accountant duly licensed in the state of your principal place of business affirming the corporation has more than \$100 million of business taxable assets in this state for the previous tax year. In lieu of submitting a formal opinion letter from a Certified Public Accountant, the applicant may submit its Florida Corporate Income/Franchise and Emergency Excise Tax Return (Form F-1120, Effective 01/09).

3) Privacy Statement and Attestation

In order for the Board of Pharmacy Office to receive your Livescan electronic fingerprinting results, you must affirm that you have been provided with and read the attached statement from the Florida Department of Law Enforcement regarding the sharing, retention, and right to challenge incorrect criminal history records, and the "Privacy Statement" document from the Federal Bureau of Investigation. The appropriate form(s) to provide this affirmation are included within Items #1 and #2 of the application.

Licensure Process

Once the application is deemed complete, the board staff authorizes an inspection. Upon completion of the inspection, the inspector notifies the board office as to whether the inspection was satisfactory or unsatisfactory. If the inspection is satisfactory, a permit number is issued within 10 days. <u>Please wait 7 - 14 days from your satisfactory inspection before checking on the status of your permit</u>.

You may look up your license number on our website at <u>http://www.flhealthsource.com/</u> under "Verify a License."

Drug Enforcement Administration (DEA)

Please note that the DEA will not issue a registration until the Florida Board of Pharmacy has issued a pharmacy permit. More information is available by visiting the DEA website at http://www.DEAdiversion.usdoj.gov, or by contacting them at (800) 667-9752.

IMPORTANT NOTICE: Pursuant to Section 465.022(5), F.S., the Department or Board shall deny an application for a pharmacy permit if the applicant or an affiliated person, partner, officer, director, or prescription department manager or consultant pharmacist of record of the applicant:

(a) Has obtained a permit by misrepresentation or fraud.

(b) Has attempted to procure, or has procured, a permit for any other person by making, or causing to be made, any false representation.

(c) Has been convicted of, or entered a plea of guilty or nolo contendere to, regardless of adjudication, a crime in any jurisdiction which relates to the practice of, or the ability to practice, the profession of pharmacy.

(d) Has been convicted of, or entered a plea of guilty or nolo contendere to, regardless of adjudication, a crime in any jurisdiction which relates to health care fraud.

(e) Has been convicted of, or entered a plea of guilty or nolo contendere to, regardless of adjudication, a felony under chapter 409, chapter 817, or chapter 893, or a similar felony offense committed in another state or jurisdiction, since July 1, 2009.

(f) Has been convicted of, or entered a plea of guilty or nolo contendere to, regardless of adjudication, a felony under 21 U.S.C. ss. 801-970 or 42 U.S.C. ss. 1395-1396 since July 1, 2009.

(g) Has been terminated for cause from the Florida Medicaid program pursuant to s. 409.913, unless the applicant has been in good standing with the Florida Medicaid program for the most recent 5-year period.

(h) Has been terminated for cause, pursuant to the appeals procedures established by the state, from any other state Medicaid program, unless the applicant has been in good standing with a state Medicaid program for the most recent 5-year period and the termination occurred at least 20 years before the date of the application.

(i) Is currently listed on the United States Department of Health and Human Services Office of Inspector General's List of Excluded Individuals and Entities.

(j) Has dispensed any medicinal drug based upon a communication that purports to be a prescription as defined by s. 465.003(14) or s. 893.02 when the pharmacist knows or has reason to believe that the purported prescription is not based upon a valid practitioner-patient relationship that includes a documented patient evaluation, including history and a physical examination adequate to establish the diagnosis for which any drug is prescribed and any other requirement established by board rule under chapter 458, chapter 459, chapter 461, chapter 463, chapter 464, or chapter 466.

For felonies in which the defendant entered a plea of guilty or nolo contendere in an agreement with the court to enter a pretrial intervention or drug diversion program, the department shall deny the application if upon final resolution of the case the licensee has failed to successfully complete the program.

If applicable to you, please provide the documentation to the Florida Board of Pharmacy.

PHARMACY PERMIT APPLICATION CHECKLIST

Keep a copy of the completed application for your records.

It is recommended that you use the following checklist to help ensure that your application is complete. Failure to attach any required document, or to have required documentation sent to the Board, will result in an incomplete application. <u>Final approval for inspection cannot be granted until the application is complete</u>.

INSTITUTIONAL PHARMACY PERMIT

All A	pplication	Questions	Answered?

- \$255.00 Fee Attached (Permit fee includes \$250 application fee and \$5.00 unlicensed activity fee)
- Articles of Incorporation paperwork from the Secretary of State provided?
- _____COR Designation and Privacy Statement Acknowledgement provided (Application Item #1)?
 - ____Affiliate/Owner Privacy Statement Acknowledgement provided for each affiliate/owner (Application Item #2)?
 - Answers to Policy and Procedure Questions provided for **Institutional Pharmacy** applicants (Application Item #3)?
 - Answers to Policy and Procedure Questions provided for **Modified Class II Institutional Pharmacy** applicants (Application Item #4)?
 - ____Applicant/Affiliate/Owner supplemental documents provided for explaining any previous ownership, disciplinary actions, voluntary relinquishments and/or criminal activity?
 - __Applicant/Affiliate/Owner pharmacy permit questions answered and supplemental documents provided?
 - Policies and Procedures for preventing controlled substance dispensing based on fraudulent representations or invalid practitioner-patient relationships submitted?



FLORIDA BOARD OF PHARMACY P.O. Box 6320 Tallahassee, FL 32314-6320 850-245-4474 http://www.floridaspharmacy.gov

APPLICATION

Application Type – Please choose one of the following:				
New Establishment (\$255.00 fee) Complete: Section A <u>only</u> , along with Items #1 and 2.		Change of Location (\$100.00 fee) Complete: Sections A and B <u>only</u> .		
Change of Ownership (\$255.00 fee) Complete: Sections A and C <u>only</u> , along with Items #1 and 2.			Stock Transfer (no fee) Complete: Section A, pages 2-3 and Section D <u>only</u> .	
Pharmacy Permit Type – Please ch	noose only one of	the following:		
Institutional Class I		Modified Institutional Class II AClass II B Class II C Complete: Item #4 <u>only</u> ,		
Institutional Class II Complete: Item #3 <u>only</u> ,	Modifie			
SECTION A. Please comple	te for all Appl	lication Types		
Please list your Federal Employer	Identification Nu	mber:		
1. Corporate Name			Telephone Number	
2. Doing Business As (d/b/a)			E-Mail Address** (see note below)	
3. Mailing Address				
City	State		Zip	
4. Physical Address			·	
City	State		Zip	
5. Consultant Pharmacist of Reco	ord (COR) Informa	ation		
Name			License Number	
	1			
Email Address ** (see note below)		Telephone Number		
6. Contact Person		Title		
Email Address ** (see note below)		Telephone Number		
			e-mail address released in response to a public stead contact the office by phone or in writing.**	

7. Ownership Information			
a. Type of Ownership:Indi	ividual	CorporationPartnership	
NOTE: If the applicant is a corpor of the Articles of Incorporation on	ation or limi file with the	ted partnership you must include with your application a Florida Secretary of State's office.	<u> copy</u>
b. Are the applicants, officers,	directors,	shareholders, members and partners over the age c	of 18?
Yes No			
	blic Accounta	100 million of business taxable assets in this state ant for previous tax year or Florida Corporate Income /France	
Yes No			
interest of 5% or greater and a operation of the applicant inclu fingerprints and fees unless y only submit fingerprints for th file with DOH or AHCA and ava	iny person uding office ou answere e Consulta ailable to th the % of C	corporation. Each person listed below having ar who, directly or indirectly, manages, oversees, or ers and members of the board of directors must sul ed yes to 7c. If 7c. is "Yes", please list the owners nt Pharmacist of Record. If 7c. is "Yes" and the p ne Board of Pharmacy, the requirement to submit the ownership column does not add to 100%, please sary.	controls the bmit a set of s below and prints are on he prints for
Owner/Officer-Title	Date of Birth	Mailing Address, City, State, Zip Code	% of Ownership
			%
			%
			%
			%
			%
business permit which was dis	ciplined, s	ship interest of 5% or more in a pharmacy or any of uspended, revoked, or closed involuntarily within t t disclosing the reason the entity was closed.	
Yes No	_		
	untarily rel	rship interest of 5% or more in a pharmacy or any c inquished or closed voluntarily within the past 5 yearing the reason the entity was closed.	
Yes No			

Pursuant to Section 465.022(5), the questions $9 - 17$ are being asked. If you answer "Yes" to any of the following questions, explain on <u>a separate sheet</u> providing accurate details and submit copies of supporting documentation.
9. Has the applicant or any principal, officer, agent, managing employee, or affiliated person of the applicant obtained a permit by misrepresentation or fraud?
Yes No
10. Has the applicant or any principal, officer, agent, managing employee, or affiliated person of the applicant attempted to procure, or has procured, a permit for any other person by making, or causing to be made, any false representation?
Yes No
11. Has the applicant or any principal, officer, agent, managing employee, or affiliated person of the applicant been convicted of, or entered a plea of guilty or nolo contendere to, regardless of adjudication, a crime in any jurisdiction which relates to the practice of, or the ability to practice, the profession of pharmacy?
Yes No
12. Has the applicant or any principal, officer, agent, managing employee, or affiliated person of the applicant been convicted of, or entered a plea of guilty or nolo contendere to, regardless of adjudication, a crime in any jurisdiction which relates to health care fraud?
YesNo
13. Has the applicant or any principal, officer, agent, managing employee, or affiliated person of the applicant been convicted of, or entered a plea of guilty or nolo contendere to, regardless of adjudication, a felony under 21 U.S.C. ss. 801-970 or 42 U.S.C. ss. 1395-1396 since July 1, 2009?
Yes No
14. Has the applicant or any principal, officer, agent, managing employee, or affiliated person of the applicant ever been terminated for cause from the Florida Medicaid program pursuant to s. 409.913, unless the applicant has been in good standing with the Florida Medicaid program for the most recent 5-year period?
Yes No
15. Has the applicant or any principal, officer, agent, managing employee, or affiliated person of the applicant been terminated for cause, pursuant to the appeals procedures established by the state, from any other state Medicaid program, unless the applicant has been in good standing with a state Medicaid program for the most recent 5-year period and the termination occurred at least 20 years before the date of the application?
Yes No
16. Is the applicant or any principal, officer, agent, managing employee, or affiliated person of the applicant currently listed on the United States Department of Health Human Services Office of Inspector General's List of Excluded Individuals and Entities? (If yes, please submit proof.)
Yes No

16b. If response is "yes" to question 16a, is the student loan default or delinquency the only reason you
are listed on the LEIE?

Yes No _____

17. Has the applicant or any principal, officer, agent, managing employee, or affiliated person of the applicant dispensed any medicinal drug based upon a communication that purports to be a prescription as defined by s. 465.003(14) or s. 893.02 when the pharmacist knows or has reason to believe that the purported prescription is not based upon a valid practitioner-patient relationship that includes a documented patient evaluation, including history and a physical examination adequate to establish the diagnosis for which any drug is prescribed and any other requirement established by board rule under chapter 458, chapter 459, chapter 461, chapter 463, chapter 464, or chapter 466?

Yes _____ No ___

18. Are you currently registered or permitted in any other states? (If yes, provide the state, permit type and	d
permit number for each permit. Attach a separate sheet if necessary.)	

Yes____No____

State	Permit Type	Permit Number

19. Has the applicant, affiliated pe	rsons, partners, officer, d	irectors, or Consultant Phar	nacist of Record
ever owned a pharmacy? (If yes, pr	ovide the name of the pharma	cy, the state where the pharmacy	is located and the
status of the pharmacy. Attach a separa	te sheet if necessary.)		

Yes	No
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No

Individual's Name	Pharmacy Name	State	Status

20. Has any disciplinary action ever been taken against any license, permit or registration issued to the applicant, affiliated persons, partners, officers, directors or consultant pharmacist of record?

Yes _____

21. Has the applicant, affiliated person, partner, officer, or director ever been convicted of a felony or misdemeanor, excluding minor traffic convictions? You must include all misdemeanors and felonies, even if adjudication was withheld by the court, so that you would not have a record of conviction.

Yes _____ No _____

22. Does the applicant, affiliated person, partner, officer, director have any outstanding fines, liens or overpayments assessed by a final order of the department? (*If yes, please answer question #22a,*)

Yes _____ No ____

22a. Does the applicant, affiliated pers department?	on, partner, offic	er, director have a	repayment plan approved by the
Yes No			
23. Is the applicant, affiliated persons prosecution for a crime in any jurisdic		ers, or directors, ui	nder investigation or
Yes No			
24. Is the applicant, affiliated persons administrative action by the licensing subdivisions?			
Yes No			
SECTION B. Please complete		of Location <u>only</u>	<u>V</u> .
1. Current Practice Location Address	3		
City	State		Zip
		Tolophone Numb	
E-Mail Address** (see note below)		Telephone Numb	er
2. New Practice Location Address			
City	State		Zip
E-Mail Address** (see note below)		Telephone Numb	er
Please provide your existing Pharma	cy Permit Numbe	er:	
Please provide your existing federal I	DEA Number:		
** <u>NOTE:</u> Under Florida law, email addresses are public records. If you do not want your e-mail address released in response to a public records request, do not provide an email address or send electronic mail to our office. Instead contact the office by phone or in writing.**			
SECTION C. Please complete	for Change of	of Ownership <u>o</u>	nly.
1. Are you changing physical locatio	ons with this cha	nge of ownership?	
Yes No <u>NOTE: If yes, please complete Section B above.</u>			
2. Please provide date when busines	s transaction fo	r the change of ow	nership will be completed?
Date:			
3. Do you have a signed letter from both the buyer and seller which indicates dates that pharmacy permit license should be transferred? <u>NOTE: A copy of the signed letter should be provided with your application.</u>			
Yes No			

SECTION D. Please complete for Stock Transfer of Ownership Interests only.		
1. Please provide the date when the transfer of ownership interest took place?		
Date:		
2. Did your company's FEIN change as a result of the transfer of ownership interest referenced in Section D, Question 1 above?		
Yes No	NOTE: If yes, please complete Section C above and include necessary fee.	

ALL QUESTIONS MUST BE ANSWERED OR YOUR APPLICATION WILL BE RETURNED

Section 456.013(1), F.S., requires that applicants supplement their applications as needed to reflect any material change in any circumstances or conditions stated in the application, which takes place between the initial filing of the application and the final grant or denial of the license, which might affect the decision of the department.

I swear and affirm that the statements contained in this application are true, complete, and correct and I agree that said statements shall form the basis of my application and I do authorize the Florida Board of Pharmacy to make any investigations that they deem appropriate and to secure any additional information concerning me, and I further authorize them to furnish any information they may have or have in the future concerning me to any person, corporation, institution, association, Board, or any municipal, county, state, or federal governmental agencies or units, and I understand according to the Florida Board of Pharmacy Statutes that a Pharmacy Permit may be revoked or suspended for presenting any false, fraudulent, or forged statement, certificate, diploma, or other item, in connection with an application for a license or permit, as set forth in Section 465.015(2)(a), F.S.

Under penalty of perjury I have read the foregoing document and that the facts stated in it are true. I recognize that providing false information may result in disciplinary action against my license or criminal penalties.

SIGNA	TURE			
		-		

(Owner or officer of establishment)

DATE	

FLORIDA DEPARTMENT OF LAW ENFORCEMENT

NOTICE FOR APPLICANTS SUBMITTING FINGERPRINTS WHERE CRIMINAL RECORD RESULTS WILL BECOME PART OF THE CARE PROVIDER BACKGROUND SCREENING CLEARINGHOUSE

NOTICE OF:

- SHARING OF CRIMINAL HISTORY RECORD INFORMATION WITH SPECIFIED AGENCIES,
- **RETENTION OF FINGERPRINTS**,
- PRIVACY POLICY, AND
- RIGHT TO CHALLENGE AN INCORRECT CRIMINAL HISTORY RECORD

This notice is to inform you that when you submit a set of fingerprints to the Florida Department of Law Enforcement (FDLE) for the purpose of conducting a search for any Florida and national criminal history records that may pertain to you, the results of that search will be returned to the Care Provider Background Screening Clearinghouse. By submitting fingerprints, you are authorizing the dissemination of any state and national criminal history record that may pertain to you to the Specified Agency or Agencies from which you are seeking approval to be employed, licensed, work under contract, or to serve as a volunteer, pursuant to the National Child Protection Act of 1993, as amended, and Section 943.0542, Florida Statutes. "Specified agency" means the Department of Health, the Department of Children and Family Services, the Division of Vocational Rehabilitation within the Department of Education, the Agency for Health Care Administration, the Department of Elder Affairs, the Department of Juvenile Justice, and the Agency for Persons with Disabilities when these agencies are conducting state and national criminal history background screening on persons who provide care for children or persons who are elderly or disabled. The fingerprints submitted will be retained by FDLE and the Clearinghouse will be notified if FDLE receives Florida arrest information on you.

Your Social Security Number (SSN) is needed to keep records accurate because other people may have the same name and birth date. Disclosure of your SSN is imperative for the performance of the Clearinghouse agencies' duties in distinguishing your identity from that of other persons whose identification information may be the same as or similar to yours.

Licensing and employing agencies are allowed to release a copy of the state and national criminal record information to a person who requests a copy of his or her own record if the identification of the record was based on submission of the person's fingerprints. Therefore, if you wish to review your record, you may request that the agency that is screening the record provide you with a copy. After you have reviewed the criminal history record, if you believe it is incomplete or inaccurate, you may conduct a personal review as provided in s. 943.056, F.S., and Rule 11C8.001, F.A.C. If national information is believed to be in error, the FBI should be contacted at 304-625-2000. You can receive any national criminal history record that may pertain to you directly from the FBI, pursuant to 28 CFR Sections 16.30-16.34. You have the right to obtain a prompt determination as to the validity of your challenge before a final decision is made about your status as an employee, volunteer, contractor, or subcontractor.

Until the criminal history background check is completed, you may be denied unsupervised access to children, the elderly, or persons with disabilities.

The FBI's Privacy Statement follows on a separate page and contains additional information.

US Department of Justice, Federal Bureau of Investigation, Criminal Justice Information Services Division

Privacy Statement

Authority: The FBI's acquisition, preservation and exchange of information requested by this form is generally authorized under 28 U.S.C. 534. Depending on the nature of your application, supplemental authorities include numerous Federal statutes, hundreds of State statutes pursuant to Pub.L.92-544, Presidential executive orders, regulations and/or orders of the Attorney General of the United States, or other authorized authorities. Examples include, but are not limited to: 5 U.S.C. 9101; Pub.L.94-29; Pub.L.101-604; and Executive Orders 10450 and 12968. Providing the requested information is voluntary; however, failure to furnish the information may affect timely completion of approval of your application.

Social Security Account Number (SSAN): Your SSAN is needed to keep records accurate because other people may have the same name and birth date. Pursuant to the Federal Privacy Act of 1974 (5 USC 552a), the requesting agency is responsible for informing you whether disclosure is mandatory or voluntary, by what statutory or other authority your SSAN is solicited, and what uses will be made of it. Executive Order 9397 also asks Federal Agencies to use this number to help identify individuals in agency records.

Principal Purpose: Certain determinations, such as employment, security, licensing and adoption, may be predicated on fingerprint based checks. Your fingerprints and other information contained on (and along with) this form may be submitted to the requesting agency, the agency conducting the application investigation, and/or FBI for the purpose of comparing the submitted information to available records in order to identify other information that may be pertinent to the application. During the processing of this application, and for as long hereafter as my be relevant to the activity for which this application is being submitted, the FBI(may disclose any potentially pertinent information to the requesting agency and/or to the agency conducting the investigation. The FBI may also retain the submitted information in the FBI's permanent collection of fingerprints and related information, where it will be subject to comparisons against other submissions received by the FBI. Depending on the nature of your application, the requesting agency and/or the agency conducting the application investigation may also retain the fingerprints and other submitted information for other authorized purposes of such agency(ies).

Routine Uses: The fingerprints and information reported on this form may be disclosed pursuant to your consent, and may also be disclosed by the FBI without your consent as permitted by the Federal Privacy Act of 1974 (5 USC 552a(b)) and all applicable routine uses as many be published at any time in the Federal Register, including the routine uses for the FBI Fingerprint Identification Records System (Justice, FBI-009) and the FBI's Blanket Routine Uses (Justice/FBI-BRU). Routine uses include, but are not limited to, disclosures to: appropriate governmental authorities responsible for civil or criminal law enforcement counterintelligence, national security or public safety matters to which the information may be relevant; to State and local governmental agencies and nongovernmental entities for application processing as authorized by Federal and State legislation, executive order, or regulation, including employment, security, licensing, and adoption checks; and as otherwise authorized by law, treaty, executive order, regulation, or other lawful authority. If other agencies are involved in processing the application, they may have additional routine uses.

Additional Information: The requesting agency and/or the agency conducting the application investigation will provide you additional information pertinent to the specific circumstances of this application, which may include identification of other authorities, purposes, uses, and consequences of not providing requested information. In addition, any such agency in the Federal Executive Branch has also published notice.

Electronic Fingerprinting

Take this form with you to the Live Scan service provider. Please check the service provider's requirements to see if you need to bring any additional items.

- Background screening results are obtained from the Florida Department of Law Enforcement and the Federal Bureau of Investigation by submitting to a fingerprint scan using the Livescan method;
- You can find a Livescan service provider at: <u>http://www.floridahealth.gov/licensing-and-regulation/background-screening/livescan-service-providers.html</u>
- Failure to submit background screening will delay your application;
- Applicants may use any Livescan service provider approved by the Florida Department of Law Enforcement to submit their background screening to the department;
- If you do not provide the correct Originating Agency Identification (ORI) number to the livescan service provider the Board office <u>will not</u> receive your background screening results;
- You must provide accurate demographic information to the livescan service provider at the time your fingerprints are taken, *including your Social Security number (SSN)*;
- The ORI number for the Board of Pharmacy is **EDOH4680Z**.
- Typically background screening results submitted through a Livescan service provider are received by the Board within 24-72 hours of being processed.
- If you obtain your livescan from a service provider who does not capture your photo you may be required to be reprinted by another agency in the future.

Name:		SSN#:
Aliases:		
Address:		Apt. Number:
City:	State:	Zip Code:
Date of Birth: ////Place (MM/DD/YYYY)	of Birth:	
Weight: Height:	Eye Color:	Hair Color:
Race:	Sex: (M=Male; F=Female)	
Citizenship:		
Transaction Control Number (TCN#): (This will be provided to you by the Live Scan S		

Keep this form for your records.



Item #1- Consultant Pharmacist of Record Designation and Privacy Statement Acknowledgement

To: Florida Board of Pharmacy Post Office Box 6320 Tallahassee, FL 32314-6320 (850) 245-4474 - phone (850) 921-5389 - fax MQAPharmPDMAffiliate@flhealth.gov File #: (if known)

License #: (if applicable)

Section A. Consultant I	Pharmacist of Record (COR) I	Designation
Applicant/Pharmacy Name:		
Applicant/Pharmacy Mailing	Address:	
City	State	Zip
Incoming COR Name:		License#:
		PU
Date Beginning as COR:	Incoming COR Signature	
Transaction Control Number	· (TCN) – related to Livescan Fingerpr	ints (optional if known):
	on regarding Livescan Fingerprints go to: <u>http://fl</u>	
OBTIONAL: Only provide the	following information if there is an Outg	oing COB at the current pharmacu
Outgoing COR Name:	Tonowing information in there is an outgo	License#:
		PU
Date Ending as COR:	Outgoing COR Signature (optional)	
Section B. Incoming C	COR Privacy Statement Ackno	owledgement
	be completed by same person listed in <u>Sec</u>	
regarding the sharing, retention	d the statement from the Florida Depart on, privacy and right to challenge incor ment from the Federal Bureau of Inves	rect criminal history records and
Date:	Incoming COR Signature	



Item #2- Affiliate/Owner Privacy Statement Acknowledgement

To be completed by EACH Affiliate/Owner listed in the application.

To: Florida Board of Pharmacy, Post Office Box 6320, Tallahassee, FL 32314-6320 (850) 245-4474 - phone * (850) 921-5389 – fax * <u>MQAPharmPDMAffiliate@flhealth.gov</u>

Applicant Name:		
Affiliate/Owner Mailing Address:		
City	State	Zip
		P
Affiliate/Owner E-Mail ** (see note below)	Affiliate/Owner	Telephone Number
Affiliate/Owner Transaction Control Nur ** For more information regarding Livescan Fingerp		

I have been provided and read the statement from the Florida Department of Law Enforcement regarding the sharing, retention, privacy and right to challenge incorrect criminal history records and the "Privacy Statement" document from the Federal Bureau of Investigation."

Affiliate/Owner Signature (Required)

Date (of signature)



Item #3 - Policy and Procedure Questions

To be completed by Institutional Class II Pharmacy Permit Applicants

The Consultant Pharmacist of Record is responsible for developing and maintaining a current policy and procedure manual. A copy of the permittee's policy and procedure manual as provided herein shall accompany the permit application. The original policy and procedure manual shall be kept within the Class II Institutional Pharmacy and shall be available for inspection by the Department of Health. The board office will approve the policy and procedure manual based upon answers submitted for the following questions, where applicable, by using excerpts or summaries from the policy and procedure manual.

- List the following: Firm Name: Doing business as (d/b/a): Telephone number: Address: Permit number (if already licensed as an institutional pharmacy):
- 2) Explain the practice setting of the proposed facility.
- 3) What are the objectives and purpose of the permittee? Give detailed explanation of the services of the facility scope and practice.
- 4) What are the experience, qualifications, special education, and/or training of the compounding pharmacist? Please provide a resume.
- 5) Address the ratio of supportive personnel to each pharmacist. How will the supportive personnel be utilized? Include a job description for any such supportive personnel.
- 6) Describe the drug delivery system. Begin with the ordering of medications and track your procedures up to delivery to the patient. If utilizing remote medication order processing and the pharmacist is not an employee of the institution, describe the pharmacist and institution's responsibility.
- 7) What categories of parenteral/enteral products will be prepared (i.e. IV, enteral, irrigating, and oncology products)? Include sample labels.
- 8) What is the policy regarding the delivery of parenteral/enteral products to the patient? Describe methods used and trace the path the product takes from the time it leaves the permittee until it reaches the patient. Describe how products are protected from extreme temperature conditions.

- 9) Address the policy and procedure, special equipment and special techniques to dispense sterile preparations for parenteral therapy/nutrition. If this type of dispensing will not be performed, please state so accordingly.
- 10) Address the policy and procedure, special equipment and special techniques to dispense sterile jejunostomy feeding/sterile irrigation solutions. If this type of dispensing will not be performed, please state so accordingly.
- 11) Address the policy and procedure, special equipment and special techniques to dispense cytotoxic or anti-neoplastic agents. If this type of dispensing will not be performed, please state so accordingly.
- 12) What is the procedure for the annual review and updating of the policy and procedure manual?
- 13) Include the layout/floor plan of the pharmacy. The drawing must include the dimensions of the clean room and the pharmacy, location of the hood, sink and other equipment. The drawing must also show the location of the clean room relative to other pharmacy and storage areas.
- 14) Include a sample copy of a patient profile.
- 15) Address the use of aseptic techniques.
- 16) Describe the Quality Assurance Program.
- 17) Describe with detail the policy and procedure for patient education, including the personnel involved.
- 18) Address the policy and procedure for handling waste and returns.
- 19) Describe the type of certified laminar flow hood(s) used and the frequency of certification.
- 20) Describe the refrigerator/freezer to be used.
- 21) Describe appropriate waste containers for:
 - a. Used needles and syringes.
 - b. Cytotoxic waste including disposable apparel used in preparation.
- 22) Address the following supplies to be used: gloves, mask, gowns, needles, syringes, disinfectant cleaning agents, clean towels, hand-washing materials with bactericidal properties, vacuum containers/transfer sets, and spill kits for cytotoxic agent spills.
- 23) Address the following references to be used:
 - a. Chapters 465 and 893, F.S., and Rule Title 64B16, F.A.C.
 - b. Authoritative Therapeutic Reference.
 - c. Handbook of injectable drugs by American Society of Health-System Pharmacists.
- 24) Occupational Safety and Health Administration guidelines for safe handling of cytotoxic drugs.



Item #4 - Policy and Procedure Questions

To be completed by Modified Institutional Class II Pharmacy Permit Applicants

Modified Institutional Class II pharmacies are those institutional pharmacies in short-term, primary care treatment centers that meet all the requirements for a Class II permit, except space and equipment requirements. Modified Class II Institutional pharmacies are designated as Type "A", Type "B" and Type "C" according to the type of specialized pharmaceutical delivery system utilized. Please review Rule 64B16-28.702, Florida Administrative Code for specific requirements.

The Consultant Pharmacist of Record is responsible for developing and maintaining a current policy and procedure manual. A copy of the permittee's policy and procedure manual as provided herein shall accompany the permit application. The original policy and procedure manual shall be kept within the Modified Class II Institutional Pharmacy and shall be available for inspection by the Department of Health. The board office will approve the policy and procedure manual based upon answers submitted for the following questions, where applicable, by using excerpts or summaries from the policy and procedure manual

- List the following: Firm Name: Doing business as (d/b/a): Telephone number: Address: Consultant pharmacist of record:
- 2) Describe the purpose of the establishment. What sector of the community are you serving?
- 3) Is this is an inpatient facility? If so, how many beds are housed in the facility? What is the average length of stay?
- 4) List the drug formulary to be used.
- 5) Include a diagram of pharmacy storage space and a description of drug security measures.
- 6) Describe the consultant pharmacist of record's responsibilities.
- 7) Under whose DEA registration will controlled substances be ordered?
- 8) Describe the drug delivery system. Begin with the ordering of medications and track your procedures up to delivery to the patient.
- 9) Include a statement that perpetual inventory records will be maintained for controlled substances and injectable inventory.
- 10) Include a statement to the effect that no drugs will be dispensed from the facility.

If compounding sterile preparations, please answer the additional questions below.

- 11) If compounding sterile preparations, describe compliance with Rule 64B16-27.797, F.A.C.
- 12) What categories of parenteral/enteral products will be prepared (i.e. IV, enteral, irrigating, and oncology products)? Include sample labels.
- 13) What is the policy regarding the delivery of parenteral/enteral products to the patient? Describe methods used and trace the path the product takes from the time it leaves the permittee until it reaches the patient. Describe how this product is protected from extreme temperature conditions.
- 14) Address the policy and procedure, special equipment and special techniques to dispense sterile preparations for parenteral therapy/nutrition. If this type of dispensing will not be performed, please state so accordingly.
- 15) Address the policy and procedure, special equipment and special techniques to dispense sterile jejunostomy feeding/sterile irrigation solutions. If this type of dispensing will not be performed, please state so accordingly.
- 16) Address the policy and procedure, special equipment and special techniques to dispense cytotoxic or anti-neoplastic agents. If this type of dispensing will not be performed, please state so accordingly.
- 17) What is the procedure for the annual review and updating of the policy and procedure manual?
- 18) Include the layout/floor plan of the pharmacy. The drawing must include the dimensions of the clean room and the pharmacy, location of the hood, sink and other equipment. The drawing must also show the location of the clean room relative to other pharmacy and storage areas.
- 19) Include a sample copy of a patient profile.
- 20) Address the use of aseptic techniques.
- 21) Describe the Quality Assurance Program.
- 22) Describe with detail the policy and procedure for patient education, including the personnel involved.
- 23) Address the policy and procedure for handling waste and returns.
- 24) Describe the type of certified laminar flow hood(s) to be used and the frequency of certification.
- 25) Describe the refrigerator/freezer to be used.
- 26) Describe appropriate waste containers for:
 - a. Used needles and syringes.
 - b. Cytotoxic waste including disposable apparel used in preparation.
- 27) Address the following supplies to be used: gloves, mask, gowns, needles, syringes, disinfectant cleaning agents, clean towels, hand-washing materials with bactericidal properties, vacuum containers/transfer sets, and spill kits for cytotoxic agent spills.
- 28) Address the following references to be used:
 - a. Chapters 465 and 893, F.S., and Title 64B16, F.A.C.
 - b. Authoritative Therapeutic Reference.
 - c. Handbook of Injectable Drugs by American Society of Health-System Pharmacists.
 - d. Occupational Safety and Health Administration guidelines for safe handling of cytotoxic drugs.