

Teaching Institution Application for Registration (Form DHHS 224-C)

NC Department of Health and Human Services

Division of Mental Health, Developmental Disabilities, and Substance Abuse Services - Drug Control Unit

3008 Mail Center Service Center Raleigh, North Carolina 27699-3008 (919) 733-1765

Application Instructions – PLEASE READ THESE INSTRUCTIONS CAREFULLY

This application will be used by the North Carolina Department of Health and Human Services' Drug Control Unit to initiate a registration renewal under the North Carolina Controlled Substances Act of 1971 as well as assist in determining whether or not the registrant is in compliance with State and Federal laws pertaining to controlled substances. Therefore, please fill out this application in its entirety. Do not leave any fields blank, rather indicate that a field is not applicable by typing "N/A" in the space provided. Failure to complete the entire form will result in the application being returned to the registrant along with a request for additional information. To submit this Application for Reregistration, e-mail both the completed electronic PDF and a signed PDF copy to <u>nccsareg@dhhs.nc.gov</u> along with a signed PDF copy of a Registrant Disclosure of Loss, Diversion, or Destruction of Controlled Substances (Addendum to Forms DHHS 226 and 227).

Attestation

By signing below, you attest that the information provided on this form is true, accurate, and complete to the best of your knowledge. All responses are subject to verification by the North Carolina Department of Health and Human Services' Drug Control Unit. Furthermore, you acknowledge that you have read and understand NC GS 90-101(a1):

"(a1) Any physician who prescribes or dispenses Buprenorphine for the treatment of opiate dependence shall annually register with the Department, in accordance with rules adopted by the Commission. In the application for registration under this subsection, the applicant shall document plans to ensure that patients are directly engaged or referred to a qualified provider to receive counseling and case management, as appropriate, and shall acknowledge the application of federal confidentiality regulations to patient information. Applicant plans for referral to appropriate services shall be a written document and may include either an executed memorandum of agreement, contractual arrangement, or linkage agreement with qualified providers. The Department shall provide assistance upon request to physicians registered under this subsection to identify and establish linkages with qualified providers of counseling and case management. The Department shall provide the North Carolina Medical Board with any evidence of noncompliance with this subsection by a qualified physician prior to taking action to rescind the physician's registration to prescribe or dispense Buprenorphine for the treatment of opiate dependency."

	Date
Signature	Phone Number
Name and Title	E-Mail Address

Section A - Registrant Information			
Registrant's Name		-	
Physical Address	Physical County		
Facility's State, City, Zip			
Mailing Address	Phone Number		
Mailing State, City, Zip			

Cection B Registration Olassinea				
B1. Check all applicable drug schedules in which y	ou are applying for:			
Schedule II (Narcotic)	\Box Schedule III (Narcotic)	□ Schedule	IV	
□ Schedule IIN (Non-narcotic)	□ Schedule IIIN (Non-narcotic)	□ Schedule V		
B2. Are you currently authorized to manufacture, otherwise handle controlled substances in the sch North Carolina or the Federal Government?	· · · ·		□ Yes	□ No
B3. Has the registrant been convicted of a felony of possession, distribution, or dispensing of controlled	-	anufacture,	□ Yes	🗆 No
B4. Has any previous registration held by the regis under Federal CSA or NCCSA been surrendered, re		J. J	□ Yes	🗆 No

Section B - Registration Classification

If you answered "Yes" to questions B3 and/or B4, please submit a letter along with this application setting forth the circumstances of such action.

Section C - Point of Contact

A Drug Control Inspector will conduct an unannounced inspection of the applicant's facility at some point during the registration period. Please provide a list of up to three individuals for whom the Inspector should ask for upon arrival at the facility. The names and titles provided should be listed in the desired order of contact and should include individuals who are knowledgeable of and possess some degree of responsibility for the disposition of controlled substances at the facility. Any phone numbers provided for points of contact in Section C should be a direct line in order to assist the Drug Control Unit with reaching the correct individual(s) if needed – the central phone number provided in Section A will serve as a backup. Please note that the Inspector may also interview other persons other than those listed below at his/her discretion.

Primary Contact	Name: E-mail:	Title: Phone:
Secondary Contact	Name: E-mail:	Title: Phone:
Tertiary Contact	Name: E-mail:	Title: Phone:

Section D - State Registration History

D1. Please select the event below that best describes your reason for submitting an Application for Registration (Form DHHS 224) and provide an answer to each supporting question for that event (choose only one answer from below)

\Box The application is for a new facility / first time registrant	\Box The application reflects a name change for a registrant		
Anticipated Opening Date:	Name on Previous Registration:		
	Previous DHHS Registration No:		
\Box The application reflects a change of location/address for a registrant	\Box The application reflects a change in ownership		
Name on Previous Registration:	Name on Previous Registration:		
Previous Address (Line 1):	Previous DHHS Registration No:		
Previous Address (Line 2):	Was Business Sold or Merged:		
Previous City:	Percentage of Ownership Sold:		
Previous DHHS Registration No:	Corporate or Branch Level Sold:		

Section E - Drug Enforceme	ent Administration (DEA) Registration	on		
E1. Does the applicant currently posses	s any controlled substances?		□ Yes	□ No
E2. What is the current status of the app	plicant's DEA Registration? (choose only one answ	ver from below and	provide the request	ted information)
\Box Valid Registration in pos	session Name on Registration:		DEA Number:	
\Box Applied for Registration	Applicant's Name:		Date Applied:	
\Box DEA Registration will be	applied for pending approval of NC DHHS Registra	tion		
□ Other <i>(explain)</i> :				
E3. Who is responsible for controlled su	bstances? (this is the individual who signed DEA F	orm 224):		
E4. Has the applicant granted Power of	Attorney to any individuals for ordering controlle	d substances?	□ Yes	🗆 No
If yes, please provide the n	ame(s) of the individual(s):			

Section F - Primary Supplier of Controlled Substances

Supplier Name	
Address	City
State	Zip Code
Sales Rep's Name	Phone Number

Section G - Secondary Supplier of Controlled Substances			
Supplier Name			
Address		City	
State		Zip Code	
Sales Rep's Name		Phone Number	

Section H - Storage and Security

H1. How many total storage locations are utilized for the storage of controlled substances at the facility? Describe the type of storage equipment for each location (i.e. wall cabinet, combination safe, keyed safe, etc.).

H2. How is access to the controlled substance inventory location(s) controlled? List the persons and/or titles and number of individuals with access, describe how key control is practiced, and provide any other information deemed pertinent to assuring the security of controlled substances at the facility.

H3. How are unexecuted controlled substance order forms stored?

Section I - Records

I1. Biennial Inventory Date

12. Describe the procedure for purchasing and receiving Schedule II controlled substances. How are DEA Form-222s, invoices, and any other documents acknowledging the purchase and receipt of Schedule II controlled substances recorded and maintained? If the applicant is not registered for Schedule II, please write/type "N/A" for this question.

I3. Describe the procedure for purchasing and receiving Schedule III, IV, and V controlled substances. How are pharmacy provider requisition forms, invoices, and any other documents acknowledging the purchase and receipt of Schedule III, IV, and V controlled substances recorded and maintained? If the applicant is not registered for Schedule III, IV, and V, please write/type "N/A" for this question.

14. Describe how you intend to use controlled substances are used for training purposes? What records will be kept to document any activity involving the use of a controlled substance?

Section J - Effective Controls for the Prevention of Diversion

J1. Other than physical security measures that have already been discussed in previous sections of this document, what steps is the applicant taking to maintain effective controls for the prevention of diversion of controlled substances? Answers should include, but are not limited to, software reporting systems being utilized to monitor user and drug activity as well as the frequency and individuals involved in the review of such material.