

GENERAL ASSEMBLY OF NORTH CAROLINA
SESSION 2003

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SENATE DRS55113-LNf-106 (3/25)

Short Title: Controlled Substance/Physician Registration. (Public)

Sponsors: Senator Purcell.

Referred to:

A BILL TO BE ENTITLED

AN ACT TO REQUIRE PHYSICIANS WHO DISPENSE THE CONTROLLED
SUBSTANCE BUPRENORPHINE FOR THE TREATMENT OF OPIATE
DEPENDENCE TO REGISTER WITH THE DEPARTMENT OF HEALTH AND
HUMAN SERVICES.

The General Assembly of North Carolina enacts:

SECTION 1. G.S. 90-101 reads as rewritten:

"§ 90-101. Annual registration and fee to engage in listed activities with controlled substances; effect of registration; exceptions; waiver; inspection.

(a) Every person who manufactures, distributes, dispenses, or conducts research with any controlled substance within this State or who proposes to engage in any of these activities shall annually register with the North Carolina Department of Health and Human Services, in accordance with rules adopted by the Commission, and shall pay the registration fee set by the Commission for the category to which the applicant belongs. An applicant for registration shall file an application for registration with the Department of Health and Human Services and submit the required fee with the application. The categories of applicants and the maximum fee for each category are as follows:

<u>CATEGORY</u>	<u>MAXIMUM FEE</u>
Clinic	\$150.00
Hospital	350.00
Nursing Home	150.00
Teaching Institution	150.00
Researcher	150.00
Analytical Laboratory	150.00
Distributor	600.00

1 Manufacturer..... 700.00.

2 (a1) Any physician who dispenses Buprenorphine for the treatment of opiate
3 dependence shall annually register with the Department, in accordance with rules
4 adopted by the Commission, and shall pay an annual registration fee of two hundred
5 fifty dollars (\$250.00). In the application for registration under this subsection, the
6 applicant shall document plans to ensure that patients are directly engaged or referred to
7 a qualified provider to receive counseling and case management, as appropriate, and
8 shall acknowledge the application of federal confidentiality regulations to patient
9 information.

10 (b) Persons registered by the North Carolina Department of Health and Human
11 Services under this Article (including research facilities) to manufacture, distribute,
12 dispense or conduct research with controlled substances may possess, manufacture,
13 distribute, dispense or conduct research with those substances to the extent authorized
14 by their registration and in conformity with the other provisions of this Article.

15 (c) The following persons shall not be required to register and may lawfully
16 possess controlled substances under the provisions of this Article:

17 (1) An agent, or an employee thereof, of any registered manufacturer,
18 distributor, or dispenser of any controlled substance if such agent is
19 acting in the usual course of his business or employment;

20 (2) The State courier service operated by the Department of
21 Administration, a common or contract carrier, or a public
22 warehouseman, or an employee thereof, whose possession of any
23 controlled substance is in the usual course of his business or
24 employment;

25 (3) An ultimate user or a person in possession of any controlled substance
26 pursuant to a lawful order of a practitioner;

27 (4) Repealed by Session Laws 1977, c. 891, s. 4.

28 (5) Any law-enforcement officer acting within the course and scope of
29 official duties, or any person employed in an official capacity by, or
30 acting as an agent of, any law-enforcement agency or other agency
31 charged with enforcing the provisions of this Article when acting
32 within the course and scope of official duties; and

33 (6) A practitioner, as defined in G.S. 90-87(22)a., who is required to be
34 licensed in North Carolina by his respective licensing board.

35 (d) The Commission may, by rule, waive the requirement for registration of
36 certain classes of manufacturers, distributors, or dispensers if it finds it consistent with
37 the public health and safety.

38 (e) A separate registration shall be required at each principal place of business,
39 research or professional practice where the registrant manufactures, distributes,
40 dispenses or uses controlled substances.

41 (f) The North Carolina Department of Health and Human Services is authorized
42 to inspect the establishment of a registrant, applicant for registration, or practitioner in
43 accordance with rules adopted by the Commission.

1 (g) Practitioners licensed in North Carolina by their respective licensing boards
2 may possess, dispense or administer controlled substances to the extent authorized by
3 law and by their boards.

4 (h) A physician licensed by the North Carolina Medical Board pursuant to
5 Article 1 of this Chapter may possess, dispense or administer tetrahydrocannabinols in
6 duly constituted pharmaceutical form for human administration for treatment purposes
7 pursuant to rules adopted by the Commission.

8 (i) A physician licensed by the North Carolina Medical Board pursuant to
9 Article 1 of this Chapter may dispense or administer Dronabinol or Nabilone as
10 scheduled in G.S. 90-90(5) only as an antiemetic agent in cancer chemotherapy."

11 **SECTION 2.** This act becomes effective October 1, 2003.