§ 90-88. Authority to control.

(a) The Commission may add, delete, or reschedule substances within Schedules I through VI of this Article on the petition of any interested party, or its own motion. In every case the Commission shall give notice of and hold a public hearing pursuant to Chapter 150B of the General Statutes prior to adding, deleting or rescheduling a controlled substance within Schedules I through VI of this Article, except as provided in subsection (d) of this section. A petition by the Commission, the North Carolina Department of Justice, or the North Carolina Board of Pharmacy to add, delete, or reschedule a controlled substance within Schedules I through VI of this Article shall be placed on the agenda, for consideration, at the next regularly scheduled meeting of the Commission, as a matter of right.

(a1) In making a determination regarding a substance, the Commission shall consider the following:

(1) The actual or relative potential for abuse;
(2) The scientific evidence of its pharmacological effect, if known;
(3) The state of current scientific knowledge regarding the substance;
(4) The history and current pattern of abuse;
(5) The scope, duration, and significance of abuse;
(6) The risk to the public health;
(7) The potential of the substance to produce psychic or physiological dependence liability; and
(8) Whether the substance is an immediate precursor of a substance already controlled under this Article.

(b) After considering the required factors, the Commission shall make findings with respect thereto and shall issue an order adding, deleting or rescheduling the substance within Schedules I through VI of this Article.

(c) If the Commission designates a substance as an immediate precursor, substances which are precursors of the controlled precursor shall not be subject to control solely because they are precursors of the controlled precursor.

(d) If any substance is designated, rescheduled or deleted as a controlled substance under federal law, the Commission shall similarly control or cease control of, the substance under this Article unless the Commission objects to such inclusion. The Commission, at its next regularly scheduled meeting that takes place 30 days after publication in the Federal Register of a final order scheduling a substance, shall determine either to adopt a rule to similarly control the substance under this Article or to object to such action. No rule-making notice or hearing as specified by Chapter 150B of the General Statutes is required if the Commission makes a decision to similarly control a substance. However, if the Commission makes a decision to object to adoption of the federal action, it shall initiate rule-making procedures pursuant to Chapter 150B of the General Statutes within 180 days of its decision to object.

(e) The Commission shall exclude any nonnarcotic substance from the provisions of this Article if such substance may, under the federal Food, Drug and Cosmetic Act, lawfully be sold over-the-counter without prescription.

(f) Authority to control under this Article does not include distilled spirits, wine, malt beverages, or tobacco.

(g) The Commission shall similarly exempt from the provisions of this Article any chemical agents and diagnostic reagents not intended for administration to humans or other animals, containing controlled substances which either (i) contain additional adulterant or denaturing agents so that the resulting mixture has no significant abuse potential, or (ii) are packaged in such a form or concentration that the particular form as packaged has no

significant abuse potential, where such substance was exempted by the Federal Bureau of Narcotics and Dangerous Drugs.

(h) Repealed by Session Laws 1987, c. 413, s. 4.

(i) The North Carolina Department of Health and Human Services shall maintain a list of all preparations, compounds, or mixtures which are excluded, exempted and excepted from control under any schedule of this Article by the United States Drug Enforcement Administration and/or the Commission. This list and any changes to this list shall be mailed to the North Carolina Board of Pharmacy, the State Bureau of Investigation and each district attorney of this State. (1971, c. 919, s. 1; 1973, c. 476, s. 128; cc. 524, 541; c. 1358, ss. 2, 3, 15; 1977, c. 667, s. 3; 1981, c. 51, s. 9; 1987, c. 413, ss. 1-4; 1989, c. 770, s. 16; 1997-443, s. 11A.118(a); 2000-189, s. 4; 2001-487, s. 22.)