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- 1 AN ACT concerning public health.
- 2 Be it enacted by the People of the State of Illinois,
- 3 represented in the General Assembly:
- 4 Section 1. Short title. This Act may be cited as the
- 5 Ephedra Prohibition Act.
- 6 Section 5. Findings. The General Assembly finds the 7 following:
- 8 (1) Over 3 billion servings of ephedra are consumed 9 by Americans each year.
 - (2) Ephedra, or ma huang, has been associated with a wide range of severe adverse events, including death.
 - (3) The U.S. Food and Drug Administration has received over 18,000 reports of adverse reactions by ephedra users, including strokes, seizures, heart attacks, and deaths.
 - (4) The Inspector General of the U.S. Department of Health and Human Services noted in a report on ephedra adverse events that 60% of those adverse events were experienced by people under the age of 40.
 - (5) A study reported in the Annals of Internal Medicine concluded that, compared with other herbs, ephedra is associated with a greatly increased risk for adverse reactions and that the use of ephedra should be restricted.
 - (6) The American Medical Association and the consumer group, Public Citizen, have called for a nationwide ban on ephedra.
 - (7) The National Collegiate Athletics Association, the National Football League, and the International Olympic Committee have all banned ephedra use by their athletes because of concerns about the safety of this

- 1 dietary supplement.
- 2 (8) The U.S. Army has banned the sale of ephedra
- 3 products in army commissaries worldwide after 33 army
- 4 personnel died from consuming ephedra products.
- 5 (9) Canada, Britain, Germany, and Australia have
- 6 all taken steps to restrict the sale of ephedra products.
- 7 Section 10. Purpose. The purpose of this Act is to ban
- 8 the sale of all dietary supplements containing ephedrine
- 9 alkaloids in the State of Illinois regardless of the age of
- 10 the purchaser in order to protect the health and public
- 11 safety of Illinois residents.
- 12 Section 15. Definitions:
- "Ephedra" means herbs and herbal products that contain
- 14 ephedrine alkaloids, including ma huang, Chinese ephedra,
- ephedra sinica, ephedra herb powder, epitonin, or any extract
- 16 of those substances, but does not include any drug that
- 17 contains ephedrine and is lawfully sold, transferred, or
- 18 furnished over the counter with or without a prescription
- 19 pursuant to the federal Food, Drug, and Cosmetic Act (21
- 20 U.S.C. 301 and following) or regulations adopted under that
- 21 Act.
- 22 "Person" means any natural person, individual,
- 23 corporation, unincorporated association, proprietorship,
- 24 firm, partnership, joint venture, joint stock association, or
- any other business organization or entity.
- 26 Section 20. Prohibition.
- 27 (a) No person may sell or offer for sale any dietary
- 28 supplement containing any quantity of ephedra or ephedrine
- 29 alkaloids to any person located within the State or to any
- 30 person making the purchase from within the State.
- 31 (b) The prohibition in subsection (a) of this Section

- 1 does not apply to the sale of any product that receives
- 2 explicit approval as safe and effective for its intended use
- 3 under the federal Food, Drug, and Cosmetic Act (21 U.S.C.
- 4 355) or is lawfully marketed under an over-the-counter
- 5 monograph issued by the U.S. Food and Drug Administration.
- 6 Section 25. Penalties.
- 7 (a) Any person who violates this Act is guilty of a Class
- 8 A misdemeanor. The penalty is imprisonment for less than one
- 9 year or a fine of not more than \$5,000 or both for a first
- 10 offense.
- 11 (b) For a subsequent violation of this Act, a person is
- 12 guilty of a Class 3 felony, and the penalty is imprisonment
- for less than 5 years or a fine of not more than \$20,000 or
- 14 both.
- 15 Section 99. Effective date. This Act shall take effect
- 16 upon becoming law.