THE GENERAL ASSEMBLY OF PENNSYLVANIA

SENATE BILL

No. 955

Session of 2005

INTRODUCED BY KASUNIC, COSTA, BOSCOLA, STOUT, FONTANA, TARTAGLIONE, WENGER, ORIE, WOZNIAK, RAFFERTY AND RHOADES, OCTOBER 26, 2005

REFERRED TO PUBLIC HEALTH AND WELFARE, OCTOBER 26, 2005

AN ACT

Amending the act of April 14, 1972 (P.L.233, No.64), entitled "An act relating to the manufacture, sale and possession of 3 controlled substances, other drugs, devices and cosmetics; 4 conferring powers on the courts and the secretary and Department of Health, and a newly created Pennsylvania Drug, 6 Device and Cosmetic Board; establishing schedules of 7 controlled substances; providing penalties; requiring registration of persons engaged in the drug trade and for the 8 9 revocation or suspension of certain licenses and registrations; and repealing an act," further providing for 10 schedules of controlled substances, for professional 11 12 prescription administration and dispensing, for prohibited 13 acts and penalties, for prohibitions relating to liquefied ammonia gas, precursors and chemicals and for effect on local 14 15 ordinances. 16 The General Assembly of the Commonwealth of Pennsylvania hereby enacts as follows: 17 18 Section 1. Section 4(5) of the act of April 14, 1972 19 (P.L.233, No.64), known as The Controlled Substance, Drug, Device and Cosmetic Act, amended July 3, 1985 (P.L.138, No.39), 20 21 is amended to read: 22 Section 4. Schedules of Controlled Substances. -- The following schedules include the controlled substances listed or

to be listed by whatever official name, common or usual name,

24

- 1 chemical name, or trade name designated.
- 2 * * *
- 3 (5) Schedule V--In determining that a substance comes within
- 4 this schedule, the secretary shall find: a low potential for
- 5 abuse relative to the substances listed in Schedule IV;
- 6 currently accepted medical use in the United States; and limited
- 7 physical dependence and/or psychological dependence liability
- 8 relative to the substances listed in Schedule IV. The following
- 9 controlled substances are included in this schedule:
- 10 (i) Any compound, mixture, or preparation containing limited
- 11 quantities of any of the following narcotics or any of their
- 12 salts, which shall include one or more nonnarcotic active
- 13 medicinal ingredients in sufficient proportion to confer upon
- 14 the compound, mixture, or preparation, valuable medicinal
- 15 qualities other than those possessed by the narcotic alone:
- 16 1. Not more than 200 milligrams of codeine, or any of its
- 17 salts, per 100 milliliter or per 100 grams and not more than 10
- 18 milligrams per dosage unit.
- 19 2. Not more than 100 milligrams of dihydrocodeine, or any of
- 20 its salts, per 100 milliliters or per 100 grams and not more
- 21 than 5 milligrams per dosage unit.
- 22 3. Not more than 100 milligrams of ethylmorphine, or any of
- 23 its salts, per 100 milliliters or per 100 grams and not more
- 24 than 5 milligrams per dosage unit.
- 25 4. Not more than 2.5 milligrams of diphenoxylate and not
- 26 less than 25 micrograms of atropine sulfate per dosage unit.
- 5. Not more than 100 milligrams of opium per 100 milliliters
- 28 or per 100 grams, or not more than 5 milligrams per dosage unit.
- 29 <u>6. Any detectable quantity of ephedrine, its salts or</u>
- 30 optical isomers, or salts of optical isomers, except for

- 1 pediatric products in liquid form that:
- 2 (A) are labeled pursuant to Federal regulation primarily
- 3 intended for administration to children under 12 years of age
- 4 according to label instructions; and
- 5 (B) <u>according to label instructions do not exceed 15</u>
- 6 <u>milligrams of ephedrine per 5 milliliters of liquid product.</u>
- 7. Any detectable quantity of psuedoephedrine, its salts or
- 8 optical isomers, or salts of optical isomers, except for
- 9 pediatric products in liquid form that:
- 10 (A) are labeled pursuant to Federal regulation primarily
- 11 <u>intended for administration to children under 12 years of age</u>
- 12 according to label instructions; and
- 13 (B) according to label instructions do not exceed 15
- 14 milligrams of ephedrine per 5 milliliters of liquid product.
- 15 (ii) Buprenorphine.
- 16 Section 2. Section 11(c) of the act is amended to read:
- 17 Section 11. Professional Prescription, Administration, and
- 18 Dispensing. --
- 19 * * *
- 20 (c) (1) No controlled substance in Schedule V may be
- 21 distributed or dispensed for other than a medicinal purpose.
- 22 (2) If a substance described in section 4(5)(i)6 or 7 of
- 23 this act is dispensed, sold or distributed in a pharmacy:
- 24 (i) The substance shall be dispensed, sold or distributed
- 25 only by a licensed pharmacist or a licensed pharmacy technician.
- 26 (ii) The substance shall not be dispensed, sold or
- 27 distributed to any person under 18 years of age.
- 28 (iii) Any person purchasing, receiving or otherwise
- 29 <u>acquiring the substance shall:</u>
- 30 (A) Produce a government-issued photo identification showing

- 1 the date of birth of the person.
- 2 (B) Sign a written log or receipt showing the date of the
- 3 transaction, the name of the person and the name and the amount
- 4 of the substance purchased, received or otherwise acquired.
- 5 (3) (i) No person shall purchase, receive or otherwise
- 6 acquire more than 7.5 grams of a substance described in section
- 7 4(5)(i)6 or 7 of this act within any 30-day period.
- 8 (ii) No licensed pharmacist or licensed pharmacy technician
- 9 shall dispense, sell or distribute more than 7.5 grams of the
- 10 <u>substances described in section 4(5)(i)6 or 7 of this act within</u>
- 11 any 30-day period.
- 12 (iii) The limits described in subclauses (i) and (ii) shall
- 13 not apply to any quantity of the substance dispensed under a
- 14 valid prescription.
- 15 (4) The secretary, upon application of a manufacturer of a
- 16 drug product, may exempt the product from section 4(5)(i)6 and 7
- 17 of this act if the secretary determines that the product has
- 18 been formulated in such a way as to effectively prevent the
- 19 conversion of the active ingredient into methamphetamine.
- 20 * * *
- 21 Section 3. Section 13(a) of the act is amended by adding
- 22 clauses and the section is amended by adding a subsection to
- 23 read:
- 24 Section 13. Prohibited Acts; Penalties. -- (a) The following
- 25 acts and the causing thereof within the Commonwealth are hereby
- 26 prohibited:
- 27 * * *
- 28 (39) The knowing or intentional dispensing, sale or
- 29 distribution of a substance in violation of section 11(c).
- 30 (40) The knowing or intentional entry of false information

- 1 in the log required under section 11(c)(2)(iii)(B).
- 2 * * *
- 3 (q) Any person who violates subsection (a)(39) or (40) shall
- 4 be quilty of a misdemeanor of the first degree and shall, upon
- 5 conviction, be sentenced to imprisonment not exceeding five
- 6 years or to pay a fine not exceeding ten thousand (\$10,000), or
- 7 both.
- 8 Section 4. Section 13.1 of the act, added July 15, 2004
- 9 (P.L.729, No.84), is amended to read:
- 10 Section 13.1. Liquefied Ammonia Gas; Precursors and
- 11 Chemicals.--(a) The following acts are prohibited:
- 12 (1) Possessing or transporting liquefied ammonia gas:
- (i) for any purpose other than legitimate agricultural or
- 14 industrial use; or
- 15 (ii) in a container not approved by the Department of
- 16 Agriculture or the Department of Transportation or both.
- 17 (2) Possessing or transporting liquefied ammonia gas with
- 18 intent to manufacture a controlled substance.
- 19 (3) Possessing red phosphorous, hypophosphoric acid,
- 20 ammonium sulfate, phosphorous, iodine, hydriodic acid,
- 21 ephedrine, pseudoephedrine, lithium, sodium, potassium,
- 22 sassafras oil, safrole oil or other oil containing safrole or
- 23 equivalent, whether in powder or liquid form, with intent to
- 24 manufacture a controlled substance.
- 25 (4) Possessing or transporting a substance containing any
- 26 <u>detectable quantity of ephedrine, its salts or optical isomers,</u>
- 27 or salts of optical isomers, or any detectable quantity of
- 28 pseudoephedrine, its salts or optical isomers, or salts of
- 29 optical of optical isomers, in a vehicle with knowledge or
- 30 intent it will be used to manufacture a controlled substance.

- 1 (b) A person who violates subsection (a)(1) commits a
- 2 misdemeanor and upon conviction shall be sentenced to
- 3 imprisonment not exceeding five years and to pay a fine not
- 4 exceeding ten thousand dollars (\$10,000).
- 5 (c) A person who violates subsection (a)(2) or (3) commits a
- 6 felony and upon conviction shall be sentenced to imprisonment
- 7 not exceeding seven years and to pay a fine not exceeding
- 8 fifteen thousand dollars (\$15,000).
- 9 (d) A person who violates subsection (a)(4) commits a felony
- 10 of the second degree and shall be sentenced to a term of
- 11 imprisonment not exceeding ten years or a fine not exceeding
- 12 <u>twenty-five thousand (\$25,000), or both.</u>
- 13 Section 5. Section 41.1 of the act, added December 4, 1980
- 14 (P.L.1093, No.186), is amended to read:
- 15 Section 41.1. Effect on Local Ordinances.--(a) Nothing in
- 16 this act relating to drug paraphernalia shall be deemed to
- 17 supersede or invalidate any consistent local ordinance,
- 18 including zoning and nuisance ordinances, relating to the
- 19 possession, sale or use of drug paraphernalia.
- 20 (b) The provisions of section 11 of this act shall preempt
- 21 any inconsistent local ordinance, including zoning and nuisance
- 22 ordinances, relating to the disbursement, sale or distribution
- 23 of ephedrine or pseudoephedrine.
- 24 Section 6. This act shall take effect in 60 days.