

109TH CONGRESS
1ST SESSION

H. R. 1378

To amend the Controlled Substances Act with respect to the regulation of ephedrine alkaloids, including ephedrine and pseudoephedrine, and for other purposes.

IN THE HOUSE OF REPRESENTATIVES

MARCH 17, 2005

Mrs. EMERSON (for herself and Mr. BERRY) introduced the following bill; which was referred to the Committee on Energy and Commerce, and in addition to the Committee on the Judiciary, for a period to be subsequently determined by the Speaker, in each case for consideration of such provisions as fall within the jurisdiction of the committee concerned

A BILL

To amend the Controlled Substances Act with respect to the regulation of ephedrine alkaloids, including ephedrine and pseudoephedrine, and for other purposes.

1 *Be it enacted by the Senate and House of Representa-*
2 *tives of the United States of America in Congress assembled,*

3 **SECTION 1. SHORT TITLE.**

4 This Act may be cited as the “Ephedrine Alkaloids
5 Regulation Act of 2005”.

6 **SEC. 2. FINDINGS.**

7 The Congress finds as follows:

1 (1) The United States faces increasing danger
2 related to methamphetamine trafficking, production,
3 and abuse.

4 (2) Methamphetamine is a highly addictive drug
5 that can be readily made from products and precursors
6 purchased from retail stores. Step-by-step recipes
7 can easily be found on the Internet, which is a
8 factor in the dramatic increase in the number of
9 clandestine labs in recent years.

10 (3) Methamphetamine-producing clandestine
11 laboratories have been identified by the Drug Enforcement
12 Administration as a significant threat to
13 the Nation's public health and safety. The manufacture
14 of methamphetamine produces highly toxic and
15 unstable chemicals that threaten the well-being of
16 first responders, law enforcement officers, and the
17 community at-large.

18 (4) Methamphetamine production, once exclusively
19 found in West Coast States, has rapidly
20 moved eastward to the Midwest. Production can now
21 be found on the East Coast, in the States of New
22 York and Florida.

23 (5) Methamphetamine abuse is indiscriminate
24 of age, socioeconomic level, or race.

1 (6) Pseudoephedrine is a necessary precursor
2 chemical in the production of methamphetamine,
3 which prompted the Drug Enforcement Administra-
4 tion to initiate investigations regarding the chemi-
5 cal's sale and distribution.

6 (7) Efforts to reduce access to pseudoephedrine
7 by methamphetamine producers, such as blister
8 packaging and sales thresholds, have not been effec-
9 tive deterrents, and pseudoephedrine tablets remain
10 pervasive in the illicit production of methamphet-
11 amine.

12 (8) Pseudoephedrine in liquid gel and liquid
13 forms have not been found to be used in meth-
14 amphetamine production.

15 (9) As States and communities attempt to com-
16 bat and control methamphetamine through restrict-
17 ing the sale of pseudoephedrine products, it is in-
18 cumbent upon the Congress to develop a uniform
19 standard for the distribution of pseudoephedrine in
20 tablet form.

21 **SEC. 2. CONTROLLED SUBSTANCES; ADDITION OF EPHED-**
22 **RINE ALKALOIDS AND PHENYLPROPANOLA-**
23 **MINE TO SCHEDULE V.**

24 (a) IN GENERAL.—Effective upon the expiration of
25 30 days after the date of the enactment of this Act, ephed-

1 rine alkaloids (including ephedrine and pseudoephedrine)
2 and phenylpropanolamine, and their salts, optical isomers,
3 and salts of optical isomers, whether alone or in combina-
4 tion with other substances, shall be considered to be listed
5 in schedule V of the schedules of controlled substances es-
6 tablished under section 202(c) of the Controlled Sub-
7 stances Act, subject to subsection (b) and subject to the
8 authority of the Attorney General under such Act to des-
9 ignate substances as controlled substances or listed chemi-
10 cals. The Attorney General shall amend part 1308 of title
11 21, Code of Federal Regulations, accordingly.

12 (b) CERTAIN FORMS OF PSEUDOEPHEDRINE AND
13 PHENYLPROPANOLAMINE.—Subject to the authority of
14 the Attorney General under the Controlled Substances Act
15 to designate substances as controlled substances or listed
16 chemicals—

17 (1) subsection (a) does not apply to
18 pseudoephedrine or phenylpropanolamine when con-
19 tained in a drug that is in liquid or gel form and
20 is marketed or distributed lawfully in the United
21 States under the Federal Food, Drug, and Cosmetic
22 Act; and

23 (2) pseudoephedrine or phenylpropanolamine
24 when so contained shall be considered a listed chemi-
25 cal.

1 **SEC. 3. REGULATION OF TRANSACTIONS INVOLVING LIST-**
2 **ED CHEMICALS; EXEMPTION FOR CERTAIN**
3 **DOSAGE FORMS AND QUANTITIES OF**
4 **PSEUDOEPHEDRINE OR PHENYLPROPANOLA-**
5 **MINE.**

6 (a) DEFINITION OF REGULATED TRANSACTION.—
7 Section 102(39)(A)(iv) of the Controlled Substances Act
8 (21 U.S.C. 802(39)(A)(iv)) is amended—

9 (1) in the matter preceding subclause (I), by
10 striking “unless—” and inserting “unless, subject to
11 clause (v)—”; and

12 (2) in subclause (II)—

13 (A) by inserting “in liquid or gel form”
14 after “containing pseudoephedrine or phenyl-
15 propanolamine products”; and

16 (B) by striking “shall be 9 grams of
17 pseudoephedrine or 9 grams of phenyl-
18 propanolamine” and inserting “shall be any
19 quantity in excess of 9.0 grams of
20 pseudoephedrine or 9.0 grams of phenyl-
21 propanolamine”.

22 (b) DEFINITION OF ORDINARY OVER-THE-COUNTER
23 PSEUDOEPHEDRINE OR PHENYLPROPANOLAMINE PROD-
24 UCT.—Section 102 of the Controlled Substances Act (21
25 U.S.C. 802) is amended—

1 (1) in paragraph (39)(A)(iv)(I)(aa), by striking
2 “, except that” and all that follows through
3 “1996”); and

4 (2)(A) by striking paragraph (45); and

5 (B) by redesignating paragraph (46) as para-
6 graph (45).

7 (c) CHEMICAL MIXTURES NOT EASILY USED IN IL-
8 LICIT PRODUCTION.—Section 102(39)(A)(v) of the Con-
9 trolled Substances Act (21 U.S.C. 802(39)(A)(v)) is
10 amended by inserting after “chemical mixture” the fol-
11 lowing: “(including a mixture that may be marketed or
12 distributed lawfully in the United States under the Fed-
13 eral Food, Drug, and Cosmetic Act)”.

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