1	HOUSE BILL 825
2	47TH LEGISLATURE - STATE OF NEW MEXICO - FIRST SESSION, 2005
3	INTRODUCED BY
4	John A. Heaton
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10	AN ACT
11	RELATING TO CONTROLLED SUBSTANCES; INCLUDING PSEUDOEPHEDRINE AS
12	A CONTROLLED SUBSTANCE; PREVENTING THE USE OF PSEUDOEPHEDRINE
13	FOR CONVERSION INTO METHAMPHETAMINE.
14	
15	BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF NEW MEXICO:
16	Section 1. Section 30-31-3 NMSA 1978 (being Laws 1972,
17	Chapter 84, Section 3, as amended) is amended to read:
18	"30-31-3. DUTY TO ADMINISTER
19	A. The board shall administer the Controlled
20	Substances Act and may add by regulation substances to the list
21	of substances enumerated in Schedules I through IV pursuant to
22	the procedures of the Uniform Licensing Act. In determining
23	whether a substance has the potential for abuse, the board
24	shall consider the following:
25	(1) the actual or relative abuse of the
	. 155349. 1

1 substance; the scientific evidence of the 2 (2)3 pharmacological effect of the substance, if known; the state of current scientific knowledge 4 (3) regarding the substance; 5 the history and current pattern of abuse; 6 (4) 7 (5) the scope, duration and significance of abuse: 8 9 (6) the risk to the public health; and 10 (7) the potential of the substance to produce 11 psychic or physiological dependence liability. 12 **B**. After considering the factors enumerated in 13 Subsection A of this section, the board shall make findings and 14 issue regulations controlling the substance if it finds the substance has a potential for abuse. 15 16 C. If any substance is designated as a controlled 17 substance under federal law and notice is given to the board, 18 the board may, by regulation, similarly control the substance 19 under the Controlled Substances Act after providing for a 20 hearing pursuant to the Uniform Licensing Act. 21 Authority to control under this section does not D. 22 extend to distilled spirits, wine, malt beverages, tobacco or 23 pesticides as defined in the Pesticide Control Act. 24 [E. The board shall exclude any nonnarcotic 25 substance from a schedule if such substance may, under Section . 155349. 1

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61-11-22 NMSA 1978, be lawfully sold over the counter without a prescription.]"

Section 2. Section 30-31-10 NMSA 1978 (being Laws 1972, Chapter 84, Section 10) is amended to read:

"30-31-10. SCHEDULE V. --

<u>A.</u> The following controlled substances are included in Schedule V:

[A-] (1) any compound, mixture or preparation [containing] that contains the following limited quantities of any of the following narcotic drugs [which] and that also contains one or more nonnarcotic active medicinal ingredients in sufficient proportion to confer upon the compound, mixture or preparation valuable medicinal qualities other than those possessed by the narcotic drug alone:

[<del>(1)</del>] <u>(a)</u> not more than two hundred milligrams of codeine, or any of its salts, per one hundred milliliters or per one hundred grams;

[<del>(2)</del>] <u>(b)</u> not more than one hundred milligrams of dihydrocodeine, or any of its salts, per one hundred milliliters or per one hundred grams;

[<del>(3)</del>] <u>(c)</u> not more than one hundred milligrams of ethylmorphine, or any of its salts, per one hundred milliliters or per one hundred grams;

[<del>(4)</del>] <u>(d)</u> not more than two and fivetenths milligrams of diphenoxylate and not less than twenty-.155349.1

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1	five micrograms of atropine sulfate per dosage unit; or
2	[ <del>(5)</del> ] <u>(e)</u> not more than one hundred
3	milligrams of opium per one hundred milliliters or per one
4	hundred grams; <u>and</u>
5	(2) any compound, mixture or preparation that
6	<u>contains any detectable quantity of pseudoephedrine, its salts</u>
7	<u>or its optical isomers, or salts of its optical isomers. A</u>
8	compound, mixture or preparation as specified in this
9	subsection shall:
10	<u>(a) be dispensed, sold or distributed</u>
11	<u>only by a licensed pharmacist;</u>
12	<u>(b) require that a person purchasing,</u>
13	receiving or otherwise acquiring the compound, mixture or
14	preparation shall produce a photo identification showing the
15	<u>date of birth of the person and shall sign a written log or</u>
16	receipt showing the date of the transaction, the name of the
17	person and the amount of the compound, mixture or preparation;
18	and
19	(c) be limited to no more than nine
20	grams of any product, mixture or preparation within a
21	thirty-day period; provided, however, that this limit shall not
22	apply to any quantity of such product, mixture or preparation
23	dispensed pursuant to a valid prescription.
24	B. The board may by regulation exempt any compound,
25	mixture or preparation containing any depressant or stimulant
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1 substance enumerated in Schedules III, IV or V from the 2 application of the Controlled Substances Act if: the compound, mixture or preparation 3 (1) 4 contains one or more active medicinal ingredients not having a depressant or stimulant effect on the central nervous system; 5 6 and 7 (2)such ingredients are included in such 8 combinations, quantity, proportion or concentration as to 9 vitiate the potential for abuse of the substances [which] that 10 do have a depressant or stimulant effect on the nervous system. 11 C. The board may, by rule, remove a product 12 containing pseudoephedrine from Schedule V if the board 13 determines that the product is formulated as to effectively 14 prevent the conversion of pseudoephedrine into 15 methamphetamine." 16 - 5 -17 18 19 20 21 22 23 24 25 . 155349. 1

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