H.B. NO. 2592

A BILL FOR AN ACT

RELATING TO CONTROLLED SUBSTANCES.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF HAWAII:

1	SECTION 1. Section 329-14, Hawaii Revised Statutes, is
2	amended by amending subsection (d) to read as follows:
3	"(d) Any material, compound, mixture, or preparation that
4	contains any quantity of the following hallucinogenic
5	substances, their salts, isomers, and salts of isomers, unless
6	specifically excepted, whenever the existence of these salts,
7	isomers, and salts of isomers is possible within the specific
8	chemical designation:
9	(1) Alpha-ethyltryptamine (AET);
10	(2) 2,5-dimethoxy-4-ethylamphetamine (DOET);
11	(3) 2,5-dimethoxyamphetamine (2,5-DMA);
12	(4) 3,4-methylenedioxy amphetamine;
13	(5) 3,4-methylenedioxymethamphetamine (MDMA);
14	(6) N-hydroxy-3,4-methylenedioxyamphetamine (N-hydroxy-
15	MDA);
16	(7) 3,4-methylenedioxy-N-ethylamphetamine (MDE);
17	(8) 5-methoxy-3,4-methylenedioxy-amphetamine;
18	(9) 4-bromo-2,5-dimethoxy-amphetamine(4-bromo-2,5-DMA);

1	(10)	4-Bromo-2,5-dimethoxyphenethylamine (Nexus);
2	(11)	3,4,5-trimethoxy amphetamine;
3	(12)	Bufotenine;
4	(13)	4-methoxyamphetamine (PMA);
5	(14)	Diethyltryptamine;
6	(15)	Dimethyltryptamine;
7	(16)	4-methyl-2,5-dimethoxy-amphetamine;
8	(17)	Gamma hydroxybutyrate (GHB) (some other names include
9		gamma hydroxybutyric acid; 4-hydroxybutyrate; 4-
10		hydroxybutanoic acid; sodium oxybate; sodium
11		oxybutyrate);
12	(18)	Ibogaine;
13	(19)	Lysergic acid diethylamide;
14	(20)	Marijuana;
15	(21)	Parahexyl;
16	(22)	Mescaline;
17	(23)	Peyote;
18	(24)	N-ethyl-3-piperidyl benzilate;
19	(25)	N-methyl-3-piperidyl benzilate;
20	(26)	Psilocybin;
21	(27)	Psilocyn;
22	(28)	1-[1-(2-Thienyl) cyclohexyl] Pyrrolidine (TCPy);

1	(29)	Tetrahydrocannabinols;
2	(30)	Ethylamine analog of phencyclidine (PCE);
3	(31)	Pyrrolidine analog of phencyclidine (PCPy, PHP);
4	(32)	Thiophene analog of phencyclidine (TPCP; TCP);
5	(33)	Gamma-butyrolactone, including butyrolactone;
6		butyrolactone gamma; 4-butyrolactone; 2(3H)-furanone
7		dihydro; dihydro-2(3H)-furanone; tetrahydro-2-
8		furanone; 1,2-butanolide; 1,4-butanolide; 4-
9		butanolide; gamma-hydroxybutyric acid lactone; 3-
10		hydroxybutyric acid lactone and 4-hydroxybutanoic acid
11		lactone with Chemical Abstract Service number 96-48-0
12		when any such substance is intended for human
13		ingestion;
14	(34)	1,4 butanediol, including butanediol; butane-1,4-diol;
15		1,4- butylenes glycol; butylene glycol; 1,4-
16		dihydroxybutane; 1,4- tetramethylene glycol;
1 7		tetramethylene glycol; tetramethylene 1,4- diol with
18		Chemical Abstract Service number 110-63-4 when any
19		such substance is intended for human ingestion;
20	(35)	2,5-dimethoxy-4-(n)-propylthiophenethylamine (2C-T-7),
21		its optical isomers, salts, and salts of isomers;

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1	(36)	N-benzylpiperazine (BZP; 1-benzylpiperazine) its
2		optical isomers, salts, and salts of isomers;
3	(37)	1-(3-trifluoromethylphenyl)piperazine (TFMPP), its
4		optical isomers, salts, and salts of isomers;
5	(38)	Alpha-methyltryptamine (AMT), its isomers, salts, and
6		salts of isomers; [and]
7	(39)	5-methoxy-N,N-diisopropyltryptamine (5-MeO-DIPT), its
8		isomers, salts, and salts of isomers $[-]_{\underline{j}}$
9	(40)	Salvia divinorum;
10	(41)	Salvinorin A; and
11	(42)	Divinorin A."
12	SECT	ION 2. Section 329-16, Hawaii Revised Statutes, is
13	amended b	y amending subsection (c) to read as follows:
14	"(c)	Any of the following opiates, including their isomers,
15	esters, e	thers, salts, and salts of isomers, whenever the
16	existence	of these isomers, esters, ethers, and salts is
17	possible	within the specific chemical designation:
18	(1)	Alfentanil;
19	(2)	Alphaprodine;
20	(3)	Anileridine;
21	(4)	Bezitramide;
	(-)	

1	(6)	Carfentanil;
2	(7)	Dihydrocodeine;
3	(8)	Diphenoxylate;
4	(9)	Fentanyl;
5	(10)	Isomethadone;
6	(11)	Levo-alphacetylmethadol (LAAM);
7	(12)	Levomethorphan;
8	(13)	Levorphanol;
9	(14)	Metazocine;
10	(15)	Methadone;
11	(16)	Methadone-Intermediate, 4-cyano-2-dimethylamino-4, 4-
12		dphenyl butane;
13	(17)	Moramide-Intermediate, 2-methyl-3-morpholino-1, 1-
14		diphenyl-propane-carboxylic acid;
15	(18)	Pethidine (Meperidine);
16	(19)	Pethidine-Intermediate-A, 4-cyano-1-methyl-4-
17		phenylpiperidine;
18	(20)	Pethidine-Intermediate-B, ethyl-4-phenylpiperidine-4-
19		carboxylate;
20	(21)	Pethidine-Intermediate-C, 1-methyl-4-phenylpiperidine-
21		4-carboxylic acid;
22	(22)	Phenazocine;

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- 1 (23) Piminodine;
- 2 (24) Racemethorphan;
- 3 (25) Racemorphan;

4 (26) Remifentanil; [and]

- 5 (27) Sufentanil [-]; and
- 6 (28) Tapentadol."

7 SECTION 3. Section 329-20, Hawaii Revised Statutes, is
8 amended by amending subsection (b) to read as follows:

9 "(b) Depressants. Any material, compound, mixture, or preparation which contains any quantity of the following 10 substances, including its salts, isomers, esters, ethers, and 11 12 salts of isomers, whenever the existence of these isomers, esters, ethers, and salts is possible within the specific 13 chemical designation, that has a degree of danger or probable 14 danger associated with a depressant effect on the central 15 16 nervous system:

- 17 (1) Alprazolam;
- 18 (2) Barbital;
- 19 (3) Bromazepam;
- 20 (4) Butorphanol;
- 21 (5) Camazepam;
- 22 (6) Carisoprodol;

1	(7)	Chloral betaine;
2	(8)	Chloral hydrate;
3	(9)	Chlordiazepoxide;
4	(10)	Clobazam;
5	(11)	Clonazepam;
6	(12)	Clorazepate;
7	(13)	Clotiazepam;
8	(14)	Cloxazolam;
9	(15)	Delorazepam;
10	(16)	Dichloralphenazone (Midrin);
11	(17)	Diazepam;
12	(18)	Estazolam;
13	(19)	Ethchlorvynol;
14	(20)	Ethinamate;
15	(21)	Ethyl loflazepate;
16	(22)	Fludiazepam;
17	(23)	Flunitrazepam;
18	(24)	Flurazepam;
19	(25)	Fospropofol (Lusedra);
20	[(25)] <u>(26)</u> Halazepam;
21	[(26)] <u>(27)</u> Haloxazolam;
22	[(27)] <u>(28)</u> Ketazolam;

1	[(28)] (29)	Loprazolam;
2	 [-(29)] (30)	Lorazepam;
3		Lormetazepam;
	[(30)] <u>(31)</u>	_
4	[(31)] <u>(32)</u>	Mebutamate;
5	[(32)] <u>(33)</u>	Medazepam;
6	[(33)] <u>(34)</u>	Meprobamate;
7	[(34)] <u>(35)</u>	Methohexital;
8	[(35)] <u>(36)</u>	Methylphenobarbital (mephorbarbital);
9	[(36)] <u>(37)</u>	Midazolam;
10	[-(37)] <u>(38)</u>	Nimetazepam;
11	[(38)] <u>(39)</u>	Nitrazepam;
12	[(39)] <u>(40)</u>	Nordiazepam;
13	[-(40)] <u>(41)</u>	Oxazepam;
14	[(41)] <u>(42)</u>	Oxazolam;
15	[-(42)] <u>(43)</u>	Paraldehyde;
16	[(43)] <u>(44)</u>	Petrichloral;
17	[(44)] <u>(45)</u>	Phenobarbital;
18	[(45)] <u>(46)</u>	Pinazepam;
19	[(46)] <u>(47)</u>	Prazepam;
20	[(47)] <u>(48)</u>	Quazepam;
21	[-(48)] <u>(49)</u>	Temazepam;
22	[(49)] <u>(50)</u>	Tetrazepam;

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1	[(50)] <u>(51)</u> Triazolam;
2	[(51)] <u>(52)</u> Zaleplon;
3	[(52)] <u>(53)</u> Zolpidem; and
4	[(53)] <u>(54)</u> Zopiclone (Lunesta)."
5	SECTION 4. Section 329-22, Hawaii Revised Statutes, is
6	amended by amending subsection (d) to read as follows:
7	"(d) Depressants. Unless specifically exempted or excluded
8	or unless listed in another schedule, any material, compound,
9	mixture, or preparation that contains any quantity of the
10	following substances having a depressant effect on the central
11	nervous system, including its salts, isomers, and salts of
12	isomers:
13	(1) Lacosamide [(R)-2-acetoamido-N-benzyl-3-methoxy-
14	propionamide], (Vimpat); and
15	[(1)] <u>(2)</u> Pregabalin [(S)-3-(aminomethyl)-5-methylhexanoic
16	
	acid]."
17	acid]." SECTION 5. Section 329-35, Hawaii Revised Statutes, is
17 18	
	SECTION 5. Section 329-35, Hawaii Revised Statutes, is
18	SECTION 5. Section 329-35, Hawaii Revised Statutes, is amended to read as follows:
18 19	SECTION 5. Section 329-35, Hawaii Revised Statutes, is amended to read as follows: "§329-35 Order to show cause. (a) [Before denying,

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1	registration should not be denied, revoked, or suspended, or why
2	the renewal should not be refused. The order to show cause
3	shall contain a statement of the basis therefor and shall call
4	upon the applicant or registrant to appear before the department
5	of public safety at a time and place not less than thirty days
6	after the date of service of the order, but in the case of a
7	denial or renewal of registration the show cause order shall be
8	served not later than thirty days before the expiration of the
9	registration. These proceedings shall be conducted in
10	accordance with chapter 91 without regard to any criminal
11	prosecution or other proceeding. Proceedings to refuse renewal
12	of registration shall not abate the existing registration which
13	shall remain in effect pending the outcome of the administrative
14	hearing.] If, upon examination of the application for
15	registration from any applicant and other information gathered
16	by the department regarding the applicant, the administrator is
17	unable to make the determinations required by the applicable
18	provisions of sections 329-32 and 329-33 and applicable rules to
19	register the applicant, the administrator shall serve upon the
20	applicant an order to show cause why the registration should not
21	be denied.

1	(b) If, upon information gathered by the department
2	regarding any registrant, the administrator determines that the
3	registration of such registrant is subject to suspension or
4	revocation pursuant to section 329-34 or applicable rules, the
5	department shall serve upon the registrant an order to show
6	cause why the registration should not be revoked or suspended.
7	(c) The order to show cause shall call upon the applicant
8	or registrant to appear before the department at a time and
9	place stated in the order, which shall not be less than thirty
10	days after the date of receipt of the order. The order to show
11	cause shall also contain a statement of the legal basis for such
12	hearing and for the denial, revocation, or suspension of
13	registration and a summary of the matters of fact and law
14	asserted.
15	(d) Upon receipt of an order to show cause, the applicant
16	or registrant must, (if the registrant or applicant desires a
17	hearing) file a request for a hearing with the department within
18	thirty days after service of the order to show cause. Failure
19	to request a hearing will result in the automatic termination of
20	the registrant's registration and in the case of a new
21	application or renewal the unprocessed application will be
22	returned to the applicant.

[(b)] (e) The department of public safety may suspend any 1 2 registration simultaneously with the institution of proceedings under section 329-34, or where renewal of registration is 3 refused, if it finds that there is an imminent danger to the 4 public health or safety which warrants this action. The 5 suspension shall continue in effect until the conclusion of the 6 proceedings, including judicial review thereof, unless sooner 7 withdrawn by the department of public safety or dissolved by a 8 court of competent jurisdiction. 9 [(c)] (f) The department of public safety may subpoena and 10 examine witnesses under oath upon all such charges as may be 11 [preferred] referred before it [, and the circuit court of the 12 circuit in which the hearing is held shall enforce by 13 appropriate order the attendance and testimony of witnesses so 14 15 subpoenaed]." SECTION 6. Section 329-64, Hawaii Revised Statutes, is 16 amended by amending subsection (a) to read as follows: 17 The requirements imposed by sections $329-62[_7]$ and 18 "(a)

19 329-63(a) [-, and 329-67] of this part shall not apply to any of
20 the following:

PSD-01(10)

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1	(1)	Any pharmacist or other authorized person who sells or
2		furnishes a substance upon the prescription of a
3		physician, dentist, podiatrist, or veterinarian;
4	(2)	Any physician, dentist, podiatrist, or veterinarian
5		who administers or furnishes a substance to patients;
6	(3)	Any manufacturer or wholesaler licensed by the State
7		who sells, transfers, or otherwise furnishes a
8		substance to a licensed pharmacy, physician, dentist,
9		podiatrist, or veterinarian;
10	(4)	Any sale, transfer, furnishing, or receipt of any drug
11		that contains pseudoephedrine or norpseudoephedrine
12		that is lawfully sold, transferred, or furnished over
13		the counter without a prescription pursuant to the
14		federal Food, Drug, and Cosmetic Act (21 United States
15		Code Sec. 301 et seq.) or regulations adopted
16		thereunder as long as it complies with the
17		requirements of sections 329-73, 329-74, and 329-75[$+$
18		and]
19	(5)	Any "dietary supplement" as defined by the federal
20		Food, Drug, and Cosmetic Act (21 United States Code
21		Sec. 301) containing ephedrine alkaloids extracted

1	from any species of Ephedra that meets all of the
2	following criteria:
3	(A) -It contains, per dosage unit or serving, not more
4	than-twenty-five milligrams-of ephedrine
5	alkaloids and its labeling does not suggest or
6	recommend a total daily-intake of more than one
7	hundred milligrams of ephedrine alkaloids;
8	(B) It contains no hydrochloride or sulfate salts of
9	ephedrine-alkaloids; and
10	(C) It is packaged with a prominent label securely
11	affixed to each package that states all of the
12	following:
13	(i) The amount in milligrams of ephedrine
14	alkaloids-in a dosage unit or serving;
15	(ii) The amount of the dictary supplement that
16	constitutes a dosage unit or serving; and
17	(iii) The maximum recommended dosage of cphedrine
18	alkaloids for a healthy adult human is not
19	more than one-hundred milligrams in a
20	twenty four hour period]."
21	SECTION 7. Section 329-101, Hawaii Revised Statutes, is

"(f) [Intentional or knowing failure] Failure to transmit 1 2 any information as required by this section (to include request by the designated state agency for data corrections) shall be a 3 misdemeanor, may incur administrative fines and shall result in 4 the immediate suspension of that pharmacy or practitioner's 5 6 ability to dispense controlled substances in the State until 7 authorized by the administrator." SECTION 8. Section 329-104, Hawaii Revised Statutes, is 8 amended to read as follows: 9 10 "§329-104 Confidentiality of information; disclosure of 11 information. (a) The information collected under this part shall not be available to the public or used for any commercial 12 purpose. Ownership of all data collected shall reside with the 13 State. 14 15 (b) Responsibility for limiting access to information in the system is vested in the administrator. Access to the 16 information collected at the central repository pursuant to this 17 part shall be confidential, and access to the information shall 18 19 be limited to [+

20 (1) Personnel] personnel of the designated state agency[;
 21 and

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1	(2)	The Drug Enforcement Administration diversion group
2		supervisor].
3	(c)	This section shall not prevent the disclosure, at the
4	discretio	n of the administrator, of investigative information to:
5	(1)	Law enforcement officers, investigative agents of
6		federal, state, or county law enforcement agencies,
7		United States attorneys, county prosecuting attorneys,
8		or the attorney general; provided that the
9		administrator has reasonable grounds to believe that
10		the disclosure of any information collected under this
11		part is in furtherance of an ongoing criminal <u>or</u>
12		regulatory investigation or prosecution;
13	(2)	Registrants authorized under chapters 448, 453, and
14		463E who are registered to administer, prescribe, or
15		dispense controlled substances; provided that the
16		information disclosed relates only to the registrant's
17		own patient;
18	(3)	Pharmacists, employed by a pharmacy registered under
19		section 329-32, who request prescription information
20		about a customer relating to a violation or possible
21		violation of this chapter; or

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1	(4) Other state-authorized governmental prescription-
2	monitoring programs.
3	Information disclosed to a registrant, pharmacist, or authorized
4	government agency under this section shall be transmitted by a
5	secure means determined by the designated agency.
6	(d) No person shall knowingly disclose or attempt to
7	disclose, or use or attempt to use, information in the system in
8	violation of this section. Any person who violates this section
9	is guilty of a class C felony.
10	[(c) The designated state agency shall purge or cause to b e
11	purged from the central repository system, no later than three
12	years after the date a patient's prescription data-are made
13	available to the designated state agency, the identification
14	number of the patient, unless the information is part of an
15	active investigation.]"
16	SECTION 9. Statutory material to be repealed is bracketed
17	and stricken. New statutory material is underscored.
18	SECTION 10. This Act shall take effect upon its approval.
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20	INTRODUCED BY:
21	BY REQUEST

'JAN 2 5 2010

Report Title: Controlled Substances

Description:

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Makes Hawaii's controlled substance laws consistent with that of federal law and clarifies sections of chapter 329 relating to controlled substances.

JUSTIFICATION SHEET

- DEPARTMENT: Public Safety
- TITLE: A BILL FOR AN ACT RELATING TO CONTROLLED SUBSTANCES.
- PURPOSE: Update chapter 329, Hawaii Revised Statutes (HRS), to include emergency scheduling as designated under section 329-11 and amendments made to the federal Controlled Substance Act; amend section 329-35 relating to the order to show cause; amend section 329-64 relating to exemptions for the requirements of precursor chemicals; and amend sections 329-101 and 329-104 relating to the Department's electronic prescription monitoring program.
- MEANS: Amend sections 329-14(d), 329-16(c), 329-20(b), 329-22(d), 329-35, 329-64(a), 329-101(f), and 329-104, Hawaii Revised Statutes.
- JUSTIFICATION: Proposed amendments to chapter 329, HRS, will accomplish the following:
 - (1) Amend Hawaii's Uniform Controlled Substances Act, chapter 329, HRS, in accordance with the requirements of section 329-11, by adding the hallucinogenic substance Salvia Divinorum and/or Salvinorin A to Schedule I as required by section 329-11(e), HRS.
 - (2) Update Hawaii's Uniform Controlled Substance Act, chapter 329, HRS, with a change made to the federal Controlled Substance Act, title 21, code of federal regulations, part 1308.12 by adding the narcotic drugs Tapentadol to Schedule II as required by section 329-11(d), HRS.
 - (3) Update Hawaii's Uniform Controlled Substance Act, chapter 329, HRS, with a change made to the federal Controlled Substance Act, title 21, code of federal regulations, part 1308.14 by adding the depressants drug Fospropofol (Lusedra) to Schedule IV as required by section 329-11(d), HRS.

- (4) Update Hawaii's Uniform Controlled Substance Act, chapter 329, HRS, with a change made to the federal Controlled Substance Act, title 21, code of federal regulations, part 1308.15 by adding the depressants drug Lacosamide [(R)-2acetoamido-N-benzyl-3-methoxypropionamide] to Schedule V as required by section 329-11(d), HRS.
- (5) Update section 329-35, HRS, to be consistent with federal language listed in the federal Controlled Substance Act, title 21, code of federal regulations, part 1301.37 relating to the "order to show cause."
- (6) Amend section 329-64, HRS, relating to exemptions to the requirements imposed by sections 329-62, 329-63(a), and 329-67 relating to the sales of precursor chemicals requiring that all individuals and entities that conduct retail sales of pseudoephedrine containing products obtain a precursor chemical permit. Section 329-64 is also amended to delete the exemption for the retail sales of dietary supplements that contain ephedrine. Ephedrine was designated as a drug to be dispensed by prescription only by Act 171, section 2, Session Laws of Hawaii 2006.
- (7) Amend section 329-101(f) to clarify the language relating to the penalty for failure to transmit controlled substance prescription data to the Department due to non-compliance by pharmacies.
- (8) Amend section 329-104(e) by deleting the requirement for the designated state agency to purge the patient identification number data on all controlled substance prescriptions after 3 years. Due to administrative, civil and regulatory investigations lasting longer than three years the department feels that it is necessary to permanently maintain the patient identification number with the rest of the data as a means of better identifying patients with the same name.

Impact on the public: This bill is intended to protect the public by updating Hawaii's controlled substance schedules as well as assist pharmacists and physicians in better serving their patients by streamlining the requirements for dispensing controlled substances.

Impact on the department and other agencies: These proposed amendments would assist the Department's Narcotics Enforcement Division in clarifying regulations of the Uniform Controlled Substances Act.

GENERAL FUND: None.

OTHER FUNDS: None.

PPBS PROGRAM DESIGATION: PSD 502.

OTHER AFFECTED AGENCIES:

Department of Health, Food and Drug Branch.

EFFECTIVE DATE: Upon approval.