A BILL FOR AN ACT

RELATING TO THE UNIFORM CONTROLLED SUBSTANCES ACT.

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

- 1 SECTION 1. Section 329-1, Hawaii Revised Statutes, is
- 2 amended as follows:
- 3 1. By adding three new definitions to be appropriately
- 4 inserted and to read:
- 5 ""Pharmacy delegate" means an individual employed by the
- 6 pharmacy and selected by the pharmacist to act as that
- 7 pharmacist's agent to whom the pharmacist has delegated the task
- 8 of accessing electronic prescription accountability system
- 9 information and for whose actions the pharmacist takes full
- 10 responsibility.
- "Practitioner delegate" means an agent or employee of a
- 12 practitioner (physician, dentist, veterinarian, advanced
- 13 practice registered nurse with prescriptive authority, or
- 14 physician assistant) to whom the practitioner has delegated the
- 15 task of accessing electronic prescription accountability system
- 16 information and for whose actions the practitioner takes full
- responsibility.



1	"Reverse distributor" means a registrant who is registered
2	under section 329-32 to receive controlled substances acquired
3	from another state certified controlled substance registrant for
4	the purpose of:
5	(1) Returning unwanted, unusable, or outdated controlled
6	substances to the manufacturer or the manufacturer's
7	agent; or
8	(2) Where necessary, processing such substances or
9	arranging for the processing of such substances for
10	disposal as authorized by the administrator."
11	2. By amending the definition of "dispense" to read:
12	""Dispense" means to deliver a controlled substance to an
13	ultimate user or research subject by or pursuant to the lawful
14	order of a practitioner, including the [prescribing,]
15	administering[7] of a practitioner's controlled substances, and
16	packaging, labeling, or compounding necessary to prepare the
17	substance for that delivery. A controlled substance is
18	dispensed when:
19	(1) It is compounded, prepared, labeled, and packaged
20	pursuant to the lawful order of a practitioner by a
21	licensed pharmacist acting in the usual course of

1		[his] the licensed pharmacist's professional practice
2		and who is either registered individually or employed
3	•	in a registered pharmacy or by a registered
4		institutional practitioner, for delivery to the
5		ultimate user;
6	(2)	It is compounded, prepared, labeled and packaged for
7		delivery to the ultimate user by a practitioner acting
8	\	in the usual course of [his] the practitioner's
9		professional practice;
10	(3)	It is prepared, labeled, and packaged pursuant to the
11		lawful order of a practitioner by a registered health
12		care professional acting as an agent of the
13		practitioner for delivery to the ultimate user by the
14		practitioner; or
15	(4)	It is prepackaged by a pharmacist for use in an
16		emergency facility for delivery to the ultimate user
17		by a licensed or registered health care professional
18		pursuant to the order of a physician."
19	3. I	By amending the definition of "locum tenens
20	practition	ner" to read:
21	""Loc	cum tenens practitioner" means a practitioner[÷

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1
         (1) Who] who is licensed in this State and [registered
 2
              under section 329-32 to administer, prescribe, or
 3
              dispense a controlled substance in the course of
 4
              professional practice, who temporarily substitutes
 5
              for another [registered] practitioner for a period not
 6
              to exceed sixty days at that other practitioner's
7
              registered place of business[; and
8
         (2) Whose Drug Enforcement Administration controlled
9
              substance registration number has not been transferred
10
              to the State of Hawaiil.
11
    Locum tenens practitioners are not eligible to receive an oral
12
    code number as designated by section [+]328-16(k)[+]."
13
                     Section 329-14, Hawaii Revised Statutes, is
         SECTION 2.
14
    amended by amending subsection (b) to read as follows:
15
         "(b) Any of the following opiates, including their
16
    isomers, esters, ethers, salts, and salts of isomers, esters,
17
    and ethers, unless specifically excepted, whenever the existence
18
    of these isomers, esters, ethers, and salts is possible within
19
    the specific chemical designation:
20
              Acetyl-alpha-methylfentanyl (N-[1-(1-methyl-2-
         (1)
21
              phenethyl) -4-piperidinyl] -N-phenylacetamide);
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```
1
          (2)
              Acetylmethadol;
2
          (3)
              Allylprodine;
3
              Alphacetylmethadol (except levo-alphacetylmethadol,
          (4)
4
               levomethadyl acetate, or LAAM);
5
          (5)
               Alphameprodine;
6
          (6)
              Alphamethadol;
              Alpha-methylfentanyl (N-[1-(alpha-methyl-beta-
7
          (7)
8
              phenyl)ethyl-4-piperidyl] propionanilide; 1-(1-methyl-
9
               2-phenylethyl)-4-(N-propanilido) piperidine);
10
          (8)
              Alpha-methylthiofentanyl (N-[1-methyl-2-(2-
11
              thienyl)ethyl-4-piperidinyl]-N-phenylpropanamide);
12
         (9)
              Benzethidine;
13
        (10)
              Betacetylmethadol;
14
        (11)
              Beta-hydroxyfentanyl (N-[1-(2-hydroxy-2-phenethyl)-4-
15
              piperidinyl] -N-phenylpropanamide);
16
        (12)
              Beta-hydroxy-3-methylfentanyl (N-[1-(2-hydroxy-2-
17
              phenethyl) -3-methyl-4-piperidinyl] -N-
18
              phenylpropanamide);
19
        (13)
              Betameprodine;
20
        (14)
              Betamethadol:
21
        (15)
              Betaprodine;
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1
         (16)
               Clonitazene;
 2
         (17)
               Dextromoramide;
 3
         (18)
               Diampromide;
 4
         (19)
               Diethylthiambutene;
 5
         (20)
               Difenoxin;
 6
         (21)
               Dimenoxadol;
 7
         (22)
               Dimepheptanol;
 8
         (23)
               Dimethylthiambutene;
 9
         (24)
               Dioxaphetyl butyrate;
10
         (25)
               Dipipanone;
11
               Ethylmethylthiambutene;
         (26)
12
               Etonitazene;
         (27)
13
         (28)
               Etoxeridine:
14
         (29)
               Furethidine;
15
         (30)
               Hydroxypethidine;
16
         (31)
               Ketobemidone;
17
         (32)
               Levomoramide;
18
         (33)
               Levophenacylmorphan;
19
               3-Methylfentanyl (N-[3-methyl-1-(2-phenylethyl)-4-
         (34)
20
               piperidyl]-N-phenylpropanamide);
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```
1
         (35)
               3-methylthiofentanyl (N-[3-methyl-1-(2-thienyl)ethyl-
 2
               4-piperidinyl]-N-phenylpropanamide);
 3
         (36)
               Morpheridine;
 4
         (37)
               MPPP (1-methyl-4-phenyl-4-propionoxypiperidine);
 5
         (38)
               Noracymethadol;
 6
         (39)
               Norlevorphanol;
7
         (40)
               Normethadone;
8
         (41)
               Norpipanone;
9
               Para-fluorofentanyl (N-(4-fluorophenyl)-N-[1-(2-
         (42)
10
               phenethyl) -4-piperidinyl] propanamide;
11
               PEPAP (1-(-2-phenethyl)-4-phenyl-4-acetoxypiperidine;
         (43)
12
         (44)
               Phenadoxone;
13
         (45)
               Phenampromide;
14
        (46)
               Phenomorphan;
15
        (47)
               Phenoperidine;
16
        (48)
               Piritramide;
17
        (49)
              Proheptazine:
18
        (50)
              Properidine;
19
        (51)
               Propiram;
20
        (52)
              Racemoramide;
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```
1
         (53)
               Thiofentanyl (N-phenyl-N-[1-(2-thienyl)ethyl-4-
 2
               piperidinyl]-propanamide);
 3
         (54)
               Tilidine;
 4
         (55)
               Trimeperidine:
 5
               N-[1-benzyl-4-piperidyl]-N-phenylpropanamide
         (56)
 6
               (benzylfentanyl), its optical isomers, salts, and
 7
               salts of isomers; [and]
 8
         (57)
              N-[1-(2-thienyl)methyl-4-piperidyl]-N-
 9
              phenylpropanamide (thenylfentanyl), its optical
10
               isomers, salts, and salts of isomers [-]; and
        (58) N-(1-phenethylpiperidin-4-yl)-N-phenylacetamide,
11
12
              (acetyl fentanyl) its optical, positional, and
13
              geometric isomers, salts, and salts of isomers."
14
         SECTION 3. Section 329-14, Hawaii Revised Statutes, is
15
    amended by amending subsection (g) to read as follows:
16
         "(g) Any of the following cannabinoids, their salts,
    isomers, and salts of isomers, unless specifically excepted,
17
18
    whenever the existence of these salts, isomers, and salts of
19
    isomers is possible within the specific chemical designation:
20
              Tetrahydrocannabinols; meaning tetrahydrocannabinols
         (1)
21
              naturally contained in a plant of the genus Cannabis
```

1		(cannabis plant), as well as synthetic equivalents of
2		the substances contained in the plant, or in the
3		resinous extractives of Cannabis, sp. or synthetic
4		substances, derivatives, and their isomers with
5		similar chemical structure and pharmacological
6		activity to those substances contained in the plant,
7		such as the following: Delta 1 cis or trans
8		tetrahydrocannabinol, and their optical isomers; Delta
9		6 cis or trans tetrahydrocannabinol, and their optical
10		isomers; and Delta 3,4 cis or trans-
11		tetrahydrocannabinol, and its optical isomers (since
12		nomenclature of these substances is not
13		internationally standardized, compounds of these
14		structures, regardless of numerical designation of
15		atomic positions, are covered);
16	(2)	Naphthoylindoles; meaning any compound containing a 3-
17		(1-naphthoyl) indole structure with substitution at the
18		nitrogen atom of the indole ring by a alkyl,
19		haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl,
20		1-(N-methyl-2-piperidinyl) methyl or 2-(4-morpholinyl)
21		ethyl group, whether or not further substituted in the

1		indole ring to any extent and whether or not
2		substituted in the naphthyl ring to any extent;
3	(3)	Naphthylmethylindoles; meaning any compound containing
4		a 1H-indol-3-yl-(1-naphthyl) methane structure with
5		substitution at the nitrogen atom of the indole ring
6		by a alkyl, haloalkyl, alkenyl, cycloalkylmethyl,
7		cycloalkylethyl, 1-(N-methyl-2-piperidinyl) methyl or
8		2-(4-morpholinyl) ethyl group whether or not further
9		substituted in the indole ring to any extent and
10		whether or not substituted in the naphthyl ring to any
11		extent;
12	(4)	Naphthoylpyrroles; meaning any compound containing a
13		3-(1-naphthoyl) pyrrole structure with substitution at
14		the nitrogen atom of the pyrrole ring by a alkyl,
15		haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl,
16		1-(N-methyl-2-piperidinyl) methyl or 2-(4-morpholinyl)
17		ethyl group whether or not further substituted in the
18		pyrrole ring to any extent, whether or not substituted
19		in the naphthyl ring to any extent;
20	(5)	Naphthylmethylindenes; meaning any compound containing
21		a naphthylideneindene structure with substitution at

1		the 3-position of the indene ring by a alkyl,
2		haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl,
3		1-(N-methyl-2-piperidinyl) methyl or 2-(4-morpholinyl)
4		ethyl group whether or not further substituted in the
5		indene ring to any extent, whether or not substituted
6		in the naphthyl ring to any extent;
7	(6)	Phenylacetylindoles; meaning any compound containing a
8		3-phenylacetylindole structure with substitution at
9		the nitrogen atom of the indole ring by a alkyl,
10		haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl,
11		1-(N-methyl-2-piperidinyl) methyl or 2-(4-morpholinyl)
12		ethyl group whether or not further substituted in the
13		indole ring to any extent, whether or not substituted
14		in the phenyl ring to any extent;
15	(7)	Cyclohexylphenols; meaning any compound containing a
16		2-(3-hydroxycyclohexyl) phenol structure with
17		substitution at the 5-position of the phenolic ring by
18		a alkyl, haloalkyl, alkenyl, cycloalkylmethyl,
19		cycloalkylethyl, 1-(N-methyl-2-piperidinyl) methyl or
20		2-(4-morpholinyl) ethyl group whether or not
21		substituted in the cyclohexyl ring to any extent:

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1
          (8)
               Benzoylindoles; meaning any compound containing a 3-
 2
               (benzoyl) indole structure with substitution at the
 3
               nitrogen atom of the indole ring by a alkyl,
 4
              haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl,
 5
               1-(N-methyl-2-piperidinyl) methyl or 2-(4-morpholinyl)
 6
               ethyl group whether or not further substituted in the
 7
               indole ring to any extent and whether or not
 8
               substituted in the phenyl ring to any extent;
 9
         (9)
              2,3-Dihydro-5-methyl-3-(4-morpholinylmethyl)
10
              pyrrolo[1,2,3-de]-1,4-benzoxazin-6-yl]-1-
11
              naphthalenylmethanone (another trade name is WIN
12
              55,212-2);
13
               (6a, 10a) -9-(hydroxymethyl) -6, 6-dimethyl-3-(2-
        (10)
14
              methyloctan-2-yl)-6a,7,10,10a-
15
              tetrahydrobenzo[c]chromen-1-ol (other trade names are:
16
              HU-210 and HU-211);
17
        (11)
              Tetramethylcyclopropanoylindoles; meaning any compound
18
              containing a 3-tetramethylcyclopropanoylindole
19
              structure with substitution at the nitrogen atom of
20
              the indole ring by an alkyl, haloalkyl, cyanoalkyl,
21
              alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-
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1
               methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl,
 2
               1-(N-methyl-2-pyrrolidinyl) methyl, 1-(N-methyl-3-
 3
               morpholinyl) methyl, or tetrahydropyranylmethyl group,
 4
               whether or not further substituted in the indole ring
 5
               to any extent and whether or not substituted in the
 6
               tetramethylcyclopropyl ring to any extent;
7
        (12)
              N-(1-adamantyl)-1-pentyl-1H-indazole-3-carboxamide,
8
               its optical, positional, and geometric isomers, salts,
9
               and salts of isomers (Other names: APINACA, AKB48);
10
              Quinolin-8-yl 1-pentyl-1H-indole-3-carboxylate, its
        (13)
11
               optical, positional, and geometric isomers, salts, and
12
               salts of isomers (Other names: PB-22; QUPIC);
13
              Quinolin-8-yl 1-(5fluoropentyl)-1H-indole-3-
        (14)
14
               carboxylate, its optical, positional, and geometric
15
               isomers, salts, and salts of isomers (Other names:
16
              fluoro-PB-22; 5F-PB-22);
17
              N-(1-amino-3-methyl-1-oxobutan-2-yl)-1-(4-methyl-1-oxobutan-2-yl)
        (15)
18
              fluorobenzyl)-1H-indazole-3-carboxamide, its optical,
19
              positional, and geometric isomers, salts, and salts of
20
               isomers (Other names: AB-FUBINACA);
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```
1
         (16)
               N-(1-amino-3,3-dimethyl-1-oxobutan-2-yl)-1-pentyl-1H-
 2
               indazole-3-carboxamide, its optical, positional, and
 3
               geometric isomers, salts, and salts of isomers (Other
 4
               names:
                       ADB-PINACA):
 5
         (17)
               N-(1-amino-3-methyl-1-oxobutan-2-yl)-1-
 6
               (cyclohexylmethyl) -1H-indazole-3-carboxamide, its
 7
               optical, positional, and geometric isomers, salts, and
 8
               salts of isomers (Other names: AB-CHMINACA);
 9
         (18)
               N-(1-amino-3-methyl-1-oxobutan-2-yl)-1-pentyl-1H-
10
               indazole-3-carboxamide, and geometric isomers, salts,
11
               and salts of isomers (Other names: AB-PINACA);
12
         (19)
               [1-(5-fluoropentyl)-1H-indazol-3-yl] (naphthalen-1-
13
               yl) methanone, and geometric isomers, salts, and salts
14
               of isomers (Other names: THJ-2201);
15
         (20)
               Methyl (1-(4-fluorobenzyl)-1 H-indazole-3-carbonyl)-L-
16
               valinate, and geometric isomers, salts, and salts of
17
               isomers (Other names: FUB-AMB):
18
         (21)
               (S)-methyl 2-(1-(5-fluoropentyl)-1H-indazole-3-
19
               carboxamido) - 3 - methylbutanoate, and geometric isomers,
20
               salts, and salts of isomers (Other names: 5-fluoro-
21
               AMB, 5-fluoro-AMP);
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1	(22)	N-((3s,5s,7s)-adamantan-1-yl)-1-(5-fluoropentyl)-1H-
2		indazole-3-carboxamide, and geometric isomers, salts,
3		and salts of isomers (Other names: AKB48 N-(5-
4		fluoropentyl) analog, 5F-AKB48, APINACA 5-fluoropentyl
5		analog, 5F-APINACA);
6	(23)	N-adamantyl-1-fluoropentylindole-3-Carboxamide, and
7		geometric isomers, salts, and salts of isomers (Other
8		names: STS-135, 5F-APICA; 5-fluoro-APICA); [and]
9	(24)	Naphthalen-1-yl 1-(5-fluoropentyl)-1H-indole-3-
10		caboxylate, and geometric isomers, salts, and salts of
11		isomers (Other names: NM2201) [-]; and
12	(25)	N-(1-amino-3,3-dimethyl-1-oxobutan-2-yl)-1-
13		(cyclohexylmethyl)-1H-indazole-3-carboxamide, and
14		geometric isomers, salts, and salts of isomers (Other
15		names: MAB-CHMINACA and ADB-CHMINACA)."
16	SECT	ION 4. Section 329-20, Hawaii Revised Statutes, is
17	amended by	y amending subsection (e) to read as follows:
18	"(e)	Other substances. Unless specifically excepted or
19	unless li	sted in another schedule, any material, compound,
20	mixture,	or preparation which contains any quantity of the



1 following substances, including its [salts: Pentazocine.] 2 optical isomers and its salts, isomers, and salts of isomers: 3 (1) Pentazocine; and 4 (2) Eluxadoline (5-[[[(2S)-2-amino-3-[4-aminocarbonyl)-5 2,6-dimethylphenyl]-1-oxopropyl][(1S)-1-(4-phenyl-1H-6 imidazol-2-yl)ethyl]amino]methyl]-2-methoxybenzoic 7 acid." SECTION 5. Section 329-23, Hawaii Revised Statutes, is 8 9 amended by amending subsection (a) to read as follows: 10 "(a) The department of public safety shall [republish] 11 make available to the public on the department's website the 12 schedules annually or more often, as may be necessary to update 13 the schedules." 14 SECTION 6. Section 329-31, Hawaii Revised Statutes, is 15 amended to read as follows: 16 "§329-31 Rules. The department of public safety may **17** [promulgate] adopt rules and charge reasonable fees relating to the registration and control of the manufacture, distribution, 18 [prescription, and] prescribing, dispensing [of], or reverse 19 20 distribution with controlled substances within this State."

1	SECT	ION 7. Section 329-32, Hawaii Revised Statutes, is
2	amended a	s follows:
3	1.	By amending subsections (a) and (b) to read:
4	" (a)	Every person who:
5	(1)	Manufactures, distributes, prescribes, [ex] dispenses,
6		or conducts reverse distribution with any controlled
7		substance within this State;
8	(2)	Proposes to engage in the manufacture, distribution,
9		prescription, [ex] dispensing, or reverse distribution
10		of any controlled substance within this State; or
11	(3)	Dispenses or proposes to dispense any controlled
12		substance for use in this State by shipping, mailing,
13		or otherwise delivering the controlled substance from
14		a location outside this State;
15	shall obt	ain a registration issued by the department of public
16	safety in	accordance with the department's rules. A licensed or
17	registere	d health care professional who acts as the authorized
18	agent of	a practitioner and who administers controlled
19	substance	s at the direction of the practitioner shall not be
20	required	to obtain a registration.

- (b) Persons registered by the department of public safety
- 2 under this chapter to manufacture, distribute, prescribe,
- 3 dispense, store, [ex] conduct research, or conduct reverse
- 4 <u>distribution</u> with controlled substances may possess,
- 5 manufacture, distribute, prescribe, dispense, store, or conduct
- 6 research with those substances to the extent authorized by their
- 7 registration and in conformity with this part."
- 8 2. By amending subsection (e) to read:
- 9 "(e) A separate registration shall be required at each
- 10 principal place of business or professional practice where the
- 11 applicant manufactures, distributes, prescribes, [ex] dispenses,
- or conducts reverse distribution with controlled substances,
- 13 except an office used by a practitioner (who is registered at
- 14 another location) where controlled substances are prescribed but
- 15 neither administered nor otherwise dispensed as a regular part
- 16 of the professional practice of the practitioner at such office,
- 17 and where no supplies of controlled substances are maintained."
- 18 SECTION 8. Section 329-33, Hawaii Revised Statutes, is
- 19 amended as follows:
- 20 1. By amending subsection (a) to read:

T	"(a)	The department of public safety shall register an
2	applicant	to manufacture, dispense, prescribe, [ex] distribute,
3	or conduc	t reverse distribution with controlled substances
4	included	in sections 329-14, 329-16, 329-18, 329-20, and 329-22
5	unless it	determines that the issuance of that registration
6	would be	inconsistent with the public interest. In determining
7	the public	c interest, the department of public safety shall
8	consider	the following factors:
9	(1)	Maintenance of effective controls against diversion of
10		controlled substances into other than legitimate
11		medical, scientific, or industrial channels;
12	(2)	Compliance with applicable state and local law;
13	(3)	Any convictions of the applicant under any federal and
14		state laws relating to any controlled substance;
15	(4)	Past experience in the manufacture or distribution of
16		controlled substances, and the existence in the
17		applicant's establishment of effective controls
18		against diversion;
19	(5)	Furnishing by the applicant of false or fraudulent
20		material in any application filed under this chapter;

1	(6)	Suspension, revocation, or surrender of the
2		applicant's federal registration to manufacture,
3		distribute, prescribe, or dispense controlled
4		substances as authorized by federal law; and
5	(7)	Any other factor relevant to and consistent with the
6		public health and safety."
7	2.	By amending subsection (c) to read:
8	"(c)	Practitioners [must] shall be registered to dispense
9	or to pre	scribe any controlled substances or to conduct research
10	with cont	rolled substances in schedules II through V if they are
11	authorize	d to dispense or to prescribe or conduct research under
12	the law o	f this State. The department of public safety need not
13	require s	eparate registration under this part for practitioners
14	engaging	in research with nonnarcotic controlled substances in
15	schedules	II through V where the registrant is already
16	registere	d under this part in another capacity. [Practitioners
17	registere	d under federal law to conduct research with schedule I
18	substance	s may conduct research with schedule I substances
19	within th	is State upon furnishing the department of public

safety evidence of that federal registration.] "

20

1	SECT	ION 9. Section 329-34, Hawaii Revised Statutes, is
2	amended b	y amending subsection (a) to read as follows:
3	"(a)	A registration under section 329-33 to manufacture,
4	distribut	e, [ex] dispense, or conduct reverse distribution with
5	a control	led substance may be suspended or revoked by the
6	departmen	t of public safety upon a finding that the registrant:
7	(1)	Has furnished false or fraudulent material information
8		in any application filed under this chapter;
9	(2)	Has been convicted of a felony or has been granted a
10		motion for the deferral of acceptance of a guilty plea
11		or a nolo contendere plea to a felony, pursuant to
12		chapter 853 and under any state or federal law
13		relating to any controlled substance;
14	(3)	Has had the registrant's federal registration
15		suspended or revoked to manufacture, distribute,
16		prescribe, [ex] dispense, or conduct reverse
17		distribution with controlled substances; or
18	(4)	Has had the registrant's state license to practice the
19		registrant's profession suspended or revoked by the
20		applicable governing state board."

1 SECTION 10. Section 329-36, Hawaii Revised Statutes, is 2 amended to read as follows: 3 "§329-36 Records of registrants. Persons registered to 4 manufacture, distribute, prescribe, [er] dispense, or conduct 5 reverse distribution with controlled substances under this 6 chapter shall keep records and maintain inventories in 7 conformance with the recordkeeping and inventory requirements of federal law and with any additional rules the department of 8 9 public safety issues." 10 SECTION 11. Section 329-37, Hawaii Revised Statutes, is 11 amended to read as follows: 12 "§329-37 Filing requirements. All persons registered to manufacture, distribute, conduct reverse distribution, or 13 14 dispense controlled substances and all persons who transport, 15 warehouse, or otherwise handle controlled substances, shall file with the department of public safety on forms and within the 16 17 time and manner prescribed by the department of public safety, copies of order, receipt and distribution of schedule I and 18 19 schedule II controlled substances and other controlled 20 substances designated by the department of public safety, 21

showing the amounts of such controlled substances ordered,

1	received, distributed, transported, warehoused, or otherwise
2	handled."
3	SECTION 12. Section 329-38, Hawaii Revised Statutes, is
4	amended by amending subsection (a) to read as follows:
5	"(a) No controlled substance in schedule II may be
6	dispensed without a written prescription of a practitioner,
7	except:
8	(1) In the case of an emergency situation, a pharmacist
9	may dispense a controlled substance listed in schedule
10	II upon receiving oral authorization from a
11	prescribing practitioner; provided that:
12	(A) The quantity prescribed and dispensed is limited
13	to the amount adequate to treat the patient
14	during the emergency period (dispensing beyond
15	the emergency period [must] shall be pursuant to
16	a written prescription signed by the prescribing
17	<pre>practitioner);</pre>
18	(B) If the prescribing practitioner is not known to
19	the pharmacist, the pharmacist shall make a
20	reasonable effort to determine that the oral
21	authorization came from a registered

1		practitioner, which may include a callback to the
2		prescribing practitioner using the phone number
3		in the telephone directory or other good faith
4		efforts to identify the prescriber; and
5	(C)	Within seven days after authorizing an emergency
6		oral prescription, the prescribing practitioner
7		shall cause a written prescription for the
8		emergency quantity prescribed to be delivered to
9		the dispensing pharmacist. In addition to
10		conforming to the requirements of this
11		subsection, the prescription shall have written
12		on its face "Authorization for Emergency
13		Dispensing". The written prescription may be
14		delivered to the pharmacist in person or by mail,
15		and if by mail, the prescription shall be
16		postmarked within the seven-day period. Upon
17		receipt, the dispensing pharmacist shall attach
18		this prescription to the oral emergency
19		prescription, which had earlier been reduced to
20		writing. The pharmacist shall notify the
21		administrator if the prescribing practitioner

1		fails to deliver a written prescription to the
2		pharmacy within the allotted time. Failure of
3		the pharmacist to do so shall void the authority
4		conferred by this paragraph to dispense without a
5		written prescription of a prescribing individual
6		practitioner. Any practitioner who fails to
7		deliver a written prescription within the seven-
8		day period shall be in violation of section 329-
9		41(a)(1);
10	(2)	No schedule II narcotic controlled substance may be
11		prescribed or dispensed for more than a thirty-day
12		supply;
13	[-(2)-]	(3) When dispensed directly by a practitioner, other
14		than a pharmacist, to the ultimate user. The
15		practitioner in dispensing a controlled substance in
16		schedule II shall affix to the package a label
17		showing:
18		(A) The date of dispensing;
19		(B) The name, strength, and quantity of the drug
20		dispensed;
21		(C) The dispensing practitioner's name and address.

1		(D) The name of the patient;
2		(E) The "use by" date for the drug, which shall be:
3		(i) The expiration date on the manufacturer's or
4		principal labeler's container; or
5		(ii) One year from the date the drug is
6		dispensed, whichever is earlier; and
7		(F) Directions for use, and cautionary statements, if
8		any, contained in the prescription or as required
9		by law.
10		A complete and accurate record of all schedule II
11		controlled substances ordered, administered,
12		prescribed, and dispensed shall be maintained for five
13		years. Prescriptions and records of dispensing shall
14		otherwise be retained in conformance with the
15		requirements of section 329-36. No prescription for a
16		controlled substance in schedule II may be refilled;
17		or
18	[(3)]	(4) In the case of an electronic prescription, a
19		pharmacist may dispense a controlled substance listed
20		in schedule II upon receiving an electronic
21		prescription."

1	SECTION 13. Section 329-49, Hawaii Revised Statutes, is
2	amended by amending subsection (a) to read as follows:
3	"(a) Any person who violates this chapter or any rule
4	adopted by the department pursuant to this chapter shall be
5	fined not more than \$10,000 for each separate offense. Any
6	action taken to collect the penalty provided for in this
7	subsection shall be considered a civil action and the fine shall
8	be deposited into the [state general fund.] controlled substance
9	registration revolving fund pursuant to section 329-59.
10	SECTION 14. Section 329-52, Hawaii Revised Statutes, is
11	amended by amending subsection (c) to read as follows:
12	"(c) For purposes of this section, "controlled premises"
13	means:
14	(1) Places where persons registered or exempted from
15	registration requirements under this chapter are
16	required to keep records; and
17	(2) Places, including factories, warehouses,
18	establishments, and conveyances in which persons
19	registered or exempted from registration requirements
20	under this chapter are permitted to hold, manufacture,
21	compound, process, sell dispense deliver conduct

1	chemical analysis, or otherwise dispose of any
2	controlled substance or regulated chemical designated
3	under section 329-61."
4	SECTION 15. Section 329-54, Hawaii Revised Statutes, is
5	amended by amending subsection (c) to read as follows:
6	"(c) A practitioner engaged in medical research is not
7	required or compelled to furnish the name or identity of a
8	research subject to the department of public safety, nor may the
9	practitioner be compelled in any state or local civil, criminal,
10	administrative, legislative, or other proceedings to furnish the
11	name or identity of any research subject that the practitioner
12	is obligated to keep confidential [-] unless the subject violates
13	section 329-41 or 329-46 or commits an offense pursuant to part
14	IV of chapter 712."
15	SECTION 16. Section 329-59, Hawaii Revised Statutes, is
16	amended by amending subsection (b) to read as follows:
17	"(b) The fund shall consist of all moneys derived from
18	fees collected pursuant to sections 329-31 and 329-67 [and],
19	legislative appropriations [-], and fines collected pursuant to
20	section 329-49. All fees collected pursuant to sections 329-31
21	and 329-67 and fines collected pursuant to section 329-49 shall

- be deposited in the controlled substance registration revolving
 fund."

 SECTION 17. Section 329-74. Hawaii Revised Statutes is
- 3 SECTION 17. Section 329-74, Hawaii Revised Statutes, is
- 4 amended by amending subsection (a) to read as follows:
- 5 "(a) A person commits the offense of unlawful transport of
- 6 pseudoephedrine if the person transports more than three
- 7 packages of any product the sale of which is restricted by
- 8 section 329-75 [without a permit issued from the department]."
- 9 SECTION 18. Section 329-101, Hawaii Revised Statutes, is
- 10 amended by amending subsection (b) to read as follows:
- 11 "(b) The designated state agency shall determine those
- 12 schedules of controlled substances, classes of controlled
- 13 substances, and specific controlled substances that are
- 14 purportedly being misused and abused in the State. As part of
- 15 the controlled substance registration process, all
- 16 practitioners, except veterinarians, and pharmacies shall be
- 17 registered with the department to utilize the electronic
- 18 prescription accountability system. No identified controlled
- 19 substances may be dispensed unless information relevant to the
- 20 dispensation of the substance is reported electronically or by
- 21 means indicated by the designated state agency to the central

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1 repository established under section 329-102, in accordance with 2 rules adopted by the department." SECTION 19. Section 329-104, Hawaii Revised Statutes, is 3 4 amended by amending subsection (c) to read as follows: 5 This section shall not prevent the disclosure, at the "(c) 6 discretion of the administrator, of investigative information 7 to: 8 (1) Law enforcement officers, investigative agents of 9 federal, state, or county law enforcement or **10** regulatory agencies, United States attorneys, county 11 prosecuting attorneys, or the attorney general; 12 provided that the administrator has reasonable grounds 13 to believe that the disclosure of any information 14 collected under this part is in furtherance of an 15 ongoing criminal or regulatory investigation or 16 prosecution; 17 (2) Registrants authorized under chapters 448, 453, and 18 463E who are registered to administer, prescribe, or 19 dispense controlled substances[+] and their 20 practitioner delegate; provided that the information

T		disclosed relates only to the registrant's own
2		patient;
3	(3)	Pharmacists[7] or pharmacist delegates, employed by a
4		pharmacy registered under section 329-32, who request
5		prescription information about a customer relating to
6		a violation or possible violation of this chapter;
7		[or]
8	(4)	Other state-authorized governmental prescription-
9		monitoring programs [-] :
10	(5)	The chief medical examiner or licensed physician
11		designee who requests information and certifies the
12		request is for the purpose of investigating the death
13		of an individual;
14	(6)	Qualified personnel for the purpose of bona fide
15		research or education; provided that data elements
16		that would reasonably identify a specific recipient,
17		prescriber, or dispenser shall be deleted or redacted
18		from the information prior to disclosure; provided
19		further that release of the information may be made
20		only pursuant to a written agreement between qualified

1	personnel and the administrator in order to ensure	
2	compliance with this subsection; and	
3	(7) Other entities or individuals authorized by the	
4	administrator to assist the program with projects that	
5	enhance the electronic prescription accountability	
6	system.	
7	Information disclosed to a registrant, pharmacist, or authorized	
8	government agency under this section shall be transmitted by a	
9	secure means determined by the designated agency."	
10	SECTION 20. Section 329-31.5, Hawaii Revised Statutes, is	
11	repealed.	
12	[" §329 31.5 Clinics. Registration as a clinic is required	
13	when an out patient medical facility maintains centralized	
14	ordering, storage, and record keeping of controlled substances	
15	to be administered and/or dispensed to patients. Registration	
16	of a clinic requires that:	
17	(1) Each location where controlled substances are stocked	
18	be registered by name, location, and designated	
19	principal practitioner or affiliated pharmacy. The	
20	principal practitioner or affiliated pharmacy shall be	
21	responsible for the accurate maintenance of records	

1		which document all controlled substances ordered,
2		received, administered, and dispensed within the
3		clinic;
4	(2)	Controlled substances stocked at a clinic under the
5		clinic State of Hawaii and Drug Enforcement
6		Administration registration numbers be administered to
7		clinic patients by licensed or registered health care
8		professionals under the supervision of the treating
9		practitioner;
10	(3)	Controlled substances stocked at a clinic under the
11		clinic State of Hawaii and Drug Enforcement
12		Administration registration numbers be dispensed to
13		clinic patients only by the treating practitioner for
14		emergency and urgent care, when a written prescription
15		would not be practical;
16	-(4)-	A centralized record signed and dated by the treating
17		practitioner which indicates the patient, controlled
18		substance, date and time of administration and/or
19		dispensing be maintained and stored with the current
20		controlled substance inventory, ordering, and receipt

1		records. These records shall be maintained for five
2		years; and
3	(5)	A clinic practitioner who individually maintains a
4		personal stock of controlled substances does so under
5		the practitioner's individual State and Drug
6		Enforcement Administration registration number These
7		controlled substances shall be kept separate from
8		clinic stock and cannot be accessed by other
9		practitioners.
10	The -	term "affiliated pharmacy" as used in this section
11	means a l	icensed pharmacy which supplies and monitors the
12	controlle	d substances stocked in a registered clinic.
13	The-	term "clinic" as used in this section means an out
14	patient m e	edical facility owned and operated by a legal entity
15	that emplo	bys individual practitioners for the treatment of
16	patients a	and which may or may not provide after hours emergency
17	or urgent	-care.
18	The t	term "principal physician" means the practitioner in a
19	clinic who	ose signature appears on the clinic's State of Hawaii
20	and Drug I	Inforcement Administration registrations, and who is
21	responsib	le for the proper maintenance, storage, and record

- 1 keeping of the controlled substances ordered and centrally
- 2 stocked in the clinic using the clinic Drug Enforcement
- 3 Administration registration number."]
- 4 SECTION 21. Section 329-73, Hawaii Revised Statutes, is
- 5 repealed.
- 6 ["[\$329-73] Pseudoephedrine permit. (a) Beginning
- 7 January 1, 2006, any person transporting by any means more than
- 8 three packages of any product the sale of which is restricted by
- 9 section 329 75 shall obtain a pseudoephedrine permit.
- 10 (b) The requirements imposed by [subsection] (a) shall not
- 11 apply to persons registered with the department under section
- 12 329-67. A pseudocphedrine permit shall be issued by the
- 13 department in a form and manner as prescribed by the department
- 14 by rule. A pseudocphedrine permit shall be valid for one year
- 15 and renewable annually."]
- 16 SECTION 22. Statutory material to be repealed is bracketed
- 17 and stricken. New statutory material is underscored.
- 18 SECTION 23. This Act shall take effect on July 1, 2112.

Report Title:

Uniform Controlled Substances Act; Electronic Prescriptions; Veterinarian

Description:

Updates the Uniform Controlled Substances Act to make it consistent with amendments in federal controlled substances law. Clarifies existing definitions to be consistent with federal controlled substance law; and adds new definitions to allow the use of "delegates" by practitioners and pharmacists to access the electronic prescription accountability system. Clarifies that individuals that conduct reverse distribution with controlled substances must register with the Department of Public Safety and follow appropriate controlled substance statutes and rules. Allows for the posting of updates to Hawaii's drug schedules on the department's website. Mandates that the collections of administrative fines be deposited into the controlled substance registration revolving fund to support the program. Deletes the requirement for a pseudoephedrine permit for transporting over 3 packages of pseudoephedrine. Requires that all practitioners, except veterinarians, and pharmacies register to utilize the electronic prescription accountability system when they obtain a controlled substance registration. Authorizes the Department of Public Safety Narcotics Enforcement Division Administrator to allow access to State, county, or federal regulatory agencies to the database when conducting joint regulatory investigations. (SB2915 HD1)

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