### A BILL FOR AN ACT

RELATING TO CONTROLLED SUBSTANCES.

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

1	SECTION	ON 1.	Section 329-16, Hawaii Revised Statutes, is
2	amended by	amen	ding subsection (b) to read as follows:
3	"(b)	Any	of the following substances, except those
4	narcotic d	rugs	listed in other schedules, whether produced
5	directly o	r ind	directly by extraction from substances of
6	vegetable	origi	n, or independently by means of chemical
7	synthesis,	or b	y combination of extraction and chemical
8	synthesis:		
9	(1)	Opium	a and opiate, and any salt, compound, derivative,
10	a .	or pr	reparation of opium or opiate, including the
11		follo	wing:
12		(A)	Raw opium;
13		(B)	Opium extracts;
14		(C)	Opium fluid;
15		(D)	Powdered opium;
16		(E)	Granulated opium;
17		(F)	Codeine;

1		(G) Ethylmorphine;
2		(H) Etorphine hydrochloride;
3		(I) Hydrocodone;
4		(J) Hydromorphone;
5		(K) Metopon;
6		(L) Morphine;
7		(M) Oxycodone;
8		(N) Oxymorphone; [and]
9		(O) Thebaine;
10		(P) Dihydroetorphine;
11		(Q) Oripavine; and
12		(R) Tincture of opium;
13	(2)	Any salt, compound, isomer, derivative, or preparation
14		thereof which is chemically equivalent or identical
15		with any of the substances referred to in paragraph
16		(1), but not including the isoquinoline alkaloids of
17		opium;
18	(3)	Opium poppy and poppy straw;
19	(4)	Coca leaves and any salt, compound, derivative, or
20		preparation of coca leaves, and any salt, compound,
21		derivative, or preparation thereof which is chemically
22		equivalent or identical with any of these substances,

1		but not including decocanized coca leaves or
2		extractions which do not contain cocaine or ecgonine;
3		cocaine or any salt or isomer thereof; and
4	(5)	Concentrate of poppy straw (the crude extract of poppy
5		straw in either liquid, solid, or powder form that
6		contains the phenanthrene alkaloids of the opium
7		poppy)."
8	SECT	ION 2. Section 329-16, Hawaii Revised Statutes, is
9	amended by	y amending subsection (e) to read as follows:
10	"(e)	Stimulants. Any material, compound, mixture, or
11	preparation	on which contains any quantity of the following
12	substances	s having a danger or probable danger associated with a
13	stimulant	effect on the central nervous system:
14	(1)	Amphetamine, its salts, optical isomers, and salts of
15		its optical isomers;
16	(2)	Any substance which contains any quantity of
17	g.	methamphetamine, including its salts, isomers, and
18		salts of isomers;
19	(3)	Phenmetrazine and its salts; [and]
20	(4)	Methylphenidate[.]; and
21	(5)	Lisdexamfetamine, its salts, isomers, and salts of its
22		isomers."

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1
         SECTION 3. Section 329-18, Hawaii Revised Statutes, is
 2
    amended by amending subsection (g) to read as follows:
 3
          "(g) Any anabolic steroid. The term "anabolic steroid"
 4
    means any drug or hormonal substance chemically and
 5
    pharmacologically related to testosterone (other than estrogens,
 6
    progestins, and corticosteroids) that promotes muscle growth, and
 7
    includes:
 8
          (1)
              Boldenone;
 9
         (2)
              Clostebol (4-Chlorotestosterone);
10
         (3)
               Dehydrochlormethyltestosterone;
11
         (4)
               Dihydrotestosterone (4-dihydrotestosterone);
12
              Drostanolone;
         (5)
13
         (6)
              Ethylestrenol;
14
               Fluoxymesterone;
         (7)
15
               Formebolone (Formyldienolone);
         (8)
16
         (9)
              Mesterolone;
17
        (10)
              Methandranone;
18
        (11)
             Methandriol;
19
        (12)
             Methandrostenolone (Methandienone);
20
        (13)
             Methenolone;
21
              Methyltestosterone;
        (14)
22
        (15)
              Mibolerone;
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1
        (16) Nandrolone;
2
             Norethandrolone;
        (17)
3
        (18) Oxandrolone;
 4
        (19)
              Oxymesterone;
5
        (20)
              Oxymetholone;
6
              Stanolone (Dihydrotestosterone);
        (21)
7
        (22)
              Stanozolol;
8
        (23)
              Testolactone;
9
        (24)
              Testosterone;
10
        (25)
              Trenbolone; [and]
11
        (26)
              3[beta], 17-dihydroxy-5a-androstane;
12
        (27)
              3[alpha], 17[beta]-dihydroxy-5a-androstane;
13
        (28)
             5[alpha]-androstan-3, 17-dione;
14
              1-androstenediol (3[beta], 17[beta]-dihydroxy-
        (29)
15
              5[alpha]-androst-1-ene);
16
        (30) 1-androstenediol (3[alpha], 17[beta]-dihydroxy-
17
              5[alpha]-androst-1-ene);
18
        (31) 4-androstenediol (3[beta], 17[beta]-dihydroxy-androst-
19
              4-ene);
20
        (32) 5-androstenediol (3[beta], 17[beta]-dihydroxy-androst-
21
              5-ene);
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1
              1-androstenedione ([5[alpha]]-androst-1-en-3, 17-
        (33)
2
              dione);
3
        (34)
              4-androstenedione (androst-4-en-3, 17-dione);
4
        (35)
             5-androstenedione (androst-5-en-3, 17-dione);
5
        (36)
              Bolasterone (7[alpha], 17[alpha]-dimethyl-17[beta]-
6
              hydroxyandrost-4-en-3-one);
7
        (37)
              Calusterone (7[beta], 17[alpha]-dimethyl-17[beta]-
8
              hydroxyandrost-4-en-3-one);
9
        (38)
              [Delta]1-dihydrotestosterone (a.k.a. '1-testosterone')
10
              (17[beta]-hydroxy-5[alpha]-androst-1-en-3-one);
11
        (39)
              Furazabol (17[alpha]-methyl-17[beta]-
12
              hydroxyandrostano[2,3-c]-furazan);
13
        (40)
              13[beta]-ethyl-17[beta]-hydroxygon-4-en-3-one;
14
        (41)
             4-hydroxytestosterone (4,17[beta]-dihydroxy-androst-4-
15
              en-3-one);
16
              4-hydroxy-19-nortestosterone (4,17[beta]-dihydroxy-
        (42)
17
              estr-4-en-3-one);
18
        (43)
              Mesterolone (1[alpha]methyl-17[beta]-hydroxy-
19
              [5[alpha]]-androstan-3-one);
20
        (44)
              Methandienone (17[alpha]-methyl-17[beta]-
21
              hydroxyandrost-1, 4-dien-3-one);
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1
        (45)
              Methandriol (17[alpha]-methyl-3[beta], 17[beta]-
2
              dihydroxyandrost-5-ene);
3
              Methenolone (1-methyl-17[beta]-hydroxy-5[alpha]-
        (46)
4
              androst-1-en-3-one);
5
              17[alpha]-methyl-3[beta], 17[beta]-dihydroxy-5a-
        (47)
6
              androstane;
7
              17[alpha]-methyl-3[alpha], 17[beta]-dihydroxy-5a-
        (48)
8
              androstane;
9
        (49)
              17[alpha]-methyl-3[beta], 17[beta]-dihydroxyandrost-4-
10
              ene;
11
        (50)
              17[alpha]-methyl-4-hydroxynandrolone (17[alpha]-
12
              methyl-4-hydroxy-17[beta]-hydroxyestr-4-en-3-one);
13
              Methyldienolone (17[alpha]-methyl-17[beta]-
        (51)
14
              hydroxyestra-4, 9(10)-dien-3-one);
15
        (52)
              Methyltrienolone (17[alpha]-methyl-17[beta]-
16
              hydroxyestra-4, 9-11-trien-3-one);
17
              17[alpha]-methyl-[Delta] 1-dihydrotestosterone (17b
        (53)
18
              [beta]-hydroxy-17[alpha]-methyl-5[alpha]-androst-1-en-
19
              3-one) (a.k.a. '17-[alpha]-methyl-1-testosterone');
20
              19-nor-4-androstenediol (3[beta], 17[beta]-
        (54)
21
              dihydroxyestr-4-ene);
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1
              19-nor-4-androstenediol (3[alpha], 17[beta]-
        (55)
2
              dihvdroxyestr-4-ene);
3
        (56)
             19-nor-5-androstenediol (3[beta], 17[beta]-
              dihydroxyestr-5-ene);
4
5
              19-nor-5-androstenediol (3[alpha], 17[beta]-
        (57)
              dihydroxyestr-5-ene);
6
7
        (58)
              19-nor-4-androstenedione (estr-4-en-3, 17-dione);
8
              19-nor-5-androstenedione (estr-5-en-3, 17-dione;
        (59)
9
              Norbolethone (13[beta], 17[alpha]-diethyl-17[beta]-
        (60)
10
              hydroxygon-4-en-3-one);
              Norclostebol (4-chloro-17[beta]-hydroxyestr-4-en-3-
11
        (61)
12
              one);
              Normethandrolone (17[alpha]-methyl-17[beta]-
13
        (62)
14
              hydroxyestr-4-en-3-one);
              Stenbolone (17[beta]-hydroxy-2-methyl-[5[alpha]]-
15
        (63)
16
              androst-1-en-3-one);
              Tetrahydrogestrinone (13[beta], 17[alpha]-diethyl-
17
        (64)
18
              17[beta]-hydroxygon-4, 9, 11-trien-3-one); and
       [(26)] (65) Any salt, ester, or isomer of a drug or substance
19
20
              described or listed in this subsection, if that salt,
21
              ester, or isomer promotes muscle growth, except the term
              "anabolic steroid" does not include an anabolic steroid
22
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1	which is expressly intended for administration through
2	implants to cattle or other nonhuman species and which
3	has been approved by the Secretary of Health and Human
4	Services for nonhuman administration. If any person
5	prescribes, dispenses, or distributes an anabolic
6	steroid intended for administration to nonhuman species
7	for human use, the person shall be considered to have
8	prescribed, dispensed, or distributed an anabolic
9	steroid within the meaning of this paragraph."
10	SECTION 4. Section 329-33, Hawaii Revised Statutes, is
11	amended by amending subsection (a) to read as follows:
12	"(a) The department of public safety shall register an
13	applicant to manufacture, dispense, prescribe, or distribute
14	controlled substances included in sections 329-14, 329-16,
15	329-18, 329-20, and 329-22 unless it determines that the
16	issuance of that registration would be inconsistent with the
17	public interest. In determining the public interest, the
18	department of public safety shall consider the following
19	factors:
20	(1) Maintenance of effective controls against diversion of
21	controlled substances into other than legitimate
22	medical scientific or industrial channels:

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1	(2)	Compliance with applicable state and local law;
2	(3)	Any convictions of the applicant under any federal and
3		state laws relating to any controlled substance;
4	(4)	Past experience in the manufacture or distribution of
5		controlled substances, and the existence in the
6		applicant's establishment of effective controls
7		against diversion;
8	(5)	Furnishing by the applicant of false or fraudulent
9		material in any application filed under this chapter;
10	(6)	Suspension [or], revocation, or surrender of the
11		applicant's federal registration to manufacture,
12		distribute, prescribe, or dispense controlled
13		substances as authorized by federal law; and
14	(7)	Any other factor relevant to and consistent with the
15		<pre>public health and safety."</pre>
16	SECT	ION 5. Section 329-38, Hawaii Revised Statutes, is
17	amended by	y amending subsection (c) to read as follows:
18	"(c)	The transfer of original prescription information for
19	a control	led substance listed in schedule III, IV, or V for the
20	purpose o	f [refill] dispensing is permissible between pharmacies
21	on a one	time basis[, subject to the following requirements:]

only. However, pharmacies electronically sharing a real-time,

22

1	online da	tabas	e may transfer up to the maximum refills permitted
2	by law and	d the	prescriber's authorization. Transfers are
3	subject t	o the	following requirements:
4	(1)	The	transfer shall be communicated directly between
5		two	licensed pharmacists, and the transferring
6		phar	macist shall:
7		(A)	Write or otherwise place the word "VOID" on the
8			face of the invalidated prescription;
9		(B)	Record on the reverse of the invalidated
10			prescription the name, address, and DEA
11			registration number of the pharmacy to which it
12			was transferred and the name of the pharmacist
13			receiving the prescription information; and
14		(C)	Record the date of the transfer and the name of
15			the pharmacist transferring the information;
16	(2)	The	pharmacist receiving the transferred prescription
17		info	ermation shall[:] reduce to writing the following:
18		(A)	Write or otherwise place the word "transfer" on
19			the face of the transferred prescription;
20		(B)	Record all information required to be on a
21			prescription, including:

1	(i)	The date of issuance of original
2		prescription;
3	(ii)	The original number of refills authorized on
4		original prescription;
5	(iii)	The date of original dispensing;
6	(iv)	The number of valid refills remaining and
7		[date of last refill; dates and locations
8		of previous refills;
9	(v)	The pharmacy's name, address, DEA
10		registration number, and original
11		prescription number from which the
12		prescription information was transferred;
13		[ <del>and</del> ]
14	(vi)	The name of transferor pharmacist; and
15	(vii)	The pharmacy's name, address, and Drug
16		Enforcement Administration registration
17		number, along with the prescription number
18		from which the prescription was originally
19		<pre>filled;</pre>
20	(3) Both the	original and transferred prescription shall
21	be mainta	ined for a period of five years from the date
22	of last r	efill;

1	[-(4)	The procedure allowing the transfer of prescription
2		information for refill purposes is permissible only
3		between pharmacies located on the same island in this
4		State; and
5	[ <del>(5)</del> ]	(4) Any pharmacy electronically accessing a
6		prescription record shall satisfy all information
7		requirements of a manual mode prescription transferal.
8	Fail	ure to comply with this subsection shall void the
9	authority	of the pharmacy to transfer prescriptions or receive a
10	transferr	ed prescription to or from another pharmacy."
11	SECT	ION 6. Section 329-41, Hawaii Revised Statutes, is
12	amended b	y amending subsection (a) to read as follows:
13	<b>"</b> (a)	It is unlawful for any person:
14	(1)	Who is subject to part III to distribute, administer,
15		prescribe, or dispense a controlled substance in
16		violation of section 329-38 or rules authorized under
17		section 329-31; however, a licensed manufacturer or
18		wholesaler may sell or dispense a controlled substance
19		to a master of a transpacific ship or a person in
20		charge of a transpacific aircraft upon which no
21		physician is regularly employed, for the actual
22		medical needs of persons on board such ship or

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1		aliciale when not in port, provided schedule i of it
2		controlled substances shall be sold to the master of
3		such ship or person in charge of such aircraft only in
4		accordance with the provisions set forth in 21 Code of
5		Federal Regulations, Sections 1301, 1305, and 1307,
6		adopted pursuant to Title 21, United States Code,
7		Section 821;
8	(2)	Who is a registrant to manufacture a controlled
9		substance not authorized by the registrant's
10		registration or to distribute or dispense a controlled
11		substance not authorized by the registrant's
12		registration to another registrant or another
13		authorized person;
14	(3)	To refuse or fail to make available, keep, or furnish
15		any record, notification, order form, prescription,
16		statement, invoice, or information in patient charts
17		relating to the administration, dispensing, or
18		prescribing of controlled substances;
19	(4)	To refuse any lawful entry into any premises for any
20		inspection authorized by this chapter;
21	(5)	Knowingly to keep or maintain any store, shop,
22		warehouse, dwelling, building, vehicle, boat,

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1		arrorard, or other structure or prace for the purpose
2		of using these substances or which is used for keeping
3		or selling them in violation of this chapter or
4		chapter 712, part IV;
5	(6)	Who is a practitioner or pharmacist to dispense a
6		controlled substance to any individual not known to
7		the practitioner or pharmacist, [without first
8		obtaining proper identification and documenting, by
9		signature on a log book kept by the practitioner or
10		pharmacist, the identity of and the type of
11		identification presented by] except under the
12		following circumstances:
13		(A) When dispensing a controlled substance directly
14		to an individual, the practitioner or pharmacist
15		shall first obtain and document, in a log book or
16		an electronic database, the full name,
17		identification number, identification type, and
18		signature, whether by actual signature or by
19		electronic signature capture device, of the
20		individual obtaining the controlled substance.
21	\$0,°0,	If the individual does not have any form of
22		proper identification, the pharmacist shall

1		verify the validity of the prescription and
2		identity of the patient with the prescriber, or
3		their authorized agent, before dispensing the
4		controlled substance[.]; and
5	<u>(B)</u>	For mail order prescriptions, the practitioner or
6		pharmacist shall not be subject to subparagraph
7		(A); provided that all other requirements of
8		chapter 329 shall apply and that the practitioner
9		or pharmacist, as part of the initial
10		registration process of an individual in a mail
11		order prescription drug plan and prior to the
12		controlled substance being dispensed, shall
13		obtain all identification information, including
14		the full name, identification number,
15		identification type, signature, and a photocopy
16		of a form of proper identification of the
17		individual obtaining the controlled substance.
18		The practitioner or pharmacist shall also comply
19		with other requirements set forth by rule.
20	For	the purpose of this section, "proper
21	ider	tification" means government-issued identification
22	cont	aining the photograph, printed name,

1		identification number, and signature of the individual
2		obtaining the controlled substance;
3	(7)	Who is a practitioner to predate or pre-sign
4		prescriptions to facilitate the obtaining or attempted
5		obtaining of controlled substances; or
6	(8)	Who is a practitioner to facilitate the issuance or
7		distribution of a written prescription or to issue an
8		oral prescription for a controlled substance when not
9		physically in the State."
10	SECT	ION 7. Section 329-52, Hawaii Revised Statutes, is
11	amended t	o read as follows:
12	"§32	9-52 Administrative inspections [and warrants]. [(a)
13	<del>Issuance</del>	and execution of administrative inspection warrants
14	shall be	as follows:
15	(1)	A judge of the circuit court, or any district judge
16		within the judge's jurisdiction, and upon proper oath
17		or affirmation showing probable cause, may issue
18		warrants for the purpose of conducting administrative
19		inspections authorized by this chapter or rules
20		hereunder, and seizures of the property appropriate to
21		the inspections. For purposes of the issuance of
22		administrative inspection warrants, probable cause

1		exist	s upon showing a valid public interest in the
2		effec	tive enforcement of this chapter or rules
3		hereu	nder, sufficient to justify administrative
4		inspe	ction of the area, premises, building or
5		conve	yance in the circumstances specified in the
6		appli	cation for the warrant;
7	<del>(2)</del>	<del>A war</del>	rant shall issue only upon an affidavit of a
8		desig	nated officer or employee having knowledge of the
9		facts	alleged, sworn to before the judge and
10		estab	lishing the grounds for issuing the warrant. If
11		the j	udge is satisfied that grounds for the
12		appli	cation exist or that there is probable cause to
13		belie	ve they exist, the judge shall issue a warrant
14		ident	ifying the area, premises, building, or
15		conve	yance to be inspected, the purpose of the
16		inspe	ection, and, if appropriate, the type of property
17		to be	inspected, if any. The warrant shall:
18		<del>(A)</del>	State the grounds for its issuance and the name
19			of each person whose affidavit has been taken in
20			support thereof;
21		<del>(B)</del>	Be directed to a person authorized by section
22			329-51 to execute it;

1		(6)	Command the person to whom it is directed to
2			inspect the area, premises, building, or
3			conveyance identified for the purpose specified
4			and, if appropriate, direct the seizure of the
5			property specified;
6		<del>(D)</del>	Identify the item or types of property to be
7			seized, if any;
8		<del>(E)</del>	Direct that it be served during normal business
9			hours and designate the judge to whom it shall be
10			returned;
11	<del>(3)</del>	<del>A wa</del>	rrant issued pursuant to this section must be
12		ехее	uted and returned within ten days of its date
13		unle	ss, upon a showing of a need for additional time,
14		the-	court orders otherwise. If property is seized
15		purs	uant to a warrant, a copy shall be given to the
16		pers	on from whom or from whose premises the property
17		<del>is t</del>	aken, together with a receipt for the property
18		take	n. The return of the warrant shall be made
19		prom	ptly, accompanied by a written inventory of any
20		prop	erty taken. The inventory shall be made in the
21		pres	ence of the person executing the warrant and of
22		the-	person from whose possession or premises the

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1		propo	erty was taken, if present, or in the presence of
2		at l	east one credible person other than the person
3		ежес	uting the warrant. A copy of the inventory shall
4		<del>be</del> de	elivered to the person from whom or from whose
5		prem	ises the property was taken and to the applicant
6		for	the warrant;
7	-(4)-	The	<del>judge who has issued a warrant shall attach</del>
8		ther	eto a copy of the return and all papers returnable
9		<del>in c</del>	onnection therewith and file them with the chief
10		cler	k of the judicial circuit in which the inspection
11		was :	made.
12	<del>(b)</del>	The	department of public safety may make
13	administr	ative	inspections of controlled premises in accordance
14	with the	follo	wing provisions:
15	<del>(1)</del>	For	purposes of this section only, "controlled
. 16		prem	ises" means:
17		<del>(A)</del>	Places where persons registered or exempted from
18			registration requirements under this chapter are
19			required to keep records; and
20		<del>(B)</del>	Places including factories, warehouses,
21			establishments, and conveyances in which persons
22			registered or exempted from registration

1		requirements under this chapter are permitted to
2		hold, manufacture, compound, process, sell,
3		deliver, or otherwise dispose of any controlled
4		substance.
5	<del>(2)</del>	When authorized by an administrative inspection
6		warrant issued pursuant to subsection (a) an officer
7		or employee designated by the department of public
8		safety, upon presenting the warrant and appropriate
9		credentials to the owner, operator, or agent in
10		charge, may enter controlled premises for the purpose
11		of conducting an administrative inspection.
12	<del>(3)</del>	When authorized by an administrative inspection
13		warrant, an officer or employee designated by the
14		department of public safety may:
15		(A) Inspect and copy records required by this chapter
16		to be kept;
17		(B) Inspect, within reasonable limits and in a
18		reasonable manner, controlled premises and all
19		pertinent equipment, finished and unfinished
20		material, containers and labeling found therein,
21		and, except as provided in subsection (b) (5), all
22		other things therein, including records, files,

1			papers, processes, controls, and facilities
2			bearing on violation of this chapter; and
3		<del>(C)</del>	Inventory any stock of any controlled substance
4			therein and obtain samples thereof.
5	(4)	This	section does not prevent the inspection without a
6		warr	ant of books and records pursuant to an
7		admi	nistrative subpoena issued in accordance with law,
8		nor	does it prevent entries and administrative
9		insp	ections, including seizures of property, without a
10		warr	ant:
11		<del>(A)</del>	If the owner, operator, or agent in charge of the
12			controlled premises consents;
13		<del>(B)</del>	In situations presenting imminent danger to
14			health or safety;
15		<del>(C)</del>	In situations involving inspection of conveyances
16			if there is reasonable cause to believe that the
17			mobility of the conveyance makes it impracticable
18			to obtain a warrant;
19		<del>(D)</del>	In any other exceptional or emergency
20			circumstance where time or opportunity to apply
21			for a warrant is lacking; or

1		(E) In all other situations in which a warrant is not
2		constitutionally required.
3	<del>(5)</del>	An inspection authorized by this section shall not
4		extend to financial data, sales data, other than
5		shipment data, or pricing data unless the owner,
6		operator, or agent in charge of the controlled
7		premises consents in writing.]
8	(a) The a	administrator or any of the administrator's agents may
9	make admin	nistrative inspections of controlled premises upon
10	presenting	g appropriate credentials to the registrant or persons
11	subject to	parts III, IV, VIII, and IX of this chapter or their
12	agents in	accordance with the following provisions:
13	(1)	Inspections shall be at reasonable times and within
14	No.	reasonable limits and in a reasonable manner of
15		controlled premises and vehicles in which persons
16		registered or exempted from registration requirements
17		under this chapter are permitted to hold, manufacture,
18		compound, process, sell, dispense, deliver, or
19		otherwise dispose of any controlled substance or
20		regulated chemical designated under section 329-61 and
21		all pertinent equipment, finished and unfinished

	materials, containers, and labeling therein to
	determine if this chapter is being violated;
(2)	The administrator or any of the administrator's agents
	shall have access to and may copy any and all records,
	books, logs, or documents pertaining to the
	administering, prescribing, dispensing, or sale of
	controlled substances or regulated chemicals
	designated under this chapter without a warrant; and
(3)	The administrator or any of the administrator's agents
	may inventory any stock of any controlled substance or
	regulated chemical designated under section 329-61 and
	secure samples or specimens of any drug, device, or
	chemical not seized as evidence by paying or offering
	to pay for the sample. The administrator shall make
	or cause to be made examinations of samples secured
	under this section to determine whether or not this
	chapter is being violated.
(b)	An inspection of records authorized by this section
shall not	extend to financial data, data relating to pricing of
items, ot	her than shipment and sale amounts, unless the owner,
operator,	or agent in charge of the controlled premises consents
in writin	g <u>.</u>
	(b) shall not items, ot

1	(C)	For purposes of this section, "controlled premises"
2	means:	
3	(1)	Places where persons registered or exempted from
4		registration requirements under this chapter are
5		required to keep records; and
6	(2)	Places, including factories, warehouses,
7		establishments, and conveyances in which persons
8		registered or exempted from registration requirements
9		under this chapter are permitted to hold, manufacture,
10		compound, process, sell, dispense, deliver, or
11		otherwise dispose of any controlled substance or
12		regulated chemical designated under section 329-61."
13	SECT	ION 8. Statutory material to be repealed is bracketed
14	and stric	ken. New statutory material is underscored.
15	SECT	ION 9. This Act shall take effect upon July 1, 2050.

### Report Title:

Controlled Substances

#### Description:

Makes Hawaii's controlled substance laws consistent with that of federal law and clarifies sections of chapter 329 relating to controlled substances. Authorizes administrative inspections of the premises and records other than financial data, for establishments such as pharmacies that are authorized to dispense controlled substances. Effective 07/01/2050. (HD2)