CONTROLLED SUBSTANCE MODIFICATIONS
2011 GENERAL SESSION
STATE OF UTAH
Chief Sponsor: Gage Froerer
Senate Sponsor: Allen M. Christensen
LONG TITLE
Committee Note:
The Health and Human Services Interim Committee recommended this bill.
General Description:
This bill modifies provisions relating to the Utah Controlled Substances Act by creating
a controlled class of listed synthetic cannabinoid substances found in products often
referred to as "spice."
Highlighted Provisions:
This bill:
 expands the definition of a controlled substance to include a list of synthetic
equivalent cannabinoid substances and their analogs and homologs found in
products commonly referred to as "spice";
 clarifies that the tetrahydrocannabinols in Schedule I of the Utah Controlled
Substances Act include those both naturally and synthetically derived;
 provides that it is an affirmative defense that the person produced, possessed, or
administered any of these listed substances if the person:
 was engaged in medical research; and
 was a holder of a license to possess controlled substances for research;
 authorizes the Controlled Substances Advisory Committee to recommend
placement of a substance on a controlled substance list if it finds that the substance
has a potential for abuse and that an accepted standard has not been established for



safe use in treatment for medical purposes;
 adds "spice" to the driver license provisions regarding driving under the influence;
and
 provides that a legislative body of a political subdivision may not enact an
ordinance that is less restrictive than any provision of the Utah Controlled
Substances Act.
Money Appropriated in this Bill:
None
Other Special Clauses:
This bill provides an effective date.
Utah Code Sections Affected:
AMENDS:
41-6a-517, as last amended by Laws of Utah 2009, Chapter 390
58-37-2 (Superseded 07/01/11), as last amended by Laws of Utah 2010, Chapters 64
and 101
58-37-2 (Effective 07/01/11), as last amended by Laws of Utah 2010, Chapters 64, 101,
and 276
58-37-3, as last amended by Laws of Utah 1997, Chapter 64
58-37-4, as last amended by Laws of Utah 2010, Chapter 106
58-37-6, as last amended by Laws of Utah 2010, Chapter 287
58-37-8, as last amended by Laws of Utah 2010, Chapter 64
58-38a-203, as enacted by Laws of Utah 2010, Chapter 231
58-38a-204, as enacted by Laws of Utah 2010, Chapter 231
ENACTS:
58-37-4.2 , Utah Code Annotated 1953
Be it enacted by the Legislature of the state of Utah:
Section 1. Section 41-6a-517 is amended to read:
41-6a-517. Definitions Driving with any measurable controlled substance in the
body Penalties Arrest without warrant.
(1) As used in this section:

59	(a) "Controlled substance" [means any substance scheduled under Section 58-37-4.]
60	has the same meaning as in Section 58-37-2.
61	(b) "Practitioner" has the same meaning as [provided] in Section 58-37-2.
62	(c) "Prescribe" has the same meaning as [provided] in Section 58-37-2.
63	(d) "Prescription" has the same meaning as [provided] in Section 58-37-2.
64	(2) In cases not amounting to a violation of Section 41-6a-502, a person may not
65	operate or be in actual physical control of a motor vehicle within this state if the person has any
66	measurable controlled substance or metabolite of a controlled substance in the person's body.
67	(3) It is an affirmative defense to prosecution under this section that the controlled
68	substance was:
69	(a) involuntarily ingested by the accused;
70	(b) prescribed by a practitioner for use by the accused; or
71	(c) otherwise legally ingested.
72	(4) (a) A person convicted of a violation of Subsection (2) is guilty of a class B
73	misdemeanor.
74	(b) A person who violates this section is subject to conviction and sentencing under
75	both this section and any applicable offense under Section 58-37-8.
76	(5) A peace officer may, without a warrant, arrest a person for a violation of this
77	section when the officer has probable cause to believe the violation has occurred, although not
78	in the officer's presence, and if the officer has probable cause to believe that the violation was
79	committed by the person.
80	(6) The Driver License Division shall:
81	(a) if the person is 21 years of age or older on the date of arrest:
82	(i) suspend, for a period of 120 days, the driver license of a person convicted under
83	Subsection (2) of an offense committed on or after July 1, 2009; or
84	(ii) revoke, for a period of two years, the driver license of a person if:
85	(A) the person has a prior conviction as defined under Subsection 41-6a-501(2); and
86	(B) the current violation under Subsection (2) is committed:
87	(I) within a period of 10 years after the date of the prior violation; and
88	(II) on or after July 1, 2009;
89	(b) if the person is under 21 years of age on the date of arrest:

90	(i) suspend, until the person is 21 years of age or for a period of 120 days, the driver
91	license of a person convicted under Subsection (2) of an offense committed on or after July 1,
92	2009; or
93	(ii) revoke, until the person is 21 years of age or for a period of two years, the driver
94	license of a person if:
95	(A) the person has a prior conviction as defined under Subsection 41-6a-501(2); and
96	(B) the current violation under Subsection (2) is committed:
97	(I) within a period of 10 years after the date of the prior violation; and
98	(II) on or after July 1, 2009;
99	(c) subtract from any suspension or revocation period the number of days for which a
100	license was previously suspended under Section 53-3-223 or 53-3-231, if the previous
101	suspension was based on the same occurrence upon which the record of conviction is based;
102	and
103	(d) deny, suspend, or revoke a person's license for the denial and suspension periods in
104	effect prior to July 1, 2009, for a conviction of a violation under Subsection (2) that was
105	committed prior to July 1, 2009.
106	(7) (a) The court shall notify the Driver License Division if a person fails to:
107	(i) complete all court ordered screening and assessment, educational series, and
108	substance abuse treatment; or
109	(ii) pay all fines and fees, including fees for restitution and treatment costs.
110	(b) Upon receiving the notification, the division shall suspend the person's driving
111	privilege in accordance with Subsections 53-3-221(2) and (3).
112	(8) The court shall order supervised probation in accordance with Section 41-6a-507
113	for a person convicted under Subsection (2).
114	Section 2. Section 58-37-2 (Superseded 07/01/11) is amended to read:
115	58-37-2 (Superseded 07/01/11). Definitions.
116	(1) As used in this chapter:
117	(a) "Administer" means the direct application of a controlled substance, whether by
118	injection, inhalation, ingestion, or any other means, to the body of a patient or research subject
119	by:
120	(i) a practitioner or, in the practitioner's presence, by the practitioner's authorized agent;

121	or
122	(ii) the patient or research subject at the direction and in the presence of the
123	practitioner.
124	(b) "Agent" means an authorized person who acts on behalf of or at the direction of a
125	manufacturer, distributor, or practitioner but does not include a motor carrier, public
126	warehouseman, or employee of any of them.
127	(c) "Consumption" means ingesting or having any measurable amount of a controlled
128	substance in a person's body, but this Subsection (1)(c) does not include the metabolite of a
129	controlled substance.
130	(d) "Continuing criminal enterprise" means any individual, sole proprietorship,
131	partnership, corporation, business trust, association, or other legal entity, and any union or
132	groups of individuals associated in fact although not a legal entity, and includes illicit as well
133	as licit entities created or maintained for the purpose of engaging in conduct which constitutes
134	the commission of episodes of activity made unlawful by Title 58, Chapter 37, Utah Controlled
135	Substances Act, Chapter 37a, Utah Drug Paraphernalia Act, Chapter 37b, Imitation Controlled
136	Substances Act, Chapter 37c, Utah Controlled Substance Precursor Act, or Chapter 37d,
137	Clandestine Drug Lab Act, which episodes are not isolated, but have the same or similar
138	purposes, results, participants, victims, methods of commission, or otherwise are interrelated
139	by distinguishing characteristics. Taken together, the episodes shall demonstrate continuing
140	unlawful conduct and be related either to each other or to the enterprise.
141	(e) "Control" means to add, remove, or change the placement of a drug, substance, or
142	immediate precursor under Section 58-37-3.
143	(f) (i) "Controlled substance" means a drug or substance:
144	(A) included in Schedules I, II, III, IV, or V of Section 58-37-4;
145	(B) included in Schedules I, II, III, IV, or V of the federal Controlled Substances Act,
146	Title II, P.L. 91-513; [or]
147	(C) that is a controlled substance analog[:]; or
148	(D) a substance listed in Section 58-37-4.2.
149	(ii) "Controlled substance" does not include:

(A) distilled spirits, wine, or malt beverages, as those terms are defined in Title 32A,

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Alcoholic Beverage Control Act;

152 (B) any drug intended for lawful use in the diagnosis, cure, mitigation, treatment, or 153 prevention of disease in human or other animals, which contains ephedrine, pseudoephedrine, 154 norpseudoephedrine, or phenylpropanolamine if the drug is lawfully purchased, sold, 155 transferred, or furnished as an over-the-counter medication without prescription; or 156 (C) dietary supplements, vitamins, minerals, herbs, or other similar substances 157 including concentrates or extracts, which: 158 (I) are not otherwise regulated by law; and 159 (II) may contain naturally occurring amounts of chemical or substances listed in this 160 chapter, or in rules adopted pursuant to Title 63G, Chapter 3, Utah Administrative Rulemaking 161 Act. 162 (g) (i) "Controlled substance analog" means a substance the chemical structure of 163 which is substantially similar to the chemical structure of a controlled substance listed in 164 Schedules I and II of Section 58-37-4, or in Schedules I and II of the federal Controlled 165 Substances Act, Title II, P.L. 91-513, or listed in Section 58-37-4.2: 166 (A) which has a stimulant, depressant, or hallucinogenic effect on the central nervous 167 system substantially similar to the stimulant, depressant, or hallucinogenic effect on the central 168 nervous system of controlled substances in the schedules set forth in Subsection (1)(f), or a 169 substance listed in Section 58-37-4.2; or 170 (B) which, with respect to a particular individual, is represented or intended to have a 171 stimulant, depressant, or hallucinogenic effect on the central nervous system substantially 172 similar to the stimulant, depressant, or hallucinogenic effect on the central nervous system of 173 controlled substances in the schedules or list set forth in this Subsection (1). 174 (ii) "Controlled substance analog" does not include: 175 (A) a controlled substance currently scheduled in Schedules I through V of Section 176 58-37-4 or listed in Section 58-37-4.2; 177 (B) a substance for which there is an approved new drug application; 178 (C) a substance with respect to which an exemption is in effect for investigational use by a particular person under Section 505 of the Food, Drug, and Cosmetic Act, 21 U.S.C. 355,

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180 to the extent the conduct with respect to the substance is permitted by the exemption;

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(D) any substance to the extent not intended for human consumption before an exemption takes effect with respect to the substance;

(E) any drug intended for lawful use in the diagnosis, cure, mitigation, treatment, or prevention of disease in man or other animals, which contains ephedrine, pseudoephedrine, norpseudoephedrine, or phenylpropanolamine if the drug is lawfully purchased, sold, transferred, or furnished as an over-the-counter medication without prescription; or

- (F) dietary supplements, vitamins, minerals, herbs, or other similar substances including concentrates or extracts, which are not otherwise regulated by law, which may contain naturally occurring amounts of chemical or substances listed in this chapter, or in rules adopted pursuant to Title 63G, Chapter 3, Utah Administrative Rulemaking Act.
- (h) "Conviction" means a determination of guilt by verdict, whether jury or bench, or plea, whether guilty or no contest, for any offense proscribed by Title 58, Chapters 37, 37a, 37b, 37c, or 37d, or for any offense under the laws of the United States and any other state which, if committed in this state, would be an offense under Title 58, Chapters 37, 37a, 37b, 37c, or 37d.
 - (i) "Counterfeit substance" means:

- (i) any controlled substance or container or labeling of any controlled substance that:
- (A) without authorization bears the trademark, trade name, or other identifying mark, imprint, number, device, or any likeness of them, of a manufacturer, distributor, or dispenser other than the person or persons who in fact manufactured, distributed, or dispensed the substance which falsely purports to be a controlled substance distributed by any other manufacturer, distributor, or dispenser; and
- (B) a reasonable person would believe to be a controlled substance distributed by an authorized manufacturer, distributor, or dispenser based on the appearance of the substance as described under Subsection (1)(i)(i)(A) or the appearance of the container of that controlled substance; or
 - (ii) any substance other than under Subsection (1)(i)(i) that:
- (A) is falsely represented to be any legally or illegally manufactured controlled substance; and
 - (B) a reasonable person would believe to be a legal or illegal controlled substance.
- (j) "Deliver" or "delivery" means the actual, constructive, or attempted transfer of a controlled substance or a listed chemical, whether or not an agency relationship exists.
 - (k) "Department" means the Department of Commerce.

214	(l) "Depressant or stimulant substance" means:
215	(i) a drug which contains any quantity of barbituric acid or any of the salts of barbituric
216	acid;
217	(ii) a drug which contains any quantity of:
218	(A) amphetamine or any of its optical isomers;
219	(B) any salt of amphetamine or any salt of an optical isomer of amphetamine; or
220	(C) any substance which the Secretary of Health and Human Services or the Attorney
221	General of the United States after investigation has found and by regulation designated
222	habit-forming because of its stimulant effect on the central nervous system;
223	(iii) lysergic acid diethylamide; or
224	(iv) any drug which contains any quantity of a substance which the Secretary of Health
225	and Human Services or the Attorney General of the United States after investigation has found
226	to have, and by regulation designated as having, a potential for abuse because of its depressant
227	or stimulant effect on the central nervous system or its hallucinogenic effect.
228	(m) "Dispense" means the delivery of a controlled substance by a pharmacist to an
229	ultimate user pursuant to the lawful order or prescription of a practitioner, and includes
230	distributing to, leaving with, giving away, or disposing of that substance as well as the
231	packaging, labeling, or compounding necessary to prepare the substance for delivery.
232	(n) "Dispenser" means a pharmacist who dispenses a controlled substance.
233	(o) "Distribute" means to deliver other than by administering or dispensing a controlled
234	substance or a listed chemical.
235	(p) "Distributor" means a person who distributes controlled substances.
236	(q) "Division" means the Division of Occupational and Professional Licensing created
237	in Section 58-1-103.
238	(r) (i) "Drug" means:
239	(A) a substance recognized in the official United States Pharmacopoeia, Official
240	Homeopathic Pharmacopoeia of the United States, or Official National Formulary, or any
241	supplement to any of them, intended for use in the diagnosis, cure, mitigation, treatment, or
242	prevention of disease in humans or animals;
243	(B) a substance that is required by any applicable federal or state law or rule to be

dispensed by prescription only or is restricted to administration by practitioners only;

(C) a substance other than food intended to affect the structure or any function of the body of humans or other animals; and

- (D) substances intended for use as a component of any substance specified in Subsections (1)(r)(i)(A), (B), and $(C)[\overline{, and (D)}]$.
 - (ii) "Drug" does not include dietary supplements.
- (s) "Drug dependent person" means any individual who unlawfully and habitually uses any controlled substance to endanger the public morals, health, safety, or welfare, or who is so dependent upon the use of controlled substances as to have lost the power of self-control with reference to the individual's dependency.
 - (t) "Food" means:

- (i) any nutrient or substance of plant, mineral, or animal origin other than a drug as specified in this chapter, and normally ingested by human beings; and
- (ii) foods for special dietary uses as exist by reason of a physical, physiological, pathological, or other condition including but not limited to the conditions of disease, convalescence, pregnancy, lactation, allergy, hypersensitivity to food, underweight, and overweight; uses for supplying a particular dietary need which exist by reason of age including but not limited to the ages of infancy and childbirth, and also uses for supplementing and for fortifying the ordinary or unusual diet with any vitamin, mineral, or other dietary property for use of a food. Any particular use of a food is a special dietary use regardless of the nutritional purposes.
- (u) "Immediate precursor" means a substance which the Attorney General of the United States has found to be, and by regulation designated as being, the principal compound used or produced primarily for use in the manufacture of a controlled substance, or which is an immediate chemical intermediary used or likely to be used in the manufacture of a controlled substance, the control of which is necessary to prevent, curtail, or limit the manufacture of the controlled substance.
 - (v) "Indian" means a member of an Indian tribe.
 - (w) "Indian religion" means any religion:
- (i) the origin and interpretation of which is from within a traditional Indian culture or community; and
 - (ii) which is practiced by Indians.

(x) "Indian tribe" means any tribe, band, nation, pueblo, or other organized group or community of Indians, including any Alaska Native village, which is legally recognized as eligible for and is consistent with the special programs, services, and entitlements provided by the United States to Indians because of their status as Indians.

- (y) "Manufacture" means the production, preparation, propagation, compounding, or processing of a controlled substance, either directly or indirectly by extraction from substances of natural origin, or independently by means of chemical synthesis or by a combination of extraction and chemical synthesis.
- (z) "Manufacturer" includes any person who packages, repackages, or labels any container of any controlled substance, except pharmacists who dispense or compound prescription orders for delivery to the ultimate consumer.
- (aa) "Marijuana" means all species of the genus cannabis and all parts of the genus, whether growing or not; the seeds of it; the resin extracted from any part of the plant; and every compound, manufacture, salt, derivative, mixture, or preparation of the plant, its seeds, or resin. The term does not include the mature stalks of the plant, fiber produced from the stalks, oil or cake made from the seeds of the plant, any other compound, manufacture, salt, derivative, mixture, or preparation of the mature stalks, except the resin extracted from them, fiber, oil or cake, or the sterilized seed of the plant which is incapable of germination. Any synthetic equivalents of the substances contained in the plant cannabis sativa or any other species of the genus cannabis which are chemically indistinguishable and pharmacologically active are also included.
- (bb) "Money" means officially issued coin and currency of the United States or any foreign country.
- (cc) "Narcotic drug" means any of the following, whether produced directly or indirectly by extraction from substances of vegetable origin, or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis:
 - (i) opium, coca leaves, and opiates;

- (ii) a compound, manufacture, salt, derivative, or preparation of opium, coca leaves, or opiates;
 - (iii) opium poppy and poppy straw; or
- 306 (iv) a substance, and any compound, manufacture, salt, derivative, or preparation of the

substance, which is chemically identical with any of the substances referred to in Subsection (1)(cc)(i), (ii), or (iii), except narcotic drug does not include decocainized coca leaves or extracts of coca leaves which do not contain cocaine or ecgonine.

- (dd) "Negotiable instrument" means documents, containing an unconditional promise to pay a sum of money, which are legally transferable to another party by endorsement or delivery.
- (ee) "Opiate" means any drug or other substance having an addiction-forming or addiction-sustaining liability similar to morphine or being capable of conversion into a drug having addiction-forming or addiction-sustaining liability.
- (ff) "Opium poppy" means the plant of the species papaver somniferum L., except the seeds of the plant.
- (gg) "Person" means any corporation, association, partnership, trust, other institution or entity or one or more individuals.
- (hh) "Poppy straw" means all parts, except the seeds, of the opium poppy, after mowing.
- (ii) "Possession" or "use" means the joint or individual ownership, control, occupancy, holding, retaining, belonging, maintaining, or the application, inhalation, swallowing, injection, or consumption, as distinguished from distribution, of controlled substances and includes individual, joint, or group possession or use of controlled substances. For a person to be a possessor or user of a controlled substance, it is not required that the person be shown to have individually possessed, used, or controlled the substance, but it is sufficient if it is shown that the person jointly participated with one or more persons in the use, possession, or control of any substances with knowledge that the activity was occurring, or the controlled substance is found in a place or under circumstances indicating that the person had the ability and the intent to exercise dominion and control over it.
- (jj) "Practitioner" means a physician, dentist, naturopathic physician, veterinarian, pharmacist, scientific investigator, pharmacy, hospital, or other person licensed, registered, or otherwise permitted to distribute, dispense, conduct research with respect to, administer, or use in teaching or chemical analysis a controlled substance in the course of professional practice or research in this state.
 - (kk) "Prescribe" means to issue a prescription:

338	(i) orally or in writing; or
339	(ii) by telephone, facsimile transmission, computer, or other electronic means of
340	communication as defined by division rule.
341	(ll) "Prescription" means an order issued:
342	(i) by a licensed practitioner, in the course of that practitioner's professional practice or
343	by collaborative pharmacy practice agreement; and
344	(ii) for a controlled substance or other prescription drug or device for use by a patient
345	or an animal.
346	(mm) "Production" means the manufacture, planting, cultivation, growing, or
347	harvesting of a controlled substance.
348	(nn) "Securities" means any stocks, bonds, notes, or other evidences of debt or of
349	property.
350	(oo) "State" means the state of Utah.
351	(pp) "Ultimate user" means any person who lawfully possesses a controlled substance
352	for the person's own use, for the use of a member of the person's household, or for
353	administration to an animal owned by the person or a member of the person's household.
354	(2) If a term used in this chapter is not defined, the definition and terms of Title 76,
355	Utah Criminal Code, shall apply.
356	Section 3. Section 58-37-2 (Effective 07/01/11) is amended to read:
357	58-37-2 (Effective 07/01/11). Definitions.
358	(1) As used in this chapter:
359	(a) "Administer" means the direct application of a controlled substance, whether by
360	injection, inhalation, ingestion, or any other means, to the body of a patient or research subject
361	by:
362	(i) a practitioner or, in the practitioner's presence, by the practitioner's authorized agent
363	or
364	(ii) the patient or research subject at the direction and in the presence of the
365	practitioner.
366	(b) "Agent" means an authorized person who acts on behalf of or at the direction of a
367	manufacturer, distributor, or practitioner but does not include a motor carrier, public
368	warehouseman, or employee of any of them.

(c) "Consumption" means ingesting or having any measurable amount of a controlled substance in a person's body, but this Subsection (1)(c) does not include the metabolite of a controlled substance.(d) "Continuing criminal enterprise" means any individual, sole proprietorship,

- partnership, corporation, business trust, association, or other legal entity, and any union or groups of individuals associated in fact although not a legal entity, and includes illicit as well as licit entities created or maintained for the purpose of engaging in conduct which constitutes the commission of episodes of activity made unlawful by Title 58, Chapter 37, Utah Controlled Substances Act, Chapter 37a, Utah Drug Paraphernalia Act, Chapter 37b, Imitation Controlled Substances Act, Chapter 37c, Utah Controlled Substance Precursor Act, or Chapter 37d, Clandestine Drug Lab Act, which episodes are not isolated, but have the same or similar purposes, results, participants, victims, methods of commission, or otherwise are interrelated by distinguishing characteristics. Taken together, the episodes shall demonstrate continuing unlawful conduct and be related either to each other or to the enterprise.
- (e) "Control" means to add, remove, or change the placement of a drug, substance, or immediate precursor under Section 58-37-3.
 - (f) (i) "Controlled substance" means a drug or substance:
 - (A) included in Schedules I, II, III, IV, or V of Section 58-37-4;
- 387 (B) included in Schedules I, II, III, IV, or V of the federal Controlled Substances Act, 388 Title II, P.L. 91-513; [or]
 - (C) [that is] a controlled substance analog[-]; or
 - (D) a substance listed in Section 58-37-4.2.

- (ii) "Controlled substance" does not include:
- (A) distilled spirits, wine, or malt beverages, as those terms are defined in Title 32B, Alcoholic Beverage Control Act;
 - (B) any drug intended for lawful use in the diagnosis, cure, mitigation, treatment, or prevention of disease in human or other animals, which contains ephedrine, pseudoephedrine, norpseudoephedrine, or phenylpropanolamine if the drug is lawfully purchased, sold, transferred, or furnished as an over-the-counter medication without prescription; or
- 398 (C) dietary supplements, vitamins, minerals, herbs, or other similar substances 399 including concentrates or extracts, which:

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400	(I) are not otherwise regulated by law; and
401	(II) may contain naturally occurring amounts of chemical or substances listed in this
402	chapter, or in rules adopted pursuant to Title 63G, Chapter 3, Utah Administrative Rulemaking
403	Act.
404	(g) (i) "Controlled substance analog" means a substance the chemical structure of
405	which is substantially similar to the chemical structure of a controlled substance listed in
406	Schedules I and II of Section 58-37-4, <u>a substance listed in Section 58-37-4.2</u> , or in Schedules
407	I and II of the federal Controlled Substances Act, Title II, P.L. 91-513:
408	(A) which has a stimulant, depressant, or hallucinogenic effect on the central nervous
409	system substantially similar to the stimulant, depressant, or hallucinogenic effect on the central
410	nervous system of controlled substances in the schedules set forth in Subsection (1)(f), a
411	substance listed in Section 58-37-4.2; or
412	(B) which, with respect to a particular individual, is represented or intended to have a
413	stimulant, depressant, or hallucinogenic effect on the central nervous system substantially
414	similar to the stimulant, depressant, or hallucinogenic effect on the central nervous system of
415	controlled substances in the schedules or list set forth in this Subsection (1).
416	(ii) "Controlled substance analog" does not include:
417	(A) a controlled substance currently scheduled in Schedules I through V of Section
418	58-37-4;
419	(B) a substance for which there is an approved new drug application;
420	(C) a substance with respect to which an exemption is in effect for investigational use
421	by a particular person under Section 505 of the Food, Drug, and Cosmetic Act, 21 U.S.C. 355,
422	to the extent the conduct with respect to the substance is permitted by the exemption;
423	(D) any substance to the extent not intended for human consumption before an
424	exemption takes effect with respect to the substance;
425	(E) any drug intended for lawful use in the diagnosis, cure, mitigation, treatment, or

- (E) any drug intended for lawful use in the diagnosis, cure, mitigation, treatment, or prevention of disease in man or other animals, which contains ephedrine, pseudoephedrine, norpseudoephedrine, or phenylpropanolamine if the drug is lawfully purchased, sold,
- 428 transferred, or furnished as an over-the-counter medication without prescription; or

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(F) dietary supplements, vitamins, minerals, herbs, or other similar substances including concentrates or extracts, which are not otherwise regulated by law, which may

contain naturally occurring amounts of chemical or substances listed in this chapter, or in rules adopted pursuant to Title 63G, Chapter 3, Utah Administrative Rulemaking Act.

- (h) "Conviction" means a determination of guilt by verdict, whether jury or bench, or plea, whether guilty or no contest, for any offense proscribed by Title 58, Chapters 37, 37a, 37b, 37c, or 37d, or for any offense under the laws of the United States and any other state which, if committed in this state, would be an offense under Title 58, Chapters 37, 37a, 37b, 37c, or 37d.
 - (i) "Counterfeit substance" means:

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- (i) any controlled substance or container or labeling of any controlled substance that:
- (A) without authorization bears the trademark, trade name, or other identifying mark, imprint, number, device, or any likeness of them, of a manufacturer, distributor, or dispenser other than the person or persons who in fact manufactured, distributed, or dispensed the substance which falsely purports to be a controlled substance distributed by any other manufacturer, distributor, or dispenser; and
- (B) a reasonable person would believe to be a controlled substance distributed by an authorized manufacturer, distributor, or dispenser based on the appearance of the substance as described under Subsection (1)(i)(i)(A) or the appearance of the container of that controlled substance; or
 - (ii) any substance other than under Subsection (1)(i)(i) that:
- (A) is falsely represented to be any legally or illegally manufactured controlled substance; and
 - (B) a reasonable person would believe to be a legal or illegal controlled substance.
- (j) "Deliver" or "delivery" means the actual, constructive, or attempted transfer of a controlled substance or a listed chemical, whether or not an agency relationship exists.
 - (k) "Department" means the Department of Commerce.
- (l) "Depressant or stimulant substance" means:
- 457 (i) a drug which contains any quantity of barbituric acid or any of the salts of barbituric 458 acid;
- (ii) a drug which contains any quantity of:
- 460 (A) amphetamine or any of its optical isomers;
- (B) any salt of amphetamine or any salt of an optical isomer of amphetamine; or

(C) any substance which the Secretary of Health and Human Services or the Attorney
General of the United States after investigation has found and by regulation designated
habit-forming because of its stimulant effect on the central nervous system;
(iii) lysergic acid diethylamide; or
(iv) any drug which contains any quantity of a substance which the Secretary of Health
and Human Services or the Attorney General of the United States after investigation has found

- and Human Services or the Attorney General of the United States after investigation has found to have, and by regulation designated as having, a potential for abuse because of its depressant or stimulant effect on the central nervous system or its hallucinogenic effect.
- (m) "Dispense" means the delivery of a controlled substance by a pharmacist to an ultimate user pursuant to the lawful order or prescription of a practitioner, and includes distributing to, leaving with, giving away, or disposing of that substance as well as the packaging, labeling, or compounding necessary to prepare the substance for delivery.
 - (n) "Dispenser" means a pharmacist who dispenses a controlled substance.
- (o) "Distribute" means to deliver other than by administering or dispensing a controlled substance or a listed chemical.
 - (p) "Distributor" means a person who distributes controlled substances.
- (q) "Division" means the Division of Occupational and Professional Licensing created in Section 58-1-103.
 - (r) (i) "Drug" means:

- (A) a substance recognized in the official United States Pharmacopoeia, Official Homeopathic Pharmacopoeia of the United States, or Official National Formulary, or any supplement to any of them, intended for use in the diagnosis, cure, mitigation, treatment, or prevention of disease in humans or animals;
- (B) a substance that is required by any applicable federal or state law or rule to be dispensed by prescription only or is restricted to administration by practitioners only;
- (C) a substance other than food intended to affect the structure or any function of the body of humans or other animals; and
- (D) substances intended for use as a component of any substance specified in Subsections (1)(r)(i)(A), (B), and (C)[, and (D)[.
 - (ii) "Drug" does not include dietary supplements.
- 492 (s) "Drug dependent person" means any individual who unlawfully and habitually uses

any controlled substance to endanger the public morals, health, safety, or welfare, or who is so dependent upon the use of controlled substances as to have lost the power of self-control with reference to the individual's dependency.

(t) "Food" means:

- (i) any nutrient or substance of plant, mineral, or animal origin other than a drug as specified in this chapter, and normally ingested by human beings; and
- (ii) foods for special dietary uses as exist by reason of a physical, physiological, pathological, or other condition including but not limited to the conditions of disease, convalescence, pregnancy, lactation, allergy, hypersensitivity to food, underweight, and overweight; uses for supplying a particular dietary need which exist by reason of age including but not limited to the ages of infancy and childbirth, and also uses for supplementing and for fortifying the ordinary or unusual diet with any vitamin, mineral, or other dietary property for use of a food. Any particular use of a food is a special dietary use regardless of the nutritional purposes.
- (u) "Immediate precursor" means a substance which the Attorney General of the United States has found to be, and by regulation designated as being, the principal compound used or produced primarily for use in the manufacture of a controlled substance, or which is an immediate chemical intermediary used or likely to be used in the manufacture of a controlled substance, the control of which is necessary to prevent, curtail, or limit the manufacture of the controlled substance.
 - (v) "Indian" means a member of an Indian tribe.
 - (w) "Indian religion" means any religion:
- (i) the origin and interpretation of which is from within a traditional Indian culture or community; and
 - (ii) which is practiced by Indians.
- (x) "Indian tribe" means any tribe, band, nation, pueblo, or other organized group or community of Indians, including any Alaska Native village, which is legally recognized as eligible for and is consistent with the special programs, services, and entitlements provided by the United States to Indians because of their status as Indians.
- (y) "Manufacture" means the production, preparation, propagation, compounding, or processing of a controlled substance, either directly or indirectly by extraction from substances

of natural origin, or independently by means of chemical synthesis or by a combination of extraction and chemical synthesis.

- (z) "Manufacturer" includes any person who packages, repackages, or labels any container of any controlled substance, except pharmacists who dispense or compound prescription orders for delivery to the ultimate consumer.
- (aa) "Marijuana" means all species of the genus cannabis and all parts of the genus, whether growing or not; the seeds of it; the resin extracted from any part of the plant; and every compound, manufacture, salt, derivative, mixture, or preparation of the plant, its seeds, or resin. The term does not include the mature stalks of the plant, fiber produced from the stalks, oil or cake made from the seeds of the plant, any other compound, manufacture, salt, derivative, mixture, or preparation of the mature stalks, except the resin extracted from them, fiber, oil or cake, or the sterilized seed of the plant which is incapable of germination. Any synthetic equivalents of the substances contained in the plant cannabis sativa or any other species of the genus cannabis which are chemically indistinguishable and pharmacologically active are also included.
- (bb) "Money" means officially issued coin and currency of the United States or any foreign country.
- (cc) "Narcotic drug" means any of the following, whether produced directly or indirectly by extraction from substances of vegetable origin, or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis:
 - (i) opium, coca leaves, and opiates;

- (ii) a compound, manufacture, salt, derivative, or preparation of opium, coca leaves, or opiates;
 - (iii) opium poppy and poppy straw; or
- (iv) a substance, and any compound, manufacture, salt, derivative, or preparation of the substance, which is chemically identical with any of the substances referred to in Subsection (1)(cc)(i), (ii), or (iii), except narcotic drug does not include decocainized coca leaves or extracts of coca leaves which do not contain cocaine or ecgonine.
- (dd) "Negotiable instrument" means documents, containing an unconditional promise to pay a sum of money, which are legally transferable to another party by endorsement or delivery.

(ee) "Opiate" means any drug or other substance having an addiction-forming or addiction-sustaining liability similar to morphine or being capable of conversion into a drug having addiction-forming or addiction-sustaining liability.

- (ff) "Opium poppy" means the plant of the species papaver somniferum L., except the seeds of the plant.
- (gg) "Person" means any corporation, association, partnership, trust, other institution or entity or one or more individuals.
- (hh) "Poppy straw" means all parts, except the seeds, of the opium poppy, after mowing.
- (ii) "Possession" or "use" means the joint or individual ownership, control, occupancy, holding, retaining, belonging, maintaining, or the application, inhalation, swallowing, injection, or consumption, as distinguished from distribution, of controlled substances and includes individual, joint, or group possession or use of controlled substances. For a person to be a possessor or user of a controlled substance, it is not required that the person be shown to have individually possessed, used, or controlled the substance, but it is sufficient if it is shown that the person jointly participated with one or more persons in the use, possession, or control of any substances with knowledge that the activity was occurring, or the controlled substance is found in a place or under circumstances indicating that the person had the ability and the intent to exercise dominion and control over it.
- (jj) "Practitioner" means a physician, dentist, naturopathic physician, veterinarian, pharmacist, scientific investigator, pharmacy, hospital, or other person licensed, registered, or otherwise permitted to distribute, dispense, conduct research with respect to, administer, or use in teaching or chemical analysis a controlled substance in the course of professional practice or research in this state.
 - (kk) "Prescribe" means to issue a prescription:
 - (i) orally or in writing; or

- (ii) by telephone, facsimile transmission, computer, or other electronic means of communication as defined by division rule.
 - (ll) "Prescription" means an order issued:
- (i) by a licensed practitioner, in the course of that practitioner's professional practice or by collaborative pharmacy practice agreement; and

586	(ii) for a controlled substance or other prescription drug or device for use by a patient
587	or an animal.
588	(mm) "Production" means the manufacture, planting, cultivation, growing, or
589	harvesting of a controlled substance.
590	(nn) "Securities" means any stocks, bonds, notes, or other evidences of debt or of
591	property.
592	(oo) "State" means the state of Utah.
593	(pp) "Ultimate user" means any person who lawfully possesses a controlled substance
594	for the person's own use, for the use of a member of the person's household, or for
595	administration to an animal owned by the person or a member of the person's household.
596	(2) If a term used in this chapter is not defined, the definition and terms of Title 76,
597	Utah Criminal Code, shall apply.
598	Section 4. Section 58-37-3 is amended to read:
599	58-37-3. Controlled substances.
600	(1) All substances listed in Section 58-37-4 or Section 58-37-4.2 are [considered]
601	controlled.
602	(2) All substances listed in the federal Controlled Substances Act, Title II, P.L. 91-513,
603	are [considered] controlled.
604	Section 5. Section 58-37-4 is amended to read:
605	58-37-4. Schedules of controlled substances Schedules I through V Findings
606	required Specific substances included in schedules.
607	(1) There are established five schedules of controlled substances known as Schedules I,
608	II, III, IV, and V which [shall] consist of substances listed in this section.
609	(2) Schedules I, II, III, IV, and V consist of the following drugs or other substances by
610	the official name, common or usual name, chemical name, or brand name designated:
611	(a) Schedule I:
612	(i) Unless specifically excepted or unless listed in another schedule, any of the
613	following opiates, including their isomers, esters, ethers, salts, and salts of isomers, esters, and
614	ethers, when the existence of the isomers, esters, ethers, and salts is possible within the specific
615	chemical designation:
616	(A) Acetyl-alpha-methylfentanyl

617	(N-[1-(1-methyl-2-phenethyl)-4-piperidinyl]-N-phenylacetamide);
618	(B) Acetylmethadol;
619	(C) Allylprodine;
620	(D) Alphacetylmethadol, except levo-alphacetylmethadol also known as
621	levo-alpha-acetylmethadol, levomethadyl acetate, or LAAM;
622	(E) Alphameprodine;
623	(F) Alphamethadol;
624	(G) Alpha-methylfentanyl (N-[1-(alpha-methyl-beta-phenyl)ethyl-4-piperidyl]
625	propionanilide; 1-(1-methyl-2-phenylethyl)-4-(N-propanilido) piperidine);
626	(H) Alpha-methylthiofentanyl (N-[1-methyl-2-(2-thienyl)ethyl-4-
627	piperidinyl]-N-phenylpropanamide);
628	(I) Benzethidine;
629	(J) Betacetylmethadol;
630	(K) Beta-hydroxyfentanyl (N-[1-(2-hydroxy-2-phenethyl)-4-
631	piperidinyl]-N-phenylpropanamide);
632	(L) Beta-hydroxy-3-methylfentanyl, other name: N-[1-(2-hydroxy-2-
633	phenethyl)-3-methyl-4-piperidinyl]-N-phenylpropanamide;
634	(M) Betameprodine;
635	(N) Betamethadol;
636	(O) Betaprodine;
637	(P) Clonitazene;
638	(Q) Dextromoramide;
639	(R) Diampromide;
640	(S) Diethylthiambutene;
641	(T) Difenoxin;
642	(U) Dimenoxadol;
643	(V) Dimepheptanol;
644	(W) Dimethylthiambutene;
645	(X) Dioxaphetyl butyrate;
646	(Y) Dipipanone;
647	(Z) Ethylmethylthiambutene;

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648
             (AA) Etonitazene;
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             (BB) Etoxeridine;
650
             (CC) Furethidine;
651
             (DD) Hydroxypethidine;
652
             (EE) Ketobemidone;
653
             (FF) Levomoramide;
654
             (GG) Levophenacylmorphan;
655
             (HH) Morpheridine;
656
             (II) MPPP (1-methyl-4-phenyl-4-propionoxypiperidine);
657
             (JJ) Noracymethadol;
658
             (KK) Norlevorphanol;
659
             (LL) Normethadone;
660
             (MM) Norpipanone;
661
             (NN) Para-fluorofentanyl (N-(4-fluorophenyl)-N-[1-(2-phenethyl)-4-piperidinyl]
662
      propanamide;
663
             (OO) PEPAP (1-(-2-phenethyl)-4-phenyl-4-acetoxypiperidine);
664
             (PP) Phenadoxone;
665
             (QQ) Phenampromide;
666
             (RR) Phenomorphan;
667
             (SS) Phenoperidine;
668
             (TT) Piritramide;
669
             (UU) Proheptazine;
670
             (VV) Properidine;
671
             (WW) Propiram;
672
             (XX) Racemoramide;
673
             (YY) Thiofentanyl (N-phenyl-N-[1-(2-thienyl)ethyl-4-piperidinyl]- propanamide;
674
             (ZZ) Tilidine;
675
             (AAA) Trimeperidine;
676
             (BBB) 3-methylfentanyl, including the optical and geometric isomers
677
      (N-[3-methyl-1-(2-phenylethyl)-4-piperidyl]- N-phenylpropanamide); and
678
             (CCC) 3-methylthiofentanyl
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679 (N-[(3-methyl-1-(2-thienyl)ethyl-4-piperidinyl]-N-phenylpropanamide). 680 (ii) Unless specifically excepted or unless listed in another schedule, any of the 681 following opium derivatives, their salts, isomers, and salts of isomers when the existence of the 682 salts, isomers, and salts of isomers is possible within the specific chemical designation: 683 (A) Acetorphine; 684 (B) Acetyldihydrocodeine; 685 (C) Benzylmorphine; 686 (D) Codeine methylbromide; 687 (E) Codeine-N-Oxide; 688 (F) Cyprenorphine; 689 (G) Desomorphine; 690 (H) Dihydromorphine; 691 (I) Drotebanol; 692 (J) Etorphine (except hydrochloride salt); 693 (K) Heroin; 694 (L) Hydromorphinol; 695 (M) Methyldesorphine; 696 (N) Methylhydromorphine; 697 (O) Morphine methylbromide; 698 (P) Morphine methylsulfonate; 699 (Q) Morphine-N-Oxide; 700 (R) Myrophine; 701 (S) Nicocodeine; 702 (T) Nicomorphine; 703 (U) Normorphine; 704 (V) Pholcodine; and 705 (W) Thebacon. 706 (iii) Unless specifically excepted or unless listed in another schedule, any material, 707 compound, mixture, or preparation which contains any quantity of the following hallucinogenic 708 substances, or which contains any of their salts, isomers, and salts of isomers when the

existence of the salts, isomers, and salts of isomers is possible within the specific chemical

710 designation; as used in this Subsection (2)(iii) only, "isomer" includes the optical, position, and 711 geometric isomers: 712 (A) Alpha-ethyltryptamine, some trade or other names: etryptamine: Monase: 713 α -ethyl-1H-indole-3-ethanamine; 3-(2-aminobutyl) indole; α -ET; and AET; 714 (B) 4-bromo-2,5-dimethoxy-amphetamine, some trade or other names: 715 4-bromo-2,5-dimethoxy-α-methylphenethylamine; 4-bromo-2,5-DMA; 716 (C) 4-bromo-2,5-dimethoxypenethylamine, some trade or other names: 717 2-(4-bromo-2,5-dimethoxyphenyl)-1-aminoethane; alpha-desmethyl DOB; 2C-B, Nexus; 718 (D) 2,5-dimethoxyamphetamine, some trade or other names: 719 2,5-dimethoxy-α-methylphenethylamine; 2,5-DMA; 720 (E) 2,5-dimethoxy-4-ethylamphetamine, some trade or other names: DOET; 721 (F) 4-methoxyamphetamine, some trade or other names: 722 4-methoxy-α-methylphenethylamine; paramethoxyamphetamine, PMA; 723 (G) 5-methoxy-3,4-methylenedioxyamphetamine; 724 (H) 4-methyl-2,5-dimethoxy-amphetamine, some trade and other names: 725 4-methyl-2,5-dimethoxy-α-methylphenethylamine; "DOM"; and "STP"; 726 (I) 3,4-methylenedioxy amphetamine; 727 (J) 3,4-methylenedioxymethamphetamine (MDMA); 728 (K) 3,4-methylenedioxy-N-ethylamphetamine, also known as N-ethyl-729 alpha-methyl-3,4(methylenedioxy)phenethylamine, N-ethyl MDA, MDE, MDEA; 730 (L) N-hydroxy-3,4-methylenedioxyamphetamine, also known as 731 N-hydroxy-alpha-methyl-3,4(methylenedioxy)phenethylamine, and N-hydroxy MDA; 732 (M) 3,4,5-trimethoxy amphetamine; 733 (N) Bufotenine, some trade and other names: 734 3-(β-Dimethylaminoethyl)-5-hydroxyindole; 3-(2-dimethylaminoethyl)-5-indolol; N, 735 N-dimethylserotonin; 5-hydroxy-N,N-dimethyltryptamine; mappine; 736 (O) Diethyltryptamine, some trade and other names: N,N-Diethyltryptamine; DET; 737 (P) Dimethyltryptamine, some trade or other names: DMT; 738 (Q) Ibogaine, some trade and other names: 739 7-Ethyl-6,6β,7,8,9,10,12,13-octahydro-2-methoxy-6,9-methano-5H-pyrido [1', 2':1,2] azepino

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[5,4-b] indole; Tabernanthe iboga;

741	(R) Lysergic acid diethylamide;
742	(S) Marijuana;
743	(T) Mescaline;
744	(U) Parahexyl, some trade or other names:
745	3-Hexyl-1-hydroxy-7,8,9,10-tetrahydro-6,6,9-trimethyl-6H-dibenzo[b,d]pyran; Synhexyl;
746	(V) Peyote, meaning all parts of the plant presently classified botanically as
747	Lophophora williamsii Lemaire, whether growing or not, the seeds thereof, any extract from
748	any part of such plant, and every compound, manufacture, salts, derivative, mixture, or
749	preparation of such plant, its seeds or extracts (Interprets 21 USC 812(c), Schedule I(c) (12));
750	(W) N-ethyl-3-piperidyl benzilate;
751	(X) N-methyl-3-piperidyl benzilate;
752	(Y) Psilocybin;
753	(Z) Psilocyn;
754	(AA) Tetrahydrocannabinols, naturally contained in a plant of the genus Cannabis
755	(cannabis plant), as well as synthetic equivalents of the substances contained in the cannabis
756	plant, or in the resinous extractives of Cannabis, sp. and/or synthetic substances, derivatives,
757	and their isomers with similar chemical structure and pharmacological activity to those
758	substances contained in the plant, such as the following: $\Delta 1$ cis or trans tetrahydrocannabinol,
759	and their optical isomers $\Delta 6$ cis or trans tetrahydrocannabinol, and their optical isomers $\Delta 3,4$
760	cis or trans tetrahydrocannabinol, and its optical isomers, and since nomenclature of these
761	substances is not internationally standardized, compounds of these structures, regardless of
762	numerical designation of atomic positions covered;
763	(BB) Ethylamine analog of phencyclidine, some trade or other names:
764	N-ethyl-1-phenylcyclohexylamine, (1-phenylcyclohexyl)ethylamine,
765	N-(1-phenylcyclohexyl)ethylamine, cyclohexamine, PCE;
766	(CC) Pyrrolidine analog of phencyclidine, some trade or other names:
767	1-(1-phenylcyclohexyl)-pyrrolidine, PCPy, PHP;
768	(DD) Thiophene analog of phencyclidine, some trade or other names:
769	1-[1-(2-thienyl)-cyclohexyl]-piperidine, 2-thienylanalog of phencyclidine, TPCP, TCP; and
770	(EE) 1-[1-(2-thienyl)cyclohexyl]pyrrolidine, some other names: TCPy.
771	(iv) Unless specifically excepted or unless listed in another schedule, any material

772 compound, mixture, or preparation which contains any quantity of the following substances 773 having a depressant effect on the central nervous system, including its salts, isomers, and salts 774 of isomers when the existence of the salts, isomers, and salts of isomers is possible within the 775 specific chemical designation:

- (A) Mecloqualone; and
- 777 (B) Methaqualone.

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- 778 (v) Any material, compound, mixture, or preparation containing any quantity of the 779 following substances having a stimulant effect on the central nervous system, including their 780 salts, isomers, and salts of isomers:
- 781 (A) Aminorex, some other names; aminoxaphen; 2-amino-5-phenyl-2-oxazoline; or 782 4,5-dihydro-5-phenyl-2-oxazolamine;
- 783 (B) Cathinone, some trade or other names: 2-amino-1-phenyl-1-propanone, alpha-aminopropiophenone, 2-aminopropiophenone, and norephedrone;
- 785 (C) Fenethylline;
- 786 (D) Methcathinone, some other names: 2-(methylamino)-propiophenone; 787 alpha-(methylamino)propiophenone; 2-(methylamino)-1-phenylpropan-1-one; 788 alpha-N-methylaminopropiophenone; monomethylpropion; ephedrone; N-methylcathinone; 789 methylcathinone; AL-464; AL-422; AL-463 and UR1432, its salts, optical isomers, and salts of 790 optical isomers;
 - (E) (±)cis-4-methylaminorex ((±)cis-4,5-dihydro-4-methyl-5-phenyl-2-oxazolamine);
- 792 (F) N-ethylamphetamine; and
- 793 (G) N,N-dimethylamphetamine, also known as
- 794 N,N-alpha-trimethyl-benzeneethanamine; N,N-alpha-trimethylphenethylamine.
 - (vi) Any material, compound, mixture, or preparation which contains any quantity of the following substances, including their optical isomers, salts, and salts of isomers, subject to temporary emergency scheduling:
 - (A) N-[1-benzyl-4-piperidyl]-N-phenylpropanamide (benzylfentanyl); and
- 799 (B) N-[1- (2-thienyl)methyl-4-piperidyl]-N-phenylpropanamide (thenylfentanyl).
- (vii) Unless specifically excepted or unless listed in another schedule, any material, 800 801 compound, mixture, or preparation which contains any quantity of gamma hydroxy butyrate 802 (gamma hydrobutyric acid), including its salts, isomers, and salts of isomers.

803	(b) Schedule II:
804	(i) Unless specifically excepted or unless listed in another schedule, any of the
805	following substances whether produced directly or indirectly by extraction from substances of
806	vegetable origin, or independently by means of chemical synthesis, or by a combination of
807	extraction and chemical synthesis:
808	(A) Opium and opiate, and any salt, compound, derivative, or preparation of opium or
809	opiate, excluding apomorphine, dextrorphan, nalbuphine, nalmefene, naloxone, and naltrexone
810	and their respective salts, but including:
811	(I) Raw opium;
812	(II) Opium extracts;
813	(III) Opium fluid;
814	(IV) Powdered opium;
815	(V) Granulated opium;
816	(VI) Tincture of opium;
817	(VII) Codeine;
818	(VIII) Ethylmorphine;
819	(IX) Etorphine hydrochloride;
820	(X) Hydrocodone;
821	(XI) Hydromorphone;
822	(XII) Metopon;
823	(XIII) Morphine;
824	(XIV) Oxycodone;
825	(XV) Oxymorphone; and
826	(XVI) Thebaine;
827	(B) Any salt, compound, derivative, or preparation which is chemically equivalent or
828	identical with any of the substances referred to in Subsection (2)(b)(i)(A), except that these
829	substances may not include the isoquinoline alkaloids of opium;
830	(C) Opium poppy and poppy straw;
831	(D) Coca leaves and any salt, compound, derivative, or preparation of coca leaves, and
832	any salt, compound, derivative, or preparation which is chemically equivalent or identical with
833	any of these substances, and includes cocaine and ecgonine, their salts, isomers, derivatives,

834 and salts of isomers and derivatives, whether derived from the coca plant or synthetically 835 produced, except the substances may not include decocainized coca leaves or extraction of coca 836 leaves, which extractions do not contain cocaine or ecgonine; and 837 (E) Concentrate of poppy straw, which means the crude extract of poppy straw in either 838 liquid, solid, or powder form which contains the phenanthrene alkaloids of the opium poppy. 839 (ii) Unless specifically excepted or unless listed in another schedule, any of the 840 following opiates, including their isomers, esters, ethers, salts, and salts of isomers, esters, and 841 ethers, when the existence of the isomers, esters, ethers, and salts is possible within the specific 842 chemical designation, except dextrorphan and levopropoxyphene: 843 (A) Alfentanil; 844 (B) Alphaprodine; 845 (C) Anileridine; 846 (D) Bezitramide; 847 (E) Bulk dextropropoxyphene (nondosage forms); 848 (F) Carfentanil; 849 (G) Dihydrocodeine; 850 (H) Diphenoxylate; 851 (I) Fentanyl; 852 (J) Isomethadone; 853 (K) Levo-alphacetylmethadol, some other names: levo-alpha-acetylmethadol, 854 levomethadyl acetate, or LAAM; 855 (L) Levomethorphan; 856 (M) Levorphanol; 857 (N) Metazocine; 858 (O) Methadone; 859 (P) Methadone-Intermediate, 4-cyano-2-dimethylamino-4, 4-diphenyl butane; 860 (Q) Moramide-Intermediate, 2-methyl-3-morpholino-1, 1-diphenylpropane-carboxylic 861 acid; 862 (R) Pethidine (meperidine); 863 (S) Pethidine-Intermediate-A, 4-cyano-1-methyl-4-phenylpiperidine;

(T) Pethidine-Intermediate-B, ethyl-4-phenylpiperidine-4-carboxylate;

865	(U) Pethidine-Intermediate-C, 1-methyl-4-phenylpiperidine-4-carboxylic acid;
866	(V) Phenazocine;
867	(W) Piminodine;
868	(X) Racemethorphan;
869	(Y) Racemorphan;
870	(Z) Remifentanil; and
871	(AA) Sufentanil.
872	(iii) Unless specifically excepted or unless listed in another schedule, any material,
873	compound, mixture, or preparation which contains any quantity of the following substances
874	having a stimulant effect on the central nervous system:
875	(A) Amphetamine, its salts, optical isomers, and salts of its optical isomers;
876	(B) Methamphetamine, its salts, isomers, and salts of its isomers;
877	(C) Phenmetrazine and its salts; and
878	(D) Methylphenidate.
879	(iv) Unless specifically excepted or unless listed in another schedule, any material,
880	compound, mixture, or preparation which contains any quantity of the following substances
881	having a depressant effect on the central nervous system, including its salts, isomers, and salts
882	of isomers when the existence of the salts, isomers, and salts of isomers is possible within the
883	specific chemical designation:
884	(A) Amobarbital;
885	(B) Glutethimide;
886	(C) Pentobarbital;
887	(D) Phencyclidine;
888	(E) Phencyclidine immediate precursors: 1-phenylcyclohexylamine and
889	1-piperidinocyclohexanecarbonitrile (PCC); and
890	(F) Secobarbital.
891	(v) (A) Unless specifically excepted or unless listed in another schedule, any material,
892	compound, mixture, or preparation which contains any quantity of Phenylacetone.
893	(B) Some of these substances may be known by trade or other names:
894	phenyl-2-propanone; P2P; benzyl methyl ketone; and methyl benzyl ketone.
895	(vi) Nabilone, another name for nabilone:

896 (±)-trans-3-(1,1-dimethylheptyl)-6,6a,7,8,10,10a-hexahydro-1-hydroxy-6, 897 6-dimethyl-9H-dibenzo[b,d]pyran-9-one.

(c) Schedule III:

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- (i) Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation which contains any quantity of the following substances having a stimulant effect on the central nervous system, including its salts, isomers whether optical, position, or geometric, and salts of the isomers when the existence of the salts, isomers, and salts of isomers is possible within the specific chemical designation:
- (A) Those compounds, mixtures, or preparations in dosage unit form containing any stimulant substances listed in Schedule II, which compounds, mixtures, or preparations were listed on August 25, 1971, as excepted compounds under Section 1308.32 of Title 21 of the Code of Federal Regulations, and any other drug of the quantitive composition shown in that list for those drugs or which is the same except that it contains a lesser quantity of controlled substances;
- 910 (B) Benzphetamine;
 - (C) Chlorphentermine;
- 912 (D) Clortermine; and
- 913 (E) Phendimetrazine.
 - (ii) Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation which contains any quantity of the following substances having a depressant effect on the central nervous system:
 - (A) Any compound, mixture, or preparation containing amobarbital, secobarbital, pentobarbital, or any salt of any of them, and one or more other active medicinal ingredients which are not listed in any schedule;
 - (B) Any suppository dosage form containing amobarbital, secobarbital, or pentobarbital, or any salt of any of these drugs which is approved by the Food and Drug Administration for marketing only as a suppository;
- 923 (C) Any substance which contains any quantity of a derivative of barbituric acid or any salt of any of them;
- 925 (D) Chlorhexadol;
- 926 (E) Buprenorphine;

927 (F) Any drug product containing gamma hydroxybutyric acid, including its salts, 928 isomers, and salts of isomers, for which an application is approved under the federal Food, 929 Drug, and Cosmetic Act, Section 505; 930 (G) Ketamine, its salts, isomers, and salts of isomers, some other names for ketamine: 931 ± -2-(2-chlorophenyl)-2-(methylamino)-cyclohexanone; 932 (H) Lysergic acid; 933 (I) Lysergic acid amide; 934 (J) Methyprylon; 935 (K) Sulfondiethylmethane; 936 (L) Sulfonethylmethane; 937 (M) Sulfonmethane; and 938 (N) Tiletamine and zolazepam or any of their salts, some trade or other names for a 939 tiletamine-zolazepam combination product: Telazol, some trade or other names for tiletamine: 940 2-(ethylamino)-2-(2-thienyl)-cyclohexanone, some trade or other names for zolazepam: 941 4-(2-fluorophenyl)-6,8-dihydro-1,3,8-trimethylpyrazolo-[3,4-e] [1,4]-diazepin-7(1H)-one, 942 flupyrazapon. 943 (iii) Dronabinol (synthetic) in sesame oil and encapsulated in a soft gelatin capsule in a 944 U.S. Food and Drug Administration approved drug product, some other names for dronabinol: 945 (6aR-trans)-6a,7,8,10a-tetrahydro-6,6,9-trimethyl-3-pentyl-6H-dibenzo[b,d]pyran-1-ol, or 946 (-)-delta-9-(trans)-tetrahydrocannabinol. 947 (iv) Nalorphine. 948 (v) Unless specifically excepted or unless listed in another schedule, any material, 949 compound, mixture, or preparation containing limited quantities of any of the following 950 narcotic drugs, or their salts calculated as the free anhydrous base or alkaloid: 951 (A) Not more than 1.8 grams of codeine per 100 milliliters or not more than 90 952 milligrams per dosage unit, with an equal or greater quantity of an isoquinoline alkaloid of 953 opium; 954 (B) Not more than 1.8 grams of codeine per 100 milliliters or not more than 90 955 milligrams per dosage unit, with one or more active non-narcotic ingredients in recognized 956 therapeutic amounts; 957 (C) Not more than 300 milligrams of dihydrocodeinone per 100 milliliters or not more

than 15 milligrams per dosage unit, with a fourfold or greater quantity of an isoquinoline alkaloid of opium;

- (D) Not more than 300 milligrams of dihydrocodeinone per 100 milliliters or not more than 15 milligrams per dosage unit, with one or more active, non-narcotic ingredients in recognized therapeutic amounts;
- (E) Not more than 1.8 grams of dihydrocodeine per 100 milliliters or not more than 90 milligrams per dosage unit, with one or more active non-narcotic ingredients in recognized therapeutic amounts;
- (F) Not more than 300 milligrams of ethylmorphine per 100 milliliters or not more than 15 milligrams per dosage unit, with one or more active, non-narcotic ingredients in recognized therapeutic amounts;
- (G) Not more than 500 milligrams of opium per 100 milliliters or per 100 grams, or not more than 25 milligrams per dosage unit, with one or more active, non-narcotic ingredients in recognized therapeutic amounts; and
- (H) Not more than 50 milligrams of morphine per 100 milliliters or per 100 grams with one or more active, non-narcotic ingredients in recognized therapeutic amounts.
- (vi) Unless specifically excepted or unless listed in another schedule, anabolic steroids including any of the following or any isomer, ester, salt, or derivative of the following that promotes muscle growth:
- 977 (A) Boldenone;

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- 978 (B) Chlorotestosterone (4-chlortestosterone);
- 979 (C) Clostebol;
- 980 (D) Dehydrochlormethyltestosterone;
- 981 (E) Dihydrotestosterone (4-dihydrotestosterone);
- 982 (F) Drostanolone;
- 983 (G) Ethylestrenol;
- 984 (H) Fluoxymesterone;
- 985 (I) Formebulone (formebolone);
- 986 (J) Mesterolone;
- 987 (K) Methandienone;
- 988 (L) Methandranone;

989	(M) Methandriol;
990	(N) Methandrostenolone;
991	(O) Methenolone;
992	(P) Methyltestosterone;
993	(Q) Mibolerone;
994	(R) Nandrolone;
995	(S) Norethandrolone;
996	(T) Oxandrolone;
997	(U) Oxymesterone;
998	(V) Oxymetholone;
999	(W) Stanolone;
1000	(X) Stanozolol;
1001	(Y) Testolactone;
1002	(Z) Testosterone; and
1003	(AA) Trenbolone.
1004	(vii) Anabolic steroids expressly intended for administration through implants to cattle
1005	or other nonhuman species, and approved by the Secretary of Health and Human Services for
1006	use, may not be classified as a controlled substance.
1007	(d) Schedule IV:
1008	(i) Unless specifically excepted or unless listed in another schedule, any material,
1009	compound, mixture, or preparation containing not more than 1 milligram of difenoxin and not
1010	less than 25 micrograms of atropine sulfate per dosage unit, or any salts of any of them.
1011	(ii) Unless specifically excepted or unless listed in another schedule, any material,
1012	compound, mixture, or preparation which contains any quantity of the following substances,
1013	including its salts, isomers, and salts of isomers when the existence of the salts, isomers, and
1014	salts of isomers is possible within the specific chemical designation:
1015	(A) Alprazolam;
1016	(B) Barbital;
1017	(C) Bromazepam;
1018	(D) Butorphanol;
1019	(E) Camazepam;

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1020	(F) Carisoprodol;
1021	(G) Chloral betaine;
1022	(H) Chloral hydrate;
1023	(I) Chlordiazepoxide;
1024	(J) Clobazam;
1025	(K) Clonazepam;
1026	(L) Clorazepate;
1027	(M) Clotiazepam;
1028	(N) Cloxazolam;
1029	(O) Delorazepam;
1030	(P) Diazepam;
1031	(Q) Dichloralphenazone;
1032	(R) Estazolam;
1033	(S) Ethchlorvynol;
1034	(T) Ethinamate;
1035	(U) Ethyl loflazepate;
1036	(V) Fludiazepam;
1037	(W) Flunitrazepam;
1038	(X) Flurazepam;
1039	(Y) Halazepam;
1040	(Z) Haloxazolam;
1041	(AA) Ketazolam;
1042	(BB) Loprazolam;
1043	(CC) Lorazepam;
1044	(DD) Lormetazepam;
1045	(EE) Mebutamate;
1046	(FF) Medazepam;
1047	(GG) Meprobamate;
1048	(HH) Methohexital;
1049	(II) Methylphenobarbital (mephobarbital);

(JJ) Midazolam;

1051	(KK) Nimetazepam;
1052	(LL) Nitrazepam;
1053	(MM) Nordiazepam;
1054	(NN) Oxazepam;
1055	(OO) Oxazolam;
1056	(PP) Paraldehyde;
1057	(QQ) Pentazocine;
1058	(RR) Petrichloral;
1059	(SS) Phenobarbital;
1060	(TT) Pinazepam;
1061	(UU) Prazepam;
1062	(VV) Quazepam;
1063	(WW) Temazepam;
1064	(XX) Tetrazepam;
1065	(YY) Triazolam;
1066	(ZZ) Zaleplon; and
1067	(AAA) Zolpidem.
1068	(iii) Any material, compound, mixture, or preparation of fenfluramine which contains
1069	any quantity of the following substances, including its salts, isomers whether optical, position,
1070	or geometric, and salts of the isomers when the existence of the salts, isomers, and salts of
1071	isomers is possible.
1072	(iv) Unless specifically excepted or unless listed in another schedule, any material,
1073	compound, mixture, or preparation which contains any quantity of the following substances
1074	having a stimulant effect on the central nervous system, including its salts, isomers whether
1075	optical, position, or geometric isomers, and salts of the isomers when the existence of the salts,
1076	isomers, and salts of isomers is possible within the specific chemical designation:
1077	(A) Cathine ((+)-norpseudoephedrine);
1078	(B) Diethylpropion;
1079	(C) Fencamfamine;
1080	(D) Fenproprex;
1081	(E) Mazindol;

1082	(F) Mefenorex;
1083	(G) Modafinil;
1084	(H) Pemoline, including organometallic complexes and chelates thereof;
1085	(I) Phentermine;
1086	(J) Pipradrol;
1087	(K) Sibutramine; and
1088	(L) SPA ((-)-1-dimethylamino-1,2-diphenylethane).
1089	(v) Unless specifically excepted or unless listed in another schedule, any material,
1090	compound, mixture, or preparation which contains any quantity of dextropropoxyphene
1091	(alpha-(+)-4-dimethylamino-1, 2-diphenyl-3-methyl-2-propionoxybutane), including its salts.
1092	(e) Schedule V: Any compound, mixture, or preparation containing any of the
1093	following limited quantities of narcotic drugs, or their salts calculated as the free anhydrous
1094	base or alkaloid, which includes one or more non-narcotic active medicinal ingredients in
1095	sufficient proportion to confer upon the compound, mixture, or preparation valuable medicinal
1096	qualities other than those possessed by the narcotic drug alone:
1097	(i) not more than 200 milligrams of codeine per 100 milliliters or per 100 grams;
1098	(ii) not more than 100 milligrams of dihydrocodeine per 100 milliliters or per 100
1099	grams;
1100	(iii) not more than 100 milligrams of ethylmorphine per 100 milliliters or per 100
1101	grams;
1102	(iv) not more than 2.5 milligrams of diphenoxylate and not less than 25 micrograms of
1103	atropine sulfate per dosage unit;
1104	(v) not more than 100 milligrams of opium per 100 milliliters or per 100 grams;
1105	(vi) not more than 0.5 milligram of difenoxin and not less than 25 micrograms of
1106	atropine sulfate per dosage unit; and
1107	(vii) unless specifically exempted or excluded or unless listed in another schedule, any
1108	material, compound, mixture, or preparation which contains Pyrovalerone having a stimulant
1109	effect on the central nervous system, including its salts, isomers, and salts of isomers.
1110	Section 6. Section 58-37-4.2 is enacted to read:
1111	58-37-4.2. Listed controlled substances.
1112	The following substances, their analogs, homologs, and synthetic equivalents are listed

1113	controlled substances:
1114	(1) AM-694;1-[(5-fluoropentyl)-1H-indol-3-yl]-(2-iodophenyl)methanone;
1115	(2) CP 47,497 and its C6, C8, and C9 homologs; 2-[(1R,3S)-3-hydroxycyclohexyl]
1116	-5-(2-methyloctan-2-yl)phenol;
1117	(3) HU-210; (6aR,10aR)-9-(hydroxymethyl)-6,6-dimethyl-3-(2-methyloctan-2-yl)
1118	-6a,7,10,10a-tetrahydrobenzo[c]chromen-1-ol;
1119	(4) HU-211; Dexanabinol,(6aS,10aS)-9-(hydroxymethyl)-6,6-dimethyl-3-(2-
1120	methyloctan-2-yl)-6a,7,10,10a-tetrahydrobenzo[c]chromen-1-ol;
1121	(5) JWH-015; (2-methyl-1-propyl-1H-indol-3-yl)-1-naphthalenyl-methanone;
1122	(6) JWH-018; Naphthalen-1-yl-(pentylindol-3-yl)methanone {also known as
1123	1-Pentyl-3-(1-naphthoyl)indole};
1124	(7) JWH-019; 1-hexyl-3-(1-naphthoyl)indole;
1125	(8) JWH-073; Naphthalen-1-yl(1-butylindol-3-yl)methanone {also known as
1126	1-Butyl-3-(1-naphthoyl)indole};
1127	(9) JWH-081; 4-methoxynaphthalen-1-yl-(1-pentylindol- 3-yl)methanone;
1128	(10) JWH-122; CAS#619294-47-2; (1-Pentyl-3-(4-methyl-1-naphthoyl)indole);
1129	(11) JWH-200; 1-(2-(4-(morpholinyl)ethyl))-3-(1-naphthoyl) indole;
1130	(12) JWH-250; 1-pentyl-3-(2-methoxyphenylacetyl)indole;
1131	(13) JWH-251; 2-(2-methylphenyl)-1-(1-pentyl-1H-indol-3-yl)-ethanone;
1132	(14) JWH-398; 1-pentyl-3-(4-chloro-1-naphthoyl)indole; and
1133	(15) RCS-8; 1-(2-cyclohexylethyl)-3-(2-methoxyphenylacetyl)indole {also known as
1134	BTW-8 and SR-18}.
1135	Section 7. Section 58-37-6 is amended to read:
1136	58-37-6. License to manufacture, produce, distribute, dispense, administer, or
1137	conduct research Issuance by division Denial, suspension, or revocation Records
1138	required Prescriptions.
1139	(1) (a) The division may adopt rules relating to the licensing and control of the
1140	manufacture, distribution, production, prescription, administration, dispensing, conducting of
1141	research with, and performing of laboratory analysis upon controlled substances within this
1142	state.
1143	(b) The division may assess reasonable fees to defray the cost of issuing original and

renewal licenses under this chapter pursuant to Section 63J-1-504.

(2) (a) (i) Every person who manufactures, produces, distributes, prescribes, dispenses, administers, conducts research with, or performs laboratory analysis upon any controlled substance in Schedules II through V within this state, or who proposes to engage in manufacturing, producing, distributing, prescribing, dispensing, administering, conducting research with, or performing laboratory analysis upon controlled substances included in Schedules II through V within this state shall obtain a license issued by the division.

- (ii) The division shall issue each license under this chapter in accordance with a two-year renewal cycle established by rule. The division may by rule extend or shorten a renewal period by as much as one year to stagger the renewal cycles it administers.
- (b) Persons licensed to manufacture, produce, distribute, prescribe, dispense, administer, conduct research with, or perform laboratory analysis upon controlled substances in Schedules II through V within this state may possess, manufacture, produce, distribute, prescribe, dispense, administer, conduct research with, or perform laboratory analysis upon those substances to the extent authorized by their license and in conformity with this chapter.
- (c) The following persons are not required to obtain a license and may lawfully possess controlled substances under this section:
- (i) an agent or employee, except a sales representative, of any registered manufacturer, distributor, or dispenser of any controlled substance, if the agent or employee is acting in the usual course of the person's business or employment; however, nothing in this subsection shall be interpreted to permit an agent, employee, sales representative, or detail man to maintain an inventory of controlled substances separate from the location of the person's employer's registered and licensed place of business;
- (ii) a motor carrier or warehouseman, or an employee of a motor carrier or warehouseman, who possesses any controlled substance in the usual course of the person's business or employment; and
- (iii) an ultimate user, or any person who possesses any controlled substance pursuant to a lawful order of a practitioner.
- (d) The division may enact rules waiving the license requirement for certain manufacturers, producers, distributors, prescribers, dispensers, administrators, research practitioners, or laboratories performing analysis if consistent with the public health and safety.

(e) A separate license is required at each principal place of business or professional practice where the applicant manufactures, produces, distributes, dispenses, conducts research with, or performs laboratory analysis upon controlled substances.

- (f) The division may enact rules providing for the inspection of a licensee or applicant's establishment, and may inspect the establishment according to those rules.
- (3) (a) Upon proper application, the division shall license a qualified applicant to manufacture, produce, distribute, conduct research with, or perform laboratory analysis upon controlled substances included in Schedules I through V, unless it determines that issuance of a license is inconsistent with the public interest. The division shall not issue a license to any person to prescribe, dispense, or administer a Schedule I controlled substance. In determining public interest, the division shall consider whether or not the applicant has:
- (i) maintained effective controls against diversion of controlled substances and any Schedule I or II substance compounded from any controlled substance into other than legitimate medical, scientific, or industrial channels;
 - (ii) complied with applicable state and local law;

- (iii) been convicted under federal or state laws relating to the manufacture, distribution, or dispensing of substances;
 - (iv) past experience in the manufacture of controlled dangerous substances;
 - (v) established effective controls against diversion; and
- (vi) complied with any other factors that the division establishes that promote the public health and safety.
- (b) Licenses granted under Subsection (3)(a) do not entitle a licensee to manufacture, produce, distribute, conduct research with, or perform laboratory analysis upon controlled substances in Schedule I other than those specified in the license.
- (c) (i) Practitioners shall be licensed to administer, dispense, or conduct research with substances in Schedules II through V if they are authorized to administer, dispense, or conduct research under the laws of this state.
- (ii) The division need not require a separate license for practitioners engaging in research with nonnarcotic controlled substances in Schedules II through V where the licensee is already licensed under this chapter in another capacity.
 - (iii) With respect to research involving narcotic substances in Schedules II through V,

or where the division by rule requires a separate license for research of nonnarcotic substances in Schedules II through V, a practitioner shall apply to the division prior to conducting research.

- (iv) Licensing for purposes of bona fide research with controlled substances by a practitioner considered qualified may be denied only on a ground specified in Subsection (4), or upon evidence that the applicant will abuse or unlawfully transfer or fail to safeguard adequately the practitioner's supply of substances against diversion from medical or scientific use.
- (v) Practitioners registered under federal law to conduct research in Schedule I substances may conduct research in Schedule I substances within this state upon furnishing the division evidence of federal registration.
- (d) Compliance by manufacturers, producers, and distributors with the provisions of federal law respecting registration, excluding fees, entitles them to be licensed under this chapter.
- (e) The division shall initially license those persons who own or operate an establishment engaged in the manufacture, production, distribution, dispensation, or administration of controlled substances prior to April 3, 1980, and who are licensed by the state.
- (4) (a) Any license pursuant to Subsection (2) or (3) may be denied, suspended, placed on probation, or revoked by the division upon finding that the applicant or licensee has:
 - (i) materially falsified any application filed or required pursuant to this chapter;
- (ii) been convicted of an offense under this chapter or any law of the United States, or any state, relating to any substance defined as a controlled substance;
- (iii) been convicted of a felony under any other law of the United States or any state within five years of the date of the issuance of the license;
- (iv) had a federal license denied, suspended, or revoked by competent federal authority and is no longer authorized to engage in the manufacturing, distribution, or dispensing of controlled substances;
- (v) had the licensee's license suspended or revoked by competent authority of another state for violation of laws or regulations comparable to those of this state relating to the manufacture, distribution, or dispensing of controlled substances;

(vi) violated any division rule that reflects adversely on the licensee's reliability and integrity with respect to controlled substances;

- (vii) refused inspection of records required to be maintained under this chapter by a person authorized to inspect them; or
- (viii) prescribed, dispensed, administered, or injected an anabolic steroid for the purpose of manipulating human hormonal structure so as to:
- (A) increase muscle mass, strength, or weight without medical necessity and without a written prescription by any practitioner in the course of the practitioner's professional practice; or
 - (B) improve performance in any form of human exercise, sport, or game.
- (b) The division may limit revocation or suspension of a license to a particular controlled substance with respect to which grounds for revocation or suspension exist.
- (c) (i) Proceedings to deny, revoke, or suspend a license shall be conducted pursuant to this section and in accordance with the procedures set forth in Title 58, Chapter 1, Division of Occupational and Professional Licensing Act, and conducted in conjunction with the appropriate representative committee designated by the director of the department.
- (ii) Nothing in this Subsection (4)(c) gives the Division of Occupational and Professional Licensing exclusive authority in proceedings to deny, revoke, or suspend licenses, except where the division is designated by law to perform those functions, or, when not designated by law, is designated by the executive director of the Department of Commerce to conduct the proceedings.
- (d) (i) The division may suspend any license simultaneously with the institution of proceedings under this section if it finds there is an imminent danger to the public health or safety.
- (ii) Suspension shall continue in effect until the conclusion of proceedings, including judicial review, unless withdrawn by the division or dissolved by a court of competent jurisdiction.
- (e) (i) If a license is suspended or revoked under this Subsection (4), all controlled substances owned or possessed by the licensee may be placed under seal in the discretion of the division.
 - (ii) Disposition may not be made of substances under seal until the time for taking an

appeal has lapsed, or until all appeals have been concluded, unless a court, upon application, orders the sale of perishable substances and the proceeds deposited with the court.

- (iii) If a revocation order becomes final, all controlled substances shall be forfeited.
- (f) The division shall notify promptly the Drug Enforcement Administration of all orders suspending or revoking a license and all forfeitures of controlled substances.
- (5) (a) Persons licensed under Subsection (2) or (3) shall maintain records and inventories in conformance with the record keeping and inventory requirements of federal and state law and any additional rules issued by the division.
- (b) (i) Every physician, dentist, naturopathic physician, veterinarian, practitioner, or other person who is authorized to administer or professionally use a controlled substance shall keep a record of the drugs received by him and a record of all drugs administered, dispensed, or professionally used by him otherwise than by a prescription.
- (ii) A person using small quantities or solutions or other preparations of those drugs for local application has complied with this Subsection (5)(b) if the person keeps a record of the quantity, character, and potency of those solutions or preparations purchased or prepared by him, and of the dates when purchased or prepared.
- (6) Controlled substances in Schedules I through V may be distributed only by a licensee and pursuant to an order form prepared in compliance with division rules or a lawful order under the rules and regulations of the United States.
- (7) (a) A person may not write or authorize a prescription for a controlled substance unless the person is:
- (i) a practitioner authorized to prescribe drugs and medicine under the laws of this state or under the laws of another state having similar standards; and
- (ii) licensed under this chapter or under the laws of another state having similar standards.
- (b) A person other than a pharmacist licensed under the laws of this state, or the pharmacist's licensed intern, as required by Sections 58-17b-303 and 58-17b-304, may not dispense a controlled substance.
- (c) (i) A controlled substance may not be dispensed without the written prescription of a practitioner, if the written prescription is required by the federal Controlled Substances Act.
 - (ii) That written prescription shall be made in accordance with Subsection (7)(a) and in

1299 conformity with Subsection (7)(d).

(iii) In emergency situations, as defined by division rule, controlled substances may be dispensed upon oral prescription of a practitioner, if reduced promptly to writing on forms designated by the division and filed by the pharmacy.

- (iv) Prescriptions reduced to writing by a pharmacist shall be in conformity with Subsection (7)(d).
- (d) Except for emergency situations designated by the division, a person may not issue, fill, compound, or dispense a prescription for a controlled substance unless the prescription is signed by the prescriber in ink or indelible pencil or is signed with an electronic signature of the prescriber as authorized by division rule, and contains the following information:
 - (i) the name, address, and registry number of the prescriber;
- 1310 (ii) the name, address, and age of the person to whom or for whom the prescription is 1311 issued;
 - (iii) the date of issuance of the prescription; and
 - (iv) the name, quantity, and specific directions for use by the ultimate user of the controlled substance.
 - (e) A prescription may not be written, issued, filled, or dispensed for a Schedule I controlled substance.
 - (f) Except when administered directly to an ultimate user by a licensed practitioner, controlled substances are subject to the following restrictions:
 - (i) (A) A prescription for a Schedule II substance may not be refilled.
 - (B) A Schedule II controlled substance may not be filled in a quantity to exceed a one-month's supply, as directed on the daily dosage rate of the prescriptions.
 - (ii) A Schedule III or IV controlled substance may be filled only within six months of issuance, and may not be refilled more than six months after the date of its original issuance or be refilled more than five times after the date of the prescription unless renewed by the practitioner.
 - (iii) All other controlled substances in Schedule V may be refilled as the prescriber's prescription directs, but they may not be refilled one year after the date the prescription was issued unless renewed by the practitioner.
- (iv) Any prescription for a Schedule II substance may not be dispensed if it is not

presented to a pharmacist for dispensing by a pharmacist or a pharmacy intern within 30 days after the date the prescription was issued, or 30 days after the dispensing date, if that date is specified separately from the date of issue.

- (v) A practitioner may issue more than one prescription at the same time for the same Schedule II controlled substance, but only under the following conditions:
- (A) no more than three prescriptions for the same Schedule II controlled substance may be issued at the same time;
 - (B) no one prescription may exceed a 30-day supply;

- (C) a second or third prescription shall include the date of issuance and the date for dispensing; and
- (D) unless the practitioner determines there is a valid medical reason to the contrary, the date for dispensing a second or third prescription may not be fewer than 30 days from the dispensing date of the previous prescription.
- (vi) Each prescription for a controlled substance may contain only one controlled substance per prescription form and may not contain any other legend drug or prescription item.
- (g) An order for a controlled substance in Schedules II through V for use by an inpatient or an outpatient of a licensed hospital is exempt from all requirements of this Subsection (7) if the order is:
- (i) issued or made by a prescribing practitioner who holds an unrestricted registration with the federal Drug Enforcement Administration, and an active Utah controlled substance license in good standing issued by the division under this section, or a medical resident who is exempted from licensure under Subsection 58-1-307(1)(c);
- (ii) authorized by the prescribing practitioner treating the patient and the prescribing practitioner designates the quantity ordered;
- (iii) entered upon the record of the patient, the record is signed by the prescriber affirming the prescriber's authorization of the order within 48 hours after filling or administering the order, and the patient's record reflects the quantity actually administered; and
- (iv) filled and dispensed by a pharmacist practicing the pharmacist's profession within the physical structure of the hospital, or the order is taken from a supply lawfully maintained by the hospital and the amount taken from the supply is administered directly to the patient

authorized to receive it.

(h) A practitioner licensed under this chapter may not prescribe, administer, or dispense a controlled substance to a child, without first obtaining the consent required in Section 78B-3-406 of a parent, guardian, or person standing in loco parentis of the child except in cases of an emergency. For purposes of this Subsection (7)(h), "child" has the same meaning as defined in Section 78A-6-105, and "emergency" means any physical condition requiring the administration of a controlled substance for immediate relief of pain or suffering.

- (i) A practitioner licensed under this chapter may not prescribe or administer dosages of a controlled substance in excess of medically recognized quantities necessary to treat the ailment, malady, or condition of the ultimate user.
- (j) A practitioner licensed under this chapter may not prescribe, administer, or dispense any controlled substance to another person knowing that the other person is using a false name, address, or other personal information for the purpose of securing the controlled substance.
- (k) A person who is licensed under this chapter to manufacture, distribute, or dispense a controlled substance may not manufacture, distribute, or dispense a controlled substance to another licensee or any other authorized person not authorized by this license.
- (l) A person licensed under this chapter may not omit, remove, alter, or obliterate a symbol required by this chapter or by a rule issued under this chapter.
- (m) A person licensed under this chapter may not refuse or fail to make, keep, or furnish any record notification, order form, statement, invoice, or information required under this chapter.
- (n) A person licensed under this chapter may not refuse entry into any premises for inspection as authorized by this chapter.
- (o) A person licensed under this chapter may not furnish false or fraudulent material information in any application, report, or other document required to be kept by this chapter or willfully make any false statement in any prescription, order, report, or record required by this chapter.
- (8) (a) (i) Any person licensed under this chapter who is found by the division to have violated any of the provisions of Subsections (7)(k) through (7)(o) or Subsection (10) is subject to a penalty not to exceed \$5,000. The division shall determine the procedure for adjudication of any violations in accordance with Sections 58-1-106 and 58-1-108.

1392	(ii) The division shall deposit all penalties collected under Subsection (8)(a)(i) in the
1393	General Fund as a dedicated credit to be used by the division under Subsection 58-37f-502(1).
1394	(b) Any person who knowingly and intentionally violates Subsections (7)(h) through
1395	(7)(j) or Subsection (10) is:
1396	(i) upon first conviction, guilty of a class B misdemeanor;
1397	(ii) upon second conviction, guilty of a class A misdemeanor; and
1398	(iii) on third or subsequent conviction, guilty of a third degree felony.
1399	(c) Any person who knowingly and intentionally violates Subsections (7)(k) through
1400	(7)(o) shall upon conviction be guilty of a third degree felony.
1401	(9) Any information communicated to any licensed practitioner in an attempt to
1402	unlawfully procure, or to procure the administration of, a controlled substance is not considered
1403	to be a privileged communication.
1404	(10) A person holding a valid license under this chapter who is engaged in medical
1405	research may produce, possess, or administer, but may not prescribe or dispense, a controlled
1406	substance listed in Section 58-37-4.2.
1407	Section 8. Section 58-37-8 is amended to read:
1408	58-37-8. Prohibited acts Penalties.
1409	(1) Prohibited acts A Penalties:
1410	(a) Except as authorized by this chapter, it is unlawful for any person to knowingly and
1411	intentionally:
1412	(i) produce, manufacture, or dispense, or to possess with intent to produce,
1413	manufacture, or dispense, a controlled or counterfeit substance;
1414	(ii) distribute a controlled or counterfeit substance, or to agree, consent, offer, or
1415	arrange to distribute a controlled or counterfeit substance;
1416	(iii) possess a controlled or counterfeit substance with intent to distribute; or
1417	(iv) engage in a continuing criminal enterprise where:
1418	(A) the person participates, directs, or engages in conduct which results in any
1419	violation of any provision of Title 58, Chapters 37, 37a, 37b, 37c, or 37d that is a felony; and
1420	(B) the violation is a part of a continuing series of two or more violations of Title 58,
1421	Chapters 37, 37a, 37b, 37c, or 37d on separate occasions that are undertaken in concert with
1422	five or more persons with respect to whom the person occupies a position of organizer,

supervisor, or any other position of management.

- (b) Any person convicted of violating Subsection (1)(a) with respect to:
- (i) a substance or a counterfeit of a substance classified in Schedule I or II, a controlled substance analog, or gammahydroxybutyric acid as listed in Schedule III is guilty of a second degree felony and upon a second or subsequent conviction is guilty of a first degree felony;
- (ii) a substance or a counterfeit of a substance classified in Schedule III or IV, or marijuana, or a substance listed in Section 58-37-4.2 is guilty of a third degree felony, and upon a second or subsequent conviction is guilty of a second degree felony; or
- (iii) a substance or a counterfeit of a substance classified in Schedule V is guilty of a class A misdemeanor and upon a second or subsequent conviction is guilty of a third degree felony.
- (c) Any person who has been convicted of a violation of Subsection (1)(a)(ii) or (iii) may be sentenced to imprisonment for an indeterminate term as provided by law, but if the trier of fact finds a firearm as defined in Section 76-10-501 was used, carried, or possessed on his person or in his immediate possession during the commission or in furtherance of the offense, the court shall additionally sentence the person convicted for a term of one year to run consecutively and not concurrently; and the court may additionally sentence the person convicted for an indeterminate term not to exceed five years to run consecutively and not concurrently.
- (d) Any person convicted of violating Subsection (1)(a)(iv) is guilty of a first degree felony punishable by imprisonment for an indeterminate term of not less than seven years and which may be for life. Imposition or execution of the sentence may not be suspended, and the person is not eligible for probation.
 - (2) Prohibited acts B -- Penalties:
 - (a) It is unlawful:
- (i) for any person knowingly and intentionally to possess or use a controlled substance analog or a controlled substance, unless it was obtained under a valid prescription or order, directly from a practitioner while acting in the course of the person's professional practice, or as otherwise authorized by this chapter;
- (ii) for any owner, tenant, licensee, or person in control of any building, room, tenement, vehicle, boat, aircraft, or other place knowingly and intentionally to permit them to

be occupied by persons unlawfully possessing, using, or distributing controlled substances in any of those locations; or

(iii) for any person knowingly and intentionally to possess an altered or forged prescription or written order for a controlled substance.

- (b) Any person convicted of violating Subsection (2)(a)(i) with respect to:
- (i) marijuana, if the amount is 100 pounds or more, is guilty of a second degree felony;
- (ii) a substance classified in Schedule I or II, marijuana, if the amount is more than 16 ounces, but less than 100 pounds, or a controlled substance analog, is guilty of a third degree felony; or
 - (iii) marijuana, if the marijuana is not in the form of an extracted resin from any part of the plant, and the amount is more than one ounce but less than 16 ounces, is guilty of a class A misdemeanor.
 - (c) Upon a person's conviction of a violation of this Subsection (2) subsequent to a conviction under Subsection (1)(a), that person shall be sentenced to a one degree greater penalty than provided in this Subsection (2).
 - (d) Any person who violates Subsection (2)(a)(i) with respect to all other controlled substances not included in Subsection (2)(b)(i), (ii), or (iii), including <u>a substance listed in Section 58-37-4.2</u>, or less than one ounce of marijuana, is guilty of a class B misdemeanor. Upon a second conviction the person is guilty of a class A misdemeanor, and upon a third or subsequent conviction the person is guilty of a third degree felony.
 - (e) Any person convicted of violating Subsection (2)(a)(i) while inside the exterior boundaries of property occupied by any correctional facility as defined in Section 64-13-1 or any public jail or other place of confinement shall be sentenced to a penalty one degree greater than provided in Subsection (2)(b), and if the conviction is with respect to controlled substances as listed in:
 - (i) Subsection (2)(b), the person may be sentenced to imprisonment for an indeterminate term as provided by law, and:
 - (A) the court shall additionally sentence the person convicted to a term of one year to run consecutively and not concurrently; and
- 1483 (B) the court may additionally sentence the person convicted for an indeterminate term 1484 not to exceed five years to run consecutively and not concurrently; and

1485	(ii) Subsection (2)(d), the person may be sentenced to imprisonment for an
1486	indeterminate term as provided by law, and the court shall additionally sentence the person
1487	convicted to a term of six months to run consecutively and not concurrently.
1488	(f) Any person convicted of violating Subsection (2)(a)(ii) or [(2)(a)](iii) is:
1489	(i) on a first conviction, guilty of a class B misdemeanor;
1490	(ii) on a second conviction, guilty of a class A misdemeanor; and
1491	(iii) on a third or subsequent conviction, guilty of a third degree felony.
1492	(g) A person is subject to the penalties under Subsection (2)(h) who, in an offense not
1493	amounting to a violation of Section 76-5-207:
1494	(i) violates Subsection (2)(a)(i) by knowingly and intentionally having in the person's
1495	body any measurable amount of a controlled substance; and
1496	(ii) operates a motor vehicle as defined in Section 76-5-207 in a negligent manner,
1497	causing serious bodily injury as defined in Section 76-1-601 or the death of another.
1498	(h) A person who violates Subsection (2)(g) by having in the person's body:
1499	(i) a controlled substance classified under Schedule I, other than those described in
1500	Subsection (2)(h)(ii), or a controlled substance classified under Schedule II is guilty of a second
1501	degree felony;
1502	(ii) marijuana, tetrahydrocannabinols, or equivalents described in Subsection
1503	58-37-4(2)(a)(iii)(S) or (AA), or a substance listed in Section 58-37-4.2 is guilty of a third
1504	degree felony; or
1505	(iii) any controlled substance classified under Schedules III, IV, or V is guilty of a class
1506	A misdemeanor.
1507	(i) A person is guilty of a separate offense for each victim suffering serious bodily
1508	injury or death as a result of the person's negligent driving in violation of Subsection
1509	58-37-8(2)(g) whether or not the injuries arise from the same episode of driving.
1510	(3) Prohibited acts C Penalties:
1511	(a) It is unlawful for any person knowingly and intentionally:
1512	(i) to use in the course of the manufacture or distribution of a controlled substance a
1513	license number which is fictitious, revoked, suspended, or issued to another person or, for the
1514	purpose of obtaining a controlled substance, to assume the title of, or represent oneself to be, a
1515	manufacturer, wholesaler, apothecary, physician, dentist, veterinarian, or other authorized

1516	person
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(ii) to acquire or obtain possession of, to procure or attempt to procure the administration of, to obtain a prescription for, to prescribe or dispense to any person known to be attempting to acquire or obtain possession of, or to procure the administration of any controlled substance by misrepresentation or failure by the person to disclose receiving any controlled substance from another source, fraud, forgery, deception, subterfuge, alteration of a prescription or written order for a controlled substance, or the use of a false name or address;

- (iii) to make any false or forged prescription or written order for a controlled substance, or to utter the same, or to alter any prescription or written order issued or written under the terms of this chapter; or
- (iv) to make, distribute, or possess any punch, die, plate, stone, or other thing designed to print, imprint, or reproduce the trademark, trade name, or other identifying mark, imprint, or device of another or any likeness of any of the foregoing upon any drug or container or labeling so as to render any drug a counterfeit controlled substance.
- (b) Any person convicted of violating Subsection (3)(a) is guilty of a third degree felony.
 - (4) Prohibited acts D -- Penalties:
- (a) Notwithstanding other provisions of this section, a person not authorized under this chapter who commits any act declared to be unlawful under this section, Title 58, Chapter 37a, Utah Drug Paraphernalia Act, or under Title 58, Chapter 37b, Imitation Controlled Substances Act, is upon conviction subject to the penalties and classifications under this Subsection (4) if the trier of fact finds the act is committed:
- (i) in a public or private elementary or secondary school or on the grounds of any of those schools;
- (ii) in a public or private vocational school or postsecondary institution or on the grounds of any of those schools or institutions;
- (iii) in those portions of any building, park, stadium, or other structure or grounds which are, at the time of the act, being used for an activity sponsored by or through a school or institution under Subsections (4)(a)(i) and (ii);
 - (iv) in or on the grounds of a preschool or child-care facility;
- (v) in a public park, amusement park, arcade, or recreation center;

1547	(vi) in or on the grounds of a house of worship as defined in Section 76-10-501;
1548	(vii) in a shopping mall, sports facility, stadium, arena, theater, movie house,
1549	playhouse, or parking lot or structure adjacent thereto;
1550	(viii) in or on the grounds of a library;
1551	(ix) within any area that is within 1,000 feet of any structure, facility, or grounds
1552	included in Subsections (4)(a)(i), (ii), (iv), (vi), and (vii);
1553	(x) in the presence of a person younger than 18 years of age, regardless of where the act
1554	occurs; or
1555	(xi) for the purpose of facilitating, arranging, or causing the transport, delivery, or
1556	distribution of a substance in violation of this section to an inmate or on the grounds of any
1557	correctional facility as defined in Section 76-8-311.3.
1558	(b) (i) A person convicted under this Subsection (4) is guilty of a first degree felony
1559	and shall be imprisoned for a term of not less than five years if the penalty that would
1560	otherwise have been established but for this Subsection (4) would have been a first degree
1561	felony.
1562	(ii) Imposition or execution of the sentence may not be suspended, and the person is
1563	not eligible for probation.
1564	(c) If the classification that would otherwise have been established would have been
1565	less than a first degree felony but for this Subsection (4), a person convicted under this
1566	Subsection (4) is guilty of one degree more than the maximum penalty prescribed for that
1567	offense. This Subsection (4)(c) does not apply to a violation of Subsection (2)(g).
1568	(d) (i) If the violation is of Subsection (4)(a)(xi):
1569	(A) the person may be sentenced to imprisonment for an indeterminate term as
1570	provided by law, and the court shall additionally sentence the person convicted for a term of
1571	one year to run consecutively and not concurrently; and
1572	(B) the court may additionally sentence the person convicted for an indeterminate term
1573	not to exceed five years to run consecutively and not concurrently; and
1574	(ii) the penalties under this Subsection (4)(d) apply also to any person who, acting with
1575	the mental state required for the commission of an offense, directly or indirectly solicits,
1576	requests, commands, coerces, encourages, or intentionally aids another person to commit a

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violation of Subsection (4)(a)(xi).

(e) It is not a defense to a prosecution under this Subsection (4) that the actor mistakenly believed the individual to be 18 years of age or older at the time of the offense or was unaware of the individual's true age; nor that the actor mistakenly believed that the location where the act occurred was not as described in Subsection (4)(a) or was unaware that the location where the act occurred was as described in Subsection (4)(a).

(5) Any violation of this chapter for which no penalty is specified is a class B misdemeanor.

- (6) For purposes of penalty enhancement under Subsections (1)(b) and (2)(c), a plea of guilty or no contest to a violation of this section which is held in abeyance under Title 77, Chapter 2a, Pleas in Abeyance, is the equivalent of a conviction, even if the charge has been subsequently reduced or dismissed in accordance with the plea in abeyance agreement.
- (7) A person may be charged and sentenced for a violation of this section, notwithstanding a charge and sentence for a violation of any other section of this chapter.
- (8) (a) Any penalty imposed for violation of this section is in addition to, and not in lieu of, any civil or administrative penalty or sanction authorized by law.
- (b) Where violation of this chapter violates a federal law or the law of another state, conviction or acquittal under federal law or the law of another state for the same act is a bar to prosecution in this state.
- (9) In any prosecution for a violation of this chapter, evidence or proof which shows a person or persons produced, manufactured, possessed, distributed, or dispensed a controlled substance or substances, is prima facie evidence that the person or persons did so with knowledge of the character of the substance or substances.
- (10) This section does not prohibit a veterinarian, in good faith and in the course of the veterinarian's professional practice only and not for humans, from prescribing, dispensing, or administering controlled substances or from causing the substances to be administered by an assistant or orderly under the veterinarian's direction and supervision.
 - (11) Civil or criminal liability may not be imposed under this section on:
- (a) any person registered under this chapter who manufactures, distributes, or possesses an imitation controlled substance for use as a placebo or investigational new drug by a registered practitioner in the ordinary course of professional practice or research; or
 - (b) any law enforcement officer acting in the course and legitimate scope of the

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(12) (a) Civil or criminal liability may not be imposed under this section on any Indian, as defined in Subsection 58-37-2(1)(v), who uses, possesses, or transports peyote for bona fide traditional ceremonial purposes in connection with the practice of a traditional Indian religion as defined in Subsection 58-37-2(1)(w).

- (b) In a prosecution alleging violation of this section regarding peyote as defined in Subsection 58-37-4(2)(a)(iii)(V), it is an affirmative defense that the peyote was used, possessed, or transported by an Indian for bona fide traditional ceremonial purposes in connection with the practice of a traditional Indian religion.
- (c) (i) The defendant shall provide written notice of intent to claim an affirmative defense under this Subsection (12) as soon as practicable, but not later than 10 days prior to trial.
 - (ii) The notice shall include the specific claims of the affirmative defense.
- (iii) The court may waive the notice requirement in the interest of justice for good cause shown, if the prosecutor is not unfairly prejudiced by the lack of timely notice.
- (d) The defendant shall establish the affirmative defense under this Subsection (12) by a preponderance of the evidence. If the defense is established, it is a complete defense to the charges.
- (13) (a) It is an affirmative defense that the person produced, possessed, or administered a controlled substance listed in Section 58-37-4.2 if the person:
- (i) was engaged in medical research; and
- (ii) was a holder of a valid license to possess controlled substances under Section
 58-37-6.
- 1632 (b) It is not a defense under Subsection 58-37-8(13)(a) that the person prescribed or
 1633 dispensed a controlled substance listed in 58-37-4.2.
 - (14) It is an affirmative defense that the person possessed, in the person's body, a controlled substance listed in Section 58-37-4.2 if:
 - (a) the person was the subject of medical research conducted by a holder of a valid license to possess controlled substances under Section 58-37-6; and
- 1638 (b) the substance was administered to the person by the medical researcher.
- [(13)] (15) If any provision of this chapter, or the application of any provision to any

1640	person or circumstances, is held invalid, the remainder of this chapter shall be given effect
1641	without the invalid provision or application.
1642	(16) A legislative body of a political subdivision may not enact an ordinance that is
1643	less restrictive than any provision of this chapter.
1644	Section 9. Section 58-38a-203 is amended to read:
1645	58-38a-203. Duties of the committee.
1646	(1) The committee serves as a consultative and advisory body to the Legislature
1647	regarding:
1648	(a) the movement of a controlled substance from one schedule or list to another;
1649	(b) the removal of a controlled substance from any schedule or list; and
1650	(c) the designation of a substance as a controlled substance and the placement of the
1651	substance in a designated schedule <u>or list</u> .
1652	(2) On or before September 30 of each year, the committee shall submit to the Health
1653	and Human Services Interim Committee a written report:
1654	(a) [listing] describing any substances recommended by the committee for scheduling,
1655	rescheduling, <u>listing</u> , or deletion from the schedules <u>or list</u> by the Legislature; and
1656	(b) stating the reasons for the recommendation.
1657	(3) In advising the Legislature regarding the need to add, delete, <u>relist</u> , or reschedule a
1658	substance, the committee shall consider:
1659	(a) the actual or probable abuse of the substance, including:
1660	(i) the history and current pattern of abuse both in Utah and in other states;
1661	(ii) the scope, duration, and significance of abuse;
1662	(iii) the degree of actual or probable detriment to public health which may result from
1663	abuse of the substance; and
1664	(iv) the probable physical and social impact of widespread abuse of the substance;
1665	(b) the biomedical hazard of the substance, including:
1666	(i) its pharmacology, including the effects and modifiers of the effects of the substance;
1667	(ii) its toxicology, acute and chronic toxicity, interaction with other substances,
1668	whether controlled or not, and the degree to which it may cause psychological or physiological
1669	dependence; and
1670	(iii) the risk to public health and the particular susceptibility of segments of the

1671	population;
1672	(c) whether the substance is an immediate precursor, as defined in Section 58-37-2, of
1673	a substance that is currently a controlled substance;
1674	(d) the current state of scientific knowledge regarding the substance, including whether
1675	there is any acceptable means to safely use the substance under medical supervision;
1676	(e) the relationship between the use of the substance and criminal activity, including
1677	whether:
1678	(i) persons engaged in illicit trafficking of the substance are also engaged in other
1679	criminal activity;
1680	(ii) the nature and relative profitability of manufacturing or delivering the substance
1681	encourages illicit trafficking in the substance;
1682	(iii) the commission of other crimes is one of the recognized effects of abuse of the
1683	substance; and
1684	(iv) addiction to the substance relates to the commission of crimes to facilitate the
1685	continued use of the substance;
1686	(f) whether the substance has been scheduled by other states; and
1687	(g) whether the substance has any accepted medical use in treatment in the United
1688	States.
1689	(4) The committee's duties under this chapter do not include tobacco products as
1690	defined in Section 59-14-102 or alcoholic beverages as defined in Section 32A-1-105.
1691	Section 10. Section 58-38a-204 is amended to read:
1692	58-38a-204. Guidelines for scheduling or listing drugs.
1693	(1) (a) The committee shall recommend placement of a substance in Schedule I if it
1694	finds:
1695	(i) that the substance has high potential for abuse; and
1696	(ii) that an accepted standard has not been established for safe use in treatment for
1697	medical purposes.
1698	(b) The committee may recommend placement of a substance in Schedule I under
1699	Section 58-37-4 if it finds that the substance is classified as a controlled substance in Schedule
1700	I under federal law.

(2) (a) The committee shall recommend placement of a substance in Schedule II if it

1702	finds that:
1703	(i) the substance has high potential for abuse;
1704	(ii) the substance has a currently accepted medical use in treatment in the United
1705	States, or a currently accepted medical use subject to severe restrictions; and
1706	(iii) the abuse of the substance may lead to severe psychological or physiological
1707	dependence.
1708	(b) The committee may recommend placement of a substance in Schedule II if it finds
1709	that the substance is classified as a controlled substance in Schedule II under federal law.
1710	(3) (a) The committee shall recommend placement of a substance in Schedule III if it
1711	finds that:
1712	(i) the substance has a potential for abuse that is less than the potential for substances
1713	listed in Schedules I and II;
1714	(ii) the substance has a currently accepted medical use in treatment in the United
1715	States; and
1716	(iii) abuse of the substance may lead to moderate or low physiological dependence or
1717	high psychological dependence.
1718	(b) The committee may recommend placement of a substance in Schedule III if it finds
1719	that the substance is classified as a controlled substance in Schedule III under federal law.
1720	(4) (a) The committee shall recommend placement of a substance in Schedule IV if it
1721	finds that:
1722	(i) the substance has a low potential for abuse relative to substances in Schedule III;
1723	(ii) the substance has currently accepted medical use in treatment in the United States;
1724	and
1725	(iii) abuse of the substance may lead to limited physiological dependence or
1726	psychological dependence relative to the substances in Schedule III.
1727	(b) The committee may recommend placement of a substance in Schedule IV if it find
1728	that the substance is classified as a controlled substance in Schedule IV under federal law.
1729	(5) (a) The committee shall recommend placement of a substance in Schedule V if it
1730	finds that:

(i) the substance has low potential for abuse relative to the controlled substances listed

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in Schedule IV;

1733	(11) the substance has currently accepted medical use in treatment in the United States;
1734	and
1735	(iii) the substance has limited physiological dependence or psychological dependence
1736	liability relative to the controlled substances listed in Schedule IV.
1737	(b) The committee may recommend placement of a substance in Schedule V under this
1738	chapter if it finds that the substance is classified as a controlled substance in Schedule V under
1739	federal law.
1740	(6) The committee may recommend placement of a substance on a controlled substance
1741	list if it finds that the substance has a potential for abuse and that an accepted standard has not
1742	been established for safe use in treatment for medical purposes.
1743	Section 11. Effective date.
1744	If approved by two-thirds of all the members elected to each house, this bill takes effect
1745	upon approval by the governor, or the day following the constitutional time limit of Utah
1746	Constitution Article VII, Section 8, without the governor's signature, or in the case of a veto,
1747	the date of veto override, except that the amendments to Section 58-37-2 (Effective 07/01/11)
1748	take effect on July 1, 2011.

Legislative Review Note as of 11-18-10 10:26 AM

Office of Legislative Research and General Counsel

FISCAL NOTE

H.B. 23, 2011 General Session

SHORT TITLE: Controlled Substance Modifications

SPONSOR: Froerer, G. STATE OF UTAH

STATE GOVERNMENT (UCA 36-12-13(2)(b))

Enactment of this bill likely will not materially impact the state budget. To the extent that additional convictions occur as a result of this legislatiion, the Department of Public Safety would collect additional driving reinstatement fee revenue. DPS reports that such additional revenue will cover the costs of reinstating those driving licenses.

LOCAL GOVERNMENTS (UCA 36-12-13(2)(c))

Enactment of this bill likely will not result in direct, measurable costs and/or benefits for local governments.

DIRECT EXPENDITURES BY UTAH RESIDENTS AND BUSINESSES (UCA 36-12-13(2)(d)) Individuals convicted of a violation under this legislation would pay the existing \$65 fee to reinstate their driving license.

1/22/2011, 01:30 PM, Lead Analyst: Frandsen, R./Attorney: SCA

Office of the Legislative Fiscal Analyst