12-23 11:17 1 **Kratom Revisions** 2026 GENERAL SESSION STATE OF UTAH Chief Sponsor: Evan J. Vickers House Sponsor: Jake Sawyer 2

## 3 LONG TITLE 4 **Committee Note:** 5 The Business and Labor Interim Committee recommended this bill. 6 Legislative Vote: 16 voting for 0 voting against 4 absent 7 **General Description:** 8 This bill amends provisions related to kratom. 9 **Highlighted Provisions:** This bill: 10 11 defines terms; 12 schedules 7-hydroxymitragynine, including synthetics, if the 7-hydroxymitragynine 13 concentration exceeds a certain percentage as a schedule I controlled substance; and 14 schedules Mitragynine pseudoindoxyl, including synthetics. 15 Money Appropriated in this Bill: 16 None 17 **Other Special Clauses:** 18 This bill provides a special effective date. **Utah Code Sections Affected:** 19 20 **AMENDS:** 21 **4-45-104**, as enacted by Laws of Utah 2019, Chapter 329 22 **58-37-2**, as last amended by Laws of Utah 2025, Chapter 396 23 58-37-4, as last amended by Laws of Utah 2025, Chapter 216 24

Be it enacted by the Legislature of the state of Utah:

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Section 1. Section **4-45-104** is amended to read:

4-45-104. Kratom processor requirements -- Criminal penalty.

- (1) A kratom processor may not prepare, distribute, sell, or offer for sale a kratom product:
  - (a) that is mixed or packed with a nonkratom substance that affects the quality or strength of the kratom product to such a degree as to render the kratom product

31	injurious to a consumer;
32	(b) that contains a poisonous or otherwise deleterious nonkratom ingredient, including a
33	controlled substance as defined in Section 58-37-2;
34	(c) [containing a level of 7-hydroxymitragynine in the alkaloid fraction that is greater
35	than 2% of the alkaloid composition of the kratom product] that would be considered
36	a controlled substance under Section 58-37-4;
37	(d) containing a synthetic alkaloid, including synthetic mitragynine, synthetic
38	7-hydroxymitragynine, or any other synthetically derived compound of the kratom
39	plant; or
40	(e) that does not include a product label on the kratom product packaging that states the
41	amount of mitragynine and 7-hydroxymitragynine contained in the packaged kratom
42	product.
43	(2) A kratom processor who violates Subsection (1) is guilty of a class C misdemeanor for
44	each violation.
45	(3) A kratom processor does not violate Subsection (1) if the kratom processor shows by a
46	preponderance of the evidence that the kratom processor relied in good faith upon the
47	representation of a manufacturer, processor, packer, or distributor of food represented to
48	be a kratom product.
49	(4) A kratom processor may not prepare, distribute, sell, or offer for sale a kratom product
50	that is not registered with the department in accordance with this chapter.
51	(5) A kratom processor shall register as a food establishment in accordance with Section
52	4-5-301.
53	Section 2. Section <b>58-37-2</b> is amended to read:
54	58-37-2 . Definitions.
55	(1) As used in this chapter:
56	(a) "Administer" means the direct application of a controlled substance, whether by
57	injection, inhalation, ingestion, or any other means, to the body of a patient or
58	research subject by:
59	(i) a practitioner or, in the practitioner's presence, by the practitioner's authorized
60	agent; or
61	(ii) the patient or research subject at the direction and in the presence of the
62	practitioner.
63	(b) "Agent" means an authorized person who acts on behalf of or at the direction of a
64	manufacturer, distributor, or practitioner but does not include a motor carrier, public

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 this chapter, 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;
  - (B) included in Schedules I, II, III, IV, or V of the federal Controlled Substances Act, Title II, P.L. 91-513;
  - (C) that is a controlled substance analog; or
  - (D) 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
    - (C) dietary supplements, vitamins, minerals, herbs, or other similar substances including concentrates or extracts, which:

99	(I) are not otherwise regulated by law; and
100	(II) may contain naturally occurring amounts of chemical or substances listed
101	in this chapter, or in rules adopted pursuant to Title 63G, Chapter 3, Utah
102	Administrative Rulemaking Act.
103	(g)(i) "Controlled substance analog" means:
104	(A) a substance the chemical structure of which is substantially similar to the
105	chemical structure of a controlled substance listed in Schedules I and II of
106	Section 58-37-4, a substance listed in Section 58-37-4.2, or in Schedules I and
107	II of the federal Controlled Substances Act, Title II, P.L. 91-513;
108	(B) a substance that has a stimulant, depressant, or hallucinogenic effect on the
109	central nervous system substantially similar to the stimulant, depressant, or
110	hallucinogenic effect on the central nervous system of controlled substances
111	listed in Schedules I and II of Section 58-37-4, substances listed in Section
112	58-37-4.2, or substances listed in Schedules I and II of the federal Controlled
113	Substances Act, Title II, P.L. 91-513; or
114	(C) A substance that, with respect to a particular individual, is represented or
115	intended to have a stimulant, depressant, or hallucinogenic effect on the central
116	nervous system substantially similar to the stimulant, depressant, or
117	hallucinogenic effect on the central nervous system of controlled substances
118	listed in Schedules I and II of Section 58-37-4, substances listed in Section
119	58-37-4.2, or substances listed in Schedules I and II of the federal Controlled
120	Substances Act, Title II, P.L. 91-513.
121	(ii) "Controlled substance analog" does not include:
122	(A) a controlled substance currently scheduled in Schedules I through V of
123	Section 58-37-4;
124	(B) a substance for which there is an approved new drug application;
125	(C) a substance with respect to which an exemption is in effect for investigational
126	use by a particular person under Section 505 of the Food, Drug, and Cosmetic
127	Act, 21 U.S.C. 355, to the extent the conduct with respect to the substance is
128	permitted by the exemption;
129	(D) any substance to the extent not intended for human consumption before an
130	exemption takes effect with respect to the substance;
131	(E) any drug intended for lawful use in the diagnosis, cure, mitigation, treatment,
132	or prevention of disease in man or other animals, which contains ephedrine.

133	pseudoephedrine, norpseudoephedrine, or phenylpropanolamine if the drug is
134	lawfully purchased, sold, transferred, or furnished as an over-the-counter
135	medication without prescription; or
136	(F) dietary supplements, vitamins, minerals, herbs, or other similar substances
137	including concentrates or extracts, which are not otherwise regulated by law,
138	which may contain naturally occurring amounts of chemical or substances
139	listed in this chapter, or in rules adopted pursuant to Title 63G, Chapter 3, Utah
140	Administrative Rulemaking Act.
141	(h)(i) "Conviction" means a determination of guilt by verdict, whether jury or bench,
142	or plea, whether guilty or no contest, for any offense proscribed by:
143	(A) this chapter;
144	(B) Chapter 37a, Utah Drug Paraphernalia Act;
145	(C) Chapter 37b, Imitation Controlled Substances Act;
146	(D) Chapter 37c, Utah Controlled Substance Precursor Act; or
147	(E) Chapter 37d, Clandestine Drug Lab Act; or
148	(ii) for any offense under the laws of the United States and any other state which, if
149	committed in this state, would be an offense under:
150	(A) this chapter;
151	(B) Chapter 37a, Utah Drug Paraphernalia Act;
152	(C) Chapter 37b, Imitation Controlled Substances Act;
153	(D) Chapter 37c, Utah Controlled Substance Precursor Act; or
154	(E) Chapter 37d, Clandestine Drug Lab Act.
155	(i) "Counterfeit substance" means:
156	(i) any controlled substance or container or labeling of any controlled substance that:
157	(A) without authorization bears the trademark, trade name, or other identifying
158	mark, imprint, number, device, or any likeness of them, of a manufacturer,
159	distributor, or dispenser other than the person or persons who in fact
160	manufactured, distributed, or dispensed the substance which falsely purports to
161	be a controlled substance distributed by any other manufacturer, distributor, or
162	dispenser; and
163	(B) a reasonable person would believe to be a controlled substance distributed by
164	an authorized manufacturer, distributor, or dispenser based on the appearance
165	of the substance as described under Subsection (1)(i)(i)(A) or the appearance of
166	the container of that controlled substance; or

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167	(ii) any substance other than under Subsection (1)(i)(i) that:
168	(A) is falsely represented to be any legally or illegally manufactured controlled
169	substance; and
170	(B) a reasonable person would believe to be a legal or illegal controlled substance
171	(j) "Deliver" or "delivery" means the actual, constructive, or attempted transfer of a
172	controlled substance or a listed chemical, whether or not an agency relationship exists.
173	(k) "Department" means the Department of Commerce.
174	(l) "Depressant or stimulant substance" means:
175	(i) a drug which contains any quantity of barbituric acid or any of the salts of
176	barbituric acid;
177	(ii) a drug which contains any quantity of:
178	(A) amphetamine or any of its optical isomers;
179	(B) any salt of amphetamine or any salt of an optical isomer of amphetamine; or
180	(C) any substance which the Secretary of Health and Human Services or the
181	Attorney General of the United States after investigation has found and by
182	regulation designated habit-forming because of its stimulant effect on the
183	central nervous system;
184	(iii) lysergic acid diethylamide; or
185	(iv) any drug which contains any quantity of a substance which the Secretary of
186	Health and Human Services or the Attorney General of the United States after
187	investigation has found to have, and by regulation designated as having, a
188	potential for abuse because of its depressant or stimulant effect on the central
189	nervous system or its hallucinogenic effect.
190	(m) "Dispense" means the delivery of a controlled substance by a pharmacist to an
191	ultimate user pursuant to the lawful order or prescription of a practitioner, and
192	includes distributing to, leaving with, giving away, or disposing of that substance as
193	well as the packaging, labeling, or compounding necessary to prepare the substance
194	for delivery.
195	(n) "Dispenser" means a pharmacist who dispenses a controlled substance.
196	(o) "Distribute" means to deliver other than by administering or dispensing a controlled
197	substance or a listed chemical.
198	(p) "Distributor" means a person who distributes controlled substances.
199	(q) "Division" means the Division of Professional Licensing created in Section 58-1-103.
200	(r)(i) "Drug" means:

201	(A) a substance recognized in the official United States Pharmacopoeia, Official
202	Homeopathic Pharmacopoeia of the United States, or Official National
203	Formulary, or any supplement to any of them, intended for use in the
204	diagnosis, cure, mitigation, treatment, or prevention of disease in humans or
205	animals;
206	(B) a substance that is required by any applicable federal or state law or rule to be
207	dispensed by prescription only or is restricted to administration by practitioners
208	only;
209	(C) a substance other than food intended to affect the structure or any function of
210	the body of humans or other animals; and
211	(D) substances intended for use as a component of any substance specified in
212	Subsections $(1)(r)(i)(A)$ , $(B)$ , and $(C)$ .
213	(ii) "Drug" does not include dietary supplements.
214	(iii) "Drug" includes a food intended for human consumption that intentionally
215	contains a vaccine or vaccine material as provided in Section 4-5-107.
216	(s) "Drug dependent person" means any individual who unlawfully and habitually uses
217	any controlled substance to endanger the public morals, health, safety, or welfare, or
218	who is so dependent upon the use of controlled substances as to have lost the power
219	of self-control with reference to the individual's dependency.
220	(t)(i) "Food" means:
221	(A) any nutrient or substance of plant, mineral, or animal origin other than a drug
222	as specified in this chapter, and normally ingested by human beings; and
223	(B) foods for special dietary uses as exist by reason of a physical, physiological,
224	pathological, or other condition including the conditions of disease,
225	convalescence, pregnancy, lactation, allergy, hypersensitivity to food,
226	underweight, and overweight; uses for supplying a particular dietary need
227	which exist by reason of age including the ages of infancy and childbirth, and
228	also uses for supplementing and for fortifying the ordinary or unusual diet with
229	any vitamin, mineral, or other dietary property for use of a food.
230	(ii) Any particular use of a food is a special dietary use regardless of the nutritional
231	purposes.
232	(u) "Immediate precursor" means a substance which the Attorney General of the United
233	States has found to be, and by regulation designated as being, the principal compound
234	used or produced primarily for use in the manufacture of a controlled substance, or

235	which is an immediate chemical intermediary used or likely to be used in the
236	manufacture of a controlled substance, the control of which is necessary to prevent,
237	curtail, or limit the manufacture of the controlled substance.
238	(v) "Indian" means a member of an Indian tribe.
239	(w) "Indian religion" means a religion:
240	(i) the origin and interpretation of which is from within a traditional Indian culture or
241	community; and
242	(ii) that is practiced by Indians.
243	(x) "Indian tribe" means any tribe, band, nation, pueblo, or other organized group or
244	community of Indians, including any Alaska Native village, which is legally
245	recognized as eligible for and is consistent with the special programs, services, and
246	entitlements provided by the United States to Indians because of their status as
247	Indians.
248	(y) "Manufacture" means the production, preparation, propagation, compounding, or
249	processing of a controlled substance, either directly or indirectly by extraction from
250	substances of natural origin, or independently by means of chemical synthesis or by a
251	combination of extraction and chemical synthesis.
252	(z) "Manufacturer" includes any person who packages, repackages, or labels any
253	container of any controlled substance, except pharmacists who dispense or compound
254	prescription orders for delivery to the ultimate consumer.
255	(aa)(i) "Marijuana" means all species of the genus cannabis and all parts of the genus,
256	whether growing or not, including:
257	(A) seeds;
258	(B) resin extracted from any part of the plant, including the resin extracted from
259	the mature stalks;
260	(C) every compound, manufacture, salt, derivative, mixture, or preparation of the
261	plant, seeds, or resin;
262	(D) any synthetic equivalents of the substances contained in the plant cannabis
263	sativa or any other species of the genus cannabis which are chemically
264	indistinguishable and pharmacologically active; and
265	(E) any component part or cannabinoid extracted or isolated from the plant,
266	including extracted or isolated tetrahydrocannabinols.
267	(ii) "Marijuana" does not include:
268	(A) the mature stalks of the plant;

269	(B) fiber produced from the stalks;
270	(C) oil or cake made from the seeds of the plant;
271	(D) except as provided in Subsection (1)(aa)(i), any other compound,
272	manufacture, salt, derivative, mixture, or preparation of the mature stalks,
273	fiber, oil or cake;
274	(E) the sterilized seed of the plant which is incapable of germination;
275	(F) any compound, mixture, or preparation approved by the federal Food and
276	Drug Administration under the federal Food, Drug, and Cosmetic Act, 21
277	U.S.C. Sec. 301 et seq. that is not listed in a schedule of controlled substances
278	in Section 58-37-4 or in the federal Controlled Substances Act, Title II, P.L.
279	91-513; or
280	(G) transportable industrial hemp concentrate as that term is defined in Section
281	4-41-102.
282	(bb) "Money" means officially issued coin and currency of the United States or any
283	foreign country.
284	(cc) "Narcotic drug" means any of the following, whether produced directly or indirectly
285	by extraction from substances of vegetable origin, or independently by means of
286	chemical synthesis, or by a combination of extraction and chemical synthesis:
287	(i) opium, coca leaves, and opiates;
288	(ii) a compound, manufacture, salt, derivative, or preparation of opium, coca leaves,
289	or opiates;
290	(iii) opium poppy and poppy straw; or
291	(iv) a substance, and any compound, manufacture, salt, derivative, or preparation of
292	the substance, which is chemically identical with any of the substances referred to
293	in Subsection (1)(cc)(i), (ii), or (iii), except narcotic drug does not include
294	decocainized coca leaves or extracts of coca leaves which do not contain cocaine
295	or ecgonine.
296	(dd) "Negotiable instrument" means documents, containing an unconditional promise to
297	pay a sum of money, which are legally transferable to another party by endorsement
298	or delivery.
299	(ee) "Opiate" means any drug or other substance having an addiction-forming or
300	addiction-sustaining liability similar to morphine or being capable of conversion into
301	a drug having addiction-forming or addiction-sustaining liability.
302	(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 the controlled substance.
- (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
  - (ii) for a controlled substance or other prescription drug or device for use by a patient or an animal.
- (mm) "Production" means the manufacture, planting, cultivation, growing, or harvesting of a controlled substance.
- (nn) "Securities" means any stocks, bonds, notes, or other evidences of debt or of property.
- (oo) "State" means the state of Utah.

337	(pp) "Total kratom alkaloid composition" means the total amount of all alkaloids derived
338	from the Mitragyna speciosa plant.
339	[(pp)] (qq) "Ultimate user" means any person who lawfully possesses a controlled
340	substance for the person's own use, for the use of a member of the person's
341	household, or for administration to an animal owned by the person or a member of
342	the person's household.
343	(2) If a term used in this chapter is not defined, the definition and terms of Title 76, Utah
344	Criminal Code, shall apply.
345	Section 3. Section <b>58-37-4</b> is amended to read:
346	58-37-4 . Schedules of controlled substances Schedules I through $V$ Findings
347	required Specific substances included in schedules.
348	(1) There are established five schedules of controlled substances known as Schedules I, II,
349	III, IV, and V which consist of substances listed in this section.
350	(2) Schedules I, II, III, IV, and V consist of the following drugs or other substances by the
351	official name, common or usual name, chemical name, or brand name designated:
352	(a) Schedule I:
353	(i) Unless specifically excepted or unless listed in another schedule, any of the
354	following opiates, including their isomers, esters, ethers, salts, and salts of
355	isomers, esters, and ethers, when the existence of the isomers, esters, ethers, and
356	salts is possible within the specific chemical designation:
357	(A) Acetyl-alpha-methylfentanyl
358	(N-[1-(1-methyl-2-phenethyl)-4-piperidinyl]-N-phenylacetamide);
359	(B) Acetyl fentanyl: (N-(1-phenethylpiperidin-4-yl)-N-phenylacetamide);
360	(C) Acetylmethadol;
361	(D) Acryl fentanyl (N-(1-Phenethylpiperidin-4-yl)-N-phenylacrylamide);
362	(E) Allylprodine;
363	(F) Alphacetylmethadol, except levo-alphacetylmethadol also known as
364	levo-alpha-acetylmethadol, levomethadyl acetate, or LAAM;
365	(G) Alphameprodine;
366	(H) Alphamethadol;
367	(I) Alpha-methylfentanyl (N-[1-(alpha-methyl-beta-phenyl)ethyl-4-piperidyl]
368	propionanilide; 1-(1-methyl-2-phenylethyl)-4-(N-propanilido) piperidine);
369	(J) Alpha-methylthiofentanyl (N-[1-methyl-2-(2-thienyl)ethyl-4-
370	piperidinyl]-N-phenylpropanamide);

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371	(K) Benzylpiperazine;
372	(L) Benzethidine;
373	(M) Betacetylmethadol;
374	(N) Beta-hydroxyfentanyl (N-[1-(2-hydroxy-2-phenethyl)-4-
375	piperidinyl]-N-phenylpropanamide);
376	(O) Beta-hydroxy-3-methylfentanyl, other name: N-[1-(2-hydroxy-2-
377	phenethyl)-3-methyl-4-piperidinyl]-N-phenylpropanamide;
378	(P) Betameprodine;
379	(Q) Betamethadol;
380	(R) Betaprodine;
381	(S) Butyryl fentanyl (N-(1-(2-phenylethyl)-4-piperidinyl)-N-phenylbutyramide);
382	(T) Clonitazene;
383	(U) Cyclopropyl fentanyl
384	(N-(1-Phenethylpiperidin-4-yl)-N-phenylcyclopropanecarboxamide);
385	(V) Dextromoramide;
386	(W) Diampromide;
387	(X) Diethylthiambutene;
388	(Y) Difenoxin;
389	(Z) Dimenoxadol;
390	(AA) Dimepheptanol;
391	(BB) Dimethylthiambutene;
392	(CC) Dioxaphetyl butyrate;
393	(DD) Dipipanone;
394	(EE) Ethylmethylthiambutene;
395	(FF) Etizolam
396	(1-Methyl-6-o-chlorophenyl-8-ethyl-4H-s-triazolo[3,4-c] thieno[2,3-e]1, 4-diazepine);
397	(GG) Etonitazene;
398	(HH) Etoxeridine;
399	(II) Furanyl fentanyl (N-phenyl-N-[1-(2-phenylethyl)piperidin-4-yl]
400	furan-2-carboxamide);
401	(JJ) Furethidine;
402	(KK) Hydroxypethidine;
403	(LL) Ketobemidone;
404	(MM) Levomoramide;

405	(NN) Levophenacylmorphan;
406	(OO) Methoxyacetyl fentanyl
407	(2-Methoxy-N-(1-phenylethylpiperidinyl-4-yl)-N-acetamide);
408	(PP) Morpheridine;
409	(QQ) MPPP (1-methyl-4-phenyl-4-propionoxypiperidine);
410	(RR) Noracymethadol;
411	(SS) Norlevorphanol;
412	(TT) Normethadone;
413	(UU) Norpipanone;
414	(VV) Para-fluorofentanyl (N-(4-fluorophenyl)-N-[1-(2-phenethyl)-4- piperidinyl]
415	propanamide);
416	(WW) Para-fluoroisobutyryl fentanyl
417	(N-(4-Fluorophenyl)-N-(1-phenethylpiperidin-4-yl)isobutyramide);
418	(XX) PEPAP (1-(-2-phenethyl)-4-phenyl-4-acetoxypiperidine);
419	(YY) Phenadoxone;
420	(ZZ) Phenampromide;
421	(AAA) Phenibut;
422	(BBB) Phenomorphan;
423	(CCC) Phenoperidine;
424	(DDD) Piritramide;
425	(EEE) Proheptazine;
426	(FFF) Properidine;
427	(GGG) Propiram;
428	(HHH) Racemoramide;
429	(III) Tetrahydrofuran fentanyl
430	(N-(1-Phenethylpiperidin-4-yl)-N-phenyltetrahydrofuran-2-carboxamide);
431	(JJJ) Thiofentanyl (N-phenyl-N-[1-(2-thienyl)ethyl-4-piperidinyl]- propanamide;
432	(KKK) Tianeptine;
433	(LLL) Tilidine;
434	(MMM) Trimeperidine;
435	(NNN) 3-methylfentanyl, including the optical and geometric isomers
436	(N-[3-methyl-1-(2-phenylethyl)-4-piperidyl]- N-phenylpropanamide);
437	(OOO) 3-methylthiofentanyl
438	(N-[(3-methyl-1-(2-thienyl)ethyl-4-piperidinyl]-N-phenylpropanamide);

439	(PPP) 3,4-dichloro-N-[2-(dimethylamino)cyclohexyl]-N-methylbenzamide also
440	known as U-47700; and
441	(QQQ) 4-cyano CUMYL-BUTINACA.
442	(ii) Unless specifically excepted or unless listed in another schedule, any of the
443	following opium derivatives, their salts, isomers, and salts of isomers when the
444	existence of the salts, isomers, and salts of isomers is possible within the specific
445	chemical designation:
446	(A) Acetorphine;
447	(B) Acetyldihydrocodeine;
448	(C) Benzylmorphine;
449	(D) Codeine methylbromide;
450	(E) Codeine-N-Oxide;
451	(F) Cyprenorphine;
452	(G) Desomorphine;
453	(H) Dihydromorphine;
454	(I) Drotebanol;
455	(J) Etorphine (except hydrochloride salt);
456	(K) Heroin;
457	(L) Hydromorphinol;
458	(M) Methyldesorphine;
459	(N) Methylhydromorphine;
460	(O) Morphine methylbromide;
461	(P) Morphine methylsulfonate;
462	(Q) Morphine-N-Oxide;
463	(R) Myrophine;
464	(S) Nicocodeine;
465	(T) Nicomorphine;
466	(U) Normorphine;
467	(V) Pholcodine; and
468	(W) Thebacon.
469	(iii) Unless specifically excepted or unless listed in another schedule, any material,
470	compound, mixture, or preparation which contains any quantity of the following
471	hallucinogenic substances, or which contains any of their salts, isomers, and salts
472	of isomers when the existence of the salts, isomers, and salts of isomers is possible

473	within the specific chemical designation; as used in this Subsection (2)(a)(iii)
474	only, "isomer" includes the optical, position, and geometric isomers:
475	(A) Alpha-ethyltryptamine, some trade or other names: etryptamine; Monase; $\alpha$
476	-ethyl-1H-indole-3-ethanamine; 3-(2-aminobutyl) indole; $\alpha$ -ET; and AET;
477	(B) 4-bromo-2,5-dimethoxy-amphetamine, some trade or other names:
478	4-bromo-2,5-dimethoxy-α-methylphenethylamine; 4-bromo-2,5-DMA;
479	(C) 4-bromo-2,5-dimethoxyphenethylamine, some trade or other names:
480	2-(4-bromo-2,5-dimethoxyphenyl)-1-aminoethane; alpha-desmethyl DOB;
481	2C-B, Nexus;
482	(D) 2,5-dimethoxyamphetamine, some trade or other names: 2,5-dimethoxy- $\alpha$
483	-methylphenethylamine; 2,5-DMA;
484	(E) 2,5-dimethoxy-4-ethylamphetamine, some trade or other names: DOET;
485	(F) 4-methoxyamphetamine, some trade or other names: 4-methoxy- $\alpha$
486	-methylphenethylamine; paramethoxyamphetamine, PMA;
487	(G) 5-methoxy-3,4-methylenedioxyamphetamine;
488	(H) 4-methyl-2,5-dimethoxy-amphetamine, some trade and other names:
489	4-methyl-2,5-dimethoxy-α-methylphenethylamine; "DOM"; and "STP";
490	(I) 3,4-methylenedioxy amphetamine;
491	(J) 3,4-methylenedioxymethamphetamine (MDMA);
492	(K) 3,4-methylenedioxy-N-ethylamphetamine, also known as N-ethyl-
493	alpha-methyl-3,4(methylenedioxy)phenethylamine, N-ethyl MDA, MDE,
494	MDEA;
495	(L) N-hydroxy-3,4-methylenedioxyamphetamine, also known as
496	N-hydroxy-alpha-methyl-3,4(methylenedioxy)phenethylamine, and N-hydroxy
497	MDA;
498	(M) 3,4,5-trimethoxy amphetamine;
499	(N) Bufotenine, some trade and other names: $3-(\beta$
500	-Dimethylaminoethyl)-5-hydroxyindole; 3-(2-dimethylaminoethyl)-5-indolol;
501	N, N-dimethylserotonin; 5-hydroxy-N,N-dimethyltryptamine; mappine;
502	(O) Diethyltryptamine, some trade and other names: N,N-Diethyltryptamine; DET;
503	(P) Dimethyltryptamine, some trade or other names: DMT;
504	(Q) Ibogaine, some trade and other names: 7-Ethyl-6,6β
505	,7,8,9,10,12,13-octahydro-2-methoxy-6,9-methano-5H-pyrido [1', 2':1,2]

506	azepino [5,4-b] indole; Tabernanthe iboga;
507	(R) Lysergic acid diethylamide;
508	(S) Marijuana;
509	(T) Mescaline;
510	(U) Parahexyl, some trade or other names:
511	3-Hexyl-1-hydroxy-7,8,9,10-tetrahydro-6,6,9-trimethyl-6H-dibenzo[b,d]pyran;
512	Synhexyl;
513	(V) Peyote, meaning all parts of the plant presently classified botanically as
514	Lophophora williamsii Lemaire, whether growing or not, the seeds thereof, any
515	extract from any part of such plant, and every compound, manufacture, salts,
516	derivative, mixture, or preparation of such plant, its seeds or extracts
517	(Interprets 21 USC 812(c), Schedule I(c) (12));
518	(W) N-ethyl-3-piperidyl benzilate;
519	(X) N-methyl-3-piperidyl benzilate;
520	(Y) Psilocybin;
521	(Z) Psilocyn;
522	(AA) Tetrahydrocannabinols, naturally contained in a plant of the genus Cannabis
523	(cannabis plant), except for marijuana as defined in Subsection 58-37-2
524	(1)(aa)(i)(E), as well as synthetic equivalents of the substances contained in the
525	cannabis plant, or in the resinous extractives of Cannabis, sp. and/or synthetic
526	substances, derivatives, and their isomers with similar chemical structure and
527	pharmacological activity to those substances contained in the plant, such as the
528	following: $\Delta 1$ cis or trans tetrahydrocannabinol, and their optical isomers $\Delta 6$ cis or
529	trans tetrahydrocannabinol, and their optical isomers $\Delta 3,4$ cis or trans
530	tetrahydrocannabinol, and its optical isomers, and since nomenclature of these
531	substances is not internationally standardized, compounds of these structures,
532	regardless of numerical designation of atomic positions covered;
533	(BB) Ethylamine analog of phencyclidine, some trade or other names:
534	N-ethyl-1-phenylcyclohexylamine, (1-phenylcyclohexyl)ethylamine,
535	N-(1-phenylcyclohexyl)ethylamine, cyclohexamine, PCE;
536	(CC) Pyrrolidine analog of phencyclidine, some trade or other names:
537	1-(1-phenylcyclohexyl)-pyrrolidine, PCPy, PHP;
538	(DD) Thiophene analog of phencyclidine, some trade or other names:
539	1-[1-(2-thienyl)-cyclohexyl]-piperidine, 2-thienylanalog of phencyclidine,

540	TPCP, TCP; and
541	(EE) 1-[1-(2-thienyl)cyclohexyl]pyrrolidine, some other names: TCPy.
542	(iv) Unless specifically excepted or unless listed in another schedule, any material
543	compound, mixture, or preparation which contains any quantity of the following
544	substances having a depressant effect on the central nervous system, including its
545	salts, isomers, and salts of isomers when the existence of the salts, isomers, and
546	salts of isomers is possible within the specific chemical designation:
547	(A) Mecloqualone; and
548	(B) Methaqualone.
549	(v) Any material, compound, mixture, or preparation containing any quantity of the
550	following substances having a stimulant effect on the central nervous system,
551	including their salts, isomers, and salts of isomers:
552	(A) Aminorex, some other names: aminoxaphen; 2-amino-5-phenyl-2-oxazoline;
553	or 4,5-dihydro-5-phenyl-2-oxazolamine;
554	(B) Cathinone, some trade or other names: 2-amino-1-phenyl-1-propanone,
555	alpha-aminopropiophenone, 2-aminopropiophenone, and norephedrone;
556	(C) Fenethylline;
557	(D) Methcathinone, some other names: 2-(methylamino)-propiophenone;
558	alpha-(methylamino)propiophenone; 2-(methylamino)-1-phenylpropan-1-one;
559	alpha-N-methylaminopropiophenone; monomethylpropion; ephedrone;
560	N-methylcathinone; methylcathinone; AL-464; AL-422; AL-463 and UR1432,
561	its salts, optical isomers, and salts of optical isomers;
562	(E) (±)cis-4-methylaminorex ((±)cis-4,5-dihydro-4-methyl-5-phenyl-2-oxazolamine)
563	(F) N-ethylamphetamine; and
564	(G) N,N-dimethylamphetamine, also known as
565	$N, N-alpha-trimethyl-benzene ethanamine;\ N, N-alpha-trimethyl phenethylamine.$
566	(vi) Any material, compound, mixture, or preparation which contains any quantity of
567	the following substances, including their optical isomers, salts, and salts of
568	isomers, subject to temporary emergency scheduling:
569	(A) N-[1-benzyl-4-piperidyl]-N-phenylpropanamide (benzylfentanyl); and
570	(B) N-[1- (2-thienyl)methyl-4-piperidyl]-N-phenylpropanamide (thenylfentanyl).
571	(vii) Unless specifically excepted or unless listed in another schedule, any material,
572	compound, mixture, or preparation which contains any quantity of gamma
573	hydroxy butyrate (gamma hydrobutyric acid) including its salts, isomers, and

574	salts of isomers.
575	(viii) Unless specifically excepted or unless listed in another schedule, the following
576	substances commonly found in Mitragyna speciosa, including synthetic versions,
577	isomers, esters, ethers, salts, and salts of isomers, esters, and ethers, when the
578	existence of the isomers, esters, ethers, and salts is possible within the specific
579	chemical designation:
580	(A) 7-hydroxymitragynine (methyl
581	(E)-2[(2S,3S,7aS,12bS)-3-ethyl-7a-hydroxy-8-methoxy-2,3,4,6,7,12b-hexahydro-
582	1H-indolo[2,3-a]quinolizin-2-yl]-3-methoxyprop-2-enoate), that is
583	concentrated at a level greater than .4% of the total kratom alkaloid
584	composition of any substance or product; and
585	(B) Mitragynine pseudoindoxyl (methyl
586	(E)-2-[(2S,6'S,7'S,8'aS)-6'-ethyl-4-methoxy-3-oxospiro[1H-indole-2,1'-3,5,6,7,8,8a-
587	hexahydro-2H-indolizine]-7'-yl]-3-methoxyprop-2-enoate).
588	(b) Schedule II:
589	(i) Unless specifically excepted or unless listed in another schedule, any of the
590	following substances whether produced directly or indirectly by extraction from
591	substances of vegetable origin, or independently by means of chemical synthesis,
592	or by a combination of extraction and chemical synthesis:
593	(A) Opium and opiate, and any salt, compound, derivative, or preparation of
594	opium or opiate, excluding apomorphine, dextrorphan, nalbuphine, nalmefene,
595	naloxone, and naltrexone, and their respective salts, but including:
596	(I) Raw opium;
597	(II) Opium extracts;
598	(III) Opium fluid;
599	(IV) Powdered opium;
600	(V) Granulated opium;
601	(VI) Tincture of opium;
602	(VII) Codeine;
603	(VIII) Ethylmorphine;
604	(IX) Etorphine hydrochloride;
605	(X) Hydrocodone;
606	(XI) Hydromorphone;
607	(XII) Metopon:

608	(XIII) Morphine;
609	(XIV) Oxycodone;
610	(XV) Oxymorphone; and
611	(XVI) Thebaine;
612	(B) Any salt, compound, derivative, or preparation which is chemically equivalent
613	or identical with any of the substances referred to in Subsection (2)(b)(i)(A),
614	except that these substances may not include the isoquinoline alkaloids of
615	opium;
616	(C) Opium poppy and poppy straw;
617	(D) Coca leaves and any salt, compound, derivative, or preparation of coca leaves,
618	and any salt, compound, derivative, or preparation which is chemically
619	equivalent or identical with any of these substances, and includes cocaine and
620	ecgonine, their salts, isomers, derivatives, and salts of isomers and derivatives,
621	whether derived from the coca plant or synthetically produced, except the
622	substances may not include decocainized coca leaves or extraction of coca
623	leaves, which extractions do not contain cocaine or ecgonine; and
624	(E) Concentrate of poppy straw, which means the crude extract of poppy straw in
625	either liquid, solid, or powder form which contains the phenanthrene alkaloids
626	of the opium poppy.
627	(ii) Unless specifically excepted or unless listed in another schedule, any of the
628	following opiates, including their isomers, esters, ethers, salts, and salts of
629	isomers, esters, and ethers, when the existence of the isomers, esters, ethers, and
630	salts is possible within the specific chemical designation, except dextrorphan and
631	levopropoxyphene:
632	(A) Alfentanil;
633	(B) Alphaprodine;
634	(C) Anileridine;
635	(D) Bezitramide;
636	(E) Bulk dextropropoxyphene (nondosage forms);
637	(F) Carfentanil;
638	(G) Dihydrocodeine;
639	(H) Diphenoxylate;
640	(I) Fentanyl;
641	(J) Isomethadone:

642	(K) Levo-alphacetylmethadol, some other names: levo-alpha-acetylmethadol,
643	levomethadyl acetate, or LAAM;
644	(L) Levomethorphan;
645	(M) Levorphanol;
646	(N) Metazocine;
647	(O) Methadone;
648	(P) Methadone-Intermediate, 4-cyano-2-dimethylamino-4, 4-diphenyl butane;
649	(Q) Moramide-Intermediate, 2-methyl-3-morpholino-1,
650	1-diphenylpropane-carboxylic acid;
651	(R) Pethidine (meperidine);
652	(S) Pethidine-Intermediate-A, 4-cyano-1-methyl-4-phenylpiperidine;
653	(T) Pethidine-Intermediate-B, ethyl-4-phenylpiperidine-4-carboxylate;
654	(U) Pethidine-Intermediate-C, 1-methyl-4-phenylpiperidine-4-carboxylic acid;
655	(V) Phenazocine;
656	(W) Piminodine;
657	(X) Racemethorphan;
658	(Y) Racemorphan;
659	(Z) Remifentanil; and
660	(AA) Sufentanil.
661	(iii) Unless specifically excepted or unless listed in another schedule, any material,
662	compound, mixture, or preparation which contains any quantity of the following
663	substances having a stimulant effect on the central nervous system:
664	(A) Amphetamine, its salts, optical isomers, and salts of its optical isomers;
665	(B) Methamphetamine, its salts, isomers, and salts of its isomers;
666	(C) Phenmetrazine and its salts; and
667	(D) Methylphenidate.
668	(iv) Unless specifically excepted or unless listed in another schedule, any material,
669	compound, mixture, or preparation which contains any quantity of the following
670	substances having a depressant effect on the central nervous system, including its
671	salts, isomers, and salts of isomers when the existence of the salts, isomers, and
672	salts of isomers is possible within the specific chemical designation:
673	(A) Amobarbital;
674	(B) Glutethimide;
675	(C) Pentobarbital;

676	(D) Phencyclidine;
677	(E) Phencyclidine immediate precursors: 1-phenylcyclohexylamine and
678	1-piperidinocyclohexanecarbonitrile (PCC); and
679	(F) Secobarbital.
680	(v)(A) Unless specifically excepted or unless listed in another schedule, any
681	material, compound, mixture, or preparation which contains any quantity of
682	Phenylacetone.
683	(B) Some of these substances may be known by trade or other names:
684	phenyl-2-propanone; P2P; benzyl methyl ketone; and methyl benzyl ketone.
685	(vi) Nabilone, another name for nabilone: (±
686	)-trans-3-(1,1-dimethylheptyl)-6,6a,7,8,10,10a-hexahydro-1-hydroxy-6,
687	6-dimethyl-9H-dibenzo[b,d]pyran-9-one.
688	(vii) A drug product or preparation that contains any component of marijuana,
689	including tetrahydrocannabinol, and is approved by the United States Food and
690	Drug Administration and scheduled by the Drug Enforcement Administration in
691	Schedule II of the federal Controlled Substances Act, Title II, P.L. 91-513.
692	(c) Schedule III:
693	(i) Unless specifically excepted or unless listed in another schedule, any material,
694	compound, mixture, or preparation which contains any quantity of the following
695	substances having a stimulant effect on the central nervous system, including its
696	salts, isomers whether optical, position, or geometric, and salts of the isomers
697	when the existence of the salts, isomers, and salts of isomers is possible within the
698	specific chemical designation:
699	(A) Those compounds, mixtures, or preparations in dosage unit form containing
700	any stimulant substances listed in Schedule II, which compounds, mixtures, or
701	preparations were listed on August 25, 1971, as excepted compounds under
702	Section 1308.32 of Title 21 of the Code of Federal Regulations, and any other
703	drug of the quantitive composition shown in that list for those drugs or which
704	is the same except that it contains a lesser quantity of controlled substances;
705	(B) Benzphetamine;
706	(C) Chlorphentermine;
707	(D) Clortermine; and
708	(E) Phendimetrazine.
709	(ii) Unless specifically excepted or unless listed in another schedule, any material,

710	compound, mixture, or preparation which contains any quantity of the following
711	substances having a depressant effect on the central nervous system:
712	(A) Any compound, mixture, or preparation containing amobarbital, secobarbital,
713	pentobarbital, or any salt of any of them, and one or more other active
714	medicinal ingredients which are not listed in any schedule;
715	(B) Any suppository dosage form containing amobarbital, secobarbital, or
716	pentobarbital, or any salt of any of these drugs which is approved by the United
717	States Food and Drug Administration for marketing only as a suppository;
718	(C) Any substance which contains any quantity of a derivative of barbituric acid
719	or any salt of any of them;
720	(D) Chlorhexadol;
721	(E) Buprenorphine;
722	(F) Any drug product containing gamma hydroxybutyric acid, including its salts,
723	isomers, and salts of isomers, for which an application is approved under the
724	federal Food, Drug, and Cosmetic Act, Section 505;
725	(G) Ketamine, its salts, isomers, and salts of isomers, some other names for
726	ketamine: ± -2-(2-chlorophenyl)-2-(methylamino)-cyclohexanone;
727	(H) Lysergic acid;
728	(I) Lysergic acid amide;
729	(J) Methyprylon;
730	(K) Sulfondiethylmethane;
731	(L) Sulfonethylmethane;
732	(M) Sulfonmethane; and
733	(N) Tiletamine and zolazepam or any of their salts, some trade or other names for
734	a tiletamine-zolazepam combination product: Telazol, some trade or other
735	names for tiletamine: 2-(ethylamino)-2-(2-thienyl)-cyclohexanone, some trade
736	or other names for zolazepam:
737	4-(2-fluorophenyl)-6,8-dihydro-1,3,8-trimethylpyrazolo-[3,4-e]
738	[1,4]-diazepin-7(1H)-one, flupyrazapon.
739	(iii) Dronabinol (synthetic) in sesame oil and encapsulated in a soft gelatin capsule in
740	a U.S. Food and Drug Administration approved drug product, some other names
741	for dronabinol:
742	(6aR-trans)-6a,7,8,10a-tetrahydro-6,6,9-trimethyl-3-pentyl-6H-dibenzo[b,d]pyran-1-ol
743	or (-)-delta-9-(trans)-tetrahydrocannabinol.

744	(iv) Nalorphine.
745	(v) Unless specifically excepted or unless listed in another schedule, any material,
746	compound, mixture, or preparation containing limited quantities of any of the
747	following narcotic drugs, or their salts calculated as the free anhydrous base or
748	alkaloid:
749	(A) Not more than 1.8 grams of codeine per 100 milliliters or not more than 90
750	milligrams per dosage unit, with an equal or greater quantity of an isoquinoline
751	alkaloid of opium;
752	(B) Not more than 1.8 grams of codeine per 100 milliliters or not more than 90
753	milligrams per dosage unit, with one or more active non-narcotic ingredients in
754	recognized therapeutic amounts;
755	(C) Not more than 300 milligrams of dihydrocodeinone per 100 milliliters or not
756	more than 15 milligrams per dosage unit, with a fourfold or greater quantity of
757	an isoquinoline alkaloid of opium;
758	(D) Not more than 300 milligrams of dihydrocodeinone per 100 milliliters or not
759	more than 15 milligrams per dosage unit, with one or more active, non-narcotic
760	ingredients in recognized therapeutic amounts;
761	(E) Not more than 1.8 grams of dihydrocodeine per 100 milliliters or not more
762	than 90 milligrams per dosage unit, with one or more active non-narcotic
763	ingredients in recognized therapeutic amounts;
764	(F) Not more than 300 milligrams of ethylmorphine per 100 milliliters or not more
765	than 15 milligrams per dosage unit, with one or more active, non-narcotic
766	ingredients in recognized therapeutic amounts;
767	(G) Not more than 500 milligrams of opium per 100 milliliters or per 100 grams,
768	or not more than 25 milligrams per dosage unit, with one or more active,
769	non-narcotic ingredients in recognized therapeutic amounts; and
770	(H) Not more than 50 milligrams of morphine per 100 milliliters or per 100 grams
771	with one or more active, non-narcotic ingredients in recognized therapeutic
772	amounts.
773	(vi) Unless specifically excepted or unless listed in another schedule, anabolic
774	steroids including any of the following or any isomer, ester, salt, or derivative of
775	the following that promotes muscle growth:
776	(A) Boldenone;
777	(B) Chlorotestosterone (4-chlortestosterone);

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778	(C) Clostebol;
779	(D) Dehydrochlormethyltestosterone;
780	(E) Dihydrotestosterone (4-dihydrotestosterone);
781	(E) Dinydrotestosterone (4-dinydrotestosterone),  (F) Drostanolone;
782	(G) Ethylestrenol;
783	(H) Fluoxymesterone;
784	(I) Formebulone (formebolone);
785	(J) Mesterolone;
786	(K) Methandienone;
787	(L) Methandranone;
788	(M) Methandriol;
789	(N) Methandrostenolone;
790	(O) Methenolone;
791	(P) Methyltestosterone;
792	(Q) Mibolerone;
793	(R) Nandrolone;
794	(S) Norethandrolone;
795	(T) Oxandrolone;
796	(U) Oxymesterone;
797	(V) Oxymetholone;
798	(W) Stanolone;
799	(X) Stanozolol;
800	(Y) Testolactone;
801	(Z) Testosterone; and
802	(AA) Trenbolone.
803	(vii) Anabolic steroids expressly intended for administration through implants to
804	cattle or other nonhuman species, and approved by the Secretary of Health and
805	Human Services for use, may not be classified as a controlled substance.
806	(viii) A drug product or preparation that contains any component of marijuana,
807	including tetrahydrocannabinol, and is approved by the United States Food and
808	Drug Administration and scheduled by the Drug Enforcement Administration in
809	Schedule III of the federal Controlled Substances Act, Title II, P.L. 91-513.
810	(ix) Nabiximols.
811	(d) Schedule IV:
011	(a) Sollowing I i i

812	(i) Unless specifically excepted or unless listed in another schedule, any material,
813	compound, mixture, or preparation containing not more than 1 milligram of
814	difenoxin and not less than 25 micrograms of atropine sulfate per dosage unit, or
815	any salts of any of them.
816	(ii) Unless specifically excepted or unless listed in another schedule, any material,
817	compound, mixture, or preparation which contains any quantity of the following
818	substances, including its salts, isomers, and salts of isomers when the existence of
819	the salts, isomers, and salts of isomers is possible within the specific chemical
820	designation:
821	(A) Alprazolam;
822	(B) Barbital;
823	(C) Bromazepam;
824	(D) Butorphanol;
825	(E) Camazepam;
826	(F) Carisoprodol;
827	(G) Chloral betaine;
828	(H) Chloral hydrate;
829	(I) Chlordiazepoxide;
830	(J) Clobazam;
831	(K) Clonazepam;
832	(L) Clorazepate;
833	(M) Clotiazepam;
834	(N) Cloxazolam;
835	(O) Delorazepam;
836	(P) Diazepam;
837	(Q) Dichloralphenazone;
838	(R) Estazolam;
839	(S) Ethchlorvynol;
840	(T) Ethinamate;
841	(U) Ethyl loflazepate;
842	(V) Fludiazepam;
843	(W) Flunitrazepam;
844	(X) Flurazepam;
845	(Y) Halazepam;

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846	(Z) Haloxazolam;
847	(AA) Ketazolam;
848	(BB) Loprazolam;
849	(CC) Lorazepam;
850	(DD) Lormetazepam;
851	(EE) Mebutamate;
852	(FF) Medazepam;
853	(GG) Meprobamate;
854	(HH) Methohexital;
855	(II) Methylphenobarbital (mephobarbital);
856	(JJ) Midazolam;
857	(KK) Nimetazepam;
858	(LL) Nitrazepam;
859	(MM) Nordiazepam;
860	(NN) Oxazepam;
861	(OO) Oxazolam;
862	(PP) Paraldehyde;
863	(QQ) Pentazocine;
864	(RR) Petrichloral;
865	(SS) Phenobarbital;
866	(TT) Pinazepam;
867	(UU) Prazepam;
868	(VV) Quazepam;
869	(WW) Temazepam;
870	(XX) Tetrazepam;
871	(YY) Tramadol;
872	(ZZ) Triazolam;
873	(AAA) Zaleplon; and
874	(BBB) Zolpidem.
875	(iii) Any material, compound, mixture, or preparation of fenfluramine which contains
876	any quantity of the following substances, including its salts, isomers whether
877	optical, position, or geometric, and salts of the isomers when the existence of the
878	salts, isomers, and salts of isomers is possible.
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 881 substances having a stimulant effect on the central nervous system, including its 882 salts, isomers whether optical, position, or geometric isomers, and salts of the 883 isomers when the existence of the salts, isomers, and salts of isomers is possible 884 within the specific chemical designation: 885 (A) Cathine ((+)-norpseudoephedrine); 886 (B) Diethylpropion; 887 (C) Fencamfamine; 888 (D) Fenproprex; 889 (E) Mazindol; 890 (F) Mefenorex; 891 (G) Modafinil; 892 (H) Pemoline, including organometallic complexes and chelates thereof; 893 (I) Phentermine; 894 (J) Pipradrol; 895 (K) Sibutramine; and 896 (L) SPA ((-)-1-dimethylamino-1,2-diphenylethane). 897 (v) Unless specifically excepted or unless listed in another schedule, any material, 898 compound, mixture, or preparation which contains any quantity of 899 dextropropoxyphene (alpha-(+)-4-dimethylamino-1, 900 2-diphenyl-3-methyl-2-propionoxybutane), including its salts. 901 (vi) A drug product or preparation that contains any component of marijuana and is 902 approved by the United States Food and Drug Administration and scheduled by 903 the Drug Enforcement Administration in Schedule IV of the federal Controlled 904 Substances Act, Title II, P.L. 91-513. 905 (e) Schedule V: 906 (i) Any compound, mixture, or preparation containing any of the following limited 907 quantities of narcotic drugs, or their salts calculated as the free anhydrous base or 908 alkaloid, which includes one or more non-narcotic active medicinal ingredients in 909 sufficient proportion to confer upon the compound, mixture, or preparation 910 valuable medicinal qualities other than those possessed by the narcotic drug alone: 911 (A) not more than 200 milligrams of codeine per 100 milliliters or per 100 grams; 912 (B) not more than 100 milligrams of dihydrocodeine per 100 milliliters or per 100 913 grams;

914	(C) not more than 100 milligrams of ethylmorphine per 100 milliliters or per 100
915	grams;
916	(D) not more than 2.5 milligrams of diphenoxylate and not less than 25
917	micrograms of atropine sulfate per dosage unit;
918	(E) not more than 100 milligrams of opium per 100 milliliters or per 100 grams;
919	(F) not more than 0.5 milligram of difenoxin and not less than 25 micrograms of
920	atropine sulfate per dosage unit; and
921	(G) unless specifically exempted or excluded or unless listed in another schedule
922	any material, compound, mixture, or preparation which contains Pyrovalerone
923	having a stimulant effect on the central nervous system, including its salts,
924	isomers, and salts of isomers.
925	(ii) A drug product or preparation that contains any component of marijuana,
926	including cannabidiol, and is approved by the United States Food and Drug
927	Administration and scheduled by the Drug Enforcement Administration in
928	Schedule V of the federal Controlled Substances Act, Title II, P.L. 91-513.
929	(iii) Gabapentin.
930	Section 4. Effective Date.
931	This bill takes effect:
932	(1) except as provided in Subsection (2), May 6, 2026; or
933	(2) if approved by two-thirds of all members elected to each house:
934	(a) upon approval by the governor;
935	(b) without the governor's signature, the day following the constitutional time limit of
936	Utah Constitution, Article VII, Section 8; or
937	(c) in the case of a veto, the date of veto override