

1 AN ACT relating to cannabinoid products.

2 ***Be it enacted by the General Assembly of the Commonwealth of Kentucky:***

3 ➔Section 1. KRS 218A.010 is amended to read as follows:

4 As used in this chapter:

- 5 (1) "Administer" means the direct application of a controlled substance, whether by  
6 injection, inhalation, ingestion, or any other means, to the body of a patient or  
7 research subject by:
- 8 (a) A practitioner or by his or her authorized agent under his or her immediate  
9 supervision and pursuant to his or her order; or
- 10 (b) The patient or research subject at the direction and in the presence of the  
11 practitioner;
- 12 (2) "Anabolic steroid" means any drug or hormonal substance chemically and  
13 pharmacologically related to testosterone that promotes muscle growth and includes  
14 those substances classified as Schedule III controlled substances pursuant to KRS  
15 218A.020 but does not include estrogens, progestins, and anticosteroids;
- 16 (3) "Cabinet" means the Cabinet for Health and Family Services;
- 17 (4) "Carfentanil" means any substance containing any quantity of carfentanil, or any of  
18 its salts, isomers, or salts of isomers;
- 19 (5) "Certified community based palliative care program" means a palliative care  
20 program which has received certification from the Joint Commission;
- 21 (6) "Child" means any person under the age of majority as specified in KRS 2.015;
- 22 (7) "Cocaine" means a substance containing any quantity of cocaine, its salts, optical  
23 and geometric isomers, and salts of isomers;
- 24 (8) "Controlled substance" means methamphetamine, or a drug, substance, or  
25 immediate precursor in Schedules I through V and includes a controlled substance  
26 analogue;
- 27 (9) (a) "Controlled substance analogue," except as provided in paragraph (b) of this

1 subsection, means a substance:

- 2 1. The chemical structure of which is substantially similar to the structure  
3 of a controlled substance in Schedule I or II; and
- 4 2. Which has a stimulant, depressant, or hallucinogenic effect on the  
5 central nervous system that is substantially similar to or greater than the  
6 stimulant, depressant, or hallucinogenic effect on the central nervous  
7 system of a controlled substance in Schedule I or II; or
- 8 3. With respect to a particular person, which such person represents or  
9 intends to have a stimulant, depressant, or hallucinogenic effect on the  
10 central nervous system that is substantially similar to or greater than the  
11 stimulant, depressant, or hallucinogenic effect on the central nervous  
12 system of a controlled substance in Schedule I or II.

13 (b) Such term does not include:

- 14 1. Any substance for which there is an approved new drug application;
- 15 2. With respect to a particular person, any substance if an exemption is in  
16 effect for investigational use for that person pursuant to federal law to  
17 the extent conduct with respect to such substance is pursuant to such  
18 exemption; or
- 19 3. Any substance to the extent not intended for human consumption before  
20 the exemption described in subparagraph 2. of this paragraph takes  
21 effect with respect to that substance;

22 (10) "Counterfeit substance" means a controlled substance which, or the container or  
23 labeling of which, without authorization, bears the trademark, trade name, or other  
24 identifying mark, imprint, number, or device, or any likeness thereof, of a  
25 manufacturer, distributor, or dispenser other than the person who in fact  
26 manufactured, distributed, or dispensed the substance;

27 (11) "Dispense" means to deliver a controlled substance to an ultimate user or research

1 subject by or pursuant to the lawful order of a practitioner, including the packaging,  
2 labeling, or compounding necessary to prepare the substance for that delivery;

3 (12) "Dispenser" means a person who lawfully dispenses a Schedule II, III, IV, or V  
4 controlled substance to or for the use of an ultimate user;

5 (13) "Distribute" means to deliver other than by administering or dispensing a controlled  
6 substance;

7 (14) "Dosage unit" means a single pill, capsule, ampule, liquid, or other form of  
8 administration available as a single unit;

9 (15) "Drug" means:

10 (a) Substances recognized as drugs in the official United States Pharmacopoeia,  
11 official Homeopathic Pharmacopoeia of the United States, or official National  
12 Formulary, or any supplement to any of them;

13 (b) Substances intended for use in the diagnosis, care, mitigation, treatment, or  
14 prevention of disease in man or animals;

15 (c) Substances (other than food) intended to affect the structure or any function of  
16 the body of man or animals; and

17 (d) Substances intended for use as a component of any article specified in this  
18 subsection.

19 It does not include devices or their components, parts, or accessories;

20 (16) "Fentanyl" means a substance containing any quantity of fentanyl, or any of its salts,  
21 isomers, or salts of isomers;

22 (17) "Fentanyl derivative" means a substance containing any quantity of any chemical  
23 compound, except compounds specifically scheduled as controlled substances by  
24 statute or by administrative regulation pursuant to this chapter, which is structurally  
25 derived from 1-ethyl-4-(N-phenylamido) piperadine:

26 (a) By substitution:

27 1. At the 2-position of the 1-ethyl group with a phenyl, furan, thiophene, or

- 1                    ethyloxotetrazole ring system; and
- 2                    2. Of the terminal amido hydrogen atom with an alkyl, alkoxy, cycloalkyl,
- 3                    or furanyl group; and
- 4                    (b) Which may be further modified in one (1) or more of the following ways:
- 5                    1. By substitution on the N-phenyl ring to any extent with alkyl, alkoxy,
- 6                    haloalkyl, hydroxyl, or halide substituents;
- 7                    2. By substitution on the piperadine ring to any extent with alkyl, allyl,
- 8                    alkoxy, hydroxy, or halide substituents at the 2-, 3-, 5-, and/or 6-
- 9                    positions;
- 10                    3. By substitution on the piperadine ring to any extent with a phenyl,
- 11                    alkoxy, or carboxylate ester substituent at the 4- position; or
- 12                    4. By substitution on the 1-ethyl group to any extent with alkyl, alkoxy, or
- 13                    hydroxy substituents;
- 14                    (18) "Good faith prior examination," as used in KRS Chapter 218A and for criminal
- 15                    prosecution only, means an in-person medical examination of the patient conducted
- 16                    by the prescribing practitioner or other health-care professional routinely relied
- 17                    upon in the ordinary course of his or her practice, at which time the patient is
- 18                    physically examined and a medical history of the patient is obtained. "In-person"
- 19                    includes telehealth examinations. This subsection shall not be applicable to hospice
- 20                    providers licensed pursuant to KRS Chapter 216B;
- 21                    (19) "Hazardous chemical substance" includes any chemical substance used or intended
- 22                    for use in the illegal manufacture of a controlled substance as defined in this section
- 23                    or the illegal manufacture of methamphetamine as defined in KRS 218A.1431,
- 24                    which:
- 25                    (a) Poses an explosion hazard;
- 26                    (b) Poses a fire hazard; or
- 27                    (c) Is poisonous or injurious if handled, swallowed, or inhaled;

- 1 (20) "Heroin" means a substance containing any quantity of heroin, or any of its salts,  
2 isomers, or salts of isomers;
- 3 (21) "Hydrocodone combination product" means a drug with:
- 4 (a) Not more than three hundred (300) milligrams of dihydrocodeinone, or any of  
5 its salts, per one hundred (100) milliliters or not more than fifteen (15)  
6 milligrams per dosage unit, with a fourfold or greater quantity of an  
7 isoquinoline alkaloid of opium; or
- 8 (b) Not more than three hundred (300) milligrams of dihydrocodeinone, or any of  
9 its salts, per one hundred (100) milliliters or not more than fifteen (15)  
10 milligrams per dosage unit, with one (1) or more active, nonnarcotic  
11 ingredients in recognized therapeutic amounts;
- 12 (22) "Immediate precursor" means a substance which is the principal compound  
13 commonly used or produced primarily for use, and which is an immediate chemical  
14 intermediary used or likely to be used in the manufacture of a controlled substance  
15 or methamphetamine, the control of which is necessary to prevent, curtail, or limit  
16 manufacture;
- 17 (23) "Industrial hemp" has the same meaning as in KRS 260.850;
- 18 (24) "Industrial hemp products" has the same meaning as in KRS 260.850;
- 19 (25) "Intent to manufacture" means any evidence which demonstrates a person's  
20 conscious objective to manufacture a controlled substance or methamphetamine.  
21 Such evidence includes but is not limited to statements and a chemical substance's  
22 usage, quantity, manner of storage, or proximity to other chemical substances or  
23 equipment used to manufacture a controlled substance or methamphetamine;
- 24 (26) "Isomer" means the optical isomer, except the Cabinet for Health and Family  
25 Services may include the optical, positional, or geometric isomer to classify any  
26 substance pursuant to KRS 218A.020;
- 27 (27) "Manufacture," except as provided in KRS 218A.1431, means the production,

1 preparation, propagation, compounding, conversion, or processing of a controlled  
2 substance, either directly or indirectly by extraction from substances of natural  
3 origin or independently by means of chemical synthesis, or by a combination of  
4 extraction and chemical synthesis, and includes any packaging or repackaging of the  
5 substance or labeling or relabeling of its container except that this term does not  
6 include activities:

7 (a) By a practitioner as an incident to his or her administering or dispensing of a  
8 controlled substance in the course of his or her professional practice;

9 (b) By a practitioner, or by his or her authorized agent under his supervision, for  
10 the purpose of, or as an incident to, research, teaching, or chemical analysis  
11 and not for sale; or

12 (c) By a pharmacist as an incident to his or her dispensing of a controlled  
13 substance in the course of his or her professional practice;

14 (28) "Marijuana" means all parts of the plant *Cannabis* sp., whether growing or not; the  
15 seeds thereof; the resin extracted from any part of the plant; and every compound,  
16 manufacture, salt, derivative, mixture, or preparation of the plant, its seeds or resin  
17 or any compound, mixture, or preparation which contains any quantity of these  
18 substances. The term "marijuana" does not include:

19 (a) Industrial hemp that is in the possession, custody, or control of a person who  
20 holds a license issued by the Department of Agriculture permitting that person  
21 to cultivate, handle, or process industrial hemp;

22 (b) Industrial hemp products that do not include any living plants, viable seeds,  
23 leaf materials, or floral materials;

24 (c) The substance cannabidiol, when transferred, dispensed, or administered  
25 pursuant to the written order of a physician practicing at a hospital or  
26 associated clinic affiliated with a Kentucky public university having a college  
27 or school of medicine;

- 1 (d) For persons participating in a clinical trial or in an expanded access program,  
2 a drug or substance approved for the use of those participants by the United  
3 States Food and Drug Administration;
- 4 (e) A cannabidiol product derived from industrial hemp, as defined in KRS  
5 260.850;~~[-or]~~
- 6 (f) **For the purpose of conducting scientific research, a cannabinoid product**  
7 **derived from industrial hemp, as defined in KRS 260.850; or**
- 8 (g) A **cannabinoid**~~cannabidiol~~ product approved as a prescription medication  
9 by the United States Food and Drug Administration;
- 10 (29) "Medical history," as used in KRS Chapter 218A and for criminal prosecution only,  
11 means an accounting of a patient's medical background, including but not limited to  
12 prior medical conditions, prescriptions, and family background;
- 13 (30) "Medical order," as used in KRS Chapter 218A and for criminal prosecution only,  
14 means a lawful order of a specifically identified practitioner for a specifically  
15 identified patient for the patient's health-care needs. "Medical order" may or may  
16 not include a prescription drug order;
- 17 (31) "Medical record," as used in KRS Chapter 218A and for criminal prosecution only,  
18 means a record, other than for financial or billing purposes, relating to a patient,  
19 kept by a practitioner as a result of the practitioner-patient relationship;
- 20 (32) "Methamphetamine" means any substance that contains any quantity of  
21 methamphetamine, or any of its salts, isomers, or salts of isomers;
- 22 (33) "Narcotic drug" means any of the following, whether produced directly or indirectly  
23 by extraction from substances of vegetable origin, or independently by means of  
24 chemical synthesis, or by a combination of extraction and chemical synthesis:
- 25 (a) Opium and opiate, and any salt, compound, derivative, or preparation of  
26 opium or opiate;
- 27 (b) Any salt, compound, isomer, derivative, or preparation thereof which is

- 1 chemically equivalent or identical with any of the substances referred to in  
2 paragraph (a) of this subsection, but not including the isoquinoline alkaloids  
3 of opium;
- 4 (c) Opium poppy and poppy straw;
- 5 (d) Coca leaves, except coca leaves and extracts of coca leaves from which  
6 cocaine, ecgonine, and derivatives of ecgonine or their salts have been  
7 removed;
- 8 (e) Cocaine, its salts, optical and geometric isomers, and salts of isomers;
- 9 (f) Ecgonine, its derivatives, their salts, isomers, and salts of isomers; and
- 10 (g) Any compound, mixture, or preparation which contains any quantity of any of  
11 the substances referred to in paragraphs (a) to (f) of this subsection;
- 12 (34) "Opiate" means any substance having an addiction-forming or addiction-sustaining  
13 liability similar to morphine or being capable of conversion into a drug having  
14 addiction-forming or addiction-sustaining liability. It does not include, unless  
15 specifically designated as controlled under KRS 218A.020, the dextrorotatory  
16 isomer of 3-methoxy-n-methylmorphinan and its salts (dextromethorphan). It does  
17 include its racemic and levorotatory forms;
- 18 (35) "Opium poppy" means the plant of the species *papaver somniferum* L., except its  
19 seeds;
- 20 (36) "Person" means individual, corporation, government or governmental subdivision  
21 or agency, business trust, estate, trust, partnership or association, or any other legal  
22 entity;
- 23 (37) "Physical injury" has the same meaning it has in KRS 500.080;
- 24 (38) "Poppy straw" means all parts, except the seeds, of the opium poppy, after mowing;
- 25 (39) "Pharmacist" means a natural person licensed by this state to engage in the practice  
26 of the profession of pharmacy;
- 27 (40) "Practitioner" means a physician, dentist, podiatrist, veterinarian, scientific



1 investigator, optometrist as authorized in KRS 320.240, advanced practice  
2 registered nurse as authorized under KRS 314.011, physician assistant as authorized  
3 under KRS 311.858, or other person licensed, registered, or otherwise permitted by  
4 state or federal law to acquire, distribute, dispense, conduct research with respect to,  
5 or to administer a controlled substance in the course of professional practice or  
6 research in this state. "Practitioner" also includes a physician, dentist, podiatrist,  
7 veterinarian, or advanced practice registered nurse authorized under KRS 314.011  
8 who is a resident of and actively practicing in a state other than Kentucky and who  
9 is licensed and has prescriptive authority for controlled substances under the  
10 professional licensing laws of another state, unless the person's Kentucky license  
11 has been revoked, suspended, restricted, or probated, in which case the terms of the  
12 Kentucky license shall prevail;

13 (41) "Practitioner-patient relationship," as used in KRS Chapter 218A and for criminal  
14 prosecution only, means a medical relationship that exists between a patient and a  
15 practitioner or the practitioner's designee, after the practitioner or his or her  
16 designee has conducted at least one (1) good faith prior examination;

17 (42) "Prescription" means a written, electronic, or oral order for a drug or medicine, or  
18 combination or mixture of drugs or medicines, or proprietary preparation, signed or  
19 given or authorized by a medical, dental, chiropody, veterinarian, optometric  
20 practitioner, or advanced practice registered nurse, and intended for use in the  
21 diagnosis, cure, mitigation, treatment, or prevention of disease in man or other  
22 animals;

23 (43) "Prescription blank," with reference to a controlled substance, means a document  
24 that meets the requirements of KRS 218A.204 and 217.216;

25 (44) "Presumptive probation" means a sentence of probation not to exceed the maximum  
26 term specified for the offense, subject to conditions otherwise authorized by law,  
27 that is presumed to be the appropriate sentence for certain offenses designated in

1 this chapter, notwithstanding contrary provisions of KRS Chapter 533. That  
2 presumption shall only be overcome by a finding on the record by the sentencing  
3 court of substantial and compelling reasons why the defendant cannot be safely and  
4 effectively supervised in the community, is not amenable to community-based  
5 treatment, or poses a significant risk to public safety;

6 (45) "Production" includes the manufacture, planting, cultivation, growing, or harvesting  
7 of a controlled substance;

8 (46) "Recovery program" means an evidence-based, nonclinical service that assists  
9 individuals and families working toward sustained recovery from substance use and  
10 other criminal risk factors. This can be done through an array of support programs  
11 and services that are delivered through residential and nonresidential means;

12 (47) "Salvia" means *Salvia divinorum* or Salvinorin A and includes all parts of the plant  
13 presently classified botanically as *Salvia divinorum*, whether growing or not, the  
14 seeds thereof, any extract from any part of that plant, and every compound,  
15 manufacture, derivative, mixture, or preparation of that plant, its seeds, or its  
16 extracts, including salts, isomers, and salts of isomers whenever the existence of  
17 such salts, isomers, and salts of isomers is possible within the specific chemical  
18 designation of that plant, its seeds, or extracts. The term shall not include any other  
19 species in the genus *salvia*;

20 (48) "Second or subsequent offense" means that for the purposes of this chapter an  
21 offense is considered as a second or subsequent offense, if, prior to his or her  
22 conviction of the offense, the offender has at any time been convicted under this  
23 chapter, or under any statute of the United States, or of any state relating to  
24 substances classified as controlled substances or counterfeit substances, except that  
25 a prior conviction for a nontrafficking offense shall be treated as a prior offense  
26 only when the subsequent offense is a nontrafficking offense. For the purposes of  
27 this section, a conviction voided under KRS 218A.275 or 218A.276 shall not

1 constitute a conviction under this chapter;

2 (49) "Sell" means to dispose of a controlled substance to another person for  
3 consideration or in furtherance of commercial distribution;

4 (50) "Serious physical injury" has the same meaning it has in KRS 500.080;

5 (51) "Synthetic cannabinoids or piperazines" means any chemical compound which is  
6 not approved by the United States Food and Drug Administration or, if approved,  
7 which is not dispensed or possessed in accordance with state and federal law, that  
8 contains Benzylpiperazine (BZP); Trifluoromethylphenylpiperazine (TFMPP); 1,1-  
9 Dimethylheptyl-11-hydroxytetrahydrocannabinol (HU-210); 1-Butyl-3-(1-  
10 naphthoyl)indole; 1-Pentyl-3-(1-naphthoyl)indole; dexanabinol (HU-211); or any  
11 compound in the following structural classes:

12 (a) Naphthoylindoles: Any compound containing a 3-(1-naphthoyl)indole  
13 structure with substitution at the nitrogen atom of the indole ring by an alkyl,  
14 haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-  
15 piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group, whether or not further  
16 substituted in the indole ring to any extent and whether or not substituted in  
17 the naphthyl ring to any extent. Examples of this structural class include but  
18 are not limited to JWH-015, JWH-018, JWH-019, JWH-073, JWH-081,  
19 JWH-122, JWH-200, and AM-2201;

20 (b) Phenylacetylindoles: Any compound containing a 3-phenylacetylindole  
21 structure with substitution at the nitrogen atom of the indole ring by an alkyl,  
22 haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-  
23 piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group whether or not further  
24 substituted in the indole ring to any extent and whether or not substituted in  
25 the phenyl ring to any extent. Examples of this structural class include but are  
26 not limited to JWH-167, JWH-250, JWH-251, and RCS-8;

27 (c) Benzoylindoles: Any compound containing a 3-(benzoyl)indole structure with

1 substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl,  
2 alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl,  
3 or 2-(4-morpholinyl)ethyl group whether or not further substituted in the  
4 indole ring to any extent and whether or not substituted in the phenyl ring to  
5 any extent. Examples of this structural class include but are not limited to  
6 AM-630, AM-2233, AM-694, Pravadoline (WIN 48,098), and RCS-4;

7 (d) Cyclohexylphenols: Any compound containing a 2-(3-  
8 hydroxycyclohexyl)phenol structure with substitution at the 5-position of the  
9 phenolic ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl,  
10 cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl  
11 group whether or not substituted in the cyclohexyl ring to any extent.  
12 Examples of this structural class include but are not limited to CP 47,497 and  
13 its C8 homologue (cannabicyclohexanol);

14 (e) Naphthylmethylindoles: Any compound containing a 1H-indol-3-yl-(1-  
15 naphthyl)methane structure with substitution at the nitrogen atom of the indole  
16 ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-  
17 methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group whether or not  
18 further substituted in the indole ring to any extent and whether or not  
19 substituted in the naphthyl ring to any extent. Examples of this structural class  
20 include but are not limited to JWH-175, JWH-184, and JWH-185;

21 (f) Naphthoypyrroles: Any compound containing a 3-(1-naphthoypyrrole  
22 structure with substitution at the nitrogen atom of the pyrrole ring by an alkyl,  
23 haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-  
24 piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group whether or not further  
25 substituted in the pyrrole ring to any extent and whether or not substituted in  
26 the naphthyl ring to any extent. Examples of this structural class include but  
27 are not limited to JWH-030, JWH-145, JWH-146, JWH-307, and JWH-368;

- 1 (g) Naphthylmethylenes: Any compound containing a 1-(1-  
2 naphthylmethyl)indene structure with substitution at the 3-position of the  
3 indene ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl,  
4 1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group whether  
5 or not further substituted in the indene ring to any extent and whether or not  
6 substituted in the naphthyl ring to any extent. Examples of this structural class  
7 include but are not limited to JWH-176;
- 8 (h) Tetramethylcyclopropanoylindoles: Any compound containing a 3-(1-  
9 tetramethylcyclopropoyl)indole structure with substitution at the nitrogen  
10 atom of the indole ring by an alkyl, haloalkyl, cycloalkylmethyl,  
11 cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl  
12 group, whether or not further substituted in the indole ring to any extent and  
13 whether or not further substituted in the tetramethylcyclopropyl ring to any  
14 extent. Examples of this structural class include but are not limited to UR-144  
15 and XLR-11;
- 16 (i) Adamantoylindoles: Any compound containing a 3-(1-adamantoyl)indole  
17 structure with substitution at the nitrogen atom of the indole ring by an alkyl,  
18 haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-  
19 piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group, whether or not further  
20 substituted in the indole ring to any extent and whether or not substituted in  
21 the adamantyl ring system to any extent. Examples of this structural class  
22 include but are not limited to AB-001 and AM-1248; or
- 23 (j) Any other synthetic cannabinoid or piperazine which is not approved by the  
24 United States Food and Drug Administration or, if approved, which is not  
25 dispensed or possessed in accordance with state and federal law;
- 26 (52) "Synthetic cathinones" means any chemical compound which is not approved by the  
27 United States Food and Drug Administration or, if approved, which is not dispensed

1 or possessed in accordance with state and federal law (not including bupropion or  
2 compounds listed under a different schedule) structurally derived from 2-  
3 aminopropan-1-one by substitution at the 1-position with either phenyl, naphthyl, or  
4 thiophene ring systems, whether or not the compound is further modified in one (1)  
5 or more of the following ways:

6 (a) By substitution in the ring system to any extent with alkyl, alkylendioxy,  
7 alkoxy, haloalkyl, hydroxyl, or halide substituents, whether or not further  
8 substituted in the ring system by one (1) or more other univalent substituents.  
9 Examples of this class include but are not limited to 3,4-  
10 Methylenedioxcathinone (bk-MDA);

11 (b) By substitution at the 3-position with an acyclic alkyl substituent. Examples of  
12 this class include but are not limited to 2-methylamino-1-phenylbutan-1-one  
13 (buphedrone);

14 (c) By substitution at the 2-amino nitrogen atom with alkyl, dialkyl, benzyl, or  
15 methoxybenzyl groups, or by inclusion of the 2-amino nitrogen atom in a  
16 cyclic structure. Examples of this class include but are not limited to  
17 Dimethylcathinone, Ethcathinone, and  $\alpha$ -Pyrrolidinopropiophenone ( $\alpha$ -PPP);  
18 or

19 (d) Any other synthetic cathinone which is not approved by the United States  
20 Food and Drug Administration or, if approved, is not dispensed or possessed  
21 in accordance with state or federal law;

22 (53) "Synthetic drugs" means any synthetic cannabinoids or piperazines or any synthetic  
23 cathinones;

24 (54) "Telehealth" has the same meaning it has in KRS 311.550;

25 (55) "Tetrahydrocannabinols" means synthetic equivalents of the substances contained in  
26 the plant, or in the resinous extractives of the plant Cannabis, sp. or synthetic  
27 substances, derivatives, and their isomers with similar chemical structure and

1 pharmacological activity such as the following:

2 (a) Delta 1 cis or trans tetrahydrocannabinol, and their optical isomers;

3 (b) Delta 6 cis or trans tetrahydrocannabinol, and their optical isomers; and

4 (c) Delta 3, 4 cis or trans tetrahydrocannabinol, and its optical isomers;

5 (56) "Traffic," except as provided in KRS 218A.1431, means to manufacture, distribute,  
6 dispense, sell, transfer, or possess with intent to manufacture, distribute, dispense,  
7 or sell a controlled substance;

8 (57) "Transfer" means to dispose of a controlled substance to another person without  
9 consideration and not in furtherance of commercial distribution; and

10 (58) "Ultimate user" means a person who lawfully possesses a controlled substance for  
11 his or her own use or for the use of a member of his or her household or for  
12 administering to an animal owned by him or her or by a member of his or her  
13 household.