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STATE OF RHODE ISLAND

IN GENERAL ASSEMBLY

JANUARY SESSION, A.D. 2015

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A N A C T

RELATING TO FOOD AND DRUGS -- UNIFORM CONTROLLED SUBSTANCES ACT--  
EXCLUSION OF HEMP

Introduced By: Representatives Newberry, and Price

Date Introduced: May 22, 2015

Referred To: House Health, Education & Welfare

It is enacted by the General Assembly as follows:

1           SECTION 1. Section 21-28-1.02 of the General Laws in Chapter 21-28 entitled "Uniform  
2   Controlled Substances Act" is hereby amended to read as follows:

3           **21-28-1.02. Definitions.** -- Unless the context otherwise requires, the words and phrases  
4   as defined in this section are used in this chapter in the sense given them in the following  
5   definitions:

6           (1) "Administer" refers to the direct application of controlled substances to the body of a  
7   patient or research subject by:

8           (i) A practitioner, or, in his or her presence by his or her authorized agent; or

9           (ii) The patient or research subject at the direction and in the presence of the practitioner  
10   whether the application is by injection, inhalation, ingestion, or any other means.

11          (2) "Agent" means an authorized person who acts on behalf of or at the direction of a  
12   manufacturer, wholesaler, distributor, or dispenser; except that these terms do not include a  
13   common or contract carrier or warehouse operator, when acting in the usual and lawful course of  
14   the carrier's or warehouse operator's business.

15          (3) "Apothecary" means a registered pharmacist as defined by the laws of this state and,  
16   where the context requires, the owner of a licensed pharmacy or other place of business where  
17   controlled substances are compounded or dispensed by a registered pharmacist; and includes  
18   registered assistant pharmacists as defined by existing law, but nothing in this chapter shall be

1 construed as conferring on a person who is not registered as a pharmacist any authority, right, or  
2 privilege that is not granted to him or her by the pharmacy laws of the state.

3 (4) "Automated data processing system" means a system utilizing computer software and  
4 hardware for the purposes of record keeping.

5 (5) "Computer" means programmable electronic device capable of multi-functions,  
6 including, but not limited to, storage, retrieval, and processing of information.

7 (6) "Control" means to add a drug or other substance or immediate precursor to a  
8 schedule under this chapter, whether by transfer from another schedule or otherwise.

9 (7) "Controlled substance" means a drug, substance, immediate precursor, or synthetic  
10 drug in schedules I -- V of this chapter. The term shall not include distilled spirits, wine, or malt  
11 beverages, as those terms are defined or used in chapter 1 of title 3, nor tobacco.

12 (8) "Counterfeit substance" means a controlled substance which, or the container or  
13 labeling of which, without authorization bears the trademark, trade name, or other identifying  
14 mark, imprint, number, or device, or any likeness of them, of a manufacturer, distributor, or  
15 dispenser, other than the person or persons who in fact manufactured, distributed, or dispensed  
16 the substance and which thereby falsely purports or is represented to be the product of, or to have  
17 been distributed by, the other manufacturer, distributor, or dispenser, or which substance is  
18 falsely purported to be or represented to be one of the controlled substances by a manufacturer,  
19 distributor, or dispenser.

20 (9) "CRT" means cathode ray tube used to impose visual information on a screen.

21 (10) "Deliver" or "delivery" means the actual, constructive, or attempted transfer of a  
22 controlled substance or imitation controlled substance, whether or not there exists an agency  
23 relationship.

24 (11) "Department" means the department of health of this state.

25 (12) "Depressant or stimulant drug" means:

26 (i) A drug which contains any quantity of:

27 (A) Barbituric acid or derivatives, compounds, mixtures, or preparations of barbituric  
28 acid; and

29 (B) "Barbiturate" or "barbiturates" includes all hypnotic and/or somnifacient drugs,  
30 whether or not derivatives of barbituric acid, except that this definition shall not include bromides  
31 and narcotics.

32 (ii) A drug which contains any quantity of:

33 (A) Amphetamine or any of its optical isomers;

34 (B) Any salt of amphetamine and/or desoxyephedrine or any salt of an optical isomer of

1 amphetamine and/or desoxyephedrine, or any compound, mixture, or preparation of them.

2 (iii) A drug which contains any quantity of coca leaves. "Coca leaves" includes cocaine,  
3 or any compound, manufacture, salt, derivative, mixture, or preparation of coca leaves, except  
4 derivatives of coca leaves, which do not contain cocaine, ecgonine, or substance from which  
5 cocaine or ecgonine may be synthesized or made.

6 (iv) Any other drug or substance which contains any quantity of a substance which the  
7 attorney general of the United States, or the director of health, after investigation, has found to  
8 have, or by regulation designates as having, a potential for abuse because of its depressant or  
9 stimulant effect on the central nervous system.

10 (13) "Director" means the director of health.

11 (14) "Dispense" means to deliver, distribute, leave with, give away, or dispose of a  
12 controlled substance to the ultimate user or human research subject by or pursuant to the lawful  
13 order of a practitioner, including the packaging, labeling, or compounding necessary to prepare  
14 the substance for that delivery.

15 (15) "Dispenser" is a practitioner who delivers a controlled substance to the ultimate user  
16 or human research subject.

17 (16) "Distribute" means to deliver (other than by administering or dispensing) a  
18 controlled substance or an imitation controlled substance and includes actual constructive, or  
19 attempted transfer. "Distributor" means a person who so delivers a controlled substance or an  
20 imitation controlled substance.

21 (17) "Downtime" means that period of time when a computer is not operable.

22 (18) "Drug addicted person" means a person who exhibits a maladaptive pattern of  
23 behavior resulting from drug use, including one or more of the following: impaired control over  
24 drug use; compulsive use; and/or continued use despite harm, and craving.

25 (19) "Drug Enforcement Administration" means the Drug Enforcement Administration  
26 United States Department of Justice or its successor.

27 (20) "Federal law" means the Comprehensive Drug Abuse Prevention and Control Act of  
28 1970, (84 stat. 1236)(see generally 21 U.S.C. § 801 et seq.), and all regulations pertaining to that  
29 federal act.

30 (21) "Hardware" means the fixed component parts of a computer.

31 (22) "Hospital" means an institution as defined in chapter 17 of title 23.

32 (23) "Imitation controlled substance" means a substance that is not a controlled  
33 substance, which by dosage unit, appearance (including color, shape, size, and markings), or by  
34 representations made, would lead a reasonable person to believe that the substance is a controlled

1 substance and, which imitation controlled substances contain substances which if ingested, could  
2 be injurious to the health of a person. In those cases when the appearance of the dosage unit is not  
3 reasonably sufficient to establish that the substance is an "imitation controlled substance" (for  
4 example in the case of powder or liquid), the court or authority concerned should consider, in  
5 addition to all other logically relevant factors, the following factors as related to "representations  
6 made" in determining whether the substance is an "imitation controlled substance":

7 (i) Statement made by an owner, possessor, transferor, recipient, or by anyone else in  
8 control of the substance concerning the nature of the substance, or its use or effect.

9 (ii) Statements made by the owner, possessor, or transferor, to the recipient that the  
10 substance may be resold for substantial profit.

11 (iii) Whether the substance is packaged in a manner reasonably similar to packaging of  
12 illicit controlled substances.

13 (iv) Whether the distribution or attempted distribution included an exchange of or  
14 demand for money or other property as consideration, and whether the amount of the  
15 consideration was substantially greater than the reasonable value of the non-controlled substance.

16 (24) "Immediate precursor" means a substance:

17 (i) Which the director of health has found to be and by regulation designated as being the  
18 principal compound used, or produced primarily for use, in the manufacture of a controlled  
19 substance;

20 (ii) Which is an immediate chemical intermediary used or likely to be used in the  
21 manufacture of those controlled substances; and

22 (iii) The control of which is necessary to prevent, curtail, or limit the manufacture of that  
23 controlled substance.

24 (25) "Laboratory" means a laboratory approved by the department of health as proper to  
25 be entrusted with controlled substances and the use of controlled substances for scientific and  
26 medical purposes and for the purposes of instruction.

27 (26) "Marijuana" means all parts of the plant *cannabis sativa* L., whether growing or not;  
28 the seeds of the plant; the resin extracted from any part of the plant; and every compound,  
29 manufacture, salt, derivative, mixture, or preparation of the plant, its seeds or resin, but shall not  
30 include the mature stalks of the plant, fiber produced from the stalks, oil or cake made from the  
31 seeds of the plant, any other compound, manufacture, salt, derivative, mixture, or preparation of  
32 mature stalks, (except the resin extracted from it), fiber, oil or cake, or the sterilized seed from the  
33 plant which is incapable of germination, nor "hemp", hereby defined as a tall widely cultivated  
34 Asian herb (*Cannabis sativa* of the family Cannabaceae, the hemp family) that has a tough bast

1 [fiber used especially for cordage and that is often separated into a tall loosely branched species](#)  
2 [\(C. sativa\) and a low-growing densely branched species \(C. indica\), nor the fiber of hemp.](#)

3 (27) "Manufacture" means the production, preparation, propagation, cultivation,  
4 compounding, or processing of a drug or other substance, including an imitation controlled  
5 substance, either directly or indirectly or by extraction from substances of natural origin, or  
6 independently by means of chemical synthesis or by a combination of extraction and chemical  
7 synthesis and includes any packaging or repackaging of the substance or labeling or relabeling of  
8 its container in conformity with the general laws of this state except by a practitioner as an  
9 incident to his or her administration or dispensing of the drug or substance in the course of his or  
10 her professional practice.

11 (28) "Manufacturer" means a person who manufactures but does not include an  
12 apothecary who compounds controlled substances to be sold or dispensed on prescriptions.

13 (29) "Narcotic drug" means any of the following, whether produced directly or indirectly  
14 by extraction from substances of vegetable origin, or independently by means of chemical  
15 synthesis or by a combination of extraction and chemical synthesis:

16 (i) Opium and opiates.

17 (ii) A compound, manufacture, salt, derivative, or preparation of opium or opiates.

18 (iii) A substance (and any compound, manufacture, salt, derivative, or preparation of it)  
19 which is chemically identical with any of the substances referred to in paragraphs (i) and (ii) of  
20 this subdivision.

21 (iv) Any other substance which the attorney general of the United States, or his or her  
22 successor, or the director of health, after investigation, has found to have, and by regulation  
23 designates as having, a potential for abuse similar to opium and opiates.

24 (30) "Official written order" means an order written on a form provided for that purpose  
25 by the Drug Enforcement Administration under any laws of the United States making provision  
26 for an official form, if order forms are authorized and required by federal law, and if no order  
27 form is provided then on an official form provided for that purpose by the director of health.

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

31 (32) "Opium poppy" means the plant of the species *papaver somniferum* L., except the  
32 seeds of the plant.

33 (33) "Ounce" means an avoirdupois ounce as applied to solids and semi-solids, and a  
34 fluid ounce as applied to liquids.

1 (34) "Person" means any corporation, association, partnership, or one or more  
2 individuals.

3 (35) "Physical dependence" means a state of adaptation that is manifested by a drug class  
4 specific withdrawal syndrome that can be produced by abrupt cessation, rapid dose reduction,  
5 decreasing blood level of the drug, and/or administration of an antagonist.

6 (36) "Poppy straw" means all parts, except the seeds, of the opium poppy, after mowing.

7 (37) "Practitioner" means:

8 (i) A physician, osteopath, dentist, chiropract, veterinarian, scientific investigator, or  
9 other person licensed, registered or permitted to distribute, dispense, conduct research with  
10 respect to or to administer a controlled substance in the course of professional practice or research  
11 in this state.

12 (ii) A pharmacy, hospital, or other institution licensed, registered or permitted to  
13 distribute, dispense, conduct research with respect to, or to administer a controlled substance in  
14 the course of professional practice or research in this state.

15 (38) "Printout" means a hard copy produced by computer that is readable without the aid  
16 of any special device.

17 (39) "Production" includes the manufacture, planting, cultivation, growing, or harvesting  
18 of a controlled substance.

19 (40) "Researcher" means a person authorized by the director of health to conduct a  
20 laboratory as defined in this chapter.

21 (41) "Sell" includes sale, barter, gift, transfer, or delivery in any manner to another, or to  
22 offer or agree to do the same.

23 (42) "Software" means programs, procedures and storage of required information data.

24 (43) "Synthetic drugs" means any synthetic cannabinoids or piperazines or any synthetic  
25 cathinones as provided for in schedule I.

26 (44) "Ultimate user" means a person who lawfully possesses a controlled substance for  
27 his or her own use or for the use of a member of his or her household, or for administering to an  
28 animal owned by him or her or by a member of his or her household.

29 (45) "Wholesaler" means a person who sells, vends, or distributes at wholesale, or as a  
30 jobber, broker agent, or distributor, or for resale in any manner in this state any controlled  
31 substance.

32 SECTION 2. This act shall take effect upon passage.

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EXPLANATION  
BY THE LEGISLATIVE COUNCIL  
OF

A N A C T

RELATING TO FOOD AND DRUGS -- UNIFORM CONTROLLED SUBSTANCES ACT--  
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1           This act would clearly specify that "hemp" would not be subject to the provisions of the  
2 "uniform controlled substances act", thereby legalizing the cultivation and possession of such  
3 plant.

4           This act would take effect upon passage.

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