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SENATE BILL 94

**52ND LEGISLATURE - STATE OF NEW MEXICO - FIRST SESSION, 2015**

INTRODUCED BY

Cisco McSorley

AN ACT

RELATING TO AGRICULTURE; PROVIDING FOR LICENSING THE GROWING,  
SELLING AND PROCESSING OF INDUSTRIAL HEMP; ESTABLISHING FEES;  
PROVIDING PENALTIES; MAKING AN APPROPRIATION.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF NEW MEXICO:

SECTION 1. [NEW MATERIAL] SHORT TITLE.--Sections 1  
through 8 of this act may be cited as the "Industrial Hemp  
Farming Act".

SECTION 2. [NEW MATERIAL] LEGISLATIVE FINDINGS AND  
PURPOSE.--

A. Industrial hemp is a suitable crop for New  
Mexico, and its production will contribute to the future  
viability of New Mexico agriculture.

B. Allowing industrial hemp production will provide  
farmers an opportunity to sell their products to a marketplace

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1 that pays them a reasonable rate of return for their labor and  
2 capital investments. Farmers in Canada report a rate of return  
3 of eight hundred dollars (\$800) per acre for the crop.

4 C. The infrastructure needed to process industrial  
5 hemp will result in increased business opportunities and new  
6 jobs in communities.

7 D. As a food crop, industrial hemp seeds and oil  
8 produced from the seeds have high nutritional value, including  
9 healthy fats and protein.

10 E. As a fiber crop, industrial hemp can be used in  
11 the manufacture of products such as clothing, building supplies  
12 and animal bedding.

13 F. As a fuel crop, industrial hemp seeds can be  
14 processed into biodiesel, and stalks can be pelletized or  
15 flaked for burning or processed for cellulosic ethanol.  
16 Industrial hemp also expands opportunities for on-farm  
17 renewable energy production.

18 G. The production of industrial hemp can play a  
19 useful agronomic role in farm land management as part of a crop  
20 rotation system.

21 H. In addition to being an efficient  
22 photosynthesizer for converting the greenhouse gases carbon  
23 dioxide and carbon monoxide to oxygen, industrial hemp is fast-  
24 growing and drought-tolerant, making it suitable for the arid  
25 southwest.

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1 I. The purpose of the Industrial Hemp Farming Act  
2 is to establish policy and procedures for growing industrial  
3 hemp in New Mexico so that farmers and other businesses in the  
4 New Mexico agricultural industry can take advantage of this  
5 market opportunity.

6 SECTION 3. [NEW MATERIAL] DEFINITIONS.--As used in the  
7 Industrial Hemp Farming Act:

8 A. "grower" means a licensed industrial hemp  
9 grower; and

10 B. "industrial hemp" means any plant that produces  
11 not more than three-tenths of one percent of delta-9-  
12 tetrahydrocannabinol per weighted unit of flowering tops and  
13 leaves and has a delta-9-tetrahydrocannabinol concentration of  
14 not more than one percent on a dry weight basis.

15 SECTION 4. [NEW MATERIAL] ADMINISTRATIVE DISCOVERY  
16 PROCESS TO DETERMINE RULES TO ENCOURAGE GROWTH AND SALES OF  
17 INDUSTRIAL HEMP--ADMINISTRATION.--The New Mexico department of  
18 agriculture shall:

19 A. monitor the initial phase of research and  
20 development necessary to ensure a viable and legal industrial  
21 hemp industry in the state; and

22 B. ensure the participation by and inclusion of  
23 individual farmers, agricultural cooperatives and businesses in  
24 the rulemaking process.

25 SECTION 5. [NEW MATERIAL] IMPLEMENTATION--FEES.--

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1           A. A person or business planning to grow and sell  
2 industrial hemp seed or industrial hemp fiber shall obtain a  
3 grower's license by submitting an application to the New Mexico  
4 department of agriculture containing the following:

5                   (1) the name and address of the applicant;

6                   (2) the location and legal description of the  
7 land to be used for the production of industrial hemp and the  
8 name and address of the person holding title to the land on  
9 which the industrial hemp will be planted;

10                  (3) any other information required for  
11 completion of a nationwide criminal background check; and

12                  (4) a nonrefundable application or renewal fee  
13 of no more than one hundred fifty dollars (\$150).

14           B. A grower shall maintain records showing:

15                   (1) the origin of the seed purchased and  
16 planted;

17                   (2) the quantity of the seed purchased and  
18 planted;

19                   (3) the amount of industrial hemp harvested  
20 and sold; and

21                   (4) buyers and recipients of the industrial  
22 hemp plants, fiber and seed.

23           C. The New Mexico department of agriculture shall  
24 help to ensure availability of seed. The department shall:

25                   (1) maintain an authorized list of certified

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1 seed sources for industrial hemp;

2 (2) certify industrial hemp seed obtained from  
3 other sources;

4 (3) maintain a list of growers and processors  
5 for whom seed has been provided; and

6 (4) maintain a list of growers and processors.

7 D. The New Mexico department of agriculture may  
8 collaborate with individual farmers, agricultural cooperatives  
9 or businesses to establish an industrial hemp seed bank and  
10 provide seed for a fee that does not exceed ten percent more  
11 than the cost of the seed to growers upon request.

12 E. The New Mexico department of agriculture may  
13 enter into joint powers agreements with an Indian nation, tribe  
14 or pueblo to share information, to provide technical assistance  
15 and to generally cooperate with the Indian nation, tribe or  
16 pueblo to facilitate the production of industrial hemp on  
17 tribal land.

18 F. The New Mexico department of agriculture may  
19 revoke or suspend a license of a grower if there is substantial  
20 evidence of violations of the provisions of the Industrial Hemp  
21 Farming Act or rules adopted pursuant to that act. The  
22 department shall impose fines subsequent to the implementation  
23 of the Industrial Hemp Farming Act.

24 G. Fees collected pursuant to this section are  
25 appropriated to the New Mexico department of agriculture to

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1 carry out the provisions of the Industrial Hemp Farming Act.

2 SECTION 6. [NEW MATERIAL] DEPARTMENT OF PUBLIC  
3 SAFETY--DUTIES AND POWERS.--The department of public safety:

4 A. shall conduct background checks on applicants  
5 requesting licenses upon request by the New Mexico department  
6 of agriculture;

7 B. shall inspect growing fields and processing  
8 facilities upon verifiable evidence that a designated  
9 industrial hemp field is unlicensed and is in violation of the  
10 Industrial Hemp Farming Act;

11 C. shall train law enforcement officers to identify  
12 industrial hemp;

13 D. shall inform the New Mexico department of  
14 agriculture of any criminal offenses regarding the growing or  
15 processing of industrial hemp; and

16 E. may enter into joint powers agreements with an  
17 Indian nation, tribe or pueblo to share information, to provide  
18 technical assistance and to generally cooperate with the Indian  
19 nation, tribe or pueblo to facilitate the production of  
20 industrial hemp on tribal land.

21 SECTION 7. [NEW MATERIAL] COOPERATION BETWEEN  
22 AGENCIES.--The New Mexico department of agriculture and the  
23 department of public safety shall cooperate fully with one  
24 another to implement and enforce the provisions of the  
25 Industrial Hemp Farming Act.

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1           SECTION 8.   ~~[NEW MATERIAL]~~ PENALTY.--A person who  
2 fraudulently obtains a license pursuant to the Industrial Hemp  
3 Farming Act or violates the provisions of the license is guilty  
4 of a fourth degree felony and shall be sentenced pursuant to  
5 the provisions of Section 31-18-15 NMSA 1978.

6           SECTION 9.   Section 30-31-2 NMSA 1978 (being Laws 1972,  
7 Chapter 84, Section 2, as amended) is amended to read:

8           "30-31-2.   DEFINITIONS.--As used in the Controlled  
9 Substances Act:

10           A.   "administer" means the direct application of a  
11 controlled substance by any means to the body of a patient or  
12 research subject by a practitioner or the practitioner's agent;

13           B.   "agent" includes an authorized person who acts  
14 on behalf of a manufacturer, distributor or dispenser.  It does  
15 not include a common or contract carrier, public  
16 warehouseperson or employee of the carrier or warehouseperson;

17           C.   "board" means the board of pharmacy;

18           D.   "bureau" means the narcotic and dangerous drug  
19 section of the criminal division of the United States  
20 department of justice, or its successor agency;

21           E.   "controlled substance" means a drug or substance  
22 listed in Schedules I through V of the Controlled Substances  
23 Act or rules adopted thereto;

24           F.   "counterfeit substance" means a controlled  
25 substance that bears the unauthorized trademark, trade name,

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1 imprint, number, device or other identifying mark or likeness  
2 of a manufacturer, distributor or dispenser other than the  
3 person who in fact manufactured, distributed or dispensed the  
4 controlled substance;

5 G. "deliver" means the actual, constructive or  
6 attempted transfer from one person to another of a controlled  
7 substance or controlled substance analog, whether or not there  
8 is an agency relationship;

9 H. "dispense" means to deliver a controlled  
10 substance to an ultimate user or research subject pursuant to  
11 the lawful order of a practitioner, including the  
12 administering, prescribing, packaging, labeling or compounding  
13 necessary to prepare the controlled substance for that  
14 delivery;

15 I. "dispenser" means a practitioner who dispenses  
16 and includes hospitals, pharmacies and clinics where controlled  
17 substances are dispensed;

18 J. "distribute" means to deliver other than by  
19 administering or dispensing a controlled substance or  
20 controlled substance analog;

21 K. "drug" or "substance" means substances  
22 recognized as drugs in the official United States  
23 pharmacopoeia, official homeopathic pharmacopoeia of the United  
24 States or official national formulary or any respective  
25 supplement to those publications. It does not include devices

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1 or their components, parts or accessories;

2 L. "hashish" means the resin extracted from any  
3 part of marijuana, whether growing or not, and every compound,  
4 manufacture, salt, derivative, mixture or preparation of such  
5 resins;

6 M. "manufacture" means the production, preparation,  
7 compounding, conversion or processing of a controlled substance  
8 or controlled substance analog by extraction from substances of  
9 natural origin or independently by means of chemical synthesis  
10 or by a combination of extraction and chemical synthesis and  
11 includes any packaging or repackaging of the substance or  
12 labeling or relabeling of its container, except that this term  
13 does not include the preparation or compounding of a controlled  
14 substance:

15 (1) by a practitioner as an incident to  
16 administering or dispensing a controlled substance in the  
17 course of the practitioner's professional practice; or

18 (2) by a practitioner, or by the  
19 practitioner's agent under the practitioner's supervision, for  
20 the purpose of or as an incident to research, teaching or  
21 chemical analysis and not for sale;

22 N. "marijuana" means all parts of the plant  
23 cannabis, including any and all varieties, species and  
24 subspecies of the genus Cannabis, whether growing or not, the  
25 seeds thereof and every compound, manufacture, salt,

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1 derivative, mixture or preparation of the plant or its seeds.  
2 It does not include the mature stalks of the plant, hashish,  
3 tetrahydrocannabinols extracted or isolated from marijuana,  
4 fiber produced from the stalks, oil or cake made from the seeds  
5 of the plant, any other compound, manufacture, salt,  
6 derivative, mixture or preparation of the mature stalks, fiber,  
7 oil or cake, or the sterilized seed of the plant that is  
8 incapable of germination or any variety of the species sativa  
9 of the genus Cannabis that produces not more than three-tenths  
10 of one percent of delta-9-tetrahydrocannabinol per weighted  
11 unit of flowering tops and leaves and has a delta-9-  
12 tetrahydrocannabinol concentration of not more than one percent  
13 on a dry weight basis;

14 0. "narcotic drug" means any of the following,  
15 whether produced directly or indirectly by extraction from  
16 substances of vegetable origin or independently by means of  
17 chemical synthesis or by a combination of extraction and  
18 chemical synthesis:

19 (1) opium and opiate and any salt, compound,  
20 derivative or preparation of opium or opiate;

21 (2) any salt, compound, isomer, derivative or  
22 preparation that is a chemical equivalent of any of the  
23 substances referred to in Paragraph (1) of this subsection,  
24 except the isoquinoline alkaloids of opium;

25 (3) opium poppy and poppy straw, including all

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1 parts of the plant of the species *Papaver somniferum* L. except  
2 its seeds; or

3 (4) coca leaves and any salt, compound,  
4 derivative or preparation of coca leaves, any salt, compound,  
5 isomer, derivative or preparation that is a chemical equivalent  
6 of any of these substances except decocainized coca leaves or  
7 extractions of coca leaves that do not contain cocaine or  
8 ecgonine;

9 P. "opiate" means any substance having an  
10 addiction-forming or addiction-sustaining liability similar to  
11 morphine or being capable of conversion into a drug having  
12 addiction-forming or addiction-sustaining liability. "Opiate"  
13 does not include, unless specifically designated as controlled  
14 under Section 30-31-5 NMSA 1978, the dextrorotatory isomer of  
15 3-methoxy-n-methylmorphinan and its salts, dextromethorphan.  
16 "Opiate" does include its racemic and levorotatory forms;

17 Q. "person" means an individual, partnership,  
18 corporation, association, institution, political subdivision,  
19 government agency or other legal entity;

20 R. "practitioner" means a physician, certified  
21 advanced practice chiropractic physician, doctor of oriental  
22 medicine, dentist, physician assistant, certified nurse  
23 practitioner, clinical nurse specialist, certified nurse-  
24 midwife, prescribing psychologist, veterinarian, euthanasia  
25 technician, pharmacist, pharmacist clinician or other person

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1 licensed or certified to prescribe and administer drugs that  
2 are subject to the Controlled Substances Act;

3 S. "prescription" means an order given individually  
4 for the person for whom is prescribed a controlled substance,  
5 either directly from a licensed practitioner or the  
6 practitioner's agent to the pharmacist, including by means of  
7 electronic transmission, or indirectly by means of a written  
8 order signed by the prescriber, bearing the name and address of  
9 the prescriber, the prescriber's license classification, the  
10 name and address of the patient, the name and quantity of the  
11 drug prescribed, directions for use and the date of issue and  
12 in accordance with the Controlled Substances Act or rules  
13 adopted thereto;

14 T. "scientific investigator" means a person  
15 registered to conduct research with controlled substances in  
16 the course of the person's professional practice or research  
17 and includes analytical laboratories;

18 U. "ultimate user" means a person who lawfully  
19 possesses a controlled substance for the person's own use or  
20 for the use of a member of the person's household or for  
21 administering to an animal under the care, custody and control  
22 of the person or by a member of the person's household;

23 V. "drug paraphernalia" means all equipment,  
24 products and materials of any kind that are used, intended for  
25 use or designed for use in planting, propagating, cultivating,

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1 growing, harvesting, manufacturing, compounding, converting,  
2 producing, processing, preparing, testing, analyzing,  
3 packaging, repackaging, storing, containing, concealing,  
4 injecting, ingesting, inhaling or otherwise introducing into  
5 the human body a controlled substance or controlled substance  
6 analog in violation of the Controlled Substances Act. It  
7 includes:

8 (1) kits used, intended for use or designed  
9 for use in planting, propagating, cultivating, growing or  
10 harvesting any species of plant that is a controlled substance  
11 or controlled substance analog or from which a controlled  
12 substance can be derived;

13 (2) kits used, intended for use or designed  
14 for use in manufacturing, compounding, converting, producing,  
15 processing or preparing controlled substances or controlled  
16 substance analogs;

17 (3) isomerization devices used, intended for  
18 use or designed for use in increasing the potency of any  
19 species of plant that is a controlled substance;

20 (4) testing equipment used, intended for use  
21 or designed for use in identifying or in analyzing the  
22 strength, effectiveness or purity of controlled substances or  
23 controlled substance analogs;

24 (5) scales or balances used, intended for use  
25 or designed for use in weighing or measuring controlled

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1 substances or controlled substance analogs;

2 (6) diluents and adulterants, such as quinine  
3 hydrochloride, mannitol, mannite dextrose and lactose, used,  
4 intended for use or designed for use in cutting controlled  
5 substances or controlled substance analogs;

6 (7) separation gins and sifters used, intended  
7 for use or designed for use in removing twigs and seeds from,  
8 or in otherwise cleaning and refining, marijuana;

9 (8) blenders, bowls, containers, spoons and  
10 mixing devices used, intended for use or designed for use in  
11 compounding controlled substances or controlled substance  
12 analogs;

13 (9) capsules, balloons, envelopes and other  
14 containers used, intended for use or designed for use in  
15 packaging small quantities of controlled substances or  
16 controlled substance analogs;

17 (10) containers and other objects used,  
18 intended for use or designed for use in storing or concealing  
19 controlled substances or controlled substance analogs;

20 (11) hypodermic syringes, needles and other  
21 objects used, intended for use or designed for use in  
22 parenterally injecting controlled substances or controlled  
23 substance analogs into the human body;

24 (12) objects used, intended for use or  
25 designed for use in ingesting, inhaling or otherwise

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1 introducing marijuana, cocaine, hashish or hashish oil into the  
2 human body, such as:

3 (a) metal, wooden, acrylic, glass,  
4 stone, plastic or ceramic pipes, with or without screens,  
5 permanent screens, hashish heads or punctured metal bowls;

6 (b) water pipes;

7 (c) carburetion tubes and devices;

8 (d) smoking and carburetion masks;

9 (e) roach clips, meaning objects used to  
10 hold burning material, such as a marijuana cigarette, that has  
11 become too small to hold in the hand;

12 (f) miniature cocaine spoons and cocaine  
13 vials;

14 (g) chamber pipes;

15 (h) carburetor pipes;

16 (i) electric pipes;

17 (j) air-driven pipes;

18 (k) chilams;

19 (l) bongs; or

20 (m) ice pipes or chillers; and

21 (13) in determining whether an object is drug  
22 paraphernalia, a court or other authority should consider, in  
23 addition to all other logically relevant factors, the  
24 following:

25 (a) statements by the owner or by anyone

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1 in control of the object concerning its use;

2 (b) the proximity of the object, in time  
3 and space, to a direct violation of the Controlled Substances  
4 Act or any other law relating to controlled substances or  
5 controlled substance analogs;

6 (c) the proximity of the object to  
7 controlled substances or controlled substance analogs;

8 (d) the existence of any residue of a  
9 controlled substance or controlled substance analog on the  
10 object;

11 (e) instructions, written or oral,  
12 provided with the object concerning its use;

13 (f) descriptive materials accompanying  
14 the object that explain or depict its use;

15 (g) the manner in which the object is  
16 displayed for sale; and

17 (h) expert testimony concerning its use;

18 W. "controlled substance analog" means a substance  
19 other than a controlled substance that has a chemical structure  
20 substantially similar to that of a controlled substance in  
21 Schedule I, II, III, IV or V or that was specifically designed  
22 to produce effects substantially similar to that of controlled  
23 substances in Schedule I, II, III, IV or V. Examples of  
24 chemical classes in which controlled substance analogs are  
25 found include the following:

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- 1 (1) phenethylamines;
- 2 (2) N-substituted piperidines;
- 3 (3) morphinans;
- 4 (4) ecgonines;
- 5 (5) quinazolinones;
- 6 (6) substituted indoles; and
- 7 (7) arylcycloalkylamines.

8 Specifically excluded from the definition of "controlled  
9 substance analog" are those substances that are generally  
10 recognized as safe and effective within the meaning of the  
11 Federal Food, Drug and Cosmetic Act or have been manufactured,  
12 distributed or possessed in conformance with the provisions of  
13 an approved new drug application or an exemption for  
14 investigational use within the meaning of Section 505 of the  
15 Federal Food, Drug and Cosmetic Act;

16 X. "human consumption" includes application,  
17 injection, inhalation, ingestion or any other manner of  
18 introduction;

19 Y. "drug-free school zone" means a public school,  
20 parochial school or private school or property that is used for  
21 a public, parochial or private school purpose and the area  
22 within one thousand feet of the school property line, but it  
23 does not mean any post-secondary school; and

24 Z. "valid practitioner-patient relationship" means  
25 a professional relationship, as defined by the practitioner's

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1 licensing board, between the practitioner and the patient."

2 SECTION 10. APPROPRIATION.--

3 A. One hundred fifty thousand dollars (\$150,000) is  
4 appropriated from the general fund for expenditure in fiscal  
5 year 2016 in the following amounts for the following purposes:

6 (1) one hundred thousand dollars (\$100,000) to  
7 the board of regents of New Mexico state university to  
8 establish and maintain databases, a seed bank and a seed  
9 certification program pursuant to the Industrial Hemp Farming  
10 Act; and

11 (2) fifty thousand dollars (\$50,000) to the  
12 department of public safety to train law enforcement officers  
13 to identify industrial hemp and to implement a law enforcement  
14 program regarding the growth, sale and processing of industrial  
15 hemp pursuant to the Industrial Hemp Farming Act.

16 B. Any unexpended or unencumbered balance remaining  
17 at the end of fiscal year 2016 shall revert to the general  
18 fund.

19 SECTION 11. EFFECTIVE DATE.--The effective date of the  
20 provisions of this act is July 1, 2015.

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