1	HOUSE BILL 565
2	50TH LEGISLATURE - STATE OF NEW MEXICO - FIRST SESSION, 2011
3	INTRODUCED BY
4	Ray Begaye
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10	AN ACT
11	RELATING TO AGRICULTURE; PROVIDING FOR LICENSING THE GROWING,
12	SELLING AND PROCESSING OF INDUSTRIAL HEMP; ESTABLISHING FEES;
13	PROVIDING PENALTIES; MAKING AN APPROPRIATION.
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15	BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF NEW MEXICO:
16	SECTION 1. [<u>NEW MATERIAL</u>] SHORT TITLESections 1
17	through 8 of this act may be cited as the "Industrial Hemp
18	Farming Act".
19	SECTION 2. [<u>NEW MATERIAL</u>] LEGISLATIVE FINDINGS AND
20	PURPOSE
21	A. Industrial hemp is a suitable crop for New
22	Mexico, and its production will contribute to the future
23	viability of New Mexico agriculture.
24	B. Allowing industrial hemp production will provide
25	farmers with an opportunity to sell their products to a
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marketplace that pays them a reasonable rate of return for their labor and capital investments. Farmers in Canada report a rate of return of eight hundred dollars (\$800) per acre for the crop.

C. The infrastructure needed to process industrial hemp will result in increased business opportunities and new 6 7 jobs in our communities.

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

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

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

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

н. In addition to being an efficient photosynthesizer for converting the greenhouse gases carbon dioxide and carbon monoxide to oxygen, industrial hemp is fast growing and drought tolerant, making it suitable for the arid .182570.1 - 2 -

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

SECTION 3. [<u>NEW MATERIAL</u>] DEFINITIONS.--As used in the Industrial Hemp Farming Act:

9 A. "grower" means a licensed industrial hemp 10 grower; and

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

SECTION 4. [<u>NEW MATERIAL</u>] ADMINISTRATIVE DISCOVERY PROCESS TO DETERMINE RULES TO ENCOURAGE GROWTH AND SALES OF INDUSTRIAL HEMP--ADMINISTRATION.--The New Mexico department of agriculture shall:

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

B. ensure the participation by and inclusion of individual farmers, agricultural cooperatives and businesses in .182570.1

1 the rulemaking process.

2 SECTION 5. [NEW MATERIAL] IMPLEMENTATION--FEES.--3 A person or business planning to grow and sell Α. 4 industrial hemp seed or industrial hemp fiber shall obtain a 5 grower's license by submitting an application to the New Mexico department of agriculture containing the following: 6 7 (1)the name and address of the applicant; (2) the location and legal description of the 8 9 land to be used for the production of industrial hemp and the name and address of the person holding title to the land on 10 which the industrial hemp will be planted; 11 12 (3) any other information required for 13 completion of a nationwide criminal background check; and 14 (4) a nonrefundable application or renewal fee of no more than one hundred fifty dollars (\$150). 15 A grower shall maintain records showing: 16 Β. the origin of the seed purchased and 17 (1)18 planted; 19 (2) the quantity of the seed purchased and 20 planted; (3) the amount of industrial hemp harvested 21 and sold; and 22 buyers and recipients of the industrial (4) 23 hemp plants, fiber and seed. 24 The New Mexico department of agriculture shall 25 С. .182570.1 - 4 -

1 help to ensure availability of seed. The department shall: 2 (1)maintain an authorized list of certified 3 seed sources for industrial hemp; certify industrial hemp seed obtained from 4 (2) 5 other sources: maintain a list of growers and processors 6 (3) 7 for whom seed has been provided; and maintain a list of growers and processors. 8 (4) 9 D. The New Mexico department of agriculture may collaborate with individual farmers, agricultural cooperatives 10 or businesses to establish an industrial hemp seed bank and 11 12 provide seed for a fee that does not exceed ten percent more than the cost of the seed to growers upon request. 13 14 Ε. The New Mexico department of agriculture may enter into joint powers agreements with an Indian nation, tribe 15 or pueblo to share information, to provide technical assistance 16 and to generally cooperate with the Indian nation, tribe or 17 pueblo to facilitate the production of industrial hemp on 18 19 tribal land. 20 F. The New Mexico department of agriculture may revoke or suspend the license of a grower if there is 21 substantial evidence of violations of the provisions of the 22 Industrial Hemp Farming Act or rules adopted pursuant to that 23 The department shall impose fines subsequent to the act. 24 implementation of the Industrial Hemp Farming Act. 25

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1 G. Fees collected pursuant to this section are 2 appropriated to the New Mexico department of agriculture to carry out the provisions of the Industrial Hemp Farming Act. 3 SECTION 6. [NEW MATERIAL] DEPARTMENT OF PUBLIC 4 5 SAFETY--DUTIES AND POWERS.--The department of public safety: shall conduct background checks on applicants 6 Α. 7 requesting licenses upon request by the New Mexico department 8 of agriculture; 9 Β. shall inspect growing fields and processing facilities upon verifiable evidence that a designated 10 11 industrial hemp field is unlicensed and is in violation of the 12 Industrial Hemp Farming Act; 13 C. shall train law enforcement officers to identify 14 industrial hemp; shall inform the New Mexico department of 15 D. agriculture of any criminal offenses regarding the growing or 16 17 processing of industrial hemp; and 18 Ε. may enter into joint powers agreements with an 19 Indian nation, tribe or pueblo to share information, to provide 20 technical assistance and to generally cooperate with the Indian nation, tribe or pueblo to facilitate the production of 21 industrial hemp on tribal land. 22 SECTION 7. [NEW MATERIAL] COOPERATION BETWEEN 23 AGENCIES. -- The New Mexico department of agriculture and the 24 25 department of public safety shall cooperate fully with one .182570.1

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another to implement and enforce the provisions of the
 Industrial Hemp Farming Act.

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

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

10 "30-31-2. DEFINITIONS.--As used in the Controlled 11 Substances Act:

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

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

C. "board" means the board of pharmacy;

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

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

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F. "counterfeit substance" means a controlled substance that bears the unauthorized trademark, trade name, imprint, number, device or other identifying mark or likeness of a manufacturer, distributor or dispenser other than the person who in fact manufactured, distributed or dispensed the controlled substance;

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

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

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

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

K. "drug" or "substance" means substances recognized as drugs in the official United States pharmacopoeia, official homeopathic pharmacopoeia of the United .182570.1

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States or official national formulary or any respective supplement to those publications. It does not include devices or their components, parts or accessories;

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

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

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

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

N. "marijuana" means all parts of the plant cannabis, including any and all varieties, species and .182570.1 - 9 -

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1 subspecies of the genus Cannabis, whether growing or not, the 2 seeds thereof and every compound, manufacture, salt, 3 derivative, mixture or preparation of the plant or its seeds. It does not include the mature stalks of the plant, hashish, 4 5 tetrahydrocannabinols extracted or isolated from marijuana, fiber produced from the stalks, oil or cake made from the seeds 6 7 of the plant, any other compound, manufacture, salt, derivative, mixture or preparation of the mature stalks, fiber, 8 9 oil or cake, or the sterilized seed of the plant that is incapable of germination; or any variety of the species sativa 10 of the genus Cannabis that produces not more than three-tenths 11 12 of one percent of delta 9 tetrahydrocannabinol per weighted unit of flowering tops and leaves and that has a delta 9 13 tetrahydrocannabinol concentration of not more than one percent 14 on a dry weight basis; 15

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

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

(2) any salt, compound, isomer, derivative or preparation that is a chemical equivalent of any of the substances referred to in Paragraph (1) of this subsection, .182570.1

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except the isoquinoline alkaloids of opium;

2 (3) opium poppy and poppy straw, including all
3 parts of the plant of the species Papaver somniferum L. except
4 its seeds; or

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

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

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

R. "practitioner" means a physician, certified advanced practice chiropractic physician, doctor of oriental medicine, dentist, physician assistant, certified nurse practitioner, clinical nurse specialist, certified nurse-.182570.1

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midwife, prescribing psychologist, veterinarian, euthanasia technician, pharmacist, pharmacist clinician or other person licensed or certified to prescribe and administer drugs that are subject to the Controlled Substances Act;

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

"scientific investigator" means a person т. registered to conduct research with controlled substances in the course of the person's professional practice or research and includes analytical laboratories;

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

"drug paraphernalia" means all equipment, V. .182570.1

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1 products and materials of any kind that are used, intended for 2 use or designed for use in planting, propagating, cultivating, 3 growing, harvesting, manufacturing, compounding, converting, producing, processing, preparing, testing, analyzing, 4 packaging, repackaging, storing, containing, concealing, 5 injecting, ingesting, inhaling or otherwise introducing into 6 7 the human body a controlled substance or controlled substance 8 analog in violation of the Controlled Substances Act. Tt. 9 includes:

10 (1) kits used, intended for use or designed 11 for use in planting, propagating, cultivating, growing or 12 harvesting any species of plant that is a controlled substance 13 or controlled substance analog or from which a controlled 14 substance can be derived;

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

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

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

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scales or balances used, intended for use 1 (5) 2 or designed for use in weighing or measuring controlled 3 substances or controlled substance analogs; diluents and adulterants, such as quinine 4 (6) 5 hydrochloride, mannitol, mannite dextrose and lactose, used, intended for use or designed for use in cutting controlled 6 7 substances or controlled substance analogs; separation gins and sifters used, intended 8 (7) 9 for use or designed for use in removing twigs and seeds from, or in otherwise cleaning and refining, marijuana; 10 blenders, bowls, containers, spoons and (8) 11 12 mixing devices used, intended for use or designed for use in compounding controlled substances or controlled substance 13 14 analogs; capsules, balloons, envelopes and other (9) 15 containers used, intended for use or designed for use in 16 packaging small quantities of controlled substances or 17 controlled substance analogs; 18 19 (10)containers and other objects used, 20 intended for use or designed for use in storing or concealing controlled substances or controlled substance analogs; 21 (11)hypodermic syringes, needles and other 22 objects used, intended for use or designed for use in 23 parenterally injecting controlled substances or controlled 24 substance analogs into the human body; 25 .182570.1 - 14 -

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1	(12) objects used, intended for use or
2	designed for use in ingesting, inhaling or otherwise
3	introducing marijuana, cocaine, hashish or hashish oil into the
4	human body, such as:
5	(a) metal, wooden, acrylic, glass,
6	stone, plastic or ceramic pipes, with or without screens,
7	permanent screens, hashish heads or punctured metal bowls;
8	(b) water pipes;
9	(c) carburetion tubes and devices;
10	(d) smoking and carburetion masks;
11	(e) roach clips, meaning objects used to
12	hold burning material, such as a marijuana cigarette, that has
13	become too small to hold in the hand;
14	(f) miniature cocaine spoons and cocaine
15	vials;
16	(g) chamber pipes;
17	(h) carburetor pipes;
18	(i) electric pipes;
19	(j) air-driven pipes;
20	(k) chilams;
21	(1) bongs; or
22	(m) ice pipes or chillers; and
23	(13) in determining whether an object is drug
24	paraphernalia, a court or other authority should consider, in
25	addition to all other logically relevant factors, the
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1 following: 2 (a) statements by the owner or by anyone in control of the object concerning its use; 3 the proximity of the object, in time 4 (b) and space, to a direct violation of the Controlled Substances 5 Act or any other law relating to controlled substances or 6 7 controlled substance analogs; 8 (c) the proximity of the object to controlled substances or controlled substance analogs; 9 (d) the existence of any residue of a 10 controlled substance or controlled substance analog on the 11 12 object; instructions, written or oral, (e) 13 14 provided with the object concerning its use; descriptive materials accompanying (f) 15 the object that explain or depict its use; 16 the manner in which the object is 17 (g) displayed for sale; and 18 19 (h) expert testimony concerning its use; "controlled substance analog" means a substance 20 W. other than a controlled substance that has a chemical structure 21 substantially similar to that of a controlled substance in 22 Schedule I, II, III, IV or V or that was specifically designed 23 to produce effects substantially similar to that of controlled 24 substances in Schedule I, II, III, IV or V. Examples of 25 .182570.1 - 16 -

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1	chemical classes in which controlled substance analogs are
2	found include the following:
3	<pre>(1) phenethylamines;</pre>
4	(2) N-substituted piperidines;
5	<pre>(3) morphinans;</pre>
6	(4) ecgonines;
7	(5) quinazolinones;
8	(6) substituted indoles; and
9	(7) arylcycloalkylamines.
10	Specifically excluded from the definition of "controlled
11	substance analog" are those substances that are generally
12	recognized as safe and effective within the meaning of the
13	Federal Food, Drug and Cosmetic Act or have been manufactured,
14	distributed or possessed in conformance with the provisions of
15	an approved new drug application or an exemption for
16	investigational use within the meaning of Section 505 of the
17	Federal Food, Drug and Cosmetic Act;
18	X. "human consumption" includes application,
19	injection, inhalation, ingestion or any other manner of
20	introduction;
21	Y. "drug-free school zone" means a public school,
22	parochial school or private school or property that is used for
23	a public, parochial or private school purpose and the area
24	within one thousand feet of the school property line, but it
25	does not mean any post-secondary school; and
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1 Ζ. "valid practitioner-patient relationship" means 2 a professional relationship, as defined by the practitioner's licensing board, between the practitioner and the patient." 3 SECTION 10. APPROPRIATION.--4 5 One hundred fifty thousand dollars (\$150,000) is Α. appropriated from the general fund for expenditure in fiscal 6 7 year 2012 for the following: 8 one hundred thousand dollars (\$100,000) to (1)9 the board of regents of New Mexico state university to establish and maintain databases, a seed bank and a seed 10 11 certification program pursuant to the Industrial Hemp Farming 12 Act; and fifty thousand dollars (\$50,000) to the 13 (2) 14 department of public safety to train law enforcement officers to identify industrial hemp and to implement a law enforcement 15 program regarding the growth, sale and processing of industrial 16 hemp pursuant to the Industrial Hemp Farming Act. 17 18 Β. Any unexpended or unencumbered balance remaining 19 at the end of fiscal year 2012 shall revert to the general 20 fund. EFFECTIVE DATE.--The effective date of the SECTION 11. 21 provisions of this act is July 1, 2011. 22 - 18 -23 24 25

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