SENATE BILL 300

45TH LEGISLATURE - STATE OF NEW MEXICO - SECOND SESSION, 2002

INTRODUCED BY

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AN ACT

RELATING TO PSYCHOLOGISTS; GRANTING PRESCRIPTIVE AUTHORITY TO CERTAIN PSYCHOLOGISTS; PROVIDING QUALIFICATIONS AND LIMITATIONS; REQUIRING MALPRACTICE INSURANCE.

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

Section 1. Section 26-1-2 NMSA 1978 (being Laws 1967, Chapter 23, Section 2, as amended) is amended to read:

"26-1-2. DEFINITIONS.--As used in the New Mexico Drug, Device and Cosmetic Act:

A. "board" means the board of pharmacy or its duly authorized agent;

B. "person" includes individual, partnership, corporation, association, institution or establishment;

C. "biological product" means any virus, therapeutic serum, toxin, antitoxin or analogous product

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applicable to the prevention, treatment or cure of diseases or injuries of man and domestic animals and, as used within the meaning of this definition:

(1) a "virus" is interpreted to be a product containing the minute living cause of an infectious disease and includes filterable viruses, bacteria, rickettsia, fungi and protozoa;

(2) a "therapeutic serum" is a product obtained from blood by removing the clot or clot components and the blood cells;

(3) a "toxin" is a product containing a soluble substance poisonous to laboratory animals or man in doses of one milliliter or less of the product and having the property, following the injection of nonfatal doses into an animal, or causing to be produced therein another soluble substance that specifically neutralizes the poisonous substance and that is demonstrable in the serum of the animal thus immunized; and

(4) an "antitoxin" is a product containing the soluble substance in serum or other body fluid of an immunized animal that specifically neutralizes the toxin against which the animal is immune;

D. "controlled substance" means any drug, substance or immediate precursor enumerated in Schedules I through V of the Controlled Substances Act;

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E. "drug" means:

(1) articles recognized in an official compendium;

(2) articles intended for use in the diagnosis, cure, mitigation, treatment or prevention of disease in man or other animals and includes the domestic animal biological products regulated under the federal Virus-Serum-Toxin Act, 37 Stat 832-833, 21 U.S.C. 151-158 and the biological products applicable to man regulated under Federal 58 Stat 690, as amended, 42 U.S.C. 216, Section 351, 58 Stat 702, as amended, and 42 U.S.C. 262;

(3) articles other than food that affect the structure or any function of the body of man or other animals; and

(4) articles intended for use as a component of Paragraph (1), (2) or (3) of this subsection, but does not include devices or their component parts or accessories;

F. "dangerous drug" means a drug, other than a controlled substance enumerated in Schedule I of the Controlled Substances Act, that because of a potentiality for harmful effect or the method of its use or the collateral measures necessary to its use is not safe except under the supervision of a practitioner licensed by law to direct the use of such drug and hence for which adequate directions for use cannot be prepared. "Adequate directions for use" means

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directions under which the layman can use a drug or device safely and for the purposes for which it is intended. A drug shall be dispensed only upon the prescription of a practitioner licensed by law to administer or prescribe such drug if it:

(1) is a habit-forming drug and contains any quantity of a narcotic or hypnotic substance or a chemical derivative of such substance that has been found under the federal act and the board to be habit forming;

(2) because of its toxicity or other potential for harmful effect or the method of its use or the collateral measures necessary to its use is not safe for use except under the supervision of a practitioner licensed by law to administer or prescribe the drug;

(3) is limited by an approved application by Section 505 of the federal act to the use under the professional supervision of a practitioner licensed by law to administer or prescribe the drug;

(4) bears the legend: "Caution: federal law prohibits dispensing without prescription.";

(5) bears the legend: "Caution: federal law restricts this drug to use by or on the order of a licensed veterinarian."; or

(6) bears the legend "RX only";

G. "counterfeit drug" means a drug other than a

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controlled substance that, or the container or labeling of which, without authorization, bears the trademark, trade name or other identifying mark, imprint or device or any likeness of a drug manufacturer, processor, packer or distributor other than the person who manufactured, processed, packed or distributed the drug and that falsely purports or is represented to be the product of or to have been packed or distributed by such other drug manufacturer, processor, packer or distributor;

H. "device", except when used in Subsection P of this section and in Subsection G of Section 26-1-3, Subsection L and Paragraph (4) of Subsection A of Section 26-1-11 and Subsection C of Section 26-1-24 NMSA 1978, means an instrument, apparatus, implement, machine, contrivance, implant, in vitro reagent or other similar or related article, including any component, part or accessory, that is:

(1) recognized in an official compendium;

(2) intended for use in the diagnosis of disease or other conditions or in the cure, mitigation, treatment or prevention of disease in man or other animals; or

(3) intended to affect the structure or a function of the body of man or other animals and that does not achieve any of its principal intended purposes through chemical action within or on the body of man or other animals and that is not dependent on being metabolized for achievement

of any of its principal intended purposes;

I. "prescription" means an order given individually for the person for whom prescribed, either directly from the prescriber to the pharmacist or indirectly by means of a written order signed by the prescriber, and bearing the name and address of the prescriber, his 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. No person other than a practitioner shall prescribe or write a prescription;

J. "practitioner" means a physician, doctor of oriental medicine, dentist, veterinarian, certified nurse practitioner, clinical nurse specialist, pharmacist, pharmacist clinician, certified nurse-midwife, prescribing psychologist or other person licensed or certified to prescribe and administer drugs that are subject to the New Mexico Drug, Device and Cosmetic Act;

K. "cosmetic" means:

(1) articles intended to be rubbed, poured, sprinkled or sprayed on, introduced into or otherwise applied to the human body or any part thereof for cleansing, beautifying, promoting attractiveness or altering the appearance; and

(2) articles intended for use as a component of any articles enumerated in Paragraph (1) of this

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subsection, except that the term shall not include soap;

L. "official compendium" means the official United States pharmacopoeia national formulary or the official homeopathic pharmacopoeia of the United States or any supplement to either of them;

M. "label" means a display of written, printed or graphic matter upon the immediate container of an article. A requirement made by or under the authority of the New Mexico Drug, Device and Cosmetic Act that any word, statement or other information appear on the label shall not be considered to be complied with unless the word, statement or other information also appears on the outside container or wrapper, if any, of the retail package of the article or is easily legible through the outside container or wrapper;

N. "immediate container" does not include package liners;

O. "labeling" means all labels and other written, printed or graphic matter:

(1) on an article or its containers or wrappers; or

(2) accompanying an article;

P. "misbranded" means a label to an article that is misleading. In determining whether the label is misleading, there shall be taken into account, among other things, not only representations made or suggested by

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statement, word, design, device or any combination of the foregoing, but also the extent to which the label fails to reveal facts material in the light of such representations or material with respect to consequences that may result from the use of the article to which the label relates under the conditions of use prescribed in the label or under such conditions of use as are customary or usual;

Q. "advertisement" means all representations disseminated in any manner or by any means, other than by labeling, for the purpose of inducing, or that are likely to induce, directly or indirectly, the purchase of drugs, devices or cosmetics;

R. "antiseptic", when used in the labeling or advertisement of an antiseptic, shall be considered to be a representation that it is a germicide, except in the case of a drug purporting to be or represented as an antiseptic for inhibitory use as a wet dressing, ointment, dusting powder or such other use as involves prolonged contact with the body;

S. "new drug" means any drug:

(1) the composition of which is such that the drug is not generally recognized, among experts qualified by scientific training and experience to evaluate the safety and efficacy of drugs, as safe and effective for use under the conditions prescribed, recommended or suggested in the labeling thereof; or

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(2) the composition of which is such that the drug, as a result of investigation to determine its safety and efficacy for use under such conditions, has become so recognized, but that has not, otherwise than in such investigations, been used to a material extent or for a material time under such conditions;

T. "contaminated with filth" applies to a drug, device or cosmetic not securely protected from dirt, dust and, as far as may be necessary by all reasonable means, from all foreign or injurious contaminations, or a drug, device or cosmetic found to contain dirt, dust, foreign or injurious contamination or infestation;

U. "selling of drugs, devices or cosmetics" shall be considered to include the manufacture, production, processing, packing, exposure, offer, possession and holding of any such article for sale and the sale and the supplying or applying of any such article in the conduct of a drug or cosmetic establishment;

V. "color additive" means a material that:

(1) is a dye, pigment or other substance made by a process of synthesis or similar artifice or extracted, isolated or otherwise derived, with or without intermediate or final change of identity, from a vegetable, mineral, animal or other source; or

(2) when added or applied to a drug or

cosmetic or to the human body or a part thereof, is capable, alone or through reaction with other substances, of imparting color thereto; except that such term does not include any material that has been or hereafter is exempted under the federal act;

W. "federal act" means the Federal Food, Drug and Cosmetic Act;

X. "restricted device" means a device for which the sale, distribution or use is lawful only upon the written or oral authorization of a practitioner licensed by law to administer, prescribe or use the device and for which the federal food and drug administration requires special training or skills of the practitioner to use or prescribe. This definition does not include custom devices defined in the federal act and exempt from performance standards or premarket approval requirements under Section 520(b) of the federal act; and

Y. "prescription device" means a device that, because of its potential for harm, the method of its use or the collateral measures necessary to its use, is not safe except under the supervision of a practitioner licensed in this state to direct the use of such device and for which "adequate directions for use" cannot be prepared, but that bears the label: "Caution: federal law restricts this device to sale by or on the order of a _____", the blank to be

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filled with the word "physician", "doctor of oriental medicine", "dentist", "veterinarian", "certified nurse practitioner", "clinical nurse specialist", "pharmacist", "pharmacist clinician", "certified nurse-midwife" or with the descriptive designation of any other practitioner licensed in this state to use or order the use of the device."

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

"30-31-2. DEFINITIONS.--As used in the Controlled 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 his 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 warehouseman or employee of the carrier or warehouseman;

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;

F. "counterfeit substance" means a controlled

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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 States or official national formulary or any respective

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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 his administering or dispensing of a controlled substance in the course of his professional practice; or

(2) by a practitioner, or by his agent under his 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 subspecies of the genus cannabis, whether growing or not, the seeds thereof and every compound, manufacture, salt,

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derivative, mixture or preparation of the plant or its seeds. It does not include the mature stalks of the plant, hashish, tetrahydrocannabinols extracted or isolated from marijuana, fiber produced from the stalks, oil or cake made from the seeds of the plant, any other compound, manufacture, salt, derivative, mixture or preparation of the mature stalks, fiber, oil or cake, or the sterilized seed of the plant that is incapable of germination;

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, except the isoquinoline alkaloids of opium;

(3) opium poppy and poppy straw, including all parts of the plant of the species Papaver somniferum L. except 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

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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, doctor of oriental medicine, dentist, physician assistant, certified nurse practitioner, clinical nurse specialist, certified nurse-midwife, prescribing psychologist, veterinarian, pharmacist, pharmacist clinician or other person licensed or certified to prescribe and administer drugs that are subject to the Controlled Substances Act;

S. "prescription" means an order given individually for the person for whom is prescribed a controlled substance, either directly from the prescriber to the pharmacist or indirectly by means of a written order

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signed by the prescriber, in accordance with the Controlled Substances Act or rules adopted thereto;

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

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

V. "drug paraphernalia" means all equipment, products and materials of any kind that are used, intended for use or designed for use in planting, propagating, cultivating, growing, harvesting, manufacturing, compounding, converting, producing, processing, preparing, testing, analyzing, packaging, repackaging, storing, containing, concealing, injecting, ingesting, inhaling or otherwise introducing into the human body a controlled substance or controlled substance analog in violation of the Controlled Substances Act. It includes:

(1) kits used, intended for use or designed for use in planting, propagating, cultivating, growing or harvesting any species of plant that is a controlled substance or controlled substance analog or from which a controlled

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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;

(5) scales or balances used, intended for use or designed for use in weighing or measuring controlled substances or controlled substance analogs;

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

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

(8) blenders, bowls, containers, spoons and mixing devices used, intended for use or designed for use in compounding controlled substances or controlled substance

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analogs;

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

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

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

(12) objects used, intended for use or designed for use in ingesting, inhaling or otherwise introducing marijuana, cocaine, hashish or hashish oil into the human body, such as:

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

(b) water pipes;

(c) carburetion tubes and devices;

(d) smoking and carburetion masks;

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

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(f) miniature cocaine spoons and cocaine vials;

(g) chamber pipes;

(h) carburetor pipes;

(i) electric pipes;

(j) air-driven pipes;

(k) chilams;

(l) bongs; or

(m) ice pipes or chillers; and

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

(a) statements by the owner or by anyone in control of the object concerning its use;

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

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

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

(e) instructions, written or oral,

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provided with the object concerning its use;

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

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

(h) expert testimony concerning its use;

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

(1) phenethylamines;

(2) N-substituted piperidines;

(3) morphinans;

(4) ecgonines;

(5) quinazolinones;

(6) substituted indoles; and

(7) arylcycloalkylamines.

Specifically excluded from the definition of "controlled substance analog" are those substances that are generally recognized as safe and effective within the meaning of the

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Federal Food, Drug and Cosmetic Act or have been manufactured, distributed or possessed in conformance with the provisions of an approved new drug application or an exemption for investigational use within the meaning of Section 505 of the Federal Food, Drug and Cosmetic Act;

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

Y. "drug-free school zone" means a public school or property that is used for public school purposes and the area within one thousand feet of the school property line, but it does not mean any post-secondary school."

Section 3. Section 61-3-3 NMSA 1978 (being Laws 1991, Chapter 190, Section 2, as amended) is amended to read:

"61-3-3. DEFINITIONS.--As used in the Nursing Practice Act:

A. "advanced practice" means the practice of professional registered nursing by a registered nurse who has been prepared through additional formal education as provided in Sections 61-3-23.2 through 61-3-23.4 NMSA 1978 to function beyond the scope of practice of professional registered nursing, including certified nurse practitioners, certified registered nurse anesthetists and clinical nurse specialists;

B. "board" means the board of nursing;

C. "certified nurse practitioner" means a

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registered nurse who is licensed by the board for advanced practice as a certified nurse practitioner and whose name and pertinent information are entered on the list of certified nurse practitioners maintained by the board;

D. "certified registered nurse anesthetist" means a registered nurse who is licensed by the board for advanced practice as a certified registered nurse anesthetist and whose name and pertinent information are entered on the list of certified registered nurse anesthetists maintained by the board;

E. "clinical nurse specialist" means a registered nurse who is licensed by the board for advanced practice as a clinical nurse specialist and whose name and pertinent information are entered on the list of clinical nurse specialists maintained by the board;

F. "collaboration" means the cooperative working relationship with another health care provider in the provision of patient care, and such collaborative practice includes the discussion of patient diagnosis and cooperation in the management and delivery of health care;

G. "emergency procedures" means airway and vascular access procedures;

H. "licensed practical nurse" means a nurse who practices licensed practical nursing and whose name and pertinent information are entered in the register of licensed

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practical nurses maintained by the board;

I. "licensed practical nursing" means the practice of a directed scope of nursing requiring basic knowledge of the biological, physical, social and behavioral sciences and nursing procedures, which practice is at the direction of a registered nurse, physician, prescribing psychologist or dentist licensed to practice in this state. This practice includes but is not limited to:

(1) contributing to the assessment of the health status of individuals, families and communities;

(2) participating in the development and modification of the plan of care;

(3) implementing appropriate aspects of the plan of care commensurate with education and verified competence;

(4) collaborating with other health care professionals in the management of health care; and

(5) participating in the evaluation of responses to interventions;

J. "nursing diagnosis" means a clinical judgment about individual, family or community responses to actual or potential health problems or life processes, which judgment provides a basis for the selection of nursing interventions to achieve outcomes for which the person making the judgment is accountable;

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K. "practice of nursing" means assisting individuals, families or communities in maintaining or attaining optimal health, assessing and implementing a plan of care to accomplish defined goals and evaluating responses to care and treatment. This practice is based on specialized knowledge, judgment and nursing skills acquired through educational preparation in nursing and in the biological, physical, social and behavioral sciences and includes but is not limited to:

(1) initiating and maintaining comfort measures;

(2) promoting and supporting optimal human functions and responses;

(3) establishing an environment conducive to well-being or to the support of a dignified death;

(4) collaborating on the health care regimen;

(5) administering medications and performing treatments prescribed by a person authorized in this state or in any other state in the United States to prescribe them;

(6) recording and reporting nursing observations, assessments, interventions and responses to health care;

(7) providing counseling and health teaching;

(8) delegating and supervising nursing interventions that may be performed safely by others and are

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not in conflict with the Nursing Practice Act; and

(9) maintaining accountability for safe and effective nursing care;

L. "professional registered nursing" means the practice of the full scope of nursing requiring substantial knowledge of the biological, physical, social and behavioral sciences and of nursing theory and may include advanced practice pursuant to the Nursing Practice Act. This practice includes but is not limited to:

(1) assessing the health status of individuals, families and communities;

(2) establishing a nursing diagnosis;

(3) establishing goals to meet identified health care needs;

(4) developing a plan of care;

(5) determining nursing intervention to implement the plan of care;

(6) implementing the plan of care commensurate with education and verified competence;

(7) evaluating responses to interventions;

(8) teaching based on the theory and practice of nursing;

(9) managing and supervising the practice of nursing;

(10) collaborating with other health care

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professionals in the management of health care; and

(11) conducting nursing research;

M. "registered nurse" means a nurse who practices professional registered nursing and whose name and pertinent information are entered in the register of licensed registered nurses maintained by the board; and

N. "scope of practice" means the parameters within which nurses practice based upon education, experience, licensure, certification and expertise."

Section 4. Section 61-9-1 NMSA 1978 (being Laws 1963, Chapter 92, Section 1) is amended to read:

"61-9-1. SHORT TITLE.--[~~This act~~] Chapter 61, Article 9 NMSA 1978 may be cited as the "Professional Psychologist Act"."

Section 5. Section 61-9-3 NMSA 1978 (being Laws 1963, Chapter 92, Section 3, as amended) is amended to read:

"61-9-3. DEFINITIONS.--As used in the Professional Psychologist Act:

A. "board" means the New Mexico state board of psychologist examiners;

B. "guidelines or protocol" means a written agreement between a prescribing psychologist and a physician authorized by law in New Mexico to prescribe controlled substances;

C. "prescribing psychologist" means a doctoral

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level licensed psychologist who has been certified by the board pursuant to the Professional Psychologist Act to prescribe psychotropic medications in accordance with guidelines or protocol;

D. "psychotropic medication" means a controlled substance or dangerous drug that may not be dispensed or administered without a prescription and whose primary indication for use has been approved by the federal food and drug administration for the treatment of mental disorders and is listed as a psychotherapeutic agent in drug facts and comparisons or in the American hospital formulary service;

[~~B.~~] E. "person" includes an individual, firm, partnership, association or corporation;

[~~C.~~] F. "psychologist" means [~~any~~] a person who engages in the practice of psychology or holds himself out to the public by any title or description of services representing himself as a psychologist, which incorporates the words "psychological", "psychologist", "psychology", or when a person describes himself as above and, under such title or description, offers to render or renders services involving the application of principles, methods and procedures of the science and profession of psychology to persons for compensation or other personal gain;

[~~D.~~] G. "practice of psychology" means the observation, description, evaluation, interpretation and

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modification of human behavior by the application of psychological principles, methods and procedures for the purpose of preventing or eliminating symptomatic, maladaptive or undesired behavior and of enhancing interpersonal relationships, work and life adjustment, personal effectiveness, behavioral health and mental health, and further means the rendering of such psychological services to individuals, families or groups regardless of whether payment is received for services rendered. The "practice of psychology" includes psychological testing or neuropsychological testing and the evaluation or assessment of personal characteristics such as intelligence, personality, abilities, interests, aptitudes and neuropsychological functioning; counseling, psychoanalysis, psychotherapy, hypnosis, biofeedback, behavior analysis and therapy; diagnosis and treatment of any mental and emotional disorder or disability, alcoholism and substance abuse, disorders of habit or conduct and the psychological aspects of physical illness, accident, injury and disability; and psychoeducational evaluation, therapy, remediation and consultation; and

[~~E.~~] H. "school" or "college" means [~~any~~] a university or other institution of higher education that is regionally accredited and that offers a full-time graduate course of study in psychology as defined by rule of the board

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or that is approved by the American psychological association."

Section 6. A new section of the Professional Psychologist Act, Section 61-9-12.1 NMSA 1978, is enacted to read:

"61-9-12.1. [NEW MATERIAL] PRESCRIBING PSYCHOLOGIST--CERTIFICATION--GUIDELINES OR PROTOCOL.--

A. The board may certify a psychologist as a prescribing psychologist upon determining that the psychologist:

(1) holds a current license to practice psychology in New Mexico;

(2) has completed a doctoral program in psychology from an accredited institution of higher education or professional school, or, if the program was not accredited at the time of the psychologist's graduation, that the program meets professional standards determined acceptable by the board;

(3) meets the requirements for additional training that have been adopted by the board and by the New Mexico board of medical examiners, which requirements shall be at least equivalent to those required for a pharmacist clinician in New Mexico;

(4) has passed a national examination

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approved by the board and by the New Mexico board of medical examiners that demonstrates professional competency for prescriptive authority; and

(5) has medical malpractice insurance equal to at least one million dollars ($1,000,000) per occurrence. Evidence of the insurance shall be submitted to the board annually.

B. A prescribing psychologist planning to exercise prescriptive authority in his practice shall have on file at his place of practice the guidelines or protocol. The guidelines or protocol shall authorize the prescribing psychologist to prescribe psychotropic medication and shall be established and approved by a licensed physician in accordance with rules adopted by the board and by the New Mexico board of medical examiners. A copy of the guidelines or protocol shall be on file with the board and with the New Mexico board of medical examiners. The physician who is a party to the guidelines or protocol shall be in active practice in New Mexico.

C. The guidelines or protocol required by Subsection B of this section shall include:

(1) a statement identifying the licensed physician and the prescribing psychologist who are the parties to the guidelines or protocol;

(2) a statement of the types of prescriptive

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authority decisions that the prescribing psychologist is authorized to make, which may include:

(a) a statement of the types of mental disorders or psychotropic drug categories involved and the type of prescriptive authority authorized in each case; and

(b) a general statement of the procedures, decision criteria or plan the prescribing psychologist is to follow when exercising prescriptive authority;

(3) a statement of the activities the prescribing psychologist is to follow in the course of exercising prescriptive authority, including documentation of decisions made and a plan for communication or feedback to the authorizing physician concerning specific decisions made. Documentation may occur on the prescriptive record, patient profile, patient medical chart or in a separate log book; and

(4) a statement that describes appropriate mechanisms for reporting to the physician monitoring activities and results.

D. The guidelines or protocol shall be reviewed by the board and by the New Mexico board of medical examiners and shall be revised every two years if necessary.

E. The New Mexico board of medical examiners shall adopt rules relating to the acts and duties of licensed physicians pursuant to this section.

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F. The board shall, by rule, provide for continuing education requirements of not fewer than twenty-five hours of classroom work per year for prescribing psychologists.

G. The board shall adopt rules establishing the grounds for denial, suspension or revocation of a certification as a prescribing psychologist, including a provision for suspension or revocation of a license to practice psychology upon suspension or revocation of a certification. Actions of denial, suspension or revocation of a certification shall be in accordance with the Uniform Licensing Act.

H. The board shall contract with an independent peer review organization to perform in-office chart review of prescribing psychologists, and shall report its findings to the board, the New Mexico board of medical examiners, the legislature and the governor no later than November 1, 2006."

Section 7. A new section of the Professional Psychologist Act, Section 61-9-12.2 NMSA 1978, is enacted to read:

"61-9-12.2. [NEW MATERIAL] PRESCRIBING PRACTICES.--

A. A prescription written by a prescribing psychologist shall:

(1) comply with applicable state and federal

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laws;

(2) be identified as issued by the psychologist as "prescribing psychologist"; and

(3) include the prescribing psychologist's board-assigned identification number.

B. A prescribing psychologist shall not delegate prescriptive authority to any other person. Records of all prescriptions shall be maintained in patient records.

C. When authorized to prescribe controlled substances, a prescribing psychologist shall file with the board in a timely manner all individual federal drug enforcement agency registrations and numbers. The board shall maintain current records on every prescribing psychologist, including federal registrations and numbers.

D. The board shall provide to the board of pharmacy an annual list of prescribing psychologists that contains the information agreed upon. The board shall promptly notify the board of pharmacy of prescribing psychologists who are added or deleted from the list."

Section 8. Section 61-9-17 NMSA 1978 (being Laws 1963, Chapter 92, Section 16, as amended) is amended to read:

"61-9-17. DRUGS--MEDICINES.--[~~Nothing in the Professional Psychologist Act shall be construed as permitting~~] Except as provided in Sections 61-9-12.1 and 61-9-12.2 NMSA 1978, psychologists or psychologist associates

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licensed under the Professional Psychologist Act [~~to~~] shall not administer or prescribe drugs or medicine or in any manner engage in the practice of medicine as defined by the laws of this state."

Section 9. EFFECTIVE DATE.--The effective date of the provisions of this act is July 1, 2002.

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