

LEGISLATURE OF NEBRASKA  
ONE HUNDREDTH LEGISLATURE  
SECOND SESSION

**LEGISLATIVE BILL 885**

Introduced by Johnson, 37.

Read first time January 11, 2008

Committee: Health and Human Services

A BILL

1 FOR AN ACT relating to pharmacy; to amend sections 38-2840 and  
2 71-1,142, Revised Statutes Supplement, 2007; to redefine  
3 prescription; to harmonize provisions; to provide  
4 operative dates; to repeal the original sections; and  
5 to outright repeal section 71-1,142, Revised Statutes  
6 Supplement, 2007, as amended by section 2 of this  
7 legislative bill.

8 Be it enacted by the people of the State of Nebraska,

1           Section 1. Section 38-2840, Revised Statutes Supplement,  
2 2007, is amended to read:

3           38-2840 Prescription means an order for a drug or device  
4 issued by a practitioner for a specific patient, for emergency  
5 use, ~~or~~ for use in immunizations, or for a single-dose sample for  
6 research purposes. Prescription does not include a chart order.

7           Sec. 2. Section 71-1,142, Revised Statutes Supplement,  
8 2007, is amended to read:

9           71-1,142 For purposes of sections 71-1,142 to 71-1,151  
10 and elsewhere in the Uniform Licensing Law, unless the context  
11 otherwise requires:

12           (1) Practice of pharmacy means (a) the interpretation,  
13 evaluation, and implementation of a medical order, (b) the  
14 dispensing of drugs and devices, (c) drug product selection,  
15 (d) the administration of drugs or devices, (e) drug utilization  
16 review, (f) patient counseling, (g) the provision of pharmaceutical  
17 care, and (h) the responsibility for compounding and labeling of  
18 dispensed or repackaged drugs and devices, proper and safe storage  
19 of drugs and devices, and maintenance of proper records. The active  
20 practice of pharmacy means the performance of the functions set  
21 out in this subdivision by a pharmacist as his or her principal or  
22 ordinary occupation;

23           (2) Administer means to directly apply a drug or device  
24 by injection, inhalation, ingestion, or other means to the body of  
25 a patient or research subject;

1           (3) Administration means the act of (a) administering,  
2           (b) keeping a record of such activity, and (c) observing,  
3           monitoring, reporting, and otherwise taking appropriate action  
4           regarding desired effect, side effect, interaction, and  
5           contraindication associated with administering the drug or device;

6           (4) Board means the Board of Pharmacy;

7           (5) Caregiver means any person acting as an agent on  
8           behalf of a patient or any person aiding and assisting a patient;

9           (6) Chart order means an order for a drug or device  
10          issued by a practitioner for a patient who is in the hospital  
11          where the chart is stored or for a patient receiving detoxification  
12          treatment or maintenance treatment pursuant to section 28-412.  
13          Chart order does not include a prescription;

14          (7) Compounding means the preparation of components into  
15          a drug product (a) as the result of a practitioner's medical order  
16          or initiative occurring in the course of practice based upon the  
17          relationship between the practitioner, patient, and pharmacist or  
18          (b) for the purpose of, or as an incident to, research, teaching,  
19          or chemical analysis and not for sale or dispensing. Compounding  
20          includes the preparation of drugs or devices in anticipation of  
21          receiving medical orders based upon routine, regularly observed  
22          prescribing patterns;

23          (8) Delegated dispensing means the practice of pharmacy  
24          by which one or more pharmacists have jointly agreed, on a  
25          voluntary basis, to work in conjunction with one or more persons

1 pursuant to sections 71-1,147.42 to 71-1,147.64 under a protocol  
2 which provides that such person may perform certain dispensing  
3 functions authorized by the pharmacist or pharmacists under certain  
4 specified conditions and limitations;

5 (9) Deliver or delivery means to actually,  
6 constructively, or attempt to transfer a drug or device from one  
7 person to another, whether or not for consideration;

8 (10) Department means the Division of Public Health of  
9 the Department of Health and Human Services;

10 (11) Device means an instrument, apparatus, implement,  
11 machine, contrivance, implant, in vitro reagent, or other similar  
12 or related article, including any component, part, or accessory,  
13 which is prescribed by a practitioner and dispensed by a pharmacist  
14 or other person authorized by law to do so;

15 (12) Dialysis drug or device distributor means a  
16 manufacturer or wholesaler who provides dialysis drugs, solutions,  
17 supplies, or devices, to persons with chronic kidney failure for  
18 self-administration at the person's home or specified address,  
19 pursuant to a prescription;

20 (13) Dialysis drug or device distributor worker means a  
21 person working for a dialysis drug or device distributor with a  
22 delegated dispensing permit who has completed the approved training  
23 and has demonstrated proficiency to perform the task or tasks of  
24 assembling, labeling, or delivering drugs or devices pursuant to a  
25 prescription;

1           (14) Dispense or dispensing means interpreting,  
2 evaluating, and implementing a medical order, including preparing  
3 and delivering a drug or device to a patient or caregiver  
4 in a suitable container appropriately labeled for subsequent  
5 administration to, or use by, a patient. Dispensing includes (a)  
6 dispensing incident to practice, (b) dispensing pursuant to a  
7 delegated dispensing permit, (c) dispensing pursuant to a medical  
8 order, and (d) any transfer of a prescription drug or device to a  
9 patient or caregiver other than by administering;

10           (15) Distribute means to deliver a drug or device, other  
11 than by administering or dispensing;

12           (16) Facility means a health care facility as defined in  
13 section 71-413;

14           (17) Hospital has the same meaning as in section 71-419;

15           (18) Person means an individual, corporation,  
16 partnership, limited liability company, association, or other legal  
17 entity;

18           (19) Labeling means the process of preparing and affixing  
19 a label to any drug container or device container, exclusive  
20 of the labeling by a manufacturer, packer, or distributor of  
21 a nonprescription drug or commercially packaged legend drug or  
22 device. Any such label shall include all information required by  
23 federal and state law or regulation;

24           (20) Medical order means a prescription, a chart order,  
25 or an order for pharmaceutical care issued by a practitioner;

1           (21) Pharmaceutical care means the provision of drug  
2 therapy for the purpose of achieving therapeutic outcomes that  
3 improve a patient's quality of life. Such outcomes include (a) the  
4 cure of disease, (b) the elimination or reduction of a patient's  
5 symptomatology, (c) the arrest or slowing of a disease process, or  
6 (d) the prevention of a disease or symptomatology. Pharmaceutical  
7 care includes the process through which the pharmacist works in  
8 concert with the patient and his or her caregiver, physician, or  
9 other professionals in designing, implementing, and monitoring a  
10 therapeutic plan that will produce specific therapeutic outcomes  
11 for the patient;

12           (22) Pharmacist means any person who is licensed by the  
13 State of Nebraska to practice pharmacy;

14           (23) Pharmacy has the same meaning as in section 71-425;

15           (24) Drugs, medicines, and medicinal substances means (a)  
16 articles recognized in the official United States Pharmacopoeia,  
17 the Homeopathic Pharmacopoeia of the United States, the official  
18 National Formulary, or any supplement to any of them, (b) articles  
19 intended for use in the diagnosis, cure, mitigation, treatment, or  
20 prevention of diseases in humans or animals, (c) articles, except  
21 food, intended to affect the structure or any function of the  
22 body of a human or an animal, (d) articles intended for use as a  
23 component of any articles specified in subdivision (a), (b), or (c)  
24 of this subdivision, except any device or its components, parts, or  
25 accessories, and (e) prescription drugs or devices as defined in

1 subdivision (31) of this section;

2 (25) Patient counseling means the verbal communication  
3 by a pharmacist, pharmacist intern, or practitioner, in a manner  
4 reflecting dignity and the right of the patient to a reasonable  
5 degree of privacy, of information to the patient or caregiver in  
6 order to improve therapeutic outcomes by maximizing proper use of  
7 prescription drugs and devices and also includes the duties set out  
8 in section 71-1,147.35;

9 (26) Pharmacist in charge means a pharmacist who is  
10 designated on a pharmacy license or designated by a hospital as  
11 being responsible for the practice of pharmacy in the pharmacy  
12 for which a pharmacy license is issued and who works within the  
13 physical confines of such pharmacy for a majority of the hours  
14 per week that the pharmacy is open for business averaged over a  
15 twelve-month period or thirty hours per week, whichever is less;

16 (27) Pharmacist intern means a person who meets the  
17 requirements of section 71-1,144;

18 (28) Pharmacy technician means an individual registered  
19 under sections 71-1,147.65 to 71-1,147.72;

20 (29) Practitioner means a certified registered nurse  
21 anesthetist, a certified nurse midwife, a dentist, an optometrist,  
22 a nurse practitioner, a physician assistant, a physician, a  
23 podiatrist, or a veterinarian;

24 (30) Prescribe means to issue a medical order;

25 (31) Prescription drug or device or legend drug or

1 device means (a) a drug or device which is required under  
2 federal law to be labeled with one of the following statements  
3 prior to being dispensed or delivered: (i) Caution: Federal law  
4 prohibits dispensing without prescription; (ii) Caution: Federal  
5 law restricts this drug to use by or on the order of a licensed  
6 veterinarian; or (iii) "Rx Only" or (b) a drug or device which is  
7 required by any applicable federal or state law to be dispensed  
8 pursuant only to a prescription or chart order or which is  
9 restricted to use by practitioners only;

10 (32) Prescription means an order for a drug or device  
11 issued by a practitioner for a specific patient, for emergency  
12 use, ~~or~~ for use in immunizations, or for a single-dose sample for  
13 research purposes. Prescription does not include a chart order;

14 (33) Nonprescription drugs means nonnarcotic medicines or  
15 drugs which may be sold without a medical order and which are  
16 prepackaged for use by the consumer and labeled in accordance with  
17 the requirements of the laws and regulations of this state and the  
18 federal government;

19 (34) Public health clinic worker means a person in a  
20 public health clinic with a delegated dispensing permit who has  
21 completed the approved training and has demonstrated proficiency  
22 to perform the task of dispensing authorized refills of oral  
23 contraceptives pursuant to a written prescription;

24 (35) Public health clinic means the department, any  
25 county, city-county, or multicounty health department, or any

1 private not-for-profit family planning clinic licensed as a health  
2 clinic as defined in section 71-416;

3 (36) Signature means the name, word, or mark of a person  
4 written in his or her own hand with the intent to authenticate a  
5 writing or other form of communication or a digital signature which  
6 complies with section 86-611 or an electronic signature;

7 (37) Supervision means the immediate personal guidance  
8 and direction by the licensed pharmacist on duty in the facility of  
9 the performance by a pharmacy technician of authorized activities  
10 or functions subject to verification by such pharmacist, except  
11 that when a pharmacy technician performs authorized activities or  
12 functions to assist a pharmacist on duty in the facility when the  
13 prescribed drugs or devices will be administered by a licensed  
14 staff member or consultant or by a licensed physician assistant to  
15 persons who are patients or residents of a facility, the activities  
16 or functions of such pharmacy technician shall only be subject to  
17 verification by a pharmacist on duty in the facility;

18 (38) Verification means the confirmation by a supervising  
19 pharmacist of the accuracy and completeness of the acts, tasks,  
20 or functions undertaken by a pharmacy technician to assist the  
21 pharmacist in the practice of pharmacy;

22 (39) Written control procedures and guidelines means  
23 the document prepared and signed by the pharmacist in charge  
24 and approved by the board which specifies the manner in which  
25 basic levels of competency of pharmacy technicians employed by

1 the pharmacy are determined, the manner in which supervision is  
2 provided, the manner in which the functions of pharmacy technicians  
3 are verified, the maximum ratio of pharmacy technicians to one  
4 pharmacist used in the pharmacy, and guidelines governing the use  
5 of pharmacy technicians and the functions which they may perform;

6 (40) Medical gas distributor means a person who dispenses  
7 medical gases to a patient or ultimate user but does not include a  
8 person who manufactures medical gases or a person who distributes,  
9 transfers, delivers, dispenses, or sells medical gases to a person  
10 other than a patient or ultimate user;

11 (41) Facsimile means a copy generated by a system that  
12 encodes a document or photograph into electrical signals, transmits  
13 those signals over telecommunications lines, and reconstructs the  
14 signals to create an exact duplicate of the original document at  
15 the receiving end;

16 (42) Electronic signature has the same definition found  
17 in section 86-621; and

18 (43) Electronic transmission means transmission of  
19 information in electronic form. Electronic transmission may  
20 include computer-to-computer transmission or computer-to-facsimile  
21 transmission.

22 Sec. 3. Sections 1, 5, and 6 of this act become operative  
23 on December 1, 2008. Sections 2, 3, and 4 of this act become  
24 operative on their effective date.

25 Sec. 4. Original section 71-1,142, Revised Statutes

1 Supplement, 2007, is repealed.

2           Sec. 5. Original section 38-2840, Revised Statutes  
3 Supplement, 2007, is repealed.

4           Sec. 6. The following section is outright repealed:  
5 Section 71-1,142, Revised Statutes Supplement, 2007, as amended by  
6 section 2 of this legislative bill.