

1.1 moves to amend H.F. No. 3138, the delete everything amendment
1.2 (A18-0776), as follows:

1.3 Page 90, after line 10, insert:

1.4 "Section 1. Minnesota Statutes 2016, section 151.01, subdivision 23, is amended to read:

1.5 Subd. 23. **Practitioner.** "Practitioner" means a licensed doctor of medicine, licensed
1.6 doctor of osteopathic medicine duly licensed to practice medicine, licensed doctor of
1.7 dentistry, licensed doctor of optometry, licensed podiatrist, licensed veterinarian, or licensed
1.8 advanced practice registered nurse. For purposes of sections 151.15, subdivision 4; 151.252,
1.9 subdivision 3; 151.37, subdivision 2, paragraphs (b), (e), and (f); and 151.461, "practitioner"
1.10 also means a physician assistant authorized to prescribe, dispense, and administer under
1.11 chapter 147A. For purposes of sections 151.15, subdivision 4; 151.252, subdivision 3;
1.12 151.37, subdivision 2, paragraph (b); and 151.461, "practitioner" also means a dental therapist
1.13 authorized to dispense and administer under chapter 150A. For purposes of sections 151.216;
1.14 151.252, subdivision 3; and 151.461, practitioner also means a pharmacist authorized to
1.15 prescribe under section 151.216: tobacco cessation medications approved by the Food and
1.16 Drug Administration (FDA) but excluding bupropion; opiate antagonists; and travel
1.17 medications.

1.18 Sec. 2. Minnesota Statutes 2016, section 151.01, subdivision 27, is amended to read:

1.19 Subd. 27. **Practice of pharmacy.** "Practice of pharmacy" means:

1.20 (1) interpretation and evaluation of prescription drug orders;

1.21 (2) compounding, labeling, and dispensing drugs and devices (except labeling by a
1.22 manufacturer or packager of nonprescription drugs or commercially packaged legend drugs
1.23 and devices);

2.1 (3) participation in clinical interpretations and monitoring of drug therapy for assurance
2.2 of safe and effective use of drugs, including the performance of laboratory tests that are
2.3 waived under the federal Clinical Laboratory Improvement Act of 1988, United States Code,
2.4 title 42, section 263a et seq., provided that a pharmacist may interpret the results of laboratory
2.5 tests but may modify drug therapy only pursuant to a protocol or collaborative practice
2.6 agreement;

2.7 (4) participation in drug and therapeutic device selection; drug administration for first
2.8 dosage and medical emergencies; drug regimen reviews; and drug or drug-related research;

2.9 (5) participation in administration of influenza vaccines to all eligible individuals six
2.10 years of age and older and all other vaccines to patients 13 years of age and older by written
2.11 protocol with a physician licensed under chapter 147, a physician assistant authorized to
2.12 prescribe drugs under chapter 147A, or an advanced practice registered nurse authorized to
2.13 prescribe drugs under section 148.235, provided that:

2.14 (i) the protocol includes, at a minimum:

2.15 (A) the name, dose, and route of each vaccine that may be given;

2.16 (B) the patient population for whom the vaccine may be given;

2.17 (C) contraindications and precautions to the vaccine;

2.18 (D) the procedure for handling an adverse reaction;

2.19 (E) the name, signature, and address of the physician, physician assistant, or advanced
2.20 practice registered nurse;

2.21 (F) a telephone number at which the physician, physician assistant, or advanced practice
2.22 registered nurse can be contacted; and

2.23 (G) the date and time period for which the protocol is valid;

2.24 (ii) the pharmacist has successfully completed a program approved by the Accreditation
2.25 Council for Pharmacy Education specifically for the administration of immunizations or a
2.26 program approved by the board;

2.27 (iii) the pharmacist utilizes the Minnesota Immunization Information Connection to
2.28 assess the immunization status of individuals prior to the administration of vaccines, except
2.29 when administering influenza vaccines to individuals age nine and older;

2.30 (iv) the pharmacist reports the administration of the immunization to the Minnesota
2.31 Immunization Information Connection; and

3.1 (v) the pharmacist complies with guidelines for vaccines and immunizations established
3.2 by the federal Advisory Committee on Immunization Practices, except that a pharmacist
3.3 does not need to comply with those portions of the guidelines that establish immunization
3.4 schedules when administering a vaccine pursuant to a valid, patient-specific order issued
3.5 by a physician licensed under chapter 147, a physician assistant authorized to prescribe
3.6 drugs under chapter 147A, or an advanced practice nurse authorized to prescribe drugs
3.7 under section 148.235, provided that the order is consistent with the United States Food
3.8 and Drug Administration approved labeling of the vaccine;

3.9 (6) participation in the initiation, management, modification, and discontinuation of
3.10 drug therapy according to a written protocol or collaborative practice agreement between:
3.11 (i) one or more pharmacists and one or more dentists, optometrists, physicians, podiatrists,
3.12 or veterinarians; or (ii) one or more pharmacists and one or more physician assistants
3.13 authorized to prescribe, dispense, and administer under chapter 147A, or advanced practice
3.14 nurses authorized to prescribe, dispense, and administer under section 148.235. Any changes
3.15 in drug therapy made pursuant to a protocol or collaborative practice agreement must be
3.16 documented by the pharmacist in the patient's medical record or reported by the pharmacist
3.17 to a practitioner responsible for the patient's care;

3.18 (7) participation in the storage of drugs and the maintenance of records;

3.19 (8) patient counseling on therapeutic values, content, hazards, and uses of drugs and
3.20 devices;

3.21 (9) offering or performing those acts, services, operations, or transactions necessary in
3.22 the conduct, operation, management, and control of a pharmacy; ~~and~~

3.23 (10) participation in the initiation, management, modification, and discontinuation of
3.24 therapy with opiate antagonists, as defined in section 604A.04, subdivision 1, pursuant to:

3.25 (i) a written protocol as allowed under clause (6); or

3.26 (ii) a written protocol with a community health board medical consultant or a practitioner
3.27 designated by the commissioner of health, as allowed under section 151.37, subdivision 13;
3.28 and

3.29 (11) prescribing under section 151.216: tobacco cessation medications approved by the
3.30 FDA but excluding bupropion; opiate antagonists; and travel medications.

4.1 Sec. 3. Minnesota Statutes 2016, section 151.01, is amended by adding a subdivision to
4.2 read:

4.3 Subd. 42. **Travel medication.** "Travel medication" means a medication that is: (1)
4.4 recommended by the federal Centers for Disease Control and Prevention for individuals
4.5 traveling outside of the United States; (2) included on the list of travel medications approved
4.6 by the Board of Pharmacy under section 151.216, subdivision 2, paragraph (a); and (3)
4.7 prescribed to an individual for the specific purpose of traveling outside the United States."

4.8 Page 90, after line 21, insert:

4.9 "Sec. 5. **[151.216] PHARMACIST PRESCRIBING.**

4.10 Subdivision 1. **Tobacco cessation medications, opiate antagonists, and travel**
4.11 **medications.** (a) A pharmacist is authorized to prescribe: tobacco cessation medications
4.12 approved by the Food and Drug Administration (FDA) but excluding bupropion; opiate
4.13 antagonists for the purpose of treating opioid overdose; and travel medications.

4.14 (b) A pharmacist who prescribes products or medications under this subdivision must
4.15 provide the patient with a written record of the product or medication prescribed by the
4.16 pharmacist.

4.17 (c) A pharmacist who prescribes products or medications under this subdivision is
4.18 prohibited from delegating the prescribing of the product or medication to any other person,
4.19 but may allow a pharmacist intern who is registered pursuant to section 151.101 to prepare
4.20 a prescription for the product or medication, provided that the prescription shall not be
4.21 processed or dispensed until it is reviewed, approved, and signed by the pharmacist.

4.22 Subd. 2. Protocols. (a) The board shall develop standardized protocols that pharmacists
4.23 must follow in order to prescribe products and medications under this subdivision. The
4.24 protocol for travel medications must include a list of travel medications approved by the
4.25 board as appropriate to be prescribed by pharmacists for individuals traveling outside of
4.26 the United States. In developing the protocols, the board shall consult with the Minnesota
4.27 Board of Medical Practice, the Minnesota Board of Nursing, the commissioner of health,
4.28 professional pharmacy associations, and professional associations of physicians, physician
4.29 assistants, and advanced practice registered nurses.

4.30 (b) Nothing in this subdivision prohibits a pharmacist from participating in the initiation,
4.31 management, modification, and discontinuation of therapy through a protocol as allowed
4.32 in this section or section 151.37, subdivisions 2 and 13.

5.1 Sec. 6. Minnesota Statutes 2016, section 151.253, is amended by adding a subdivision to
5.2 read:

5.3 Subd. 4. **Emergency veterinary compounding.** A pharmacist working in a pharmacy
5.4 licensed by the board in the veterinary pharmacy license category may compound and
5.5 provide a drug product to a veterinarian without first receiving a patient-specific prescription
5.6 only when:

5.7 (1) the compounded drug product is needed to treat an animal in an urgent or emergency
5.8 situation. For the purpose of this clause, "urgent or emergency situation" means a situation
5.9 where the health of an animal is threatened, or where suffering or death of an animal is
5.10 likely to result from failure to immediately treat;

5.11 (2) timely access to a compounding pharmacy is not available, as determined by the
5.12 prescribing veterinarian;

5.13 (3) there is no commercially manufactured drug approved by the United States Food
5.14 and Drug Administration that is suitable for treating the animal, or there is a documented
5.15 shortage of a commercially manufactured drug;

5.16 (4) the compounded drug is to be administered by a veterinarian or a bona fide employee
5.17 of the veterinarian or dispensed to a client of a veterinarian in an amount not to exceed what
5.18 is necessary to treat an animal for a period of ten days;

5.19 (5) the pharmacy has selected the sterile or nonsterile compounding license category,
5.20 in addition to the veterinary pharmacy licensing category; and

5.21 (6) the pharmacy is appropriately registered by the United States Drug Enforcement
5.22 Administration when providing compounded products that contain controlled substances."

5.23 Page 99, after line 13, insert:

5.24 "Sec. 10. Minnesota Statutes 2016, section 256B.0625, subdivision 13h, is amended to
5.25 read:

5.26 Subd. 13h. **Medication therapy management services.** (a) Medical assistance covers
5.27 medication therapy management services for a recipient taking prescriptions to treat or
5.28 prevent one or more chronic medical conditions. For purposes of this subdivision,
5.29 "medication therapy management" means the provision of the following pharmaceutical
5.30 care services by a licensed pharmacist to optimize the therapeutic outcomes of the patient's
5.31 medications:

5.32 (1) performing or obtaining necessary assessments of the patient's health status;

- 6.1 (2) formulating a medication treatment plan;
- 6.2 (3) monitoring and evaluating the patient's response to therapy, including safety and
6.3 effectiveness;
- 6.4 (4) performing a comprehensive medication review to identify, resolve, and prevent
6.5 medication-related problems, including adverse drug events;
- 6.6 (5) documenting the care delivered and communicating essential information to the
6.7 patient's other primary care providers;
- 6.8 (6) providing verbal education and training designed to enhance patient understanding
6.9 and appropriate use of the patient's medications;
- 6.10 (7) providing information, support services, and resources designed to enhance patient
6.11 adherence with the patient's therapeutic regimens; ~~and~~
- 6.12 (8) coordinating and integrating medication therapy management services within the
6.13 broader health care management services being provided to the patient; and
- 6.14 (9) prescribing products or medications as permitted under section 151.216, in accordance
6.15 with standardized protocols developed by the Board of Pharmacy.

6.16 Nothing in this subdivision shall be construed to expand or modify the scope of practice of
6.17 the pharmacist as defined in section 151.01, subdivision 27.

6.18 (b) To be eligible for reimbursement for services under this subdivision, a pharmacist
6.19 must meet the following requirements:

6.20 (1) have a valid license issued by the Board of Pharmacy of the state in which the
6.21 medication therapy management service is being performed;

6.22 (2) have graduated from an accredited college of pharmacy on or after May 1996, or
6.23 completed a structured and comprehensive education program approved by the Board of
6.24 Pharmacy and the American Council of Pharmaceutical Education for the provision and
6.25 documentation of pharmaceutical care management services that has both clinical and
6.26 didactic elements;

6.27 (3) be practicing in an ambulatory care setting as part of a multidisciplinary team or
6.28 have developed a structured patient care process that is offered in a private or semiprivate
6.29 patient care area that is separate from the commercial business that also occurs in the setting,
6.30 or in home settings, including long-term care settings, group homes, and facilities providing
6.31 assisted living services, but excluding skilled nursing facilities; and

6.32 (4) make use of an electronic patient record system that meets state standards.

7.1 (c) For purposes of reimbursement for medication therapy management services, the
7.2 commissioner may enroll individual pharmacists as medical assistance providers. The
7.3 commissioner may also establish contact requirements between the pharmacist and recipient,
7.4 including limiting the number of reimbursable consultations per recipient.

7.5 (d) If there are no pharmacists who meet the requirements of paragraph (b) practicing
7.6 within a reasonable geographic distance of the patient, a pharmacist who meets the
7.7 requirements may provide the services via two-way interactive video. Reimbursement shall
7.8 be at the same rates and under the same conditions that would otherwise apply to the services
7.9 provided. To qualify for reimbursement under this paragraph, the pharmacist providing the
7.10 services must meet the requirements of paragraph (b), and must be located within an
7.11 ambulatory care setting that meets the requirements of paragraph (b), clause (3). The patient
7.12 must also be located within an ambulatory care setting that meets the requirements of
7.13 paragraph (b), clause (3). Services provided under this paragraph may not be transmitted
7.14 into the patient's residence.

7.15 (e) Medication therapy management services may be delivered into a patient's residence
7.16 via secure interactive video if the medication therapy management services are performed
7.17 electronically during a covered home care visit by an enrolled provider. Reimbursement
7.18 shall be at the same rates and under the same conditions that would otherwise apply to the
7.19 services provided. To qualify for reimbursement under this paragraph, the pharmacist
7.20 providing the services must meet the requirements of paragraph (b) and must be located
7.21 within an ambulatory care setting that meets the requirements of paragraph (b), clause (3)."

7.22 Renumber the sections in sequence and correct the internal references

7.23 Amend the title accordingly