

1.1 moves to amend H.F. No. 2932, the first engrossment, as follows:

1.2 Delete everything after the enacting clause and insert:

1.3 "Section 1. Minnesota Statutes 2012, section 152.02, subdivision 8b, is amended to read:

1.4 Subd. 8b. **Board of Pharmacy; expedited scheduling of additional substances.**

1.5 (a) The state Board of Pharmacy may, by rule, add a substance to Schedule I provided that
1.6 it finds that the substance has a high potential for abuse, has no currently accepted medical
1.7 use in the United States, has a lack of accepted safety for use under medical supervision,
1.8 has known adverse health effects, and is currently available for use within the state. For
1.9 the purposes of this subdivision only, the board may use the expedited rulemaking process
1.10 under section 14.389. ~~The scheduling of a substance under this subdivision expires the~~
1.11 ~~day after the adjournment of the legislative session immediately following the substance's~~
1.12 ~~scheduling unless the legislature by law ratifies the action.~~

1.13 (b) ~~If the board schedules a substance under this subdivision, the board shall notify~~
1.14 ~~in a timely manner the chairs and ranking minority members of the senate and house of~~
1.15 ~~representatives committees having jurisdiction over criminal justice and health policy~~
1.16 ~~and finance of the action and the reasons for it. The notice must include a copy of the~~
1.17 ~~administrative law judge's decision on the matter.~~

1.18 (c) ~~This subdivision expires August 1, 2014.~~

1.19 Sec. 2. Minnesota Statutes 2012, section 152.126, as amended by Laws 2013, chapter
1.20 113, article 3, section 3, is amended to read:

1.21 **~~152.126 CONTROLLED SUBSTANCES PRESCRIPTION ELECTRONIC~~**
1.22 **~~REPORTING SYSTEM~~ PRESCRIPTION MONITORING PROGRAM.**

1.23 Subdivision 1. **Definitions.** (a) For purposes of this section, the terms defined in
1.24 this subdivision have the meanings given.

2.1 (a) (b) "Board" means the Minnesota State Board of Pharmacy established under
2.2 chapter 151.

2.3 (b) (c) "Controlled substances" means those substances listed in section 152.02,
2.4 subdivisions 3 to 5 6, and those substances defined by the board pursuant to section
2.5 152.02, subdivisions 7, 8, and 12. For the purposes of this section, controlled substances
2.6 includes tramadol and butalbital.

2.7 (e) (d) "Dispense" or "dispensing" has the meaning given in section 151.01,
2.8 subdivision 30. Dispensing does not include the direct administering of a controlled
2.9 substance to a patient by a licensed health care professional.

2.10 (d) (e) "Dispenser" means a person authorized by law to dispense a controlled
2.11 substance, pursuant to a valid prescription. For the purposes of this section, a dispenser does
2.12 not include a licensed hospital pharmacy that distributes controlled substances for inpatient
2.13 hospital care or a veterinarian who is dispensing prescriptions under section 156.18.

2.14 (e) (f) "Prescriber" means a licensed health care professional who is authorized to
2.15 prescribe a controlled substance under section 152.12, subdivision 1 or 2.

2.16 (f) (g) "Prescription" has the meaning given in section 151.01, subdivision 16.

2.17 Subd. 1a. **Treatment of intractable pain.** This section is not intended to limit or
2.18 interfere with the legitimate prescribing of controlled substances for pain. No prescriber
2.19 shall be subject to disciplinary action by a health-related licensing board for prescribing a
2.20 controlled substance according to the provisions of section 152.125.

2.21 Subd. 2. **Prescription electronic reporting system.** (a) The board shall establish
2.22 by January 1, 2010, an electronic system for reporting the information required under
2.23 subdivision 4 for all controlled substances dispensed within the state.

2.24 (b) The board may contract with a vendor for the purpose of obtaining technical
2.25 assistance in the design, implementation, operation, and maintenance of the electronic
2.26 reporting system.

2.27 Subd. 3. **Prescription Electronic Reporting Monitoring Program Advisory**
2.28 **Committee Task Force.** (a) The board ~~shall convene~~ shall appoint an advisory committee.
2.29 ~~The committee must include~~ task force consisting of at least one representative of:

- 2.30 (1) the Department of Health;
- 2.31 (2) the Department of Human Services;
- 2.32 (3) each health-related licensing board that licenses prescribers;
- 2.33 (4) a professional medical association, which may include an association of pain
2.34 management and chemical dependency specialists;
- 2.35 (5) a professional pharmacy association;
- 2.36 (6) a professional nursing association;

- 3.1 (7) a professional dental association;
- 3.2 (8) a consumer privacy or security advocate; and
- 3.3 (9) a consumer or patient rights organization.

3.4 (b) The advisory ~~committee~~ task force shall advise the board on the development and

3.5 operation of the ~~electronic reporting system~~ prescription monitoring program, including,

3.6 but not limited to:

- 3.7 (1) technical standards for electronic prescription drug reporting;
- 3.8 (2) proper analysis and interpretation of prescription monitoring data; and
- 3.9 (3) an evaluation process for the program.

3.10 (c) The task force is governed by section 15.059. Notwithstanding section 15.059,

3.11 subdivision 5, the task force shall not expire.

3.12 Subd. 4. **Reporting requirements; notice.** (a) Each dispenser must submit the

3.13 following data to the board or its designated vendor, ~~subject to the notice required under~~

3.14 ~~paragraph (d)~~:

- 3.15 (1) name of the prescriber;
- 3.16 (2) national provider identifier of the prescriber;
- 3.17 (3) name of the dispenser;
- 3.18 (4) national provider identifier of the dispenser;
- 3.19 (5) prescription number;
- 3.20 (6) name of the patient for whom the prescription was written;
- 3.21 (7) address of the patient for whom the prescription was written;
- 3.22 (8) date of birth of the patient for whom the prescription was written;
- 3.23 (9) date the prescription was written;
- 3.24 (10) date the prescription was filled;
- 3.25 (11) name and strength of the controlled substance;
- 3.26 (12) quantity of controlled substance prescribed;
- 3.27 (13) quantity of controlled substance dispensed; and
- 3.28 (14) number of days supply.

3.29 (b) The dispenser must submit the required information by a procedure and in a

3.30 format established by the board. The board may allow dispensers to omit data listed in this

3.31 subdivision or may require the submission of data not listed in this subdivision provided

3.32 the omission or submission is necessary for the purpose of complying with the electronic

3.33 reporting or data transmission standards of the American Society for Automation in

3.34 Pharmacy, the National Council on Prescription Drug Programs, or other relevant national

3.35 standard-setting body.

4.1 (c) A dispenser is not required to submit this data for those controlled substance
4.2 prescriptions dispensed for:

4.3 ~~(1) individuals residing in licensed skilled nursing or intermediate care facilities;~~

4.4 ~~(2) individuals receiving assisted living services under chapter 144G or through a
4.5 medical assistance home and community-based waiver;~~

4.6 ~~(3) individuals receiving medication intravenously;~~

4.7 ~~(4) individuals receiving hospice and other palliative or end-of-life care; and~~

4.8 ~~(5) individuals receiving services from a home care provider regulated under chapter
4.9 144A.~~

4.10 (1) individuals residing in a health care facility as defined in section 151.58,
4.11 subdivision 2, paragraph (b), when a drug is distributed through the use of an automated
4.12 drug distribution system according to section 151.58; and

4.13 (2) individuals receiving a drug sample that was packaged by a manufacturer and
4.14 provided to the dispenser for dispensing as a professional sample pursuant to Code of
4.15 Federal Regulations, title 21, section 203, subpart D.

4.16 ~~(d) A dispenser must not submit data under this subdivision unless provide to the
4.17 patient for whom the prescription was written a conspicuous notice of the reporting
4.18 requirements of this section is given to the patient for whom the prescription was written
4.19 and notice that the information may be used for program administration purposes.~~

4.20 **Subd. 5. Use of data by board.** (a) The board shall develop and maintain a database
4.21 of the data reported under subdivision 4. The board shall maintain data that could identify
4.22 an individual prescriber or dispenser in encrypted form. Except as otherwise allowed
4.23 under subdivision 6, the database may be used by permissible users identified under
4.24 subdivision 6 for the identification of:

4.25 (1) individuals receiving prescriptions for controlled substances from prescribers
4.26 who subsequently obtain controlled substances from dispensers in quantities or with a
4.27 frequency inconsistent with generally recognized standards of use for those controlled
4.28 substances, including standards accepted by national and international pain management
4.29 associations; and

4.30 (2) individuals presenting forged or otherwise false or altered prescriptions for
4.31 controlled substances to dispensers.

4.32 (b) No permissible user identified under subdivision 6 may access the database
4.33 for the sole purpose of identifying prescribers of controlled substances for unusual or
4.34 excessive prescribing patterns without a valid search warrant or court order.

5.1 (c) No personnel of a state or federal occupational licensing board or agency may
5.2 access the database for the purpose of obtaining information to be used to initiate or
5.3 substantiate a disciplinary action against a prescriber.

5.4 (d) Data reported under subdivision 4 shall be ~~retained by the board in the database~~
5.5 ~~for a 12-month period, and shall be removed from the database no later than 12 months~~
5.6 ~~from the last day of the month during which the data was received.~~ made available to
5.7 permissible users for a 12-month period beginning the day the data was received and
5.8 ending 12 months from the last day of the month in which the data was received, except
5.9 that permissible users defined in subdivision 6, paragraph (b), clauses (5) and (6), may
5.10 use all data collected under this section for the purposes of administering, operating,
5.11 and maintaining the prescription monitoring program and conducting trend analyses
5.12 and other studies necessary to evaluate the effectiveness of the program. Data retained
5.13 beyond 12 months must be de-identified.

5.14 (e) The board shall not retain data reported under subdivision 4 for a period longer
5.15 than five years from the date the data was received.

5.16 Subd. 6. **Access to reporting system data.** (a) Except as indicated in this
5.17 subdivision, the data submitted to the board under subdivision 4 is private data on
5.18 individuals as defined in section 13.02, subdivision 12, and not subject to public disclosure.

5.19 (b) Except as specified in subdivision 5, the following persons shall be considered
5.20 permissible users and may access the data submitted under subdivision 4 in the same or
5.21 similar manner, and for the same or similar purposes, as those persons who are authorized
5.22 to access similar private data on individuals under federal and state law:

5.23 (1) a prescriber or an agent or employee of the prescriber to whom the prescriber has
5.24 delegated the task of accessing the data, to the extent the information relates specifically to
5.25 a current patient, to whom the prescriber is:

5.26 (i) prescribing or considering prescribing any controlled substance;

5.27 (ii) providing emergency medical treatment for which access to the data may be
5.28 necessary; or

5.29 (iii) providing other medical treatment for which access to the data may be necessary
5.30 and the patient has consented to access to the submitted data, and with the provision that
5.31 the prescriber remains responsible for the use or misuse of data accessed by a delegated
5.32 agent or employee;

5.33 (2) a dispenser or an agent or employee of the dispenser to whom the dispenser has
5.34 delegated the task of accessing the data, to the extent the information relates specifically
5.35 to a current patient to whom that dispenser is dispensing or considering dispensing any

6.1 controlled substance and with the provision that the dispenser remains responsible for the
6.2 use or misuse of data accessed by a delegated agent or employee;

6.3 (3) an individual who is the recipient of a controlled substance prescription for
6.4 which data was submitted under subdivision 4, or a guardian of the individual, parent or
6.5 guardian of a minor, or health care agent of the individual acting under a health care
6.6 directive under chapter 145C;

6.7 (4) personnel of the board specifically assigned to conduct a bona fide investigation
6.8 of a specific licensee;

6.9 (5) personnel of the board engaged in the collection, review, and analysis
6.10 of controlled substance prescription information as part of the assigned duties and
6.11 responsibilities under this section;

6.12 (6) authorized personnel of a vendor under contract with the ~~board~~ state of
6.13 Minnesota who are engaged in the design, implementation, operation, and maintenance of
6.14 the ~~electronic reporting system~~ prescription monitoring program as part of the assigned
6.15 duties and responsibilities of their employment, provided that access to data is limited to
6.16 the minimum amount necessary to carry out such duties and responsibilities, and subject
6.17 to the requirement of de-identification and time limit on retention of data specified in
6.18 subdivision 5, paragraphs (d) and (e);

6.19 (7) federal, state, and local law enforcement authorities acting pursuant to a valid
6.20 search warrant;

6.21 (8) personnel of the ~~medical assistance program~~ Minnesota health care programs
6.22 assigned to use the data collected under this section to identify recipients whose usage of
6.23 controlled substances may warrant restriction to a single primary care physician provider,
6.24 a single outpatient pharmacy, ~~or~~ and a single hospital; ~~and~~

6.25 (9) personnel of the Department of Human Services assigned to access the data
6.26 pursuant to paragraph (h); and

6.27 (10) personnel of the health professionals services program established under section
6.28 214.31, to the extent that the information relates specifically to an individual who is
6.29 currently enrolled in and being monitored by the program, and the individual consents to
6.30 access to that information. The health professionals services program personnel shall not
6.31 provide this data to a health-related licensing board or the Emergency Medical Services
6.32 Regulatory Board, except as permitted under section 214.33, subdivision 3.

6.33 For purposes of clause ~~(3)~~ (4), access by an individual includes persons in the
6.34 definition of an individual under section 13.02.

6.35 (c) ~~Any~~ A permissible user identified in paragraph (b), ~~who~~ clauses (1), (2), (5), (6),
6.36 and (8) may directly accesses access the data electronically; If the data is directly accessed

7.1 electronically, the permissible user shall implement and maintain a comprehensive
7.2 information security program that contains administrative, technical, and physical
7.3 safeguards that are appropriate to the user's size and complexity, and the sensitivity of the
7.4 personal information obtained. The permissible user shall identify reasonably foreseeable
7.5 internal and external risks to the security, confidentiality, and integrity of personal
7.6 information that could result in the unauthorized disclosure, misuse, or other compromise
7.7 of the information and assess the sufficiency of any safeguards in place to control the risks.

7.8 (d) The board shall not release data submitted under ~~this section~~ subdivision 4 unless
7.9 it is provided with evidence, satisfactory to the board, that the person requesting the
7.10 information is entitled to receive the data.

7.11 ~~(e) The board shall not release the name of a prescriber without the written consent~~
7.12 ~~of the prescriber or a valid search warrant or court order. The board shall provide a~~
7.13 ~~mechanism for a prescriber to submit to the board a signed consent authorizing the release~~
7.14 ~~of the prescriber's name when data containing the prescriber's name is requested.~~

7.15 ~~(f)~~ (e) The board shall maintain a log of all persons who access the data for a period
7.16 of at least three years and shall ensure that any permissible user complies with paragraph
7.17 (c) prior to attaining direct access to the data.

7.18 ~~(g)~~ (f) Section 13.05, subdivision 6, shall apply to any contract the board enters into
7.19 pursuant to subdivision 2. A vendor shall not use data collected under this section for
7.20 any purpose not specified in this section.

7.21 (g) The board may participate in an interstate prescription monitoring program data
7.22 exchange system provided that permissible users in other states have access to the data
7.23 only as allowed under this section, and that section 13.05, subdivision 6, applies to any
7.24 contract or memorandum of understanding that the board enters into under this paragraph.

7.25 (h) With available appropriations, the commissioner of human services shall
7.26 establish and implement a system through which the Department of Human Services shall
7.27 routinely access the data for the purpose of determining whether any client enrolled in
7.28 an opioid treatment program licensed according to chapter 245A has been prescribed or
7.29 dispensed a controlled substance in addition to that administered or dispensed by the
7.30 opioid treatment program. When the commissioner determines there have been multiple
7.31 prescribers or multiple prescriptions of controlled substances, the commissioner shall:

7.32 (1) inform the medical director of the opioid treatment program only that the
7.33 commissioner determined the existence of multiple prescribers or multiple prescriptions of
7.34 controlled substances; and

8.1 (2) direct the medical director of the opioid treatment program to access the data
8.2 directly, review the effect of the multiple prescribers or multiple prescriptions, and
8.3 document the review.

8.4 If determined necessary, the commissioner of human services shall seek a federal waiver
8.5 of, or exception to, any applicable provision of Code of Federal Regulations, title 42, part
8.6 2.34, item (c), prior to implementing this paragraph.

8.7 (i) The board may provide de-identified data submitted under subdivision 4 for public
8.8 research, policy, or education purposes, that does not involve information that is likely to
8.9 reveal the identity of the patient, prescriber, or dispenser who is the subject of the data.

8.10 **Subd. 7. Disciplinary action.** (a) A dispenser who knowingly fails to submit data to
8.11 the board as required under this section is subject to disciplinary action by the appropriate
8.12 health-related licensing board.

8.13 (b) A prescriber or dispenser authorized to access the data who knowingly discloses
8.14 the data in violation of state or federal laws relating to the privacy of health care data
8.15 shall be subject to disciplinary action by the appropriate health-related licensing board,
8.16 and appropriate civil penalties.

8.17 ~~Subd. 8. Evaluation and reporting.~~ (a) ~~The board shall evaluate the prescription~~
8.18 ~~electronic reporting system to determine if the system is negatively impacting appropriate~~
8.19 ~~prescribing practices of controlled substances. The board may contract with a vendor to~~
8.20 ~~design and conduct the evaluation.~~

8.21 ~~(b) The board shall submit the evaluation of the system to the legislature by July~~
8.22 ~~15, 2011.~~

8.23 **Subd. 9. Immunity from liability; no requirement to obtain information.** (a) A
8.24 pharmacist, prescriber, or other dispenser making a report to the program in good faith
8.25 under this section is immune from any civil, criminal, or administrative liability, which
8.26 might otherwise be incurred or imposed as a result of the report, or on the basis that the
8.27 pharmacist or prescriber did or did not seek or obtain or use information from the program.

8.28 (b) Nothing in this section shall require a pharmacist, prescriber, or other dispenser
8.29 to obtain information about a patient from the program, and the pharmacist, prescriber,
8.30 or other dispenser, if acting in good faith, is immune from any civil, criminal, or
8.31 administrative liability that might otherwise be incurred or imposed for requesting,
8.32 receiving, or using information from the program.

8.33 **Subd. 10. Funding.** (a) The board may seek grants and private funds from nonprofit
8.34 charitable foundations, the federal government, and other sources to fund the enhancement
8.35 and ongoing operations of the prescription electronic reporting system monitoring
8.36 program established under this section. Any funds received shall be appropriated to the

9.1 board for this purpose. The board may not expend funds to enhance the program in a way
 9.2 that conflicts with this section without seeking approval from the legislature.

9.3 (b) Notwithstanding any other section, the administrative services unit for the
 9.4 health-related licensing boards shall apportion between the Board of Medical Practice, the
 9.5 Board of Nursing, the Board of Dentistry, the Board of Podiatric Medicine, the Board of
 9.6 Optometry, the Board of Veterinary Medicine, and the Board of Pharmacy an amount to
 9.7 be paid through fees by each respective board. The amount apportioned to each board
 9.8 shall equal each board's share of the annual appropriation to the Board of Pharmacy
 9.9 from the state government special revenue fund for operating the prescription ~~electronic~~
 9.10 ~~reporting system~~ monitoring program under this section. Each board's apportioned share
 9.11 shall be based on the number of prescribers or dispensers that each board identified in
 9.12 this paragraph licenses as a percentage of the total number of prescribers and dispensers
 9.13 licensed collectively by these boards. Each respective board may adjust the fees that the
 9.14 boards are required to collect to compensate for the amount apportioned to each board by
 9.15 the administrative services unit.

9.16 Sec. 3. Minnesota Statutes 2012, section 214.32, is amended to read:

9.17 **214.32 PROGRAM OPERATIONS AND RESPONSIBILITIES.**

9.18 Subdivision 1. **Management.** (a) A Health Professionals Services Program
 9.19 Committee is established, consisting of ~~one person appointed by each participating~~
 9.20 ~~board, with each participating board having one vote.~~ no fewer than three, or more than
 9.21 six, executive directors of health-related licensing boards or their designees, and two
 9.22 members of the advisory committee established in paragraph (d). Program committee
 9.23 members from the health-related licensing boards shall be appointed by a means agreeable
 9.24 to the executive directors of the health-related licensing boards in July of odd-numbered
 9.25 years. Members from the advisory committee shall be appointed by a means agreeable to
 9.26 advisory committee members in July of odd-numbered years. The program committee
 9.27 ~~shall designate one board to provide administrative management of the program, set the~~
 9.28 ~~program budget and the pro rata share of~~ administrative costs under paragraph (b) and
 9.29 program expenses to be borne by each participating board, set the program budget, and
 9.30 ensure the program is meeting its statutory charge. The program committee shall establish
 9.31 uniform criteria and procedures governing termination and discharge for all health
 9.32 professionals served by the health professionals services program.

9.33 (b) The commissioner of administration shall provide guidance on the general
 9.34 operation of the program, including hiring of program personnel, and ensure that the
 9.35 program's direction is in accord with its authority. If the participating boards change

10.1 ~~which board is designated to provide administrative management of the program, any~~
 10.2 ~~appropriation remaining for the program shall transfer to the newly designated board on~~
 10.3 ~~the effective date of the change. The participating boards must inform the appropriate~~
 10.4 ~~legislative committees and the commissioner of management and budget of any change~~
 10.5 ~~in the administrative management of the program, and the amount of any appropriation~~
 10.6 ~~transferred under this provision.~~

10.7 ~~(b) (c)~~ The designated board, upon recommendation of the Health Professional
 10.8 ~~Services Program Committee, commissioner of administration~~ shall hire the program
 10.9 manager and employees and pay expenses of the program from funds appropriated for that
 10.10 purpose. The designated board commissioner of administration may apply for grants to
 10.11 pay program expenses and may enter into contracts on behalf of the program to carry out
 10.12 the purposes of the program. The participating boards shall enter into written agreements
 10.13 with the ~~designated board~~ commissioner of administration.

10.14 ~~(e) (d)~~ An advisory committee is established to advise the program committee
 10.15 consisting of:

10.16 (1) one member appointed by each of the following: ~~the Minnesota Academy of~~
 10.17 ~~Physician Assistants, the Minnesota Dental Association, the Minnesota Chiropractic~~
 10.18 ~~Association, the Minnesota Licensed Practical Nurse Association, the Minnesota Medical~~
 10.19 ~~Association, the Minnesota Nurses Association, and the Minnesota Podiatric Medicine~~
 10.20 ~~Association~~ of the professional associations whose members are eligible for health
 10.21 professionals services program services; and

10.22 (2) ~~one member appointed by each of the professional associations of the other~~
 10.23 ~~professions regulated by a participating board not specified in clause (1); and~~

10.24 ~~(3) (2)~~ two public members, as defined by section 214.02.

10.25 Members of the advisory committee shall be appointed for two years and members
 10.26 may be reappointed.

10.27 Subd. 2. **Services.** (a) The program shall provide the following services to program
 10.28 participants:

10.29 (1) referral of eligible regulated persons to qualified professionals for evaluation,
 10.30 treatment, and a written plan for continuing care consistent with the regulated person's
 10.31 illness. The referral shall take into consideration the regulated person's financial resources
 10.32 as well as specific needs;

10.33 (2) development of individualized program participation agreements between
 10.34 participants and the program to meet the needs of participants and protect the public. An
 10.35 agreement may include, but need not be limited to, recommendations from the continuing
 10.36 care plan, practice monitoring, health monitoring, practice restrictions, random drug

11.1 screening, support group participation, filing of reports necessary to document compliance,
 11.2 and terms for successful completion of the regulated person's program; and

11.3 (3) monitoring of compliance by participants with individualized program
 11.4 participation agreements or board orders.

11.5 (b) The program may develop services related to sections 214.31 to 214.37 for
 11.6 employers and colleagues of regulated persons from participating boards.

11.7 Subd. 3. **Participant costs.** Each program participant shall be responsible for
 11.8 paying for the costs of physical, psychosocial, or other related evaluation, treatment,
 11.9 laboratory monitoring, and random drug screens.

11.10 Subd. 4. **Eligibility.** Admission to the health professional services program is
 11.11 available to a person regulated by a participating board who is unable to practice with
 11.12 reasonable skill and safety by reason of illness, use of alcohol, drugs, chemicals, or
 11.13 any other materials, or as a result of any mental, physical, or psychological condition.
 11.14 Admission in the health professional services program shall be denied to persons:

11.15 (1) who have diverted controlled substances for other than self-administration;

11.16 (2) who have been terminated from this or any other state professional services
 11.17 program for noncompliance in the program, unless referred by a participating board or the
 11.18 commissioner of health;

11.19 (3) currently under a board disciplinary order or corrective action agreement, unless
 11.20 referred by a board;

11.21 (4) ~~regulated under sections 214.17 to 214.25, unless referred by a board or by the~~
 11.22 ~~commissioner of health;~~

11.23 (5) accused of sexual misconduct; or

11.24 (6) (5) whose continued practice would create a serious risk of harm to the public.

11.25 Subd. 5. **Completion; voluntary termination; discharge.** (a) A regulated person
 11.26 completes the program when the terms of the program participation agreement are fulfilled.

11.27 (b) A regulated person may voluntarily terminate participation in the health
 11.28 professionals service program at any time ~~by reporting to the person's board~~ which shall
 11.29 result in the program manager making a report to the regulated person's board under
 11.30 section 214.33, subdivision 3.

11.31 (c) The program manager may choose to discharge a regulated person from the
 11.32 program and make a referral to the person's board at any time for reasons including but not
 11.33 limited to: the degree of cooperation and compliance by the regulated person, the inability
 11.34 to secure information or the medical records of the regulated person, or indication of other
 11.35 possible violations of the regulated person's practice act. The regulated person shall be
 11.36 notified in writing by the program manager of any change in the person's program status.

12.1 A regulated person who has been terminated or discharged from the program may be
 12.2 referred back to the program for monitoring.

12.3 Subd. 6. **Duties of a health related licensing board.** (a) Upon receiving notice from
 12.4 the program manager that a regulated person has been discharged due to noncompliance
 12.5 or voluntary withdrawal, when the appropriate licensing board has probable cause to
 12.6 believe continued practice by the regulated person presents an imminent risk of harm, the
 12.7 licensing board shall temporarily suspend the regulated person's professional license. The
 12.8 suspension shall take effect upon written notice to the regulated person and shall specify
 12.9 the reason for the suspension.

12.10 (b) The suspension shall remain in effect until the appropriate licensing board
 12.11 completes an investigation and issues a final order in the matter after a hearing.

12.12 (c) At the time it issues the suspension notice, the appropriate licensing board shall
 12.13 schedule a disciplinary hearing to be held pursuant to the Administrative Procedure Act.
 12.14 The regulated person shall be provided with at least 20 days' notice of any hearing held
 12.15 pursuant to this subdivision. The hearing shall be scheduled to being no later than 60
 12.16 days after issuance of the suspension order.

12.17 Sec. 4. Minnesota Statutes 2013 Supplement, section 254A.035, subdivision 2, is
 12.18 amended to read:

12.19 Subd. 2. **Membership terms, compensation, removal and expiration.** The
 12.20 membership of this council shall be composed of 17 persons who are American Indians
 12.21 and who are appointed by the commissioner. The commissioner shall appoint one
 12.22 representative from each of the following groups: Red Lake Band of Chippewa Indians;
 12.23 Fond du Lac Band, Minnesota Chippewa Tribe; Grand Portage Band, Minnesota
 12.24 Chippewa Tribe; Leech Lake Band, Minnesota Chippewa Tribe; Mille Lacs Band,
 12.25 Minnesota Chippewa Tribe; Bois Forte Band, Minnesota Chippewa Tribe; White Earth
 12.26 Band, Minnesota Chippewa Tribe; Lower Sioux Indian Reservation; Prairie Island Sioux
 12.27 Indian Reservation; Shakopee Mdewakanton Sioux Indian Reservation; Upper Sioux
 12.28 Indian Reservation; International Falls Northern Range; Duluth Urban Indian Community;
 12.29 and two representatives from the Minneapolis Urban Indian Community and two from the
 12.30 St. Paul Urban Indian Community. The terms, compensation, and removal of American
 12.31 Indian Advisory Council members shall be as provided in section 15.059. Notwithstanding
 12.32 section 15.059, subdivision 5, the council expires June 30, 2014 does not expire.

12.33 **EFFECTIVE DATE.** This section is effective the day following final enactment.

12.34 Sec. 5. Minnesota Statutes 2013 Supplement, section 254A.04, is amended to read:

13.1 **254A.04 CITIZENS ADVISORY COUNCIL.**

13.2 There is hereby created an Alcohol and Other Drug Abuse Advisory Council to
 13.3 advise the Department of Human Services concerning the problems of alcohol and
 13.4 other drug dependency and abuse, composed of ten members. Five members shall be
 13.5 individuals whose interests or training are in the field of alcohol dependency and abuse;
 13.6 and five members whose interests or training are in the field of dependency and abuse of
 13.7 drugs other than alcohol. The terms, compensation and removal of members shall be as
 13.8 provided in section 15.059. Notwithstanding section 15.059, subdivision 5, the council
 13.9 expires June 30, 2014 does not expire. The commissioner of human services shall appoint
 13.10 members whose terms end in even-numbered years. The commissioner of health shall
 13.11 appoint members whose terms end in odd-numbered years.

13.12 **EFFECTIVE DATE.** This section is effective the day following final enactment.

13.13 Sec. 6. Minnesota Statutes 2012, section 256B.0751, is amended by adding a
 13.14 subdivision to read:

13.15 Subd. 10. **Health care homes advisory committee.** (a) The commissioners of
 13.16 health and human services shall establish a health care homes advisory committee to
 13.17 advise the commissioners on the ongoing statewide implementation of the health care
 13.18 homes program authorized in this section.

13.19 (b) The commissioners shall establish an advisory committee that includes
 13.20 representatives of the health care professions such as primary care providers; mental
 13.21 health providers; nursing and care coordinators; certified health care home clinics with
 13.22 statewide representation; health plan companies; state agencies; employers; academic
 13.23 researchers; consumers; and organizations that work to improve health care quality in
 13.24 Minnesota. At least 25 percent of the committee members must be consumers or patients
 13.25 in health care homes.

13.26 (c) The advisory committee shall advise the commissioners on ongoing
 13.27 implementation of the health care homes program, including, but not limited to, the
 13.28 following activities:

13.29 (1) implementation of certified health care homes across the state on performance
 13.30 management and implementation of benchmarking;

13.31 (2) implementation of modifications to the health care homes program based on
 13.32 results of the legislatively mandated health care home evaluation;

13.33 (3) statewide solutions for engagement of employers and commercial payers;

13.34 (4) potential modifications of the health care home rules or statutes;

14.1 (5) consumer engagement, including patient and family-centered care, patient
 14.2 activation in health care, and shared decision making;

14.3 (6) oversight for health care home subject matter task forces or workgroups; and

14.4 (7) other related issues as requested by the commissioners.

14.5 (d) The advisory committee shall have the ability to establish subcommittees on
 14.6 specific topics. The advisory committee is governed by section 15.059. Notwithstanding
 14.7 section 15.059, the advisory committee does not expire.

14.8 Sec. 7. Minnesota Statutes 2013 Supplement, section 260.835, subdivision 2, is
 14.9 amended to read:

14.10 Subd. 2. **Expiration.** Notwithstanding section 15.059, subdivision 5, the American
 14.11 Indian Child Welfare Advisory Council ~~expires June 30, 2014~~ does not expire.

14.12 **EFFECTIVE DATE.** This section is effective the day following final enactment.

14.13 Sec. 8. **MINNESOTA TANF EXPENDITURES TASK FORCE.**

14.14 Subdivision 1. **Establishment.** The Minnesota TANF Expenditures Task Force is
 14.15 established to analyze past temporary assistance for needy families (TANF) expenditures
 14.16 and make recommendations as to which, if any, programs currently receiving TANF
 14.17 funding should be funded by the general fund so that a greater portion of TANF funds
 14.18 can go directly to Minnesota families receiving assistance through the Minnesota family
 14.19 investment program under Minnesota Statutes, chapter 256J.

14.20 Subd. 2. **Membership; meetings; staff.** (a) The task force shall be composed of the
 14.21 following members who serve at the pleasure of their appointing authority:

14.22 (1) one representative of the Department of Human Services appointed by the
 14.23 commissioner of human services;

14.24 (2) one representative of the Department of Management and Budget appointed by
 14.25 the commissioner of management and budget;

14.26 (3) one representative of the Department of Health appointed by the commissioner
 14.27 of health;

14.28 (4) one representative of the Local Public Health Association of Minnesota;

14.29 (5) two representatives of county government appointed by the Association of
 14.30 Minnesota Counties, one representing counties in the seven-county metropolitan area
 14.31 and one representing all other counties;

14.32 (6) one representative of the Minnesota Legal Services Coalition;

14.33 (7) one representative of the Children's Defense Fund of Minnesota;

- 15.1 (8) one representative of the Minnesota Coalition for the Homeless;
15.2 (9) one representative of the Welfare Rights Coalition;
15.3 (10) two members of the house of representatives, one appointed by the speaker of
15.4 the house and one appointed by the minority leader; and
15.5 (11) two members of the senate, including one member of the minority party,
15.6 appointed according to the rules of the senate.

15.7 (b) Notwithstanding Minnesota Statutes, section 15.059, members of the task force
15.8 shall serve without compensation or reimbursement of expenses.

15.9 (c) The commissioner of human services must convene the first meeting of the
15.10 Minnesota TANF Expenditures Task Force by July 31, 2014. The task force must meet at
15.11 least quarterly.

15.12 (d) Staffing and technical assistance shall be provided within available resources by
15.13 the Department of Human Services, children and family services division.

15.14 Subd. 3. **Duties.** (a) The task force must report on past expenditures of the TANF
15.15 block grant, including a determination of whether or not programs for which TANF funds
15.16 have been appropriated meet the purposes of the TANF program as defined under Code of
15.17 Federal Regulations, title 45, section 260.20, and make recommendations as to which,
15.18 if any, programs currently receiving TANF funds should be funded by the general fund.
15.19 In making recommendations on program funding sources, the task force shall consider
15.20 the following:

15.21 (1) the original purpose of the TANF block grant under Code of Federal Regulations,
15.22 title 45, section 260.20;

15.23 (2) potential overlap of the population eligible for the Minnesota family investment
15.24 program cash grant and the other programs currently receiving TANF funds;

15.25 (3) the ability for TANF funds, as appropriated under current law, to effectively help
15.26 the lowest-income Minnesotans out of poverty;

15.27 (4) the impact of past expenditures on families who may be eligible for assistance
15.28 through TANF;

15.29 (5) the ability of TANF funds to support effective parenting and optimal brain
15.30 development in children under five years old; and

15.31 (6) the role of noncash assistance expenditures in maintaining compliance with
15.32 federal law.

15.33 (b) In preparing the recommendations under paragraph (a), the task force shall
15.34 consult with appropriate Department of Human Services information technology staff
15.35 regarding implementation of the recommendations.

16.1 Subd. 4. **Report.** (a) The task force must submit an initial report by November
16.2 30, 2014, on past expenditures of the TANF block grant in Minnesota to the chairs and
16.3 ranking minority members of the legislative committees with jurisdiction over health and
16.4 human services policy and finance.

16.5 (b) The task force must submit a final report by February 1, 2015, analyzing past
16.6 TANF expenditures and making recommendations as to which programs, if any, currently
16.7 receiving TANF funding should be funded by the general fund, including any phase-in
16.8 period and draft legislation necessary for implementation, to the chairs and ranking
16.9 minority members of the legislative committees with jurisdiction over health and human
16.10 services policy and finance.

16.11 Subd. 5. **Expiration.** This section expires March 1, 2015, or upon submission of the
16.12 final report required under subdivision 4, whichever is earlier."

16.13 Delete the title and insert:

16.14 "A bill for an act
16.15 relating to governmental operations; establishing and modifying health and
16.16 human services advisory councils, committees, and task forces; modifying
16.17 rulemaking authority for the Board of Pharmacy; changing the prescription
16.18 monitoring program; amending Minnesota Statutes 2012, sections 152.02,
16.19 subdivision 8b; 152.126, as amended; 214.32; 256B.0751, by adding a
16.20 subdivision; Minnesota Statutes 2013 Supplement, sections 254A.035,
16.21 subdivision 2; 254A.04; 260.835, subdivision 2."