SENATE STATE OF MINNESOTA NINETY-FOURTH SESSION

S.F. No. 2370

(SENATE AUTI	HORS: DIBB	LE and Port)
DATE	D-PG	OFFICIAL STATUS
03/10/2025	714	Introduction and first reading
		Referred to Commerce and Consumer Protection
04/03/2025	1287a	Comm report: To pass as amended
	1344	Second reading
04/07/2025	1715	Author added Port
04/28/2025	4115a	Special Order: Amended
	4126	Third reading Passed as amended
05/06/2025	4452	Returned from House with amendment
	4453	Senate not concur, conference committee of 3 requested
	4511	Senate conferees Dibble; Port; Rasmusson
05/08/2025	4603	House conferees Stephenson; Hanson, J; West; Allen
05/17/2025		Conference committee report, delete everything
		Senate adopted CC report and repassed bill
		Third Reading Repassed

1.1 A bill for an act

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relating to cannabis; including the Office of Cannabis Management as an agency for the purpose of having a government-to-government relationship with Tribal governments; modifying provisions regarding the sale of cannabinoids derived from hemp; modifying medical cannabis provisions; modifying hemp-derived topical product provisions; modifying cannabis license application requirements; modifying the limits of delta-9 tetrahydrocannabinol in edible cannabinoid products and lower-potency hemp edibles when intended to be consumed as beverages; allowing samples at cannabis events; modifying expungement and resentencing provisions for felony cannabis offenses; amending Minnesota Statutes 2024, sections 10.65, subdivision 2; 151.72, subdivisions 3, 5a; 152.22, subdivisions 4, 7, 10, 13; 152.24; 152.25; 152.26; 152.261; 152.27, subdivisions 2, 7; 152.28, subdivisions 1, 3; 152.29, subdivisions 1, 2, 3a, 4; 152.31; 152.32, subdivision 2; 152.33, subdivisions 1a, 4; 152.35; 152.37; 342.01, subdivisions 9, 47, 50, 71, by adding subdivisions; 342.02, subdivision 3; 342.09, subdivision 2; 342.12; 342.14, subdivisions 1, 3, 6; 342.151, subdivisions 2, 3; 342.22, subdivision 3; 342.28, subdivisions 1, 8; 342.29, subdivisions 1, 7; 342.30, subdivision 1; 342.32, subdivisions 4, 5; 342.33, subdivision 1; 342.40, subdivision 7, by adding a subdivision; 342.43, by adding a subdivision; 342.44, subdivision 1; 342.45, by adding a subdivision; 342.46, subdivision 6; 342.51, subdivision 2, by adding a subdivision; 342.52, subdivision 9, by adding a subdivision; 342.56, subdivision 2; 342.57; 342.59, subdivision 2; 342.61, subdivision 4; 342.63, subdivisions 2, 3, 5, 6; 342.66, subdivision 6; 609A.06, subdivisions 3, 7, 10, 12; repealing Minnesota Statutes 2024, sections 152.22, subdivision 2; 342.151, subdivision 1.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF MINNESOTA:

- Section 1. Minnesota Statutes 2024, section 10.65, subdivision 2, is amended to read:
- 1.27 Subd. 2. **Definitions.** As used in this section, the following terms have the meanings given:
- (1) "agency" means the Department of Administration; Department of Agriculture;
- Department of Children, Youth, and Families; Department of Commerce; Department of
- 1.31 Corrections; Department of Education; Department of Employment and Economic

Section 1.

- (4) "Minnesota Tribal governments" means the federally recognized Indian Tribes located in Minnesota including: Bois Forte Band; Fond Du Lac Band; Grand Portage Band; Leech Lake Band; Mille Lacs Band; White Earth Band; Red Lake Nation; Lower Sioux Indian Community; Prairie Island Indian Community; Shakopee Mdewakanton Sioux Community; and Upper Sioux Community; and
- (5) "timely and meaningful" means done or occurring at a favorable or useful time that allows the result of consultation to be included in the agency's decision-making process for a matter that has Tribal implications.

Section 1. 2

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Sec. 2. Minnesota Statutes 2024, section 151.72, subdivision 3, is amended to read:

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- Subd. 3. Sale of cannabinoids derived from hemp. (a) Notwithstanding any other section of this chapter, a product containing nonintoxicating cannabinoids, including an edible cannabinoid product, may be sold for human or animal consumption only if all of the requirements of this section are met. A product sold for human or animal consumption must not contain more than 0.3 percent of any tetrahydrocannabinol and an edible cannabinoid product must not contain an amount of any tetrahydrocannabinol that exceeds the limits established in subdivision 5a, paragraph (f).
- (b) A product containing nonintoxicating cannabinoids, other than an edible cannabinoid product, may be sold for human or animal consumption only if it is intended for application externally to a part of the body of a human or animal. Such a product must not be manufactured, marketed, distributed, or intended to be consumed:
- (1) by combustion or vaporization of the product and inhalation of smoke, aerosol, or vapor from the product;
 - (2) through chewing, drinking, or swallowing; or
- (3) through injection or application to nonintact skin or a mucous membrane or nonintact skin, except for products applied sublingually.
- (c) No other substance extracted or otherwise derived from hemp may be sold for human consumption if the substance is intended:
- (1) for external or internal use in the diagnosis, cure, mitigation, treatment, or prevention of disease in humans or other animals; or
 - (2) to affect the structure or any function of the bodies of humans or other animals.
- (d) No product containing any cannabinoid or tetrahydrocannabinol extracted or otherwise derived from hemp may be sold to any individual who is under the age of 21.
- (e) Products that meet the requirements of this section are not controlled substances under section 152.02.
- (f) Products may be sold for on-site consumption if all of the following conditions are 3.27 3.28 met:
 - (1) the retailer must also hold an on-sale license issued under chapter 340A;
 - (2) products, other than products that are intended to be consumed as a beverage, must be served in original packaging, but may be removed from the products' packaging by customers and consumed on site;

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4.1	(3) products must not be sold to a customer who the retailer knows or reasonably should
4.2	know is intoxicated;
4.3	(4) products must not be permitted to be mixed with an alcoholic beverage; and
4.4	(5) products that have been removed from packaging must not be removed from the
4.5	premises.
4.6	(g) Edible cannabinoid products that are intended to be consumed as a beverage may be
4.7	served outside of the products' packaging if the information that is required to be contained
4.8	on the label of an edible cannabinoid product is posted or otherwise displayed by the retailer.
4.9	Sec. 3. Minnesota Statutes 2024, section 151.72, subdivision 5a, is amended to read:
4.10	Subd. 5a. Additional requirements for edible cannabinoid products. (a) In addition
4.11	to the testing and labeling requirements under subdivisions 4 and 5, an edible cannabinoid
4.12	must meet the requirements of this subdivision.
4.13	(b) An edible cannabinoid product must not:
4.14	(1) bear the likeness or contain cartoon-like characteristics of a real or fictional person,
4.15	animal, or fruit that appeals to children;
4.16	(2) be modeled after a brand of products primarily consumed by or marketed to children;
4.17	(3) be made by applying an extracted or concentrated hemp-derived cannabinoid to a
4.18	commercially available candy or snack food item;
4.19	(4) be substantively similar to a meat food product; poultry food product as defined in
4.20	section 31A.02, subdivision 10; or a dairy product as defined in section 32D.01, subdivision
4.21	7;
4.22	(5) contain an ingredient, other than a hemp-derived cannabinoid, that is not approved
4.23	by the United States Food and Drug Administration for use in food;
4.24	(6) be packaged in a way that resembles the trademarked, characteristic, or
4.25	product-specialized packaging of any commercially available food product; or
4.26	(7) be packaged in a container that includes a statement, artwork, or design that could
4.27	reasonably mislead any person to believe that the package contains anything other than an
4.28	edible cannabinoid product.
4.29	(c) An edible cannabinoid product must be prepackaged in packaging or a container that

is child-resistant, tamper-evident, and opaque or placed in packaging or a container that is

child-resistant, tamper-evident, and opaque at the final point of sale to a customer. The

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requirement that packaging be child-resistant does not apply to an edible cannabinoid product that is intended to be consumed as a beverage.

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- (d) If an edible cannabinoid product, other than a product that is intended to be consumed as a beverage, is intended for more than a single use or contains multiple servings, each serving must be indicated by scoring, wrapping, or other indicators designating the individual serving size that appear on the edible cannabinoid product. If it is not possible to indicate a single serving by scoring or use of another indicator that appears on the product, the edible cannabinoid product may not be packaged in a manner that includes more than a single serving in each container, except that a calibrated dropper, measuring spoon, or similar device for measuring a single serving, when sold with the product, may be used for any edible cannabinoid products that are intended to be combined with food or beverage products prior to consumption.
- (e) A label containing at least the following information must be affixed to the packaging or container of all edible cannabinoid products sold to consumers:
- (1) the serving size;
 - (2) the cannabinoid profile per serving and in total;
- (3) a list of ingredients, including identification of any major food allergens declared 5.17 by name; and 5.18
 - (4) the following statement: "Keep this product out of reach of children."
 - (f) An edible cannabinoid product must not contain more than five milligrams of any tetrahydrocannabinol in a single serving, except that an edible cannabinoid product that is intended to be consumed as a beverage may contain no more than ten milligrams of any tetrahydrocannabinol in a single-serving container. An edible cannabinoid product, other than a product that is intended to be consumed as a beverage, may not contain more than a total of 50 milligrams of any tetrahydrocannabinol per package. An edible cannabinoid product that is intended to be consumed as a beverage may not contain more than two servings per container.
 - (g) An edible cannabinoid product may contain delta-8 tetrahydrocannabinol or delta-9 tetrahydrocannabinol that is extracted from hemp plants or hemp plant parts or is an artificially derived cannabinoid. Edible cannabinoid products are prohibited from containing any other artificially derived cannabinoid, including but not limited to THC-P, THC-O, and HHC, unless the office authorizes use of the artificially derived cannabinoid in edible

5 Sec. 3

cannabinoid products. Edible cannabinoid products are prohibited from containing synthetic
cannabinoids.

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- (h) Every person selling edible cannabinoid products to consumers, other than products that are intended to be consumed as a beverage, must ensure that all edible cannabinoid products are displayed behind a checkout counter where the public is not permitted or in a locked case.
- Sec. 4. Minnesota Statutes 2024, section 152.22, subdivision 4, is amended to read:
- Subd. 4. **Health care practitioner.** "Health care practitioner" means a Minnesota licensed Minnesota-licensed doctor of medicine, a Minnesota licensed Minnesota-licensed physician assistant acting within the scope of authorized practice, or a Minnesota licensed Minnesota-licensed advanced practice registered nurse who has an active license in good standing and the primary responsibility for the care and treatment of the qualifying medical condition of a person an individual diagnosed with a qualifying medical condition.
- Sec. 5. Minnesota Statutes 2024, section 152.22, subdivision 7, is amended to read:
- Subd. 7. **Medical cannabis manufacturer.** "Medical cannabis manufacturer" or

 "manufacturer" means an entity registered by the commissioner office to cultivate, acquire,
 manufacture, possess, prepare, transfer, transport, supply, or dispense medical cannabis,
 delivery devices, or related supplies and educational materials.
- Sec. 6. Minnesota Statutes 2024, section 152.22, subdivision 10, is amended to read:
- Subd. 10. **Patient registry number.** "Patient registry number" means a unique identification number assigned by the <u>commissioner office</u> to a patient enrolled in the registry program.
- 6.23 Sec. 7. Minnesota Statutes 2024, section 152.22, subdivision 13, is amended to read:
- Subd. 13. **Registry verification.** "Registry verification" means the verification provided by the <u>commissioner office</u> that a patient is enrolled in the registry program and that includes the patient's name, registry number, and, if applicable, the name of the patient's registered designated caregiver or parent, legal guardian, or spouse.

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Sec. 8. Minnesota Statutes 2024, section 152.24, is amended to read:

152.24 FEDERALLY APPROVED CLINICAL TRIALS.

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The <u>commissioner office</u> may prohibit enrollment of a patient in the registry program if the patient is simultaneously enrolled in a federally approved clinical trial for the treatment of a qualifying medical condition with medical cannabis. The <u>commissioner office</u> shall provide information to all patients enrolled in the registry program on the existence of federally approved clinical trials for the treatment of the patient's qualifying medical condition with medical cannabis as an alternative to enrollment in the patient registry program.

Sec. 9. Minnesota Statutes 2024, section 152.25, is amended to read:

152.25 COMMISSIONER OFFICE DUTIES.

- Subdivision 1. **Medical cannabis manufacturer registration.** (a) The eommissioner office shall register two in-state manufacturers for the production of all medical cannabis within the state. A registration agreement between the eommissioner office and a manufacturer is nontransferable. The eommissioner office shall register new manufacturers or reregister the existing manufacturers by December 1 every two years, using the factors described in this subdivision. The eommissioner office shall accept applications after December 1, 2014, if one of the manufacturers registered before December 1, 2014, ceases to be registered as a manufacturer. The eommissioner's office's determination that no manufacturer exists to fulfill the duties under sections 152.22 to 152.37 is subject to judicial review in Ramsey County District Court. Data submitted during the application process are private data on individuals or nonpublic data as defined in section 13.02 until the manufacturer is registered under this section. Data on a manufacturer that is registered are public data, unless the data are trade secret or security information under section 13.37.
 - (b) As a condition for registration, a manufacturer must agree to:
- 7.25 (1) begin supplying medical cannabis to patients by July 1, 2015; and
- 7.26 (2) comply with all requirements under sections 152.22 to 152.37.
 - (c) The <u>commissioner office</u> shall consider the following factors when determining which manufacturer to register:
 - (1) the technical expertise of the manufacturer in cultivating medical cannabis and converting the medical cannabis into an acceptable delivery method under section 152.22, subdivision 6;
 - (2) the qualifications of the manufacturer's employees;

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- (3) the long-term financial stability of the manufacturer;
- (4) the ability to provide appropriate security measures on the premises of the manufacturer;
- (5) whether the manufacturer has demonstrated an ability to meet the medical cannabis production needs required by sections 152.22 to 152.37; and
- (6) the manufacturer's projection and ongoing assessment of fees on patients with a qualifying medical condition.
- (d) If an officer, director, or controlling person of the manufacturer pleads or is found guilty of intentionally diverting medical cannabis to a person other than allowed by law under section 152.33, subdivision 1, the <u>commissioner office</u> may decide not to renew the registration of the manufacturer, provided the violation occurred while the person was an officer, director, or controlling person of the manufacturer.
- (e) The commissioner office shall require each medical cannabis manufacturer to contract with an independent laboratory to test medical cannabis produced by the manufacturer. The commissioner office shall approve the laboratory chosen by each manufacturer and require that the laboratory report testing results to the manufacturer in a manner determined by the commissioner office.
- registration. If the commissioner office intends to revoke or not renew a registration issued under this section, the commissioner office must first notify in writing the manufacturer against whom the action is to be taken and provide the manufacturer with an opportunity to request a hearing under the contested case provisions of chapter 14. If the manufacturer does not request a hearing by notifying the commissioner office in writing within 20 days after receipt of the notice of proposed action, the commissioner office may proceed with the action without a hearing. For revocations, the registration of a manufacturer is considered revoked on the date specified in the commissioner's office's written notice of revocation.
- Subd. 1b. **Temporary suspension proceedings.** The <u>eommissioner office</u> may institute proceedings to temporarily suspend the registration of a medical cannabis manufacturer for a period of up to 90 days by notifying the manufacturer in writing if any action by an employee, agent, officer, director, or controlling person of the manufacturer:
- (1) violates any of the requirements of sections 152.22 to 152.37 or the rules adopted thereunder;

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(2) permits, aids, or abets the commission of any violation of state law at the manufacturer's location for cultivation, harvesting, manufacturing, packaging, and processing or at any site for distribution of medical cannabis;

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- (3) performs any act contrary to the welfare of a registered patient or registered designated caregiver; or
 - (4) obtains, or attempts to obtain, a registration by fraudulent means or misrepresentation.
- Subd. 1c. Notice to patients. Upon the revocation or nonrenewal of a manufacturer's registration under subdivision 1a or implementation of an enforcement action under subdivision 1b that may affect the ability of a registered patient, registered designated caregiver, or a registered patient's parent, legal guardian, or spouse to obtain medical cannabis from the manufacturer subject to the enforcement action, the eommissioner office shall notify in writing each registered patient and the patient's registered designated caregiver or registered patient's parent, legal guardian, or spouse about the outcome of the proceeding and information regarding alternative registered manufacturers. This notice must be provided two or more business days prior to the effective date of the revocation, nonrenewal, or other enforcement action.
- Subd. 2. Range of compounds and dosages; report. The office shall review and publicly report the existing medical and scientific literature regarding the range of recommended dosages for each qualifying condition and the range of chemical compositions of any plant of the genus cannabis that will likely be medically beneficial for each of the qualifying medical conditions. The office shall make this information available to patients with qualifying medical conditions beginning December 1, 2014, and update the information every three years. The office may consult with the independent laboratory under contract with the manufacturer or other experts in reporting the range of recommended dosages for each qualifying medical condition, the range of chemical compositions that will likely be medically beneficial, and any risks of noncannabis drug interactions. The office shall consult with each manufacturer on an annual basis on medical cannabis offered by the manufacturer. The list of medical cannabis offered by a manufacturer shall be published on the Office of Cannabis Management website.
- Subd. 3. Deadlines. The commissioner office shall adopt rules necessary for the manufacturer to begin distribution of medical cannabis to patients under the registry program by July 1, 2015, and have notice of proposed rules published in the State Register prior to January 1, 2015.

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Subd. 4. Reports. (a) The commissioner office shall provide regular updates to the task force on medical cannabis therapeutic research and; to the chairs and ranking minority members of the legislative committees with jurisdiction over health and human services, public safety, judiciary, and civil law; and to the Cannabis Advisory Council under section 342.03 regarding: (1) any changes in federal law or regulatory restrictions regarding the use of medical cannabis or hemp; and (2) the market demand and supply in this state for products made from hemp that can be used for medicinal purposes.

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- (b) The commissioner office may submit medical research based on the data collected under sections 152.22 to 152.37 to any federal agency with regulatory or enforcement authority over medical cannabis to demonstrate the effectiveness of medical cannabis for treating a qualifying medical condition.
- Sec. 10. Minnesota Statutes 2024, section 152.26, is amended to read:

152.26 RULEMAKING.

- (a) The commissioner office may adopt rules to implement sections 152.22 to 152.37. 10.14 Rules for which notice is published in the State Register before January 1, 2015, may be 10.15
- adopted using the process in section 14.389. 10.16
- 10.17 (b) The commissioner office may adopt or amend rules, using the procedure in section 14.386, paragraph (a), to implement the addition of dried raw cannabis as an allowable form 10.18 of medical cannabis under section 152.22, subdivision 6, paragraph (a), clause (4). Section 10.19 14.386, paragraph (b), does not apply to these rules. 10.20
- Sec. 11. Minnesota Statutes 2024, section 152.261, is amended to read: 10.21

152.261 RULES; ADVERSE INCIDENTS.

- (a) The commissioner of health office shall adopt rules to establish requirements for reporting incidents when individuals who are not authorized to possess medical cannabis under sections 152.22 to 152.37 are found in possession of medical cannabis. The rules must identify professionals required to report, the information they are required to report, and actions the reporter must take to secure the medical cannabis.
- (b) The commissioner of health office shall adopt rules to establish requirements for law enforcement officials and health care professionals to report incidents involving an overdose of medical cannabis to the commissioner of health office.
- (c) Rules must include the method by which the commissioner office will collect and tabulate reports of unauthorized possession and overdose.

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Sec. 12. Minnesota Statutes 2024, section 152.27, subdivision 2, is amended to read:

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- Subd. 2. **Office duties.** (a) The office shall:
- (1) give notice of the program to health care practitioners in the state who are eligible to serve as health care practitioners and explain the purposes and requirements of the program;
- (2) allow each health care practitioner who meets or agrees to meet the program's requirements and who requests to participate, to be included in the registry program to collect data for the patient registry;
- (3) provide explanatory information and assistance to each health care practitioner in understanding the nature of therapeutic use of medical cannabis within program requirements;
- (4) create and provide a certification to be used by a health care practitioner for the practitioner to certify whether a patient has been diagnosed with a qualifying medical condition;
- (5) supervise the participation of the health care practitioner in conducting patient treatment and health records reporting in a manner that ensures stringent security and record-keeping requirements and that prevents the unauthorized release of private data on individuals as defined by section 13.02;
- (6) develop safety criteria for patients with a qualifying medical condition as a requirement of the patient's participation in the program, to prevent the patient from undertaking any task under the influence of medical cannabis that would constitute negligence or professional malpractice on the part of the patient; and
- (7) conduct research and studies based on data from health records submitted to the registry program and submit reports on intermediate or final research results to the legislature and major scientific journals. The office may contract with a third party to complete the requirements of this clause. Any reports submitted must comply with section 152.28, subdivision 2.
- (b) The office may add a delivery method under section 152.22, subdivision 6, upon a petition from a member of the public or the Cannabis Advisory Council under section 342.03 or as directed by law. If the office wishes to add a delivery method under section 152.22, subdivision 6, the office must notify the chairs and ranking minority members of the legislative policy committees having jurisdiction over health and public safety of the addition and the reasons for its addition, including any written comments received by the office from the public and any guidance received from the Cannabis Advisory Council under section

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342.03, by January 15 of the year in which the office wishes to make the change. The change shall be effective on August 1 of that year, unless the legislature by law provides otherwise.

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- Sec. 13. Minnesota Statutes 2024, section 152.27, subdivision 7, is amended to read: 12.3
 - Subd. 7. Notice requirements. Patients and registered designated caregivers shall notify the commissioner office of any address or name change within 30 days of the change having occurred. A patient or registered designated caregiver is subject to a \$100 fine for failure to notify the commissioner office of the change.
- Sec. 14. Minnesota Statutes 2024, section 152.28, subdivision 1, is amended to read: 12.8
- Subdivision 1. Health care practitioner duties. (a) Prior to a patient's enrollment in 12.9 the registry program, a health care practitioner shall: 12.10
 - (1) determine, in the health care practitioner's medical judgment, whether a patient suffers from a qualifying medical condition, and, if so determined, provide the patient with a certification of that diagnosis;
 - (2) advise patients, registered designated caregivers, and parents, legal guardians, or spouses who are acting as caregivers of the existence of any nonprofit patient support groups or organizations;
 - (3) provide explanatory information from the office to patients with qualifying medical conditions, including disclosure to all patients about the experimental nature of therapeutic use of medical cannabis; the possible risks, benefits, and side effects of the proposed treatment; the application and other materials from the office; and provide patients with the Tennessen warning as required by section 13.04, subdivision 2; and
 - (4) agree to continue treatment of the patient's qualifying medical condition and report medical findings to the office.
- (b) Upon notification from the office of the patient's enrollment in the registry program, 12.24 the health care practitioner shall: 12.25
- (1) participate in the patient registry reporting system under the guidance and supervision 12.26 of the office; 12.27
- (2) report health records of the patient throughout the ongoing treatment of the patient 12.28 to the office in a manner determined by the commissioner office and in accordance with 12.29 subdivision 2; 12.30

Sec. 14. 12 (3) determine, every three years, if the patient continues to suffer from a qualifying medical condition and, if so, issue the patient a new certification of that diagnosis; and (4) otherwise comply with all requirements developed by the office.

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- (c) A health care practitioner may utilize telehealth, as defined in section 62A.673, subdivision 2, for certifications and recertifications.
- (d) Nothing in this section requires a health care practitioner to participate in the registry program.
- Sec. 15. Minnesota Statutes 2024, section 152.28, subdivision 3, is amended to read:
- Subd. 3. **Advertising restrictions.** (a) A health care practitioner shall not publish or cause to be published any advertisement that:
- 13.11 (1) contains false or misleading statements about medical cannabis or about the medical cannabis registry program;
- 13.13 (2) uses colloquial terms to refer to medical cannabis, such as pot, weed, or grass;
- 13.14 (3) states or implies the health care practitioner is endorsed by the Department of Health
 13.15 office or by the medical cannabis registry program;
- 13.16 (4) includes images of cannabis in its plant or leaf form or of cannabis-smoking
 13.17 paraphernalia; or
 - (5) contains medical symbols that could reasonably be confused with symbols of established medical associations or groups.
 - (b) A health care practitioner found by the <u>commissioner office</u> to have violated this subdivision is prohibited from certifying that patients have a qualifying medical condition for purposes of patient participation in the registry program. The <u>commissioner's office's</u> decision that a health care practitioner has violated this subdivision is a final decision of the <u>commissioner</u> office and is not subject to the contested case procedures in chapter 14.
 - Sec. 16. Minnesota Statutes 2024, section 152.29, subdivision 1, is amended to read:
 - Subdivision 1. **Manufacturer; requirements.** (a) A manufacturer may operate eight distribution facilities, which may include the manufacturer's single location for cultivation, harvesting, manufacturing, packaging, and processing but is not required to include that location. The eommissioner office shall designate the geographical service areas to be served by each manufacturer based on geographical need throughout the state to improve patient access. A manufacturer shall not have more than two distribution facilities in each

Sec. 16.

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geographical service area assigned to the manufacturer by the commissioner office. A manufacturer shall operate only one location where all cultivation, harvesting, manufacturing, packaging, and processing of medical cannabis shall be conducted. This location may be one of the manufacturer's distribution facility sites. The additional distribution facilities may dispense medical cannabis and medical cannabis products but may not contain any medical cannabis in a form other than those forms allowed under section 152.22, subdivision 6, and the manufacturer shall not conduct any cultivation, harvesting, manufacturing, packaging, or processing at the other distribution facility sites. Any distribution facility operated by the manufacturer is subject to all of the requirements applying to the manufacturer under sections 152.22 to 152.37, including, but not limited to, security and distribution requirements.

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- (b) A manufacturer may acquire hemp grown in this state from a hemp grower, and may acquire hemp products produced by a hemp processor. A manufacturer may manufacture or process hemp and hemp products into an allowable form of medical cannabis under section 152.22, subdivision 6. Hemp and hemp products acquired by a manufacturer under this paragraph are subject to the same quality control program, security and testing requirements, and other requirements that apply to medical cannabis under sections 152.22 to 152.37 and Minnesota Rules, chapter 4770.
- (c) A medical cannabis manufacturer shall contract with a laboratory approved by the commissioner office, subject to any additional requirements set by the commissioner office, for purposes of testing medical cannabis manufactured or hemp or hemp products acquired by the medical cannabis manufacturer as to content, contamination, and consistency to verify the medical cannabis meets the requirements of section 152.22, subdivision 6. The cost of laboratory testing shall be paid by the manufacturer.
 - (d) The operating documents of a manufacturer must include:
- (1) procedures for the oversight of the manufacturer and procedures to ensure accurate record keeping;
- (2) procedures for the implementation of appropriate security measures to deter and prevent the theft of medical cannabis and unauthorized entrance into areas containing medical cannabis; and
- (3) procedures for the delivery and transportation of hemp between hemp growers and manufacturers and for the delivery and transportation of hemp products between hemp processors and manufacturers.

Sec. 16. 14

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detection systems, and a personnel identification system.

(f) A manufacturer shall not share office space with, refer patients to a health care practitioner, or have any financial relationship with a health care practitioner.

a fully operational security alarm system, facility access controls, perimeter intrusion

- (g) A manufacturer shall not permit any person to consume medical cannabis on the property of the manufacturer.
 - (h) A manufacturer is subject to reasonable inspection by the commissioner office.
- (i) For purposes of sections 152.22 to 152.37, a medical cannabis manufacturer is not subject to the Board of Pharmacy licensure or regulatory requirements under chapter 151.
- (j) A medical cannabis manufacturer may not employ any person who is under 21 years of age or who has been convicted of a disqualifying felony offense. An employee of a medical cannabis manufacturer must submit a completed criminal history records check consent form, a full set of classifiable fingerprints, and the required fees for submission to the Bureau of Criminal Apprehension before an employee may begin working with the manufacturer. The bureau must conduct a Minnesota criminal history records check and the superintendent is authorized to exchange the fingerprints with the Federal Bureau of Investigation to obtain the applicant's national criminal history record information. The bureau shall return the results of the Minnesota and federal criminal history records checks to the commissioner office.
- (k) A manufacturer may not operate in any location, whether for distribution or cultivation, harvesting, manufacturing, packaging, or processing, within 1,000 feet of a public or private school existing before the date of the manufacturer's registration with the commissioner office.
- (l) A manufacturer shall comply with reasonable restrictions set by the commissioner office relating to signage, marketing, display, and advertising of medical cannabis.
- (m) Before a manufacturer acquires hemp from a hemp grower or hemp products from a hemp processor, the manufacturer must verify that the hemp grower or hemp processor has a valid license issued by the commissioner of agriculture under chapter 18K.
- (n) Until a state-centralized, seed-to-sale system is implemented that can track a specific medical cannabis plant from cultivation through testing and point of sale, the commissioner

Sec. 16. 15

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16.1	office shall conduct at least one unannounced inspection per year of each manufacturer that
16.2	includes inspection of:
16.3	(1) business operations;
16.4	(2) physical locations of the manufacturer's manufacturing facility and distribution
16.5	facilities;
16.6	(3) financial information and inventory documentation, including laboratory testing
16.7	results; and
16.8	(4) physical and electronic security alarm systems.
16.9	Sec. 17. Minnesota Statutes 2024, section 152.29, subdivision 2, is amended to read:
16.10	Subd. 2. Manufacturer; production. (a) A manufacturer of medical cannabis shall
16.11	provide a reliable and ongoing supply of all medical cannabis needed for the registry program
16.12	through cultivation by the manufacturer and through the purchase of hemp from hemp
16.13	growers.
16.14	(b) All cultivation, harvesting, manufacturing, packaging, and processing of medical
16.15	cannabis must take place in an enclosed, locked facility at a physical address provided to
16.16	the eommissioner office during the registration process.
16.17	(c) A manufacturer must process and prepare any medical cannabis plant material or
16.18	hemp plant material into a form allowable under section 152.22, subdivision 6, prior to
16.19	distribution of any medical cannabis.
16.20	Sec. 18. Minnesota Statutes 2024, section 152.29, subdivision 3a, is amended to read:
16.21	Subd. 3a. Transportation of medical cannabis; transport staffing. (a) A medical
16.22	cannabis manufacturer may staff a transport motor vehicle with only one employee if the
16.23	medical cannabis manufacturer is transporting medical cannabis to either a certified
16.24	laboratory for the purpose of testing or a facility for the purpose of disposal. If the medical

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16.28 (b) Notwithstanding paragraph (a), a medical cannabis manufacturer that is only transporting hemp for any purpose may staff the transport motor vehicle with only one 16.29 employee. 16.30

cannabis manufacturer is transporting medical cannabis for any other purpose or destination,

the transport motor vehicle must be staffed with a minimum of two employees as required

Sec. 18. 16

by rules adopted by the commissioner office.

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- (c) A medical cannabis manufacturer may contract with a third party for armored car services for deliveries of medical cannabis from its production facility to distribution facilities. A medical cannabis manufacturer that contracts for armored car services remains responsible for the transportation manifest and inventory tracking requirements in rules adopted by the <u>commissioner office</u>.
- (d) Department of Health Office staff may transport medical cannabis for the purposes of delivering medical cannabis and other samples to a laboratory for testing under rules adopted by the commissioner office and in cases of special investigations when the commissioner office has determined there is a potential threat to public health. The transport motor vehicle must be staffed with a minimum of two Department of Health office employees. The employees must carry with them their Department of Health office identification card and a transport manifest.
- 17.13 Sec. 19. Minnesota Statutes 2024, section 152.29, subdivision 4, is amended to read:
- Subd. 4. **Report.** (a) Each manufacturer shall report to the <u>commissioner office</u> on a monthly basis the following information on each individual patient for the month prior to the report:
- 17.17 (1) the amount and dosages of medical cannabis distributed;
- 17.18 (2) the chemical composition of the medical cannabis; and
- 17.19 (3) the tracking number assigned to any medical cannabis distributed.
- 17.20 (b) For transactions involving Tribal medical cannabis program patients, each
 17.21 manufacturer shall report to the <u>commissioner office</u> on a weekly basis the following
 17.22 information on each individual Tribal medical cannabis program patient for the week prior
 17.23 to the report:
- 17.24 (1) the name of the Tribal medical cannabis program in which the Tribal medical cannabis program patient is enrolled;
- 17.26 (2) the amount and dosages of medical cannabis distributed;
- 17.27 (3) the chemical composition of the medical cannabis distributed; and
- 17.28 (4) the tracking number assigned to the medical cannabis distributed.

Sec. 19. 17

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Sec. 20. Minnesota Statutes 2024, section 152.31, is amended to read:

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152.31 DATA PRACTICES.

- (a) Government data in patient files maintained by the commissioner office and the health care practitioner, and data submitted to or by a medical cannabis manufacturer, are private data on individuals, as defined in section 13.02, subdivision 12, or nonpublic data, as defined in section 13.02, subdivision 9, but may be used for purposes of complying with chapter 13 and complying with a request from the legislative auditor or the state auditor in the performance of official duties. The provisions of section 13.05, subdivision 11, apply to a registration agreement entered between the commissioner office and a medical cannabis manufacturer under section 152.25.
- (b) Not public data maintained by the commissioner office may not be used for any purpose not provided for in sections 152.22 to 152.37, and may not be combined or linked in any manner with any other list, dataset, or database.
- (c) The commissioner office may execute data sharing arrangements with the commissioner of agriculture to verify licensing, inspection, and compliance information related to hemp growers and hemp processors under chapter 18K.
- 18.17 Sec. 21. Minnesota Statutes 2024, section 152.32, subdivision 2, is amended to read:
 - Subd. 2. Criminal and civil protections. (a) Subject to section 152.23, the following are not violations under this chapter:
 - (1) use or possession of medical cannabis or medical cannabis products by a patient enrolled in the registry program; possession by a registered designated caregiver or the parent, legal guardian, or spouse of a patient if the parent, legal guardian, or spouse is listed on the registry verification; or use or possession of medical cannabis or medical cannabis products by a Tribal medical cannabis program patient;
 - (2) possession, dosage determination, or sale of medical cannabis or medical cannabis products by a medical cannabis manufacturer, employees of a manufacturer, a Tribal medical cannabis program manufacturer, employees of a Tribal medical cannabis program manufacturer, a laboratory conducting testing on medical cannabis, or employees of the laboratory; and
 - (3) possession of medical cannabis or medical cannabis products by any person while carrying out the duties required under sections 152.22 to 152.37.

Sec. 21. 18

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- (b) Medical cannabis obtained and distributed pursuant to sections 152.22 to 152.37 and associated property is not subject to forfeiture under sections 609.531 to 609.5316.
- (c) The commissioner office, members of a Tribal medical cannabis board, the eommissioner's office's or Tribal medical cannabis board's staff, the eommissioner's office's or Tribal medical cannabis board's agents or contractors, and any health care practitioner are not subject to any civil or disciplinary penalties by the Board of Medical Practice, the Board of Nursing, or by any business, occupational, or professional licensing board or entity, solely for participation in the registry program under sections 152.22 to 152.37 or in a Tribal medical cannabis program. A pharmacist licensed under chapter 151 is not subject to any civil or disciplinary penalties by the Board of Pharmacy when acting in accordance with the provisions of sections 152.22 to 152.37. Nothing in this section affects a professional licensing board from taking action in response to violations of any other section of law.
- (d) Notwithstanding any law to the contrary, the commissioner office, the governor of Minnesota, or an employee of any state agency may not be held civilly or criminally liable for any injury, loss of property, personal injury, or death caused by any act or omission while acting within the scope of office or employment under sections 152.22 to 152.37.
- (e) Federal, state, and local law enforcement authorities are prohibited from accessing the patient registry under sections 152.22 to 152.37 except when acting pursuant to a valid search warrant.
- (f) Notwithstanding any law to the contrary, neither the commissioner office nor a public employee may release data or information about an individual contained in any report, document, or registry created under sections 152.22 to 152.37 or any information obtained about a patient participating in the program, except as provided in sections 152.22 to 152.37.
- (g) No information contained in a report, document, or registry or obtained from a patient under sections 152.22 to 152.37 or from a Tribal medical cannabis program patient may be admitted as evidence in a criminal proceeding unless independently obtained or in connection with a proceeding involving a violation of sections 152.22 to 152.37.
- (h) Notwithstanding section 13.09, any person who violates paragraph (e) or (f) is guilty of a gross misdemeanor.
- (i) An attorney may not be subject to disciplinary action by the Minnesota Supreme Court, a Tribal court, or the professional responsibility board for providing legal assistance to prospective or registered manufacturers or others related to activity that is no longer subject to criminal penalties under state law pursuant to sections 152.22 to 152.37, or for

Sec. 21. 19

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providing legal assistance to a Tribal medical cannabis program or a Tribal medical cannabis program manufacturer.

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- (j) The following do not constitute probable cause or reasonable suspicion, and shall not be used to support a search of the person or property of the person possessing or applying for the registry verification or equivalent, or otherwise subject the person or property of the person to inspection by any governmental agency:
- (1) possession of a registry verification or application for enrollment in the registry program by a person entitled to possess a registry verification or apply for enrollment in the registry program; or
- (2) possession of a verification or equivalent issued by a Tribal medical cannabis program or application for enrollment in a Tribal medical cannabis program by a person entitled to possess such a verification or application.
- Sec. 22. Minnesota Statutes 2024, section 152.33, subdivision 1a, is amended to read:
- Subd. 1a. **Intentional diversion outside the state; penalties.** (a) In addition to any other applicable penalty in law, the <u>commissioner office</u> may levy a fine of \$250,000 against a manufacturer and may immediately initiate proceedings to revoke the manufacturer's registration, using the procedure in section 152.25, if:
 - (1) an officer, director, or controlling person of the manufacturer pleads or is found guilty under subdivision 1 of intentionally transferring medical cannabis, while the person was an officer, director, or controlling person of the manufacturer, to a person other than allowed by law; and
 - (2) in intentionally transferring medical cannabis to a person other than allowed by law, the officer, director, or controlling person transported or directed the transport of medical cannabis outside of Minnesota.
- 20.25 (b) All fines collected under this subdivision shall be deposited in the state government special revenue fund.
- Sec. 23. Minnesota Statutes 2024, section 152.33, subdivision 4, is amended to read:
 - Subd. 4. **Submission of false records; criminal penalty.** A person who knowingly submits false records or documentation required by the <u>commissioner office</u> to register as a manufacturer of medical cannabis under sections 152.22 to 152.37 is guilty of a felony and may be sentenced to imprisonment for not more than two years or by payment of a fine of not more than \$3,000, or both.

Sec. 23. 20

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Sec. 24. Minnesota Statutes 2024, section 152.35, is amended to read:

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152.35 FEES; DEPOSIT OF REVENUE.

- (a) The commissioner office shall collect an application fee of \$20,000 from each entity submitting an application for registration as a medical cannabis manufacturer. Revenue from the fee shall be deposited in the state treasury and credited to the state government special revenue fund.
- (b) The commissioner office shall establish and collect an annual fee from a medical cannabis manufacturer equal to the cost of regulating and inspecting the manufacturer in that year. Revenue from the fee amount shall be deposited in the state treasury and credited to the state government special revenue fund.
- (c) A medical cannabis manufacturer may charge patients enrolled in the registry program a reasonable fee for costs associated with the operations of the manufacturer. The manufacturer may establish a sliding scale of patient fees based upon a patient's household income and may accept private donations to reduce patient fees.
- Sec. 25. Minnesota Statutes 2024, section 152.37, is amended to read:

152.37 FINANCIAL EXAMINATIONS; PRICING REVIEWS.

- Subdivision 1. Financial records. A medical cannabis manufacturer shall maintain detailed financial records in a manner and format approved by the commissioner office, and shall keep all records updated and accessible to the commissioner office when requested.
- Subd. 2. Certified annual audit. A medical cannabis manufacturer shall submit the results of an annual certified financial audit to the commissioner office no later than May 1 of each year for the calendar year beginning January 2015. The annual audit shall be conducted by an independent certified public accountant and the costs of the audit are the responsibility of the medical cannabis manufacturer. Results of the audit shall be provided to the medical cannabis manufacturer and the commissioner office. The commissioner office may also require another audit of the medical cannabis manufacturer by a certified public accountant chosen by the commissioner office with the costs of the audit paid by the medical cannabis manufacturer.
- Subd. 3. **Power to examine.** (a) The commissioner office or designee may examine the 21.29 business affairs and conditions of any medical cannabis manufacturer, including but not 21.30 limited to a review of the financing, budgets, revenues, sales, and pricing. 21.31

Sec. 25. 21

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22.1	(b) An ex	amination may cove	r the medical ca	nnabis manufacture	r's business affairs,
22.2	practices, and	d conditions includin	g but not limited	d to a review of the	financing, budgets,
22.3	revenues, sal	es, and pricing. The	commissioner of	ffice shall determine	e the nature and scope
22.4	of each exam	ination and in doing	so shall take int	to account all availa	able relevant factors
22.5	concerning th	ne financial and busi	ness affairs, prac	ctices, and condition	ns of the examinee.
22.6	The costs inc	urred by the departm	nent in conducting	ng an examination sl	hall be paid for by the
22.7	medical cann	abis manufacturer.			
22.8	(c) When	making an examinat	ion under this se	ction, the commissi	oner office may retain
22.9	attorneys, ap	praisers, independen	t economists, inc	dependent certified p	public accountants, or
22.10	other profess	ionals and specialist	s as designees. A	A certified public ac	ecountant retained by
22.11	the commissi	ioner office may not	be the same cert	tified public accoun	tant providing the
22.12	certified annu	ual audit in subdivisi	on 2.		
22.13	(d) The ea	ommissioner office sl	hall make a repo	rt of an examination	conducted under this
22.14	section and p	rovide a copy to the	medical cannabi	s manufacturer. The	e commissioner office
22.15	shall then pos	st a copy of the report	on the departme	nt's website. All wor	rking papers, recorded
22.16	information,	documents, and copi	ies produced by,	obtained by, or disc	closed to the
22.17	commissione	er office or any other	person in the co	ourse of an examina	tion, other than the
22.18	information of	contained in any con	nmissioner office	e official report, ma	de under this section
22.19	are private da	ata on individuals or	nonpublic data,	as defined in section	on 13.02.

- Sec. 26. Minnesota Statutes 2024, section 342.01, subdivision 9, is amended to read: 22.20
- 22.21 Subd. 9. Bona fide labor organization. "Bona fide labor organization" means a labor union that represents or is actively seeking to represent eannabis workers. of: 22.22
- 22.23 (1) a cannabis business; or
- (2) a lower-potency hemp edible manufacturer. 22.24
- Sec. 27. Minnesota Statutes 2024, section 342.01, subdivision 47, is amended to read: 22.25
 - Subd. 47. Labor peace agreement. "Labor peace agreement" means an agreement between a cannabis business and a bona fide labor organization or an agreement between a lower-potency hemp edible manufacturer and a bona fide labor organization that protects the state's interests by, at minimum, prohibiting the labor organization from engaging in picketing, work stoppages, or boycotts against the cannabis business or lower-potency hemp edible manufacturer.

Sec. 27. 22 BD

Sec. 28. Minnesota Statutes 2024, section 342.01, subdivision 50, is amended to read: 23.1 Subd. 50. Lower-potency hemp edible. (a) "Lower-potency hemp edible" means any 23.2 product that: 23.3 (1) is intended to be eaten or consumed as a beverage by humans; 23.4 (2) contains hemp concentrate or an artificially derived cannabinoid, in combination 23.5 with food ingredients; 23.6 23.7 (3) is not a drug; (4) does not contain a cannabinoid derived from cannabis plants or cannabis flower; 23.8 (5) is a type of product approved for sale by the office or is substantially similar to a 23.9 product approved by the office, including but not limited to products that resemble 23.10 nonalcoholic beverages, candy, and baked goods; and 23.11 (6) meets either of the requirements in paragraph (b). 23.12 (b) A lower-potency hemp edible includes: 23.13 (1) a product that: 23.14 (i) consists of servings that contain no more than five milligrams of delta-9 23.15 tetrahydrocannabinol; no more than 25 milligrams of cannabidiol, cannabigerol, cannabinol, 23.16 or cannabichromene; any other cannabinoid authorized by the office; or any combination 23.17 of those cannabinoids that does not exceed the identified amounts, except that a 23.18 lower-potency hemp edible that is intended to be consumed as a beverage may contain no 23.19 more than ten milligrams of delta-9 tetrahydrocannabinol in a single-serving container; 23.20 (ii) does not contain more than a combined total of 0.5 milligrams of all other 23.21 cannabinoids per serving; and 23.22 (iii) does not contain an artificially derived cannabinoid other than delta-9 23.23 tetrahydrocannabinol, except that a product may include artificially derived cannabinoids 23.24 created during the process of creating the delta-9 tetrahydrocannabinol that is added to the 23.25 product, if no artificially derived cannabinoid is added to the ingredient containing delta-9 23.26 tetrahydrocannabinol and the ratio of delta-9 tetrahydrocannabinol to all other artificially 23.27 derived cannabinoids is no less than 20 to one; or 23.28 (2) a product that: 23.29 (i) contains hemp concentrate processed or refined without increasing the percentage of 23.30

targeted cannabinoids or altering the ratio of cannabinoids in the extracts or resins of a hemp

Sec. 28. 23

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of Minnesota. A valid registration verification card must verify that the card holder is

enrolled in or authorized to participate in a Tribal medical cannabis program.

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Sec. 32. 24

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Sec. 33. Minnesota Statutes 2024, section 342.01, subdivision 71, is amended to read:

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- Subd. 71. **Visiting patient.** "Visiting patient" means an individual who is not a Minnesota resident and who possesses a valid registration verification card or its equivalent that is issued under the laws or regulations of another state, district, commonwealth, or territory of the United States verifying that the individual is enrolled in or authorized to participate in that jurisdiction's medical cannabis or medical marijuana program or in a Tribal medical cannabis program.
- Sec. 34. Minnesota Statutes 2024, section 342.02, subdivision 3, is amended to read: 25.8
- Subd. 3. Medical cannabis program. (a) The powers and duties of the Department of 25.9 Health with respect to the medical cannabis program under Minnesota Statutes 2022, sections 25.10 152.22 to 152.37, are transferred to the Office of Cannabis Management under section 25.11 15.039. 25.12
 - (b) The following protections shall apply to employees who are transferred from the Department of Health to the Office of Cannabis Management:
- (1) the employment status and job classification of a transferred employee shall not be 25.15 altered as a result of the transfer; 25.16
 - (2) transferred employees who were represented by an exclusive representative prior to the transfer shall continue to be represented by the same exclusive representative after the transfer;
 - (3) the applicable collective bargaining agreements with exclusive representatives shall continue in full force and effect for such transferred employees after the transfer;
 - (4) the state must meet and negotiate with the exclusive representatives of the transferred employees about any proposed changes affecting or relating to the transferred employees' terms and conditions of employment to the extent such changes are not addressed in the applicable collective bargaining agreement; and
 - (5) for an employee in a temporary unclassified position transferred to the Office of Cannabis Management, the total length of time that the employee has served in the appointment shall include all time served in the appointment and the transferring agency and the time served in the appointment at the Office of Cannabis Management. An employee in a temporary unclassified position who was hired by a transferring agency through an open competitive selection process in accordance with a policy enacted by Minnesota Management and Budget shall be considered to have been hired through such process after the transfer.

Sec. 34. 25

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- Sec. 35. Minnesota Statutes 2024, section 342.09, subdivision 2, is amended to read:
- Subd. 2. **Home cultivation of cannabis for personal adult use.** (a) Up to eight cannabis plants, with no more than four being mature, flowering plants may be grown at a single residence, including the curtilage or yard, without a license to cultivate cannabis issued under this chapter provided that cultivation takes place at the primary residence of an individual 21 years of age or older and in an enclosed, locked space that is not open to public view.
- (b) Pursuant to section 342.52, subdivision 9, paragraph (d), a registered designated caregiver may cultivate up to eight cannabis plants for not more than one patient household.

 In addition to eight cannabis plants for one patient household, a registered designated caregiver may cultivate up to eight cannabis plants for the caregiver's personal adult use of cannabis. Of the 16 or fewer total cannabis plants being grown in the registered caregiver's residence, no more than eight may be mature, flowering plants.
- Sec. 36. Minnesota Statutes 2024, section 342.12, is amended to read:

342.12 LICENSES; TRANSFERS; ADJUSTMENTS.

- 26.17 (a) Licenses issued under this chapter that are available to all applicants pursuant to
 26.18 section 342.14, subdivision 1b, paragraph (c), may be freely transferred subject to the prior
 26.19 written approval of the office unless the license holder has not received a final site inspection
 26.20 or the license holder is a social equity applicant.
 - (b) Licenses issued as social equity licenses pursuant to either section 342.14, subdivision 1b, paragraph (b), or section 342.175, paragraph (b), may only be transferred to another social equity applicant for three years after the date on which the office issues the license. Three years after the date of issuance, a license holder may transfer a license to any entity. Transfer of a license that was issued as a social equity license must be reviewed by the
- 26.26 Division of Social Equity and is subject to the prior written approval of the office.
- 26.27 (c) <u>Preliminary license preapproval approval issued pursuant to section 342.125 342.14,</u>
 26.28 <u>subdivision 5, may not be transferred.</u>
- 26.29 (d) A new license must be obtained when:
- 26.30 (1) the form of the licensee's legal business structure converts or changes to a different type of legal business structure; or

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(2) the licensee dissolves; consolidates; reorganizes; undergoes bankruptcy, insolvency, or receivership proceedings; merges with another legal organization; or assigns all or substantially all of its assets for the benefit of creditors.

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(e) Licenses must be renewed annually.

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- (f) License holders may petition the office to adjust the tier of a license issued within a license category if the license holder meets all applicable requirements.
- (g) The office by rule may permit the relocation of a licensed cannabis business; permit the relocation of an approved operational location, including a cultivation, manufacturing, processing, or retail location; adopt requirements for the submission of a license relocation application; establish standards for the approval of a relocation application; and charge a fee not to exceed \$250 for reviewing and processing applications. Relocation of a licensed premises pursuant to this paragraph does not extend or otherwise modify the license term of the license subject to relocation.
- Sec. 37. Minnesota Statutes 2024, section 342.14, subdivision 1, is amended to read:
- Subdivision 1. **Application; contents.** (a) The office shall establish procedures for the processing of cannabis licenses issued under this chapter. At a minimum, any application to obtain or renew a cannabis license shall include the following information, if applicable:
 - (1) the name, address, and date of birth of the applicant;
- 27.19 (2) the disclosure of ownership and control required under paragraph (b);
- 27.20 (3) the disclosure of whether the applicant or, if the applicant is a business, any officer, 27.21 director, manager, and general partner of the business has ever filed for bankruptcy;
- 27.22 (4) the address and legal property description of the business, if applicable, except an applicant is not required to secure a physical premises for the business at the time of application;
- 27.25 (5) a general description of the location or locations that the applicant plans to operate, 27.26 including the planned square feet of space for cultivation, wholesaling, and retailing, as 27.27 applicable;
- 27.28 (6) a copy of the security plan, including security monitoring, security equipment, and facility maps if applicable, except an applicant is not required to secure a physical premises for the business at the time of application;
 - (7) proof of trade name registration;

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(8) a copy of the applicant's business plan showing the expected size of the business;
anticipated growth; the methods of record keeping; the knowledge and experience of the
applicant and any officer, director, manager, and general partner of the business; the
environmental plan; and other relevant financial and operational components;
(9) standard operating procedures for:
(i) quality assurance;
(ii) inventory control, storage, and diversion prevention; and
(iii) accounting and tax compliance;
(10) an attestation signed by a bona fide labor organization stating that the applicant has entered into a labor peace agreement;
(11) a description of any training and education that the applicant will provide to employees of the business;
(12) a disclosure of any violation of a license agreement or a federal, state, or local law or regulation committed by the applicant or any true party of interest in the applicant's business that is relevant to business and working conditions;
(13) certification that the applicant will comply with the requirements of this chapter;
(14) identification of one or more controlling persons or managerial employees as agents who shall be responsible for dealing with the office on all matters;
(15) a statement that the applicant agrees to respond to the office's supplemental requests for information; and
(16) a release of information for the applicant and every true party of interest in the applicant's business license for the office to perform the background checks required under section 342.15-;
(17) proof that the applicant is a social equity applicant; and
(18) an attestation that the applicant's business policies governing business operations
comply with this chapter.
(b) An applicant must file and update as necessary a disclosure of ownership and control identifying any true party of interest as defined in section 342.185, subdivision 1, paragraph (g). The office shall establish the contents of the disclosure. Except as provided in paragraph
(f) (d), the disclosure shall, at a minimum, include the following:

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29.1	(1) the management structure, ownership, and control of the applicant or license holder,
29.2	including the name of each cooperative member, officer, director, manager, general partner,
29.3	or business entity; the office or position held by each person; each person's percentage
29.4	ownership interest, if any; and, if the business has a parent company, the name of each
29.5	owner, board member, and officer of the parent company and the owner's, board member's,
29.6	or officer's percentage ownership interest in the parent company and the cannabis business;
29.7	(2) a statement from the applicant and, if the applicant is a business, from every officer,
29.8	director, manager, and general partner of the business, indicating whether that person has
29.9	previously held, or currently holds, an ownership interest in a cannabis business in Minnesota,
29.10	any other state or territory of the United States, or any other country;
29.11	(3) if the applicant is a corporation, copies of the applicant's articles of incorporation
29.12	and bylaws and any amendments to the applicant's articles of incorporation or bylaws;
29.13	(4) copies of any partnership agreement, operating agreement, or shareholder agreement;
29.14	(5) copies of any promissory notes, security instruments, or other similar agreements;
29.15	(6) an explanation detailing the funding sources used to finance the business;
29.16	(7) a list of operating and investment accounts for the business, including any applicable
29.17	financial institution and account number; and
29.18	(8) a list of each outstanding loan and financial obligation obtained for use in the business,
29.19	including the loan amount, loan terms, and name and address of the creditor.
29.20	(c) An application may include:
29.21	(1) proof that the applicant is a social equity applicant;
29.22	(2) a description of the training and education that will be provided to any employee;
29.23	Of
29.24	(3) a copy of business policies governing operations to ensure compliance with this
29.25	chapter.
29.26	(d) (c) Commitments made by an applicant in its application, including but not limited
29.27	to the maintenance of a labor peace agreement, shall be an ongoing material condition of
29.28	maintaining and renewing the license.

(e) An application on behalf of a corporation or association shall be signed by at least

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two officers or managing agents of that entity.

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(f) (d) The office may establish exceptions to the disclosures required under paragraph 30.1 (b) for members of a cooperative who hold less than a five percent ownership interest in 30.2 30.3 the cooperative. Sec. 38. Minnesota Statutes 2024, section 342.14, subdivision 3, is amended to read: 30.4 Subd. 3. Review. (a) After an applicant submits an application that contains all required 30.5 information and pays the applicable licensing application fee, the office must review the 30.6 application. 30.7 (b) The office may deny an application if: 30.8 (1) the application is incomplete; 30.9 (2) the application contains a materially false statement about the applicant or omits 30.10 information required under subdivision 1; 30.11 (3) the applicant does not meet the qualifications under section 342.16; 30.12 (4) the applicant is prohibited from holding the license under section 342.18, subdivision 30.13 2; 30.14 (5) the application does not meet the minimum requirements under section 342.18, 30.15 subdivision 3; 30.16 30.17 (6) the applicant fails to pay the applicable application fee; (7) the application was not submitted by the application deadline; 30.18 (8) the applicant submitted more than one application for a license type; or 30.19 (9) the office determines that the applicant would be prohibited from holding a license 30.20 for any other reason. 30.21 (c) If the office denies an application, the office must notify the applicant of the denial 30.22 and the basis for the denial. 30.23 (d) The office may request additional information from any applicant if the office 30.24 determines that the information is necessary to review or process the application. If the 30.25 applicant does not provide the additional requested information within 14 calendar days of 30.26 the office's request for information, the office may deny the application. 30.27 (e) An applicant whose application is not denied under this subdivision is a qualified 30.28

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applicant.

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Sec. 39. Minnesota Statutes 2024, section 342.14, subdivision 6, is amended to read: 31.1 Subd. 6. Completed application; final authorization; issuance of license. (a) Within 31.2 18 months of receiving notice of preliminary license approval, an applicant must provide: 31.3 (1) the address and legal property description of the location where the business will 31.4 31.5 operate; (2) the name of the local unit of government where the business will be located; and 31.6 31.7 (3) if applicable, an updated description of the location where the business will operate, an updated security plan, and any other additional information required by the office. 31.8 (b) Upon receipt of the information required under paragraph (a) from an applicant that 31.9 has received preliminary license approval, the office must: 31.10 (1) forward a copy of the application to the local unit of government in which the business 31.11 operates or intends to operate with a form for certification as to whether a proposed cannabis 31.12 business complies with local zoning ordinances and, if applicable, whether the proposed 31.13 business complies with the state fire code and building code; 31.14 (2) schedule a site inspection; and 31.15 (3) require the applicant to pay the applicable license fee. 31.16 (c) The office may deny final authorization if: 31.17 (1) an applicant fails to submit any required information; 31.18 (2) the applicant submits a materially false statement about the applicant or fails to 31.19 provide any required information; 31.20 31.21 (3) the office confirms that the cannabis business for which the office granted a preliminary license preapproval approval does not meet local zoning and land use laws; 31.22 31.23 (4) the applicant fails to pay the applicable license fee; or (5) the office determines that the applicant is disqualified from holding the license or 31.24 31.25 would operate in violation of the provisions of this chapter. (d) Within 90 days of receiving the information required under paragraph (a) and the 31.26 results of any required background check, the office shall grant final authorization and issue 31.27 the appropriate license or send the applicant a notice of rejection setting forth specific 31.28 reasons that the office did not approve the application. 31.29

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Sec. 40. Minnesota Statutes 2024, section 342.151, subdivision 2, is amended to read:

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Subd. 2. Criminal history check. A license holder cannabis business may employ or contract with as many unlicensed individuals as may be necessary, provided that the license holder cannabis business is at all times accountable for the good conduct of every individual employed by or contracted with the license holder cannabis business. Before hiring an individual as a cannabis worker, the license holder cannabis business must submit to the Bureau of Criminal Apprehension the individual's full set of fingerprints and written consent for the bureau to conduct a state and national criminal history check. The bureau may exchange an individual's fingerprints with the Federal Bureau of Investigation. The Bureau of Criminal Apprehension must determine whether the individual is qualified to be employed as a cannabis worker and must notify the license holder cannabis business of the bureau's determination. The license holder cannabis business must not employ an individual who is disqualified from being employed as a cannabis worker.

- 32.14 Sec. 41. Minnesota Statutes 2024, section 342.151, subdivision 3, is amended to read:
- Subd. 3. **Disqualification.** (a) A license holder cannabis business must not employ an 32.15 32.16 individual as a cannabis worker if the individual has been convicted of any of the following crimes that would constitute a felony: 32.17
- (1) human trafficking; 32.18
- (2) noncannabis controlled substance crimes in the first or second degree; 32.19
- (3) labor trafficking; 32.20
- (4) fraud; 32.21
- (5) embezzlement; 32.22
- (6) extortion; 32.23
- (7) money laundering; or 32.24
- (8) insider trading; 32.25
- if committed in this state or any other jurisdiction for which a full pardon or similar relief 32.26 has not been granted. 32.27
- (b) A license holder cannabis business must not employ an individual as a cannabis 32.28 worker if the individual made any false statement in an application for employment. 32.29

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33.1	Sec. 42. M	innesota Statutes 202	4, section 342.	22, subdivision 3, is a	mended to read:
33.2	Subd. 3.	Issuance of registrat	cion. (a) A loca	l unit of government	shall issue a retail
33.3	registration t	o a cannabis microbu	siness with a re	etail operations endor	sement, cannabis
33.4	mezzobusine	ess with a retail opera	tions endorsem	ent, cannabis retailer	, medical cannabis
33.5	combination	business operating a	retail location,	or lower-potency hem	p edible retailer that:
33.6	(1) has a	valid license or prelin	minary license 1	preapproval approval	issued by the office;
33.7	(2) has pa	aid the registration fe	e or renewal fe	e pursuant to subdivis	sion 2;
33.8	(3) is four	nd to be in complianc	e with the requi	irements of this chapte	er at any preliminary
33.9	compliance of	check that the local u	nit of governme	ent performs; and	
33.10	(4) if app	licable, is current on	all property tax	tes and assessments a	t the location where
33.11	the retail esta	ablishment is located.			
33.12	(b) Befor	e issuing a retail regi	stration, the loc	cal unit of governmen	t may conduct a
33.13	preliminary of	compliance check to	ensure that the	cannabis business or	hemp business is in
33.14	compliance v	with any applicable lo	ocal ordinance	established pursuant t	so section 342.13.
33.15	(c) A loca	al unit of government	shall renew th	e retail registration of	a cannabis business
33.16	or hemp bus	iness when the office	renews the lice	ense of the cannabis b	ousiness or hemp
33.17	business.				
33.18	(d) A reta	ail registration issued	under this sect	ion may not be transf	erred.
33.19	Sec. 43. M	innesota Statutes 202	4, section 342.	28, subdivision 1, is ε	mended to read:
33.20	Subdivisi	ion 1. Authorized ac	tions. A canna	bis microbusiness lice	ense, consistent with
33.21	the specific l	icense endorsement o	r endorsements	, entitles the license h	older to perform any
33.22	or all of the	following within the	imits establish	ed by this section:	
33.23	(1) grow	cannabis plants from	seed or immat	ure plant to mature pl	ant and harvest
33.24	cannabis flov	wer from a mature pla	ant;		
33.25	(2) make	cannabis concentrate	·••		
33.26	(3) make	hemp concentrate, in	cluding hemp	concentrate with a del	lta-9
33.27	tetrahydroca	nnabinol concentration	on of more than	0.3 percent as measu	ared by weight;
33.28	(4) manu	facture artificially de	rived cannabin	oids;	

(5) manufacture adult-use cannabis products, lower-potency hemp edibles, and

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hemp-derived consumer products for public consumption;

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(6) purchase immature cannabis plants and seedlings and, cannabis flower, cannabis
products, lower-potency hemp edibles, and hemp-derived consumer products from another
cannabis microbusiness, a cannabis mezzobusiness, <u>a cannabis cultivator</u> , a cannabis
manufacturer, or a cannabis wholesaler, or a lower-potency hemp edible manufacturer;
(7) purchase hemp plant parts and propagules from an industrial hemp grower licensed
under chapter 18K;
(8) purchase hemp concentrate from an industrial hemp processor licensed under chapter
18K;
(9) purchase cannabis concentrate, hemp concentrate, and artificially derived cannabinoids
from another cannabis microbusiness, a cannabis mezzobusiness, a cannabis manufacturer,
or a cannabis wholesaler for use in manufacturing adult-use cannabis products, lower-potency
hemp edibles, or hemp-derived consumer products;
(10) package and label adult-use cannabis flower, adult-use cannabis products,
lower-potency hemp edibles, and hemp-derived consumer products for sale to customers;
(11) sell immature cannabis plants and seedlings, adult-use cannabis flower, adult-use
cannabis products, lower-potency hemp edibles, hemp-derived consumer products, and
other products authorized by law to other cannabis businesses and to customers;
(12) operate an establishment that permits on-site consumption of edible cannabis
products and lower-potency hemp edibles; and
(13) perform other actions approved by the office.
(13) perioriii omer denons approved by the office.
Sec. 44. Minnesota Statutes 2024, section 342.28, subdivision 8, is amended to read:
Subd. 8. Production of customer consumer products endorsement. A cannabis
microbusiness that manufactures edible cannabis products, lower-potency hemp products,
or hemp-derived consumer products must comply with the requirements in section 342.26,
subdivisions 2 and 4.
Sec. 45. Minnesota Statutes 2024, section 342.29, subdivision 1, is amended to read:
Subdivision 1. Authorized actions. A cannabis mezzobusiness license, consistent with
the specific license endorsement or endorsements, entitles the license holder to perform any
or all of the following within the limits established by this section:

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35.1	(1) grow cannabis plants from seed or immature plant to mature plant and harvest
35.2	cannabis flower from a mature plant for use as adult-use cannabis flower or for use in
35.3	adult-use cannabis products;
35.4	(2) grow cannabis plants from seed or immature plant to mature plant and harvest
35.5	cannabis flower from a mature plant for use as medical cannabis flower or for use in medical
35.6	cannabinoid products;
35.7	(3) make cannabis concentrate;
35.8	(4) make hemp concentrate, including hemp concentrate with a delta-9
35.9	tetrahydrocannabinol concentration of more than 0.3 percent as measured by weight;
35.10	(5) manufacture artificially derived cannabinoids;
35.11	(6) manufacture adult-use cannabis products, lower-potency hemp edibles, and
35.12	hemp-derived consumer products for public consumption;
35.13	(7) process medical cannabinoid products;
35.14	(8) purchase immature cannabis plants and seedlings and, cannabis flower, cannabis
35.15	products, lower-potency hemp edibles, and hemp-derived consumer products from a cannabis
35.16	microbusiness, another cannabis mezzobusiness, a cannabis cultivator, a cannabis
35.17	manufacturer, or a cannabis wholesaler, or a lower-potency hemp edible manufacturer;
35.18	(9) purchase cannabis concentrate, hemp concentrate, and synthetically artificially derived
35.19	cannabinoids from a cannabis microbusiness, another cannabis mezzobusiness, a cannabis
35.20	manufacturer, or a cannabis wholesaler for use in manufacturing adult-use cannabis products,
35.21	lower-potency hemp edibles, or hemp-derived consumer products;
35.22	(10) purchase hemp plant parts and propagules from a licensed hemp grower licensed
35.23	under chapter 18K;
35.24	(11) purchase hemp concentrate from an industrial hemp processor licensed under chapter
35.25	18K;
35.26	(12) package and label adult-use cannabis flower, adult-use cannabis products,
35.27	lower-potency hemp edibles, and hemp-derived consumer products for sale to customers;
35.28	(13) sell immature cannabis plants and seedlings, adult-use cannabis flower, adult-use
35.29	cannabis products, lower-potency hemp edibles, hemp-derived consumer products, and
35.30	other products authorized by law to other cannabis businesses and to customers; and
35.31	(14) perform other actions approved by the office.

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36.1	Sec. 46. Minnesota Statutes 2024, section 342.29, subdivision 7, is amended to read:
36.2	Subd. 7. Production of customer consumer products endorsement. A cannabis
36.3	mezzobusiness that manufactures edible cannabis products, lower-potency hemp products,
36.4	or hemp-derived consumer products must comply with the requirements in section 342.26,
36.5	subdivisions 2 and 4.
36.6	Sec. 47. Minnesota Statutes 2024, section 342.30, subdivision 1, is amended to read:
36.7	Subdivision 1. Authorized actions. A cannabis cultivator license entitles the license
36.8	holder to:
36.9	(1) grow cannabis plants within the approved amount of space from seed or immature
36.10	plant to mature plant;
36.11	(2) harvest cannabis flower from a mature plant;
36.12	(3) package and label immature cannabis plants and seedlings and cannabis flower for
36.13	sale to other cannabis businesses;
36.14	(4) sell immature cannabis plants and seedlings and cannabis flower to other cannabis
36.15	businesses;
36.16	(5) transport cannabis flower to a cannabis manufacturer located on the same premises;
36.17	and
36.18	(6) perform other actions approved by the office.
36.19	Sec. 48. Minnesota Statutes 2024, section 342.32, subdivision 4, is amended to read:
36.20	Subd. 4. Multiple licenses; limits. (a) A person, cooperative, or business holding a
36.21	cannabis retailer license may also hold a cannabis delivery service license and a cannabis
36.22	event organizer license.
36.23	(b) Except as provided in paragraph (a) and subdivision 5, no person, cooperative, or
36.24	business holding a cannabis retailer license may own or operate any other cannabis business
36.25	or hemp business.
36.26	(c) No person, cooperative, or business may hold a license to own or operate more than
36.27	one cannabis retail business in one city and three retail businesses in one county.
36.28	(d) The office by rule may limit the number of cannabis retailer licenses a person,

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cooperative, or business may hold.

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37.1	(e) For purposes of this subdivision, a restriction on the number or type of license a
37.2	business may hold applies to every cooperative member or every director, manager, and
37.3	general partner of a cannabis business.
27.4	See 40 Minnesote Statutes 2024 section 242.22 subdivision 5 is amonded to made
37.4	Sec. 49. Minnesota Statutes 2024, section 342.32, subdivision 5, is amended to read:
37.5	Subd. 5. Municipal or county cannabis store. A city or county may establish, own,
37.6	and operate a municipal cannabis store subject to the restrictions in this chapter.
37.7	Notwithstanding any law to the contrary, a city or county that establishes, owns, or operates
37.8	a municipal cannabis store may also hold a lower-potency hemp edible retailer license.
37.9	Sec. 50. Minnesota Statutes 2024, section 342.33, subdivision 1, is amended to read:
37.10	Subdivision 1. Authorized actions. A cannabis wholesaler license entitles the license
37.11	holder to:
37.12	(1) purchase immature cannabis plants and seedlings, cannabis flower, cannabis products,
37.13	lower-potency hemp edibles, and hemp-derived consumer products from cannabis
37.14	microbusinesses, cannabis mezzobusinesses, cannabis cultivators, cannabis manufacturers,
37.15	and eannabis microbusinesses lower-potency hemp edible manufacturers;
37.16	(2) purchase hemp plant parts and propagules from industrial hemp growers licensed
37.17	under chapter 18K;
37.18	(3) purchase hemp concentrate from an industrial hemp processor licensed under chapter
37.19	18K;
37.20	(4) sell immature cannabis plants and seedlings, cannabis flower, cannabis products,
37.21	lower-potency hemp edibles, and hemp-derived consumer products to cannabis
37.22	microbusinesses, cannabis mezzobusinesses, cannabis manufacturers, and cannabis retailers;
37.23	(5) sell lower-potency hemp edibles to lower-potency hemp edible retailers;
37.24	(6) import hemp-derived consumer products and lower-potency hemp edibles that contain
37.25	hemp concentrate or artificially derived cannabinoids that are derived from hemp plants or
37.26	hemp plant parts; and
37.27	(7) perform other actions approved by the office.
37.28	Sec. 51. Minnesota Statutes 2024, section 342.40, subdivision 7, is amended to read:
37.29	Subd. 7. Cannabis event sales. (a) Cannabis microbusinesses with a retail endorsement,

cannabis mezzobusinesses with a retail endorsement, cannabis retailers, medical cannabis

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combination businesses operating a retail location, and lower-potency hemp edible retailers, including the cannabis event organizer, may be authorized to sell cannabis plants, adult-use cannabis flower, adult-use cannabis products, lower-potency hemp edibles, and hemp-derived consumer products to customers at a cannabis event.

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- (b) All sales of cannabis plants, adult-use cannabis flower, adult-use cannabis products, lower-potency hemp edibles, and hemp-derived consumer products at a cannabis event must take place in a retail area as designated in the premises diagram.
 - (c) Authorized retailers may only conduct sales within their specifically assigned area.
- (d) Authorized retailers must verify the age of all customers pursuant to section 342.27, subdivision 4, before completing a sale and may not sell cannabis plants, adult-use cannabis flower, adult-use cannabis products, lower-potency hemp edibles, or hemp-derived consumer products to an individual under 21 years of age.
- (e) Authorized retailers may display one sample of each type of cannabis plant, adult-use cannabis flower, adult-use cannabis product, lower-potency hemp edible, and hemp-derived consumer product available for sale. Samples of adult-use cannabis and adult-use cannabis products must be stored in a sample jar or display case and be accompanied by a label or notice containing the information required to be affixed to the packaging or container containing adult-use cannabis flower and adult-use cannabis products sold to customers. A sample may not consist of more than eight grams of adult-use cannabis flower or adult-use cannabis concentrate, or an edible cannabis product infused with more than 100 milligrams of tetrahydrocannabinol. A cannabis retailer may allow customers to smell the adult-use cannabis flower or adult-use cannabis product before purchase.
- (f) The notice requirements under section 342.27, subdivision 6, apply to authorized retailers offering cannabis plants, adult-use cannabis flower, adult-use cannabinoid products, and hemp-derived consumer products for sale at a cannabis event.
 - (g) Authorized retailers may not:
- (1) sell adult-use cannabis flower, adult-use cannabis products, lower-potency hemp edibles, or hemp-derived consumer products to a person who is visibly intoxicated;
- (2) knowingly sell more cannabis plants, adult-use cannabis flower, adult-use cannabis products, lower-potency hemp edibles, or hemp-derived consumer products than a customer is legally permitted to possess;
- (3) sell medical cannabis flower or medical cannabinoid products; or

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39.1	(4) give a	way cannabis plants,	, cannabis flower	;, cannabis products, l	lower-potency hemp
39.2	edibles, or he	emp-derived consum	er products; or		
39.3	(5) (4) alle	ow for the dispensin	g of cannabis pl	ants, cannabis flower	c, cannabis products,
39.4	lower-potenc	y hemp edibles, or h	nemp-derived co	nsumer products in v	ending machines.
39.5	(h) Excep	t for samples of a car	nnabis plant, adu	lt-use cannabis flowe	r, adult-use cannabis
39.6	product, lower	er-potency hemp edi	ble, and hemp-d	erived consumer pro	duct, all cannabis
39.7	plants, adult-	use cannabis flower,	adult-use cannab	ois products, lower-po	otency hemp edibles,
39.8	and hemp-den	rived consumer prod	ucts for sale at a	cannabis event must l	be stored in a secure,
39.9	locked contai	ner that is not acces	sible to the publ	ic. Such items being	stored at a cannabis
39.10	event shall no	ot be left unattended	•		
39.11	(i) All car	ınabis plants, adult-ı	use cannabis flov	wer, adult-use cannat	ois products,
39.12	lower-potenc	y hemp edibles, and	hemp-derived c	onsumer products fo	r sale at a cannabis
39.13	event must co	omply with this chap	oter and rules add	opted pursuant to this	s chapter regarding
39.14	the testing, pa	ackaging, and labeli	ng of those items	S.	
39.15	(j) All can	ınabis plants, adult-u	se cannabis flow	ver, and adult-use can	nabis products sold,
39.16	damaged, or	destroyed at a canna	bis event must b	be recorded in the star	tewide monitoring
39.17	system.				
39.18	Sec. 52. Mi	nnesota Statutes 202	24, section 342.4	0, is amended by add	ling a subdivision to
39.19	read:				
39.20	<u>Subd. 7a.</u>	Cannabis sample p	oroducts. (a) Not	withstanding any oth	er provisions of law,
39.21	an authorized	l retailer may give av	way samples of c	annabis plants, canna	bis flower, cannabis
39.22	products, low	er-potency hemp edi	bles, or hemp-der	rived consumer produ	cts during a cannabis
39.23	event. A labe	l or notice containin	g the informatio	n required to be affix	xed to the packaging
39.24	or container of	containing cannabis	flower, adult-use	e cannabis products, l	ower-potency hemp
39.25	edibles, or he	mp-derived consum	er products sold	to customers must b	e displayed and
39.26	available for	consumers.			
39.27	(b) Produc	cts given away as sa	mples must not	consist of more than:	<u> </u>
39.28	(1) one gr	am of adult-use can	nabis flower or a	adult-use cannabis co	encentrate;
39.29	(2) ten mi	lligrams of tetrahyd	rocannabinol inf	used in an edible car	mabis product; and

(3) five milligrams of delta-9 tetrahydrocannabinol, five milligrams of cannabidiol, five

milligrams of cannabigerol, or any combination of those cannabinoids that does not exceed

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the identified amounts in a lower-potency hemp edible.

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(c) Authorized retailers must not give away samples to an individual who is visibly 40.1 intoxicated. 40.2 (d) Samples must be recorded in the statewide monitoring system. 40.3 Sec. 53. Minnesota Statutes 2024, section 342.43, is amended by adding a subdivision to 40.4 read: 40.5 Subd. 3. Exception; municipal or county licenses. Notwithstanding any law to the 40.6 contrary, a city or county that establishes, owns, or operates a municipal cannabis store may 40.7 also hold a lower-potency hemp edible retailer license. 40.8 Sec. 54. Minnesota Statutes 2024, section 342.44, subdivision 1, is amended to read: 40.9 Subdivision 1. Application; contents. (a) Except as otherwise provided in this 40.10 subdivision, the provisions of this chapter relating to license applications, license selection 40.11 criteria, general ownership disqualifications and requirements, and general operational 40.12 requirements do not apply to hemp businesses. 40.13 (b) The office, by rule, shall establish forms and procedures for the processing of hemp 40.14 licenses issued under this chapter. At a minimum, any application to obtain or renew a hemp 40.15 license shall include the following information, if applicable: 40.16 (1) the name, address, and date of birth of the applicant; 40.17 (2) the address and legal property description of the business; 40.18 (3) proof of trade name registration; 40.19 (4) certification that the applicant will comply with the requirements of this chapter 40.20 relating to the ownership and operation of a hemp business; 40.21 (5) identification of one or more controlling persons or managerial employees as agents 40.22 who shall be responsible for dealing with the office on all matters; and 40.23 (6) a statement that the applicant agrees to respond to the office's supplemental requests 40.24 40.25 for information. (c) An applicant for a lower-potency hemp edible manufacturer license must submit an 40.26 attestation signed by a bona fide labor organization stating that the applicant has entered 40.27 into a labor peace agreement. 40.28 (d) An application on behalf of a corporation or association shall be signed by at least 40.29 two officers or managing agents of that entity. 40.30

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Sec. 55. Minnesota Statutes 2024, section 342.45, is amended by adding a subdivision to 41.1 41.2 read:

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- Subd. 6. Building conditions. (a) A lower-potency hemp edible manufacturer must comply with state and local building, fire, and zoning codes, requirements, and regulations.
- 41.5 (b) A lower-potency hemp edible manufacturer must ensure that licensed premises are maintained in a clean and sanitary condition and are free from infestation by insects, rodents, 41.6 or other pests. 41.7
- Sec. 56. Minnesota Statutes 2024, section 342.46, subdivision 6, is amended to read: 41.8
 - Subd. 6. Compliant products. (a) A lower-potency hemp edible retailer shall ensure that all lower-potency hemp edibles offered for sale comply with the limits on the amount and types of cannabinoids that a lower-potency hemp edible can contain, including but not limited to the requirement that lower-potency hemp edibles:
 - (1) consist of servings that contain no more than five milligrams of delta-9 tetrahydrocannabinol, no more than 25 milligrams of cannabidiol, no more than 25 milligrams of cannabigerol, or any combination of those cannabinoids that does not exceed the identified amounts, except that a lower-potency hemp edible that is intended to be consumed as a beverage may contain no more than ten milligrams of delta-9 tetrahydrocannabinol in a single-serving container;
- (2) do not contain more than a combined total of 0.5 milligrams of all other cannabinoids 41.19 41.20 per serving; and
- (3) do not contain an artificially derived cannabinoid other than delta-9 41.21 tetrahydrocannabinol. 41.22
 - (b) If a lower-potency hemp edible is packaged in a manner that includes more than a single serving, the lower-potency hemp edible must indicate each serving by scoring, wrapping, or other indicators that appear on the lower-potency hemp edible designating the individual serving size. If it is not possible to indicate a single serving by scoring or use of another indicator that appears on the product, the lower-potency hemp edible may not be packaged in a manner that includes more than a single serving in each container, except that a calibrated dropper, measuring spoon, or similar device for measuring a single serving may be used for any edible cannabinoid products that are intended to be combined with food or beverage products prior to consumption. If the lower-potency hemp edible is meant to be consumed as a beverage, the beverage container may not contain more than two

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product that includes recommended dosage requirements and other information as required

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by the office; and

43.1	(5) provide the patient with any other information required by the office.
43.2	(b) A cannabis business with a medical cannabis retail endorsement may not deliver
43.3	medical cannabis flower or medical cannabinoid products to a person enrolled in the registry
43.4	program unless the cannabis business with a medical cannabis retail endorsement also holds
43.5	a cannabis delivery service license. The delivery of medical cannabis flower and medical
43.6	cannabinoid products are subject to the provisions of section 342.42.
43.7	Sec. 58. Minnesota Statutes 2024, section 342.51, is amended by adding a subdivision to
43.8	read:
43.9	Subd. 2a. Distribution to visiting patients. (a) A cannabis business with a medical
43.10	cannabis retail endorsement may distribute medical cannabis flower or medical cannabinoic
43.11	products to a visiting patient.
43.12	(b) Before receiving a distribution of medical cannabis, a visiting patient must provide
43.13	to an employee of the cannabis business:
43.14	(1) a valid medical cannabis registration verification card or equivalent document issued
43.15	by a Tribal medical cannabis program that indicates that the visiting patient is authorized
43.16	to use medical cannabis on Indian lands over which the Tribe has jurisdiction; and
43.17	(2) a valid photographic identification card issued by the Tribal medical cannabis
43.18	program, a valid driver's license, or a valid state identification card.
43.19	(c) Prior to the distribution of medical cannabis flower or medical cannabinoid products
43.20	to a visiting patient, an employee of a cannabis business must:
43.21	(1) ensure that a patient-specific label has been applied to all medical cannabis flower
43.22	and medical cannabinoid products. The label must include the recommended dosage
43.23	requirements and other information required by the office; and
43.24	(2) provide the patient with any other information required by the office.
43.25	(d) For each transaction that involves a visiting patient, a cannabis business with a
43.26	medical cannabis retail endorsement must report to the office on a weekly basis:
43.27	(1) the name of the visiting patient;
43.28	(2) the name of the Tribal medical cannabis program in which the visiting patient is
43.29	enrolled;
43.30	(3) the amount and dosages of medical cannabis distributed;

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(4) the chemical composition of the medical cannabis distributed; and

4.1	(5) the tracking number assigned to the medical cannabis that was distributed to the
4.2	visiting patient.
14.3	(e) A cannabis business with a medical cannabis retail endorsement may distribute
4.4	medical cannabis flower and medical cannabinoid products to a visiting patient in a motor
4.5	vehicle if:
4.6	(1) an employee of the cannabis business with a medical cannabis retail endorsement
4.7	receives payment and distributes medical cannabis flower and medical cannabinoid products
4.8	in a designated zone that is as close as feasible to the front door of the facility where the
4.9	cannabis business is located;
4.10	(2) the cannabis business with a medical cannabis retail endorsement ensures that the
4.11	receipt of payment and distribution of medical cannabis flower and medical cannabinoid
4.12	products are visually recorded by a closed-circuit television surveillance camera and provides
4.13	any other necessary security safeguards required by the office;
4.14	(3) the cannabis business with a medical cannabis retail endorsement does not store
4.15	medical cannabis flower or medical cannabinoid products outside a restricted access area;
4.16	(4) an employee of the cannabis business with a medical cannabis retail endorsement
4.17	transports medical cannabis flower and medical cannabinoid products from a restricted
4.18	access area to the designated zone for distribution to patients only after confirming that the
4.19	visiting patient has arrived in the designated zone;
4.20	(5) the payment for and distribution of medical cannabis flower and medical cannabinoid
4.21	products to a patient only occurs after meeting the requirements in paragraph (b);
4.22	(6) immediately following the distribution of medical cannabis flower or medical
4.23	cannabinoid products to a patient, an employee of the cannabis business with a medical
4.24	cannabis retail endorsement records the transaction in the statewide monitoring system; and
4.25	(7) immediately following the distribution of medical cannabis flower and medical
4.26	cannabinoid products, an employee of the cannabis business with a medical cannabis retail
4.27	endorsement transports all payments received into the facility where the cannabis business
4.28	is located.
4.29	Sec. 59. Minnesota Statutes 2024, section 342.52, is amended by adding a subdivision to
4.29	read:
4.31	Subd. 7a. Allowable delivery methods. A patient in the registry program may receive
4.32	medical cannabis flower and medical cannabinoid products. The office may approve

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additional delivery methods to expand the types of products that qualify as medical 45.1 cannabinoid products. 45.2

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- Sec. 60. Minnesota Statutes 2024, section 342.52, subdivision 9, is amended to read:
- Subd. 9. Registered designated caregiver. (a) The office must register a designated caregiver for a patient if the patient requires assistance in administering medical cannabis flower or medical cannabinoid products; obtaining medical cannabis flower, medical cannabinoid products, or medical cannabis paraphernalia from a cannabis business with a medical cannabis retail endorsement; or cultivating cannabis plants as permitted by section 342.09, subdivision 2.
 - (b) In order to serve as a designated caregiver, a person must:
- (1) be at least 18 years of age; 45.11
 - (2) agree to only possess the patient's medical cannabis flower and medical cannabinoid products for purposes of assisting the patient; and
 - (3) agree that if the application is approved, the person will not serve as a registered designated caregiver for more than six registered patients at one time. Patients who reside in the same residence count as one patient.
 - (c) Nothing in this section shall be construed to prevent a registered designated caregiver from being enrolled in the registry program as a patient and possessing and administering medical cannabis flower or medical cannabinoid products as a patient.
 - (d) Notwithstanding any law to the contrary, a registered designated caregiver approved to assist a patient enrolled in the registry program with obtaining medical cannabis flower may cultivate cannabis plants on behalf of one patient. A registered designated caregiver may grow up to eight cannabis plants for the patient household that the registered designated caregiver is approved to assist with obtaining medical cannabis flower. If a patient enrolled in the registry program directs the patient's registered designated caregiver to cultivate cannabis plants on behalf of the patient, the patient must assign the patient's right to cultivate cannabis plants to the registered designated caregiver and the notify the office. A patient who assigns the patient's right to cultivate cannabis plants to a registered caregiver is prohibited from cultivating cannabis plants for personal use. Nothing in this paragraph limits the right of a registered designated caregiver cultivating cannabis plants on behalf of a patient enrolled in the registry program to also cultivate cannabis plants for personal use pursuant to section 342.09, subdivision 2.

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Sec. 61. Minnesota Statutes 2024, section 342.56, subdivision 2, is amended to read:

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Subd. 2. **Health care facilities.** (a) Health care facilities licensed under chapter 144A; hospice providers licensed under chapter 144A; boarding care homes or supervised living facilities licensed under section 144.50; assisted living facilities under chapter 144G; facilities owned, controlled, managed, or under common control with hospitals licensed under chapter 144; and other health care facilities licensed by the commissioner of health or the commissioner of human services may adopt reasonable restrictions on the use of medical cannabis flower or medical, cannabinoid products, lower-potency hemp edibles, hemp-derived consumer products, or hemp-derived topical products by a patient enrolled in the registry program who resides at or is actively receiving treatment or care at the facility. The restrictions may include a provision that the facility must not store or maintain a patient's supply of medical cannabis flower or medical cannabinoid products on behalf of the patient; that a patient store the patient's supply of medical cannabis flower or medicinal, cannabinoid products, lower-potency hemp edibles, hemp-derived consumer products, or hemp-derived topical products in a locked container accessible only to the patient, the patient's designated caregiver, or the patient's parent, legal guardian, or spouse; that the facility is not responsible for providing medical cannabis or hemp for patients; and that medical cannabis flower or medical, cannabinoid products, lower-potency hemp edibles, hemp-derived consumer products, or hemp-derived topical products are used only in a location specified by the facility or provider. Nothing in this subdivision requires facilities and providers listed in this subdivision to adopt such restrictions.

(b) No facility or provider listed in this subdivision may unreasonably limit a patient's access to or use of medical cannabis flower or medical cannabinoid products, lower-potency hemp edibles, hemp-derived consumer products, or hemp-derived topical products to the extent that such use is authorized under sections 342.51 to 342.59, or, in the case of a visiting patient, authorized to use medical cannabis under the laws of their state of residence. No facility or provider listed in this subdivision may prohibit a patient access to or use of medical cannabis flower or medical cannabinoid products due solely to the fact that cannabis is a controlled substance pursuant to the federal Uniform Controlled Substances Act. If a federal regulatory agency, the United States Department of Justice, or the federal Centers for Medicare and Medicaid Services takes one of the following actions, a facility or provider may suspend compliance with this paragraph until the regulatory agency, the United States Department of Justice, or the federal Centers for Medicare and Medicaid Services notifies the facility or provider that it may resume permitting the use of medical cannabis flower or medical, cannabinoid products, lower-potency hemp edibles, hemp-derived consumer

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<u>products</u>, or <u>hemp-derived topical products</u> within the facility or in the provider's service setting:

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- (1) a federal regulatory agency or the United States Department of Justice initiates enforcement action against a facility or provider related to the facility's compliance with the medical cannabis program; or
- (2) a federal regulatory agency, the United States Department of Justice, or the federal Centers for Medicare and Medicaid Services issues a rule or otherwise provides notification to the facility or provider that expressly prohibits the use of medical cannabis in health care facilities or otherwise prohibits compliance with the medical cannabis program.
- (c) An employee or agent of a facility or provider listed in this subdivision or a person licensed under chapter 144E is not violating this chapter or chapter 152 for the possession of medical cannabis flower or medical cannabinoid products while carrying out employment duties, including providing or supervising care to a patient enrolled in the registry program, or distribution of medical cannabis flower or medical cannabinoid products to a patient enrolled in the registry program who resides at or is actively receiving treatment or care at the facility or from the provider with which the employee or agent is affiliated.
- 47.17 (d) Nothing in this subdivision is intended to require a facility covered by this subdivision
 47.18 to permit violations of sections 144.411 to 144.417.
- (e) This subdivision does not apply to sober homes under section 254B.181.
- Sec. 62. Minnesota Statutes 2024, section 342.57, is amended to read:

342.57 PROTECTIONS FOR REGISTRY PROGRAM PARTICIPANTS.

- Subdivision 1. **Presumption.** (a) There is a presumption that a patient or other person
 an individual enrolled in the registry program or a Tribal medical cannabis program patient
 is engaged in the authorized use or possession of medical cannabis flower and medical
 cannabinoid products.
- (b) This presumption may be rebutted by evidence that:
- 47.27 (1) the use or possession of medical cannabis flower or medical cannabinoid products
 47.28 by a patient or other person enrolled in the registry program was not for the purpose of
 47.29 assisting with, treating, or alleviating the patient's qualifying medical condition or symptoms
 47.30 associated with the patient's qualifying medical condition=; or
- 47.31 (2) a Tribal medical cannabis program patient's use of medical cannabis was not for a purpose authorized by the Tribal medical cannabis program.

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Subd. 2. Criminal and civil protections. (a) Subject to section 342.56, the following are not violations of this chapter or chapter 152:

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- (1) use or possession of medical cannabis flower, medical cannabinoid products, or medical cannabis paraphernalia by a patient enrolled in the registry program or by a visiting patient or a Tribal medical cannabis program patient to whom medical cannabis flower or medical cannabinoid products are distributed under section 342.51, subdivision 5;
- (2) possession of medical cannabis flower, medical cannabinoid products, or medical cannabis paraphernalia by a registered designated caregiver or a parent, legal guardian, or spouse of a patient enrolled in the registry program; or
- (3) possession of medical cannabis flower, medical cannabinoid products, or medical cannabis paraphernalia by any person while carrying out duties required under sections 342.51 to 342.60.
- (b) The Office of Cannabis Management, members of the Cannabis Advisory Council, Office of Cannabis Management employees, agents or contractors of the Office of Cannabis Management, members of a Tribal medical cannabis board, a Tribal medical cannabis board's staff, a Tribal medical cannabis board's agents or contractors, and health care practitioners participating in the registry program are not subject to any civil penalties or disciplinary action by the Board of Medical Practice, the Board of Nursing, or any business, occupational, or professional licensing board or entity solely for participating in the registry program or in a Tribal medical cannabis program either in a professional capacity or as a patient. A pharmacist licensed under chapter 151 is not subject to any civil penalties or disciplinary action by the Board of Pharmacy when acting in accordance with sections 342.51 to 342.60 either in a professional capacity or as a patient. Nothing in this section prohibits a professional licensing board from taking action in response to a violation of law.
- (c) Notwithstanding any law to the contrary, a Cannabis Advisory Council member, the governor, or an employee of a state agency must not be held civilly or criminally liable for any injury, loss of property, personal injury, or death caused by any act or omission while acting within the scope of office or employment under sections 342.51 to 342.60.
- (d) Federal, state, and local law enforcement authorities are prohibited from accessing the registry except when acting pursuant to a valid search warrant. Notwithstanding section 13.09, a violation of this paragraph is a gross misdemeanor.
- (e) Notwithstanding any law to the contrary, the office and employees of the office must not release data or information about an individual contained in any report or document or in the registry and must not release data or information obtained about a patient enrolled in

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the registry program, except as provided in sections 342.51 to 342.60. Notwithstanding section 13.09, a violation of this paragraph is a gross misdemeanor.

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- (f) No information contained in a report or document, contained in the registry, or obtained from a patient under sections 342.51 to 342.60 or from a Tribal medical cannabis program patient may be admitted as evidence in a criminal proceeding, unless:
 - (1) the information is independently obtained; or
- (2) admission of the information is sought in a criminal proceeding involving a criminal violation of sections 342.51 to 342.60.
 - (g) Possession of a registry verification or an application for enrollment in the registry program and possession of a verification of enrollment or its equivalent issued by a Tribal medical cannabis program or application for enrollment in a Tribal medical cannabis program by a person entitled to possess the verification of enrollment or application for enrollment:
 - (1) does not constitute probable cause or reasonable suspicion;
- 49.14 (2) must not be used to support a search of the person or property of the person with a registry verification or application to enroll in the registry program; and 49.15
 - (3) must not subject the person or the property of the person to inspection by any government agency.
 - (h) A patient enrolled in the registry program or in a Tribal medical cannabis program must not be subject to any penalty or disciplinary action by an occupational or a professional licensing board solely because:
- (1) the patient is enrolled in the registry program or in a Tribal medical cannabis program; 49.21 or 49.22
- (2) the patient has a positive test for cannabis components or metabolites. 49.23
 - Subd. 3. School enrollment; rental property. (a) No school may refuse to enroll or otherwise penalize a patient or person enrolled in the registry program or a Tribal medical cannabis program as a pupil solely because the patient or person is enrolled in the registry program or a Tribal medical cannabis program, unless failing to do so would violate federal law or regulations or cause the school to lose a monetary or licensing-related benefit under federal law or regulations.
- (b) No landlord may refuse to lease to a patient or person enrolled in the registry program 49.30 or a Tribal medical cannabis program or otherwise penalize a patient or person enrolled in 49.31 the registry program or a Tribal medical cannabis program solely because the patient or 49.32

person is enrolled in the registry program or a Tribal medical cannabis program, unless 50.1 failing to do so would violate federal law or regulations or cause the landlord to lose a 50.2 50.3 monetary or licensing-related benefit under federal law or regulations. (c) A school must not refuse to enroll a patient as a pupil solely because cannabis is a 50.4 50.5 controlled substance according to the Uniform Controlled Substances Act, United States Code, title 21, section 812. 50.6 (d) A school must not penalize a pupil who is a patient solely because cannabis is a 50.7 controlled substance according to the Uniform Controlled Substances Act, United States 50.8 Code, title 21, section 812. 50.9 (e) A landlord must not refuse to lease a property to a patient solely because cannabis 50.10 is a controlled substance according to the Uniform Controlled Substances Act, United States 50.11 50.12 Code, title 21, section 812. (f) A landlord must not otherwise penalize a patient solely because cannabis is a controlled 50.13 substance according to the Uniform Controlled Substances Act, United States Code, title 50.14 21, section 812. 50.15 Subd. 4. Medical care. For purposes of medical care, including organ transplants, a 50.16 patient's use of medical cannabis flower or medical cannabinoid products according to 50.17 sections 342.51 to 342.60, or a Tribal medical cannabis program patient's use of medical 50.18 cannabis as authorized by a Tribal medical cannabis program, is considered the equivalent 50.19 of the authorized use of a medication used at the discretion of a health care practitioner and 50.20 does not disqualify a patient from needed medical care. 50.21 Subd. 5. Employment. (a) Unless a failure to do so would violate federal or state law 50.22 or regulations or cause an employer to lose a monetary or licensing-related benefit under 50.23 federal law or regulations, an employer may not discriminate against a person in hiring, 50.24 termination, or any term or condition of employment, or otherwise penalize a person, if the 50.25 discrimination is based on: 50.26 (1) the person's status as a patient or person an individual enrolled in the registry program; 50.27 50.28 or (2) the person's status as a Tribal medical cannabis program patient; or 50.29 (2) (3) a patient's positive drug test for cannabis components or metabolites, unless the 50.30 patient used, possessed, sold, transported, or was impaired by medical cannabis flower or 50.31 a medical cannabinoid product on work premises, during working hours, or while operating 50.32

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an employer's machinery, vehicle, or equipment.

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(b) An employee who is a patient in the registry program or a Tribal medical cannabis 51.1 program and whose employer requires the employee to undergo drug testing according to 51.2 section 181.953 may present the employee's registry verification or verification of enrollment 51.3 in a Tribal medical cannabis program as part of the employee's explanation under section 51.4 181.953, subdivision 6. 51.5 Subd. 5a. Notice. An employer, a school, or a landlord must provide written notice to 51.6 a patient at least 14 days before the employer, school, or landlord takes an action against 51.7 the patient that is prohibited under subdivision 3 or 5. The written notice must cite the 51.8 specific federal law or regulation the employer, school, or landlord believes would be 51.9 violated if the employer, school, or landlord fails to take action. The notice must specify 51.10 which monetary or licensing-related benefit under federal law or regulations the employer, 51.11 school, or landlord would lose if the employer, school, or landlord fails to take action. 51.12 Subd. 6. Custody; visitation; parenting time. A person must not be denied custody of 51.13 a minor child or visitation rights or parenting time with a minor child based solely on the 51.14 person's individual's status as a patient or person an individual enrolled in the registry 51.15 program or on the individual's status as a Tribal medical cannabis program patient. There 51.16 must be no presumption of neglect or child endangerment for conduct allowed under sections 51.17 342.51 to 342.60 or under a Tribal medical cannabis program, unless the person's individual's 51.18 behavior creates an unreasonable danger to the safety of the minor as established by clear 51.19 and convincing evidence. 51.20 Subd. 6a. Retaliation prohibited. A school, a landlord, a health care facility, or an 51.21 employer must not retaliate against a patient for asserting the patient's rights or seeking 51.22 remedies under this section or section 152.32. 51.23 Subd. 7. Action for damages; injunctive relief. In addition to any other remedy provided 51.24 by law, a patient or person an individual enrolled in the registry program or a Tribal medical 51.25 51.26 cannabis program may bring an action for damages against any person who violates subdivision 3, 4, or 5. A person who violates subdivision 3, 4, or 5 is liable to a patient or 51.27 person an individual enrolled in the registry program or a Tribal medical cannabis program 51.28 injured by the violation for the greater of the person's actual damages or a civil penalty of 51.29 \$100 \$1,000 and reasonable attorney fees. A patient may bring an action for injunctive relief 51.30 to prevent or end a violation of subdivisions 3 to 6a. 51.31 Subd. 8. Sanctions restricted for those on parole, supervised release, or conditional 51.32 release. (a) This subdivision applies to an individual placed on parole, supervised release, 51.33 or conditional release. 51.34

(b) The commissioner of corrections may not:

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- (1) prohibit an individual from participating in the registry program <u>or a Tribal medical</u> cannabis program as a condition of release; or
- (2) revoke an individual's parole, supervised release, or conditional release or otherwise sanction an individual solely:
 - (i) for participating in the registry program or a Tribal medical cannabis program; or
 - (ii) for a positive drug test for cannabis components or metabolites.
- Sec. 63. Minnesota Statutes 2024, section 342.59, subdivision 2, is amended to read:
 - Subd. 2. **Allowable use; prohibited use.** Data specified in subdivision 1 may be used to comply with chapter 13, to comply with a request from the legislative auditor or the state auditor in the performance of official duties, and for purposes specified in sections 342.47 342.51 to 342.60. Data specified in subdivision 1 and maintained by the Office of Cannabis Management or Division of Medical Cannabis must not be used for any purpose not specified in sections 342.47 342.51 to 342.60 and must not be combined or linked in any manner with any other list, dataset, or database. Data specified in subdivision 1 must not be shared with any federal agency, federal department, or federal entity unless specifically ordered to do so by a state or federal court.
 - Sec. 64. Minnesota Statutes 2024, section 342.61, subdivision 4, is amended to read:
 - Subd. 4. **Testing of samples; disclosures.** (a) On a schedule determined by the office, every cannabis microbusiness, cannabis mezzobusiness, cannabis cultivator, cannabis manufacturer, cannabis wholesaler with an endorsement to import products, lower-potency hemp edible manufacturer, or medical cannabis combination business shall make each batch of cannabis flower, cannabis products, artificially derived cannabinoids, lower-potency hemp edibles, or hemp-derived consumer products grown, manufactured, or imported by the cannabis business or hemp business available to a cannabis testing facility.
 - (b) A cannabis microbusiness, cannabis mezzobusiness, cannabis cultivator, cannabis manufacturer, cannabis wholesaler with an endorsement to import products, lower-potency hemp edible manufacturer, or medical cannabis combination business must disclose all known information regarding pesticides, fertilizers, solvents, or other foreign materials, including but not limited to catalysts used in creating artificially derived cannabinoids, applied or added to the batch of cannabis flower, cannabis products, artificially derived cannabinoids, lower-potency hemp edibles, or hemp-derived consumer products subject to

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testing. Disclosure must be made to the cannabis testing facility and must include information about all applications by any person, whether intentional or accidental.

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- (c) The A cannabis testing facility business shall select one or more representative samples from each batch, test the samples for the presence of contaminants, and test the samples for potency and homogeneity and to allow the cannabis flower, cannabis product, artificially derived cannabinoid, lower-potency hemp edible, or hemp-derived consumer product to be accurately labeled with its cannabinoid profile. Testing for contaminants must include testing for residual solvents, foreign material, microbiological contaminants, heavy metals, pesticide residue, mycotoxins, and any items identified pursuant to paragraph (b), and may include testing for other contaminants. A cannabis testing facility must destroy or return to the cannabis business or hemp business any part of the sample that remains after testing.
- Sec. 65. Minnesota Statutes 2024, section 342.63, subdivision 2, is amended to read:
- Subd. 2. **Content of label; cannabis.** All cannabis flower and hemp-derived consumer products that consist of hemp plant parts sold to customers or patients must have affixed on the packaging or container of the cannabis flower or hemp-derived consumer product a label that contains at least the following information:
 - (1) the name and license number of the cannabis microbusiness, cannabis mezzobusiness, cannabis cultivator, medical cannabis combination business, or industrial hemp grower where the cannabis flower or hemp plant part was cultivated;
 - (2) the net weight or volume of cannabis flower or hemp plant parts in the package or container;
- 53.23 (3) the batch number;
- 53.24 (4) the cannabinoid profile;
- (5) a universal symbol established by the office indicating that the package or container contains cannabis flower, a cannabis product, a lower-potency hemp edible, or a hemp-derived consumer product;
 - (6) verification that the cannabis flower or hemp plant part was tested according to section 342.61 and that the cannabis flower or hemp plant part complies with the applicable standards;
 - (7) information on the usage of the cannabis flower or hemp-derived consumer product;
- 53.32 (8) the following statement: "Keep this product out of reach of children."; and

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(9) any other statements or information required by the office.

Sec. 66. Minnesota Statutes 2024, section 342.63, subdivision 3, is amended to read:

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- Subd. 3. Content of label; cannabinoid products. (a) All cannabis products, lower-potency hemp edibles, hemp concentrate, hemp-derived consumer products other than products subject to the requirements under subdivision 2, medical cannabinoid products, and hemp-derived topical products sold to customers or patients must have affixed to the packaging or container of the cannabis product a label that contains at least the following information:
- (1) the name and license number of the cannabis microbusiness, cannabis mezzobusiness, cannabis cultivator, medical cannabis combination business, or industrial hemp grower that cultivated the cannabis flower or hemp plant parts used in the cannabis product, lower-potency hemp edible, hemp-derived consumer product, or medical cannabinoid product;
- (2) the name and license number of the cannabis microbusiness, cannabis mezzobusiness, cannabis manufacturer, lower-potency hemp edible manufacturer, medical cannabis combination business, or industrial hemp grower that manufactured the cannabis concentrate, hemp concentrate, or artificially derived cannabinoid and, if different, the name and license number of the cannabis microbusiness, cannabis mezzobusiness, cannabis manufacturer, lower-potency hemp edible manufacturer, or medical cannabis combination business that manufactured the product;
- (3) the net weight or volume of the cannabis product, lower-potency hemp edible, or hemp-derived consumer product in the package or container;
- (4) the type of cannabis product, lower-potency hemp edible, or hemp-derived consumer product;
- 54.25 (5) the batch number;
- 54.26 (6) the serving size;
- 54.27 (7) the cannabinoid profile per serving and in total;
- 54.28 (8) a list of ingredients;
- (9) a universal symbol established by the office indicating that the package or container contains cannabis flower, a cannabis product, a lower-potency hemp edible, or a hemp-derived consumer product;

Sec. 66. 54

(10) a warning symbol developed by the office in consultation with the commissioner 55.1 of health and the Minnesota Poison Control System that: 55.2 (i) is at least three-quarters of an inch tall and six-tenths of an inch wide; 55.3 (ii) is in a highly visible color; 55.4 55.5 (iii) includes a visual element that is commonly understood to mean a person should stop; 55.6 55.7 (iv) indicates that the product is not for children; and (v) includes the phone number of the Minnesota Poison Control System; 55.8 (11) verification that the cannabis product, lower-potency hemp edible, hemp-derived 55.9 consumer product, or medical cannabinoid product was tested according to section 342.61 55.10 and that the cannabis product, lower-potency hemp edible, hemp-derived consumer product, 55.11 or medical cannabinoid product complies with the applicable standards; 55.12 (12) information on the usage of the product; 55.13 (13) the following statement: "Keep this product out of reach of children."; and 55.14 (14) any other statements or information required by the office. 55.15 (b) The office may by rule establish alternative labeling requirements for lower-potency 55.16 hemp edibles that are imported into the state if those requirements provide consumers with 55.17 information that is substantially similar to the information described in paragraph (a). 55.18 Sec. 67. Minnesota Statutes 2024, section 342.63, subdivision 5, is amended to read: 55.19 Subd. 5. Content of label; hemp-derived topical products. (a) All hemp-derived topical 55.20 products sold to customers must have affixed to the packaging or container of the product 55.21 a label that contains at least the following information: 55.22 (1) the manufacturer name, location, phone number, and website; 55.23 (2) the name and address of the independent, accredited laboratory used by the 55.24 manufacturer to test the product; 55.25 (3) the net weight or volume of the product in the package or container; 55.26 (4) the type of topical product; 55.27 (5) the amount or percentage of cannabidiol, cannabigerol, or any other cannabinoid, 55.28 derivative, or extract of hemp, per serving and in total; 55.29

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(6) a list of ingredients;

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(7) a statement that the product does not claim to diagnose, treat, cure, or prevent any disease and that the product has not been evaluated or approved by the United States Food and Drug Administration, unless the product has been so approved; and (8) any other statements or information required by the office. (b) The information required in paragraph (a), clauses (1), (2), and (5), may be provided through the use of a scannable barcode or matrix barcode that links to a page on a website maintained by the manufacturer or distributor if that page contains all of the information required by this subdivision. Sec. 68. Minnesota Statutes 2024, section 342.63, subdivision 6, is amended to read: Subd. 6. Additional information. (a) A cannabis microbusiness, cannabis mezzobusiness, cannabis retailer, or medical cannabis combination business must provide customers and patients with the following information: (1) factual information about impairment effects and the expected timing of impairment effects, side effects, adverse effects, and health risks of cannabis flower, cannabis products, lower-potency hemp edibles, and hemp-derived consumer products; (2) a statement that customers and patients must not operate a motor vehicle or heavy machinery while under the influence of cannabis flower, cannabis products, lower-potency hemp edibles, and hemp-derived consumer products; (3) resources customers and patients may consult to answer questions about cannabis flower, cannabis products, lower-potency hemp edibles, and hemp-derived consumer products, and any side effects and adverse effects; (4) contact information for the poison control center and a safety hotline or website for customers to report and obtain advice about side effects and adverse effects of cannabis flower, cannabis products, lower-potency hemp edibles, and hemp-derived consumer products; (5) substance use disorder treatment options; and (6) any other information specified by the office. (b) A cannabis microbusiness, cannabis mezzobusiness, cannabis retailer, or medical cannabis combination business may include the information described in paragraph (a) by: (1) including the information on the label affixed to the packaging or container of cannabis

flower, cannabis products, lower-potency hemp edibles, and hemp-derived consumer products

Sec. 68. 56

(1) (2) posting the information in the premises of the cannabis microbusiness, cannabis 57.1 mezzobusiness, cannabis retailer, or medical cannabis combination business; or 57.2 (2) (3) providing the information on a separate document or pamphlet provided to 57.3 customers or patients when the customer purchases cannabis flower, a cannabis product, a 57.4 57.5 lower-potency hemp edible, or a hemp-derived consumer product. Sec. 69. Minnesota Statutes 2024, section 342.66, subdivision 6, is amended to read: 57.6 Subd. 6. Prohibitions. (a) A product sold to consumers under this section must not be 57.7 manufactured, marketed, distributed, or intended: 57.8 57.9 (1) for external or internal use in the diagnosis, cure, mitigation, treatment, or prevention of disease in humans or other animals; 57.10 (2) to affect the structure or any function of the bodies of humans or other animals; 57.11 (3) to be consumed by combustion or vaporization of the product and inhalation of 57.12 smoke, aerosol, or vapor from the product; 57.13 57.14 (4) to be consumed through chewing; or 57.15 (5) to be consumed through injection or application to nonintact skin or a mucous membrane or nonintact skin, except for products applied sublingually. 57.16 57.17 (b) A product manufactured, marketed, distributed, or sold to consumers under this section must not: 57.18 57.19 (1) consist, in whole or in part, of any filthy, putrid, or decomposed substance; (2) have been produced, prepared, packed, or held under unsanitary conditions where 57.20 the product may have been rendered injurious to health, or where the product may have 57.21 been contaminated with filth; 57.22 57.23 (3) be packaged in a container that is composed, in whole or in part, of any poisonous or deleterious substance that may render the contents injurious to health; 57.24 57.25 (4) contain any additives or excipients that have been found by the United States Food and Drug Administration to be unsafe for human or animal consumption; 57.26 57.27 (5) contain a cannabinoid or an amount or percentage of cannabinoids that is different than the information stated on the label; 57.28 (6) contain a cannabinoid, other than cannabidiol, cannabigerol, or a cannabinoid 57.29 approved by the office, in an amount that exceeds the standard established in subdivision 57.30

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23, paragraph (c); or

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(7) contain any contaminants for which testing is required by the office in amounts that 58.1 exceed the acceptable minimum standards established by the office. 58.2 (c) No product containing any cannabinoid may be sold to any individual who is under 58.3 21 years of age. 58.4 Sec. 70. Minnesota Statutes 2024, section 609A.06, subdivision 3, is amended to read: 58.5 Subd. 3. Eligibility; cannabis offense. (a) A person is eligible for an expungement or 58.6 resentencing to a lesser offense if: 58.7 (1) the person was convicted of, or adjudication was stayed for, a violation of any of the 58.8 following a first-, second-, third-, fourth-, or fifth-degree controlled substance crime involving 58.9 the sale or possession of marijuana or tetrahydrocannabinols: 58.10 (i) section 152.021, subdivision 1, clause (6); 58.11 (ii) section 152.021, subdivision 2, clause (6); 58.12 (iii) section 152.022, subdivision 1, clause (5), or clause (7), item (iii); 58.13 (iv) section 152.022, subdivision 2, clause (6); 58.14 58.15 (v) section 152.023, subdivision 1, clause (5); (vi) section 152.023, subdivision 2, clause (5); 58.16 (vii) section 152.024, subdivision (4); or 58.17 (viii) section 152.025, subdivision 2, clause (1) under Minnesota Statutes 2023 58.18 Supplement, section 152.021, 152.022, 152.023, 152.024, or 152.025, or a previous version 58.19 of those or any other statutes criminalizing the possession, sale, transportation, or cultivation 58.20 of marijuana or tetrahydrocannabinols; 58.21 (2) the offense did not involve a dangerous weapon, the intentional infliction of bodily 58.22 harm on another, an attempt to inflict bodily harm on another, or an act committed with the 58.23 intent to cause fear in another of immediate bodily harm or death; 58.24 (3) the act on which the charge was based would either be a lesser offense or no longer 58.25 be a crime after August 1, 2023; and 58.26 58.27 (4) the person did not appeal the conviction, any appeal was denied, or the deadline to

Sec. 70. 58

file an appeal has expired.

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(b) A person is eligible for an expungement for any other offense charged along with
the underlying crime described in paragraph (a) if the charge was either dismissed or eligible
for expungement under section 609A.055.
(c) For purposes of this subdivision, a "lesser offense" means a nonfelony offense if the
person was charged with a felony.

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

- Sec. 71. Minnesota Statutes 2024, section 609A.06, subdivision 7, is amended to read:
- Subd. 7. **Review and determination.** (a) The Cannabis Expungement Board shall review all available records to determine whether the conviction or stay of adjudication or charge is eligible for an expungement or resentencing to a lesser offense. An expungement under this section is presumed to be in the public interest unless there is clear and convincing evidence that an expungement or resentencing to a lesser offense would create a risk to public safety.
- (b) If the Cannabis Expungement Board determines that an expungement is in the public interest, the board shall determine whether a person's conviction should be vacated and charges should be dismissed.
- (c) If the Cannabis Expungement Board determines that an expungement is in the public interest, the board shall determine whether the limitations under section 609A.03, subdivision 5a, apply.
- (d) If the Cannabis Expungement Board determines that an expungement is in the public interest, the board shall determine whether the limitations under section 609A.03, subdivision 7a, paragraph (b), clause (5), apply.
- (e) If the Cannabis Expungement Board determines that an expungement is not in the public interest, the board shall determine whether the person is eligible for resentencing to a lesser offense.
- (f) In making a determination under this subdivision, the Cannabis Expungement Board shall consider:
- (1) the nature and severity of the underlying crime, including but not limited to the total amount of marijuana or tetrahydrocannabinols possessed by the person and whether the offense involved a dangerous weapon, the intentional infliction of bodily harm on another, an attempt to inflict bodily harm on another, or an act committed with the intent to cause fear in another of immediate bodily harm or death;

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60.1	(2) whether an expungement or resentencing the person a lesser offense would increase
60.2	the risk, if any, the person poses to other individuals or society;
60.3	(3) if the person is under sentence, whether an expungement or resentencing to a lesser
60.4	offense would result in the release of the person and whether release earlier than the date
60.5	that the person would be released under the sentence currently being served would present
60.6	a danger to the public or would be compatible with the welfare of society;
60.7	(4) aggravating or mitigating factors relating to the underlying crime, including the
60.8	person's level of participation and the context and circumstances of the underlying crime;
60.9	(5) statements from victims and law enforcement, if any;
60.10	(6) if an expungement or resentencing the person to a lesser offense is considered,
60.11	whether there is good cause to restore the person's right to possess firearms and ammunition;
60.12	(7) if an expungement is considered, whether an expunged record of a conviction or stay
60.13	of adjudication may be opened for purposes of a background check required under section
60.14	122A.18, subdivision 8; and
60.15	(8) whether the person was also charged with other offenses in addition to the underlying
60.16	crime, the disposition of those other charges, and other factors deemed relevant by the
60.17	Cannabis Expungement Board.
60.18	(g) In making a determination under this subdivision, the Cannabis Expungement Board
60.19	shall not consider the impact the expungement would have on the offender based on any
60.20	records held by the Department of Health; Department of Children, Youth, and Families;
60.21	or Department of Human Services.
60.22	(h) The affirmative vote of three members is required for action taken at any meeting.
60.23	EFFECTIVE DATE. This section is effective the day following final enactment.
60.24	Sec. 72. Minnesota Statutes 2024, section 609A.06, subdivision 10, is amended to read:
60.25	Subd. 10. Notice to judicial branch and offenders. (a) The Cannabis Expungement
60.26	Board shall identify any conviction or stay of adjudication or charge that qualifies for an
60.27	order of expungement or resentencing to a lesser offense and notify the judicial branch of:
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	(1) the name and date of birth of a person whose conviction or stay of adjudication is
60.29	(1) the name and date of birth of a person whose conviction or stay of adjudication is eligible for an order of expungement or resentencing to a lesser offense;
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(4) if the person is eligible for an expungement, whether the person's conviction should 61.1 be vacated and charges should be dismissed; 61.2 (5) if the person is eligible for an expungement, whether there is good cause to restore 61.3 the offender's right to possess firearms and ammunition; 61.4 61.5 (6) if the person is eligible for an expungement, whether the limitations under section 609A.03, subdivision 7a, paragraph (b), clause (5), apply; and 61.6 61.7 (7) if the person is eligible for an expungement, whether the expungement should also apply to any other offenses charged in addition to the underlying crime; and 61.8 (8) if the person is eligible for resentencing to a lesser offense, the lesser sentence to be 61.9 imposed. 61.10 (b) The Cannabis Expungement Board shall make a reasonable and good faith effort to 61.11 notify any person whose conviction or stay of adjudication qualifies for an order of 61.12 expungement that the offense qualifies and notice is being sent to the judicial branch. Notice 61.13 sent pursuant to this paragraph shall inform the person that, following the order of 61.14 expungement, any records of an arrest, conviction, or incarceration should not appear on 61.15 any background check or study. 61.16 **EFFECTIVE DATE.** This section is effective the day following final enactment. 61.17 Sec. 73. Minnesota Statutes 2024, section 609A.06, subdivision 12, is amended to read: 61.18 Subd. 12. Order of expungement. (a) Upon receiving notice that an offense qualifies 61.19 for expungement, the court shall issue an order sealing all records relating to an arrest, 61.20 indictment or information, trial, verdict, or dismissal and discharge for an offense described 61.21 in subdivision 3, and any other offenses charged in addition to the underlying crime if 61.22 identified by the Cannabis Expungement Board as eligible for expungement. In addition, 61.23 the court shall order all records, including those pertaining to probation, incarceration, or 61.24 supervision, held by the Department of Corrections or local correctional officials sealed. 61.25 The courts shall not order the Department of Health; the Department of Children, Youth, 61.26 and Families; or the Department of Human Services to seal records under this section. If 61.27 the Cannabis Expungement Board determined that the person's conviction should be vacated 61.28 and charges should be dismissed, the order shall vacate and dismiss the charges. 61.29 (b) If the Cannabis Expungement Board determined that there is good cause to restore 61.30 the person's right to possess firearms and ammunition, the court shall issue an order pursuant 61.31 to section 609.165, subdivision 1d. 61.32

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- (c) If the Cannabis Expungement Board determined that an expunged record of a conviction or stay of adjudication may not be opened for purposes of a background check required under section 122A.18, subdivision 8, the court shall direct the order specifically to the Professional Educator Licensing and Standards Board.
- (d) The court administrator shall send a copy of an expungement order issued under this section to each agency and jurisdiction whose records are affected by the terms of the order and send a letter to the last known address of the person whose offense has been expunged identifying each agency to which the order was sent.
- (e) In consultation with the commissioner of human services, the court shall establish a schedule on which it shall provide the commissioner of human services a list identifying the name and court file number or, if no court file number is available, the citation number of each record for a person who received an expungement under this section.
- (f) Data on the person whose offense has been expunged in a letter sent under this subdivision are private data on individuals as defined in section 13.02, subdivision 12.
- 62.15 **EFFECTIVE DATE.** This section is effective the day following final enactment.

62.16 Sec. 74. **REPEALER.**

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Minnesota Statutes 2024, sections 152.22, subdivision 2; and 342.151, subdivision 1, are repealed.

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APPENDIX Repealed Minnesota Statutes: S2370-2

152.22 DEFINITIONS.

Subd. 2. Commissioner. "Commissioner" means the commissioner of health.

342.151 EMPLOYEES OF LICENSE HOLDERS.

Subdivision 1. **Definitions.** For purposes of this section, a "license holder" includes a cannabis microbusiness, cannabis mezzobusiness, cannabis cultivator, cannabis manufacturer, cannabis retailer, cannabis wholesaler, cannabis transporter, cannabis testing facility, cannabis event organizer, cannabis delivery service, lower-potency hemp edible manufacturer, lower-potency hemp edible retailer, or medical cannabis combination business.