SENATE
STATE OF MINNESOTA
NINETY-FOURTH SESSION

S.F. No. 2372

(SENATE AUTHORS: DIBBLE)

DATE	D-PG	OFFICIAL STATUS
03/10/2025	714	Introduction and first reading Referred to Commerce and Consumer Protection See SF2372

A bill for an act

relating to cannabis; modifying medical cannabis and cannabis provisions; amending Minnesota Statutes 2024, sections 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, 54, by adding a subdivision; 342.02, subdivision 3; 342.12; 342.14, subdivisions 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; 342.33, subdivision 1; 342.44, subdivision 1; 342.46, subdivision 6; 342.52, by adding a subdivision; 342.57, subdivision 2; 342.59, subdivision 2; 342.61, subdivision 4; 342.63, subdivisions 2, 3, 6; repealing Minnesota Statutes 2024, sections 152.22, subdivision 2; 342.01, subdivision 71; 342.151, subdivision 1.

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

Section 1. 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. 2. 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. 3. 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 ~~commissioner~~ office to a patient enrolled in the registry program.

Sec. 4. Minnesota Statutes 2024, section 152.22, subdivision 13, is amended to read:

Subd. 13. **Registry verification.** "Registry verification" means the verification provided by the ~~commissioner~~ office 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.

Sec. 5. Minnesota Statutes 2024, section 152.24, is amended to read:

152.24 FEDERALLY APPROVED CLINICAL TRIALS.

The ~~commissioner~~ office 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 ~~commissioner~~ office 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. 6. Minnesota Statutes 2024, section 152.25, is amended to read:

152.25 ~~COMMISSIONER~~ OFFICE DUTIES.

Subdivision 1. **Medical cannabis manufacturer registration.** (a) The ~~commissioner~~ office shall register two in-state manufacturers for the production of all medical cannabis within the state. A registration agreement between the ~~commissioner~~ office and a manufacturer is nontransferable. The ~~commissioner~~ office shall register new manufacturers or reregister the existing manufacturers by December 1 every two years, using the factors described in this subdivision. The ~~commissioner~~ 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 ~~commissioner'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:

(1) begin supplying medical cannabis to patients by July 1, 2015; and

(2) comply with all requirements under sections 152.22 to 152.37.

(c) The ~~commissioner~~ office 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;

(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 ~~commissioner~~ office 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.

Subd. 1a. **Revocation or nonrenewal of a medical cannabis manufacturer 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 ~~commissioner~~ office 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;

(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;

(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 ~~commissioner~~ 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.

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~~ 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.

(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. 7. 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. Rules for which notice is published in the State Register before January 1, 2015, may be adopted using the process in section 14.389.

(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 of medical cannabis under section 152.22, subdivision 6, paragraph (a), clause (4). Section 14.386, paragraph (b), does not apply to these rules.

Sec. 8. Minnesota Statutes 2024, section 152.261, is amended to read:

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.

Sec. 9. Minnesota Statutes 2024, section 152.27, subdivision 2, is amended to read:

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 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.

Sec. 10. Minnesota Statutes 2024, section 152.27, subdivision 7, is amended to read:

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. 11. Minnesota Statutes 2024, section 152.28, subdivision 1, is amended to read:

Subdivision 1. **Health care practitioner duties.** (a) Prior to a patient's enrollment in the registry program, a health care practitioner shall:

(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, the health care practitioner shall:

(1) participate in the patient registry reporting system under the guidance and supervision of the office;

(2) report health records of the patient throughout the ongoing treatment of the patient to the office in a manner determined by the ~~commissioner~~ office and in accordance with subdivision 2;

(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.

(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. 12. 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:

(1) contains false or misleading statements about medical cannabis or about the medical cannabis registry program;

(2) uses colloquial terms to refer to medical cannabis, such as pot, weed, or grass;

(3) states or implies the health care practitioner is endorsed by the ~~Department of Health~~ office or by the medical cannabis registry program;

(4) includes images of cannabis in its plant or leaf form or of cannabis-smoking 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 ~~commissioner~~ office 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 ~~commissioner's~~ office's decision that a health care practitioner has violated this subdivision is a final decision of the ~~commissioner~~ office and is not subject to the contested case procedures in chapter 14.

Sec. 13. 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 ~~commissioner~~ 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 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.

(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.

(e) A manufacturer shall implement security requirements, including requirements for the delivery and transportation of hemp and hemp products, protection of each location by a fully operational security alarm system, facility access controls, perimeter intrusion 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.

(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~~ office shall conduct at least one unannounced inspection per year of each manufacturer that includes inspection of:

(1) business operations;

(2) physical locations of the manufacturer's manufacturing facility and distribution facilities;

(3) financial information and inventory documentation, including laboratory testing results; and

(4) physical and electronic security alarm systems.

Sec. 14. Minnesota Statutes 2024, section 152.29, subdivision 2, is amended to read:

Subd. 2. **Manufacturer; production.** (a) A manufacturer of medical cannabis shall provide a reliable and ongoing supply of all medical cannabis needed for the registry program through cultivation by the manufacturer and through the purchase of hemp from hemp growers.

(b) All cultivation, harvesting, manufacturing, packaging, and processing of medical cannabis must take place in an enclosed, locked facility at a physical address provided to the ~~commissioner~~ office during the registration process.

(c) A manufacturer must process and prepare any medical cannabis plant material or hemp plant material into a form allowable under section 152.22, subdivision 6, prior to distribution of any medical cannabis.

Sec. 15. Minnesota Statutes 2024, section 152.29, subdivision 3a, is amended to read:

Subd. 3a. **Transportation of medical cannabis; transport staffing.** (a) A medical cannabis manufacturer may staff a transport motor vehicle with only one employee if the medical cannabis manufacturer is transporting medical cannabis to either a certified laboratory for the purpose of testing or a facility for the purpose of disposal. If the medical 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 by rules adopted by the ~~commissioner~~ office.

(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 employee.

(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 ~~commissioner~~ office.

(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.

Sec. 16. Minnesota Statutes 2024, section 152.29, subdivision 4, is amended to read:

Subd. 4. **Report.** (a) Each manufacturer shall report to the ~~commissioner~~ office on a monthly basis the following information on each individual patient for the month prior to the report:

(1) the amount and dosages of medical cannabis distributed;

(2) the chemical composition of the medical cannabis; and

(3) the tracking number assigned to any medical cannabis distributed.

(b) For transactions involving Tribal medical cannabis program patients, each manufacturer shall report to the ~~commissioner~~ office on a weekly basis the following information on each individual Tribal medical cannabis program patient for the week prior to the report:

(1) the name of the Tribal medical cannabis program in which the Tribal medical cannabis program patient is enrolled;

(2) the amount and dosages of medical cannabis distributed;

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

(4) the tracking number assigned to the medical cannabis distributed.

Sec. 17. Minnesota Statutes 2024, section 152.31, is amended to read:

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.

Sec. 18. 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.

(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 ~~commissioner's~~ office's or Tribal medical cannabis board's staff, the ~~commissioner'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

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

(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. 19. 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 ~~commissioner~~ office 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.

(b) All fines collected under this subdivision shall be deposited in the state government special revenue fund.

Sec. 20. 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 ~~commissioner~~ office 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. 21. Minnesota Statutes 2024, section 152.35, is amended to read:

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. 22. 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 business affairs and conditions of any medical cannabis manufacturer, including but not limited to a review of the financing, budgets, revenues, sales, and pricing.

(b) An examination may cover the medical cannabis manufacturer's business affairs, practices, and conditions including but not limited to a review of the financing, budgets, revenues, sales, and pricing. The ~~commissioner~~ office shall determine the nature and scope of each examination and in doing so shall take into account all available relevant factors concerning the financial and business affairs, practices, and conditions of the examinee. The costs incurred by the department in conducting an examination shall be paid for by the medical cannabis manufacturer.

(c) When making an examination under this section, the ~~commissioner~~ office may retain attorneys, appraisers, independent economists, independent certified public accountants, or other professionals and specialists as designees. A certified public accountant retained by the ~~commissioner~~ office may not be the same certified public accountant providing the certified annual audit in subdivision 2.

(d) The ~~commissioner~~ office shall make a report of an examination conducted under this section and provide a copy to the medical cannabis manufacturer. The ~~commissioner~~ office shall then post a copy of the report on the department's website. All working papers, recorded information, documents, and copies produced by, obtained by, or disclosed to the ~~commissioner~~ office or any other person in the course of an examination, other than the information contained in any ~~commissioner~~ office official report, made under this section are private data on individuals or nonpublic data, as defined in section 13.02.

Sec. 23. Minnesota Statutes 2024, section 342.01, subdivision 9, is amended to read:

Subd. 9. **Bona fide labor organization.** "Bona fide labor organization" means a labor union that represents or is actively seeking to represent ~~cannabis~~ workers~~.~~ of:

(1) a cannabis business; or

(2) a lower-potency hemp edible manufacturer.

Sec. 24. Minnesota Statutes 2024, section 342.01, subdivision 47, is amended to read:

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. 25. Minnesota Statutes 2024, section 342.01, subdivision 54, is amended to read:

Subd. 54. **Medical cannabis flower.** "Medical cannabis flower" means cannabis flower provided to a patient enrolled in the registry program ~~or a visiting patient~~; a registered designated caregiver; or a parent, legal guardian, or spouse of an enrolled patient by a registered designated caregiver, cannabis retailer, or cannabis business with a medical cannabis retail endorsement to treat or alleviate the symptoms of a qualifying medical condition. Medical cannabis flower does not include adult-use cannabis flower.

Sec. 26. Minnesota Statutes 2024, section 342.01, is amended by adding a subdivision to read:

Subd. 54a. **Medical cannabis paraphernalia.** "Medical cannabis paraphernalia" means a delivery device, related supply, or educational material used by a patient enrolled in the registry program to administer medical cannabis and medical cannabinoid products.

Sec. 27. Minnesota Statutes 2024, section 342.02, subdivision 3, is amended to read:

Subd. 3. **Medical cannabis program.** (a) The powers and duties of the Department of Health with respect to the medical cannabis program under Minnesota Statutes 2022, sections 152.22 to 152.37, are transferred to the Office of Cannabis Management under section 15.039.

(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 altered as a result of the transfer;

(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.

~~(c) This subdivision is effective July 1, 2024.~~

Sec. 28. Minnesota Statutes 2024, section 342.12, is amended to read:

342.12 LICENSES; TRANSFERS; ADJUSTMENTS.

(a) Licenses issued under this chapter that are available to all applicants pursuant to section 342.14, subdivision 1b, paragraph (c), may be freely transferred subject to the prior written approval of the office unless the license holder has not received a final site inspection 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 Division of Social Equity and is subject to the prior written approval of the office.

(c) Preliminary license ~~preapproval~~ approval issued pursuant to section ~~342.125~~ 342.14, subdivision 5, may not be transferred.

(d) A new license must be obtained when:

(1) the form of the licensee's legal business structure converts or changes to a different type of legal business structure; or

(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.

(e) Licenses must be renewed annually.

(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. 29. Minnesota Statutes 2024, section 342.14, subdivision 3, is amended to read:

Subd. 3. **Review.** (a) After an applicant submits an application that contains all required information and pays the applicable ~~licensing~~ application fee, the office must review the application.

(b) The office may deny an application if:

(1) the application is incomplete;

(2) the application contains a materially false statement about the applicant or omits information required under subdivision 1;

(3) the applicant does not meet the qualifications under section 342.16;

(4) the applicant is prohibited from holding the license under section 342.18, subdivision 2;

(5) the application does not meet the minimum requirements under section 342.18, subdivision 3;

(6) the applicant fails to pay the applicable application fee;

(7) the application was not submitted by the application deadline;

(8) the applicant submitted more than one application for a license type; or

(9) the office determines that the applicant would be prohibited from holding a license for any other reason.

(c) If the office denies an application, the office must notify the applicant of the denial and the basis for the denial.

(d) The office may request additional information from any applicant if the office determines that the information is necessary to review or process the application. If the applicant does not provide the additional requested information within 14 calendar days of the office's request for information, the office may deny the application.

(e) An applicant whose application is not denied under this subdivision is a qualified applicant.

Sec. 30. Minnesota Statutes 2024, section 342.14, subdivision 6, is amended to read:

Subd. 6. **Completed application; final authorization; issuance of license.** (a) Within 18 months of receiving notice of preliminary license approval, an applicant must provide:

(1) the address and legal property description of the location where the business will operate;

(2) the name of the local unit of government where the business will be located; and

(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.

(b) Upon receipt of the information required under paragraph (a) from an applicant that has received preliminary license approval, the office must:

(1) forward a copy of the application to the local unit of government in which the business operates or intends to operate with a form for certification as to whether a proposed cannabis business complies with local zoning ordinances and, if applicable, whether the proposed business complies with the state fire code and building code;

(2) schedule a site inspection; and

(3) require the applicant to pay the applicable license fee.

(c) The office may deny final authorization if:

(1) an applicant fails to submit any required information;

(2) the applicant submits a materially false statement about the applicant or fails to provide any required information;

(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;

(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 would operate in violation of the provisions of this chapter.

(d) Within 90 days of receiving the information required under paragraph (a) and the results of any required background check, the office shall grant final authorization and issue

the appropriate license or send the applicant a notice of rejection setting forth specific reasons that the office did not approve the application.

Sec. 31. Minnesota Statutes 2024, section 342.151, subdivision 2, is amended to read:

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.

Sec. 32. 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 individual as a cannabis worker if the individual has been convicted of any of the following crimes that would constitute a felony:

(1) human trafficking;

(2) noncannabis controlled substance crimes in the first or second degree;

(3) labor trafficking;

(4) fraud;

(5) embezzlement;

(6) extortion;

(7) money laundering; or

(8) insider trading;

if committed in this state or any other jurisdiction for which a full pardon or similar relief has not been granted.

(b) A ~~license holder~~ cannabis business must not employ an individual as a cannabis worker if the individual made any false statement in an application for employment.

Sec. 33. Minnesota Statutes 2024, section 342.22, subdivision 3, is amended to read:

Subd. 3. **Issuance of registration.** (a) A local unit of government shall issue a retail registration to a cannabis microbusiness with a retail operations endorsement, cannabis mezzobusiness with a retail operations endorsement, cannabis retailer, medical cannabis combination business operating a retail location, or lower-potency hemp edible retailer that:

(1) has a valid license or preliminary license ~~preapproval~~ approval issued by the office;

(2) has paid the registration fee or renewal fee pursuant to subdivision 2;

(3) is found to be in compliance with the requirements of this chapter at any preliminary compliance check that the local unit of government performs; and

(4) if applicable, is current on all property taxes and assessments at the location where the retail establishment is located.

(b) Before issuing a retail registration, the local unit of government may conduct a preliminary compliance check to ensure that the cannabis business or hemp business is in compliance with any applicable local ordinance established pursuant to section 342.13.

(c) A local unit of government shall renew the retail registration of a cannabis business or hemp business when the office renews the license of the cannabis business or hemp business.

(d) A retail registration issued under this section may not be transferred.

Sec. 34. Minnesota Statutes 2024, section 342.28, subdivision 1, is amended to read:

Subdivision 1. **Authorized actions.** A cannabis microbusiness 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:

(1) grow cannabis plants from seed or immature plant to mature plant and harvest cannabis flower from a mature plant;

(2) make cannabis concentrate;

(3) make hemp concentrate, including hemp concentrate with a delta-9 tetrahydrocannabinol concentration of more than 0.3 percent as measured by weight;

(4) manufacture artificially derived cannabinoids;

(5) manufacture adult-use cannabis products, lower-potency hemp edibles, and hemp-derived consumer products for public consumption;

(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, a cannabis cultivator, 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.

Sec. 35. 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. 36. 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:

(1) grow cannabis plants from seed or immature plant to mature plant and harvest cannabis flower from a mature plant for use as adult-use cannabis flower or for use in adult-use cannabis products;

(2) grow cannabis plants from seed or immature plant to mature plant and harvest cannabis flower from a mature plant for use as medical cannabis flower or for use in medical cannabinoid products;

(3) make cannabis concentrate;

(4) make hemp concentrate, including hemp concentrate with a delta-9 tetrahydrocannabinol concentration of more than 0.3 percent as measured by weight;

(5) manufacture artificially derived cannabinoids;

(6) manufacture adult-use cannabis products, lower-potency hemp edibles, and hemp-derived consumer products for public consumption;

(7) process medical cannabinoid products;

(8) purchase immature cannabis plants and seedlings ~~and~~, cannabis flower, cannabis products, lower-potency hemp edibles, and hemp-derived consumer products from a cannabis microbusiness, another cannabis mezzobusiness, a cannabis cultivator, a cannabis manufacturer, ~~or~~ a cannabis wholesaler, or a lower-potency hemp edible manufacturer;

(9) purchase cannabis concentrate, hemp concentrate, and ~~synthetically~~ artificially derived cannabinoids from a cannabis microbusiness, another 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) purchase hemp plant parts and propagules from a licensed hemp grower licensed under chapter 18K;

(11) purchase hemp concentrate from an industrial hemp processor licensed under chapter 18K;

(12) package and label adult-use cannabis flower, adult-use cannabis products, lower-potency hemp edibles, and hemp-derived consumer products for sale to customers;

(13) 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; and

(14) perform other actions approved by the office.

Sec. 37. Minnesota Statutes 2024, section 342.29, subdivision 7, is amended to read:

Subd. 7. **Production of ~~customer~~ consumer products endorsement.** A cannabis mezzobusiness 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. 38. Minnesota Statutes 2024, section 342.30, subdivision 1, is amended to read:

Subdivision 1. **Authorized actions.** A cannabis cultivator license entitles the license holder to:

(1) grow cannabis plants within the approved amount of space from seed or immature plant to mature plant~~,~~;

(2) harvest cannabis flower from a mature plant~~,~~;

(3) package and label immature cannabis plants and seedlings and cannabis flower for sale to other cannabis businesses~~,~~;

(4) sell immature cannabis plants and seedlings and cannabis flower to other cannabis businesses;

(5) transport cannabis flower to a cannabis manufacturer located on the same premises~~,~~; and

(6) perform other actions approved by the office.

Sec. 39. Minnesota Statutes 2024, section 342.33, subdivision 1, is amended to read:

Subdivision 1. **Authorized actions.** A cannabis wholesaler license entitles the license holder to:

(1) purchase immature cannabis plants and seedlings, cannabis flower, cannabis products, lower-potency hemp edibles, and hemp-derived consumer products from cannabis microbusinesses, cannabis mezzobusinesses, cannabis cultivators, cannabis manufacturers, and ~~cannabis microbusinesses~~ lower-potency hemp edible manufacturers;

(2) purchase hemp plant parts and propagules from industrial hemp growers licensed under chapter 18K;

(3) purchase hemp concentrate from an industrial hemp processor licensed under chapter 18K;

(4) sell immature cannabis plants and seedlings, cannabis flower, cannabis products, lower-potency hemp edibles, and hemp-derived consumer products to cannabis microbusinesses, cannabis mezzobusinesses, cannabis manufacturers, and cannabis retailers;

(5) sell lower-potency hemp edibles to lower-potency hemp edible retailers;

(6) import hemp-derived consumer products and lower-potency hemp edibles that contain hemp concentrate or artificially derived cannabinoids that are derived from hemp plants or hemp plant parts; and

(7) perform other actions approved by the office.

Sec. 40. Minnesota Statutes 2024, section 342.44, subdivision 1, is amended to read:

Subdivision 1. **Application; contents.** (a) Except as otherwise provided in this subdivision, the provisions of this chapter relating to license applications, license selection criteria, general ownership disqualifications and requirements, and general operational requirements do not apply to hemp businesses.

(b) The office~~, by rule,~~ shall establish forms and procedures for the processing of hemp licenses issued under this chapter. At a minimum, any application to obtain or renew a hemp license shall include the following information, if applicable:

(1) the name, address, and date of birth of the applicant;

(2) the address and legal property description of the business;

(3) proof of trade name registration;

(4) certification that the applicant will comply with the requirements of this chapter relating to the ownership and operation of a hemp business;

(5) identification of one or more controlling persons or managerial employees as agents who shall be responsible for dealing with the office on all matters; and

(6) a statement that the applicant agrees to respond to the office's supplemental requests for information.

(c) An applicant for a lower-potency hemp edible manufacturer license must submit an attestation signed by a bona fide labor organization stating that the applicant has entered into a labor peace agreement.

(d) An application on behalf of a corporation or association shall be signed by at least two officers or managing agents of that entity.

Sec. 41. Minnesota Statutes 2024, section 342.46, subdivision 6, is amended to read:

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;

(2) do not contain more than a combined total of 0.5 milligrams of all other cannabinoids per serving; and

(3) do not contain an artificially derived cannabinoid other than delta-9 tetrahydrocannabinol.

(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 servings per container.~~ If the lower-potency hemp edible is meant to be consumed as a beverage, the beverage container must not contain more than two servings.

(c) Notwithstanding paragraph (b), any edible cannabinoid products that are intended to be combined with food or beverage products before consumption must indicate a single serving using one of the following methods:

(1) the product is packaged in individual servings;

(2) the product indicates a single serving by scoring or use of another indicator that appears on the product; or

(3) the product is sold with a calibrated dropper, measuring spoon, or similar device for measuring a single serving.

~~(c)~~ (d) A single package containing multiple servings of a lower-potency hemp edible must contain no more than 50 milligrams of delta-9 tetrahydrocannabinol, 250 milligrams of cannabidiol, 250 milligrams of cannabigerol, or any combination of those cannabinoids that does not exceed the identified amounts.

Sec. 42. Minnesota Statutes 2024, section 342.52, is amended by adding a subdivision to read:

Subd. 7a. **Allowable delivery methods.** A patient in the registry program may receive medical cannabis flower and medical cannabinoid products. The office may approve additional delivery methods to expand the types of products that qualify as medical cannabinoid products.

Sec. 43. Minnesota Statutes 2024, section 342.57, subdivision 2, is amended to read:

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

(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~~ 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, 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 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 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.

(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 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:

(1) does not constitute probable cause or reasonable suspicion;

(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

(3) must not subject the person or the property of the person to inspection by any government agency.

Sec. 44. 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. 45. 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 testing. Disclosure must be made to the cannabis testing facility and must include information about all applications by any person, whether intentional or accidental.

(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. 46. 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;

(3) the batch number;

(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;

(8) the following statement: "Keep this product out of reach of children."; and

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

Sec. 47. Minnesota Statutes 2024, section 342.63, subdivision 3, is amended to read:

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;

(5) the batch number;

(6) the serving size;

(7) the cannabinoid profile per serving and in total;

(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;

(10) a warning symbol developed by the office in consultation with the commissioner of health and the Minnesota Poison Control System that:

(i) is at least three-quarters of an inch tall and six-tenths of an inch wide;

(ii) is in a highly visible color;

(iii) includes a visual element that is commonly understood to mean a person should stop;

(iv) indicates that the product is not for children; and

(v) includes the phone number of the Minnesota Poison Control System;

(11) verification that the cannabis product, lower-potency hemp edible, hemp-derived consumer product, or medical cannabinoid product was tested according to section 342.61

and that the cannabis product, lower-potency hemp edible, hemp-derived consumer product, or medical cannabinoid product complies with the applicable standards;

(12) information on the usage of the product;

(13) the following statement: "Keep this product out of reach of children."; and

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

(b) The office may by rule establish alternative labeling requirements for lower-potency hemp edibles that are imported into the state if those requirements provide consumers with information that is substantially similar to the information described in paragraph (a).

Sec. 48. 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 ~~by:~~;

~~(1)~~ (2) posting the information in the premises of the cannabis microbusiness, cannabis mezzobusiness, cannabis retailer, or medical cannabis combination business; or

~~(2)~~ (3) providing the information on a separate document or pamphlet provided to customers or patients when the customer purchases cannabis flower, a cannabis product, a lower-potency hemp edible, or a hemp-derived consumer product.

Sec. 49. **REPEALER.**

Minnesota Statutes 2024, sections 152.22, subdivision 2; 342.01, subdivision 71; and 342.151, subdivision 1, are repealed.

152.22 DEFINITIONS.

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

342.01 DEFINITIONS.

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.

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.