REVISOR

23-02009

This Document can be made available in alternative formats upon request

State of Minnesota

HOUSE OF REPRESENTATIVES H. F. No. 773

NINETY-THIRD SESSION

01/25/2023

Authored by Edelson, Curran and Tabke The bill was read for the first time and referred to the Committee on Health Finance and Policy

1.1	A bill for an act
1.2	relating to cannabis; providing for the regulation of the concentration and
1.3	conversion of cannabinoids derived from hemp; providing for regulation of the
1.4	transportation and testing of concentrated cannabinoids and artificially derived
1.5	cannabinoids derived from hemp; providing for the licensing of edible cannabinoid
1.6	product manufacturers and distributors; providing for regulation of the
1.7	transportation, testing, and labeling of hemp-derived consumer products and edible
1.8	cannabinoid products; providing for the regulation of the sale of hemp-derived
1.9	consumer products and edible cannabinoid products; providing for enforcement
1.10	of regulations; establishing guidelines for local licensing of certain retailers;
1.11	establishing a gross receipts tax on edible cannabinoid products; establishing
1.12	criminal penalties; authorizing exclusive liquor stores to sell certain products;
1.13	requiring reports; appropriating money; amending Minnesota Statutes 2022, sections
1.14	13.3806, by adding a subdivision; 18K.02, subdivision 5, by adding subdivisions;
1.15	18K.03, by adding a subdivision; 18K.04, subdivisions 1, 4, by adding a
1.16	subdivision; 18K.06; 34A.01, subdivision 4; 144.99, subdivision 1; 152.027, by
1.17	adding a subdivision; 181.938, subdivision 2; 297A.99, by adding a subdivision;
1.18	340A.412, subdivision 14; proposing coding for new law in Minnesota Statutes,
1.19	chapters 18K; 152; 295; repealing Minnesota Statutes 2022, section 151.72.
1.20	BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF MINNESOTA:
1.21	ARTICLE 1
1.22	AGRICULTURE POLICY
1.23	Section 1. Minnesota Statutes 2022, section 18K.02, is amended by adding a subdivision
1.23	Section 1. Winnesota Statutes 2022, section 16K.02, is amended by adding a subdivision
1.24	to read:
1.25	Subd. 3a. Artificially derived cannabinoid. "Artificially derived cannabinoid" means
1.26	a cannabinoid extracted from hemp plants or hemp plant parts with a chemical makeup that
1.27	is changed after extraction to create a different cannabinoid or other chemical compound
1.28	by applying a catalyst other than heat or light. Artificially derived cannabinoid includes but
1.29	is not limited to any tetrahydrocannabinol created from cannabidiol. Artificially derived

	01/20/23	REVISOR	BD/HL	23-02009
2.1	cannabinoid does not include a product c	ontaining cannab	pinoids as defined in sec	ction 152.50,
2.2	subdivision 15.			
2.3	Sec. 2. Minnesota Statutes 2022, secti	on 18K.02, is an	nended by adding a su	bdivision to
2.4	read:			
2.5	Subd. 3b. Concentrated cannabinoi	d. "Concentrated	l cannabinoid" means a	cannabinoid
2.6	extracted from hemp plants or hemp pla	nt parts that is eit	ther isolated from othe	r substances
2.7	and exists in a pure form or is present in	a mixture in an a	amount greater than the	e percentage
2.8	that naturally occurs in the hemp plant o	r hemp plant par	ts. Concentrated canna	abinoid does
2.9	not include a product containing cannab	inoids as defined	l in section 152.50, sub	odivision 15.
2 10	Sec. 3. Minnesota Statutes 2022, secti	on 18K 02 is an	nended by adding a su	bdivision to
2.10 2.11	read:	011 10 K .02, 18 all	nended by adding a su	
2.11				
2.12	Subd. 3c. Conversion of cannabing			
2.13	cannabinoids" or "convert cannabinoids	" means the proc	cess of creating artific	ally derived
2.14	cannabinoids.			
2.15	Sec. 4. Minnesota Statutes 2022, secti	on 18K.02, subd	livision 5, is amended	to read:
2.16	Subd. 5. Processing. "Processing" m	neans rendering b	by refinement hemp pla	ants or hemp
2.17	plant parts from their natural or original	state after harve	est. Processing include	s but is not
2.18	limited to decortication, devitalization,	chopping, crushi	ng, extraction, <u>conver</u>	sion of
2.19	cannabinoids, and packaging. Processin	g does not inclu	de typical farm operat	ions such as
2.20	sorting, grading, baling, and harvesting.			
		1012 02 :	1 11 11	1 1
2.21	Sec. 5. Minnesota Statutes 2022, secti	on 18K.03, 18 an	nended by adding a su	bdivision to
2.22	read:			
2.23	Subd. 3. Possession of concentrate	d cannabinoids	or artificially derived	<u>d</u>
2.24	cannabinoids. (a) Notwithstanding the	provisions of ch	apter 152 or any other	law to the
2.25	contrary, a licensee may possess and tra	nsport concentra	ated cannabinoids or a	rtificially
2.26	derived cannabinoids provided that the	licensee:		
2.27	(1) is authorized to concentrate or $concentrate$	onvert cannabing	oids;	
2.28	(2) complies with an approved plan	to secure, store,	and dispose of concen	trated
2.29	cannabinoids or artificially derived can	nabinoids;		

BD/HL

3.1	(3) complies with section 18K.045 and any applicable rules regarding the transportation
3.2	of concentrated cannabinoids or artificially derived cannabinoids; and
3.3	(4) complies with any additional requirements or rules adopted by the commissioner.
3.4	(b) Notwithstanding the provisions of chapter 152 or any other law to the contrary, an
3.5	approved laboratory may possess concentrated cannabinoids or artificially derived
3.6	cannabinoids during the period that the laboratory is approved by the commissioner to
3.7	perform testing on concentrated cannabinoids or artificially derived cannabinoids and the
3.8	laboratory maintains any required accreditation.
3.9	(c) A licensee or laboratory that possesses concentrated cannabinoids or artificially
3.10	derived cannabinoids in violation of this subdivision may be subject to any applicable
3.11	licensing penalty, criminal penalty, or both.
3.12	Sec. 6. Minnesota Statutes 2022, section 18K.04, subdivision 1, is amended to read:
3.13	Subdivision 1. Requirement; issuance; presumption. (a) A person must obtain a license
3.14	from the commissioner before (1) growing industrial hemp for commercial or research
3.15	purposes, and (2) before processing industrial hemp for commercial purposes.
3.16	(b) To obtain a license under paragraph (a), a person must apply to the commissioner
3.17	in the form prescribed by the commissioner and must pay the annual registration and
3.18	inspection fee established by the commissioner in accordance with section 16A.1285,
3.19	subdivision 2.
3.20	(c) For a license to grow industrial hemp for commercial or research purposes, the license
3.21	application must include the name and address of the applicant and the legal description of
3.22	the land area or areas where industrial hemp will be grown by the applicant and any other
3.23	information required under Code of Federal Regulations, title 7, part 990.
3.24	(d) For a license to process industrial hemp for commercial purposes, the license
3.25	application must include the name and address of the applicant, the legal description of the
3.26	processing location, whether the applicant intends to concentrate cannabinoids or convert
3.27	cannabinoids into any other type of cannabinoid or other chemical compound, and any other
3.28	information required by the commissioner.
3.29	(e) A licensee is responsible for compliance with the license requirements irrespective
3.30	of the acts or omissions of an authorized representative acting on behalf of the licensee.

23-02009

- (f) When an applicant has paid the fee and completed the application process to the 4.1 satisfaction of the commissioner, the commissioner must issue a license which is valid until 4.2 4.3 December 31 of the year of application. (g) A person licensed under paragraph (a) to grow industrial hemp is presumed to be 4.4 4.5 growing industrial hemp for commercial or research purposes. (h) The commissioner may issue an applicant a full license to grow and process industrial 4.6 hemp, including an authorization to concentrate cannabinoids or convert cannabinoids, or 4.7 may issue an applicant a partial license that permits only certain specified actions. 4.8 Sec. 7. Minnesota Statutes 2022, section 18K.04, is amended by adding a subdivision to 4.9 4.10 read: 4.11 Subd. 1a. Concentration or conversion of cannabinoids. (a) An applicant or a licensee must notify the commissioner if the applicant or licensee intends to concentrate cannabinoids 4.12 or convert extracted cannabinoids into any other type of cannabinoid or other chemical 4.13 compound, including but not limited to the concentration of any tetrahydrocannabinol or 4.14 conversion of cannabidiol into any tetrahydrocannabinol. A licensee may not concentrate 4.15 or convert cannabinoids without the express permission of the commissioner. 4.16 (b) An applicant or a licensee seeking permission to convert cannabinoids must disclose: 4.17 (1) the method of conversion that will be used, including any specific catalysts that will 4.18 be employed; 4.19 (2) the manner in which artificially derived cannabinoids will be secured and stored; 4.20 (3) the molecular nomenclature of all cannabinoids or other chemical compound that 4.21 will be created; 4.22 (4) the amount of each cannabinoid or other chemical compound that the applicant or 4.23 licensee expects to hold in storage at any time; 4.24 (5) a plan for the disposal and destruction of any waste products generated in the 4.25 4.26 conversion process, including any tetrahydrocannabinol or other artificially derived cannabinoids that are contaminated or that will not be used for commercial or research 4.27 4.28 purposes; and (6) any other information required by the commissioner. 4.29 (c) An applicant or a licensee may identify information provided pursuant to paragraph 4.30 (b), clause (1), as a trade secret. 4.31
 - Article 1 Sec. 7.

	01/20/23	REVISOR	BD/HL	23-02009
5.1	(d) An applicant or licensee seeking	permission to concer	ntrate cannabinoids	must
5.2	disclose:			
5.3	(1) the manner in which concentrated	d cannabinoids will b	be secured and store	ed;
5.4	(2) the amount of each cannabinoid t	hat the licensee expe	ects to hold in storag	ge at any
5.5	time; and			
5.6	(3) any other information required by	y the commissioner.		
5.7	(e) On a schedule and in the form and	manner established b	y the commissioner,	a licensee
5.8	must notify the commissioner of:			
5.9	(1) the amount of each cannabinoid of	or other chemical con	mpound that the lice	ensee
5.10	concentrated or created during the repor	ting period;		
5.11	(2) the amount of each concentrated	cannabinoid or artifi	cially derived canna	abinoid
5.12	being stored by the licensee; and			
5.13	(3) the amount of concentrated cannal	pinoid or artificially d	lerived cannabinoid,	, including
5.14	any tetrahydrocannabinol, that the licens	see disposed of or de	stroyed.	
5.15	Sec. 8. Minnesota Statutes 2022, section	on 18K.04, subdivisi	on 4, is amended to	read:
5.16	Subd. 4. Industrial hemp licensing of	lata classification. (a) In addition to data	classified
5.17	pursuant to section 13.41, the following	data collected, creat	ed, or maintained by	y the
5.18	commissioner under this chapter is class	ified as private data,	as defined in section	on 13.02,
5.19	subdivision 12, or nonpublic data, as de	fined in section 13.02	2, subdivision 9:	
5.20	(1) nondesignated addresses provide	d by licensees and a	oplicants; and	
5.21	(2) data that identify the specific loca	ations where licensee	es and applicants gro	ow or
5.22	process, or will grow or process, industr	ial hemp, including	but not limited to le	gal
5.23	descriptions, street addresses, geospatial	locations, maps, and	d property boundari	es and
5.24	dimensions; and			
5.25	(3) information provided pursuant to	subdivision 1a that	is identified as a tra-	de secret,
5.26	except that the commissioner is authoriz	ed to provide a list c	of all catalysts identi	ified as
5.27	being used to convert cannabinoids to app	proved independent la	aboratories in order to	o facilitate
5.28	testing.			
5.29	(b) The commissioner may disclose	data classified as priv	vate data or nonpub	lic data
5.30	under this subdivision if the commission	ner determines that th	nere is a substantive	threat to
5.31	human health or safety or to the environ	ment, or to aid in the	e law enforcement p	rocess.

6.1	Sec. 9. [18K.045] TRANSPORTATION.
6.2	(a) A licensee or any other person transporting industrial hemp in a form other than as
6.3	a concentrated cannabinoid or artificially derived cannabinoid must comply with rules
6.4	adopted by the commissioner.
6.5	(b) A licensee authorized to concentrate cannabinoids or convert cannabinoids may
6.6	transport concentrated cannabinoids or artificially derived cannabinoids on public roadways
6.7	provided that:
6.8	(1) all concentrated cannabinoids or artificially derived cannabinoids are packaged in
6.9	tamper-evident containers that are not visible or recognizable from outside the transporting
6.10	vehicle;
6.11	(2) the licensee has a shipping manifest in the licensee's possession that describes the
6.12	contents of all tamper-evident containers, discloses all concentrated cannabinoids or
6.13	artificially derived cannabinoids, and identifies all other items being transported;
6.14	(3) the transporting vehicle does not bear any markings to indicate that the vehicle
6.15	contains industrial hemp, cannabinoids, or any other form of cannabis and does not bear
6.16	the name or logo of the licensee;
6.17	(4) all departures, arrivals, and stops are appropriately documented;
6.18	(5) at least two designated employees staff any vehicle used to transport concentrated
6.19	cannabinoids or artificially derived cannabinoids and at least one employee remains with
6.20	the vehicle at all times that the vehicle is transporting concentrated cannabinoids or artificially
6.21	derived cannabinoids;
6.22	(6) no person other than a designated employee enters a vehicle at any time that the
6.23	vehicle is transporting concentrated cannabinoids or artificially derived cannabinoids except
6.24	that a licensed manufacturer or approved laboratory receiving a shipment may assist in
6.25	unloading concentrated cannabinoids or artificially derived cannabinoids from a vehicle;
6.26	and
6.27	(7) the licensee complies with any other rules adopted by the commissioner regarding
6.28	the transportation of concentrated cannabinoids or artificially derived cannabinoids.
6.29	(c) Notwithstanding section 221.025, transportation of concentrated cannabinoids or
6.30	artificially derived cannabinoids by a person or entity other than a licensee may only be
6.31	performed by a motor carrier of property that is registered with the commissioner of
6.32	transportation. A motor carrier of property that is transporting concentrated cannabinoids
6.33	or artificially derived cannabinoids must comply with the requirements of paragraph (b).

7.1	Sec. 10. [18K.046] TESTING.
7.2	(a) Testing of industrial hemp other than a concentrated cannabinoid or an artificially
7.3	derived cannabinoid must comply with rules adopted by the commissioner.
7.4	(b) Testing of a concentrated cannabinoid or an artificially derived cannabinoid must
7.5	comply with rules adopted by the commissioner and, at a minimum, must:
7.6	(1) identify contaminants, including residual solvents, foreign material, microbiological
7.7	contaminants, heavy metals, pesticide residue, and mycotoxins; and
7.8	(2) provide a cannabinoid profile that identifies and quantifies the cannabinoids in a
7.9	testing sample.
7.10	(c) The commissioner may require that testing is performed by an independent laboratory
7.11	and shall establish a process for laboratory approval. At a minimum, a laboratory must
7.12	operate formal management systems under the International Organization for Standardization
7.13	to qualify for approval.
7.14	(d) A licensee must disclose all known information regarding pesticides, fertilizers,
7.15	solvents, or other foreign materials applied to industrial hemp or added to industrial hemp
7.16	during any production or processing stages. Disclosure must be made to any entity performing
7.17	testing or sampling and, upon request, to the commissioner. Disclosure must include all
7.18	information known to the licensee regardless of whether the application or addition was
7.19	made intentionally or accidentally or by the licensee or any other person or entity.
7.20	(e) A licensee must allow a sampling agent, the commissioner, or the commissioner's
7.21	designee to collect regulatory samples of industrial hemp, including concentrated
7.22	cannabinoids and artificially derived cannabinoids.
7.23	(f) The commissioner shall consult with the commissioner of health to determine the
7.24	contaminants that must be identified in testing and to establish standards for allowable levels
7.25	of contaminants in concentrated cannabinoids and artificially derived cannabinoids.
7.26	Sec. 11. [18K.047] MONITORING SYSTEM.
7.27	(a) The commissioner shall coordinate with the commissioner of health to identify an
7.28	approved monitoring system for the integrated tracking, inventory, and verification of
7.29	industrial hemp, including concentrated cannabinoids, as defined in section 18K.02,
7.30	subdivision 3b; artificially derived cannabinoids, as defined in section 18K.02, subdivision
7.31	3a; and edible cannabinoid products, as defined in section 152.50, subdivision 7.

23-02009

8.1	(b) A licensee must use the monitoring system to track all industrial hemp, including
8.2	concentrated cannabinoids and artificially derived cannabinoids, and edible cannabinoid
8.3	products in the licensee's possession, to the point of disposal, transfer, or sale. For the
8.4	purposes of this section, a licensee possesses the industrial hemp, including concentrated
8.5	cannabinoids and artificially derived cannabinoids, and edible cannabinoid products that
8.6	the licensee cultivates from seed or immature plant, converts from any other cannabinoid,
8.7	receives from another licensee, manufactures, or receives from an entity licensed to
8.8	manufacture or distribute products containing cannabinoids derived from industrial hemp.
8.9	This paragraph does not apply to products lawfully purchased by a licensee for personal
8.10	use.
8.11	Sec. 12. Minnesota Statutes 2022, section 18K.06, is amended to read:
8.12	18K.06 RULEMAKING.
8.13	(a) The commissioner shall adopt rules governing the production, testing, processing,
8.14	and licensing of industrial hemp. Notwithstanding section 14.125, the commissioner's
8.15	authority to adopt these rules expires June 30, 2022.
9 16	(b) Rules adopted under paragraph (a) must include, but not be limited to, provisions
8.16 8.17	governing:
0.17	governing.
8.18	(1) the supervision and inspection of industrial hemp during its growth and harvest;
8.19	(2) the testing of industrial hemp to determine delta-9 tetrahydrocannabinol levels;
8.20	(3) the use of background check results required under section 18K.04 to approve or
8.21	deny a license application; and
8.22	(4) any other provision or procedure necessary to carry out the purposes of this chapter.
8.23	(c) Rules issued under this section must be consistent with federal law regarding the
8.24	production, distribution, and sale of industrial hemp.
8.25	(d) The commissioner may adopt rules governing artificially derived cannabinoids,
8.26	including the methods of conversion and the storage, transportation, testing, monitoring,
8.27	and disposal of artificially derived cannabinoids. Rules governing artificially derived
8.28	cannabinoids for which notice is published in the State Register before July 1, 2024, may
8.29	be adopted using the expedited rulemaking process in section 14.389.

01/20/23REVISOR **BD**/HL 23-02009 **ARTICLE 2** 9.1 **HEALTH POLICY** 9.2 Section 1. [152.50] DEFINITIONS. 9.3 Subdivision 1. Applicability. For the purposes of sections 152.50 to 152.65, the terms 9.4 defined in this section have the meanings given. 9.5 Subd. 2. Artificially derived cannabinoid. "Artificially derived cannabinoid" has the 9.6 meaning given in section 18K.02, subdivision 3a. 9.7 Subd. 3. Batch. "Batch" means a specific quantity of a specific edible cannabinoid 9.8 product that is manufactured by an edible cannabinoid product manufacturer at the same 9.9 time and using the same methods, equipment, and ingredients. A batch is uniform and 9.10 intended to meet specifications for identity, strength, purity, and composition, and is 9.11 manufactured, numbered, and stored according to a single batch production record executed 9.12 and documented during the same cycle of manufacture and produced by a continuous 9.13 9.14 process. 9.15 Subd. 4. Batch number. "Batch number" means a unique numeric or alphanumeric identifier assigned to a batch of edible cannabinoid products by an edible cannabinoid 9.16 product manufacturer. 9.17 Subd. 5. Commissioner. "Commissioner" means the commissioner of health. 9.18 Subd. 6. Concentrated cannabinoid. "Concentrated cannabinoid" means a cannabinoid 9.19 extracted from hemp plants or hemp plant parts that is either isolated from other substances 9.20 and exists in a pure form or is present in a mixture in an amount greater than the percentage 9.21 that naturally occurs in the hemp plant or hemp plant parts. Concentrated cannabinoid does 9.22 not include a product containing cannabinoids. 9.23 Subd. 7. Edible cannabinoid product. "Edible cannabinoid product" means any product 9.24 that is: 9.25 (1) intended to be eaten or consumed as a beverage by humans; 9.26 (2) contains a cannabinoid in combination with food ingredients or is intended or 9.27 generally expected to be added to food ingredients; and 9.28 (3) is not a drug or medical cannabis product. 9.29 Subd. 8. Hemp-derived consumer product. "Hemp-derived consumer product" means 9.30 any product intended for human or animal consumption that is harvested from hemp plants 9.31 or hemp plant parts or contains cannabinoids extracted from hemp plants or hemp plant 9.32

	01/20/23	REVISOR	BD/HL	23-02009
10.1	parts. Hemp-derived consumer produc	t does not include e	dible cannabinoid p	products or
10.2	products for which the United States F	ood and Drug Adm	inistration has issue	ed generally
10.3	recognized as safe notices.			
10.4	Subd. 9. Hemp plants or hemp pla	nt parts. "Hemp pla	nts" or "hemp plant	parts" means
10.5	any part of the industrial hemp plant, w	whether growing or	not, including the s	talk, leaves,
10.6	buds, and seeds, but does not include ar	ny derivatives, extrac	ets, cannabinoids, is	omers, acids,
10.7	salts, and salts of isomers that are sepa	rated from the plant		
10.8	Subd. 10. Industrial hemp. "Indus	trial hemp" has the n	neaning given in sec	ction 18K.02,
10.9	subdivision 3.			
10.10	Subd. 11. Label. "Label" means a c	display of written, p	rinted, or graphic n	natter upon
10.11	the immediate container of any produc	t that contains a can	nabinoid.	
10.12	Subd. 12. Local unit of governme	nt. "Local unit of go	overnment" means	a home rule
10.13	charter or statutory city, county, town,	or other political su	bdivision.	
10.14	Subd. 13. Marijuana. "Marijuana"	has the meaning giv	en in section 152.01	, subdivision
10.15	<u>9.</u>			
10.16	Subd. 14. Matrix barcode. "Matrix	x barcode" means a	code that stores dat	ta in a
10.17	two-dimensional array of geometricall	y shaped dark and li	ight cells capable o	f being read
10.18	by the camera of a smartphone or othe	r mobile device.		
10.19	Subd. 15. Product containing can	nabinoids. "Product	containing cannabi	noids" means
10.20	hemp-derived consumer products and	edible cannabinoid	products.	
10.21	Sec. 2. [152.51] DUTIES OF COMM	IISSIONED INTE	PACENCY COO	PERATION
10.21	Sec. 2. [132.31] DU HES OF COMM		NAGENCICOU	
10.22	Subdivision 1. Regulation of canna	binoid products; po	owers and duties. (a	a) To promote
10.23	the public safety and welfare, ensure the	hat consumers have	access to relevant i	nformation,
10.24	and support responsible businesses, the	e commissioner shal	ll make rules, estab	lish policy,
10.25	issue licenses, and take enforcement ac	ction to regulate the	manufacturing, dist	ribution, and
10.26	sale of products containing cannabinoi	<u>ds.</u>		
10.27	(b) The commissioner shall exercise	e the following pow	vers and duties:	
10.28	(1) establish and regularly update s	tandards for produc	t testing, packaging	<u>g, marketing,</u>
10.29	and labeling;			
10.30	(2) approve cannabinoid product ty	pes and identify spe	cific cannabinoids	that products
10.31	may contain;			

	01/20/23	REVISOR	BD/HL	23-02009
11.1	(3) ensure that products are manufa	actured using meth	ods and in facilities that	at meet
11.2	appropriate health and safety standards			
11.3	(4) ensure that artificially derived c	annabinoids are st	ored and disposed of in	n a secure
11.4	manner;			
11.5	(5) ensure that products identified a	as posing a risk to	public health or safety	can be
11.6	identified and recalled;			
11.7	(6) issue and renew licenses;			
11.8	(7) prevent unauthorized access to	products containin	g cannabinoids by indi	ividuals
11.9	under 21 years of age;			
11.10	(8) impose and collect civil and add	ninistrative penalt	ies;	
11.11	(9) publish such information as may	y be deemed neces	sary for the welfare of	businesses
11.12	that manufacture, distribute, or sell car	nabinoid products	s; employees of those b	usinesses;
11.13	and the health and safety of the genera	l public;		
11.14	(10) remain informed regarding dev	velopments in laws	s, policies, and practice	es affecting
11.15	industrial hemp, marijuana, and cannal	pinoid products;		
11.16	(11) provide reports as required by	law; and		
11.17	(12) exercise other powers and authors	ority and perform o	other duties required of	or imposed
11.18	upon the commissioner by law.			
11.19	Subd. 2. Interagency agreements.	(a) The commissi	oner and the commissi	oner of
11.20	agriculture shall enter into interagency a	greements to ensur	re that edible cannabino	id products
11.21	are handled, manufactured, and inspec	ted in a manner the	at is consistent with the	e relevant
11.22	food safety requirements in chapters 2	8A, 31, and 34A, a	and associated rules.	
11.23	(b) The commissioner may coopera	te and enter into ag	greements with the com	missioners
11.24	and directors of other state agencies an	d departments to p	promote the beneficial	interests of
11.25	the state.			
11.26	Sec. 3. [152.52] RULEMAKING.			
11.27	(a) The commissioner shall adopt r	ules governing the	manufacture, distribut	tion, and
11.28	sale of products containing cannabinoi	ds.		
11.29	(b) Rules adopted under paragraph	(a) must include b	out not be limited to pro	ovisions
11.30	governing:			

BD/HL

12.1	(1) the supervision and inspection of the manufacture, distribution, and retail sale of
12.2	products containing cannabinoids;
12.3	(2) the secure storage and disposal of artificially derived cannabinoids;
12.4	(3) the testing of products containing cannabinoids;
12.5	(4) the use of background check results required to approve or deny a license application;
12.6	and
12.7	(5) any other provision or procedure necessary to carry out the purposes of sections
12.8	<u>152.50 to 152.65.</u>
12.9	(c) Rules governing the manufacture, distribution, and sale of products containing
12.10	cannabinoids for which notice is published in the State Register before July 1, 2024, may
12.11	be adopted using the expedited rulemaking process in section 14.389.
12.12	Sec. 4. [152.53] HEMP-DERIVED CONSUMER PRODUCTS.
12.13	Subdivision 1. Scope. This section applies to the manufacture, marketing, distribution,
12.14	and sale of hemp-derived consumer products.
12.15	Subd. 2. Approved cannabinoids. (a) Products manufactured, marketed, distributed,
12.16	and sold under this section may contain cannabidiol, cannabigerol, or both. Except as
12.17	provided in paragraph (c), products may not contain any other cannabinoid unless approved
12.18	by the commissioner.
12.19	(b) The commissioner may approve any cannabinoid and authorize the cannabinoid's
12.20	use in manufacturing, marketing, distribution, and sales under this section if the commissioner
12.21	determines that the cannabinoid does not impair the human central nervous system; muscles;
12.22	or audio, visual, or mental processes.
12.23	(c) Products manufactured, marketed, distributed, and sold under this section may contain
12.24	cannabinoids other than cannabidiol, cannabigerol, or any other cannabinoid approved by
12.25	the commissioner provided that the cannabinoids are naturally occurring in hemp plants or
12.26	hemp plant parts and the total of all other cannabinoids present in a product does not exceed
12.27	one milligram per package.
12.28	Subd. 3. Approved products. Products sold to consumers under this section may only
12.29	be manufactured, marketed, distributed, intended, or generally expected to be used by
12.30	applying the product externally to a part of the body of a human or animal.
12.31	Subd. 4. Testing requirements. (a) The commissioner shall establish the types of
12.32	cannabinoids and contaminants for which testing must be completed and the acceptable

23-02009

- minimum standards of any contaminant for which testing is required. The commissioner 13.1 must make testing requirement information available to the general public and any entity 13.2 13.3 manufacturing, testing, marketing, distributing, or selling products that contain cannabinoids. (b) A manufacturer of a product regulated under this section must submit representative 13.4 13.5 samples of each batch of the product to an independent, approved laboratory in order to certify that the product complies with the standards in this section and established by the 13.6 commissioner. Testing must be consistent with generally accepted industry standards for 13.7 herbal and botanical substances, and, at a minimum, the testing must determine: 13.8 (1) if the product contains the amount or percentage of cannabinoids that is stated on 13.9 13.10 the label of the product; (2) the types and amounts of cannabinoids present in a product for which testing is 13.11 required by the commissioner; 13.12 (3) if the product contains any cannabinoid, other than cannabidiol, cannabigerol, or a 13.13 cannabinoid approved by the commissioner, in an amount that exceeds the standard 13.14 established in subdivision 2, paragraph (c); 13.15 (4) if the product contains any contaminants for which testing is required by the 13.16 commissioner, including residual solvents, foreign material, microbiological contaminants, 13.17 heavy metals, pesticide residue, and mycotoxins in amounts that exceed the acceptable 13.18 minimum standards established by the commissioner; and 13.19 13.20 (5) if the product contains any tetrahydrocannabinol and, if so, if the product contains more than a total of 0.3 percent by weight of all tetrahydrocannabinols. 13.21 (c) Upon the request of the commissioner, the manufacturer of the product must provide 13.22 the commissioner with the results of the testing required in this section. 13.23 (d) A manufacturer must allow a sampling agent, the commissioner, or the commissioner's 13.24 designee to collect regulatory samples of products regulated under this section. 13.25 (e) A product whose test results are inconsistent with the product's label or show that 13.26 the product's contents do not meet the standards established in law or by the commissioner 13.27 must not be sold to consumers, except that the commissioner may allow a product to be 13.28 13.29 sold after being relabeled to be consistent with the test results if the product otherwise 13.30 complies with this section. (f) Testing of the industrial hemp from which the cannabinoid was derived, or possession 13.31 of a certificate of analysis for such industrial hemp, does not meet the testing requirements 13.32
- 13.33 of this section.

	01/20/23	REVISOR	BD/HL	23-02009
14.1	Subd. 5. Labeling. (a) A product re	egulated under this	s section must bear a l	label that
14.2	contains, at a minimum:	0		
14.3	(1) the name, address or location, co	ontact phone numbe	er, and website of the r	nanufacturer
14.4	of the product;	I		
14.5	(2) the name and address of the inc	lependent. accredit	ted laboratory used by	v the
14.6	manufacturer to test the product; and		<u></u>	<u>,</u>
14.7	(3) an accurate statement of the am	ount or percentage	of cannabinoids for v	vhich testing
14.8	is required by the commissioner found			
14.9	(b) The information in paragraph (a	a) may be provided	d on an outer package	e if the
14.10	immediate container that holds the pro-	· - ·		
14.11	(c) The information required in par	agraph (a) may be	provided through the	e use of a
14.12	scannable barcode or matrix barcode t			
14.13	that page contains all of the information			
14.14	(d) The label must also include a st	atement that the pr	oduct does not claim	to diagnose,
14.15	treat, cure, or prevent any disease, doe	s not claim that the	e product may be use	d to alter the
14.16	structure or function of human or anin	nal bodies, and has	not been evaluated o	r approved
14.17	by the United States Food and Drug Ad	ministration unless	the product has been	so approved.
14.18	The labeling must not contain any stat	ement, artwork, or	design that is incons	istent with
14.19	the required statement.			
14.20	(e) The information required by this	subdivision must	be prominently and cc	onspicuously
14.21	placed on the label or displayed on the v	vebsite in terms tha	t can be easily read and	d understood
14.22	by the consumer.			
14.23	Subd. 6. Prohibitions. (a) A produ	ct sold to consume	ers under this section	must not be
14.24	manufactured, marketed, distributed, o	or intended:		
14.25	(1) for external or internal use in the	e diagnosis, cure, n	nitigation, treatment, o	or prevention
14.26	of disease in humans or other animals;	<u>.</u>		
14.27	(2) to affect the structure or any fur	nction of the bodie	s of humans or other	animals;
14.28	(3) to be consumed by combustion	or vaporization of	the product and inha	lation of
14.29	smoke, aerosol, or vapor from the proc	luct;		
14.30	(4) to be consumed through chewin	ng, drinking, or sw	allowing; or	
14.31	(5) to be consumed through injection	on or application to	a mucous membrane	or nonintact
14.32	skin.			

Article 2 Sec. 4.

	01/20/23	REVISOR	BD/HL	23-02009
15.1	(b) A product manufactured, marke	eted, distributed, or so	ld to consumers und	ler this
15.2	section must not:			
15.3	(1) consist, in whole or in part, of a	ny filthy, putrid, or de	composed substanc	<u>e;</u>
15.4	(2) have been produced, prepared,	packed, or held under	unsanitary condition	ns where
15.5	the product may have been rendered in	jurious to health, or v	where the product ma	ay have
15.6	been contaminated with filth;			
15.7	(3) be packaged in a container that	is composed, in whol	e or in part, of any p	oisonous
15.8	or deleterious substance that may rend	er the contents of the	product injurious to	health;
15.9	(4) contain any additives or excipie	nts that have been for	and by the United St	ates Food
15.10	and Drug Administration to be unsafe	for human or animal o	consumption;	
15.11	(5) contain a cannabinoid or an am	ount or percentage of	cannabinoids that is	different
15.12	than the information stated on the labe	<u>l;</u>		
15.13	(6) contain a cannabinoid, other that	n cannabidiol, cannal	pigerol, or a cannabi	noid
15.14	approved by the commissioner, in an a	mount that exceeds th	e standard establish	ed in
15.15	subdivision 2, paragraph (c); or			
15.16	(7) contain any contaminants for w	hich testing is require	d by the commission	ner in
15.17	amounts that exceed the acceptable mi	nimum standards esta	blished by the comm	nissioner.
15.18	(c) No product containing any cann	abinoid may be sold t	o any individual wh	o is under
15.19	21 years of age.			
15.20	Subd. 7. Enforcement; penalties.	a) The commissioner	may enforce this sec	tion under
15.21	the relevant provisions of sections 144	.989 to 144.993.		
15.22	(b) Notwithstanding section 144.99	, subdivision 11, a pe	rson who commits a	ny of the
15.23	following acts regarding a product reg	ulated under this secti	on is guilty of a gro	<u>SS</u>
15.24	misdemeanor:			
15.25	(1) knowingly altering or otherwise	e falsifying testing res	<u>ults;</u>	
15.26	(2) knowingly providing false infor	mation on a product l	abel; or	
15.27	(3) intentionally making a false ma	terial statement to the	commissioner.	
15.28	(c) Any person who sells a product	regulated under this	section to a person u	nder 21
15.29	years of age, except a sale for no remune	eration, is guilty of a m	isdemeanor. It is an a	ıffirmative
15.30	defense to a charge under this paragrap	h if the defendant pro	ves by a prepondera	nce of the

BD/HL

16.1	evidence that the defendant reasonably and in good faith relied on proof of age as described
16.2	in section 152.64, subdivision 5.
16.3	Sec. 5. [152.54] LICENSING AND ENFORCEMENT; EDIBLE CANNABINOID
16.4	PRODUCTS.
16.5	Subdivision 1. Licenses; types. The commissioner shall issue the following types of
16.6	licenses:
16.7	(1) edible cannabinoid product manufacturer licenses; and
16.8	(2) edible cannabinoid product distributor licenses.
16.9	Subd. 2. Record of retail licensees. The commissioner shall maintain a list of the name,
16.10	address, trade name, license effective date, and license expiration date of any entity licensed
16.11	by a local government to sell edible cannabinoid products. The commissioner shall update
16.12	the record to reflect any license suspension, revocation, or cancellation.
16.13	Subd. 3. Fees; renewal; transfer. Licenses issued under sections 152.50 to 152.65 must
16.14	be renewed annually. Licenses may not be transferred. The commissioner may impose an
16.15	annual licensing fee that does not exceed \$500.
16.16	Subd. 4. Licensing disqualifications and requirements. (a) The commissioner shall,
16.17	by rule, establish a list of criminal offenses for which a conviction presumptively disqualifies
16.18	a person from receiving or maintaining a license to manufacture or distribute edible
16.19	cannabinoid products and may establish a time period after which a conviction may no
16.20	longer be used to presumptively disqualify a person. The commissioner must not include a
16.21	violation of chapter 152 involving the possession of marijuana or a conviction for a
16.22	comparable offense in another jurisdiction on the list of presumptively disqualifying offenses.
16.23	(b) A person convicted of a presumptively disqualifying offense may submit information
16.24	to demonstrate that the person does not pose a risk of harm to any person, will remain law
16.25	abiding, and will comply with the provisions of sections 152.50 to 152.65. If the
16.26	commissioner determines that the person has submitted sufficient information, the
16.27	commissioner may set aside the presumptive disqualification.
16.28	(c) The commissioner shall, by rule, establish other requirements for license holders or
16.29	applicants, including requirements that license holders or applicants must:
16.30	(1) be 21 years of age or older;

23-02009

17.1 (2) have completed an application for licensure or application for renewal and have fully and truthfully complied with all information requests of the commissioner relating to license 17.2 17.3 application and renewal; (3) have paid the applicable application or licensing fee; and 17.4 17.5 (4) not be employed by the commissioner or any state agency with regulatory authority under sections 152.50 to 152.65 or the rules adopted pursuant to those sections. 17.6 17.7 Subd. 5. Application contents, process, and fee. (a) License applications must be made in the form and manner required by the commissioner and shall include all information 17.8 required by the board. 17.9 (b) The applicant for a license must submit a completed criminal history records check 17.10 consent form, a full set of classifiable fingerprints, and the required fees to the commissioner. 17.11 Upon receipt of this information, the commissioner must submit the completed criminal 17.12 history records check consent form, full set of classifiable fingerprints, and required fees 17.13 to the Bureau of Criminal Apprehension. After receiving this information, the bureau must 17.14 conduct a Minnesota criminal history records check of the license applicant. The bureau 17.15 may exchange a license applicant's fingerprints with the Federal Bureau of Investigation to 17.16 obtain the applicant's national criminal history record information. The bureau must return 17.17 the results of the Minnesota and federal criminal history records checks to the commissioner 17.18 to determine if the applicant is disqualified under subdivision 4. 17.19 (c) If the commissioner receives an application that fails to provide the required 17.20 information, the commissioner shall issue a deficiency notice to the applicant. The applicant 17.21 shall have ten business days from the date of the deficiency notice to submit the required 17.22 information. Failure by an applicant to submit all required information will result in the 17.23 application being rejected. 17.24 (d) Within 90 days of receiving a completed application, the commissioner shall issue 17.25 the appropriate license or send the applicant a notice of rejection setting forth specific 17.26 reasons why the commissioner did not approve the application. 17.27 (e) The commissioner may charge a nonrefundable fee, not to exceed \$200, to cover the 17.28 costs associated with reviewing and processing applications. 17.29 17.30 Subd. 6. Inspection and enforcement. (a) The commissioner may enforce sections 152.50 to 152.65, including enforcement against a retailer licensed to sell edible cannabinoid 17.31 products by a local unit of government, under the provisions of sections 144.989 to 144.993. 17.32

BD/HL

18.1	(b) The authority issuing a license to sell edible cannabinoid products may take the
18.2	enforcement actions described in section 152.63, subdivision 7.
18.3	Subd. 7. Not public data. (a) The following data collected, created, or maintained by
18.4	the commissioner is classified as nonpublic data, pursuant to section 13.02, subdivision 9,
18.5	or private data on individuals, pursuant to section 13.02, subdivision 12:
18.6	(1) data submitted by an applicant for a cannabis business license, other than the
18.7	applicant's name, designated address, and trade name;
18.8	(2) the identity of a complainant who has made a report concerning a license holder or
18.9	an applicant that appears in inactive complaint data unless the complainant consents to the
18.10	disclosure;
18.11	(3) the nature or content of unsubstantiated complaints when the information is not
18.12	maintained in anticipation of legal action;
18.13	(4) inactive investigative data relating to violations of statutes or rules;
18.14	(5) the record of any disciplinary proceeding except as limited by paragraph (b);
18.15	(6) data identifying retail customers of a licensed retailer; and
18.16	(7) data identifying employees of a licensed manufacturer, distributor, or retailer of
18.17	edible cannabinoid products.
18.18	(b) Minutes, application data on license holders except nondesignated addresses, orders
18.19	for hearing, findings of fact, conclusions of law, and specifications of the final disciplinary
18.20	action contained in the record of the disciplinary action are classified as public, pursuant to
18.21	section 13.02, subdivision 15. If there is a public hearing concerning the disciplinary action,
18.22	the entire record concerning the disciplinary proceeding is public data pursuant to section
18.23	13.02, subdivision 15. If the license holder and the commissioner agree to resolve a complaint
18.24	without a hearing, the agreement and the specific reasons for the agreement are public data.
18.25	(c) The commissioner must not share data classified as nonpublic or private data on
18.26	individuals under this subdivision or other data identifying an individual applicant or license
18.27	holder with any federal agency, federal department, or federal entity unless specifically
18.28	ordered to do so by a state or federal court.
18.29	(d) The commissioner must establish written procedures to ensure that only individuals
18.30	authorized by law may enter, update, or access the data classified as nonpublic or private
18.31	data on individuals in this subdivision. An authorized individual's ability to enter, update,
18.32	or access data in the system must correspond to the official duties or training level of the

19.1	individual and to the statutory authorization granting access for that purpose. All queries
19.2	and responses and all actions in which not public data are entered, updated, accessed, shared,
19.3	or disseminated, must be recorded in a data audit trail. Data contained in the audit trail have
19.4	the same classification as the underlying data tracked by the audit trail.
10.5	See (1152 55) DOSSESSION OF CONCENTRATED CANNADINOIDS AND
19.5	Sec. 6. [152.55] POSSESSION OF CONCENTRATED CANNABINOIDS AND
19.6	ARTIFICIALLY DERIVED CANNABINOIDS.
19.7	(a) Notwithstanding any other provision of this chapter or any other law to the contrary,
19.8	a licensee may possess concentrated cannabinoids and artificially derived cannabinoids
19.9	provided the licensee:
19.10	(1) is authorized to manufacture products from concentrated cannabinoids and artificially
19.11	derived cannabinoids;
19.12	(2) complies with an approved plan to secure, store, and dispose of concentrated
19.13	cannabinoids and artificially derived cannabinoids; and
19.14	(3) complies with any additional requirements or rules adopted by the commissioner.
19.15	(b) Notwithstanding any other provision of this chapter or any other law to the contrary,
19.16	an approved laboratory may possess concentrated cannabinoids and artificially derived
19.17	cannabinoids during the period that the laboratory is approved by the commissioner to
19.18	perform testing on edible cannabinoid products and the laboratory maintains any required
19.19	accreditation.
19.20	(c) A licensee or laboratory that possesses concentrated cannabinoids or artificially
19.21	derived cannabinoids in violation of this section may be subject to any applicable licensing
19.22	penalty, criminal penalty, or both.
19.23	Sec. 7. [152.56] STATEWIDE MONITORING SYSTEM.
19.24	(a) The commissioner shall coordinate with the commissioner of agriculture to identify
19.25	an approved monitoring system for the integrated tracking, inventory, and verification of
19.26	industrial hemp, concentrated cannabinoids, artificially derived cannabinoids, and edible
19.27	cannabinoid products.
19.28	(b) A licensee must use the monitoring system to track all industrial hemp, concentrated
19.29	cannabinoids, artificially derived cannabinoids, and edible cannabinoid products in the
19.30	licensee's possession to the point of disposal, transfer, or sale. For the purposes of this
19.31	section, a licensee possesses the industrial hemp, concentrated cannabinoids, artificially
19.32	derived cannabinoids, and edible cannabinoid products that the licensee cultivates from

23-02009

- seed or immature plant, concentrates, converts from any other cannabinoid, receives from 20.1 an entity licensed to cultivate or process industrial hemp, manufactures, or receives from 20.2 20.3 an entity licensed to manufacture or distribute products containing cannabinoids derived from industrial hemp. This paragraph does not apply to products lawfully purchased by a 20.4 licensee for personal use. 20.5 Sec. 8. [152.57] EDIBLE CANNABINOID PRODUCTS; CANNABINOID LIMITS; 20.6 **APPROVAL OF PRODUCTS.** 20.7 Subdivision 1. Limits on tetrahydrocannabinol. (a) No edible cannabinoid product 20.8 20.9 may contain more than a total of 0.3 percent of all tetrahydrocannabinols, as measured by weight. 20.10
- 20.11 (b) An edible cannabinoid product that meets the requirement under paragraph (a) must

20.12 not contain more than a total of five milligrams of all tetrahydrocannabinols in a single

20.13 serving. A single package that consists of multiple servings may not contain more than a

- 20.14 total of 50 milligrams of all tetrahydrocannabinols.
- 20.15 <u>Subd. 2.</u> Limits on artificially derived cannabinoids. An edible cannabinoid product 20.16 may contain delta-8 tetrahydrocannabinol, delta-9 tetrahydrocannabinol, or both. No edible 20.17 <u>cannabinoid product may contain any other artificially derived cannabinoid, including but</u> 20.18 <u>not limited to THC-O, THC-P, or HHC unless the commissioner authorizes the use of the</u> 20.19 artificially derived cannabinoid in edible cannabinoid products.
- 20.20 Subd. 3. Approval of product types. (a) No edible cannabinoid product may be sold
- 20.21 <u>unless the product complies with a category or type of product approved by the commissioner.</u>
- 20.22 (b) The commissioner shall approve types or categories of edible cannabinoid products
 20.23 for retail sale.
- 20.24 (c) The commissioner shall not approve any edible cannabinoid product that:
- 20.25 (1) bears the likeness or contains characteristics of a real or fictional person, animal, or
- 20.26 <u>fruit;</u>
- 20.27 (2) is modeled after a brand of products primarily consumed by or marketed to children;
- 20.28 (3) is designed to appeal to persons under age 21;
- 20.29 (4) is made by applying extracted, converted, or concentrated tetrahydrocannabinol to
- 20.30 <u>a finished food product that does not contain cannabinoids and is sold to consumers, including</u>
- 20.31 **but not limited to a candy or snack food;**
- 20.32 (5) is or appears to be a lollipop or ice cream;

21.1	(6) cannot be:
21.2	(i) packaged in a single serving container;
21.3	(ii) packaged in such a way that each serving is clearly indicated through the use of
21.4	individual pieces that constitute a serving; or
21.5	(iii) prepared or packaged in such a way that an individual serving size is indicated
21.6	through the use of scoring, individual wrapping, or other indicators that appear on the edible
21.7	cannabinoid product; or
21.8	(7) contains an ingredient, other than an ingredient extracted or derived from hemp plants
21.9	or hemp plant parts, that is not approved by the United States Food and Drug Administration
21.10	for use in food.
21.11	(d) The commissioner shall not approve any product intended to be consumed by
21.12	combustion or vaporization of the product and inhalation of smoke, aerosol, or vapor from
21.13	the product.
21.14	Sec. 9. [152.58] TESTING OF EDIBLE CANNABINOID PRODUCTS.
21.15	Subdivision 1. Standards established by the commissioner. (a) The commissioner
21.16	shall establish a process for independent laboratories and permit laboratories to apply for
21.17	approval in the form and manner determined by the commissioner. At a minimum, a
21.18	laboratory must operate formal management systems under the International Organization
21.19	for Standardization and be able to enter information into the statewide monitoring system
21.20	to qualify for approval. The commissioner shall identify approved independent laboratories
21.21	on the commissioner's public-facing website.
21.22	(b) The commissioner, in consultation with the commissioner of agriculture, shall
21.23	establish and regularly update requirements for the testing of edible cannabinoid products,
21.24	including a list of:
21.25	(1) cannabinoids for which testing must be performed;
21.26	(2) contaminants, including residual solvents, foreign material, microbiological
21.27	contaminants, heavy metals, pesticide residue, and mycotoxins, for which testing must be
21.28	completed, and the acceptable minimum standards of any contaminant for which testing is
21.29	required; and
21.30	(3) all catalysts identified as being used to manufacture artificially derived cannabinoids,
21.31	indicating whether testing must be performed to determine the presence of a residual catalyst.

23-02009

22.1	(c) The commissioner must make the lists established pursuant to paragraph (b), clauses
22.2	(1) and (2), available to the general public and any entity manufacturing, testing, marketing,
22.3	distributing, or selling products that contain cannabinoids. The commissioner must make
22.4	the lists established pursuant to paragraph (b), clause (3), available to approved independent
22.5	laboratories and laboratories operated by a state government agency, office, or department.
22.6	Subd. 2. Testing by manufacturers and distributors. (a) A licensed manufacturer,
22.7	distributor, or retailer must not sell, offer for sale, or otherwise transfer edible cannabinoid
22.8	products to another licensee or customer unless a representative sample of the batch of
22.9	edible cannabinoid products has been tested according to this section and any relevant rules
22.10	adopted by the commissioner, and has been found to meet the applicable testing standards.
22.11	(b) A licensed manufacturer of edible cannabinoid products shall make each batch
22.12	available for testing by an approved independent laboratory pursuant to a schedule and in
22.13	a manner established by the commissioner. A distributor of edible cannabinoid products
22.14	manufactured by an entity that is not a licensed manufacturer may make each batch available
22.15	for testing by an approved independent laboratory if the commissioner authorizes the testing
22.16	after determining that the process used in the manufacture of the edible cannabinoid product
22.17	will assure consistency within each batch.
22.18	(c) If a certification from an approved independent laboratory verifies that an edible
22.19	cannabinoid product meets the applicable testing standards, a licensed manufacturer,
22.20	distributor, or retailer may sell, offer for sale, or otherwise transfer the batch from which
22.21	the sample was taken to another licensee or customer. If a sample does not meet the
22.22	applicable testing standards, the batch from which the sample was taken shall be subject to
22.23	procedures established by the commissioner for such batches, including destruction,
22.24	remediation, or retesting.
22.25	(d) The licensed manufacturer or distributor on whose behalf testing was performed
22.26	must retain the test results for at least five years after the date of testing. Upon request of
22.27	the commissioner, the manufacturer or distributor must make the results available for
22.28	inspection or provide a copy to the commissioner.
22.29	Subd. 3. Confirmatory tests. A licensed manufacturer, distributor, or retailer must allow
22.30	a sampling agent, the commissioner, or the commissioner's designee to collect and test

22.31 regulatory samples of edible cannabinoid products.

23-02009

23.1	Sec. 10. [152.59] SAFETY OF FOOD INGREDIENTS.
23.2	The commissioner, in consultation with the commissioner of agriculture, shall establish
23.3	rules and policies, perform inspections, and require or perform product testing to ensure
23.4	that the food ingredients and processes used in manufacturing edible cannabinoid products
23.5	comply with the requirements related to food safety that appear in chapters 28A, 31, and
23.6	34A, and associated rules.
23.7	Sec. 11. [152.60] PACKAGING, LABELING, AND MARKETING OF EDIBLE
23.8	CANNABINOID PRODUCTS.
22.0	Subdivision 1 Deckaging (a) All adible connehinoid products sold to sustamore must
23.9 23.10	Subdivision 1. Packaging. (a) All edible cannabinoid products sold to customers must be packaged as required under this section and any relevant rules adopted by the
23.10	commissioner.
23.11	
23.12	(b) An edible cannabinoid product must be prepackaged in packaging or a container that
23.13	is child-resistant, tamper-evident, and opaque or placed in packaging or a container that is
23.14	child-resistant, tamper-evident, and opaque at the final point of sale to a customer. The
23.15	requirement that packaging be child-resistant does not apply to an edible cannabinoid product
23.16	that is intended to be consumed as a beverage and that contains no more than a total of 0.25
23.17	milligrams of all tetrahydrocannabinols.
23.18	(c) An edible cannabinoid product must not be packaged:
23.19	(1) in a way that resembles the trademarked, characteristic, or product-specialized
23.20	packaging of any commercially available food product;
23.21	(2) bears the likeness or contains cartoon-like characteristics of a real or fictional person,
23.22	animal, or fruit that is designed or likely to appeal to persons under 21 years of age; or
23.23	(3) in a container that includes a statement, artwork, or design that could reasonably
23.24	mislead any person to believe that the package contains anything other than an edible
23.25	cannabinoid product.
23.26	Subd. 2. Labeling. (a) All edible cannabinoid products sold to customers must be labeled
23.27	as required under this section and any relevant rules adopted by the commissioner.
23.28	(b) An edible cannabinoid product sold to customers must bear a label that contains, at
23.29	<u>a minimum:</u>
23.30	(1) the name, address or location, contact phone number, and website of the manufacturer

23.31 <u>of the product;</u>

	01/20/23	REVISOR	BD/HL	23-02009
24.1	(2) the name and address of the appr	oved independen	at laboratory used by th	
24.1	manufacturer to test the product;			
24.3	(3) the batch number;			
24.4	(4) the cannabinoid profile;			
24.5	(5) the serving size;			
24.6	(6) the total number of milligrams of	all tetrahydroca	nnabinols in a single se	rving, and,
24.7	if a package contains more than a single	serving, the tota	al number of milligrams	s of all
24.8	tetrahydrocannabinols per package;			
24.9	(7) a list of ingredients, including ide	entification of an	y major food allergens	declared
24.10	by name;			
24.11	(8) a statement that the product does	not claim to dia	gnose, treat, cure, or pr	event any
24.12	disease, does not claim that the product	may be used to a	alter the structure or fur	nction of
24.13	human or animal bodies, and has not bee	n evaluated or ap	oproved by the United S	States Food
24.14	and Drug Administration unless the prod	duct has been so	approved. The labeling	g must not
24.15	contain any statement, artwork, or desig	n that is inconsis	stent with the required s	statement;
24.16	and			
24.17	(9) the following statement: "Keep the	nis product out o	f reach of children."	
24.18	(c) The information in paragraph (b)	may be provide	d on an outer package i	f the
24.19	immediate container that holds the production	uct is too small t	o contain all of the info	ormation.
24.20	(d) The information required in parag	graph (b), clause	es (1), (2), (3), and (4), r	may be
24.21	provided through the use of a scannable	barcode or matr	ix barcode that links to	a page on
24.22	a website maintained by the manufacture	er or distributor	if that page contains all	l of the
24.23	information required by this subdivision	<u>.</u>		
24.24	(e) The information required by this s	ubdivision must	be prominently and con	spicuously
24.25	placed on the label or displayed on the we	bsite in terms tha	t can be easily read and	understood
24.26	by the consumer.			
24.27	Subd. 3. Marketing. (a) No licensee	or other person sh	all publish or cause to be	e published
24.28	an advertisement for an edible cannabine	oid product in a	manner that:	
24.29	(1) contains false or misleading state	ments;		
24.30	(2) contains unverified claims about	the health or the	rapeutic benefits or eff	ects of
24.31	consuming edible cannabinoid products			

01/20/23 REVISOR BD/HL 23-02009 (3) promotes the overconsumption of edible cannabinoid products; 25.1 (4) depicts a person under 21 years of age consuming an edible cannabinoid product; or 25.2 (5) includes an image or phrase designed or likely to appeal to persons under 21 years 25.3 of age or encourage consumption by persons under 21 years of age. 25.4 25.5 (b) No licensee or other person shall publish or cause to be published an advertisement for edible cannabinoid products in any print publication or on radio, television, or any other 25.6 25.7 medium if 30 percent or more of the audience of that medium is reasonably expected to be individuals who are under 21 years of age, as determined by reliable, current audience 25.8 composition data. 25.9 (c) No licensee or other person shall utilize unsolicited pop-up advertisements on the 25.10 Internet or advertising directed toward location-based devices, including but not limited to 25.11 cellular telephones, to advertise edible cannabinoid products unless: 25.12 (1) the advertising occurs via a mobile device application that is installed on the device 25.13 by the device's owner and includes a permanent and easy to implement opt-out feature; and 25.14 (2) the owner of the device is 21 years of age or older. 25.15 Sec. 12. [152.61] MANUFACTURE OF EDIBLE CANNABINOID PRODUCTS. 25.16 25.17 Subdivision 1. Authorized actions. An edible cannabinoid product manufacturer license entitles the license holder to: 25.18 25.19 (1) purchase industrial hemp, concentrated cannabinoids, and artificially derived cannabinoids from a person or entity licensed to grow or process industrial hemp; 25.20 (2) manufacture edible cannabinoid products for public consumption; 25.21 (3) package and label edible cannabinoid products; 25.22 25.23 (4) sell or distribute edible cannabinoid products manufactured by the licensee to licensed distributors or retailers; and 25.24 25.25 (5) perform other actions approved by the commissioner. Subd. 2. Requirements. (a) Manufacturing of edible cannabinoid products must take 25.26 25.27 place: (1) in an enclosed facility that: 25.28 25.29 (i) has appropriate locks or other restrictions to control access; and (ii) meets the sanitary standards specified by the commissioner; and 25.30

 26.1 (2) on equipment that is used exclusively for the manufacture of products containing 26.2 cannabinoids. 26.3 (b) A licensed manufacturer must comply with all applicable testing, storage, packaging 26.4 labeling, and health and safety requirements described in sections 152.50 to 152.65. 26.5 (c) Upon the sale of any edible cannabinoid product to a licensed distributor or retailer 26.6 a licensed manufacturer must provide a label that meets the requirements of section 152.60 26.7 subdivision 2, or provide sufficient information for the distributor or retailer to properly 26.8 label the edible cannabinoid product. 26.9 (d) A licensed manufacturer must record all transactions involving industrial hemp, 26.10 artificially derived cannabinoids, and edible cannabinoid products in the statewide monitoring 26.11 system. 26.12 Subd. 3. Falsification of records; criminal penalty. Notwithstanding section 144.99, 		01/20/23	REVISOR	BD/HL	23-02009
 (b) A licensed manufacturer must comply with all applicable testing, storage, packaging labeling, and health and safety requirements described in sections 152.50 to 152.65. (c) Upon the sale of any edible cannabinoid product to a licensed distributor or retailer a licensed manufacturer must provide a label that meets the requirements of section 152.60 subdivision 2, or provide sufficient information for the distributor or retailer to properly label the edible cannabinoid product. (d) A licensed manufacturer must record all transactions involving industrial hemp, artificially derived cannabinoids, and edible cannabinoid products in the statewide monitoring system. 	26.1	(2) on equipment that is used exc	lusively for the man	ufacture of products c	ontaining
 labeling, and health and safety requirements described in sections 152.50 to 152.65. (c) Upon the sale of any edible cannabinoid product to a licensed distributor or retailer a licensed manufacturer must provide a label that meets the requirements of section 152.60 subdivision 2, or provide sufficient information for the distributor or retailer to properly label the edible cannabinoid product. (d) A licensed manufacturer must record all transactions involving industrial hemp, artificially derived cannabinoids, and edible cannabinoid products in the statewide monitoring system. 	26.2	cannabinoids.			
 labeling, and health and safety requirements described in sections 152.50 to 152.65. (c) Upon the sale of any edible cannabinoid product to a licensed distributor or retailer a licensed manufacturer must provide a label that meets the requirements of section 152.60 subdivision 2, or provide sufficient information for the distributor or retailer to properly label the edible cannabinoid product. (d) A licensed manufacturer must record all transactions involving industrial hemp, artificially derived cannabinoids, and edible cannabinoid products in the statewide monitoring system. 	26.3	(b) A licensed manufacturer must	comply with all appl	icable testing, storage,	packaging,
 26.6 <u>a licensed manufacturer must provide a label that meets the requirements of section 152.60</u> 26.7 <u>subdivision 2, or provide sufficient information for the distributor or retailer to properly</u> 26.8 <u>label the edible cannabinoid product.</u> 26.9 <u>(d) A licensed manufacturer must record all transactions involving industrial hemp,</u> 26.10 <u>artificially derived cannabinoids, and edible cannabinoid products in the statewide monitoring</u> 26.11 <u>system.</u> 	26.4	· · ·			
 26.7 subdivision 2, or provide sufficient information for the distributor or retailer to properly 26.8 label the edible cannabinoid product. 26.9 (d) A licensed manufacturer must record all transactions involving industrial hemp, 26.10 artificially derived cannabinoids, and edible cannabinoid products in the statewide monitoring 26.11 system. 	26.5	(c) Upon the sale of any edible ca	nnabinoid product to	a licensed distributor	r or retailer,
 26.8 label the edible cannabinoid product. 26.9 (d) A licensed manufacturer must record all transactions involving industrial hemp, 26.10 artificially derived cannabinoids, and edible cannabinoid products in the statewide monitoring 26.11 system. 	26.6	a licensed manufacturer must provide	e a label that meets th	e requirements of sect	ion 152.60,
 26.9 (d) A licensed manufacturer must record all transactions involving industrial hemp, 26.10 artificially derived cannabinoids, and edible cannabinoid products in the statewide monitoring 26.11 system. 	26.7	subdivision 2, or provide sufficient in	nformation for the di	stributor or retailer to	properly
 artificially derived cannabinoids, and edible cannabinoid products in the statewide monitoring 26.11 system. 	26.8	label the edible cannabinoid product.	<u>-</u>		
26.11 <u>system.</u>	26.9	(d) A licensed manufacturer must	t record all transaction	ons involving industria	al hemp,
	26.10	artificially derived cannabinoids, and	edible cannabinoid pr	oducts in the statewide	monitoring
Subd. 3. Falsification of records; criminal penalty. Notwithstanding section 144.99,	26.11	system.			
	26.12	Subd. 3. Falsification of records	; criminal penalty.	Notwithstanding secti	on 144.99,
26.13 <u>subdivision 11, a person, including a licensed manufacturer, who intentionally alters or</u>	26.13	subdivision 11, a person, including a	licensed manufactur	er, who intentionally	alters or
26.14 <u>falsifies any information required to be included on the label of an edible cannabinoid</u>	26.14	falsifies any information required to	be included on the la	ibel of an edible canna	abinoid
26.15 product is guilty of a gross misdemeanor and may be sentenced to imprisonment for not	26.15	product is guilty of a gross misdeme	anor and may be sen	tenced to imprisonme	nt for not
26.16 more than one year or to payment of a fine of not more than \$3,000, or both.	26.16	more than one year or to payment of	a fine of not more th	1an \$3,000, or both.	
26.17 Sec. 13. [152.62] DISTRIBUTION OF EDIBLE CANNABINOID PRODUCTS.	26.17	Sec. 13. [152.62] DISTRIBUTIO	N OF EDIBLE CAN	NABINOID PROD	UCTS.
26.18 Subdivision 1. Authorized actions. An edible cannabinoid product distributor license	26.18	Subdivision 1. Authorized action	ns. <u>An edible cannab</u>	oinoid product distribu	tor license
26.19 <u>entitles the license holder to:</u>	26.19	entitles the license holder to:			
26.20 (1) purchase edible cannabinoid products from manufacturers;	26.20	(1) purchase edible cannabinoid	products from manuf	àcturers;	
26.21 (2) sell edible cannabinoid products that meet the requirements of sections 152.50 to	26.21	(2) sell edible cannabinoid produ	cts that meet the requ	uirements of sections	152.50 to
26.22 152.65 to licensed retailers; and	26.22	152.65 to licensed retailers; and			
26.23 (3) perform other actions approved by the commissioner.	26.23	(3) perform other actions approve	ed by the commission	ner.	
26.24 Subd. 2. Requirements. A licensed distributor must:	26.24	Subd. 2. Requirements. A licens	ed distributor must:		
26.25 (1) ensure that edible cannabinoid products are stored in a manner that prevents any	26.25	(1) ensure that edible cannabinoid	d products are stored	in a manner that prev	ents any
26.26 <u>cross contamination;</u>	26.26	cross contamination;			
26.27 (2) ensure that any edible cannabinoid products intended for distribution are stored in	26.27	(2) ensure that any edible cannab	inoid products intend	led for distribution are	e stored in
26.28 <u>an enclosed facility with appropriate locks or other restrictions to control access;</u>	26.28	an enclosed facility with appropriate	locks or other restric	ctions to control acces	<u>s;</u>
26.29 (3) store edible cannabinoid products in clean and sanitary conditions, free from	26.29	(3) store edible cannabinoid prod	ucts in clean and san	itary conditions, free	from
26.30 infestation by insects, rodents, or other pests; and	26.30	infestation by insects, rodents, or oth	er pests; and		

BD/HL

27.1	(4) maintain accurate records and ensure that appropriate labels remain affixed to edible
27.2	cannabinoid products.
27.3	Subd. 3. Distribution of products manufactured outside of the state. (a) A licensed
27.4	distributor may perform the actions described in subdivision 1 related to edible cannabinoid
27.5	products manufactured outside of this state provided that:
27.6	(1) the manufacturer is licensed in another state and subject to regulations designed to
27.7	protect the health and safety of consumers and those regulations are substantially similar
27.8	to the regulations in this state; or
27.9	(2) the distributor establishes, to the satisfaction of the commissioner, that the
27.10	manufacturer engages in practices that are substantially similar to the practices required for
27.11	licensure of manufacturers in this state.
27.12	(b) A distributor must enter all relevant information regarding an edible cannabinoid
27.13	product manufactured in another state into the statewide monitoring system before the
27.14	product may be distributed to a licensed retailer. Relevant information includes information
27.15	regarding the cultivation, processing, and testing of the industrial hemp used in the
27.16	manufacture of the edible cannabinoid product. If information regarding the industrial hemp
27.17	or edible cannabinoid product was submitted to a statewide monitoring system used in
27.18	another state, the commissioner may require submission of any information provided to
27.19	that statewide monitoring system and shall assist in the transfer of data from another state
27.20	as needed and in compliance with any data classification established by either state.
27.21	Subd. 4. Violations in other jurisdictions. The commissioner may suspend, revoke, or
27.22	cancel the license of a distributor who is prohibited from distributing edible cannabinoid
27.23	products in any other jurisdiction, convicted of an offense involving the distribution of
27.24	edible cannabinoid products in any other jurisdiction, or found liable for distributing any
27.25	product that injured customers in any other jurisdiction. A licensee shall disclose to the
27.26	commissioner all relevant information related to the licensee's actions in another jurisdiction.
27.27	Failure to disclose relevant information may result in disciplinary action by the commissioner,
27.28	including the suspension, revocation, or cancellation of a license.
27.29	Subd. 5. Reliance on product label no defense. Notwithstanding any law to the contrary,
27.30	it is not a defense in any civil or criminal action that a licensed distributor relied on
27.31	information on a product label or otherwise provided by a manufacturer who is not a licensed
27.32	edible cannabinoid manufacturer.
27.33	Subd. 6. Unlicensed distribution; distribution of noncompliant products; distribution
27.34	to unlicensed retailers; criminal penalty. Notwithstanding section 144.99, subdivision

BD/HL

- 28.1 11, a person, including a licensed distributor, who does any of the following is guilty of a
- 28.2 gross misdemeanor and may be sentenced to imprisonment for not more than one year or
- 28.3 to payment of a fine of not more than \$3,000, or both:
- 28.4 (1) distributes an edible cannabinoid product without first obtaining a license from the
 28.5 commissioner;
- 28.6 (2) distributes an edible cannabinoid product to a retailer that does not comply with the
- 28.7 <u>limits on the amount or types of cannabinoids a product can contain;</u>
- 28.8 (3) distributes an edible cannabinoid product to a retailer that does not comply with the
- applicable testing, packaging, or labeling requirements; or
- 28.10 (4) distributes an edible cannabinoid product to a retailer who is not licensed to sell
- 28.11 <u>edible cannabinoid products.</u>

28.12 Sec. 14. [152.63] EDIBLE CANNABINOID PRODUCT RETAILER LICENSES; 28.13 LOCAL UNITS OF GOVERNMENT.

- 28.14 Subdivision 1. Issuance by local unit of government. (a) A city or town may issue
- 28.15 annual edible cannabinoid product retailer licenses to persons within the city or town's
- 28.16 jurisdiction. A county board may issue annual edible cannabinoid product retailer licenses
- 28.17 to persons in an area of the county that is unorganized or unincorporated.
- 28.18 (b) Any ordinance adopted by a local unit of government and any license issued by a
- 28.19 local unit of government must comply with the requirements and limits under this section.
- 28.20 Subd. 2. Fees. The annual license fee for an edible cannabinoid product retailer license
- 28.21 is the fee set by the local unit of government issuing the license. The fee must be set at an
- 28.22 amount equal to the median amount of the annual license fee that the local unit of government
- 28.23 charges for an off-sale intoxicating liquor license and the annual license fee that the local
- 28.24 <u>unit of government charges for a license to sell cigarettes and tobacco. The license fee is</u>
- 28.25 intended to cover the costs of issuing the license, inspecting the licensee, and other directly
- 28.26 related costs of enforcement.
- 28.27 Subd. 3. Persons eligible; transfer of licenses. (a) A local unit of government may
 28.28 issue an edible cannabinoid product retailer license to a person who:
- 28.29 (1) is at least 21 years of age;
- 28.30 (2) has completed an application for licensure or application for renewal and has fully
- 28.31 and truthfully complied with all information requests relating to license application and
- 28.32 <u>renewal;</u>

REVISOR

29.1	(3) has paid any applicable licensing fee;
29.2	(4) is not employed by the commissioner or any state agency with regulatory authority
29.3	under sections 152.50 to 152.65 or the rules adopted pursuant to those sections; and
29.4	(5) is of good moral character and repute.
29.5	(b) In determining whether a person is of good moral character and repute, the local unit
29.6	of government may rely on the list of disqualifying offenses established by the commissioner,
29.7	but must not disqualify an application for a violation of chapter 152 involving the possession
29.8	of marijuana or a conviction for a comparable offense in another jurisdiction.
29.9	(c) Licenses may not be transferred.
29.10	Subd. 4. Background check. (a) The chief of police is responsible for conducting a
29.11	background check for an applicant prior to a city or town's issuance of an edible cannabinoid
29.12	product retailer license. A county sheriff is responsible for conducting a background check
29.13	for an applicant prior to the county's issuance of an edible cannabinoid product retailer
29.14	license and for those cities and towns that do not have a police department.
29.15	(b) The applicant for a retail license must submit a completed criminal history records
29.16	check consent form, a full set of classifiable fingerprints, and the required fees to the
29.17	appropriate authority. Upon receipt of this information, the appropriate authority must
29.18	submit the completed criminal history records check consent form, full set of classifiable
29.19	fingerprints, and required fees to the Bureau of Criminal Apprehension. After receiving this
29.20	information, the bureau must conduct a Minnesota criminal history records check of the
29.21	license applicant. The bureau may exchange a license applicant's fingerprints with the
29.22	Federal Bureau of Investigation to obtain the applicant's national criminal history record
29.23	information. The bureau must return the results of the Minnesota and federal criminal history
29.24	records checks to the commissioner to determine if the applicant is disqualified under
29.25	subdivision 3.
29.26	Subd. 5. Retail locations; restrictions on the time, place, and manner of operations. A
29.27	local unit of government may adopt reasonable restrictions on the time, place, and manner
29.28	of the retail sale of edible cannabinoid products. A local unit of government may prohibit
29.29	the sale of edible cannabinoid products within 500 feet of a school, day care, nursing home,
29.30	or house of worship.
29.31	Subd. 6. Notice to commissioner. Within ten days of the issuance of an edible
29.32	cannabinoid product retailer license, a local unit of government shall inform the commissioner
29.33	of the licensee's name, address, trade name, and the effective date and expiration date of

BD/HL

30.1	the license. The local unit of government shall also inform the commissioner of a license
30.2	cancellation, suspension, or revocation during the license period.
30.3	Subd. 7. Enforcement by local unit of government. On a finding that a license holder
30.4	failed to comply with an applicable statute, rule, or ordinance relating to edible cannabinoid
30.5	products, or failed to comply with a lawful license condition duly imposed by the authority
30.6	issuing the license or agreed to by the license holder, the authority issuing a retail license
30.7	under this section may revoke the license, suspend the license for up to 60 days, impose a
30.8	civil penalty of up to \$2,000 for each violation, or impose any combination of these sanctions.
30.9	No suspension or revocation takes effect until the license holder has been given an
30.10	opportunity for a hearing under sections 14.57 to 14.69 of the Administrative Procedure
30.11	Act. This section does not require a city, town, or county to conduct the hearing before an
30.12	employee of the Office of Administrative Hearings. Imposition of a penalty or suspension
30.13	by the issuing authority does not preclude imposition of an additional penalty or suspension
30.14	by the commissioner.
30.15	Sec. 15. [152.64] EDIBLE CANNABINOID PRODUCT RETAILER OPERATIONS.
30.16	Subdivision 1. Authorized actions. An edible cannabinoid product retailer license
30.17	entitles the license holder to purchase edible cannabinoid products from a licensed
30.18	manufacturer or licensed distributor, sell and deliver edible cannabinoid products to customers
30.19	who are 21 years of age or older, and perform other actions approved by the commissioner
30.20	and licensing authority.
30.21	Subd. 2. Requirements. A licensed retailer must:
30.22	(1) ensure that all edible cannabinoid products are stored in a manner that prohibits
30.23	access by persons under 21 years of age;
30.24	(2) ensure that all edible cannabinoid products offered for sale comply with the applicable
30.25	testing, packaging, and labeling requirements;
30.26	(3) ensure that all edible cannabinoid products offered for sale comply with the limits
30.27	on the amount and types of cannabinoids that an edible cannabinoid product can contain;
30.28	(4) record all transactions involving edible cannabinoid products in the statewide
30.29	monitoring system;
30.30	(5) comply with state and local building, fire, and zoning requirements or regulations;
30.31	and

BD/HL

31.1	(6) ensure that the retail premises is maintained in clean and sanitary conditions, free
31.2	from infestation by insects, rodents, or other pests.
31.3	Subd. 3. Prohibitions. A licensed retailer must not:
31.4	(1) sell an edible cannabinoid product to a person who is visibly intoxicated;
31.5	(2) operate a drive-through window;
31.6	(3) allow for the dispensing of edible cannabinoid products from vending machines;
31.7	(4) sell edible cannabinoid products if the retailer knows that the statewide monitoring
31.8	system is not operational; or
31.9	(5) sell edible cannabinoid products to a customer without verifying that the customer
31.10	is at least 21 years of age.
31.11	Subd. 4. Signage. At each location where edible cannabinoid products are sold, the
31.12	licensee shall display a sign in plain view to provide public notice that selling any edible
31.13	cannabinoid product to any person under 21 years of age is illegal and subject to penalties.
31.14	The notice shall be placed in a conspicuous location in the licensed establishment and shall
31.15	be readily visible to any person who is purchasing or attempting to purchase edible
31.16	cannabinoid products. The sign shall provide notice that all persons responsible for selling
31.17	edible cannabinoid products must verify the age of any customer who is under 30 years of
31.18	age by means of photographic identification containing the bearer's date of birth.
31.19	Subd. 5. Age verification. (a) Prior to initiating a sale or delivery, an employee of a
31.20	licensed retailer must verify that a customer is 21 years of age or older.
31.21	(b) Proof of age may be established only by one of the following:
31.22	(1) a valid driver's license or identification card issued by Minnesota, another state, or
31.23	a province of Canada that includes the photograph and date of birth of the licensed person;
31.24	(2) a valid Tribal identification card as defined in section 171.072, paragraph (b);
31.25	(3) a valid passport issued by the United States;
31.26	(4) a valid instructional permit issued under section 171.05 to a person of legal age to
31.27	purchase adult-use cannabis or adult-use cannabis products that includes a photograph and
31.28	the date of birth of the person issued the permit; or
31.29	(5) in the case of a foreign national, a valid passport.
31.30	(c) A licensed retailer may seize a customer's form of identification listed under paragraph
31.31	(b) if the licensed retailer has reasonable grounds to believe that the form of identification

BD/HL

- has been altered or falsified or is being used to violate any law. A licensed retailer that seizes 32.1 a form of identification as authorized under this paragraph must deliver it to a law 32.2 32.3 enforcement agency within 24 hours of seizing it. (d) The commissioner may authorize the use of age-verification software or other 32.4 32.5 processes to permit the purchase of edible cannabinoid products from a licensed retailer's website. 32.6 Subd. 6. Deliveries. Only a licensed retailer may deliver edible cannabinoid products 32.7 from the retailer's store to the residence of a purchaser or other location, provided that such 32.8 delivery must be made only to a person who is 21 years of age or older. A licensed retailer 32.9 32.10 may refuse to sell or deliver edible cannabinoid products to any person whom the retailer has reason to believe is ineligible to buy edible cannabinoid products or when the retailer 32.11 believes that the person intends to deliver an edible cannabinoid product to an ineligible 32.12 consumer. The licensed retailer must verify that the person receiving the delivery is the 32.13 person who purchased the edible cannabinoid product and is 21 years of age or older. 32.14 Subd. 7. Violations; criminal penalty. Notwithstanding section 144.99, subdivision 32.15 11, a person, including a licensed retailer, who commits any of the following acts from the 32.16 premises of a licensed retailer or another business that sells retail goods to customers is 32.17 guilty of a gross misdemeanor and may be sentenced to imprisonment for not more than 32.18 one year or to payment of a fine of not more than \$3,000, or both: 32.19 (1) the person sells an edible cannabinoid product knowing that the product does not 32.20 comply with the limits on the amount or types of cannabinoids a product can contain; 32.21 (2) the person sells an edible cannabinoid product knowing that the product does not 32.22 comply with the applicable testing, packaging, or labeling requirements; or 32.23 (3) the person sells an edible cannabinoid product to a person under 21 years of age, 32.24 except that it is an affirmative defense to a charge under this clause if the defendant proves 32.25 32.26 by a preponderance of the evidence that the defendant reasonably and in good faith relied on proof of age as described in subdivision 5. 32.27
- 32.28 Sec. 16. [152.65] REPORT.
- 32.29 By January 15 of each year, the commissioner shall submit a report to the legislative
- 32.30 committees and divisions with jurisdiction over health policy and finance on the regulation
- 32.31 of edible cannabinoid products. The report shall describe all actions taken by the
- 32.32 commissioner in the previous year, identify all rules adopted by the commissioner regarding
- 32.33 edible cannabinoid products, list the total number of manufacturing licenses and distributor

	01/20/23	REVISOR	BD/HL	23-02009
33.1	licenses issued by the board and	d the total number of retai	ler licenses issued l	by local units
33.2	of government, summarize enfo	preement actions taken by	the commissioner,	and include
33.3	proposed legislative changes, if	any.		
33.4		ARTICLE 3		
33.5	TAXATION OF	FEDIBLE CANNABING	OID PRODUCTS	
33.6	Section 1. [295.81] EDIBLE	CANNABINOID PROD	UCT GROSS REC	CEIPTS TAX.
33.7	Subdivision 1. Definitions.	(a) For the purposes of this	s section, the follow	ing terms have
33.8	the meanings given.			
33.9	(b) "Commissioner" means	the commissioner of reven	nue.	
33.10	(c) "Edible cannabinoid pro	duct" has the meaning giv	en in section 152.5	0, subdivision
33.11	<u>7.</u>			
33.12	(d) "Edible cannabinoid pro	duct retailer" means a reta	ailer that sells edible	e cannabinoid
33.13	products, including a:			
33.14	(1) retailer maintaining a pla	ace of business in this stat	ie;	
33.15	(2) marketplace provider ma	aintaining a place of busin	ness in this state, as	defined in
33.16	section 297A.66, subdivision 1,	, paragraph (a);		
33.17	(3) retailer not maintaining	a place of business in this	state; and	
33.18	(4) marketplace provider no	t maintaining a place of b	ousiness in this state	, as defined in
33.19	section 297A.66, subdivision 1,	, paragraph (b).		
33.20	(e) "Gross receipts" means th	ne total amount received, i	n money or by barte	er or exchange,
33.21	for all sales at retail of edible ca	nnabinoid products as mea	asured by the sales	price, but does
33.22	not include:			
33.23	(1) any taxes imposed direct	ly on the consumer that ar	e separately stated of	on the invoice,
33.24	bill of sale, or similar documen	t given to the purchaser; a	und	
33.25	(2) discounts, including cash	n, terms, or coupons, that a	are not reimbursed b	y a third party
33.26	and that are allowed by the selle	er and taken by a purchase	er on a sale.	
33.27	(f) "On-site sale" means the	sale of edible cannabinoid	l products for consu	mption on the
33.28	premises of an establishment lie	censed under section 152.	63, including establ	ishments with
33.29	an intoxicating liquor license.			
33.30	(g) "Retail sale" has the mea	aning given in section 297	A.61, subdivision	<u>4.</u>

23-02009

34.1	Subd. 2. Gross receipts tax imposed. (a) A tax is imposed on each edible cannabinoid
34.2	retailer equal to 2.5 percent of gross receipts from retail sales in Minnesota of edible
34.3	cannabinoid products. The edible cannabinoid retailer may but is not required to collect the
34.4	tax from the purchaser. If separately stated on the invoice, bill of sale, or similar document
34.5	given to the purchaser, the tax is excluded from the sales price for purposes of the tax
34.6	imposed under chapter 297A.
34.7	(b) If a product subject to the tax imposed by this section is bundled in a single transaction
34.8	with a product or service that is not subject to the tax imposed by this section, the entire
34.9	sales price of the transaction is subject to the tax imposed by this section.
34.10	(c) The tax imposed under this section is in addition to the tax imposed by chapter 297A
34.11	on the sale or use of edible cannabinoid products.
34.12	Subd. 3. Use tax imposed; credit for taxes paid. (a) A person that receives edible
34.13	cannabinoid products for use or storage in Minnesota, other than from an edible cannabinoid
34.14	retailer that paid the tax under subdivision 2, is subject to tax at the rate imposed under
34.15	subdivision 2. Liability for the tax is incurred when the person has possession of the edible
34.16	cannabinoid product in Minnesota. The tax must be remitted to the commissioner in the
34.17	same manner prescribed for taxes imposed under chapter 297A.
34.18	(b) A person that has paid taxes to another state or any subdivision thereof on the same
34.19	transaction and is subject to tax under this section is entitled to a credit for the tax legally
34.20	due and paid to another state or subdivision thereof to the extent of the lesser of (1) the tax
34.21	actually paid to the other state or subdivision thereof, or (2) the amount of tax imposed by
34.22	Minnesota on the transaction subject to tax in the other state or subdivision thereof.
34.23	Subd. 4. Exemptions. (a) The tax imposed under this section does not apply to sales of
34.24	medical cannabis and medical cannabis products purchased by or for the patients enrolled
34.25	in the registry program.
34.26	(b) The use tax imposed under subdivision 2, paragraph (b), does not apply to the
34.27	possession, use, or storage of edible cannabinoid products if (1) the edible cannabinoid
34.28	products have an aggregate cost in any calendar month to the customer of \$100 or less, and
34.29	(2) the edible cannabinoid products were carried into this state by the customer.
34.30	(c) Unless otherwise specified in this section, the exemptions applicable to taxes imposed
34.31	under chapter 297A are not applicable to the taxes imposed under this section.
34.32	Subd. 5. Tax collection required. An edible cannabinoid retailer with nexus in
34.33	Minnesota, who is not subject to tax under subdivision 2, is required to collect the tax

23-02009

- imposed under subdivision 3 from the purchaser of the edible cannabinoid product and give 35.1 the purchaser a receipt for the tax paid. The tax collected must be remitted to the 35.2 35.3 commissioner in the same manner prescribed for the taxes imposed under chapter 207A. Subd. 6. Taxes paid to another state or any subdivision thereof; credit. An edible 35.4 cannabinoid retailer that has paid taxes to another state or any subdivision thereof measured 35.5 by gross receipts and is subject to tax under this section on the same gross receipts is entitled 35.6 to a credit for the tax legally due and paid to another state or any subdivision thereof to the 35.7 35.8 extent of the lesser of (1) the tax actually paid to the other state or any subdivision thereof, or (2) the amount of tax imposed by Minnesota on the gross receipts subject to tax in the 35.9 other taxing state or any subdivision thereof. 35.10 35.11 Subd. 7. Sourcing of sales. The provisions of section 297A.668 apply to the taxes imposed by this section. 35.12 Subd. 8. Administration. Unless specifically provided otherwise, the audit, assessment, 35.13 refund, penalty, interest, enforcement, collection remedies, appeal, and administrative 35.14 provisions of chapters 270C and 289A that are applicable to taxes imposed under chapter 35.15 297A, except the requirement to file returns and remit taxes due electronically, apply to the 35.16 tax imposed under this section. 35.17 Subd. 9. Returns; payment of tax. (a) An edible cannabinoid retailer must report the 35.18 tax on a return prescribed by the commissioner and must remit the tax with the return. The 35.19 return and the tax must be filed and paid using the filing cycle and due dates provided for 35.20 taxes imposed under section 289A.20, subdivision 4, and chapter 297A. 35.21 (b) Interest must be paid on an overpayment refunded or credited to the taxpayer from 35.22 the date of payment of the tax until the date that the refund is paid or credited. For purposes 35.23 of this subdivision, the date of payment is the due date of the return or the date of actual 35.24 payment of the tax, whichever is later. 35.25 Subd. 10. Deposit of revenue. The commissioner shall deposit all revenues, including 35.26 penalties and interest, derived from the tax imposed by this section in the general fund. 35.27 EFFECTIVE DATE. This section is effective for gross receipts received after December 35.28
- 35.29 <u>31, 2023.</u>

	01/20/23	REVISOR	BD/HL	23-02009
36.1	Sec. 2. Minnesota Statutes 2022, se	ection 297A.99, is a	mended by adding a su	ubdivision to
36.2	read:			
36.3	Subd. 4a. Edible cannabinoid p	roduct local tax pr	ohibited. A political s	subdivision
36.4	of this state is prohibited from impos	ing a tax under this	section solely on the s	ale of edible
36.5	cannabinoid products.			
36.6	EFFECTIVE DATE. This section	on is effective the d	ay following final ena	ctment.
36.7		ARTICLE 4		
36.8	М	ISCELLANEOUS		
36.9	Section 1. Minnesota Statutes 2022	, section 13.3806, is	s amended by adding a	u subdivision
36.10	to read:			
36.11	Subd. 23. Edible cannabinoid p	roducts data. Data	held by the commissio	ner of health
36.12	in connection with the licensing of n	nanufacturers, distri	butors, and retailers o	f edible
36.13	cannabinoid products are classified u	under section 152.54	4, subdivision 7.	
36.14	Sec. 2. Minnesota Statutes 2022, se	ection 34A.01, subd	livision 4, is amended	to read:
36.15	Subd. 4. Food. "Food" means eve	ery ingredient used	for, entering into the c	consumption
36.16	of, or used or intended for use in the p	preparation of food,	drink, confectionery, c	or condiment
36.17	for humans or other animals, whethe	r simple, mixed, or	compound; and article	es used as
36.18	components of these ingredients, exc	cept that edible canr	nabinoid products, as o	defined in
36.19	section 151.72, subdivision 1, parag	caph (c) 152.50, sub	division 7, are not for	od.
36.20	Sec. 3. Minnesota Statutes 2022, se	ection 144.99, subd	ivision 1, is amended	to read:
36.21	Subdivision 1. Remedies availab	le. The provisions of	Chapters 103I and 157	and sections
36.22	115.71 to 115.77; 144.12, subdivisio	n 1, paragraphs (1),	(2), (5), (6), (10), (12), (13), (14),
36.23	and (15); 144.1201 to 144.1204; 144.	121; 144.1215; 144.	1222; 144.35; 144.38	l to 144.385;
36.24	144.411 to 144.417; 144.495; 144.71	to 144.74; 144.950)1 to 144.9512; 144.97	7 to 144.98;
36.25	144.992; 152.22 to 152.37; <u>152.50 to</u>	<u>o 152.65;</u> 326.70 to	326.785; 327.10 to 32	27.131; and
36.26	327.14 to 327.28 and all rules, order	s, stipulation agreer	nents, settlements, con	mpliance
36.27	agreements, licenses, registrations, c	ertificates, and pern	nits adopted or issued	by the
36.28	department or under any other law n	ow in force or later	enacted for the preser	vation of
36.29	public health may, in addition to prov	visions in other statu	tes, be enforced under	this section.

37.1	Sec. 4. Minnesota Statutes 2022, section 152.027, is amended by adding a subdivision to
37.2	read:
37.3	Subd. 8. Sale or possession of edible cannabinoid products. (a) As used in this section,
37.4	"edible cannabinoid product" has the meaning given in section 152.50, subdivision 7.
37.5	(b) A person under 21 years of age who unlawfully possesses any amount of an edible
37.6	cannabinoid product is guilty of a petty misdemeanor.
37.7	(c) A person who unlawfully sells an edible cannabinoid product, except a sale for no
37.8	remuneration to a person who is 21 years of age or older, is guilty of a misdemeanor.
37.9	EFFECTIVE DATE. This section is effective August 1, 2023, and applies to crimes
37.10	committed on or after that date.
37.11	Sec. 5. Minnesota Statutes 2022, section 181.938, subdivision 2, is amended to read:
37.12	Subd. 2. Prohibited practice. (a) An employer may not refuse to hire a job applicant
37.13	or discipline or discharge an employee because the applicant or employee engages in or has
37.14	engaged in the use or enjoyment of lawful consumable products, if the use or enjoyment
37.15	takes place off the premises of the employer during nonworking hours. For purposes of this
37.16	section, "lawful consumable products" means products whose use or enjoyment is lawful
37.17	and which are consumed during use or enjoyment, and includes food, alcoholic or
37.18	nonalcoholic beverages, and tobacco, and edible cannabinoid products as defined in section
37.19	<u>152.50, subdivision 7.</u>
37.20	(b) An edible cannabinoid product is a lawful consumable product for the purpose of
37.21	Minnesota law, regardless of whether federal or other state law considers cannabis use,
37.22	possession, impairment, sale, or transfer to be unlawful. Nothing in this section shall be
37.23	construed to limit an employer's ability to discipline or discharge an employee for cannabis
37.24	use, possession, impairment, sale, or transfer during working hours, on work premises, or
37.25	while operating an employer's vehicle, machinery, or equipment.
37.26	Sec. 6. Minnesota Statutes 2022, section 340A.412, subdivision 14, is amended to read:
37.27	Subd. 14. Exclusive liquor stores. (a) Except as otherwise provided in this subdivision,
37.28	an exclusive liquor store may sell only the following items:
37.29	(1) alcoholic beverages;
37.30	(2) tobacco products;
37.31	(3) ice;

REVISOR

BD/HL

38.1	(4) beverages, either liquid or powder, specifically designated for mixing with intoxicating
38.2	liquor;
38.3	(5) soft drinks;
38.4	(6) liqueur-filled candies;
38.5	(7) food products that contain more than one-half of one percent alcohol by volume;
38.6	(8) cork extraction devices;
38.7	(9) books and videos on the use of alcoholic beverages;
38.8	(10) magazines and other publications published primarily for information and education
38.9	on alcoholic beverages;
38.10	(11) multiple-use bags designed to carry purchased items;
38.11	(12) devices designed to ensure safe storage and monitoring of alcohol in the home, to
38.12	prevent access by underage drinkers;
38.13	(13) home brewing equipment;
38.14	(14) clothing marked with the specific name, brand, or identifying logo of the exclusive
38.15	liquor store, and bearing no other name, brand, or identifying logo;
38.16	(15) citrus fruit; and
38.17	(16) glassware; and
38.18	(17) edible cannabinoid products as defined in section 152.50, subdivision 7.
38.19	(b) An exclusive liquor store that has an on-sale, or combination on-sale and off-sale
38.20	license may sell food for on-premise consumption when authorized by the municipality
38.21	issuing the license.
38.22	(c) An exclusive liquor store may offer live or recorded entertainment.
38.23	Sec. 7. <u>REPEALER.</u>

38.24 Minnesota Statutes 2022, section 151.72, is repealed.

REVISOR

39.1	ARTICLE 5
39.2	APPROPRIATIONS
39.3	Section 1. DEPARTMENT OF HEALTH; APPROPRIATION.
39.4	\$ in fiscal year 2024 and \$ in fiscal year 2025 are appropriated from the general
39.5	fund to the commissioner of health to perform the duties related to regulating edible
39.6	cannabinoid products described in Minnesota Statutes, sections 152.50 to 152.65.
39.7	Sec. 2. DEPARTMENT OF AGRICULTURE; APPROPRIATION.
39.8	\$ in fiscal year 2024 and \$ in fiscal year 2025 are appropriated from the general
39.9	fund to the commissioner of agriculture for the regulation of concentrated cannabinoids and
39.10	artificially derived cannabinoids as described in Minnesota Statutes, chapter 18K.

APPENDIX Repealed Minnesota Statutes: 23-02009

151.72 SALE OF CERTAIN CANNABINOID PRODUCTS.

Subdivision 1. **Definitions.** (a) For the purposes of this section, the following terms have the meanings given.

(b) "Certified hemp" means hemp plants that have been tested and found to meet the requirements of chapter 18K and the rules adopted thereunder.

(c) "Edible cannabinoid product" means any product that is intended to be eaten or consumed as a beverage by humans, contains a cannabinoid in combination with food ingredients, and is not a drug.

(d) "Hemp" has the meaning given to "industrial hemp" in section 18K.02, subdivision 3.

(e) "Label" has the meaning given in section 151.01, subdivision 18.

(f) "Labeling" means all labels and other written, printed, or graphic matter that are:

(1) affixed to the immediate container in which a product regulated under this section is sold;

(2) provided, in any manner, with the immediate container, including but not limited to outer containers, wrappers, package inserts, brochures, or pamphlets; or

(3) provided on that portion of a manufacturer's website that is linked by a scannable barcode or matrix barcode.

(g) "Matrix barcode" means a code that stores data in a two-dimensional array of geometrically shaped dark and light cells capable of being read by the camera on a smartphone or other mobile device.

(h) "Nonintoxicating cannabinoid" means substances extracted from certified hemp plants that do not produce intoxicating effects when consumed by any route of administration.

Subd. 2. **Scope.** (a) This section applies to the sale of any product that contains cannabinoids extracted from hemp and that is an edible cannabinoid product or is intended for human or animal consumption by any route of administration.

(b) This section does not apply to any product dispensed by a registered medical cannabis manufacturer pursuant to sections 152.22 to 152.37.

(c) The board must have no authority over food products, as defined in section 34A.01, subdivision 4, that do not contain cannabinoids extracted or derived from hemp.

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, provided that a product sold for human or animal consumption does not contain more than 0.3 percent of any tetrahydrocannabinol and an edible cannabinoid product does not contain an amount of any tetrahydrocannabinol that exceeds the limits established in subdivision 5a, paragraph (f).

(b) 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.

(c) 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.

(d) Products that meet the requirements of this section are not controlled substances under section 152.02.

Subd. 4. **Testing requirements.** (a) A manufacturer of a product regulated under this section must submit representative samples of the product to an independent, accredited laboratory in order to certify that the product complies with the standards adopted by the board. Testing must be consistent with generally accepted industry standards for herbal and botanical substances, and, at a minimum, the testing must confirm that the product:

(1) contains the amount or percentage of cannabinoids that is stated on the label of the product;

APPENDIX Repealed Minnesota Statutes: 23-02009

(2) does not contain more than trace amounts of any mold, residual solvents, pesticides, fertilizers, or heavy metals; and

(3) does not contain more than 0.3 percent of any tetrahydrocannabinol.

(b) Upon the request of the board, the manufacturer of the product must provide the board with the results of the testing required in this section.

(c) Testing of the hemp from which the nonintoxicating cannabinoid was derived, or possession of a certificate of analysis for such hemp, does not meet the testing requirements of this section.

Subd. 5. Labeling requirements. (a) A product regulated under this section must bear a label that contains, at a minimum:

(1) the name, location, contact phone number, and website of the manufacturer of the product;

(2) the name and address of the independent, accredited laboratory used by the manufacturer to test the product; and

(3) an accurate statement of the amount or percentage of cannabinoids found in each unit of the product meant to be consumed.

(b) The information in paragraph (a) may be provided on an outer package if the immediate container that holds the product is too small to contain all of the information.

(c) The information required in paragraph (a) may be provided through the use of a scannable barcode or matrix barcode that links to a page on the manufacturer's website if that page contains all of the information required by this subdivision.

(d) The label must also include a statement stating that the product does not claim to diagnose, treat, cure, or prevent any disease and has not been evaluated or approved by the United States Food and Drug Administration (FDA) unless the product has been so approved.

(e) The information required by this subdivision must be prominently and conspicuously placed on the label or displayed on the website in terms that can be easily read and understood by the consumer.

(f) The labeling must not contain any claim that the product may be used or is effective for the prevention, treatment, or cure of a disease or that it may be used to alter the structure or function of human or animal bodies, unless the claim has been approved by the FDA.

Subd. 5a. Additional requirements for edible cannabinoid products. (a) In addition to the testing and labeling requirements under subdivisions 4 and 5, an edible cannabinoid must meet the requirements of this subdivision.

(b) An edible cannabinoid product must not:

(1) bear the likeness or contain cartoon-like characteristics of a real or fictional person, animal, or fruit that appeals to children;

(2) be modeled after a brand of products primarily consumed by or marketed to children;

(3) be made by applying an extracted or concentrated hemp-derived cannabinoid to a commercially available candy or snack food item;

(4) contain an ingredient, other than a hemp-derived cannabinoid, that is not approved by the United States Food and Drug Administration for use in food;

(5) be packaged in a way that resembles the trademarked, characteristic, or product-specialized packaging of any commercially available food product; or

(6) be packaged in a container that includes a statement, artwork, or design that could reasonably mislead any person to believe that the package contains anything other than an edible cannabinoid product.

(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 requirement that packaging be child-resistant does not apply to an edible cannabinoid product that is intended to be consumed as a beverage and which contains no more than a trace amount of any tetrahydrocannabinol.

APPENDIX Repealed Minnesota Statutes: 23-02009

(d) If an edible cannabinoid product 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.

(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 by name; and

(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, or more than a total of 50 milligrams of any tetrahydrocannabinol per package.

Subd. 6. **Enforcement.** (a) A product regulated under this section, including an edible cannabinoid product, shall be considered an adulterated drug if:

(1) it consists, in whole or in part, of any filthy, putrid, or decomposed substance;

(2) it has been produced, prepared, packed, or held under unsanitary conditions where it may have been rendered injurious to health, or where it may have been contaminated with filth;

(3) its container is composed, in whole or in part, of any poisonous or deleterious substance that may render the contents injurious to health;

(4) it contains any food additives, color additives, or excipients that have been found by the FDA to be unsafe for human or animal consumption;

(5) it contains an amount or percentage of nonintoxicating cannabinoids that is different than the amount or percentage stated on the label;

(6) it contains more than 0.3 percent of any tetrahydrocannabinol or, if the product is an edible cannabinoid product, an amount of tetrahydrocannabinol that exceeds the limits established in subdivision 5a, paragraph (f); or

(7) it contains more than trace amounts of mold, residual solvents, pesticides, fertilizers, or heavy metals.

(b) A product regulated under this section shall be considered a misbranded drug if the product's labeling is false or misleading in any manner or in violation of the requirements of this section.

(c) The board's authority to issue cease and desist orders under section 151.06; to embargo adulterated and misbranded drugs under section 151.38; and to seek injunctive relief under section 214.11, extends to any violation of this section.