Sixtieth Legislative Assembly of North Dakota

SENATE BILL NO. 2319

Introduced by

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Senators Grindberg, Lyson, Nelson

Representatives DeKrey, Delmore, Thoreson

- 1 A BILL for an Act to amend and reenact sections 19-03.1-01 and 19-03.4-08 of the North
- 2 Dakota Century Code, relating to definitions and the sale of scheduled listed chemical products.

3 BE IT ENACTED BY THE LEGISLATIVE ASSEMBLY OF NORTH DAKOTA:

- SECTION 1. AMENDMENT. Section 19-03.1-01 of the North Dakota Century Code is amended and reenacted as follows:
 - **19-03.1-01. Definitions.** As used in this chapter and in chapters 19-03.2 and 19-03.4, unless the context otherwise requires:
 - "Administer" means to apply a controlled substance, whether by injection, inhalation, ingestion, or any other means, directly to the body of a patient or research subject by:
 - A practitioner or, in the practitioner's presence, by the practitioner's authorized agent; or
 - b. The patient or research subject at the direction and in the presence of the practitioner.
 - "Agent" means an authorized person who acts on behalf of or at the direction of a manufacturer, distributor, or dispenser. It does not include a common or contract carrier, public warehouseman, or employee of the carrier or warehouseman.
 - "Anabolic steroids" means any drug or hormonal substance, chemically and pharmacologically related to testosterone, other than estrogens, progestins, and corticosteroids.
- 21 4. "Board" means the state board of pharmacy.
- 5. "Bureau" means the drug enforcement administration in the United States department of justice or its successor agency.

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1 6. "Controlled substance" means a drug, substance, or immediate precursor in 2 schedules I through V as set out in this chapter. 3 7. "Counterfeit substance" means a controlled substance which, or the container or 4 labeling of which, without authorization, bears the trademark, trade name, or other 5 identifying mark, imprint, number or device, or any likeness thereof, of a 6 manufacturer, distributor, or dispenser other than the person who in fact 7 manufactured, distributed, or dispensed the substance. 8 8. "Deliver" or "delivery" means the actual, constructive, or attempted transfer from 9 one person to another of a controlled substance whether or not there is an agency 10 relationship. 11 9. "Dispense" means to deliver a controlled substance to an ultimate user or research 12 subject by or pursuant to the lawful order of a practitioner, including the 13 prescribing, administering, packaging, labeling, or compounding necessary to 14 prepare the substance for that delivery. 15 10. "Dispenser" means a practitioner who dispenses. 11. 16 "Distribute" means to deliver other than by administering or dispensing a controlled 17 substance. 18 12. "Distributor" means a person who distributes. 19 13. "Drug" means: 20 Substances recognized as drugs in the official United States pharmacopeia, 21 national formulary, or the official homeopathic pharmacopeia of the United 22 States, or any supplement to any of them; 23 Substances intended for use in the diagnosis, cure, mitigation, treatment, or b. 24 prevention of disease in individuals or animals: Substances, other than food, intended to affect the structure or any function of 25 C. 26 the body of individuals or animals; and 27 d. Substances intended for use as a component of any article specified in 28 subdivision a, b, or c. The term does not include devices or their components,

parts, or accessories.

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1 14. "Hashish" means the resin extracted from any part of the plant cannabis with or 2 without its adhering plant parts, whether growing or not, and every compound, 3 manufacture, salt, derivative, mixture, or preparation of the resin. 4 15. "Immediate precursor" means a substance: 5 That the board has found to be and by rule designates as being the principal 6 compound commonly used or produced primarily for use in the manufacture 7 of a controlled substance; 8 b. That is an immediate chemical intermediary used or likely to be used in the 9 manufacture of the controlled substance; and 10 The control of which is necessary to prevent, curtail, or limit the manufacture C. 11 of the controlled substance. 12 16. "Manufacture" means the production, preparation, propagation, compounding, 13 conversion, or processing of a controlled substance, either directly or indirectly by 14 extraction from substances of natural origin, or independently by means of 15 chemical synthesis, or by a combination of extraction and chemical synthesis and 16 includes any packaging or repackaging of the substance or labeling or relabeling of 17 its container. The term does not include the preparation or compounding of a 18 controlled substance by an individual for the individual's own use or the 19 preparation, compounding, packaging, or labeling of a controlled substance: 20 By a practitioner as an incident to the practitioner's administering or 21 dispensing of a controlled substance in the course of the practitioner's 22 professional practice; or 23 By a practitioner, or by the practitioner's authorized agent under the b. 24 practitioner's supervision, for the purpose of, or as an incident to, research, 25 teaching, or chemical analysis and not for sale. 26 17. "Marijuana" means all parts of the plant cannabis whether growing or not; the 27 seeds thereof; the resinous product of the combustion of the plant cannabis; and 28 every compound, manufacture, salt, derivative, mixture, or preparation of the plant 29 or its seeds. The term does not include the mature stalks of the plant, fiber

produced from the stalks, oil or cake made from the seeds of the plant, any other

compound, manufacture, salt, derivative, mixture, or preparation of mature stalks,

1 fiber, oil, or cake, or the sterilized seed of the plant which is incapable of 2 germination. 3 18. "Methamphetamine precursor drug" means a drug or product containing 4 ephedrine, pseudoephedrine, or any of their salts, optical isomers, or salts of 5 optical isomers. 6 19. "Narcotic drug" means any of the following, whether produced directly or indirectly 7 by extraction from substances of vegetable origin, or independently by means of 8 chemical synthesis, or by a combination of extraction and chemical synthesis: 9 Opium and opiate and any salt, compound, derivative, or preparation of opium a. 10 or opiate. 11 b. Any salt, compound, isomer, derivative, or preparation thereof which is 12 chemically equivalent or identical with any of the substances referred to in 13 subdivision a, but not including the isoquinoline alkaloids of opium. 14 Opium poppy and poppy straw. C. 15 d. Coca leaves and any salt, compound, derivative, or preparation of coca 16 leaves, any salt, compound, isomer, derivative, or preparation thereof which is 17 chemically equivalent or identical with any of these substances, but not 18 including decocainized coca leaves or extractions of coca leaves which do not 19 contain cocaine or ecgonine. 20 20. 19. "Opiate" means any substance having an addiction-forming or addiction-sustaining 21 liability similar to morphine or being capable of conversion into a drug having 22 addiction-forming or addiction-sustaining liability. The term does not include, 23 unless specifically designated as controlled under section 19-03.1-02, the 24 dextrorotatory isomer of 3-methoxy-n-methylmorphinan and its salts 25 (dextromethorphan). The term includes its racemic and levorotatory forms. 26 21. 20. "Opium poppy" means the plant of the species papaver somniferum L., except its 27 seeds. 28 22. 21. "Over-the-counter sale" means a retail sale of a drug or product other than a 29 controlled, or imitation controlled, substance.

1 23. 22. "Person" means individual, corporation, limited liability company, government or 2 governmental subdivision or agency, business trust, estate, trust, partnership or 3 association, or any other legal entity. 4 "Poppy straw" means all parts, except the seeds, of the opium poppy, after 24. 23. 5 mowing. 6 25. 24. "Practitioner" means: 7 A physician, dentist, veterinarian, pharmacist, scientific investigator, or other 8 person licensed, registered, or otherwise permitted by the jurisdiction in which 9 the individual is practicing to distribute, dispense, conduct research with 10 respect to, or to administer a controlled substance in the course of 11 professional practice or research. 12 b. A pharmacy, hospital, or other institution licensed, registered, or otherwise 13 permitted to distribute, dispense, conduct research with respect to, or to 14 administer a controlled substance in the course of professional practice or 15 research in this state. 16 26. 25. "Production" includes the manufacturing, planting, cultivating, growing, or 17 harvesting of a controlled substance. 18 27. 26. "Sale" includes barter, exchange, or gift, or offer therefor, and each such 19 transaction made by a person, whether as principal, proprietor, agent, servant, or 20 employee. 21 27. "Scheduled listed chemical product" means a product that contains ephedrine, 22 pseudoephedrin, or phenylpropanolamine, or each of the salts, optical isomers, 23 and salts of optical isomers of each chemical, and that may be marketed or 24 distributed in the United States under the Federal Food, Drug, and Cosmetic Act 25 [21 U.S.C. 301 et seq.] as a nonprescription drug. 26 28. "State" when applied to a part of the United States includes any state, district, 27 commonwealth, territory, insular possession thereof, and any area subject to the 28 legal authority of the United States of America. 29 29. "Ultimate user" means an individual who lawfully possesses a controlled substance 30 for the individual's own use or for the use of a member of the individual's

1		house	ehold or for administering to an animal owned by the individual or by a
2		meml	per of the individual's household.
3	SEC	CTION	2. AMENDMENT. Section 19-03.4-08 of the North Dakota Century Code is
4	amended a	nd ree	nacted as follows:
5	19-0	03.4-08	3. (Effective through July 31, 2007) Retail or over-the-counter sale of
6	methamph	etamiı	ne precursor drugs scheduled listed chemical products - Penalty.
7	1.	The r	etail sale of methamphetamine precursor drugs scheduled listed chemical
8		produ	ucts is limited to:
9		a. \$	Sales in packages containing not more than a total of two grams of one or
10		ı	more methamphetamine precursor drugs scheduled listed chemical products
11		(calculated in terms of ephedrine HCl and base, pseudoephedrine HCl base,
12		<u> </u>	and phenylpropanolamine base; and
13		b. \$	Sales in blister packs, each blister containing not more than two dosage units
14		(or when the use of blister packs is technically infeasible, sales in unit dose
15		ı	packets or pouches.
16	2.	A per	son may not deliver :
17		<u>a.</u> <u>l</u>	Deliver in a single over-the-counter sale more than two packages of a
18		+	methamphetamine precursor drug scheduled listed chemical product or a
19		(combination of methamphetamine precursor drugs scheduled listed chemical
20		1	oroducts; or
21		<u>b.</u> \	Without regard to the number of over-the-counter sales, deliver more than a
22		<u>(</u>	daily amount of three and six-tenths grams of scheduled listed chemical
23		ı	products, calculated in terms of ephedrine base, pseudoephedrine base, and
24		ı	ohenylpropanolamine base, to a purchaser.
25	<u>3.</u>	When	offering scheduled listed chemical products for sale, the person shall place
26		the p	roducts behind a counter or other barrier, or in a locked cabinet, where
27		purch	asers do not have direct access to the products before the sale is made.
28	3. <u>4.</u>	a. \	When offering a methamphetamine precursor drug scheduled listed chemical
29		1	oroducts for retail sale, a person shall require, obtain, and make a written
30		ı	record of the identification of the person purchasing the methamphetamine
31		i	erecursor drug scheduled listed chemical product, the identification being a

1 document issued by a government agency as described in subdivisions a and 2 b of subsection 5 6, and shall do at least one of the following: 3 (1) Maintain continuous recorded video surveillance of the portion of the 4 premises where the methamphetamine precursor drug is displayed for 5 sale and place signs or placards giving notice to the public of the surveillance; 6 7 (2) Place the methamphetamine precursor drug behind a counter or other 8 barrier accessible only to the person making the sale of the drug; or 9 (3) Display only one package of any brand or type of a methamphetamine 10 precursor drug for purchase in an area accessible to the public deliver 11 the product directly into the custody of the purchaser. 12 <u>b.</u> The person shall maintain a written list of sales that identifies the product by 13 name, the quantity sold, the names and addresses of the purchasers, the 14 dates and times of the sales, and a notice to a purchaser that the making of 15 false statements or misrepresentations may subject the purchaser to federal 16 and state criminal penalties. The purchaser shall sign the written list of sales 17 and enter his or her name, address, and the date and time of the sale. The 18 person making the sale shall determine that the name entered by the 19 purchaser corresponds with the name on the identification provided by the 20 purchaser and that the date and time of the purchase is correct. The person 21 making the sale shall enter the name of the product and the quantity sold on 22 the list. 23 The person shall maintain the record of identification required by this b. с. 24 subsection for three years, after which the record must be destroyed. The 25 person may not use or maintain the record for any private or commercial 26 purpose or disclose the record to any person, except as required by law. The 27 person shall disclose the record, upon request, to a law enforcement agency 28 for a law enforcement purpose. A person who in good faith releases the 29 information in the record of identification to federal, state, or local law 30 enforcement authorities is immune from civil liability for such release unless

ı				the release constitutes gross negligence or intentional, wanton, or willium
2				misconduct.
3	4.	<u>5.</u>	Ар	erson may not deliver in an over-the-counter sale a methamphetamine
4			pre	cursor drug scheduled listed chemical product to a person under the age of
5			eigł	nteen years.
6	5.	<u>6.</u>	It is	a prima facie case of a violation of subsection 4 $\underline{5}$ if the person making the sale
7			did	not require and obtain proof of age from the purchaser , unless from the
8			pur	chaser's outward appearance the person would reasonably presume the
9			pur	chaser to be twenty five years of age or older. "Proof of age" means a
10			doc	ument issued by a governmental agency which:
11			a.	Contains a description of the person or a photograph of the person, or both,
12				and gives the person's date of birth; and
13			b.	Includes a passport, military identification card, or driver's license.
14	6.	<u>7.</u>	It is	an affirmative defense to a violation of subsection 4 5 if:
15			a.	The person making the sale required and obtained proof of age from the
16				purchaser;
17			b.	The purchaser falsely represented the purchaser's proof of age by use of a
18				false, forged, or altered document;
19			c.	The appearance of the purchaser was such that an ordinary and prudent
20				person would believe the purchaser to be at least eighteen years of age; and
21			d.	The sale was made in good faith and in reliance upon the appearance and
22				representation of proof of age of the purchaser.
23	7.	<u>8.</u>	This	s section does not apply to pediatric products labeled pursuant to federal
24			regi	ulation primarily intended for administration to children under twelve years of
25			age	according to label instructions or to a product that the state board of pharmacy,
26			upo	n application of a manufacturer, exempts from this section because the product
27			has	been formulated in such a way as to effectively prevent the conversion of the
28			acti	ve ingredient into methamphetamine, or its salts or precursors.
29		<u>9.</u>	A p	erson may not:
30			<u>a.</u>	Make a false statement or misrepresentation in the written list of sale that is
31				prepared and maintained as required by subsection 4; or

ı			<u>D.</u>	Purchase more than nine grams or ephedrine base, pseudoephedrine base,
2				or phenylpropanolamine base in scheduled listed chemical products in a
3				thirty-day period.
4	8.	<u>10.</u>	Ар	erson who willfully violates subsection 1 <u>or 9</u> is guilty of a class A
5			mis	demeanor. A person who willfully violates subsection 2, 3, et 4, or 5 is guilty of
6			an i	nfraction.
7	9.	<u>11.</u>	Ар	erson who is the owner, operator, or manager of the retail outlet or who is the
8			sup	ervisor of the employee or agent committing a violation of this section of the
9			outl	et where methamphetamine precursor drugs scheduled listed chemical
10			prod	ducts are available for sale is not subject to the penalties of this section if the
11			pers	son:
12			a.	Did not have prior knowledge of, participate in, or direct the employee or
13				agent to commit, the violation of this section; and
14			b.	Documents Certifies to the attorney general that the employee or agent, at the
15				time of initial employment and each calendar year thereafter, participated in a
16				training program approved by the attorney general providing the employee or
17				agent with information regarding the state and federal regulations governing
18				the sale, possession, and packaging of such drugs products.
19			The	approval of the training program by the attorney general is not subject to
20			cha	pter 28-32.
21	10.	<u>12.</u>	A p	olitical subdivision, including a home rule city or county, may not enact any
22			ordi	nance relating to the sale by a retail distributor of over-the-counter products
23			con	taining ephedrine, pseudoephedrine, or phenylpropanolamine. Any existing
24			ordi	nance is void.
25		(Eff	ectiv	e after July 31, 2007) Retail or over-the-counter sale of methamphetamine
26	prec	ursor	drug	s - Penalty.
27		1.	The	retail sale of nonliquid methamphetamine precursor drugs is limited to:
28			a.	Sales in packages containing not more than a total of three grams of one or
29				more methamphetamine precursor drugs, calculated in terms of ephedrine
30				HCl and pseudoephedrine HCl; and

1		b. Sales in blister packs, each blister containing not more than two dosage units,
2		or when the use of blister packs is technically infeasible, sales in unit dose
3		packets or pouches.
4	2.	A person may not deliver in a single over the counter sale more than two packages
5		of a methamphetamine precursor drug or a combination of methamphetamine
6		precursor drugs.
7	3.	A person may not deliver in an over-the-counter sale a methamphetamine
8		precursor drug to a person under the age of eighteen years.
9	4.	It is a prima facie case of a violation of subsection 3 if the person making the sale
10		did not require and obtain proof of age from the purchaser, unless from the
11		purchaser's outward appearance the person would reasonably presume the
12		purchaser to be twenty-five years of age or older. "Proof of age" means a
13		document issued by a governmental agency which:
14		a. Contains a description of the person or a photograph of the person, or both,
15		and gives the person's date of birth; and
16		b. Includes a passport, military identification card, or driver's license.
17	5.	It is an affirmative defense to a violation of subsection 3 if:
18		a. The person making the sale required and obtained proof of age from the
19		purchaser;
20		b. The purchaser falsely represented the purchaser's proof of age by use of a
21		false, forged, or altered document;
22		e. The appearance of the purchaser was such that an ordinary and prudent
23		person would believe the purchaser to be at least eighteen years of age; and
24		d. The sale was made in good faith and in reliance upon the appearance and
25		representation of proof of age of the purchaser.
26	6.	This section does not apply to pediatric products labeled pursuant to federal
27		regulation primarily intended for administration to children under twelve years of
28		age according to label instructions or to a product that the state board of pharmacy,
29		upon application of a manufacturer, exempts from this section because the product
30		has been formulated in such a way as to effectively prevent the conversion of the
31		active ingredient into methamphetamine, or its salts or precursors.

1 A person who willfully violates subsection 1 is guilty of a class A misdemeanor. A 2 person who willfully violates subsection 2 or 3 is guilty of an infraction. 3 8. A person who is the owner, operator, or manager of the retail outlet or who is the 4 supervisor of the employee or agent committing a violation of this section of the 5 outlet where methamphetamine precursor drugs are available for sale is not 6 subject to the penalties of this section if the person: 7 Did not have prior knowledge of, participate in, or direct the employee or 8 agent to commit, the violation of this section; and 9 Documents that the employee or agent, at the time of initial employment and b. 10 each calendar year thereafter, participated in a training program approved by 11 the attorney general providing the employee or agent with information 12 regarding the state and federal regulations governing the sale, possession, 13 and packaging of such drugs. 14 The approval of the training program by the attorney general is not subject to chapter 28-32. 15 16 A political subdivision, including a home rule city or county, may not enact any 9. 17 ordinance relating to the sale by a retail distributor of over-the-counter products 18 containing ephedrine, pseudoephedrine, or phenylpropanolamine. Any existing 19 ordinance is void.