CHAPTER 152
DRUGS; CONTROLLED SUBSTANCES

DEFINITIONS AND SCHEDULES

OF CONTROLLED SUBSTANCES

152.01 DEFINITIONS.

Subdivision 1. **Words, terms, and phrases.** Unless the language or context clearly indicates that a different meaning is intended, the following words, terms, and phrases, for the purposes of this chapter, shall be given the meanings subjoined to them.

Subd. 2. **Drug.** The term "drug" includes all medicines and preparations recognized in the United States Pharmacopoeia or National Formulary and any substance or mixture of substances intended to be used for the cure, mitigation, or prevention of disease of either humans or other animals.

Subd. 3. MS 1967 [Repealed, 1969 c 933 s 22]

Subd. 3. **Administer.** "Administer" means to deliver by, or pursuant to the lawful order of a practitioner a single dose of a controlled substance to a patient or research subject by injection, inhalation, ingestion, or by any other immediate means.

Subd. 3a. **Cocaine.** "Cocaine" means coca leaves and any salt, compound, derivative, or preparation of coca leaves, including cocaine and ecgonine, the salts and isomers of cocaine and ecgonine, and the salts of their isomers and any salt, compound, derivative, or preparation thereof that is chemically equivalent or identical with any of those substances, except decocainized coca leaves or extraction of coca leaves, which extractions do not contain cocaine or ecgonine.

Subd. 4. MS 1967 [Repealed, 1969 c 933 s 22]

Subd. 4. **Controlled substance.** "Controlled substance" means a drug, substance, or immediate precursor in Schedules I through V of section 152.02. The term shall not include distilled spirits, wine, malt beverages, intoxicating liquors or tobacco.

Subd. 5. [Repealed, 1971 c 937 s 22]

Subd. 5a. **Hallucinogen.** "Hallucinogen" means any hallucinogen listed in section 152.02, subdivision 2, clause (3), or Minnesota Rules, part 6800.4210, item C, except marijuana and Tetrahydrocannabinols.

Subd. 6. **Pharmacist intern.** The term "pharmacist intern" means a natural person, a graduate of the College of Pharmacy, University of Minnesota, or other pharmacy college, approved by the board, or a person satisfactorily progressing toward the degree in pharmacy required for licensure, registered by the state Board of Pharmacy, for the purpose of obtaining practical experience as a requirement for licensure as a pharmacist or a qualified applicant, awaiting licensure.

Subd. 7. **Manufacture.** "Manufacture," in places other than a pharmacy, means and includes the production, cultivation, quality control, and standardization by mechanical, physical, chemical, or pharmaceutical means, packing, repacking, tableting, encapsulating, labeling, relabeling, filling, or by other process, of drugs.

Subd. 8. **Dispense.** "Dispense" means to deliver one or more doses of a controlled substance in a suitable container, properly labeled, for subsequent administration to, or use by a patient or research subject.

Subd. 9. **Marijuana.** "Marijuana" means all parts of the plant of any species of the genus Cannabis, including all agronomical varieties, whether growing or not; the seeds thereof; the resin extracted from any part of such plant; and every compound, manufacture, salt, derivative, mixture, or preparation of such plant, its seeds or resin, but shall not include the mature stalks

of such plant, fiber from such stalks, oil or cake made from the seeds of such plant, any other compound, manufacture, salt, derivative, mixture, or preparation of such mature stalks, except the resin extracted therefrom, fiber, oil, or cake, or the sterilized seed of such plant which is incapable of germination.

Subd. 9a. **Mixture.** "Mixture" means a preparation, compound, mixture, or substance containing a controlled substance, regardless of purity except as provided in subdivision 16; sections 152.021, subdivision 2, paragraph (b); 152.022, subdivision 2, paragraph (b); and 152.023, subdivision 2, paragraph (b).

Subd. 10. **Narcotic drug.** "Narcotic drug" means any of the following, whether produced directly or indirectly by extraction from substances of vegetable origin, or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis:

(1) opium, coca leaves, opiates, and methamphetamine;

(2) a compound, manufacture, salt, derivative, or preparation of opium, coca leaves, opiates, or methamphetamine;

(3) a substance, and any compound, manufacture, salt, derivative, or preparation thereof, which is chemically identical with any of the substances referred to in clauses (1) and (2), except that the words "narcotic drug" as used in this chapter shall not include decocainized coca leaves or extracts of coca leaves, which extracts do not contain cocaine or ecgonine.

Subd. 11. **Opiate.** "Opiate" means any dangerous substance having an addiction forming or addiction sustaining liability similar to morphine or being capable of conversion into a drug having such addiction forming or addiction sustaining liability.

Subd. 12. **Opium poppy.** "Opium poppy" means the plant of the species Papaver somniferum L., except the seeds thereof.

Subd. 12a. **Park zone.** "Park zone" means an area designated as a public park by the federal government, the state, a local unit of government, a park district board, or a park and recreation board in a city of the first class. "Park zone" includes the area within 300 feet or one city block, whichever distance is greater, of the park boundary.

Subd. 13. **Person.** "Person" includes every individual, copartnership, corporation or association of one or more individuals.

Subd. 14. **Poppy straw.** "Poppy straw" means all parts, except the seeds, of the opium poppy, after mowing.

Subd. 14a. **School zone.** "School zone" means:

(1) any property owned, leased, or controlled by a school district or an organization operating a nonpublic school, as defined in section 123B.41, subdivision 9, where an elementary, middle, secondary school, secondary vocational center or other school providing educational services in grade one through grade 12 is located, or used for educational purposes, or where extracurricular or cocurricular activities are regularly provided;

(2) the area surrounding school property as described in clause (1) to a distance of 300 feet or one city block, whichever distance is greater, beyond the school property; and

(3) the area within a school bus when that bus is being used to transport one or more elementary or secondary school students.

Subd. 15. **Immediate precursor.** "Immediate precursor" means a substance which the state Board of Pharmacy has found to be and by rule designates as being the principal compound

commonly used or produced for use, and which is an immediate chemical intermediary used or likely to be used in the manufacture of a controlled substance, the control of which is necessary to prevent, curtail, or limit such manufacture.

Subd. 15a. **Sell.** "Sell" means:

(1) to sell, give away, barter, deliver, exchange, distribute or dispose of to another, or to manufacture; or

(2) to offer or agree to perform an act listed in clause (1); or

(3) to possess with intent to perform an act listed in clause (1).

Subd. 16. **Small amount.** "Small amount" as applied to marijuana means 42.5 grams or less. This provision shall not apply to the resinous form of marijuana. The weight of fluid used in a water pipe may not be considered in determining a small amount except in cases where the marijuana is mixed with four or more fluid ounces of fluid.

Subd. 16a. **Subsequent controlled substance conviction.** Notwithstanding section 152.18, subdivision 1, a "subsequent controlled substance conviction" means that before commission of the offense for which the person is convicted under this chapter, the person received a disposition for a felony-level offense under section 152.18, subdivision 1, was convicted in Minnesota of a felony violation of this chapter or a felony-level attempt or conspiracy to violate this chapter, or was convicted elsewhere for conduct that would have been a felony under this chapter if committed in Minnesota. An earlier disposition for a felony-level offense under section 152.18, subdivision 1, or an earlier conviction is not relevant if ten years have elapsed since discharge from sentence or stay of adjudication.

Subd. 17. [Repealed, 1994 c 636 art 2 s 69]

Subd. 18. **Drug paraphernalia.** (a) Except as otherwise provided in paragraph (b), "drug paraphernalia" means all equipment, products, and materials of any kind, except those items used in conjunction with permitted uses of controlled substances under this chapter or the Uniform Controlled Substances Act, which are knowingly or intentionally used primarily in (1) manufacturing a controlled substance, (2) injecting, ingesting, inhaling, or otherwise introducing into the human body a controlled substance, (3) testing the strength, effectiveness, or purity of a controlled substance, or (4) enhancing the effect of a controlled substance.

(b) "Drug paraphernalia" does not include the possession, manufacture, delivery, or sale of hypodermic needles or syringes in accordance with section 151.40, subdivision 2.

Subd. 19. **Public housing zone.** "Public housing zone" means any public housing project or development administered by a local housing agency, plus the area within 300 feet of the property's boundary, or one city block, whichever distance is greater.

Subd. 20. **Unlawfully.** "Unlawfully" means selling or possessing a controlled substance in a manner not authorized by law.

Subd. 21. **Orphan drug.** "Orphan drug" means a drug for a disease or condition which is rare in the United States and has been designated as an orphan drug by the Secretary of Health and Human Services as provided in the Orphan Drug Act, Public Law 92-414, as amended.

Subd. 22. **Drug treatment facility.** "Drug treatment facility" means any facility in which a residential rehabilitation program licensed under Minnesota Rules, parts 9530.4100 to 9530.4450, is located, and includes any property owned, leased, or controlled by the facility.

Subd. 23. **Analog.** (a) Except as provided in paragraph (b), "analog" means a substance, the chemical structure of which is substantially similar to the chemical structure of a controlled substance in Schedule I or II:

(1) that has a stimulant, depressant, or hallucinogenic effect on the central nervous system that is substantially similar to or greater than the stimulant, depressant, or hallucinogenic effect on the central nervous system of a controlled substance in Schedule I or II; or

(2) with respect to a particular person, if the person represents or intends that the substance have a stimulant, depressant, or hallucinogenic effect on the central nervous system that is substantially similar to or greater than the stimulant, depressant, or hallucinogenic effect on the central nervous system of a controlled substance in Schedule I or II.

(b) "Analog" does not include:

(1) a controlled substance;

(2) any substance for which there is an approved new drug application under the Federal Food, Drug, and Cosmetic Act; or

(3) with respect to a particular person, any substance, if an exemption is in effect for investigational use, for that person, as provided by United States Code, title 21, section 355, and the person is registered as a controlled substance researcher as required under section 152.12, subdivision 3, to the extent conduct with respect to the substance is pursuant to the exemption and registration.

History: *(3899-2, 3899-5, 3899-7, 3906-12) 1921 c 190 s 2,5,7; 1939 c 102 s 2; 1967 c 408 s 1,2; 1971 c 937 s 1-11; Ex1971 c 38 s 1; Ex1971 c 48 s 17; 1973 c 693 s 1; 1979 c 157 s 1; 1981 c 37 s 2; 1981 c 295 s 1; 1982 c 557 s 1; 1982 c 642 s 22; 1985 c 248 s 70; 1986 c 444; 1987 c 298 s 1; 1989 c 290 art 3 s 1-7; 1991 c 279 s 1,2; 1992 c 359 s 1-3; 1993 c 82 s 1; 1997 c 239 art 4 s 1,2; 1998 c 397 art 11 s 3; 1999 c 98 s 1; 2005 c 136 art 7 s 2; 2011 c 53 s 1-3*

152.02 MS 1967 [Repealed, 1969 c 933 s 22]

152.02 SCHEDULES OF CONTROLLED SUBSTANCES; ADMINISTRATION OF CHAPTER.

Subdivision 1. **Five schedules.** There are established five schedules of controlled substances, to be known as Schedules I, II, III, IV, and V. The schedules consist of the substances listed in this section by whatever official name, common or usual name, chemical name, or trade name designated.

Subd. 2. **Schedule I.** (a) Schedule I consists of the substances listed in this subdivision.

(b) Opiates. Unless specifically excepted or unless listed in another schedule, any of the following substances, including their analogs, isomers, esters, ethers, salts, and salts of isomers, esters, and ethers, whenever the existence of the analogs, isomers, esters, ethers, and salts is possible:

(1) acetylmethadol;

(2) allylprodine;

(3) alphacetylmethadol (except levo-alphacetylmethadol, also known as levomethadyl acetate);

(4) alphameprodine;

(5) alphamethadol;

(6) alpha-methylfentanyl benzethidine;

(7) betacetylmethadol;

(8) betameprodine;

(9) betamethadol;

(10) betaprodine;

(11) clonitazene;

(12) dextromoramide;

(13) diampromide;

(14) diethyliambutene;

(15) difenoxin;

(16) dimenoxadol;

(17) dimepheptanol;

(18) dimethyliambutene;

(19) dioxaphetyl butyrate;

(20) dipipanone;

(21) ethylmethylthiambutene;

(22) etonitazene;

(23) etoxeridine;

(24) furethidine;

(25) hydroxypethidine;

(26) ketobemidone;

(27) levomoramide;

(28) levophenacylmorphan;

(29) 3-methylfentanyl;

(30) acetyl-alpha-methylfentanyl;

(31) alpha-methylthiofentanyl;

(32) benzylfentanyl beta-hydroxyfentanyl;

(33) beta-hydroxy-3-methylfentanyl;

(34) 3-methylthiofentanyl;

(35) thenylfentanyl;

(36) thiofentanyl;

(37) para-fluorofentanyl;

(38) morpheridine;

(39) 1-methyl-4-phenyl-4-propionoxypiperidine;

(40) noracymethadol;

(41) norlevorphanol;

(42) normethadone;

(43) norpipanone;

(44) 1-(2-phenylethyl)-4-phenyl-4-acetoxypiperidine (PEPAP);

(45) phenadoxone;

(46) phenampromide;

(47) phenomorphan;

(48) phenoperidine;

(49) piritramide;

(50) proheptazine;

(51) properidine;

(52) propiram;

(53) racemoramide;

(54) tilidine;

(55) trimeperidine.

(c) Opium derivatives. Any of the following substances, their analogs, salts, isomers, and salts of isomers, unless specifically excepted or unless listed in another schedule, whenever the existence of the analogs, salts, isomers, and salts of isomers is possible:

(1) acetorphine;

(2) acetyldihydrocodeine;

(3) benzylmorphine;

(4) codeine methylbromide;

(5) codeine-n-oxide;

(6) cyprenorphine;

(7) desomorphine;

(8) dihydromorphine;

(9) drotebanol;

(10) etorphine;

(11) heroin;

(12) hydromorphinol;

(13) methyldesorphine;

(14) methyldihydromorphine;

(15) morphine methylbromide;

(16) morphine methylsulfonate;

(17) morphine-n-oxide;

(18) myrophine;

(19) nicocodeine;

(20) nicomorphine;

(21) normorphine;

(22) pholcodine;

(23) thebacon.

(d) Hallucinogens. Any material, compound, mixture or preparation which contains any quantity of the following substances, their analogs, salts, isomers (whether optical, positional, or geometric), and salts of isomers, unless specifically excepted or unless listed in another schedule, whenever the existence of the analogs, salts, isomers, and salts of isomers is possible:

(1) methylenedioxy amphetamine;

(2) methylenedioxymethamphetamine;

(3) methylenedioxy-N-ethylamphetamine (MDEA);

(4) n-hydroxy-methylenedioxyamphetamine;

(5) 4-bromo-2,5-dimethoxyamphetamine (DOB);

(6) 2,5-dimethoxyamphetamine (2,5-DMA);

(7) 4-methoxyamphetamine;

(8) 5-methoxy-3, 4-methylenedioxy amphetamine;

(9) alpha-ethyltryptamine;

(10) bufotenine;

(11) diethyltryptamine;

(12) dimethyltryptamine;

(13) 3,4,5-trimethoxy amphetamine;

(14) 4-methyl-2, 5-dimethoxyamphetamine (DOM);

(15) ibogaine;

(16) lysergic acid diethylamide (LSD);

(17) mescaline;

(18) parahexyl;

(19) N-ethyl-3-piperidyl benzilate;

(20) N-methyl-3-piperidyl benzilate;

(21) psilocybin;

(22) psilocyn;

(23) tenocyclidine (TPCP or TCP);

(24) N-ethyl-1-phenyl-cyclohexylamine (PCE);

(25) 1-(1-phenylcyclohexyl) pyrrolidine (PCPy);

(26) 1-[1-(2-thienyl)cyclohexyl]-pyrrolidine (TCPy);

(27) 4-chloro-2,5-dimethoxyamphetamine (DOC);

(28) 4-ethyl-2,5-dimethoxyamphetamine (DOET);

(29) 4-iodo-2,5-dimethoxyamphetamine (DOI);

(30) 4-bromo-2,5-dimethoxyphenethylamine (2C-B);

(31) 4-chloro-2,5-dimethoxyphenethylamine (2C-C);

(32) 4-methyl-2,5-dimethoxyphenethylamine (2-CD);

(33) 4-ethyl-2,5-dimethoxyphenethylamine (2C-E);

(34) 4-iodo-2,5-dimethoxyphenethylamine (2C-I);

(35) 4-propyl-2,5-dimethoxyphenethylamine (2C-P);

(36) 4-isopropylthio-2,5-dimethoxyphenethylamine (2C-T-4);

(37) 4-propylthio-2,5-dimethoxyphenethylamine (2C-T-7);

(38) 2-(8-bromo-2,3,6,7-tetrahydrofuro [2,3-f][1]benzofuran-4-yl)ethanamine (2-CB-FLY);

(39) bromo-benzodifuranyl-isopropylamine (Bromo-DragonFLY);

(40) alpha-methyltryptamine (AMT);

(41) N,N-diisopropyltryptamine (DiPT);

(42) 4-acetoxy-N,N-dimethyltryptamine (4-AcO-DMT);

(43) 4-acetoxy-N,N-diethyltryptamine (4-AcO-DET);

(44) 4-hydroxy-N-methyl-N-propyltryptamine (4-HO-MPT);

(45) 4-hydroxy-N,N-dipropyltryptamine (4-HO-DPT);

(46) 4-hydroxy-N,N-diallyltryptamine (4-HO-DALT);

(47) 4-hydroxy-N,N-diisopropyltryptamine (4-HO-DiPT);

(48) 5-methoxy-N,N-diisopropyltryptamine (5-MeO-DiPT);

(49) 5-methoxy-α-methyltryptamine (5-MeO-AMT);

(50) 5-methoxy-N,N-dimethyltryptamine (5-MeO-DMT);

(51) 5-methylthio-N,N-dimethyltryptamine (5-MeS-DMT);

(52) 5-methoxy-N-methyl-N-propyltryptamine (5-MeO-MiPT);

(53) 5-methoxy-α-ethyltryptamine (5-MeO-AET);

(54) 5-methoxy-N,N-dipropyltryptamine (5-MeO-DPT);

(55) 5-methoxy-N,N-diethyltryptamine (5-MeO-DET);

(56) 5-methoxy-N,N-diallytryptamine (5-MeO-DALT);

(57) methoxetamine (MXE);

(58) 5-iodo-2-aminoindane (5-IAI);

(59) 5,6-methylenedioxy-2-aminoindane (MDAI).

(e) Peyote. All parts of the plant presently classified botanically as Lophophora williamsii Lemaire, whether growing or not, the seeds thereof, any extract from any part of the plant, and every compound, manufacture, salts, derivative, mixture, or preparation of the plant, its seeds or extracts. The listing of peyote as a controlled substance in Schedule I does not apply to the nondrug use of peyote in bona fide religious ceremonies of the American Indian Church, and members of the American Indian Church are exempt from registration. Any person who manufactures peyote for or distributes peyote to the American Indian Church, however, is required to obtain federal registration annually and to comply with all other requirements of law.

(f) Central nervous system depressants. Unless specifically excepted or unless listed in another schedule, any material compound, mixture, or preparation which contains any quantity of the following substances, their analogs, salts, isomers, and salts of isomers whenever the existence of the analogs, salts, isomers, and salts of isomers is possible:

(1) mecloqualone;

(2) methaqualone;

(3) gamma-hydroxybutyric acid (GHB), including its esters and ethers;

(4) flunitrazepam.

(g) Stimulants. Unless specifically excepted or unless listed in another schedule, any material compound, mixture, or preparation which contains any quantity of the following substances, their analogs, salts, isomers, and salts of isomers whenever the existence of the analogs, salts, isomers, and salts of isomers is possible:

(1) aminorex;

(2) cathinone;

(3) fenethylline;

(4) methcathinone;

(5) methylaminorex;

(6) N,N-dimethylamphetamine;

(7) N-benzylpiperazine (BZP);

(8) methylmethcathinone (mephedrone);

(9) 3,4-methylenedioxy-N-methylcathinone (methylone);

(10) methoxymethcathinone (methedrone);

(11) methylenedioxypyrovalerone (MDPV);

(12) fluoromethcathinone;

(13) methylethcathinone (MEC);

(14) 1-benzofuran-6-ylpropan-2-amine (6-APB);

(15) dimethylmethcathinone (DMMC);

(16) fluoroamphetamine;

(17) fluoromethamphetamine;

(18) α-methylaminobutyrophenone (MABP or buphedrone);

(19) β-keto-N-methylbenzodioxolylpropylamine (bk-MBDB or butylone);

(20) 2-(methylamino)-1-(4-methylphenyl)butan-1-one (4-MEMABP or BZ-6378);

(21) naphthylpyrovalerone (naphyrone); and

(22) any other substance, except bupropion or compounds listed under a different schedule, that is structurally derived from 2-aminopropan-1-one by substitution at the 1-position with either phenyl, naphthyl, or thiophene ring systems, whether or not the compound is further modified in any of the following ways:

(i) by substitution in the ring system to any extent with alkyl, alkylenedioxy, alkoxy, haloalkyl, hydroxyl, or halide substituents, whether or not further substituted in the ring system by one or more other univalent substituents;

(ii) by substitution at the 3-position with an acyclic alkyl substituent;

(iii) by substitution at the 2-amino nitrogen atom with alkyl, dialkyl, benzyl, or methoxybenzyl groups; or

(iv) by inclusion of the 2-amino nitrogen atom in a cyclic structure.

(h) Marijuana, tetrahydrocannabinols, and synthetic cannabinoids. Unless specifically excepted or unless listed in another schedule, any natural or synthetic material, compound, mixture, or preparation that contains any quantity of the following substances, their analogs, isomers, esters, ethers, salts, and salts of isomers, esters, and ethers, whenever the existence of the isomers, esters, ethers, or salts is possible:

(1) marijuana;

(2) tetrahydrocannabinols naturally contained in a plant of the genus Cannabis, synthetic equivalents of the substances contained in the cannabis plant or in the resinous extractives of the plant, or synthetic substances with similar chemical structure and pharmacological activity to those substances contained in the plant or resinous extract, including, but not limited to, 1 cis or trans tetrahydrocannabinol, 6 cis or trans tetrahydrocannabinol, and 3,4 cis or trans tetrahydrocannabinol;

(3) synthetic cannabinoids, including the following substances:

(i) Naphthoylindoles, which are any compounds containing a 3-(1-napthoyl)indole structure with substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl or 2-(4-morpholinyl)ethyl group, whether or not further substituted in the indole ring to any extent and whether or not substituted in the naphthyl ring to any extent. Examples of naphthoylindoles include, but are not limited to:

(A) 1-Pentyl-3-(1-naphthoyl)indole (JWH-018 and AM-678);

(B) 1-Butul-3-(1-naphthoyl)indole (JWH-073);

(C) 1-Pentyl-3-(4-methoxy-1-naphthoyl)indole (JWH-081);

(D) 1-[2-(4-morpholinyl)ethyl]-3-(1-naphthoyl)indole (JWH-200);

(E) 1-Propyl-2-methyl-3-(1-naphthoyl)indole (JWH-015);

(F) 1-Hexyl-3-(1-naphthoyl)indole (JWH-019);

(G) 1-Pentyl-3-(4-methyl-1-naphthoyl)indole (JWH-122);

(H) 1-Pentyl-3-(4-ethyl-1-naphthoyl)indole (JWH-210);

(I) 1-Pentyl-3-(4-chloro-1-naphthoyl)indole (JWH-398);

(J) 1-(5-fluoropentyl)-3-(1-naphthoyl)indole (AM-2201).

(ii) Napthylmethylindoles, which are any compounds containing a 1H-indol-3-yl-(1-naphthyl)methane structure with substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl or 2-(4-morpholinyl)ethyl group, whether or not further

substituted in the indole ring to any extent and whether or not substituted in the naphthyl ring to any extent. Examples of naphthylmethylindoles include, but are not limited to:

(A) 1-Pentyl-1H-indol-3-yl-(1-naphthyl)methane (JWH-175);

(B) 1-Pentyl-1H-indol-3-yl-(4-methyl-1-naphthyl)methan (JWH-184).

(iii) Naphthoylpyrroles, which are any compounds containing a 3-(1-naphthoyl)pyrrole structure with substitution at the nitrogen atom of the pyrrole ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl or 2-(4-morpholinyl)ethyl group whether or not further substituted in the pyrrole ring to any extent, whether or not substituted in the naphthyl ring to any extent. Examples of naphthoylpyrroles include, but are not limited to, (5-(2-fluorophenyl)-1-pentylpyrrol-3-yl)-naphthalen-1-ylmethanone (JWH-307).

(iv) Naphthylmethylindenes, which are any compounds containing a naphthylideneindene structure with substitution at the 3-position of the indene ring by an allkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl or 2-(4-morpholinyl)ethyl group whether or not further substituted in the indene ring to any extent, whether or not substituted in the naphthyl ring to any extent. Examples of naphthylemethylindenes include, but are not limited to, E-1-[1-(1-naphthalenylmethylene)-1H-inden-3-yl]pentane (JWH-176).

(v) Phenylacetylindoles, which are any compounds containing a 3-phenylacetylindole structure with substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl or 2-(4-morpholinyl)ethyl group whether or not further substituted in the indole ring to any extent, whether or not substituted in the phenyl ring to any extent. Examples of phenylacetylindoles include, but are not limited to:

(A) 1-(2-cyclohexylethyl)-3-(2-methoxyphenylacetyl)indole (RCS-8);

(B) 1-pentyl-3-(2-methoxyphenylacetyl)indole (JWH-250);

(C) 1-pentyl-3-(2-methylphenylacetyl)indole (JWH-251);

(D) 1-pentyl-3-(2-chlorophenylacetyl)indole (JWH-203).

(vi) Cyclohexylphenols, which are compounds containing a 2-(3-hydroxycyclohexyl)phenol structure with substitution at the 5-position of the phenolic ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl or 2-(4-morpholinyl)ethyl group whether or not substituted in the cyclohexyl ring to any extent. Examples of cyclohexylphenols include, but are not limited to:

(A) 5-(1,1-dimethylheptyl)-2-[(1R,3S)-3-hydroxycyclohexyl]-phenol (CP 47,497);

(B) 5-(1,1-dimethyloctyl)-2-[(1R,3S)-3-hydroxycyclohexyl]-phenol (Cannabicyclohexanol or CP 47,497 C8 homologue);

(C) 5-(1,1-dimethylheptyl)-2-[(1R,2R)-5-hydroxy-2-(3-hydroxypropyl)cyclohexyl] -phenol (CP 55,940).

(vii) Benzoylindoles, which are any compounds containing a 3-(benzoyl)indole structure with substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl or 2-(4-morpholinyl)ethyl group whether or not further substituted in the indole ring to any extent and whether or not substituted in the phenyl ring to any extent. Examples of benzoylindoles include, but are not limited to:

(A) 1-Pentyl-3-(4-methoxybenzoyl)indole (RCS-4);

(B) 1-(5-fluoropentyl)-3-(2-iodobenzoyl)indole (AM-694);

(C) (4-methoxyphenyl-[2-methyl-1-(2-(4-morpholinyl)ethyl)indol-3-yl]methanone (WIN 48,098 or Pravadoline).

(viii) Others specifically named:

(A) (6aR,10aR)-9-(hydroxymethyl)-6,6-dimethyl-3-(2-methyloctan-2-yl) -6a,7,10,10a-tetrahydrobenzo[c]chromen-1-ol (HU-210);

(B) (6aS,10aS)-9-(hydroxymethyl)-6,6-dimethyl-3-(2-methyloctan-2-yl) -6a,7,10,10a-tetrahydrobenzo[c]chromen-1-ol (Dexanabinol or HU-211);

(C) 2,3-dihydro-5-methyl-3-(4-morpholinylmethyl)pyrrolo[1,2,3-de] -1,4-benzoxazin-6-yl-1-naphthalenylmethanone (WIN 55,212-2).

(i) A controlled substance analog, to the extent that it is implicitly or explicitly intended for human consumption.

Subd. 3. **Schedule II.** (a) Schedule II consists of the substances listed in this subdivision.

(b) Unless specifically excepted or unless listed in another schedule, any of the following substances whether produced directly or indirectly by extraction from substances of vegetable origin or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis:

(1) Opium and opiate, and any salt, compound, derivative, or preparation of opium or opiate.

(i) Excluding:

(A) apomorphine;

(B) thebaine-derived butorphanol;

(C) dextrophan;

(D) nalbuphine;

(E) nalmefene;

(F) naloxone;

(G) naltrexone;

(H) and their respective salts;

(ii) but including the following:

(A) opium, in all forms and extracts;

(B) codeine;

(C) dihydroetorphine;

(D) ethylmorphine;

(E) etorphine hydrochloride;

(F) hydrocodone;

(G) hydromorphone;

(H) metopon;

(I) morphine;

(J) oxycodone;

(K) oxymorphone;

(L) thebaine;

(M) oripavine;

(2) any salt, compound, derivative, or preparation thereof which is chemically equivalent or identical with any of the substances referred to in clause (1), except that these substances shall not include the isoquinoline alkaloids of opium;

(3) opium poppy and poppy straw;

(4) coca leaves and any salt, cocaine compound, derivative, or preparation of coca leaves (including cocaine and ecgonine and their salts, isomers, derivatives, and salts of isomers and derivatives), and any salt, compound, derivative, or preparation thereof which is chemically equivalent or identical with any of these substances, except that the substances shall not include decocainized coca leaves or extraction of coca leaves, which extractions do not contain cocaine or ecgonine;

(5) concentrate of poppy straw (the crude extract of poppy straw in either liquid, solid, or powder form which contains the phenanthrene alkaloids of the opium poppy).

(c) Any of the following opiates, including their isomers, esters, ethers, salts, and salts of isomers, esters and ethers, unless specifically excepted, or unless listed in another schedule, whenever the existence of such isomers, esters, ethers and salts is possible within the specific chemical designation:

(1) alfentanil;

(2) alphaprodine;

(3) anileridine;

(4) bezitramide;

(5) bulk dextropropoxyphene (nondosage forms);

(6) carfentanil;

(7) dihydrocodeine;

(8) dihydromorphinone;

(9) diphenoxylate;

(10) fentanyl;

(11) isomethadone;

(12) levo-alpha-acetylmethadol (LAAM) levomethorphan;

(13) levorphanol;

(14) metazocine;

(15) methadone;

(16) methadone - intermediate, 4-cyano-2-dimethylamino-4, 4-diphenylbutane;

(17) moramide - intermediate, 2-methyl-3-morpholino-1, 1-diphenyl-propane-carboxylic acid;

(18) pethidine;

(19) pethidine - intermediate - a, 4-cyano-1-methyl-4-phenylpiperidine;

(20) pethidine - intermediate - b, ethyl-4-phenylpiperidine-4-carboxylate;

(21) pethidine - intermediate - c, 1-methyl-4-phenylpiperidine-4-carboxylic acid;

(22) phenazocine;

(23) piminodine;

(24) racemethorphan;

(25) racemorphan;

(26) remifentanil;

(27) sufentanil;

(28) tapentadol.

(d) Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation which contains any quantity of the following substances having a stimulant effect on the central nervous system:

(1) amphetamine, its salts, optical isomers, and salts of its optical isomers;

(2) methamphetamine, its salts, isomers, and salts of its isomers;

(3) phenmetrazine and its salts;

(4) methylphenidate;

(5) lisdexamfetamine.

(e) Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation which contains any quantity of the following substances having a depressant effect on the central nervous system, including its salts, isomers, and salts of isomers whenever the existence of such salts, isomers, and salts of isomers is possible within the specific chemical designation:

(1) amobarbital;

(2) glutethimide;

(3) secobarbital;

(4) pentobarbital;

(5) phencyclidine;

(6) phencyclidine immediate precursors:

(i) 1-phenylcyclohexylamine;

(ii) 1-piperidinocyclohexanecarbonitrile;

(7) phenylacetone.

(f) Hallucinogenic substances: nabilone.

Subd. 4. **Schedule III.** (a) Schedule III consists of the substances listed in this subdivision.

(b) Stimulants. Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation which contains any quantity of the following substances having a potential for abuse associated with a stimulant effect on the central nervous system, including its salts, isomers, and salts of such isomers whenever the existence of such salts, isomers, and salts of isomers is possible within the specific chemical designation:

(1) benzphetamine;

(2) chlorphentermine;

(3) clortermine;

(4) phendimetrazine.

(c) Depressants. Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation which contains any quantity of the following substances having a potential for abuse associated with a depressant effect on the central nervous system:

(1) any compound, mixture, or preparation containing amobarbital, secobarbital, pentobarbital or any salt thereof and one or more other active medicinal ingredients which are not listed in any schedule;

(2) any suppository dosage form containing amobarbital, secobarbital, pentobarbital, or any salt of any of these drugs and approved by the food and drug administration for marketing only as a suppository;

(3) any substance which contains any quantity of a derivative of barbituric acid, or any salt of a derivative of barbituric acid, except those substances which are specifically listed in other schedules;

(4) any drug product containing gamma hydroxybutyric acid, including its salts, isomers, and salts of isomers, for which an application is approved under section 505 of the federal Food, Drug, and Cosmetic Act;

(5) any of the following substances:

(i) chlorhexadol;

(ii) ketamine, its salts, isomers and salts of isomers;

(iii) lysergic acid;

(iv) lysergic acid amide;

(v) methyprylon;

(vi) sulfondiethylmethane;

(vii) sulfonenthylmethane;

(viii) sulfonmethane;

(ix) tiletamine and zolazepam and any salt thereof;

(x) embutramide.

(d) Nalorphine.

(e) Narcotic drugs. Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation containing any of the following narcotic drugs, or their salts calculated as the free anhydrous base or alkaloid, in limited quantities as follows:

(1) not more than 1.80 grams of codeine per 100 milliliters or not more than 90 milligrams per dosage unit, with an equal or greater quantity of an isoquinoline alkaloid of opium;

(2) not more than 1.80 grams of codeine per 100 milliliters or not more than 90 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts;

(3) not more than 300 milligrams of dihydrocodeinone per 100 milliliters or not more than 15 milligrams per dosage unit, with a fourfold or greater quantity of an isoquinoline alkaloid of opium;

(4) not more than 300 milligrams of dihydrocodeinone per 100 milliliters or not more than 15 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts;

(5) not more than 1.80 grams of dihydrocodeine per 100 milliliters or not more than 90 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts;

(6) not more than 300 milligrams of ethylmorphine per 100 milliliters or not more than 15 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts;

(7) not more than 500 milligrams of opium per 100 milliliters or per 100 grams, or not more than 25 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts;

(8) not more than 50 milligrams of morphine per 100 milliliters or per 100 grams with one or more active, nonnarcotic ingredients in recognized therapeutic amounts;

(f) Anabolic steroids and human growth hormone.

(1) Anabolic steroids, for purposes of this subdivision, means any drug or hormonal substance, chemically and pharmacologically related to testosterone, other than estrogens, progestins, corticosteroids, and dehydroepiandrosterone, and includes:

(i) 3[beta],17[beta]-dihydroxy-5[alpha]-androstane;

(ii) 3[alpha],17[beta]-dihydroxy-5[alpha]-androstane;

(iii) androstanedione (5[alpha]-androstan-3,17-dione);

(iv) 1-androstenediol (3[beta],17[beta]-dihydroxy-5[alpha]-androst-l-ene;

(v) 3[alpha],17[beta]-dihydroxy-5[alpha]-androst-1-ene);

(vi) 4-androstenediol (3[beta],17[beta]-dihydroxy-androst-4-ene);

(vii) 5-androstenediol (3[beta],17[beta]-dihydroxy-androst-5-ene);

(viii) 1-androstenedione (5[alpha]-androst-1-en-3,17-dione);

(ix) 4-androstenedione (androst-4-en-3,17-dione);

(x) 5-androstenedione (androst-5-en-3,17-dione);

(xi) bolasterone (7[alpha],17[alpha]-dimethyl-17[beta]-hydroxyandrost-4-en-3-one);

(xii) boldenone (17[beta]-hydroxyandrost-1,4-diene-3-one);

(xiii) boldione (androsta-1,4-diene-3,17-dione);

(xiv) calusterone (7[beta],17[alpha]-dimethyl-17[beta]-hydroxyandrost-4-en-3-one);

(xv) clostebol (4-chloro-17[beta]-hydroxyandrost-4-en-3-one);

(xvi) dehydrochloromethyltestosterone (4-chloro-17[beta]-hydroxy-17[alpha]-methylandrost-1,4-dien-3-one);

(xvii) desoxymethyltestosterone (17[alpha]-methyl-5[alpha]-androst-2-en-17[beta]-ol);

(xviii) [delta]1-dihydrotestosterone- (17[beta]-hydroxy-5[alpha]-androst-1-en-3-one);

(xix) 4-dihydrotestosterone (17[beta]-hydroxy-androstan-3-one);

(xx) drostanolone (17[beta]hydroxy-2[alpha]-methyl-5[alpha]-androstan-3-one);

(xxi) ethylestrenol (17[alpha]-ethyl-17[beta]-hydroxyestr-4-ene);

(xxii) fluoxymesterone (9-fluoro-17[alpha]-methyl-11[beta],17[beta]-dihydroxyandrost-4-en-3-one);

(xxiii) formebolone (2-formyl-17[alpha]-methyl-11[alpha],17[beta]-dihydroxyandrost-1,4-dien-3-one);

(xxiv) furazabol (17[alpha]-methyl-17[beta]-hydroxyandrostano[2,3-c]-furazan)13[beta]-ethyl-17[beta] -hydroxygon-4-en-3-one;

(xxv) 4-hydroxytestosterone (4,17[beta]-dihydroxyandrost-4-en-3-one);

(xxvi) 4-hydroxy-19-nortestosterone (4,17[beta]-dihydroxyestr-4-en-3-one);

(xxvii) mestanolone (17[alpha]-methyl-17[beta]-hydroxy-5[alpha]-androstan-3-one);

(xxviii) mesterolone (1[alpha]-methyl-17[beta]-hydroxy-5[alpha]-androstan-3-one);

(xxix) methandienone (17[alpha]-methyl-17[beta]-hydroxyandrost-1,4-dien-3-one);

(xxx) methandriol (17[alpha]-methyl-3[beta],17[beta]-dihydroxyandrost-5-ene);

(xxxi) methenolone (1-methyl-17[beta]-hydroxy-5[alpha]-androst-1-en-3-one);

(xxxii) 17[alpha]-methyl-3[beta],17[beta]-dihydroxy-5[alpha]-androstane;

(xxxiii) 17[alpha]-methyl-3[alpha],17[beta]-dihydroxy-5[alpha]-androstane;

(xxxiv) 17[alpha]-methyl-3[beta],17[beta]-dihydroxyandrost-4-ene;

(xxxv) 17[alpha]-methyl-4-hydroxynandrolone (17[alpha]-methyl-4-hydroxy-17[beta]-hydroxyestr-4-en-3-one);

(xxxvi) methyldienolone (17[alpha]-methyl-17[beta]-hydroxyestra-4,9(10)-dien-3-one);

(xxxvii) methyltrienolone (17[alpha]-methyl-17[beta]-hydroxyestra-4,9-11-trien-3-one);

(xxxviii) methyltestosterone (17[alpha]-methyl-17[beta]-hydroxyandrost-4-en-3-one);

(xxxix) mibolerone (7[alpha],17[alpha]-dimethyl-17[beta]-hydroxyestr-4-en-3-one);

(xl) 17[alpha]-methyl-[delta]1-dihydrotestosterone (17[beta]-hydroxy-17[alpha]-methyl-5[alpha]-androst-1-en-3-one);

(xli) nandrolone (17[beta]-hydroxyestr-4-en-3-one);

(xlii) 19-nor-4-androstenediol (3[beta],17[beta]-dihydroxyestr-4-ene;

(xliii) 3[alpha],17[beta]-dihydroxyestr-4-ene); 19-nor-5-androstenediol (3[beta],17[beta]-dihydroxyestr-5-ene;

(xliv) 3[alpha],17[beta]-dihydroxyestr-5-ene);

(xlv) 19-nor-4,9(10)-androstadienedione (estra-4,9(10)-diene-3,17-dione);

(xlvi) 19-nor-5-androstenedione (estr-5-en-3,17-dione);

(xlvii) norbolethone (13[beta],17[alpha]-diethyl-17[beta]-hydroxygon-4-en-3-one);

(xlviii) norclostebol (4-chloro-17[beta]-hydroxyestr-4-en-3-one);

(xlix) norethandrolone (17[alpha]-ethyl-17[beta]-hydroxyestr-4-en-3-one);

(l) normethandrolone (17[alpha]-methyl-17[beta]-hydroxyestr-4-en-3-one);

(li) oxandrolone (17[alpha]-methyl-17[beta]-hydroxy-2-oxa-5[alpha]-androstan-3-one);

(lii) oxymesterone (17[alpha]-methyl-4,17[beta]-dihydroxyandrost-4-en-3-one);

(liii) oxymetholone (17[alpha]-methyl-2-hydroxymethylene-17[beta]-hydroxy-5[alpha]-androstan-3-one);

(liv) stanozolol (17[alpha]-methyl-17[beta]-hydroxy-5[alpha]-androst-2-eno[3,2-c]-pyrazole);

(lv) stenbolone (17[beta]-hydroxy-2-methyl-5[alpha]-androst-1-en-3-one);

(lvi) testolactone (13-hydroxy-3-oxo-13,17-secoandrosta-1,4-dien-17-oic acid lactone);

(lvii) testosterone (17[beta]-hydroxyandrost-4-en-3-one);

(lviii) tetrahydrogestrinone (13[beta],17[alpha]-diethyl-17[beta]-hydroxygon-4,9,11-trien-3-one);

(lix) trenbolone (17[beta]-hydroxyestr-4,9,11-trien-3-one);

(lx) any salt, ester, or ether of a drug or substance described in this paragraph.

Anabolic steroids are not included if they are: (A) expressly intended for administration through implants to cattle or other nonhuman species; and (B) approved by the United States Food and Drug Administration for that use;

(2) Human growth hormones.

(g) Hallucinogenic substances. Dronabinol (synthetic) in sesame oil and encapsulated in a soft gelatin capsule in a United States Food and Drug Administration approved product.

(h) Any material, compound, mixture, or preparation containing the following narcotic drug or its salt: buprenorphine.

Subd. 5. **Schedule IV.** (a) Schedule IV consists of the substances listed in this subdivision.

(b) Narcotic drugs. Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation containing any of the following narcotic drugs, or their salts calculated as the free anhydrous base or alkaloid, in limited quantities as follows:

(1) not more than one milligram of difenoxin and not less than 25 micrograms of atropine sulfate per dosage unit;

(2) dextropropoxyphene (Darvon and Darvocet).

(c) Depressants. Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation containing any quantity of the following substances, including its salts, isomers, and salts of isomers whenever the existence of the salts, isomers, and salts of isomers is possible:

(1) alprazolam;

(2) barbital;

(3) bromazepam;

(4) camazepam;

(5) carisoprodol;

(6) chloral betaine;

(7) chloral hydrate;
(8) chlordiazepoxide;
(9) clobazam;
(10) clonazepam;
(11) clorazepate;
(12) clotiazepam;
(13) cloxazolam;
(14) delorazepam;
(15) diazepam;
(16) dichloralphenazone;
(17) estazolam;
(18) ethchlorvynol;
(19) ethinamate;
(20) ethyl loflazepate;
(21) fludiazepam;
(22) flurazepam;
(23) halazepam;
(24) haloxazolam;
(25) ketazolam;
(26) loprazolam;
(27) lorazepam;
(28) lormetazepam mebutamate;
(29) medazepam;
(30) meprobamate;
(31) methohexital;
(32) methylphenobarbital;
(33) midazolam;
(34) nimetazepam;
(35) nitrazepamnordiazepam;
(36) oxazepam;
(37) oxazolam;
(38) paraldehydepetrichloral;
(39) phenobarbital;
(40) pinazepam;
(41) prazepam;
(42) quazepam;

(43) temazepam;

(44) tetrazepam;

(45) triazolam;

(46) zaleplon;

(47) zolpidem;

(48) zopiclone.

(d) Any material, compound, mixture, or preparation which contains any quantity of the following substance including its salts, isomers, and salts of such isomers, whenever the existence of such salts, isomers, and salts of isomers is possible: fenfluramine.

(e) Stimulants. Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation which contains any quantity of the following substances having a stimulant effect on the central nervous system, including its salts, isomers, and salts of isomers:

(1) cathine (norpseudoephedrine);

(2) diethylpropion;

(3) fencamfamine;

(4) fenproporex;

(5) mazindol;

(6) mefenorex;

(7) modafinil;

(8) pemoline (including organometallic complexes and chelates thereof);

(9) phentermine;

(10) pipradol;

(11) sibutramine;

(12) SPA (1-dimethylamino-1,2-diphenylethane).

Subd. 6. **Schedule V; restrictions on methamphetamine precursor drugs.** (a) As used in this subdivision, the following terms have the meanings given:

(1) "methamphetamine precursor drug" means any compound, mixture, or preparation intended for human consumption containing ephedrine or pseudoephedrine as its sole active ingredient or as one of its active ingredients; and

(2) "over-the-counter sale" means a retail sale of a drug or product but does not include the sale of a drug or product pursuant to the terms of a valid prescription.

(b) The following items are listed in Schedule V:

(1) any compound, mixture, or preparation containing any of the following limited quantities of narcotic drugs, which shall include one or more nonnarcotic active medicinal ingredients in sufficient proportion to confer upon the compound, mixture or preparation valuable medicinal qualities other than those possessed by the narcotic drug alone:

(i) not more than 100 milligrams of dihydrocodeine per 100 milliliters or per 100 grams;

(ii) not more than 100 milligrams of ethylmorphine per 100 milliliters or per 100 grams;

(iii) not more than 2.5 milligrams of diphenoxylate and not less than 25 micrograms of atropine sulfate per dosage unit;

(iv) not more than 100 milligrams of opium per 100 milliliters or per 100 grams; or

(v) not more than 0.5 milligrams of difenoxin and not less than 25 micrograms of atropine sulfate per dosage unit.

(2) Stimulants. Unless specifically exempted or excluded or unless listed in another schedule, any material, compound, mixture, or preparation that contains any quantity of the following substance having a stimulant effect on the central nervous system, including its salts, isomers, and salts of isomers: pyrovalerone.

(3) Depressants. Unless specifically exempted or excluded or unless listed in another schedule, any material, compound, mixture, or preparation that contains any quantity of the following substance having a depressant effect on the central nervous system, including its salts, isomers, and salts of isomers:

(i) pregabalin;

(ii) lacosamide.

(4) Any compound, mixture, or preparation containing ephedrine or pseudoephedrine as its sole active ingredient or as one of its active ingredients.

(c) No person may sell in a single over-the-counter sale more than two packages of a methamphetamine precursor drug or a combination of methamphetamine precursor drugs or any combination of packages exceeding a total weight of six grams, calculated as the base.

(d) Over-the-counter sales of methamphetamine precursor drugs are limited to:

(1) packages containing not more than a total of three grams of one or more methamphetamine precursor drugs, calculated in terms of ephedrine base or pseudoephedrine base; or

(2) for nonliquid products, sales in blister packs, where each blister contains not more than two dosage units, or, if the use of blister packs is not technically feasible, sales in unit dose packets or pouches.

(e) A business establishment that offers for sale methamphetamine precursor drugs in an over-the-counter sale shall ensure that all packages of the drugs are displayed behind a checkout counter where the public is not permitted and are offered for sale only by a licensed pharmacist, a registered pharmacy technician, or a pharmacy clerk. The establishment shall ensure that the person making the sale requires the buyer:

(1) to provide photographic identification showing the buyer's date of birth; and

(2) to sign a written or electronic document detailing the date of the sale, the name of the buyer, and the amount of the drug sold.

A document described under clause (2) must be retained by the establishment for at least three years and must at all reasonable times be open to the inspection of any law enforcement agency.

Nothing in this paragraph requires the buyer to obtain a prescription for the drug's purchase.

(f) No person may acquire through over-the-counter sales more than six grams of methamphetamine precursor drugs, calculated as the base, within a 30-day period.

(g) No person may sell in an over-the-counter sale a methamphetamine precursor drug to a person under the age of 18 years. It is an affirmative defense to a charge under this paragraph if the defendant proves by a preponderance of the evidence that the defendant reasonably and in good faith relied on proof of age as described in section 340A.503, subdivision 6.

(h) A person who knowingly violates paragraph (c), (d), (e), (f), or (g) is guilty of a misdemeanor and may be sentenced to imprisonment for not more than 90 days, or to payment of a fine of not more than $1,000, or both.

(i) An owner, operator, supervisor, or manager of a business establishment that offers for sale methamphetamine precursor drugs whose employee or agent is convicted of or charged with violating paragraph (c), (d), (e), (f), or (g) is not subject to the criminal penalties for violating any of those paragraphs if the person:

(1) did not have prior knowledge of, participate in, or direct the employee or agent to commit the violation; and

(2) documents that an employee training program was in place to provide the employee or agent with information on the state and federal laws and regulations regarding methamphetamine precursor drugs.

(j) Any person employed by a business establishment that offers for sale methamphetamine precursor drugs who sells such a drug to any person in a suspicious transaction shall report the transaction to the owner, supervisor, or manager of the establishment. The owner, supervisor, or manager may report the transaction to local law enforcement. A person who reports information under this subdivision in good faith is immune from civil liability relating to the report.

(k) Paragraphs (b) to (j) do not apply to:

(1) pediatric products labeled pursuant to federal regulation primarily intended for administration to children under 12 years of age according to label instructions;

(2) methamphetamine precursor drugs that are certified by the Board of Pharmacy as being manufactured in a manner that prevents the drug from being used to manufacture methamphetamine;

(3) methamphetamine precursor drugs in gel capsule or liquid form; or

(4) compounds, mixtures, or preparations in powder form where pseudoephedrine constitutes less than one percent of its total weight and is not its sole active ingredient.

(l) The Board of Pharmacy, in consultation with the Department of Public Safety, shall certify methamphetamine precursor drugs that meet the requirements of paragraph (k), clause (2), and publish an annual listing of these drugs.

(m) Wholesale drug distributors licensed and regulated by the Board of Pharmacy pursuant to sections 151.42 to 151.51 and registered with and regulated by the United States Drug Enforcement Administration are exempt from the methamphetamine precursor drug storage requirements of this section.

(n) This section preempts all local ordinances or regulations governing the sale by a business establishment of over-the-counter products containing ephedrine or pseudoephedrine. All ordinances enacted prior to the effective date of this act are void.

Subd. 7. **Board of Pharmacy; regulation of substances.** The Board of Pharmacy is authorized to regulate and define additional substances which contain quantities of a substance possessing abuse potential in accordance with the following criteria:

(1) The Board of Pharmacy shall place a substance in Schedule I if it finds that the substance has: A high potential for abuse, no currently accepted medical use in the United States, and a lack of accepted safety for use under medical supervision.

(2) The Board of Pharmacy shall place a substance in Schedule II if it finds that the substance has: A high potential for abuse, currently accepted medical use in the United States, or currently accepted medical use with severe restrictions, and that abuse may lead to severe psychological or physical dependence.

(3) The Board of Pharmacy shall place a substance in Schedule III if it finds that the substance has: A potential for abuse less than the substances listed in Schedules I and II, currently accepted medical use in treatment in the United States, and that abuse may lead to moderate or low physical dependence or high psychological dependence.

(4) The Board of Pharmacy shall place a substance in Schedule IV if it finds that the substance has: A low potential for abuse relative to the substances in Schedule III, currently accepted medical use in treatment in the United States, and that abuse may lead to limited physical dependence or psychological dependence relative to the substances in Schedule III.

(5) The Board of Pharmacy shall place a substance in Schedule V if it finds that the substance has: A low potential for abuse relative to the substances listed in Schedule IV, currently accepted medical use in treatment in the United States, and limited physical dependence and/or psychological dependence liability relative to the substances listed in Schedule IV.

Subd. 8. **Add, delete, or reschedule substances.** The state Board of Pharmacy may, by rule, add substances to or delete or reschedule substances listed in this section. The Board of Pharmacy may not delete or reschedule a drug that is in Schedule I, except as provided in subdivision 12.

In making a determination regarding a substance, the Board of Pharmacy shall consider the following: The actual or relative potential for abuse, the scientific evidence of its pharmacological effect, if known, the state of current scientific knowledge regarding the substance, the history and current pattern of abuse, the scope, duration, and significance of abuse, the risk to public health, the potential of the substance to produce psychic or physiological dependence liability, and whether the substance is an immediate precursor of a substance already controlled under this section. The state Board of Pharmacy may include any nonnarcotic drug authorized by federal law for medicinal use in a schedule only if such drug must, under either federal or state law or rule, be sold only on prescription.

Subd. 8a. [Repealed by amendment, 2012 c 240 s 1]

Subd. 8b. **Board of Pharmacy; expedited scheduling of additional substances.** (a) The state Board of Pharmacy may, by rule, add a substance to Schedule I provided that it finds that the substance has a high potential for abuse, has no currently accepted medical use in the United States, has a lack of accepted safety for use under medical supervision, has known adverse health effects, and is currently available for use within the state. For the purposes of this subdivision only, the board may use the expedited rulemaking process under section 14.389. The scheduling of a substance under this subdivision expires the day after the adjournment of the legislative session immediately following the substance's scheduling unless the legislature by law ratifies the action.

(b) If the board schedules a substance under this subdivision, the board shall notify in a timely manner the chairs and ranking minority members of the senate and house of representatives committees having jurisdiction over criminal justice and health policy and finance of the action

and the reasons for it. The notice must include a copy of the administrative law judge's decision on the matter.

(c) This subdivision expires August 1, 2014.

Subd. 9. **Except substances by rule.** The state Board of Pharmacy may by rule except any compound, mixture, or preparation containing any stimulant or depressant substance listed in subdivision 4, paragraphs (b) and (c), or in subdivisions 5 and 6 from the application of all or any part of this chapter, if the compound, mixture, or preparation contains one or more active medicinal ingredients not having a stimulant or depressant effect on the central nervous system; provided, that such admixtures shall be included therein in such combinations, quantity, proportion, or concentration as to vitiate the potential for abuse of the substances which do have a stimulant or depressant effect on the central nervous system.

Subd. 10. **Dextromethorphan.** Dextromethorphan shall not be deemed to be included in any schedule by reason of the enactment of Laws 1971, chapter 937, unless controlled pursuant to the foregoing provisions of this section.

Subd. 11. [Repealed, 1993 c 337 s 20]

Subd. 12. **Coordination of controlled substance regulation with federal law and state statute.** If any substance is designated, rescheduled, or deleted as a controlled substance under federal law and notice thereof is given to the state Board of Pharmacy, the state Board of Pharmacy shall similarly control the substance under this chapter, after the expiration of 30 days from publication in the Federal Register of a final order designating a substance as a controlled substance or rescheduling or deleting a substance. Such order shall be filed with the secretary of state. If within that 30-day period, the state Board of Pharmacy objects to inclusion, rescheduling, or deletion, it shall publish the reasons for objection and afford all interested parties an opportunity to be heard. At the conclusion of the hearing, the state Board of Pharmacy shall publish its decision, which shall be subject to the provisions of chapter 14.

In exercising the authority granted by this chapter, the state Board of Pharmacy shall be subject to the provisions of chapter 14.

The state Board of Pharmacy shall annually submit a report to the legislature on or before December 1 that specifies what changes the board made to the controlled substance schedules maintained by the board in Minnesota Rules, parts 6800.4210 to 6800.4250, in the preceding 12 months. The report must include specific recommendations for amending the controlled substance schedules contained in subdivisions 2 to 6, so that they conform with the controlled substance schedules maintained by the board in Minnesota Rules, parts 6800.4210 to 6800.4250.

Subd. 13. [Repealed by amendment, 2012 c 240 s 1]

History: *1971 c 937 s 12; 1973 c 693 s 2-4; 1976 c 338 s 1-4; 1979 c 157 s 2-4; 1979 c 243 s 2; 1982 c 424 s 130; 1983 c 260 s 39,40; 1985 c 248 s 70; 1987 c 14 s 1; 1987 c 298 s 2; 1987 c 384 art 2 s 40; 1989 c 230 s 1; 1994 c 465 art 1 s 20-22; 1996 c 408 art 11 s 2; 1997 c 7 art 2 s 21; 1997 c 187 art 5 s 21; 1997 c 239 art 4 s 3,4,15; 1998 c 367 art 4 s 7; 1999 c 9 s 1; 1999 c 163 s 1; 2000 c 262 s 1; 2001 c 173 s 1; 1Sp2001 c 8 art 8 s 1; 2005 c 136 art 7 s 3,4; art 17 s 1,2; 1Sp2005 c 7 s 25; 2009 c 59 art 5 s 3,4; 2011 c 53 s 4,5; 2012 c 240 s 1*

CONTROLLED SUBSTANCE CRIMES

152.021 CONTROLLED SUBSTANCE CRIME IN THE FIRST DEGREE.

Subdivision 1. **Sale crimes.** A person is guilty of controlled substance crime in the first degree if:

(1) on one or more occasions within a 90-day period the person unlawfully sells one or more mixtures of a total weight of ten grams or more containing cocaine, heroin, or methamphetamine;

(2) on one or more occasions within a 90-day period the person unlawfully sells one or more mixtures of a total weight of 50 grams or more containing a narcotic drug other than cocaine, heroin, or methamphetamine;

(3) on one or more occasions within a 90-day period the person unlawfully sells one or more mixtures of a total weight of 50 grams or more containing amphetamine, phencyclidine, or hallucinogen or, if the controlled substance is packaged in dosage units, equaling 200 or more dosage units; or

(4) on one or more occasions within a 90-day period the person unlawfully sells one or more mixtures of a total weight of 50 kilograms or more containing marijuana or Tetrahydrocannabinols, or one or more mixtures of a total weight of 25 kilograms or more containing marijuana or Tetrahydrocannabinols in a school zone, a park zone, a public housing zone, or a drug treatment facility.

Subd. 2. **Possession crimes.** (a) A person is guilty of a controlled substance crime in the first degree if:

(1) the person unlawfully possesses one or more mixtures of a total weight of 25 grams or more containing cocaine, heroin, or methamphetamine;

(2) the person unlawfully possesses one or more mixtures of a total weight of 500 grams or more containing a narcotic drug other than cocaine, heroin, or methamphetamine;

(3) the person unlawfully possesses one or more mixtures of a total weight of 500 grams or more containing amphetamine, phencyclidine, or hallucinogen or, if the controlled substance is packaged in dosage units, equaling 500 or more dosage units; or

(4) the person unlawfully possesses one or more mixtures of a total weight of 100 kilograms or more containing marijuana or Tetrahydrocannabinols.

(b) For the purposes of this subdivision, the weight of fluid used in a water pipe may not be considered in measuring the weight of a mixture except in cases where the mixture contains four or more fluid ounces of fluid.

Subd. 2a. **Methamphetamine manufacture crime.** (a) Notwithstanding subdivision 1, sections 152.022, subdivision 1, 152.023, subdivision 1, and 152.024, subdivision 1, a person is guilty of controlled substance crime in the first degree if the person manufactures any amount of methamphetamine.

(b) [Renumbered 152.0262, subdivision 1]

Subd. 3. **Penalty.** (a) A person convicted under subdivisions 1 to 2a, paragraph (a), may be sentenced to imprisonment for not more than 30 years or to payment of a fine of not more than $1,000,000, or both.

(b) If the conviction is a subsequent controlled substance conviction, a person convicted under subdivisions 1 to 2a, paragraph (a), shall be committed to the commissioner of corrections

for not less than four years nor more than 40 years and, in addition, may be sentenced to payment of a fine of not more than $1,000,000.

(c) In a prosecution under subdivision 1 involving sales by the same person in two or more counties within a 90-day period, the person may be prosecuted for all of the sales in any county in which one of the sales occurred.

History: *1989 c 290 art 3 s 8; 1990 c 602 art 7 s 1; 1991 c 279 s 3; 1992 c 359 s 4,5; 1993 c 326 art 13 s 5; 1995 c 244 s 1; 1997 c 239 art 4 s 5,6; 1998 c 367 art 4 s 1; 1Sp2003 c 2 art 8 s 2,3; 2005 c 136 art 7 s 5,6,21; 2011 c 53 s 6*

152.022 CONTROLLED SUBSTANCE CRIME IN THE SECOND DEGREE.

Subdivision 1. **Sale crimes.** A person is guilty of controlled substance crime in the second degree if:

(1) on one or more occasions within a 90-day period the person unlawfully sells one or more mixtures of a total weight of three grams or more containing cocaine, heroin, or methamphetamine;

(2) on one or more occasions within a 90-day period the person unlawfully sells one or more mixtures of a total weight of ten grams or more containing a narcotic drug other than cocaine, heroin, or methamphetamine;

(3) on one or more occasions within a 90-day period the person unlawfully sells one or more mixtures of a total weight of ten grams or more containing amphetamine, phencyclidine, or hallucinogen or, if the controlled substance is packaged in dosage units, equaling 50 or more dosage units;

(4) on one or more occasions within a 90-day period the person unlawfully sells one or more mixtures of a total weight of 25 kilograms or more containing marijuana or Tetrahydrocannabinols;

(5) the person unlawfully sells any amount of a Schedule I or II narcotic drug to a person under the age of 18, or conspires with or employs a person under the age of 18 to unlawfully sell the substance; or

(6) the person unlawfully sells any of the following in a school zone, a park zone, a public housing zone, or a drug treatment facility:

(i) any amount of a Schedule I or II narcotic drug, lysergic acid diethylamide (LSD), 3,4-methylenedioxy amphetamine, or 3,4-methylenedioxymethamphetamine;

(ii) one or more mixtures containing methamphetamine or amphetamine; or

(iii) one or more mixtures of a total weight of five kilograms or more containing marijuana or Tetrahydrocannabinols.

Subd. 2. **Possession crimes.** (a) A person is guilty of controlled substance crime in the second degree if:

(1) the person unlawfully possesses one or more mixtures of a total weight of six grams or more containing cocaine, heroin, or methamphetamine;

(2) the person unlawfully possesses one or more mixtures of a total weight of 50 grams or more containing a narcotic drug other than cocaine, heroin, or methamphetamine;

(3) the person unlawfully possesses one or more mixtures of a total weight of 50 grams or more containing amphetamine, phencyclidine, or hallucinogen or, if the controlled substance is packaged in dosage units, equaling 100 or more dosage units; or

(4) the person unlawfully possesses one or more mixtures of a total weight of 50 kilograms or more containing marijuana or Tetrahydrocannabinols.

(b) For the purposes of this subdivision, the weight of fluid used in a water pipe may not be considered in measuring the weight of a mixture except in cases where the mixture contains four or more fluid ounces of fluid.

Subd. 3. **Penalty.** (a) A person convicted under subdivision 1 or 2 may be sentenced to imprisonment for not more than 25 years or to payment of a fine of not more than $500,000, or both.

(b) If the conviction is a subsequent controlled substance conviction, a person convicted under subdivision 1 or 2 shall be committed to the commissioner of corrections for not less than three years nor more than 40 years and, in addition, may be sentenced to payment of a fine of not more than $500,000.

(c) In a prosecution under subdivision 1 involving sales by the same person in two or more counties within a 90-day period, the person may be prosecuted for all of the sales in any county in which one of the sales occurred.

History: *1989 c 290 art 3 s 9; 1990 c 602 art 7 s 2; 1991 c 199 art 1 s 53; 1991 c 279 s 4; 1992 c 359 s 6,7; 1993 c 326 art 3 s 1; art 13 s 6; 1995 c 244 s 2; 1997 c 239 art 4 s 7,8; 1998 c 367 art 4 s 2; 1Sp2001 c 8 art 8 s 2; 2011 c 53 s 7*

152.023 CONTROLLED SUBSTANCE CRIME IN THE THIRD DEGREE.

Subdivision 1. **Sale crimes.** A person is guilty of controlled substance crime in the third degree if:

(1) the person unlawfully sells one or more mixtures containing a narcotic drug;

(2) on one or more occasions within a 90-day period the person unlawfully sells one or more mixtures containing phencyclidine or hallucinogen, it is packaged in dosage units, and equals ten or more dosage units;

(3) the person unlawfully sells one or more mixtures containing a controlled substance classified in Schedule I, II, or III, except a Schedule I or II narcotic drug, to a person under the age of 18;

(4) the person conspires with or employs a person under the age of 18 to unlawfully sell one or more mixtures containing a controlled substance listed in Schedule I, II, or III, except a Schedule I or II narcotic drug; or

(5) on one or more occasions within a 90-day period the person unlawfully sells one or more mixtures of a total weight of five kilograms or more containing marijuana or Tetrahydrocannabinols.

Subd. 2. **Possession crimes.** (a) A person is guilty of controlled substance crime in the third degree if:

(1) on one or more occasions within a 90-day period the person unlawfully possesses one or more mixtures of a total weight of three grams or more containing cocaine, heroin, or methamphetamine;

(2) on one or more occasions within a 90-day period the person unlawfully possesses one or more mixtures of a total weight of ten grams or more containing a narcotic drug other than cocaine, heroin, or methamphetamine;

(3) on one or more occasions within a 90-day period the person unlawfully possesses one or more mixtures containing a narcotic drug, it is packaged in dosage units, and equals 50 or more dosage units;

(4) on one or more occasions within a 90-day period the person unlawfully possesses any amount of a schedule I or II narcotic drug or five or more dosage units of lysergic acid diethylamide (LSD), 3,4-methylenedioxy amphetamine, or 3,4-methylenedioxymethamphetamine in a school zone, a park zone, a public housing zone, or a drug treatment facility;

(5) on one or more occasions within a 90-day period the person unlawfully possesses one or more mixtures of a total weight of ten kilograms or more containing marijuana or Tetrahydrocannabinols; or

(6) the person unlawfully possesses one or more mixtures containing methamphetamine or amphetamine in a school zone, a park zone, a public housing zone, or a drug treatment facility.

(b) For the purposes of this subdivision, the weight of fluid used in a water pipe may not be considered in measuring the weight of a mixture except in cases where the mixture contains four or more fluid ounces of fluid.

Subd. 3. **Penalty.** (a) A person convicted under subdivision 1 or 2 may be sentenced to imprisonment for not more than 20 years or to payment of a fine of not more than $250,000, or both.

(b) If the conviction is a subsequent controlled substance conviction, a person convicted under subdivision 1 or 2 shall be committed to the commissioner of corrections for not less than two years nor more than 30 years and, in addition, may be sentenced to payment of a fine of not more than $250,000.

(c) In a prosecution under subdivision 1 or 2 involving sales or acts of possession by the same person in two or more counties within a 90-day period, the person may be prosecuted in any county in which one of the sales or acts of possession occurred.

History: *1989 c 290 art 3 s 10; 1990 c 602 art 7 s 3,4; 1991 c 199 art 1 s 54; 1991 c 279 s 5; 1992 c 359 s 8; 1993 c 326 art 3 s 2; art 13 s 7; 1995 c 244 s 3; 1997 c 239 art 4 s 9-11; 1998 c 367 art 4 s 3; 1Sp2001 c 8 art 8 s 3; 2011 c 53 s 8*

152.024 CONTROLLED SUBSTANCE CRIME IN THE FOURTH DEGREE.

Subdivision 1. **Sale crimes.** A person is guilty of controlled substance crime in the fourth degree if:

(1) the person unlawfully sells one or more mixtures containing a controlled substance classified in Schedule I, II, or III, except marijuana or Tetrahydrocannabinols;

(2) the person unlawfully sells one or more mixtures containing a controlled substance classified in Schedule IV or V to a person under the age of 18;

(3) the person conspires with or employs a person under the age of 18 to unlawfully sell a controlled substance classified in Schedule IV or V; or

(4) the person unlawfully sells any amount of marijuana or Tetrahydrocannabinols in a school zone, a park zone, a public housing zone, or a drug treatment facility, except a small amount for no remuneration.

Subd. 2. **Possession crimes.** A person is guilty of controlled substance crime in the fourth degree if:

(1) the person unlawfully possesses one or more mixtures containing phencyclidine or hallucinogen, it is packaged in dosage units, and equals ten or more dosage units; or

(2) the person unlawfully possesses one or more mixtures containing a controlled substance classified in Schedule I, II, or III, except marijuana or Tetrahydrocannabinols, with the intent to sell it.

Subd. 3. **Penalty.** (a) A person convicted under subdivision 1 or 2 may be sentenced to imprisonment for not more than 15 years or to payment of a fine of not more than $100,000, or both.

(b) If the conviction is a subsequent controlled substance conviction, a person convicted under subdivision 1 or 2 shall be committed to the commissioner of corrections or to a local correctional authority for not less than one year nor more than 30 years and, in addition, may be sentenced to payment of a fine of not more than $100,000.

History: *1989 c 290 art 3 s 11; 1990 c 602 art 7 s 5; 1991 c 279 s 6; 1993 c 326 art 13 s 8; 1995 c 244 s 4; 1997 c 239 art 4 s 12*

152.025 CONTROLLED SUBSTANCE CRIME IN THE FIFTH DEGREE.

Subdivision 1. **Sale crimes.** (a) A person is guilty of a controlled substance crime in the fifth degree and if convicted may be sentenced to imprisonment for not more than five years or to payment of a fine of not more than $10,000, or both if:

(1) the person unlawfully sells one or more mixtures containing marijuana or tetrahydrocannabinols, except a small amount of marijuana for no remuneration; or

(2) the person unlawfully sells one or more mixtures containing a controlled substance classified in Schedule IV.

(b) Except as provided in paragraph (c), if a person is guilty of a controlled substance crime in the fifth degree and the conviction is a subsequent controlled substance conviction, the person convicted shall be committed to the commissioner of corrections or to a local correctional authority for not less than six months nor more than ten years and, in addition, may be sentenced to payment of a fine of not more than $20,000 if:

(1) the person unlawfully sells one or more mixtures containing marijuana or tetrahydrocannabinols, except a small amount of marijuana for no remuneration; or

(2) the person unlawfully sells one or more mixtures containing a controlled substance classified in Schedule IV.

(c) Prior to the time of sentencing, the prosecutor may file a motion to have the person sentenced without regard to the mandatory minimum sentence established by paragraph (b). The motion must be accompanied by a statement on the record of the reasons for it. When presented with the motion, or on its own motion, the court may sentence the person without regard to the mandatory minimum sentence if the court finds, on the record, substantial and compelling reasons to do so.

Subd. 2. **Possession and other crimes.** (a) A person is guilty of controlled substance crime in the fifth degree and if convicted may be sentenced to imprisonment for not more than five years or to payment of a fine of not more than $10,000, or both if:

(1) the person unlawfully possesses one or more mixtures containing a controlled substance classified in Schedule I, II, III, or IV, except a small amount of marijuana; or

(2) the person procures, attempts to procure, possesses, or has control over a controlled substance by any of the following means:

(i) fraud, deceit, misrepresentation, or subterfuge;

(ii) using a false name or giving false credit; or

(iii) falsely assuming the title of, or falsely representing any person to be, a manufacturer, wholesaler, pharmacist, physician, doctor of osteopathy licensed to practice medicine, dentist, podiatrist, veterinarian, or other authorized person for the purpose of obtaining a controlled substance.

(b) Except as provided in paragraph (c), if a person is guilty of a controlled substance crime in the fifth degree and the conviction is a subsequent controlled substance conviction, the person convicted shall be committed to the commissioner of corrections or to a local correctional authority for not less than six months nor more than ten years and, in addition, may be sentenced to payment of a fine of not more than $20,000 if:

(1) the person unlawfully possesses one or more mixtures containing a controlled substance classified in Schedule I, II, III, or IV, except a small amount of marijuana; or

(2) the person procures, attempts to procure, possesses, or has control over a controlled substance by any of the following means:

(i) fraud, deceit, misrepresentation, or subterfuge;

(ii) using a false name or giving false credit; or

(iii) falsely assuming the title of, or falsely representing any person to be, a manufacturer, wholesaler, pharmacist, physician, doctor of osteopathy licensed to practice medicine, dentist, podiatrist, veterinarian, or other authorized person for the purpose of obtaining a controlled substance.

(c) Prior to the time of sentencing, the prosecutor may file a motion to have the person sentenced without regard to the mandatory minimum sentence established by paragraph (b). The motion must be accompanied by a statement on the record of the reasons for it. When presented with the motion, or on its own motion, the court may sentence the person without regard to the mandatory minimum sentence if the court finds, on the record, substantial and compelling reasons to do so.

Subd. 3. [Repealed, 2009 c 83 art 3 s 24]

History: *1989 c 290 art 3 s 12; 1990 c 602 art 7 s 6; 1992 c 359 s 9; 1993 c 326 art 13 s 9; 1995 c 244 s 5; 2009 c 83 art 3 s 3,4; 2010 c 382 s 35*

152.026 MANDATORY SENTENCES.

A defendant convicted and sentenced to a mandatory sentence under sections 152.021 to 152.025 and 152.0262 is not eligible for probation, parole, discharge, or supervised release until that person has served the full term of imprisonment as provided by law, notwithstanding sections 242.19, 243.05, 609.12, and 609.135. "Term of imprisonment" has the meaning given in section 244.01, subdivision 8.

History: *1989 c 290 art 3 s 13; 1993 c 326 art 13 s 10; 2005 c 136 art 7 s 21*

152.0261 IMPORTING CONTROLLED SUBSTANCES ACROSS STATE BORDERS.

Subdivision 1. **Felony.** A person who crosses a state or international border into Minnesota while in possession of an amount of a controlled substance that constitutes a first-degree controlled substance crime under section 152.021, subdivision 2, is guilty of importing controlled substances and may be sentenced as provided in subdivision 3.

Subd. 1a. **Use of person under 18 to import.** A person who conspires with or employs a person under the age of 18 to cross a state or international border into Minnesota while that person or the person under the age of 18 is in possession of an amount of a controlled substance that constitutes a controlled substance crime under sections 152.021 to 152.025 and 152.0262, with the intent to obstruct the criminal justice process, is guilty of importing controlled substances and may be sentenced as provided in subdivision 3.

Subd. 2. **Jurisdiction.** A violation of this section may be charged, indicted, and tried in any county, but not more than one county, into or through which the actor has brought the controlled substance.

Subd. 3. **Penalty.** A person convicted of violating this section is guilty of a felony and may be sentenced to imprisonment for not more than 35 years or to payment of a fine of not more than $1,250,000, or both.

History: *1990 c 602 art 7 s 7; 1998 c 367 art 4 s 4,5; 2005 c 136 art 7 s 21*

152.0262 POSSESSION OF SUBSTANCES WITH INTENT TO MANUFACTURE METHAMPHETAMINE CRIME.

Subdivision 1. **Possession of precursors.** (a) A person is guilty of a crime if the person possesses any chemical reagents or precursors with the intent to manufacture methamphetamine and if convicted may be sentenced to imprisonment for not more than ten years or to payment of a fine of not more than $20,000, or both.

(b) A person is guilty of a crime if the person possesses any chemical reagents or precursors with the intent to manufacture methamphetamine and may be sentenced to imprisonment for not more than 15 years or to payment of a fine of not more than $30,000, or both, if the conviction is for a subsequent controlled substance conviction.

As used in this section and section 152.021, "chemical reagents or precursors" includes any of the following substances, or any similar substances that can be used to manufacture methamphetamine, or the salts, isomers, and salts of isomers of a listed or similar substance:

(1) ephedrine;

(2) pseudoephedrine;

(3) phenyl-2-propanone;

(4) phenylacetone;

(5) anhydrous ammonia;

(6) organic solvents;

(7) hydrochloric acid;

(8) lithium metal;

(9) sodium metal;

(10) ether;

(11) sulfuric acid;

(12) red phosphorus;

(13) iodine;

(14) sodium hydroxide;

(15) benzaldehyde;

(16) benzyl methyl ketone;

(17) benzyl cyanide;

(18) nitroethane;

(19) methylamine;

(20) phenylacetic acid;

(21) hydriodic acid; or

(22) hydriotic acid.

Subd. 2. [Repealed, 2009 c 83 art 2 s 50]

History: *1989 c 290 art 3 s 8; 1990 c 602 art 7 s 1; 1991 c 279 s 3; 1992 c 359 s 4,5; 1993 c 326 art 13 s 5; 1995 c 244 s 1; 1997 c 239 art 4 s 5,6; 1998 c 367 art 4 s 1; 1Sp2003 c 2 art 8 s 2,3; 2005 c 136 art 7 s 5,6,21; 2009 c 83 art 2 s 8*

152.027 OTHER CONTROLLED SUBSTANCE OFFENSES.

Subdivision 1. **Sale of Schedule V controlled substance.** Except as provided in section 152.02, subdivision 6, a person who unlawfully sells one or more mixtures containing a controlled substance classified in Schedule V may be sentenced to imprisonment for not more than one year or to payment of a fine of not more than $3,000, or both.

Subd. 2. **Possession of Schedule V controlled substance.** Except as provided in section 152.02, subdivision 6, a person who unlawfully possesses one or more mixtures containing a controlled substance classified in Schedule V may be sentenced to imprisonment for not more than one year or to payment of a fine of not more than $3,000, or both. The court may order that a person who is convicted under this subdivision and placed on probation be required to take part in a drug education program as specified by the court.

Subd. 3. **Possession of marijuana in a motor vehicle.** A person is guilty of a misdemeanor if the person is the owner of a private motor vehicle, or is the driver of the motor vehicle if the owner is not present, and possesses on the person, or knowingly keeps or allows to be kept within the area of the vehicle normally occupied by the driver or passengers, more than 1.4 grams of marijuana. This area of the vehicle does not include the trunk of the motor vehicle if the vehicle is equipped with a trunk, or another area of the vehicle not normally occupied by the driver or passengers if the vehicle is not equipped with a trunk. A utility or glove compartment is deemed to be within the area occupied by the driver and passengers.

Subd. 4. **Possession or sale of small amounts of marijuana.** (a) A person who unlawfully sells a small amount of marijuana for no remuneration, or who unlawfully possesses a small amount of marijuana is guilty of a petty misdemeanor and shall be required to participate in a drug education program unless the court enters a written finding that a drug education program is inappropriate. The program must be approved by an area mental health board with a curriculum approved by the state alcohol and drug abuse authority.

(b) A person convicted of an unlawful sale under paragraph (a) who is subsequently convicted of an unlawful sale under paragraph (a) within two years is guilty of a misdemeanor and shall be required to participate in a chemical dependency evaluation and treatment if so indicated by the evaluation.

(c) A person who is convicted of a petty misdemeanor under paragraph (a) who willfully and intentionally fails to comply with the sentence imposed, is guilty of a misdemeanor. Compliance with the terms of the sentence imposed before conviction under this paragraph is an absolute defense.

Subd. 5. **Sale or possession of salvia divinorum.** (a) A person who unlawfully sells any amount of salvia divinorum or salvinorin A is guilty of a gross misdemeanor.

(b) A person who unlawfully possesses any amount of salvia divinorum or salvinorin A is guilty of a misdemeanor.

Subd. 6. **Sale or possession of synthetic cannabinoids.** (a) As used in this subdivision, "synthetic cannabinoid" includes any substance included in section 152.02, subdivision 2, paragraph (h), clause (3).

(b) A person who unlawfully sells a synthetic cannabinoid for no remuneration is guilty of a gross misdemeanor.

(c) A person who unlawfully sells a synthetic cannabinoid is guilty of a felony and if convicted may be sentenced to imprisonment for not more than five years or to payment of a fine of not more than $10,000, or both.

(d) A person who unlawfully possesses any amount of a synthetic cannabinoid is guilty of a misdemeanor.

(e) Notwithstanding any contrary provision in sections 152.021 to 152.025, this subdivision describes the exclusive penalties for the sale and possession of synthetic cannabinoid.

History: *1989 c 290 art 3 s 14; 2005 c 10 art 3 s 8; 2005 c 136 art 7 s 7,8; 2010 c 368 s 1; 2011 c 53 s 9; 2011 c 76 art 1 s 20; 2012 c 240 s 2*

LOCATION OF USE, SALE, MANUFACTURE

152.0271 NOTICE OF DRUG CONVICTIONS; DRIVER'S LICENSE REVOCATION.

When a person is convicted of violating a provision of sections 152.021 to 152.027 and 152.0262, the sentencing court shall determine whether the person unlawfully sold or possessed the controlled substance while driving a motor vehicle. If so, the court shall notify the commissioner of public safety of its determination and order the commissioner to revoke the person's driver's license for 30 days. If the person does not have a driver's license or if the person's driver's license is suspended or revoked at the time of the conviction, the commissioner shall delay the issuance or reinstatement of the person's driver's license for 30 days after the person applies for the issuance or reinstatement of the license. Upon receipt of the court's order, the commissioner is authorized to take the licensing action without a hearing.

History: *1993 c 347 s 1; 2005 c 136 art 7 s 21*

152.0275 CERTAIN CONTROLLED SUBSTANCE OFFENSES; RESTITUTION; PROHIBITIONS ON PROPERTY USE; NOTICE PROVISIONS.

Subdivision 1. **Restitution.** (a) As used in this subdivision:

(1) "clandestine lab site" means any structure or conveyance or outdoor location occupied or affected by conditions or chemicals typically associated with the manufacturing of methamphetamine;

(2) "emergency response" includes, but is not limited to, removing and collecting evidence, securing the site, removal, remediation, and hazardous chemical assessment or inspection of the site where the relevant offense or offenses took place, regardless of whether these actions are performed by the public entities themselves or by private contractors paid by the public entities, or the property owner;

(3) "remediation" means proper cleanup, treatment, or containment of hazardous substances or methamphetamine at or in a clandestine lab site, and may include demolition or disposal of structures or other property when an assessment so indicates; and

(4) "removal" means the removal from the clandestine lab site of precursor or waste chemicals, chemical containers, or equipment associated with the manufacture, packaging, or storage of illegal drugs.

(b) A court may require a person convicted of manufacturing or attempting to manufacture a controlled substance or of an illegal activity involving a precursor substance, where the response to the crime involved an emergency response, to pay restitution to all public entities that participated in the response. The restitution ordered may cover the reasonable costs of their participation in the response.

(c) In addition to the restitution authorized in paragraph (b), a court may require a person convicted of manufacturing or attempting to manufacture a controlled substance or of illegal activity involving a precursor substance to pay restitution to a property owner who incurred removal or remediation costs because of the crime.

Subd. 2. **Property-related prohibitions; notice; Web site.** (a) As used in this subdivision:

(1) "clandestine lab site" has the meaning given in subdivision 1, paragraph (a);

(2) "property" means publicly or privately owned real property including buildings and other structures, motor vehicles as defined in section 609.487, subdivision 2a, public waters, and public rights-of-way;

(3) "remediation" has the meaning given in subdivision 1, paragraph (a); and

(4) "removal" has the meaning given in subdivision 1, paragraph (a).

(b) A peace officer who arrests a person at a clandestine lab site shall notify the appropriate county or local health department, state duty officer, and child protection services of the arrest and the location of the site.

(c) A county or local health department or sheriff shall order that any property or portion of a property that has been found to be a clandestine lab site and contaminated by substances, chemicals, or items of any kind used in the manufacture of methamphetamine or any part of the manufacturing process, or the by-products or degradates of manufacturing methamphetamine be prohibited from being occupied or used until it has been assessed and remediated as provided in the Department of Health's clandestine drug labs general cleanup guidelines. The remediation shall be accomplished by a contractor who will make the verification required under paragraph (e).

(d) Unless clearly inapplicable, the procedures specified in chapter 145A and any related rules adopted under that chapter addressing the enforcement of public health laws, the removal

and abatement of public health nuisances, and the remedies available to property owners or occupants apply to this subdivision.

(e) Upon the proper removal and remediation of any property used as a clandestine lab site, the contractor shall verify to the property owner and the applicable authority that issued the order under paragraph (c) that the work was completed according to the Department of Health's clandestine drug labs general cleanup guidelines and best practices. The contractor shall provide the verification to the property owner and the applicable authority within five days from the completion of the remediation. Following this, the applicable authority shall vacate its order.

(f) If a contractor issues a verification and the property was not remediated according to the Department of Health's clandestine drug labs general cleanup guidelines, the contractor is liable to the property owner for the additional costs relating to the proper remediation of the property according to the guidelines and for reasonable attorney fees for collection of costs by the property owner. An action under this paragraph must be commenced within six years from the date on which the verification was issued by the contractor.

(g) If the applicable authority determines under paragraph (c) that a motor vehicle has been contaminated by substances, chemicals, or items of any kind used in the manufacture of methamphetamine or any part of the manufacturing process, or the by-products or degradates of manufacturing methamphetamine and if the authority is able to obtain the certificate of title for the motor vehicle, the authority shall notify the registrar of motor vehicles of this fact and in addition, forward the certificate of title to the registrar. The authority shall also notify the registrar when it vacates its order under paragraph (e).

(h) The applicable authority issuing an order under paragraph (c) shall record with the county recorder or registrar of titles of the county where the clandestine lab is located an affidavit containing the name of the owner, a legal description of the property where the clandestine lab was located, and a map drawn from available information showing the boundary of the property and the location of the contaminated area on the property that is prohibited from being occupied or used that discloses to any potential transferee:

(1) that the property, or portion of the property, was the site of a clandestine lab;

(2) the location, condition, and circumstances of the clandestine lab, to the full extent known or reasonably ascertainable; and

(3) that the use of the property or some portion of it may be restricted as provided by paragraph (c).

If an inaccurate drawing or description is filed, the authority, on request of the owner or another interested person, shall file a supplemental affidavit with a corrected drawing or description.

If the authority vacates its order under paragraph (e), the authority shall record an affidavit that contains the recording information of the above affidavit and states that the order is vacated. Upon filing the affidavit vacating the order, the affidavit and the affidavit filed under this paragraph, together with the information set forth in the affidavits, cease to constitute either actual or constructive notice.

(i) If proper removal and remediation has occurred on the property, an interested party may record an affidavit indicating that this has occurred. Upon filing the affidavit described in this paragraph, the affidavit and the affidavit filed under paragraph (h), together with the information

set forth in the affidavits, cease to constitute either actual or constructive notice. Failure to record an affidavit under this section does not affect or prevent any transfer of ownership of the property.

(j) The county recorder or registrar of titles must record all affidavits presented under paragraph (h) or (i) in a manner that ensures their disclosure in the ordinary course of a title search of the subject property.

(k) The commissioner of health shall post on the Internet contact information for each local community health services administrator.

(l) Each local community health services administrator shall maintain information related to property within the administrator's jurisdiction that is currently or was previously subject to an order issued under paragraph (c). The information maintained must include the name of the owner, the location of the property, the extent of the contamination, the status of the removal and remediation work on the property, and whether the order has been vacated. The administrator shall make this information available to the public either upon request or by other means.

(m) Before signing an agreement to sell or transfer real property, the seller or transferor must disclose in writing to the buyer or transferee if, to the seller's or transferor's knowledge, methamphetamine production has occurred on the property. If methamphetamine production has occurred on the property, the disclosure shall include a statement to the buyer or transferee informing the buyer or transferee:

(1) whether an order has been issued on the property as described in paragraph (c);

(2) whether any orders issued against the property under paragraph (c) have been vacated under paragraph (j); or

(3) if there was no order issued against the property and the seller or transferor is aware that methamphetamine production has occurred on the property, the status of removal and remediation on the property.

(n) Unless the buyer or transferee and seller or transferor agree to the contrary in writing before the closing of the sale, a seller or transferor who fails to disclose, to the best of their knowledge, at the time of sale any of the facts required, and who knew or had reason to know of methamphetamine production on the property, is liable to the buyer or transferee for:

(1) costs relating to remediation of the property according to the Department of Health's clandestine drug labs general cleanup guidelines and best practices; and

(2) reasonable attorney fees for collection of costs from the seller or transferor.

An action under this paragraph must be commenced within six years after the date on which the buyer or transferee closed the purchase or transfer of the real property where the methamphetamine production occurred.

(o) This section preempts all local ordinances relating to the sale or transfer of real property designated as a clandestine lab site.

History: *2005 c 136 art 7 s 9*

152.028 PERMISSIVE INFERENCE OF KNOWING POSSESSION.

Subdivision 1. **Residences.** The presence of a controlled substance in open view in a room, other than a public place, under circumstances evincing an intent by one or more of the persons present to unlawfully mix, compound, package, or otherwise prepare for sale the controlled substance permits the fact finder to infer knowing possession of the controlled substance by each

person in close proximity to the controlled substance when the controlled substance was found. The permissive inference does not apply to any person if:

(1) one of them legally possesses the controlled substance; or

(2) the controlled substance is on the person of one of the occupants.

Subd. 2. **Passenger automobiles.** The presence of a controlled substance in a passenger automobile permits the fact finder to infer knowing possession of the controlled substance by the driver or person in control of the automobile when the controlled substance was in the automobile. This inference may only be made if the defendant is charged with violating section 152.021, 152.022, 152.023, 152.0261, or 152.0262. The inference does not apply:

(1) to a duly licensed operator of an automobile who is at the time operating it for hire in the lawful and proper pursuit of the operator's trade;

(2) to any person in the automobile if one of them legally possesses a controlled substance; or

(3) when the controlled substance is concealed on the person of one of the occupants.

History: *1989 c 290 art 3 s 15; 1990 c 602 art 7 s 8; 2005 c 136 art 7 s 21*

152.029 PUBLIC INFORMATION; SCHOOL ZONES, PARK ZONES, PUBLIC HOUSING ZONES, AND DRUG TREATMENT FACILITIES.

The attorney general shall disseminate information to the public relating to the penalties for committing controlled substance crimes in park zones, school zones, public housing zones, and drug treatment facilities. The attorney general shall draft a plain language version of sections 152.022 and 152.023 and relevant provisions of the Sentencing Guidelines, that describes in a clear and coherent manner using words with common and everyday meanings the content of those provisions. The attorney general shall publicize and disseminate the plain language version as widely as practicable, including distributing the version to school boards, local governments, and administrators and occupants of drug treatment facilities and public housing.

History: *1989 c 290 art 3 s 16; 1991 c 279 s 7; 1997 c 239 art 4 s 13*

152.03 [Repealed, 1969 c 933 s 22]

152.04 [Repealed, 1969 c 933 s 22]

152.041 [Repealed, 1971 c 937 s 22]

152.05 [Repealed, 1969 c 933 s 22]

152.06 [Repealed, 1969 c 933 s 22]

152.07 [Repealed, 1969 c 933 s 22]

152.08 [Repealed, 1969 c 933 s 22]

152.09 [Repealed, 1989 c 290 art 3 s 37]

DRUG PARAPHERNALIA

152.092 POSSESSION OF DRUG PARAPHERNALIA PROHIBITED.

It is unlawful for any person knowingly or intentionally to use or to possess drug paraphernalia. Any violation of this section is a petty misdemeanor.

History: *1982 c 557 s 2*

152.093 MANUFACTURE OR DELIVERY OF DRUG PARAPHERNALIA PROHIBITED.

It is unlawful for any person knowingly or intentionally to deliver drug paraphernalia or knowingly or intentionally to possess or manufacture drug paraphernalia for delivery. Any violation of this section is a misdemeanor.

History: *1982 c 557 s 3*

152.094 DELIVERY OF DRUG PARAPHERNALIA TO A MINOR PROHIBITED.

Any person 18 years of age or older who violates section 152.093 by knowingly or intentionally delivering drug paraphernalia to a person under 18 years of age who is at least three years younger is guilty of a gross misdemeanor.

History: *1982 c 557 s 4; 1986 c 444*

152.095 ADVERTISEMENT OF DRUG PARAPHERNALIA PROHIBITED.

It is unlawful for any person knowingly or intentionally to place in any newspaper, magazine, handbill, or other publication any advertisement or promotion for the sale of drug paraphernalia. A violation of this section is a misdemeanor.

History: *1982 c 557 s 5*

CONSPIRACIES

152.096 CONSPIRACIES PROHIBITED.

Subdivision 1. **Prohibited acts; penalties.** Any person who conspires to commit any act prohibited by this chapter, except possession or distribution for no remuneration of a small amount of marijuana as defined in section 152.01, subdivision 16, is guilty of a felony and upon conviction may be imprisoned, fined, or both, up to the maximum amount authorized by law for the act the person conspired to commit.

Subd. 2. **Conviction of coconspirator not required.** A person liable under this section may be charged with and convicted of conspiracy although the person or persons with whom that person conspired have not been convicted or have been convicted of some other crime based on the same act.

History: *1982 c 557 s 6; 1986 c 444; 1989 c 290 art 3 s 17*

SIMULATED CONTROLLED SUBSTANCES

152.097 SIMULATED CONTROLLED SUBSTANCES.

Subdivision 1. **Prohibition.** It is unlawful for any person knowingly to manufacture, sell, transfer or deliver or attempt to sell, transfer or deliver a noncontrolled substance upon:

(1) the express representation that the noncontrolled substance is a narcotic or nonnarcotic controlled substance; or

(2) the express representation that the substance is of such nature or appearance that the recipient of the delivery will be able to sell, transfer or deliver the substance as a controlled substance; or

(3) under circumstances which would lead a reasonable person to believe that the substance was a controlled substance. Any of the following factors shall constitute relevant evidence:

(i) the noncontrolled substance was packaged in a manner normally used for the illegal delivery of controlled substances; or

(ii) the delivery or attempted delivery included an exchange of or demand for money or other valuable property as consideration for delivery of the noncontrolled substance, and the amount of the consideration was substantially in excess of the reasonable value of the noncontrolled substance; or

(iii) the physical appearance of the noncontrolled substance is substantially identical to a specified controlled substance.

Subd. 2. **No defense.** In any prosecution under this section, it is no defense that the accused believed the noncontrolled substance to actually be a controlled substance.

Subd. 3. **Exemption.** This section does not apply to the prescribing and dispensing of placebos by licensed practitioners and licensed pharmacists.

Subd. 4. **Penalty.** A person who violates this section may be sentenced to imprisonment for not more than three years or to payment of a fine of not more than $20,000, or both. Sentencing for a conviction for attempting to sell, transfer, or deliver a noncontrolled substance in violation of this section is governed by section 609.17, subdivision 4.

History: *1982 c 599 s 1; 1989 c 290 art 3 s 18*

PRECURSORS OF CONTROLLED SUBSTANCES

152.0971 TERMS.

Subdivision 1. **Terms.** For purposes of sections 152.0971 to 152.0974, the following terms have the meanings given.

Subd. 1a. **Authorized agent.** An "authorized agent" is an individual representing a business who is responsible for the disbursement or custody of precursor substances.

Subd. 2. **Furnish.** "Furnish" means to sell, transfer, deliver, send, or supply a precursor substance by any other means.

Subd. 2a. **Purchaser.** A "purchaser" is a manufacturer, wholesaler, retailer, or any other person in this state who receives or seeks to receive a precursor substance.

Subd. 2b. **Receive.** "Receive" means to purchase, receive, collect, or otherwise obtain a precursor substance from a supplier.

Subd. 3. **Supplier.** A "supplier" is a manufacturer, wholesaler, retailer, or any other person in this or any other state who furnishes a precursor substance to another person in this state.

History: *1990 c 565 s 22; 1993 c 326 art 3 s 3-6*

152.0972 PRECURSORS OF CONTROLLED SUBSTANCES.

Subdivision 1. **Precursor substances.** The following precursors of controlled substances are "precursor substances":

(1) phenyl-2-propanone;

(2) methylamine;

(3) ethylamine;

(4) d-lysergic acid;

(5) ergotamine tartrate;

(6) diethyl malonate;

(7) malonic acid;

(8) hydriodic acid;

(9) ethyl malonate;

(10) barbituric acid;

(11) piperidine;

(12) n-acetylanthranilic acid;

(13) pyrrolidine;

(14) phenylacetic acid;

(15) anthranilic acid;

(16) ephedrine;

(17) pseudoephedrine;

(18) norpseudoephedrine;

(19) phenylpropanolamine;

(20) propionic anhydride;

(21) isosafrole;

(22) safrole;

(23) piperonal;

(24) thionylchloride;

(25) benzyl cyanide;

(26) ergonovine maleate;

(27) n-methylephedrine;

(28) n-ethylpseudoephedrine;

(29) n-methylpseudoephedrine;

(30) chloroephedrine;

(31) chloropseudoephedrine; and

(32) any substance added to this list by rule adopted by the state Board of Pharmacy.

Subd. 2. **Adoption of rules.** The state Board of Pharmacy may adopt rules under chapter 14 that add a substance to the list in subdivision 1, if the substance is a precursor to a controlled substance, or delete a substance from the list. A rule adding or deleting a substance is effective only until December 31 of the year following the calendar year during which the rule was adopted.

History: *1990 c 565 s 23; 1993 c 326 art 3 s 7*

152.0973 REPORT OF TRANSACTION.

Subdivision 1. **Predelivery notice.** A supplier who furnishes a precursor substance to a person in this state shall, not less than 21 days before delivery of the substance, submit to the Bureau of Criminal Apprehension a report of the transaction that includes the identification information specified in subdivision 3.

Subd. 1a. **Report of precursor substances received from out of state.** A purchaser of a precursor substance from outside of Minnesota shall, not less than 21 days before taking possession of the substance, submit to the Bureau of Criminal Apprehension a report of the transaction that includes the identification information specified in subdivision 3.

Subd. 2. **Regular reports.** The bureau may authorize a purchaser or supplier to submit the reports on a monthly basis with respect to repeated, regular transactions between the supplier and the purchaser involving the same substance if the superintendent of the Bureau of Criminal Apprehension determines that:

(1) a pattern of regular supply of the precursor substance exists between the supplier and the purchaser of the substance; or

(2) the purchaser has established a record of utilizing the precursor substance for lawful purposes.

Subd. 2a. **Report of missing precursor substance.** A supplier or purchaser who discovers a discrepancy between the quantity of precursor substance shipped and the quantity of precursor substance received shall report the discrepancy to the Bureau of Criminal Apprehension within three days of knowledge of the discrepancy. The report must include:

(1) the complete name and address of the purchaser;

(2) the type of precursor substance missing;

(3) whether the precursor substance is missing due to theft, loss, or shipping discrepancy;

(4) the method of delivery used;

(5) the name of the common carrier or person who transported the substance; and

(6) the date of shipment.

Subd. 3. **Proper identification.** A report submitted by a supplier or purchaser under this section must include:

(1) the purchaser's driver's license number or state identification number and residential or mailing address other than a post office box number taken from the purchaser's driver's license or state identification card, if the purchaser is not an authorized agent;

(2) the motor vehicle license number of the motor vehicle operated by the purchaser at the time of sale, if the purchaser is not an authorized agent;

(3) a complete description of how the precursor substance will be used, if the purchaser is not an authorized agent;

(4) a letter of authorization from the business for which the precursor substance is being furnished, including the state tax identification number and address of the business, a full description of how the precursor substance is to be used, and the signature of the authorized agent for the purchaser;

(5) the signature of the supplier as a witness to the signature and identification of the purchaser;

(6) the type and quantity of the precursor substance;

(7) the method of delivery used; and

(8) the complete name and address of the supplier.

Subd. 4. **Retention of records.** A supplier shall retain a copy of reports filed under subdivisions 1, 2, and 2a for five years. A purchaser shall retain a copy of reports filed under subdivisions 1a and 2a for five years.

Subd. 5. **Inspections.** All records relating to sections 152.0971 to 152.0974 shall be open to inspection by the Bureau of Criminal Apprehension during regular business hours.

Subd. 6. **Penalties.** (a) A person who does not submit a report as required by this section is guilty of a misdemeanor.

(b) A person who knowingly submits a report required by this section with false or fictitious information is guilty of a gross misdemeanor.

(c) A person who is convicted a second or subsequent time of violating paragraph (a) is guilty of a gross misdemeanor if the subsequent offense occurred after the earlier conviction.

History: *1990 c 565 s 24; 1993 c 326 art 3 s 8-14*

152.0974 EXCEPTIONS.

Sections 152.0971 to 152.0974 do not apply to:

(1) a pharmacist or other authorized person who sells or furnishes a precursor substance on the prescription of a physician, dentist, podiatrist, or veterinarian;

(2) a physician, dentist, podiatrist, or veterinarian who administers or furnishes a precursor substance to patients;

(3) a manufacturer or wholesaler licensed by the state Board of Pharmacy who sells, transfers, or otherwise furnishes a precursor substance to a licensed pharmacy, physician, dentist, podiatrist, or veterinarian; or

(4) the furnishing or receipt of a drug that contains ephedrine, pseudoephedrine, norpseudoephedrine, or phenylpropanolamine and is lawfully furnished over the counter without a prescription under the Federal Food, Drug, and Cosmetic Act, United States Code, title 21, chapter 9, or regulations adopted under that act.

History: *1990 c 565 s 25*

SALES AND RECORDS

152.10 SALES, PERSONS ELIGIBLE.

No person other than a licensed pharmacist, assistant pharmacist or pharmacist intern under the supervision of a pharmacist shall sell a stimulant or depressant drug and then only as provided in sections 152.021 to 152.12 and 152.0262.

History: *(3906-13) 1939 c 102 s 3; 1967 c 408 s 5; 1991 c 199 art 2 s 1; 2005 c 136 art 7 s 21*

152.101 MANUFACTURERS, RECORDS.

Subdivision 1. **Preparation of record.** Every person engaged in manufacturing, compounding, processing, selling, delivering or otherwise disposing of any controlled substance

shall, upon July 1, 1971, May 1, 1973, and every second year thereafter, prepare a complete and accurate record of all stocks of each controlled substance on hand and shall keep such record for two years. When additional controlled substances are designated after July 1, 1971, a similar record must be prepared upon the effective date of their designation. On and after July 1, 1971, every person manufacturing, compounding or processing any controlled substance shall prepare and keep, for not less than two years, a complete and accurate record of the kind and quantity of each drug manufactured, compounded or processed and the date of such manufacture, compounding, or processing; and every person selling, delivering, or otherwise disposing of any controlled substance shall prepare or obtain, and keep for not less than two years, a complete and accurate record of the kind and quantity of each such controlled substance received, sold, delivered, or otherwise disposed of, the name and address from whom it was received and to whom it was sold, delivered or otherwise disposed of, and the date of such transaction. The form of such records shall be prescribed by the state board of pharmacy.

Subd. 2. **Application to doctors.** This section shall not apply to a licensed doctor of medicine, a doctor of osteopathy duly licensed to practice medicine, a licensed doctor of dentistry, a licensed doctor of podiatry, or licensed doctor of veterinary medicine in the course of that doctor's professional practice, unless such practitioner regularly engages in dispensing any such drugs to the practitioner's patients for which the patients are charged, either separately or together with charges for other professional services.

Subd. 3. **Research exception.** This section shall not apply to a person engaged in bona fide research conducted under an exemption granted under applicable federal law.

History: *1967 c 408 s 6; 1971 c 937 s 14; 1973 c 693 s 6; 1986 c 444*

PRESCRIPTIONS

152.11 PRESCRIPTIONS.

Subdivision 1. **General prescription requirements for controlled substances.** (a) A written prescription or an oral prescription reduced to writing, when issued for a controlled substance in Schedule II, III, IV, or V, is void unless (1) it is written in ink and contains the name and address of the person for whose use it is intended; (2) it states the amount of the controlled substance to be compounded or dispensed, with directions for its use; (3) if a written prescription, it contains the handwritten signature, address, and federal registry number of the prescriber and a designation of the branch of the healing art pursued by the prescriber; and if an oral prescription, the name and address of the prescriber and a designation of the prescriber's branch of the healing art; and (4) it shows the date when signed by the prescriber, or the date of acceptance in the pharmacy if an oral prescription.

(b) An electronic prescription for a controlled substance in Schedule II, III, IV, or V is void unless it complies with the standards established pursuant to section 62J.497 and with those portions of Code of Federal Regulations, title 21, parts 1300, 1304, 1306, and 1311, that pertain to electronic prescriptions.

(c) A prescription for a controlled substance in Schedule II, III, IV, or V that is transmitted by facsimile, either computer to facsimile machine or facsimile machine to facsimile machine, is void unless it complies with the applicable requirements of Code of Federal Regulations, title 21, part 1306.

(d) Every licensed pharmacy that dispenses a controlled substance prescription shall retain the original prescription in a file for a period of not less than two years, open to inspection by any officer of the state, county, or municipal government whose duty it is to aid and assist with the enforcement of this chapter. An original electronic or facsimile prescription may be stored in an electronic database, provided that the database provides a means by which original prescriptions can be retrieved, as transmitted to the pharmacy, for a period of not less than two years.

(e) Every licensed pharmacy shall distinctly label the container in which a controlled substance is dispensed with the directions contained in the prescription for the use of that controlled substance.

Subd. 1a. **Prescription requirements for Schedule II controlled substances.** No person may dispense a controlled substance included in Schedule II of section 152.02 without a prescription issued by a doctor of medicine, a doctor of osteopathy licensed to practice medicine, a doctor of dental surgery, a doctor of dental medicine, a doctor of podiatry, or a doctor of veterinary medicine, lawfully licensed to prescribe in this state or by a practitioner licensed to prescribe controlled substances by the state in which the prescription is issued, and having a current federal Drug Enforcement Administration registration number. The prescription must either be printed or written in ink and contain the handwritten signature of the prescriber or be transmitted electronically or by facsimile as permitted under subdivision 1. Provided that in emergency situations, as authorized by federal law, such drug may be dispensed upon oral prescription reduced promptly to writing and filed by the pharmacist. Such prescriptions shall be retained in conformity with section 152.101. No prescription for a Schedule II substance may be refilled.

Subd. 2. **Prescription requirements for Schedule III or IV controlled substances.** No person may dispense a controlled substance included in Schedule III or IV of section 152.02 without a prescription issued, as permitted under subdivision 1, by a doctor of medicine, a doctor of osteopathy licensed to practice medicine, a doctor of dental surgery, a doctor of dental medicine, a doctor of podiatry, a doctor of optometry limited to Schedule IV, or a doctor of veterinary medicine, lawfully licensed to prescribe in this state or from a practitioner licensed to prescribe controlled substances by the state in which the prescription is issued, and having a current federal drug enforcement administration registration number. Such prescription may not be dispensed or refilled except with the documented consent of the prescriber, and in no event more than six months after the date on which such prescription was issued and no such prescription may be refilled more than five times.

Subd. 2a. **Federal registration number exemption.** A prescription need not bear a federal drug enforcement administration registration number that authorizes the prescriber to prescribe controlled substances if the drug prescribed is not a controlled substance in Schedule II, III, IV, or V. No person shall impose a requirement inconsistent with this subdivision.

Subd. 2b. **Restriction on release of federal registration number.** No person or entity may offer for sale, sell, lease, or otherwise release a federal drug enforcement administration registration number for any reason, except for drug enforcement purposes authorized by this chapter and the federal controlled substances registration system. For purposes of this section, an entity includes a state governmental agency or regulatory board, a health plan company as defined under section 62Q.01, subdivision 4, a managed care organization as defined under section 62Q.01, subdivision 5, or any other entity that maintains prescription data.

Subd. 2c. **Restriction on use of federal registration number.** No entity may use a federal drug enforcement administration registration number to identify or monitor the prescribing

practices of a prescriber to whom that number has been assigned, except for drug enforcement purposes authorized by this chapter and the federal controlled substances registration system. For purposes of this section, an entity includes a health plan company as defined under section 62Q.01, subdivision 4, a managed care organization as defined under section 62Q.01, subdivision 5, or any other entity that maintains prescription data.

Subd. 2d. **Identification requirement for Schedule II or III controlled substance prescriptions.** (a) No person may dispense a controlled substance included in Schedule II or III without requiring the person purchasing the controlled substance, who need not be the person for whom the controlled substance prescription is written, to present valid photographic identification, unless the person purchasing the controlled substance, or if applicable the person for whom the controlled substance prescription is written, is known to the dispenser.

(b) This subdivision applies only to purchases of controlled substances that are not covered, in whole or in part, by a health plan company or other third-party payor.

Subd. 3. **Dispensing orphan drugs.** For the purpose of this section, nothing shall prohibit the dispensing of orphan drugs prescribed by a person practicing in and licensed by another state as a physician, dentist, veterinarian, or podiatrist; who has a current federal drug enforcement administration registration number; and who may legally prescribe Schedule II, III, IV, or V controlled substances in that state.

History: *(3906-14) 1939 c 102 s 4; 1939 c 193 s 4; 1955 c 185 s 2; 1967 c 408 s 7; 1971 c 937 s 15; 1973 c 693 s 7; 1986 c 444; 1993 c 82 s 2; 1994 c 465 art 1 s 23; 1995 c 66 s 1,2; 1998 c 316 s 1-3; 2003 c 62 s 8; 2004 c 242 s 1,2; 2007 c 147 art 11 s 6; art 12 s 8; 2012 c 246 s 1*

152.12 DOCTORS MAY PRESCRIBE.

Subdivision 1. **Prescribing, dispensing, administering controlled substances in Schedules II through V.** A licensed doctor of medicine, a doctor of osteopathy, duly licensed to practice medicine, a doctor of dental surgery, a doctor of dental medicine, a licensed doctor of podiatry, or a licensed doctor of optometry limited to Schedules IV and V, and in the course of professional practice only, may prescribe, administer, and dispense a controlled substance included in Schedules II through V of section 152.02, may cause the same to be administered by a nurse, an intern or an assistant under the direction and supervision of the doctor, and may cause a person who is an appropriately certified and licensed health care professional to prescribe and administer the same within the expressed legal scope of the person's practice as defined in Minnesota Statutes.

Subd. 2. **Doctor of veterinary medicine.** A licensed doctor of veterinary medicine, in good faith, and in the course of professional practice only, and not for use by a human being, may prescribe, administer, and dispense a controlled substance included in Schedules II through V of section 152.02, and may cause the same to be administered by an assistant under the direction and supervision of the doctor.

Subd. 3. **Research project use of controlled substances.** Any qualified person may use controlled substances in the course of a bona fide research project but cannot administer or dispense such drugs to human beings unless such drugs are prescribed, dispensed and administered by a person lawfully authorized to do so. Every person who engages in research involving the use of such substances shall apply annually for registration by the state Board of Pharmacy and shall pay any applicable fee specified in section 151.065, provided that such

registration shall not be required if the person is covered by and has complied with federal laws covering such research projects.

Subd. 4. **Sale of controlled substances not prohibited for certain persons and entities.** Nothing in this chapter shall prohibit the sale to, or the possession of, a controlled substance in Schedule II, III, IV or V by: Registered drug wholesalers, registered manufacturers, registered pharmacies, or any licensed hospital or other licensed institutions wherein sick and injured persons are cared for or treated, or bona fide hospitals wherein animals are treated; or by licensed pharmacists, licensed doctors of medicine, doctors of osteopathy duly licensed to practice medicine, licensed doctors of dental surgery, licensed doctors of dental medicine, licensed doctors of podiatry, licensed doctors of optometry limited to Schedules IV and V, or licensed doctors of veterinary medicine when such practitioners use controlled substances within the course of their professional practice only.

Nothing in this chapter shall prohibit the possession of a controlled substance in Schedule II, III, IV or V by an employee or agent of a registered drug wholesaler, registered manufacturer, or registered pharmacy, while acting in the course of employment; by a patient of a licensed doctor of medicine, a doctor of osteopathy duly licensed to practice medicine, a licensed doctor of dental surgery, a licensed doctor of dental medicine, or a licensed doctor of optometry limited to Schedules IV and V; or by the owner of an animal for which a controlled substance has been prescribed by a licensed doctor of veterinary medicine, when such controlled substances are dispensed according to law.

Subd. 5. **Analytical laboratory not prohibited from providing anonymous analysis service.** Nothing in this chapter shall prohibit an analytical laboratory from conducting an anonymous analysis service when such laboratory is registered by the Federal Drug Enforcement Administration, nor prohibit the possession of a controlled substance by an employee or agent of such analytical laboratory while acting in the course of employment.

History: *(3906-15) 1939 c 102 s 5; 1967 c 408 s 8; 1971 c 937 s 16; 1973 c 693 s 8,9; 1974 c 369 s 2; 1986 c 444; 1988 c 440 s 3; 2003 c 62 s 9,10; 1Sp2011 c 9 art 5 s 27*

152.125 INTRACTABLE PAIN.

Subdivision 1. **Definition.** For purposes of this section, "intractable pain" means a pain state in which the cause of the pain cannot be removed or otherwise treated with the consent of the patient and in which, in the generally accepted course of medical practice, no relief or cure of the cause of the pain is possible, or none has been found after reasonable efforts. Reasonable efforts for relieving or curing the cause of the pain may be determined on the basis of, but are not limited to, the following:

(1) when treating a nonterminally ill patient for intractable pain, evaluation by the attending physician and one or more physicians specializing in pain medicine or the treatment of the area, system, or organ of the body perceived as the source of the pain; or

(2) when treating a terminally ill patient, evaluation by the attending physician who does so in accordance with the level of care, skill, and treatment that would be recognized by a reasonably prudent physician under similar conditions and circumstances.

Subd. 2. **Prescription and administration of controlled substances for intractable pain.** Notwithstanding any other provision of this chapter, a physician may prescribe or administer a controlled substance in Schedules II to V of section 152.02 to an individual in the course of the physician's treatment of the individual for a diagnosed condition causing intractable pain. No

physician shall be subject to disciplinary action by the Board of Medical Practice for appropriately prescribing or administering a controlled substance in Schedules II to V of section 152.02 in the course of treatment of an individual for intractable pain, provided the physician keeps accurate records of the purpose, use, prescription, and disposal of controlled substances, writes accurate prescriptions, and prescribes medications in conformance with chapter 147.

Subd. 3. **Limits on applicability.** This section does not apply to:

(1) a physician's treatment of an individual for chemical dependency resulting from the use of controlled substances in Schedules II to V of section 152.02;

(2) the prescription or administration of controlled substances in Schedules II to V of section 152.02 to an individual whom the physician knows to be using the controlled substances for nontherapeutic purposes;

(3) the prescription or administration of controlled substances in Schedules II to V of section 152.02 for the purpose of terminating the life of an individual having intractable pain; or

(4) the prescription or administration of a controlled substance in Schedules II to V of section 152.02 that is not a controlled substance approved by the United States Food and Drug Administration for pain relief.

Subd. 4. **Notice of risks.** Prior to treating an individual for intractable pain in accordance with subdivision 2, a physician shall discuss with the individual the risks associated with the controlled substances in Schedules II to V of section 152.02 to be prescribed or administered in the course of the physician's treatment of an individual, and document the discussion in the individual's record.

History: *1997 c 124 s 1*

152.126 CONTROLLED SUBSTANCES PRESCRIPTION ELECTRONIC REPORTING SYSTEM.

Subdivision 1. **Definitions.** For purposes of this section, the terms defined in this subdivision have the meanings given.

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

(b) "Controlled substances" means those substances listed in section 152.02, subdivisions 3 to 5, and those substances defined by the board pursuant to section 152.02, subdivisions 7, 8, and 12.

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

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

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

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

Subd. 1a. **Treatment of intractable pain.** This section is not intended to limit or interfere with the legitimate prescribing of controlled substances for pain. No prescriber shall be subject

to disciplinary action by a health-related licensing board for prescribing a controlled substance according to the provisions of section 152.125.

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

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

Subd. 3. **Prescription Electronic Reporting Advisory Committee.** (a) The board shall convene an advisory committee. The committee must include at least one representative of:

(1) the Department of Health;

(2) the Department of Human Services;

(3) each health-related licensing board that licenses prescribers;

(4) a professional medical association, which may include an association of pain management and chemical dependency specialists;

(5) a professional pharmacy association;

(6) a professional nursing association;

(7) a professional dental association;

(8) a consumer privacy or security advocate; and

(9) a consumer or patient rights organization.

(b) The advisory committee shall advise the board on the development and operation of the electronic reporting system, including, but not limited to:

(1) technical standards for electronic prescription drug reporting;

(2) proper analysis and interpretation of prescription monitoring data; and

(3) an evaluation process for the program.

Subd. 4. **Reporting requirements; notice.** (a) Each dispenser must submit the following data to the board or its designated vendor, subject to the notice required under paragraph (d):

(1) name of the prescriber;

(2) national provider identifier of the prescriber;

(3) name of the dispenser;

(4) national provider identifier of the dispenser;

(5) prescription number;

(6) name of the patient for whom the prescription was written;

(7) address of the patient for whom the prescription was written;

(8) date of birth of the patient for whom the prescription was written;

(9) date the prescription was written;

(10) date the prescription was filled;

(11) name and strength of the controlled substance;

(12) quantity of controlled substance prescribed;

(13) quantity of controlled substance dispensed; and

(14) number of days supply.

(b) The dispenser must submit the required information by a procedure and in a format established by the board. The board may allow dispensers to omit data listed in this subdivision or may require the submission of data not listed in this subdivision provided the omission or submission is necessary for the purpose of complying with the electronic reporting or data transmission standards of the American Society for Automation in Pharmacy, the National Council on Prescription Drug Programs, or other relevant national standard-setting body.

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

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

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

(3) individuals receiving medication intravenously;

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

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

(d) A dispenser must not submit data under this subdivision unless a conspicuous notice of the reporting requirements of this section is given to the patient for whom the prescription was written.

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

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

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

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

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

(d) Data reported under subdivision 4 shall be retained by the board in the database for a 12-month period, and shall be removed from the database no later than 12 months from the last day of the month during which the data was received.

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

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

(1) a prescriber or an agent or employee of the prescriber to whom the prescriber has delegated the task of accessing the data, to the extent the information relates specifically to a current patient, to whom the prescriber is prescribing or considering prescribing any controlled substance and with the provision that the prescriber remains responsible for the use or misuse of data accessed by a delegated agent or employee;

(2) a dispenser or an agent or employee of the dispenser to whom the dispenser has delegated the task of accessing the data, to the extent the information relates specifically to a current patient to whom that dispenser is dispensing or considering dispensing any controlled substance and with the provision that the dispenser remains responsible for the use or misuse of data accessed by a delegated agent or employee;

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

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

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

(6) authorized personnel of a vendor under contract with the board who are engaged in the design, implementation, operation, and maintenance of the electronic reporting system as part of the assigned duties and responsibilities of their employment, provided that access to data is limited to the minimum amount necessary to carry out such duties and responsibilities;

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

(8) personnel of the medical assistance program assigned to use the data collected under this section to identify recipients whose usage of controlled substances may warrant restriction to a single primary care physician, a single outpatient pharmacy, or a single hospital.

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

(c) Any permissible user identified in paragraph (b), who directly accesses the data electronically, shall implement and maintain a comprehensive information security program that contains administrative, technical, and physical safeguards that are appropriate to the user's size and complexity, and the sensitivity of the personal information obtained. The permissible user shall identify reasonably foreseeable internal and external risks to the security, confidentiality, and integrity of personal information that could result in the unauthorized disclosure, misuse, or other compromise of the information and assess the sufficiency of any safeguards in place to control the risks.

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

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

(f) The board shall maintain a log of all persons who access the data and shall ensure that any permissible user complies with paragraph (c) prior to attaining direct access to the data.

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

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

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

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

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

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

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

Subd. 10. **Funding.** (a) The board may seek grants and private funds from nonprofit charitable foundations, the federal government, and other sources to fund the enhancement and ongoing operations of the prescription electronic reporting system established under this section. Any funds received shall be appropriated to the board for this purpose. The board may not expend funds to enhance the program in a way that conflicts with this section without seeking approval from the legislature.

(b) The administrative services unit for the health-related licensing boards shall apportion between the Board of Medical Practice, the Board of Nursing, the Board of Dentistry, the Board of Podiatric Medicine, the Board of Optometry, and the Board of Pharmacy an amount to be paid through fees by each respective board. The amount apportioned to each board shall equal each board's share of the annual appropriation to the Board of Pharmacy from the state government special revenue fund for operating the prescription electronic reporting system under this section. Each board's apportioned share shall be based on the number of prescribers or dispensers that

each board identified in this paragraph licenses as a percentage of the total number of prescribers and dispensers licensed collectively by these boards. Each respective board may adjust the fees that the boards are required to collect to compensate for the amount apportioned to each board by the administrative services unit.

History: *2007 c 147 art 11 s 7; 2008 c 321 s 7; 2009 c 79 art 11 s 9-11; 1Sp2010 c 1 art 19 s 3*

152.13 DUTIES OF STATE BOARD OF PHARMACY.

It shall be the duty of the state board to enforce the provisions of this chapter, and the power and authority of the board, as now defined by the laws of this state, are hereby extended so as to be commensurate with the duties hereby imposed.

History: *(3899-10) 1921 c 190 s 10; 1967 c 408 s 9*

152.135 RESTRICTIONS ON SALES, MARKETING, AND POSSESSION OF EPHEDRINE.

Subdivision 1. **Prescription status for ephedrine.** Except as provided in this section, a material, compound, mixture, or preparation that contains any quantity of ephedrine, a salt of ephedrine, an optical isomer of ephedrine, or a salt of an optical isomer of ephedrine, may be dispensed only upon the prescription of a duly licensed practitioner authorized by the laws of the state to prescribe prescription drugs.

Subd. 2. **Exceptions.** (a) A drug product containing ephedrine, its salts, optical isomers, and salts of optical isomers is exempt from subdivision 1 if the drug product:

(1) may be lawfully sold over the counter without a prescription under the Federal Food, Drug, and Cosmetic Act, United States Code, title 21, section 321, et seq.;

(2) is labeled and marketed in a manner consistent with the pertinent OTC Tentative Final or Final Monograph;

(3) is manufactured and distributed for legitimate medicinal use in a manner that reduces or eliminates the likelihood of abuse;

(4) is not marketed, advertised, or labeled for the indication of stimulation, mental alertness, weight loss, muscle enhancement, appetite control, or energy;

(5) is in solid oral dosage forms, including soft gelatin caplets, that combine 400 milligrams of guaifenesin and 25 milligrams of ephedrine per dose, according to label instructions; or is an anorectal preparation containing not more than five percent ephedrine; and

(6) is sold in a manner that does not conflict with section 152.02, subdivision 6.

(b) Subdivisions 1 and 3 shall not apply to products containing ephedra or ma huang and lawfully marketed as dietary supplements under federal law.

Subd. 3. **Mismarketing of ephedrine prohibited.** The marketing, advertising, or labeling of a product containing ephedrine, a salt of ephedrine, an optical isomer of ephedrine, or a salt of an optical isomer of ephedrine for the indication of stimulation, mental alertness, weight loss, appetite control, or energy, is prohibited. In determining compliance with this subdivision, the following factors may be considered:

(1) the packaging of the drug product;

(2) the name and labeling of the product;

(3) the manner of distribution, advertising, and promotion of the product;

(4) verbal representations made concerning the product; and

(5) the duration, scope, and significance of abuse or misuse of the product.

Subd. 4. [Repealed, 1Sp2003 c 2 art 8 s 19]

Subd. 5. **Sales for illicit purposes prohibited.** It is unlawful for a person to sell, distribute, or otherwise make available a product containing ephedrine, pseudoephedrine, or phenylpropanolamine or their salts, optical isomers, or salts of optical isomers if the person knows or reasonably should know that the product will be used as a precursor to an illegal substance.

Subd. 6. **Penalty.** A person who violates this section is guilty of a misdemeanor.

History: *1998 c 367 art 4 s 6; 2005 c 136 art 7 s 10*

ANHYDROUS AMMONIA

152.136 ANHYDROUS AMMONIA; PROHIBITED CONDUCT; CRIMINAL PENALTIES; CIVIL LIABILITY.

Subdivision 1. **Definitions.** As used in this section, "tamper" means action taken by a person not authorized to take that action by law or by the owner or authorized custodian of an anhydrous ammonia container or of equipment where anhydrous ammonia is used, stored, distributed, or transported.

Subd. 2. **Prohibited conduct.** (a) A person may not:

(1) steal or unlawfully take or carry away any amount of anhydrous ammonia;

(2) purchase, possess, transfer, or distribute any amount of anhydrous ammonia, knowing, or having reason to know, that it will be used to unlawfully manufacture a controlled substance;

(3) place, have placed, or possess anhydrous ammonia in a container that is not designed, constructed, maintained, and authorized to contain or transport anhydrous ammonia;

(4) transport anhydrous ammonia in a container that is not designed, constructed, maintained, and authorized to transport anhydrous ammonia;

(5) use, deliver, receive, sell, or transport a container designed and constructed to contain anhydrous ammonia without the express consent of the owner or authorized custodian of the container; or

(6) tamper with any equipment or facility used to contain, store, or transport anhydrous ammonia.

(b) For the purposes of this subdivision, containers designed and constructed for the storage and transport of anhydrous ammonia are described in rules adopted under section 18C.121, subdivision 1, or in Code of Federal Regulations, title 49.

Subd. 3. **No cause of action.** (a) Except as provided in paragraph (b), a person tampering with anhydrous ammonia containers or equipment under subdivision 2 shall have no cause of action for damages arising out of the tampering against:

(1) the owner or lawful custodian of the container or equipment;

(2) a person responsible for the installation or maintenance of the container or equipment; or

(3) a person lawfully selling or offering for sale the anhydrous ammonia.

(b) Paragraph (a) does not apply to a cause of action against a person who unlawfully obtained the anhydrous ammonia or anhydrous ammonia container or who possesses the anhydrous ammonia or anhydrous ammonia container for any unlawful purpose.

Subd. 4. **Criminal penalty.** A person who knowingly violates subdivision 2 is guilty of a felony and may be sentenced to imprisonment for not more than five years or to payment of a fine of not more than $50,000, or both.

History: *2005 c 136 art 7 s 11*

METHAMPHETAMINE-RELATED CRIMES; CHILDREN AND VULNERABLE ADULTS

152.137 METHAMPHETAMINE-RELATED CRIMES INVOLVING CHILDREN AND VULNERABLE ADULTS.

Subdivision 1. **Definitions.** (a) As used in this section, the following terms have the meanings given.

(b) "Chemical substance" means a substance intended to be used as a precursor in the manufacture of methamphetamine or any other chemical intended to be used in the manufacture of methamphetamine.

(c) "Child" means any person under the age of 18 years.

(d) "Methamphetamine paraphernalia" means all equipment, products, and materials of any kind that are used, intended for use, or designed for use in manufacturing, injecting, ingesting, inhaling, or otherwise introducing methamphetamine into the human body.

(e) "Methamphetamine waste products" means substances, chemicals, or items of any kind used in the manufacture of methamphetamine or any part of the manufacturing process, or the by-products or degradates of manufacturing methamphetamine.

(f) "Vulnerable adult" has the meaning given in section 609.232, subdivision 11.

Subd. 2. **Prohibited conduct.** (a) No person may knowingly engage in any of the following activities in the presence of a child or vulnerable adult; in the residence of a child or a vulnerable adult; in a building, structure, conveyance, or outdoor location where a child or vulnerable adult might reasonably be expected to be present; in a room offered to the public for overnight accommodation; or in any multiple unit residential building:

(1) manufacturing or attempting to manufacture methamphetamine;

(2) storing any chemical substance;

(3) storing any methamphetamine waste products; or

(4) storing any methamphetamine paraphernalia.

(b) No person may knowingly cause or permit a child or vulnerable adult to inhale, be exposed to, have contact with, or ingest methamphetamine, a chemical substance, or methamphetamine paraphernalia.

Subd. 3. **Criminal penalty.** A person who violates subdivision 2 is guilty of a felony and may be sentenced to imprisonment for not more than five years or to payment of a fine of not more than $10,000, or both.

Subd. 4. **Multiple sentences.** Notwithstanding sections 609.035 and 609.04, a prosecution for or conviction under this section is not a bar to conviction of or punishment for any other crime committed by the defendant as part of the same conduct.

Subd. 5. **Protective custody.** A peace officer may take any child present in an area where any of the activities described in subdivision 2, paragraph (a), clauses (1) to (4), are taking place into protective custody in accordance with section 260C.175, subdivision 1, clause (2), item (ii). A child taken into protective custody under this subdivision shall be provided health screening to assess potential health concerns related to methamphetamine as provided in section 260C.188. A child not taken into protective custody under this subdivision but who is known to have been exposed to methamphetamine shall be offered health screening for potential health concerns related to methamphetamine as provided in section 260C.188.

Subd. 6. **Reporting maltreatment of vulnerable adult.** (a) A peace officer shall make a report of suspected maltreatment of a vulnerable adult if the vulnerable adult is present in an area where any of the activities described in subdivision 2, paragraph (a), clauses (1) to (4), are taking place, and the peace officer has reason to believe the vulnerable adult inhaled, was exposed to, had contact with, or ingested methamphetamine, a chemical substance, or methamphetamine paraphernalia. The peace officer shall immediately report to the county common entry point as described in section 626.557, subdivision 9b.

(b) As required in section 626.557, subdivision 9b, law enforcement is the primary agency to conduct investigations of any incident when there is reason to believe a crime has been committed. Law enforcement shall initiate a response immediately. If the common entry point notified a county agency for adult protective services, law enforcement shall cooperate with that county agency when both agencies are involved and shall exchange data to the extent authorized in section 626.557, subdivision 12b, paragraph (g). County adult protection shall initiate a response immediately.

(c) The county social services agency shall immediately respond as required in section 626.557, subdivision 10, upon receipt of a report from the common entry point staff.

History: *2005 c 136 art 7 s 12*

152.14 [Repealed, 1969 c 933 s 22]

152.15 Subdivision 1. [Repealed, 1969 c 933 s 22]

Subdivision 1. [Repealed, 1989 c 290 art 3 s 37]

Subd. 2. [Repealed, 1989 c 290 art 3 s 37]

Subd. 2a. [Repealed, 1989 c 290 art 3 s 37]

Subd. 2b. [Repealed, 1989 c 290 art 3 s 37]

Subd. 3. [Repealed, 1989 c 290 art 3 s 37]

Subd. 4. [Repealed, 1987 c 330 s 4]

Subd. 4a. [Repealed, 1989 c 290 art 3 s 37]

Subd. 5. [Repealed, 1989 c 290 art 3 s 37]

MISCELLANEOUS PROVISIONS

152.151 [Repealed, 1996 c 310 s 1]

152.152 STAYED SENTENCE LIMITED.

If a person is convicted under section 152.021, 152.022, 152.023, or 152.0262, and the Sentencing Guidelines grid calls for a presumptive prison sentence for the offense, the court may stay imposition or execution of the sentence only as provided in this section. The sentence may be stayed based on amenability to probation only if the offender presents adequate evidence to the court that the offender has been accepted by, and can respond to, a treatment program that has been approved by the commissioner of human services. The court may impose a sentence that is a mitigated dispositional departure on any other ground only if the court includes as a condition of probation incarceration in a local jail or workhouse.

History: *1989 c 290 art 3 s 20; 2005 c 136 art 7 s 21*

152.16 [Repealed, 1967 c 408 s 11]

152.17 [Repealed, 1971 c 937 s 22]

152.18 DISCHARGE AND DISMISSAL.

Subdivision 1. **Deferring prosecution for certain first time drug offenders.** If any person who has not previously participated in or completed a diversion program authorized under section 401.065 or who has not previously been placed on probation without a judgment of guilty and thereafter been discharged from probation under this section is found guilty of a violation of section 152.024, subdivision 2, 152.025, subdivision 2, or 152.027, subdivision 2, 3, 4, or 6, paragraph (d), for possession of a controlled substance, after trial or upon a plea of guilty, and the court determines that the violation does not qualify as a subsequent controlled substance conviction under section 152.01, subdivision 16a, the court may, without entering a judgment of guilty and with the consent of the person, defer further proceedings and place the person on probation upon such reasonable conditions as it may require and for a period, not to exceed the maximum sentence provided for the violation. The court may give the person the opportunity to attend and participate in an appropriate program of education regarding the nature and effects of alcohol and drug abuse as a stipulation of probation. Upon violation of a condition of the probation, the court may enter an adjudication of guilt and proceed as otherwise provided. The court may, in its discretion, dismiss the proceedings against the person and discharge the person from probation before the expiration of the maximum period prescribed for the person's probation. If during the period of probation the person does not violate any of the conditions of the probation, then upon expiration of the period the court shall discharge the person and dismiss the proceedings against that person. Discharge and dismissal under this subdivision shall be without court adjudication of guilt, but a not public record of it shall be retained by the Bureau of Criminal Apprehension for the purpose of use by the courts in determining the merits of subsequent proceedings against the person. The not public record may also be opened only upon court order for purposes of a criminal investigation, prosecution, or sentencing. Upon request by law enforcement, prosecution, or corrections authorities, the bureau shall notify the requesting party of the existence of the not public record and the right to seek a court order to open it pursuant to this section. The court shall forward a record of any discharge and dismissal under this subdivision to the bureau which shall make and maintain the not public record of it as provided under this subdivision. The discharge or dismissal shall not be deemed a conviction for purposes of disqualifications or disabilities imposed by law upon conviction of a crime or for any other purpose.

For purposes of this subdivision, "not public" has the meaning given in section 13.02, subdivision 8a.

Subd. 2. [Repealed, 1996 c 408 art 9 s 10]

Subd. 3. **Expungement of certain marijuana offenses.** Any person who has been found guilty of a violation of section 152.09 with respect to a small amount of marijuana which violation occurred prior to April 11, 1976, and whose conviction would have been a petty misdemeanor under the provisions of section 152.15, subdivision 2, clause (5) in effect on April 11, 1978, but whose conviction was for an offense more serious than a petty misdemeanor under laws in effect prior to April 11, 1976, may petition the court in which the person was convicted to expunge from all official records, other than the nonpublic record retained by the Department of Public Safety pursuant to section 152.15, subdivision 2, clause (5), all recordation relating to the person's arrest, indictment or information, trial and conviction of an offense more serious than a petty misdemeanor. The court, upon being satisfied that a small amount was involved in the conviction, shall order all the recordation expunged. No person as to whom an order has been entered pursuant to this subdivision shall be held thereafter under any provision of any law to be guilty of perjury or otherwise giving a false statement by reason of the person's failure to recite or acknowledge conviction of an offense greater than a petty misdemeanor, unless possession of marijuana is material to a proceeding.

History: *1971 c 937 s 18; 1973 c 693 s 14; 1978 c 639 s 1; 1986 c 444; 1989 c 290 art 3 s 21; 1992 c 569 s 13; 1993 c 326 art 13 s 11; 1995 c 226 art 2 s 2; 1996 c 408 art 9 s 2; 2012 c 240 s 3*

152.19 [Repealed, 1988 c 665 s 17]

152.20 PENALTIES UNDER OTHER LAWS.

Any penalty imposed for violation of this chapter is in addition to, and not in lieu of, any civil or administrative penalty or sanction otherwise authorized by law.

History: *1971 c 937 s 20; 1989 c 290 art 3 s 22*

152.205 LOCAL REGULATIONS.

Sections 152.01, subdivision 18, and 152.092 to 152.095 do not preempt enforcement or preclude adoption of municipal or county ordinances prohibiting or otherwise regulating the manufacture, delivery, possession, or advertisement of drug paraphernalia.

History: *1982 c 557 s 11; 1988 c 665 s 1*

152.21 THC THERAPEUTIC RESEARCH ACT.

Subdivision 1. **Findings and purpose.** The legislature finds that scientific literature indicates promise for delta-9-tetrahydro-cannabinol (THC), the active component of marijuana, in alleviating certain side effects of cancer chemotherapy under strictly controlled medical circumstances.

The legislature also finds that further research and strictly controlled experimentation regarding the therapeutic use of THC is necessary and desirable. The intent of this section is to establish an extensive research program to investigate and report on the therapeutic effects of THC under strictly controlled circumstances in compliance with all federal laws and regulations promulgated by the federal Food and Drug Administration, the National Institute on Drug Abuse and the Drug Enforcement Administration. The intent of the legislature is to allow this research

program the greatest possible access to qualified cancer patients residing in Minnesota who meet protocol requirements. The establishment of this research program is not intended in any manner whatsoever to condone or promote the illicit recreational use of marijuana.

Subd. 2. **Definitions.** For purposes of this section, the following terms shall have the meanings given.

(a) "Commissioner" means the commissioner of health.

(b) "Marijuana" means marijuana as defined in section 152.01, subdivision 9, and delta-9-tetrahydro-cannabinol (THC), tetrahydrocannabinols or a chemical derivative of tetrahydrocannabinols, and all species of the genus Cannabis.

(c) "Principal investigator" means the individual responsible for the medical and scientific aspects of the research, development of protocol, and contacting and qualifying the clinical investigators in the state.

(d) "Clinical investigators" means those individuals who conduct the clinical trials.

(e) "Sponsor" means that individual or organization who, acting on behalf of the state, has the total responsibility for the state program.

Subd. 3. **Research grant.** The commissioner of health shall grant funds to the principal investigator selected by the commissioner pursuant to subdivision 4 for the purpose of conducting a research program under a protocol approved by the FDA regarding the therapeutic use of oral THC and other dosage forms, if available, according to the guidelines and requirements of the federal Food and Drug Administration, the Drug Enforcement Administration and the National Institute on Drug Abuse. The commissioner shall ensure that the research principal investigator complies with the requirements of subdivision 5. The commissioner may designate the principal investigator as the sponsor.

Subd. 4. **Principal investigator.** Within three months of April 25, 1980, the commissioner shall, in consultation with a representative chosen by the state Board of Pharmacy and a representative chosen by the state Board of Medical Examiners, select a person or research organization to be the principal investigator of the research program.

Subd. 5. **Duties.** The principal investigator shall:

(1) Apply to the Food and Drug Administration for a notice of "Claimed Investigational Exemption for a New Drug (IND)" pursuant to the Federal Food, Drug and Cosmetic Act, United States Code, title 21, section 301, et seq., and shall comply with all applicable laws and regulations of the federal Food and Drug Administration, the Drug Enforcement Administration, and the National Institute on Drug Abuse in establishing the program;

(2) Notify every oncologist in the state of the program, explain the purposes and requirements of the program to them, provide on request each of them with a copy of the approved protocol which shall include summaries of current papers in medical journals reporting on research concerning the safety, efficacy and appropriate use of THC in alleviating the nausea and emetic effects of cancer chemotherapy, and provide on request each of them with a bibliography of other articles published in medical journals;

(3) Allow each oncologist (clinical investigator) in the state who meets or agrees to meet all applicable federal requirements for investigational new drug research and who so requests to be included in the research program as a clinical investigator to conduct the clinical trials;

(4) Provide explanatory information and assistance to each clinical investigator in understanding the nature of therapeutic use of THC within program requirements, including the informed consent document contained in the protocol, informing and counseling patients involved in the program regarding the appropriate use and the effects of therapeutic use of THC;

(5) Apply to contract with the National Institute on Drug Abuse for receipt of dosage forms of THC, fully characterized as to contents and delivery to the human system, pursuant to regulations promulgated by the National Institute on Drug Abuse, and the federal Food and Drug Administration. The principal investigator shall ensure delivery of the THC dosages to clinical investigators as needed for participation in the program;

(6) Conduct the research program in compliance with federal laws and regulations promulgated by the federal Food and Drug Administration, the Drug Enforcement Administration, the National Institute on Drug Abuse, and the purposes and provisions of this section;

(7) Submit periodic reports as determined by the commissioner on the numbers of oncologists and patients involved in the program and the results of the program;

(8) Submit reports on intermediate or final research results, as appropriate, to the major scientific journals in the United States; and

(9) Otherwise comply with the provisions of this section.

Subd. 6. **Exemption from criminal sanctions.** For the purposes of this section, the following are not violations under this chapter:

(1) use or possession of THC, or both, by a patient in the research program;

(2) possession, prescribing use of, administering, or dispensing THC, or any combination of these actions, by the principal investigator or by any clinical investigator; and

(3) possession or distribution of THC, or both, by a pharmacy registered to handle Schedule I substances which stores THC on behalf of the principal investigator or a clinical investigator.

THC obtained and distributed pursuant to this section is not subject to forfeiture under sections 609.531 to 609.5316.

For the purposes of this section, THC is removed from Schedule I contained in section 152.02, subdivision 2, and inserted in schedule II contained in section 152.02, subdivision 3.

Subd. 7. **Citation.** This section may be cited as the "THC Therapeutic Research Act."

History: *1980 c 614 s 93; 1988 c 665 s 2; 1989 c 290 art 3 s 23; 1997 c 7 art 2 s 22*