

2. Be authorized to dispense a standard compounded formula of Sodium Pentobarbital for animal euthanasia established by the Department as follows:

i. Sodium Pentobarbital injection (for animal euthanasia), formula non-sterile solution:

U.S.P. Pentobarbital Sodium (Powder)	460 grams
Isopropyl Alcohol	250 mls.
Methyl Violet	1 drop
U.S.P. Water for injection	
Quantity sufficient to make	1000 mls.

ii. Using the formula in (a)2 above, the strength of this mixture will provide 460 mgs of Pentobarbital Sodium per milliliter.

iii. Lethal dose: one milliliter per 10 pounds of body weight for small animals; horses and other large animals—one milliliter per 10 pounds of body weight subject to a maximum dose of 100 milliliters.

iv. Package and storage: Package in tight containers with rubber stoppers and store under refrigeration. Solutions decompose on standing. Heat accelerates the decomposition.

v. Expiration date: five days from date of manufacture.

(b) Labeling: sample labeling is as follows:

- 1.
- 2.
- 3.
- 4.
- 5.
- 6.
- 7.
- 8.
- 9.

1. Name and address, city and State of registrant;
2. Name of preparation: "Pentobarbital Sodium Injection";
3. Strength of the preparation: "460 milligrams per one milliliter";
4. "Lethal dose: one milliliter per 10 pounds of body weight for small animals; horses and large animals—one milliliter per 10 pounds of body weight subject to a maximum dose of 100 milliliters";
5. "Batch number";
6. "Net contents";
7. "Expiration date";
8. "Keep under refrigeration.";
9. "Warning: Do not use the injection if it contains a precipitate."

(c) A master formula and production record shall be made and retained on file at the formulating (compounding) site. This record shall contain:

1. Name, address, city and State of registrant;
2. Name and strength of the product and a description of the dosage form;
3. The name and weight or measure of each active ingredient including the control number of each such ingredient;
4. A statement of the theoretical yield of finished product;
5. A statement describing the equipment and utensils used in the formulating (compounding);
6. A description of the finished drug product containers and closures including a specimen or copy of each label and all other labeling signed and dated by the person or persons responsible for approval of such labeling; and
7. Complete manufacturing and control instructions, procedures, special notations and precautions to be followed.

(d) Batch production records shall be prepared for each batch of drug product produced and shall include complete information relating to the production of each batch. The records shall contain:

1. An accurate reproduction of the appropriate master formula production record, checked for accuracy, dated and signed;
2. Documentation that each significant step in the manufacture, processing, packaging or holding of the batch was accomplished, including:
 - i. Dates;
 - ii. Identity of the individual equipment used;
 - iii. Specific identification of each batch of component or materials used;
 - iv. Weights and/or measures of components used in processing;
 - v. Copy of all labeling used;
 - vi. Identification of the person performing each step in the process and identification of the person checking the weights, measures and operations;
 - vii. A statement of the theoretical yield; and
 - viii. A statement of the actual yield.

New rule, R.1988 d.498, effective October 17, 1988. See: 20 N.J.R. 366(a), 20 N.J.R. 2574(b).

SUBCHAPTER 9. (RESERVED)

SUBCHAPTER 10. CONTROLLED DANGEROUS
SUBSTANCES SCHEDULES

**13:45H-10.1 Schedules of controlled dangerous
substances**

(a) The Federal controlled dangerous substance Schedules I, II, III, IV and V at 21 CFR 1308.11 through 1308.15, as amended and supplemented, promulgated by the United States Attorney General pursuant to 21 U.S.C. §§811 and 812, are incorporated herein by reference.

(b) Any reference in this chapter to controlled dangerous substance Schedules I, II, III, IV and V shall mean the Federal schedules promulgated at 21 CFR 1308.11 through 1308.15 and incorporated by reference pursuant to (a) above, unless the Director objects to the inclusion, rescheduling or deletion of a substance in accordance with the provisions of N.J.S.A. 24:21-3 and N.J.A.C. 13:45H-1.7.

(c) Any substance designated as an immediate precursor by the United States Attorney General pursuant to 21 U.S.C. §811(e), or designated a controlled dangerous substance by temporary order issued by the United States Attorney General in accordance with and subject to the provisions of 21 U.S.C. §811(d) or (h), as amended and supplemented, shall be subject to regulation under this chapter.

(d) Notwithstanding the provisions of (b) above, any substance that is an immediate precursor or that, when ingested, is metabolized or otherwise becomes a controlled dangerous substance, may be designated by the Director as a controlled dangerous substance.

(e) In accordance with (d) above, the following substances shall be designated and controlled as Schedule I controlled dangerous substances:

1. Gamma Butyrolactone
2. 1,4 Butanediol

Amended by R.1973 d.23, effective January 17, 1973.
See: 5 N.J.R. 42(b).

Tetrahydrocannabinols defined (Fed. Reg. V.36, April 28, 1975).
Amended by R.1973 d.325, effective November 20, 1973.
See: 5 N.J.R. 413(b).

Mecloqualone controlled (Fed. Reg. V.40, 28611, July 2, 1975).
Amended by R.1974 d.2, effective January 2, 1974.
See: 6 N.J.R. 63(a).

Propiram controlled (Fed. Reg. V.37, 1468, February 28, 1972).
Drotebanol controlled (Fed. Reg. V.38, 17717, August 6, 1973).
4-bromo-2,5-dimethoxy-amphetamine controlled (Fed. Reg. V.38, 26447, September 21, 1973).
2,5-dimethoxyamphetamine controlled (Fed. Reg. V.38, 26447, September 21, 1973).

4-methoxyamphetamine controlled (Fed. Reg. V.38, 26447, September 21, 1973).
Amended by R.1974 d.106, effective May 1, 1974.
See: 5 N.J.R. 413(b), 6 N.J.R. 241(b).

Mecoqualone controlled (Fed. Reg. V.40, 28611, July 2, 1975).
Amended by R.1975 d.209, effective July 23, 1975.
See: 7 N.J.R. 353(a).

Difenoxin controlled (Fed. Reg. V.40, 19813, June 1, 1975).

Etorpine HCl rescheduled from schedule I to II (Fed. Reg. V.39, 11535, April 19, 1974).

Peyote defined (Fed. Reg. V.40, April 28, 1975).
Amended by R.1977 d.441, effective November 11, 1977.
See: 9 N.J.R. 466(a), 9 N.J.R. 567(b).

Thiophene analog controlled (Fed. Reg. V.40, 28611, August 11, 1975).

Amended by R.1979 d.27, effective January 17, 1979.
See: 10 N.J.R. 534(b), 11 N.J.R. 67(c).

N-Ethyl-1-phenylcyclohexylamine controlled (Fed. Reg. V.3, 186, September 25, 1978).

1-(1-phenylcyclohexyl) pyrrolidine controlled (Fed. Reg. V.3, 186, September 25, 1978).

Amended by R.1980 d.322, effective July 17, 1980.

See: 12 N.J.R. 467(g).

Dextrophan decontrolled (Fed. Reg. V.41, 43401-43402, October 1, 1976).

Amended by R.1981 d.50, effective February 9, 1981.

See: 13 N.J.R. 132(b).

(b)1: Added Sufentanil and Tilidine.

Amended by R.1981 d.50, effective February 9, 1981.

See: 13 N.J.R. 132(b).

(b)6 added.

Amended by R.1982 d.124, effective April 19, 1982.

See: 14 N.J.R. 195(a), 14 N.J.R. 389(b).

Amended by R.1982 d.450, effective December 20, 1982, operative January 10, 1983.

See: 14 N.J.R. 1029(b), 14 N.J.R. 1457(b).

Added methaqualone to Schedule 1.

Amended by R.1983 d.339, effective August 15, 1983.

See: 15 N.J.R. 844(a), 15 N.J.R. 1375(a).

Added Parahexyl to (b)3. and also added 6. "Stimulants"—
Fenethylamine and n-Ethylamphetamine.

Amended by R.1984 d.532, effective November 19, 1984.

See: 16 N.J.R. 2332(a), 16 N.J.R. 3204(a).

(b)1: Added Alfentanil.

Amended by R.1985 d.83, effective March 4, 1985.

See: 16 N.J.R. 2900(a), 17 N.J.R. 592(a).

(b)1: Deleted "Sufentanil".

Amended by R.1985 d.458, effective September 3, 1985.

See: 17 N.J.R. 1511(a), 17 N.J.R. 2138(b).

(b)7 added.

Amended by R.1985 d.669 effective January 6, 1986.

See: 17 N.J.R. 2214(a), 18 N.J.R. 87(a).

(b)7ii added.

Amended by R.1986 d.65, effective March 17, 1986.

See: 17 N.J.R. 2721(a), 18 N.J.R. 555(c).

(b)7iii and iv added.

Amended by R.1986 d.184, effective May 19, 1986.

See: 18 N.J.R. 154(b), 18 N.J.R. 1102(a).

(b)7v.-xii. added.

Amended by R.1986 d.326, effective August 4, 1986.

See: 18 N.J.R. 603(a), 18 N.J.R. 1591(b).

(b)7xii added.

Amended by R.1986 d.457, effective November 17, 1986.

See: 18 N.J.R. 1774(a), 18 N.J.R. 2327(b).

(b)3 "excepted" substituted for "expected".

Amended by R.1987 d.324, effective August 3, 1987.

See: 19 N.J.R. 841(a), 19 N.J.R. 1454(a).

Deleted Alfentanil 9737 and moved it to section 2.

Commissioner's Order, effective March 21, 1988.

See: 20 N.J.R. 1000(c).

Added 3,4-methylenedioxy-N-ethylamphetamine (7404); N-hydroxy-3,4-methylenedioxyamphetamine (7402) and 4-methylaminorex (1590) to Schedule I at (b)3. These drugs were published in the Federal Register on October 15, 1987 (see 52 FR 38225) pursuant to N.J.S.A. 24:21-3(c).
Commissioner's Order, effective March 21, 1988.

See: 20 N.J.R. 1000(d).

Added Beta-hydroxy-3-methylfentanyl (CDS-9830) to Schedule I at (b)3. This drug was published in the Federal Register on January 8, 1988 (see 53 FR 500) pursuant to N.J.S.A. 24:21-3(c).

Commissioner's Order, effective March 23, 1988.

See: 20 N.J.R. 1001(a).

Added Methylenedioxymethamphetamine (CDS-7405) to Schedule I at (b)3. This drug was published in the Federal Register on February 22, 1988 (see FR 5156) pursuant to N.J.S.A. 24:21-3(c). Commissioner's Order, effective September 14, 1989. See: 21 N.J.R. 3297(b).

Added 1-[1-(2-thienyl)cyclohexyl]pyrrolidine (CDS 7473) to Schedule I at (b)3, pursuant to N.J.S.A. 24:21-3(c). A final order scheduling this substance was published in the Federal Register on July 6, 1989 (see 54 FR 28414).

Commissioner's order, effective May 1, 1992. See: 24 N.J.R. 2451(a).

Methcathinone (CDS 1237) added at (b)7xvi. A final order scheduling this substance was published in the Federal Register on May 1, 1992 (see 57 F.R. 18824).

Amended by R.1994 d.325, effective July 5, 1994. See: 26 N.J.R. 1630(a), 26 N.J.R. 2792(b).

Acetylmethadol 9603 added.

Repeal and New Rule, R.2008 d.58, effective March 17, 2008.

See: 39 N.J.R. 3854(a), 40 N.J.R. 1680(a).

Section was "Controlled dangerous substances; Schedule I".

Case Notes

School Board urinalysis drug/alcohol test policy for students unconstitutional: reasonableness test for permissible student search not met. *Odenheim v. Carlstadt-East Rutherford Regional School District*, 211 N.J.Super. 54 (Ch.Div.1985).

Constructive publication of addition to controlled dangerous substance schedule in New Jersey Register: lack of notice defense denied. *State v. Metcalf*, 168 N.J.Super. 375 (App.Div.1979), 81 N.J. 411 (1979).

13:45H-10.2 (Reserved)

Amended by R.1973, d.23, effective January 17, 1973.

See: 5 N.J.R. 42(b).

Naloxone decontrolled (Fed. Reg. V.36, 19116, September 29, 1971; V.39, 44392, December 24, 1974).

Amphetamine rescheduled from schedule III to II (Fed. Reg. V.36, 12734, July 7, 1971).

Methamphetamine rescheduled and schedule III to II (Fed. Reg. V.36, 12734, July 7, 1971).

Phenmetrazine rescheduled III to II (Fed. Reg. V.36, 20686, October 28, 1971).

Methylphenidate rescheduled from schedule III to II (Fed. Reg. V.36, 20686, October 28, 1971).

Amended by R.1973 d.147, effective June 5, 1973.

See: 5 N.J.R. 106(b), 5 N.J.R. 222(d).

Methaqualone controlled (Fed. Reg. V.38, 27517, October 4, 1973).

Amended by R.1974 d.2, effective January 2, 1974.

See: 6 N.J.R. 63(a).

Amobarbital rescheduled from schedule III to II (Fed. Reg. V.38, 31310, November 13, 1973).

Secobarbital rescheduled from schedule III to II (Fed. Reg. V.38, 31310, November 13, 1973).

Pentobarbital rescheduled from schedule III to II (Fed. Reg. V.38, 31310, November 13, 1973).

Amended by R.1984 d.35, effective February 21, 1984.

See: 15 N.J.R. 844(a), 16 N.J.R. 369(a).

Glutethimide-2550; added.

Amended by R.1975 d.209, effective July 23, 1975.

See: 7 N.J.R. 363(a).

Naltrexone decontrolled (Fed. Reg. V.40, 10455, March 6, 1975).

Etorphine HCl rescheduled from schedule I to II (Fed. Reg. V.39, 11535, April 19, 1974).

Concentration of poppy straw defined (Fed. Reg. V.40, 6679, February 14, 1975).

Amended by R.1978 d.247, effective July 24, 1978.

See: 10 N.J.R. 238(a), 10 N.J.R. 341(b).

Phencyclidine rescheduled from III to II (Fed. Reg. V.43, 17, January 25, 1978).

Amended by R.1984 d.35, effective February 21, 1984.

See: 15 N.J.R. 844(a), 16 N.J.R. 369(a).

Glutethimide-2550; added.

Amended by R.1978 d.390, effective November 1, 1978.

See: 10 N.J.R. 427(a), 10 N.J.R. 536(b).

Amended by R.1982 d.450, effective December 20, 1982, operative January 10, 1983.

See: 14 N.J.R. 1029(b), 14 N.J.R. 1457(b).

Deleted methaqualone from Schedule II.

Phenylcyclohexylamine and piperidinocyclohexane-carbonitrile (PCC) controlled (Fed. Reg. V.43, 21324-21325, June 16, 1978).

Amended by R.1980 d.37, effective February 11, 1980.

See: 12 N.J.R. 76(a).

Adopted paragraph (b)5.

Phenylacetone controlled (Fed. Reg. V.44, 71822-71824, effective February 11, 1980).

Amended by R.1980 d.38, effective February 11, 1980.

See: 12 N.J.R. 76(b).

Adopted subparagraph ii.

Transfer phenylcyclohexylamine and piperidinocyclohexane-carbonitrile (PCC) from paragraph (b)4 to subparagraph (b)5ii.

Amended by R.1980 d.323, effective July 17, 1980.

See: 12 N.J.R. 468(a).

Apomorphine and nalbuphine (never specifically included) decontrolled (Fed. Reg. V.41, 43401-43402, October 1, 1976).

Amended by R.1981 d.50, effective February 9, 1981.

See: 13 N.J.R. 132(b).

(b)6 added.

Amended by R.1985 d.83, effective March 4, 1985.

See: 16 N.J.R. 2900(a), 17 N.J.R. 592(a).

Amended by R.1986 d.327, effective August 4, 1986.

See: 18 N.J.R. 536(a), 18 N.J.R. 1592(a).

(b)1i: added dextrophan and nalmefene.

Amended by R.1986 d.457, effective November 17, 1986.

See: 18 N.J.R. 1774(a), 18 N.J.R. 2327(b).

(b)7 added.

Amended by R.1987 d.324, effective August 3, 1987.

See: 19 N.J.R. 841(a), 19 N.J.R. 1454(a).

Moved Alfentanil from section 1.

Amended by R.1987 d.339, effective August 17, 1987.

See: 19 N.J.R. 1050(a), 19 N.J.R. 1557(a).

(b)7iii added Nabilone.

Commissioner's Order, effective January 3, 1989.

See: 21 N.J.R. 70(b).

Added Carfentanil (CDS 9743) to Schedule II at (b)2. This drug was published in the Federal Register on October 28, 1988 (see 53 FR 43684) pursuant to N.J.S.A. 24:21-3(c).

Amended by R.1994 d.325, effective July 5, 1994.

See: 26 N.J.R. 1630(a), 26 N.J.R. 2792(b).

Levo-alphaacetylmethadol 9648 added.

Repealed by R.2008 d.58, effective March 17, 2008.

See: 39 N.J.R. 3854(a), 40 N.J.R. 1680(a).

Section was "Controlled dangerous substances; Schedule II".

13:45H-10.3 (Reserved)

As amended, R.1973 d.23, effective January 17, 1983.

See: 5 N.J.R. 42(b).

Amphetamine rescheduled from schedule III to II (Fed. Reg. V.36, 12734, July 7, 1974).

Methamphetamine rescheduled from schedule III to II (Fed. Reg. V.36, 12734, July 7, 1974).

Phenmetrazine rescheduled from schedule III to II (Fed. Reg. V.36, 20686, October 28, 1971).

Methylphenidate rescheduled from III to II (Fed. Reg. V.36, 20686, October 28, 1971).

As amended, R.1974 d.2, effective January 2, 1974.

See: 6 N.J.R. 53(a).

Amobarbital (except in suppository form or in combination with one or more active medicinal ingredients) rescheduled from schedule III to II (Fed. Reg. V.38, 31310, November 13, 1973).

Secobarbital rescheduled from schedule III to II (Fed. Reg. V.38, 31310, November 13, 1973).

Pentobarbital rescheduled from schedule III to II (Fed. Reg. V.38, 31310, November 13, 1973).

Benzphetamine controlled (Fed. Reg. V.38, 15719, June 15, 1973).

Chlorphentermine controlled (Fed. Reg. V.38, 15719, June 15, 1973).
 Clortermine controlled (Fed. Reg. V.38, 15719, June 15, 1973).
 Mazindol controlled (Fed. Reg. V.38, 15719, June 15, 1973).
 Phendimetrazine controlled (Fed. Reg. V.38, 15719, June 15, 1973).
 Amended by R.1987 d.339, effective August 17, 1987.
 See: 19 N.J.R. 911(a), 19 N.J.R. 1557(a).
 Changed code numbers.
 Schedule II stimulates in dosage form listed on August 25, 1971 as excepted compounds under section 308.32 controlled (Fed. Reg. V.38, 15719, June 15, 1973).
 As amended, R.1978 d.247, effective July 24, 1978.
 See: 10 N.J.R. 238(a), 10 N.J.R. 341(b).
 Phencyclidine rescheduled from schedule III to II (Fed. Reg. V.43, 17, January 25, 1978).
 As amended, R.1984 d.35, effective February 21, 1984.
 See: 15 N.J.R. 844(a), 16 N.J.R. 369(a).
 Glutethimide—2550; deleted.
 As amended, R.1979 d.301, effective August 6, 1979.
 See: 11 N.J.R. 440(e).
 Amended text of paragraph (b)3.
 As amended, R.1982 d.124, effective April 19, 1982.
 See: 14 N.J.R. 195(a), 14 N.J.R. 389(b).
 (b)1ii: Deleted “mazindol 1605”.
 Amended by R.1987 d.337, effective August 17, 1987.
 See: 19 N.J.R. 497(a), 19 N.J.R. 1557(b).
 Added (b)2v Tiletamine.
 Amended by R.1987 d.339, effective August 17, 1987.
 See: 19 N.J.R. 911(a), 19 N.J.R. 1557(a).
 Changed code numbers.
 Amended by Commissioner’s Order.
 See: 23 N.J.R. 1943(b).
 Anabolic steroids added at (b)4.
 Commissioner’s Order, effective June 15, 1992.
 See: 24 N.J.R. 2256(b).
 Spelling corrected in (b)4; salts, esters and isomers added.
 Amended by R.1997 d.326, effective August 4, 1997.
 See: 29 N.J.R. 2262(a), 29 N.J.R. 3450(b).
 Repealed by R.2008 d.58, effective March 17, 2008.
 See: 39 N.J.R. 3854(a), 40 N.J.R. 1680(a).
 Section was “Controlled dangerous substances; Schedule III”.

13:45H-10.4 (Reserved)

As amended, R.1974 d.2, effective January 2, 1974.
 See: 6 N.J.R. 63(a).
 Diethylpropion controlled (Fed. Reg. V.38, 18013, July 6, 1973).
 Phentermine controlled (Fed. Reg. V.38, 18013, July 6, 1973).
 Fenfluramine controlled (Fed. Reg. V.38, 15719, June 15, 1973).
 As amended, R.1975 d.209, effective July 23, 1975.
 See: 7 N.J.R. 363(a).
 Pemoline controlled (Fed. Reg. V.40, 4150, January 1, 1975).
 Chlordiazepoxide controlled (Fed. Reg. V.40, 23998, July 2, 1975).
 Clonazepam controlled (Fed. Reg. V.40, 23998, July 2, 1975).
 Clorazepate controlled (Fed. Reg. V.40, 23998, July 2, 1975).
 Diazepam controlled (Fed. Reg. V.40, 23998, July 2, 1975).
 Flurazepam controlled (Fed. Reg. V.40, 23998, July 2, 1975).
 Mebutamate controlled (Fed. Reg. V.40, 4418, January 30, 1975).
 Oxazepam controlled (Fed. Reg. V.40, 23998, July 2, 1975).
 As amended, R.1977 d.101, effective March 29, 1977.
 See: 9 N.J.R. 79(a), 9 N.J.R. 172(d).
 Prazepam controlled (Fed. Reg. V.41, 55176, December 17, 1976).
 As amended, R.1977 d.151, effective April 28, 1977.
 See: 9 N.J.R. 119(b), 9 N.J.R. 268(a).
 Dextropropoxyphene controlled (Fed. Reg. V.42, 9636, March 14, 1977).
 As amended, R.1978 d.33, effective January 24, 1978.
 See: 9 N.J.R. 562(b), 10 N.J.R. 62(b).
 Lorazepam controlled (Fed. Reg. V.42, 545747, October 8, 1977).
 As amended, R.1978 d.426, effective December 12, 1978.
 See: 10 N.J.R. 482(b), 11 N.J.R. 16(a).
 Paragraph 4 subsection (b) adopted.
 Difenoixin in combination with atropine sulfate controlled (Fed. Reg. V.43, No. 167, August 28, 1978).

As amended, R.1979 d.150, effective April 18, 1979.
 See: 11 N.J.R. 129(b), 11 N.J.R. 236(c).
 Pentazocine controlled (Fed. Reg. V.44 No. 7, January 10, 1979).
 As amended, R.1979 d.301, effective August 6, 1979.
 See: 11 N.J.R. 440(e).
 Amended text of paragraph (b)4.
 As amended, R.1980 d.327, effective July 18, 1980.
 See: 12 N.J.R. 468(b).
 Dextropropoxyphene moved from (b)3 to (b)4 (Fed. Reg. V.45, 42264-42265, July 24, 1980).
 As amended, R.1982 d.124, effective April 19, 1982.
 See: 14 N.J.R. 195(a), 14 N.J.R. 398(b).
 (b)1: Added “Mazindol 1650”;
 (b)2: Added “Halazepam 2762”;
 (b)3: Added “Alphazolam 2882”;
 As amended, R.1983 d.339, effective August 15, 1983.
 See: 15 N.J.R. 844(a), 15 N.J.R. 1375(a).
 Added Temazepam to (b)2.
 Amended by R.1985 d.191, effective April 15, 1985.
 See: 16 N.J.R. 2901(a), 17 N.J.R. 957(a).
 Temazepam added.
 As amended, R.1985, d.190, effective April 15, 1985.
 See: 16 N.J.R. 3390(a), 17 N.J.R. 956(a).
 Substantially amended.
 Amended by R.1986 d.374, effective September 8, 1986.
 See: 18 N.J.R. 1166(b), 18 N.J.R. 1827(a).
 Added midazolam and quazepam.
 Amended by R.1987 d.340, effective August 17, 1987.
 See: 19 N.J.R. 911(a), 19 N.J.R. 1557(c).
 Deleted (b)4i and substituted new.
 Commissioner’s Order, effective January 3, 1989.
 See: 21 N.J.R. 70(c).
 Added Cathine (1230), Fencamfamin (1760), Fenproporex (1575) and Mefenorex (1580) to Schedule IV at (b)1. These drugs were published in the Federal Register on May 17, 1988 (see 53 FR 17459) pursuant to N.J.S.A. 24:21-3(c).
 Repealed by R.2008 d.58, effective March 17, 2008.
 See: 39 N.J.R. 3854(a), 40 N.J.R. 1680(a).
 Section was “Controlled dangerous substances; Schedule IV”.

13:45H-10.5 (Reserved)

As amended, R.1977 d.440, effective November 4, 1977.
 See: 9 N.J.R. 465(c), 9 N.J.R. 567(a).
 Loperamide controlled (Fed. Reg. V.42, 54547, October 8, 1977).
 As amended, R.1978 d.426, effective December 12, 1978.
 See: 10 N.J.R. 482(b), 11 N.J.R. 16(a).
 Difenoixin in combination with atropine sulfate controlled (Fed. Reg. V.43, No. 167, August 28, 1978).
 As amended, R.1979 d.301, effective August 6, 1979.
 See: 11 N.J.R. 440(e).
 Amended text of paragraph (b)1.
 As amended, R.1983 d.171, effective June 6, 1983.
 See: 15 N.J.R. 126(a), 15 N.J.R. 923(a).
 Deleted Loperamide as a controlled dangerous substance.
 Amended by R.1985 d.460, effective September 3, 1985.
 See: 17 N.J.R. 1234(a), 17 N.J.R. 2138(c).
 (b)2 added. Buphenorphine added as a controlled dangerous substance.
 Amended by R.1988 d.496, effective October 11, 1988.
 See: 20 N.J.R. 1506(a), 20 N.J.R. 2575(a).
 Substantially amended.
 Commissioner’s Order, effective January 3, 1989.
 See: 21 N.J.R. 70(d).
 Added Propylhexadrine (8161) and Pyrovalerone (1485) to Schedule V at (d). The drugs were published in the Federal Register on April 4, 1988 (see 53 FR 10869) pursuant to N.J.S.A. 24:21-3(c).
 Administrative Correction to spell Buprenorphine correctly.
 See: 22 N.J.R. 3619(c).
 Public Notice, effective March 16, 1992.
 See: 24 N.J.R. 947(a).
 Deletion of Propylhexadrine.