

SENATE BILL No. 1237

March 26, 2008, Introduced by Senators SWITALSKI, GLEASON, ANDERSON, KAHN,
PAPPAGEORGE, BIRKHOLZ and JACOBS and referred to the Committee on Health Policy.

A bill to amend 1978 PA 368, entitled
"Public health code,"
by amending section 7216 (MCL 333.7216), as amended by 1999 PA
42.

THE PEOPLE OF THE STATE OF MICHIGAN ENACT:

1 Sec. 7216. (1) The following controlled substances are
2 included in schedule 3:
3 (a) Unless listed in another schedule, any material,
4 compound, mixture, or preparation containing any quantity of the
5 following substances having a potential for abuse associated with
6 a stimulant effect on the central nervous system, including their
7 salts, isomers, including optical, position, or geometric
8 isomers, and salts of the isomers if the existence of the salts,
9 isomers, and salts of isomers is possible within the specific

1 chemical designation:

2	Benzphetamine	Mediatric tabs
3	Chlorphentermine	Mediatric liquid
4	Clortermine	Phendimetrazine
5	Edrisal tabs	Special formula 711 tabs
6	Genegesic caps	Thora Dex No. 1 tab
7	Hovizyme tabs	Thora Dex No. 2 tab
8	Mazindol	

9 (b) Unless listed in another schedule, any material,
 10 compound, mixture, or preparation containing any quantity of the
 11 following substances having a potential for abuse associated with
 12 a depressant effect on the central nervous system, including
 13 their salts, isomers, including optical, position, or geometric
 14 isomers, and salts of the isomers if the existence of the salts,
 15 isomers, and salts of isomers is possible within the specific
 16 chemical designation:

17	Chlorhexadol	Phencyclidine
18	Glutethimide	Sulfondiethylmethane
19	Lysergic acid	Sulfonethylmethane
20	Lysergix acid amide	Sulfonmethane
21	Methypylon	

22 (c) Nalorphine.

23 (d) Any substance that contains any quantity of a derivative
 24 of barbituric acid, or any salt of a derivative of barbituric
 25 acid, except those substances that are specifically listed in
 26 other schedules.

(e) A compound, mixture, or preparation containing amobarbital, secobarbital, pentobarbital, or a salt of amobarbital, secobarbital, or pentobarbital, and 1 or more other active medicinal ingredients that are not listed in a schedule.

(f) A suppository dosage form containing amobarbital, secobarbital, pentobarbital, or a salt of amobarbital, secobarbital, or pentobarbital and approved by the food and drug administration for marketing only as a suppository.

(g) Any material, compound, mixture, or preparation containing limited quantities of any of the following narcotic drugs or their salts:

(i) Not more than 1.8 grams of codeine, or any of its salts, per 100 milliliters or not more than 90 milligrams per dosage unit, with an equal or greater quantity of an isoquinoline alkaloid of opium.

(ii) Not more than 1.8 grams of codeine, or any of its salts, per 100 milliliters or not more than 90 milligrams per dosage unit, with 1 or more active nonnarcotic ingredients in recognized therapeutic amounts.

(iii) Not more than 300 milligrams of dihydrocodeinone, or any of its salts, per 100 milliliters or not more than 15 milligrams per dosage unit, with a fourfold or greater quantity of an isoquinoline alkaloid of opium.

(iv) Not more than 300 milligrams of dihydrocodeinone, or any of its salts, per 100 milliliters or not more than 15 milligrams per dosage unit, with 1 or more active nonnarcotic ingredients in recognized therapeutic amounts.

(v) Not more than 1.8 grams of dihydrocodeine, or any of its salts, per 100 milliliters or not more than 90 milligrams per dosage unit, with 1 or more active nonnarcotic ingredients in recognized therapeutic amounts.

(vi) Not more than 300 milligrams of ethylmorphine, or any of its salts, per 100 milliliters or not more than 15 milligrams per dosage unit, with 1 or more ingredients in recognized therapeutic amounts.

(vii) Not more than 500 milligrams of opium per 100 milliliters or per 100 grams, or not more than 25 milligrams per dosage unit, with 1 or more active nonnarcotic ingredients in recognized therapeutic amounts.

(viii) Not more than 50 milligrams of morphine, or any of its salts, per 100 milliliters or per 100 grams, with 1 or more active nonnarcotic ingredients in recognized therapeutic amounts.

(h) Any material, compound, mixture, or preparation containing any quantity of ketamine, a salt of ketamine, an isomer of ketamine, or a salt of an isomer of ketamine.

(I) EXCEPT AS OTHERWISE PROVIDED IN THIS SECTION, ANABOLIC STEROIDS.

(2) The administrator may promulgate rules to except a compound, mixture, or preparation containing any stimulant or depressant substance listed in subsection (1)(a) and (b) from the application of all or any part of this article if the compound, mixture, or preparation contains 1 or more active medicinal ingredients not having a stimulant or depressant effect on the central nervous system and if the admixtures are in combinations,

1 quantity, proportion, or concentration that vitiate the potential
2 for abuse of the substances having a stimulant or depressant
3 effect on the central nervous system.

4 (3) AS USED IN THIS SECTION, "ANABOLIC STEROID" MEANS ANY
5 DRUG OR HORMONAL SUBSTANCE THAT IS CHEMICALLY AND
6 PHARMACOLOGICALLY RELATED TO TESTOSTERONE, OTHER THAN ESTROGENS,
7 PROGESTINS, AND CORTICOSTEROIDS, AND THAT PROMOTES MUSCLE GROWTH,
8 INCLUDING ANY SALT, ESTER, OR ISOMER OF A DRUG OR SUBSTANCE
9 LISTED IN THIS SUBSECTION, IF THAT SALT, ESTER, OR ISOMER
10 PROMOTES MUSCLE GROWTH. ANABOLIC STEROID INCLUDES ANY OF THE
11 FOLLOWING:

12 (A) BOLDENONE.

13 (B) CHLOROTESTOSTERONE (4-CHLOROTESTOSTERONE).

14 (C) CHORIONIC GONADOTROPIN.

15 (D) CLOSTEBOL.

16 (E) DEHYDROCHLORMETHYLTESTOSTERONE.

17 (F) DIHYDROTESTOSTERONE (4-DIHYDROTESTOSTERONE).

18 (G) DROSTANOLONE.

19 (H) ETHYLESTRENOL.

20 (I) FLUOXYMESTERONE.

21 (J) FORMEBOLONE.

22 (K) HUMAN GROWTH HORMONE.

23 (L) MESTEROLONE.

24 (M) METHANDIENONE.

25 (N) METHANDRANONE.

26 (O) METHANDRIOL.

27 (P) METHANDROSTENOLONE.

1 (Q) METHENOLONE.
2 (R) METHYLTESTOSTERONE.
3 (S) MIBOLERONE.
4 (T) NANDROLONE.
5 (U) NORETHANDROLONE.
6 (V) OXANDROLONE.
7 (W) OXYMESTERONE.
8 (X) OXYMETHOLONE.
9 (Y) STANOLONE.
10 (Z) STANOZOLOL.
11 (AA) TESTOLACTONE.
12 (BB) TESTOSTERONE.
13 (CC) TESTOSTERONE CYPIONATE.
14 (DD) TESTOSTERONE ENANTHATE.
15 (EE) TESTOSTERONE PROPIONATE.
16 (FF) TRENBOLONE.
17 (4) THE FOLLOWING ANABOLIC STEROID PRODUCTS ARE EXCEPTED
18 FROM ALL SCHEDULES OF CONTROLLED SUBSTANCES:
19 (A) ANABOLIC STEROIDS THAT ARE EXPRESSLY INTENDED FOR
20 ADMINISTRATION THROUGH IMPLANTS TO CATTLE OR OTHER NONHUMAN
21 SPECIES AND THAT ARE APPROVED BY THE FEDERAL FOOD AND DRUG
22 ADMINISTRATION FOR SUCH USE.
23 (B) ESTERIFIED ESTROGENS 1.25 MILLIGRAMS AND METHYL
24 TESTOSTERONE 2.5 MILLIGRAM TABLETS.
25 (C) ESTERIFIED ESTROGENS 0.625 MILLIGRAMS AND METHYL
26 TESTOSTERONE 1.25 MILLIGRAM TABLETS.
27 (D) CONJUGATED ESTROGENS 1.25 MILLIGRAMS AND METHYL

1 TESTOSTERONE 10 MILLIGRAM TABLETS.

2 (E) CONJUGATED ESTROGENS 0.625 MILLIGRAMS AND METHYL

3 TESTOSTERONE 5 MILLIGRAM TABLETS.

4 (F) TESTOSTERONE ENANTHATE 90 MILLIGRAM/MILLILITER AND

5 ESTRADIOL VALERATE 4 MILLIGRAM/MILLILITER INJECTION.

6 (G) TESTOSTERONE CYPIONATE 50 MILLIGRAM/MILLILITER AND

7 ESTRADIOL CYPIONATE 2 MILLIGRAM/MILLILITER INJECTION.