

VIRGINIA ACTS OF ASSEMBLY — CHAPTER

An Act to amend and reenact §§ 54.1-3454 and 54.1-3456.1 of the Code of Virginia, relating to Drug Control Act; Schedule V; gabapentin.

[H 2557]

Approved

Be it enacted by the General Assembly of Virginia:

1. That §§ 54.1-3454 and 54.1-3456.1 of the Code of Virginia are amended and reenacted as follows:

§ 54.1-3454. Schedule V.

The controlled substances listed in this section are included in Schedule V:

1. Any compound, mixture, or preparation containing limited quantities of any of the following narcotic drugs, which also contains 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:

Not more than 200 milligrams of codeine, or any of its salts, per 100 milliliters or per 100 grams;

Not more than 100 milligrams of dihydrocodeine, or any of its salts, per 100 milliliters or per 100 grams;

Not more than 100 milligrams of ethylmorphine, or any of its salts, per 100 milliliters or per 100 grams;

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

Not more than 100 milligrams of opium per 100 milliliters or per 100 grams;

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

The Board may except by regulation any compound, mixture, or preparation containing any depressant substance listed in subdivision 1 from the application of all or any part of this chapter and such substances so excepted may be dispensed pursuant to § 54.1-3416.

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

Pyrovalerone.

3. Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation that contains any quantity of the following substances having a depressant effect on the central nervous system, including its salts:

Brivaracetam ((2S)-2-[(4R)-2-oxo-4-propylpyrrolidin-1-yl] butanamide) (also referred to as BRV; UCB-34714; Briviact);

Ezogabine [N-[2-amino-4-(4-fluorobenzylamino)-phenyl]-carbamic acid ethyl ester]-2779;

Gabapentin [1-(aminomethyl)cyclohexanecarboxylic acid];

Lacosamide [(R)-2-acetoamido-N-benzyl-3-methoxy-propionamide];

Pregabalin [(S)-3-(aminomethyl)-5-methylhexanoic acid].

§ 54.1-3456.1. Drugs of concern.

A. The Board may promulgate regulations designating specific drugs and substances, including any controlled substance or other drug or substance where there has been or there is the actual or relative potential for abuse, as drugs of concern. Drugs or substances designated as drugs of concern shall be reported to the Department of Health Professions and shall be subject to reporting requirements for the Prescription Monitoring Program established pursuant to Chapter 25.2 (§ 54.1-2519 et seq.).

B. ~~Drugs and substances designated as drugs of concern shall include any material, compound, mixture, or preparation that contains any quantity of the substance tramadol or gabapentin, including its salts. Drugs and substances designated as drugs of concern shall not include any nonnarcotic drug that may be lawfully sold over the counter or behind the counter without a prescription.~~

2. That notwithstanding the provisions of this act or any other provision of law, any wholesale drug distributor licensed and regulated by the Board of Pharmacy and registered with and regulated by the U.S. Drug Enforcement Administration shall have until July 1, 2020, or within 6 months of final approval of compliance from the Board of Pharmacy and the U.S. Drug Enforcement Administration, whichever is earlier, to comply with the storage requirements for Schedule V controlled substances containing gabapentin.