Ohio Administrative Code

Rule 4729:9-1-05 Schedule V controlled substances.

Effective: March 22, 2020

Pursuant to section 3719.41 of the Revised Code,controlled substance schedule V is hereby established, which schedules includethe following, subject to amendment pursuant to section 3719.43 or 3719.44 ofthe Revised Code.

(A) Narcotics-narcotic preparations

Narcotic drugs containing non-narcotic active medicinal ingredients. Any compound, mixture, or preparation that contains any of the following narcotic drugs, or their salts calculated as the free anhydrous base or alkaloid, in limited quantities as set forth below, and that includes 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 narcotic drugs alone:

(1) Not more than two hundred milligrams of codeine per one hundred milliliters or per one hundred grams;

(2) Not more than one hundred milligrams of dihydrocodeine per one hundred milliliters or per one hundred grams;

(3) Not more than one hundred milligrams of ethylmorphine per one hundred milliliters or per one hundred grams;

(4) Not more than 2.5 milligrams of diphenoxylate and not less than twenty-five micrograms of atropine sulfate per dosage unit;

(5) Not more than one hundred milligrams of opium perone hundred milliliters or per one hundred grams;

(6) Not more than 0.5 milligram of difenoxin and not less than twenty-five micrograms of atropine sulfate per dosage unit.

(B) Stimulants

Unless specifically exempted or excluded under federal drug abuse control laws or unless 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 their salts, isomers, and salts of isomers:

(1) Ephedrine, except as provided in division (K) of section 3719.44 of the Revised Code;

(2) Pyrovalerone.

(C) United States food and drug administration approved cannabidiol drugs

Unless specifically exempted or excluded under federal drug abuse control laws or unless listed in another schedule, any drug product in finished dosage formulation that has been approved by the United States food and drug administration that contains cannabidiol (2-[1R-3-methyl-6R-(1-methylethenyl)-2-cyclohexen-1-yl]-5-pentyl-1,3-benzenediol) derived from cannabis and not more than 0.1 per cent (w/w) residual tetrahydrocannabinols.

(D) Depressants

Unless specifically exempted or excluded 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:

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

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

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

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