

APPROVED AND SIGNED BY THE GOVERNOR

Date 4-28-81

Time \_\_\_\_\_

No: 442

**WEST VIRGINIA LEGISLATURE**  
**REGULAR SESSION, 1981**

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**ENROLLED**

**SENATE BILL NO. 442**

(By Mr. Staggers)

—•—

PASSED April 9, 1981

In Effect ninety days from Passage



**ENROLLED**  
**Senate Bill No. 442**

(By MR. STAGGERS)

[Passed April 9, 1981; in effect ninety days from passage.]

AN ACT to amend and reenact section one hundred one, article one, chapter sixty-a of the code of West Virginia, one thousand nine hundred thirty-one, as amended, relating to the uniform controlled substances act and providing for a certain change in language to conform with federal standard.

*Be it enacted by the Legislature of West Virginia:*

That section one hundred one, article one, chapter sixty-a of the code of West Virginia, one thousand nine hundred thirty-one, as amended, be amended and reenacted to read as follows:

**ARTICLE 1. DEFINITIONS.**

**§60A-1-101. Definitions.**

1 As used in this act:

2 (a) "Administer" means the direct application of a  
3 controlled substance whether by injection, inhalation,  
4 ingestion, or any other means, to the body of a patient or  
5 research subject by:

6 (1) A practitioner (or, in his presence, by his authorized  
7 agent), or

8 (2) The patient or research subject at the direction and in  
9 the presence of the practitioner.

10 (b) "Agent" means an authorized person who acts on  
11 behalf of or at the direction of a manufacturer, distributor, or  
12 dispenser. It does not include a common or contract carrier,

13 public warehouseman, or employee of the carrier or  
14 warehouseman.

15 (c) "Bureau" means the "Bureau of Narcotics and  
16 Dangerous Drugs, United States Department of Justice," or  
17 its successor agency.

18 (d) "Controlled substance" means a drug, substance, or  
19 immediate precursor in Schedules I through V of article two.

20 (e) "Counterfeit substance" means a controlled substance  
21 which, or the container or labeling of which, without  
22 authorization, bears the trademark, trade name, or other  
23 identifying mark, imprint, number or device, or any likeness  
24 thereof, of a manufacturer, distributor, or dispenser other  
25 than the person who in fact manufactured, distributed, or  
26 dispensed the substance.

27 (f) "Deliver" or "delivery" means the actual, constructive,  
28 or attempted transfer from one person to another of a  
29 controlled substance, whether or not there is an agency  
30 relationship.

31 (g) "Dispense" means to deliver a controlled substance to  
32 an ultimate user or research subject by or pursuant to the  
33 lawful order of a practitioner, including the prescribing,  
34 administering, packaging, labeling, or compounding  
35 necessary to prepare the substance for that delivery.

36 (h) "Dispenser" means a practitioner who dispenses.

37 (i) "Distribute" means to deliver other than by  
38 administering or dispensing a controlled substance.

39 (j) "Distributor" means a person who distributes.

40 (k) "Drug" means (1) substances recognized as drugs in  
41 the official "United States Pharmacopoeia, official  
42 Homeopathic Pharmacopoeia of the United States, or official  
43 National Formulary," or any supplement to any of them; (2)  
44 substances intended for use in the diagnosis, cure, mitigation,  
45 treatment, or prevention of disease in man or animals; (3)  
46 substances (other than food) intended to affect the structure  
47 or any function of the body of man or animals; and (4)  
48 substances intended for use as a component of any article  
49 specified in clause (1), (2) or (3) of this subdivision. It does not  
50 include devices or their components, parts, or accessories.

51 (l) "Immediate precursor" means a substance which the  
52 "West Virginia Board of Pharmacy" (hereinafter in this act  
53 referred to as the state board of pharmacy) has found to be  
54 and by rule designates as being the principal compound

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

59 (m) "Manufacture" means the production, preparation,  
60 propagation, compounding, conversion, or processing of a  
61 controlled substance, either directly or indirectly or by  
62 extraction from substances of natural origin, or  
63 independently by means of chemical synthesis, or by a  
64 combination of extraction and chemical synthesis, and  
65 includes any packaging or repackaging of the substance or  
66 labeling or relabeling of its container, except that this term  
67 does not include the preparation or compounding of a  
68 controlled substance by an individual for his own use or the  
69 preparation, compounding, packaging, or labeling of a  
70 controlled substance:

71 (1) By a practitioner as an incident to his administering or  
72 dispensing of a controlled substance in the course of his  
73 professional practice, or

74 (2) By a practitioner, or by his authorized agent under his  
75 supervision, for the purpose of, or as an incident to, research,  
76 teaching, or chemical analysis and not for sale.

77 (n) "Marihuana" means all parts of the plant "Cannabis  
78 sativa L.," whether growing or not; the seeds thereof; the  
79 resin extracted from any part of the plant; and every  
80 compound, manufacture, salt, derivative, mixture, or  
81 preparation of the plant, its seeds, or resin. It does not include  
82 the mature stalks of the plant, fiber produced from the stalks,  
83 oil or cake made from the seeds of the plant, any other  
84 compound, manufacture, salt, derivative, mixture, or  
85 preparation of the mature stalks (except the resin extracted  
86 therefrom), fiber, oil, or cake, or the sterilized seed of the  
87 plant which is incapable of germination.

88 (o) "Narcotic drug" means any of the following, whether  
89 produced directly or indirectly by extraction from substances  
90 of vegetable origin, or independently by means of chemical  
91 synthesis, or by a combination of extraction and chemical  
92 synthesis:

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

95 (2) Any salt, compound, isomer, derivative, or preparation  
96 thereof which is chemically equivalent or identical with any

97 of the substances referred to in clause (1) of this subdivision,  
98 but not including the isoquinoline alkaloids of opium.

99 (3) Opium poppy and poppy straw.

100 (4) Coca leaves and any salt, compound, derivative, or  
101 preparation of coca leaves, and any salt, compound, isomer,  
102 derivative, or preparation thereof which is chemically  
103 equivalent or identical with any of these substances, but not  
104 including decocainized coca leaves or extractions of coca  
105 leaves which do not contain cocaine or ecgonine.

106 (p) "Opiate" means any substance having an  
107 addiction-forming or addiction-sustaining liability similar to  
108 morphine or being capable of conversion into a drug having  
109 addiction-forming or addiction-sustaining liability. It does  
110 not include, unless specifically designated as controlled  
111 under section 201, article two of this act, the dextrorotatory  
112 isomer of 3-methoxy-n-methylmorphinan and its salts  
113 (dextromethorphan). It does not include its racemic and  
114 levorotatory forms.

115 (q) "Opium poppy" means the plant of the species  
116 "*Papaver somniferum* L.," except its seeds.

117 (r) "Person" means individual, corporation, government  
118 or governmental subdivision or agency, business trust, estate,  
119 trust, partnership, or association, or any other legal entity.

120 (s) "Poppy straw" means all parts, except the seeds, of the  
121 opium poppy, after mowing.

122 (t) "Practitioner" means:

123 (1) A physician, dentist, veterinarian, scientific  
124 investigator, or other person licensed, registered, or  
125 otherwise permitted to distribute, dispense, conduct research  
126 with respect to, or to administer a controlled substance in the  
127 course of professional practice or research in this state.

128 (2) A pharmacy, hospital, or other institution licensed,  
129 registered, or otherwise permitted to distribute, dispense,  
130 conduct research with respect to, or to administer a  
131 controlled substance in the course of professional practice or  
132 research in this state.

133 (u) "Production" includes the manufacture, planting,  
134 cultivation, growing or harvesting of a controlled substance.

135 (v) "State," when applied to a part of the United States,  
136 includes any state, district, commonwealth, territory, insular  
137 possession thereof, and any area subject to the legal authority  
138 of the United States of America.

139 (w) "Ultimate user" means a person who lawfully  
140 possesses a controlled substance for his own use or for the  
141 use of a member of his household or for administering to an  
142 animal owned by him or by a member of his household.

The Joint Committee on Enrolled Bills hereby certifies that the foregoing bill is correctly enrolled.

*R. P. Bayler*  
Chairman Senate Committee

*Long E. Whitlow*  
Chairman House Committee

Originated in the Senate.

To take effect ninety days from passage.

*Irad C. Wallis*  
Clerk of the Senate

*C. A. Blankenship*  
Clerk of the House of Delegates

*Mont R. Rosen*  
President of the Senate

*James H. Lee, Jr.*  
Speaker House of Delegates

The within *is approved* this the *28*  
day of *April*, 1981.

*Mont R. Rosen*  
Governor



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SECY. OF STATE