GENERAL ASSEMBLY OF NORTH CAROLINA SESSION 2017

H 4

HOUSE BILL 464*

Committee Substitute Favorable 4/5/17 Committee Substitute #2 Favorable 4/19/17 Senate Judiciary Committee Substitute Adopted 6/20/17

vise Schedule of Controlled Substances.	(Public)
March 27, 2017	
FENTANYLS, DESIGNER HALLUCINOGED DIDS, SYSTEM DEPRESSANTS, AND OTHER DNFORMING CHANGES. Embly of North Carolina enacts: ION 1. This act shall be known and may be cited as rous Drug Control Act." ION 2. G.S. 90-87 reads as rewritten: tions.	NICS, SYNTHETIC SUBSTANCES AND
The term "isomer" means, except as used—G.S. 90-89(c), G.S. 90-90(1)d., and G.S. 90-95(h)(3), used in G.S. 90-89(c) the term "isomer" means the geometric isomer. As used in G.S. 90-87(17)(d), G.S. 90-95(h)(3) the term "isomer" means the diastereoisomer.means any type of isomer, including or optical isomers, and stereoisomers.	the optical isomer. As e optical, position, or G.S. 90-90(1)d., and optical isomer or
"Narcotic drug" means any of the following, whether indirectly by extraction from substances of independently by means of chemical synthesis, or extraction and chemical synthesis: a. Opium and opiate, Opium, opiate and opioid, and derivative, or preparation of opium or opioid. b. Any salt, compound, isomer, derivative, or presis chemically equivalent or identical with a referred to in clause a, but not including the isopium. c. Opium poppy and poppy straw.	vegetable origin, or by a combination of and any salt, compound, iate.opium, opiate, or eparation thereof which any of the substances
	March 27, 2017 A BILL TO BE ENTITLED SING THE SCHEDULE OF CONTROLLED SUB FENTANYLS, DESIGNER HALLUCINOGE DIDS, SYSTEM DEPRESSANTS, AND OTHER DNFORMING CHANGES. Embly of North Carolina enacts: TON 1. This act shall be known and may be cited as rous Drug Control Act." TON 2. G.S. 90-87 reads as rewritten: tions. s Article: The term "isomer" means, except as used G.S. 90-89(e), G.S. 90-90(1)d., and G.S. 90-95(h)(3), used in G.S. 90-89(e) the term "isomer" means the geometric isomer. As used in G.S. 90-87(17)(d), G.S. 90-95(h)(3) the term "isomer" means the diastereoisomer. means any type of isomer, including or optical isomers, and stereoisomers. "Narcotic drug" means any of the following, whethe indirectly by extraction from substances of independently by means of chemical synthesis, or extraction and chemical synthesis: a. Opium and opiate, Opium, opiate and opioid, a derivative, or preparation of opium or opopioid. b. Any salt, compound, isomer, derivative, or preparation of identical with a referred to in clause a, but not including the is



1 isomers, compound, derivative or preparation of coca leaves, or any 2 salt, isomer, salts of isomers, compound, derivative, or preparation 3 thereof which is chemically equivalent or identical with any of these 4 substances, except that the substances shall not include decocanized 5 coca leaves or extraction of coca leaves, which extractions do not 6 contain cocaine or ecgonine. 7 (18)"Opiate" substance having an addiction-forming means any 8 addiction-sustaining liability similar to morphine or being capable of 9 conversion into a drug having addiction-forming or addiction-sustaining 10 liability. It does not include, unless specifically designated as controlled 11 under G.S. 90-88. the dextrorotatory isomer 3-methoxy-n-methyl-morphinan and its salts (dextromethorphan). It does 12 13 include its racemic and levorotatory forms. 14 (18a) "Opioid" means any synthetic narcotic drug having opiate-like activities but 15 is not derived from opium. 16 17 **SECTION 3.** G.S. 90-89 reads as rewritten: 18 "§ 90-89. Schedule I controlled substances. 19 This schedule includes the controlled substances listed or to be listed by whatever official 20 name, common or usual name, chemical name, or trade name designated. In determining that a 21 substance comes within this schedule, the Commission shall find: a high potential for abuse, no 22 currently accepted medical use in the United States, or a lack of accepted safety for use in 23 treatment under medical supervision. The following controlled substances are included in this 24 schedule: 25 (1) Opiates. – Any of the following opiates, opiates or opioids, including the isomers, esters, ethers, salts and salts of isomers, esters, and ethers, unless 26 27 specifically excepted, or listed in another schedule, whenever the existence 28 of such isomers, esters, ethers, and salts is possible within the specific 29 chemical designation: 30 Acetyl-alpha-methylfentanyl a. 31 (N[1-(1-methyl-2-phenethyl)-4/y-piperidinyl]-N-phenylacet amide). 32 Acetylmethadol. b. 33 Repealed by Session Laws 1987, c. 412, s. 2. c. 34 d. Alpha-methylthiofentanyl 35 (N-[1-methyl-2-(2-thienyl)ethyl/y-4/y-piperidinyl]-N-phenylpropana 36 mide). 37 e. Allylprodine. 38 f. Alphacetylmethadol. Alphacetylmethadol (except 39 levo-alphacetylmethadol, also known as levomethadyl acetate and 40 LAAM). Alphameprodine. 41 g. 42 Alphamethadol. h. 43 i. Alpha-methylfentanyl (N-(1-(alpha-methyl-beta-phenyl) ethyl-4-piperidyl) 44 propionalilide; 45 1(1-methyl-2-phenyl-ethyl)-4-(N-propanilido) piperidine). Benzethidine. 46 j. 47 Betacetylmethadol. k. 48 l. Beta-hydroxfentanyl 49 (N-[1-(2-hydroxy-2-phenethyl)-4-piperidinyl]-N-phenylpropanamide

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(N-phenyl-N-[1-(2-thienyl)ethyl-4-piperidinyl]-propanamide.

1		<u>h.</u>
2		N-(1-phenethylpiperidin-4-yl)-N-phenyltetrahydrofuran-2-carbox
3		amide (also known as tetrahydrofuran fentanyl).
4		i.
5		N-(4-fluorophenyl)-2-methyl-N-[1-(2-phenylethyl)-4-piperidinyl]
6		-propanamide (also known as 4-fluoroisobutyryl fentanyl, 4-FIBF).
7		j. N-(4-fluorophenyl)-N-[1-(2-phenylethyl)-4-piperidinyl]-butanamide
8		(also known as 4-fluorobutyryl fentanyl, 4-FBF).
9	(2)	Opium derivatives. — Any of the following opium derivatives, including their
10	(-)	salts, isomers, and salts of isomers, unless specifically excepted, or listed in
11		another schedule, whenever the existence of such salts, isomers, and salts of
12		isomers is possible within the specific chemical designation:
13		a. Acetorphine.
14		b. Acetyldihydrocodeine.
15		c. Benzylmorphine.
16		d. Codeine methylbromide.
17		e. Codeine-N-Oxide.
18		f. Cyprenorphine.
19		g. Desomorphine.
20		h. Dihydromorphine.
21		i. Etorphine (except hydrochloride salt).
22		j. Heroin.
23		k. Hydromorphinol.
24		l. Methyldesorphine.
25		m. Methyldihydromorphine.
26		n. Morphine methylbromide.
27		o. Morphine methylsulfonate.
28		p. Morphine-N-Oxide.
29		q. Myrophine.
30		r. Nicocodeine.
31		s. Nicomorphine.
32		t. Normorphine.
33		u. Pholcodine.
34		v. Thebacon.
35 36	(2)	w. Drotebanol.
30 37	(3)	<u>Hallucinogenic substances. – Any material, compound, mixture, or preparation which contains any quantity of the following hallucinogenic</u>
38		substances, including their salts, isomers, and salts of isomers, unless
39		specifically excepted, or listed in another schedule, whenever the existence
40		of such salts, isomers, and salts of isomers is possible within the specific
41		chemical designation:
42		a. 3, 4-methylenedioxyamphetamine.
43		b. 5-methoxy-3, 4-methylenedioxyamphetamine.
44		c. 3, 4-Methylenedioxymethamphetamine (MDMA).
45		d. 3,4-methylenedioxy-N-ethylamphetamine (also known as
46		N-ethyl-alpha-methyl-3,4-(methylenedioxy) phenethylamine, N-ethyl
47		MDA, MDE, and MDEA).
48		e. N-hydroxy-3,4-methylenedioxyamphetamine (also known as
49		N-hydroxy/y-alpha-methyl-3,4-(methylenedioxy) phenethylamine,
50		and N-hydroxy MDA).
51		f. 3, 4, 5-trimethoxyamphetamine.
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- 3-MeO-PCP (3-methoxyphencyclidine). ij.
- kk. 4-hydroxy-MET.
 - ll.4-OH-MiPT (4-hydroxy-N-methyl-N-isopropyltryptamine).
 - 5-methoxy-N-methyl-N-propyltryptamine (5-MeO-MiPT).
- Systemic depressants. Any material compound, mixture, or preparation (4) which contains any quantity of the following substances having a depressant effect on the central nervous system, including its salts, isomers, and salts of isomers whenever the existence of such salts, isomers, and salts of isomers is possible within the specific chemical designation, unless specifically excepted or unless listed in another schedule:
 - Mecloqualone. a.

1			ific chemical designation unless specifically excepted or unless listed in
2		anot	her schedule:
3		a.	25B-NBOMe
4			(2C-B-NBOMe)-2-(4-Bromo-2,5-dimethoxyphenyl)-N-(2-methoxyb
5			enzyl)ethanamine.
6		b.	25C-NBOMe
7			(2C-C-NBOMe)-2-(4-Chloro-2,5-dimethoxyphenyl)-N-(2-methoxyb
8			enzyl)ethanamine.
9		c.	25D-NBOMe
10			(2C-D-NBOMe)-2-(2,5-dimethoxy-4-methylphenyl)-N-(2-methoxyb
11			enzyl)ethanamine.
12		d.	25E-NBOMe
13		a.	(2C-E-NBOMe)-2-(4-Ethyl-2,5-dimethoxyphenyl)-N-(2-methoxyben
14			zyl)ethanamine.
15		e.	25G-NBOMe
16		C.	(2C-G-NBOMe)-2-(2,5-dimethoxy-3,4-dimethylphenyl)-N-(2-metho
10 17			
		f.	xybenzyl)ethanamine.
18		1.	25H-NBOMe
19 20			(2C-H-NBOMe)-2-(2,5-dimethoxyphenyl)-N-(2-methoxybenzyl)etha
20			namine.
21		g.	25I-NBOMe
22			(2C-I-NBOMe)-2-(4-Iodo-2,5-dimethoxyphenyl)-N-(2-methoxybenz
23		_	yl)ethanamine.
24		h.	25N-NBOMe
25			(2C-N-NBOMe)-2-(2,5-dimethoxy-4-nitrophenyl)-N-(2-methoxyben
26			zyl)ethanamine.
27		i.	25P-NBOMe
28			(2C-P-NBOMe)-2-(4-Propyl-2,5-dimethoxyphenyl)-N-(2-methoxybe
29			nzyl)ethanamine.
30		j.	25T2-NBOMe
31			(2C-T2-NBOMe)-2,5-dimethoxy-N-[(2-methoxyphenyl)methyl]-4-(
32			methylthio)-benzeneethanamine.
33		k.	25T4-NBOMe
34			(2C-T4-NBOMe)-2,5-dimethoxy-N-[(2-methoxyphenyl)methyl]-4-[(
35			1-methylethyl)thio]-benzeneethanamine.
36		1.	25T7-NBOMe
37			(2C-T7-NBOMe)-2,5-dimethoxy-N-[(2-methoxyphenyl)methyl]-4-(p
38			ropylthio)-benzeneethanamine.
39	<u>(7)</u>	Synt	hetic cannabinoids. – Any quantity of any synthetic chemical compound
40	-1,,1		(i) is a cannabinoid receptor agonist and mimics the pharmacological
41			et of naturally occurring substances or (ii) has a stimulant, depressant, or
42			icinogenic effect on the central nervous system that is not listed as a
43			rolled substance in Schedules I through V, and is not an FDA-approved
44			s. Synthetic cannabinoids include, but are not limited to, the substances
45			d in sub-subdivisions a. through p. of this subdivision and any substance
46			contains any quantity of their salts, isomers (whether optical, positional,
47			eometric), homologues, and salts of isomers and homologues, unless
48			ifically excepted, whenever the existence of these salts, isomers,
4 9		-	ologues, and salts of isomers and homologues is possible within the
1 9 50			ific chemical designation. The following substances are examples of
<i>J</i> U		spec	the chemical designation. The following substances are examples of

1	synthe	tic cannabinoids and are not intended to be inclusive of the substances
2	-	ed in this Schedule:
3	<u>a.</u>	Naphthoylindoles. Any compound containing a
4	_	3-(1-naphthoyl)indole structure with substitution at the nitrogen atom
5		of the indole ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl,
6		cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or
7		2-(4-morpholinyl)ethyl group, whether or not further substituted in
8		the indole ring to any extent and whether or not substituted in the
9		naphthyl ring to any extent. Some trade or other names: JWH-015,
10		JWH-018, JWH-019, JWH-073, JWH-081, JWH-122, JWH-200,
11		JWH-210, JWH-398, AM-2201, and WIN 55-212.
12	<u>b.</u>	Naphthylmethylindoles. Any compound containing a
13	<u> </u>	1H-indol-3-yl-(1-naphthyl)methane structure with substitution at the
14		nitrogen atom of the indole ring by an alkyl, haloalkyl, alkenyl,
15		cycloalkylmethyl, cycloalkylethyl,
16		1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group,
17		whether or not further substituted in the indole ring to any extent and
18		whether or not substituted in the naphthyl ring to any extent.
19	<u>c.</u>	Naphthoylpyrroles. Any compound containing a
20	<u>c.</u>	3-(1-naphthoyl)pyrrole structure with substitution at the nitrogen
21		atom of the pyrrole ring by an alkyl, haloalkyl, alkenyl,
22		cycloalkylmethyl, cycloalkylethyl,
23		1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group,
24		whether or not further substituted in the pyrrole ring to any extent
25		and whether or not substituted in the naphthyl ring to any extent.
26		Another name: JWH-307.
27	<u>d.</u>	Naphthylmethylindenes. Any compound containing a
28	<u>u.</u>	naphthylideneindene structure with substitution at the 3-position of
29		the indene ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl,
30		cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or
31		2-(4-morpholinyl)ethyl group, whether or not further substituted in
32		the indene ring to any extent and whether or not substituted in the
33		naphthyl ring to any extent.
34	<u>e.</u>	Phenylacetylindoles. Any compound containing a
35	<u>c.</u>	3-phenylacetylindole structure with substitution at the nitrogen atom
36		of the indole ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl,
37		cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or
38		2-(4-morpholinyl)ethyl group, whether or not further substituted in
39		the indole ring to any extent and whether or not substituted in the
40		phenyl ring to any extent. Some trade or other names: SR-18, RCS-8,
41		JWH-250, and JWH-203.
42	<u>f.</u>	Cyclohexylphenols. Any compound containing a
43	1.	2-(3-hydroxycyclohexyl)phenol structure with substitution at the
44		5-position of the phenolic ring by an alkyl, haloalkyl, alkenyl,
45		cycloalkylmethyl, cycloalkylethyl,
46		1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group,
47		whether or not substituted in the cyclohexyl ring to any extent. Some
48		trade or other names: CP 47,497 (and homologues),
49		cannabicyclohexanol.
50	<u>g.</u>	Benzoylindoles. Any compound containing a 3-(benzoyl)indole
51	<u>⊅·</u>	structure with substitution at the nitrogen atom of the indole ring by
		or a contract of at the introgen atom of the indoic fing by

1		an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl,
2		1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group,
3		whether or not further substituted in the indole ring to any extent and
4		whether or not substituted in the phenyl ring to any extent. Some
5		trade or other names: AM-694, Pravadoline (WIN 48,098), and
6		RCS-4.
7	<u>h.</u>	2,3-Dihydro-5-methyl-3-(4-morpholinylmethyl)pyrrolo[1,2,3-de]-1,
8	<u>11.</u>	4-benzoxazin-6-yl]-1-napthalenylmethanone. Some trade or other
9		name: WIN 55,212-2.
10	i	(6aR,10aR)-9-(hydroxymethyl)-6, 6-dimethyl-3-(2-methyloctan-2-yl)
11	<u>i.</u>	- 6a,7,10,10a-tetrahydrobenzo[c]chromen-1-ol 7370. Some trade or
12		other name: HU-210.
13	;	3-(cyclopropylmethanone) indole or 3-(cyclobutylmethanone) indole
14	<u>j.</u>	or 3-(cyclopentylmethanone) indole by substitution at the nitrogen
15		
16		atom of the indole ring, whether or not further substituted in the
		indole ring to any extent, whether or not further substituted on the
17		cyclopropyl, cyclobutyl, or cyclopentyl rings to any extent.
18		Substances in this class include, but are not limited to: UR-144,
19	1_	fluoro-UR-144, XLR-11, A-796,260, and A-834,735.
20	<u>k.</u>	Indole carboxaldehydes. Any compound structurally derived from
21		1H-indole-3-carboxaldehyde or 1H-indole-2-carboxaldehyde
22		substituted in both of the following ways:
23		1. At the nitrogen atom of the indole ring by an alkyl, haloalkyl,
24		cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl,
25		1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl,
26		1-(N-methyl-2-pyrrolidinyl)methyl,
27		1-(N-methyl-3-morpholinyl)methyl,
28		tetrahydropyranylmethyl, benzyl, or halo benzyl group; and
29		2. At the carbon of the carboxaldehyde by a phenyl, benzyl,
30		naphthyl, adamantyl, cyclopropyl, or propionaldehyde group;
31		whether or not the compound is further modified to any extent in the
32		following ways: (i) substitution to the indole ring to any extent, (ii)
33		substitution to the phenyl, benzyl, naphthyl, adamantyl, cyclopropyl,
34		or propionaldehyde group to any extent, (iii) a nitrogen heterocyclic
35		analog of the indole ring, or (iv) anitrogen heterocyclic analog of the
36		phenyl, benzyl, naphthyl, adamantyl, or cyclopropyl ring. Substances
37		in this class include, but are not limited to: AB-001.
38	<u>l.</u>	Indole carboxamides. Any compound structurally derived from
39		1H-indole-3-carboxamide or 1H-indole-2-carboxamide substituted in
40		both of the following ways:
41		1. At the nitrogen atom of the indole ring by an alkyl, haloalkyl,
42		cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl,
43		1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl,
44		1-(N-methyl-2-pyrrolidinyl)methyl,
45		1-(N-methyl-3-morpholinyl)methyl,
46		tetrahydropyranylmethyl, benzyl, or halo benzyl group; and
47		2. At the nitrogen of the carboxamide by a phenyl, benzyl,
48		naphthyl, adamantyl, cyclopropyl, or propionaldehyde group;
49		whether or not the compound is further modified to any extent in the
50		following ways: (i) substitution to the indole ring to any extent, (ii)
51		substitution to the phenyl, benzyl, naphthyl, adamantyl, cyclopropyl,

1		or propionaldehyde group to any extent, (iii) a nitrogen heterocyclic
2		analog of the indole ring, or (iv) a nitrogen heterocyclic analog of the
3		phenyl, benzyl, naphthyl, adamantyl, or cyclopropyl ring. Substances
4		in this class include, but are not limited to: SDB-001 and STS-135.
5	<u>m.</u>	Indole carboxylic acids. Any compound structurally derived from
6		1H-indole-3-carboxylic acid or 1H-indole-2-carboxylic acid
7		substituted in both of the following ways:
8		1. At the nitrogen atom of the indole ring by an alkyl, haloalkyl,
9		cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl,
10		1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl,
11		1-(N-methyl-2-pyrrolidinyl)methyl,
12		1-(N-methyl-3-morpholinyl)methyl,
13		tetrahydropyranylmethyl, benzyl, or halo benzyl group; and
14		<u>At the nitrogen of the carboxamide by a phenyl, benzyl,</u>
15		<u>naphthyl</u> , adamantyl, cyclopropyl, or propionaldehyde group;
16		whether or not the compound is further modified to any
17		extent in the following ways: (i) substitution to the indole ring
18		to any extent, (ii) substitution to the phenyl, benzyl, naphthyl,
19		adamantyl, cyclopropyl, or propionaldehyde group to any
20		extent, (iii) a nitrogen heterocyclic analog of the indole ring,
21		or (iv) a nitrogen heterocyclic analog of the phenyl, benzyl,
22		naphthyl, adamantyl, or cyclopropyl ring. Substances in this
23		class include, but are not limited to: SDB-001 and STS-135.
24		whether or not the compound is further modified to any extent in the
25		following ways: (i) substitution to the indole ring to any extent, (ii)
26		substitution to the phenyl, benzyl, naphthyl, adamantyl, cyclopropyl,
27		or propionaldehyde group to any extent, (iii) a nitrogen heterocyclic
21 22 23 24 25 26 27 28		analog of the indole ring, or (iv) a nitrogen heterocyclic analog of the
29		phenyl, benzyl, naphthyl, adamantyl, or cyclopropyl ring. Substances
30		in this class include, but are not limited to: PB-22 and fluoro-PB-22.
31	<u>n.</u>	Indazole carboxaldehydes. Any compound structurally derived from
32		1H-indazole-3-carboxaldehyde or 1H-indazole-2-carboxaldehyde
33		substituted in both of the following ways:
34		1. At the nitrogen atom of the indazole ring by an alkyl,
35		haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl,
36		cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl,
37		2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl,
38		1-(N-methyl-3-morpholinyl)methyl,
39		tetrahydropyranylmethyl, benzyl, or halo benzyl group; and
40		2. At the carbon of the carboxaldehyde by a phenyl, benzyl,
41		whether or not the compound is further modified to any extent in the
42		following ways: (i) substitution to the indazole ring to any extent, (ii)
43		substitution to the phenyl, benzyl, naphthyl, adamantyl, cyclopropyl,
44		or propionaldehyde group to any extent, (iii) a nitrogen heterocyclic
45		analog of the indazole ring, or (iv) a nitrogen heterocyclic analog of
46		the phenyl, benzyl, naphthyl, adamantyl, or cyclopropyl ring.
4 0 47	0	Indazole carboxamides. Any compound structurally derived from
48	<u>0.</u>	1H-indazole-3-carboxamide or 1H-indazole-2-carboxamide
+6 49		substituted in both of the following ways:
1 9 50		
50 51		1. At the nitrogen atom of the indazole ring by an alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl,
<i>J</i> 1		maidaikyi, cyandaikyi, aikenyi, cycidaikyinnemyi,

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1		cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl,
2		2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl,
3		1-(N-methyl-3-morpholinyl)methyl,
4		tetrahydropyranylmethyl, benzyl, or halo benzyl group; and
5		2. At the nitrogen of the carboxamide by a phenyl, benzyl,
6		naphthyl, adamantyl, cyclopropyl, or propionaldehyde group;
7		whether or not the compound is further modified to any extent in the
8		following ways: (i) substitution to the indazole ring to any extent, (ii)
9		substitution to the phenyl, benzyl, naphthyl, adamantyl, cyclopropyl,
10		or propionaldehyde group to any extent, (iii) a nitrogen heterocyclic
11		analog of the indazole ring, or (iv) a nitrogen heterocyclic analog of
12		the phenyl, benzyl, naphthyl, adamantyl, or cyclopropyl ring.
13		Substances in this class include, but are not limited to: AKB-48,
14		fluoro-AKB-48, APINCACA, AB-PINACA, AB-FUBINACA,
15		ADB-FUBINACA, and ADB-PINACA.
16	<u>p.</u>	Indazole carboxylic acids. Any compound structurally derived from
17		1H-indazole-3-carboxylic acid or 1H-indazole-2-carboxylic acid
18		substituted in both of the following ways:
19		1. At the nitrogen atom of the indazole ring by an alkyl,
20		haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl,
21		cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl,
22		2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl,
23		1-(N-methyl-3-morpholinyl)methyl,
24		tetrahydropyranylmethyl, benzyl, or halo benzyl group; and
25		2. At the hydroxyl group of the carboxylic acid by a phenyl,
26		benzyl, naphthyl, adamantyl, cyclopropyl, or
27		propionaldehyde group; whether or not the compound is
28		further modified to any extent in the following ways: (i)
29		substitution to the indazole ring to any extent, (ii) substitution
30		to the phenyl, benzyl, naphthyl, adamantyl, cyclopropyl, or
31		propionaldehyde group to any extent, (iii) a nitrogen
32		heterocyclic analog of the indazole ring, or (iv) a nitrogen
33		heterocyclic analog of the phenyl, benzyl, naphthyl,
34		adamantyl, or cyclopropyl ring.
35	<u>q.</u>	Carbazoles. Any compound containing a carbazole ring system with
36		a substituent on the nitrogen atom and bearing an additional
37		substituent at the 1, 2, or 3 position of the carbazole ring system, with
38		a linkage connecting the ring system to the substituent:
39		1. Where the linkage connecting the carbazole ring system to
40		the substituent if its 1, 2, or 3 position is any of the following:
41		Alkyl, Carbonyl, Ester, Thione, Thioester, Amino,
42		Alkylamino, Amido, or Alkylamido.
43		2. Where the substituent at the 1, 2, or 3 position of the
44		carbazole ring system, disregarding the linkage, is any of the
45		following groups: Naphthyl, Quinolinyl, Adamantyl, Phenyl,
46		Cycloalkyl (limited to cyclopropyl, cyclobutyl, cyclopentyl,
47		or cyclohexyl), Biphenyl, Alkylamido (limited to ethylamido,
48		propylamido, butanamido, pentamido), Benzyl, Carboxylic
49		acid, Ester, Ether, Phenylpropylamido, or
50		Phenylpropylamino; whether or not further substituted in
51		either of the following ways: (i) the substituent at the 1, 2, or

	General Assembly Of I	ordi Carollia
1		3 position of the carbazole ring system, disregarding the
2		linkage, is further substituted to any extent (ii) further
3		substitution on the carbazole ring system to any extent. This
4		class includes, but is not limited to, the following: MDMB
5		CHMCZCA, EG-018, and EG-2201.
6	<u>r.</u>	Naphthoylnaphthalenes. Any compound structurally derived from
7		naphthalene-1-yl-(naphthalene-1-yl) methanone with substitutions on
8		either of the naphthalene rings to any extent. Substances in this class
9		include, but are not limited to: CB-13."
10	SECTION 4.	G.S. 90-90 reads as rewritten:
11	"§ 90-90. Schedule II co	ontrolled substances.
12	This schedule includ	es the controlled substances listed or to be listed by whatever official
13	name, common or usual	name, chemical name, or trade name designated. In determining that a
14	substance comes within	this schedule, the Commission shall find: a high potential for abuse;
15	currently accepted medi-	cal use in the United States, or currently accepted medical use with
16	severe restrictions; and	the abuse of the substance may lead to severe psychic or physical
17	dependence. The following	ng controlled substances are included in this schedule:
18	(1) Any o	f the following substances whether produced directly or indirectly by
19	extrac	tion from substances of vegetable origin, or independently by means
20	of che	emical synthesis, or by a combination of extraction and chemical
21	synthe	sis, unless specifically excepted or unless listed in another schedule:
22	a.	Opium and Opium, opiate, or opioid and any salt, compound,
23		derivative, or preparation of opium and opiate, excluding
24		apomorphine, nalbuphine, dextrorphan, naloxone, naltrexone and
25		nalmefene, and their respective salts, but including the following:
26		1. Raw opium.
27		2. Opium extracts.
28		3. Opium fluid extracts.
29		4. Powdered opium.
30		5. Granulated opium.
31		6. Tincture of opium.
32		7. Codeine.
33		8. Ethylmorphine.
34		9. Etorphine hydrochloride.
35		10. Hydrocodone. Any material, compound, mixture, or
36		preparation which contains any quantity of hydrocodone.
37		11. Hydromorphone.
38		12. Metopon.
39		13. Morphine.
40		14. Oxycodone.
41		15. Oxymorphone.
42		16. Thebaine.
43 44	h	17. Dihydroetorphine.
44	b.	Any salt, compound, derivative, or preparation thereof which is chemically equivalent or identical with any of the substances referred
46		to in paragraph 1 of this subdivision, except that these substances
47		shall not include the isoquinoline alkaloids of opium.
48	c.	Opium poppy and poppy straw.
49	d.	Cocaine and any salt, isomer, salts of isomers, compound, derivative,
50	u.	or preparation thereof, or coca leaves and any salt, isomer, salts of
51		isomers, compound, derivative, or preparation of coca leaves, or any
$\mathcal{I}_{\mathbf{I}}$		isomors, compound, dorrantes, or proparation of coca leaves, or any

	salt, isomer, salts of isomers, compound, derivative, or preparation
	thereof which is chemically equivalent or identical with any of these
	substances, except that the substances shall not include decocanized
	coca leaves or extraction of coca leaves, which extractions do not
	contain cocaine or ecgonine.
e.	Concentrate of poppy straw (the crude extract of poppy straw in
	either liquid, solid or powder form which contains the phenanthrine
	alkaloids of the opium poppy).
(2) Any	of the following opiates, opiates or opioids, including their isomers,
	rs, ethers, salts, and salts of isomers, whenever the existence of such
	ners, esters, ethers, and salts is possible within the specific chemical
	gnation unless specifically exempted or listed in other schedules:
	Alfentanil.
	Alphaprodine.
	Anileridine.
	Bezitramide.
	Carfentanil.
	Dihydrocodeine.
	Diphenoxylate.
	Fentanyl.
	Isomethadone.
	Levo-alphacetylmethadol. Some trade or other names:
J.	levo-alpha-acetylmethadol, levomethadyl acetate, or LAAM.
1,-	Levomethorphan.
	Levorphanol.
	Metazocine.
	Methadone.
0.	Methadone – Intermediate, 4-cyano-2-dimethylamino-4,
	4/y- diphenyl butane. Moramide – Intermediate, 2-methyl-3-morpholino-1,
р.	, J 1 ,
~	1-diphenyl-propane-carboxylic acid.
	Pethidine.
г.	Pethidine – Intermediate – A,
_	4-cyano-1-methyl-4/y-phenylpiperidine.
S.	Pethidine – Intermediate – B,
	ethyl-4-phenylpiperidine-4-carboxylate.
t.	Pethidine – Intermediate – C,
	1-methyl-4-phenylpiperidine-4-carboxylic acid.
	Phenazocine.
	Piminodine.
	Racemethorphan.
х.	Racemorphan.
у.	Remifentanil.
Z.	Sufentanil.
aa.	Tapentadol.
••••	
SECTION	5. G.S. 90-91 reads as rewritten:
	(2) Any ester isom designal. a. b. c. d. e. f. g. h. i. j. k. l. m. n. o. p. q. r. s. t. u. v. w. x. y. x. y. z. aa"

SECTION 5. G.S. 90-91 reads as rewritten:

"§ 90-91. Schedule III controlled substances.

This schedule includes the controlled substances listed or to be listed by whatever official name, common or usual name, chemical name, or trade name designated. In determining that a substance comes within this schedule, the Commission shall find: a potential for abuse less than

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the substances listed in Schedules I and II; currently accepted medical use in the United States; and abuse may lead to moderate or low physical dependence or high psychological dependence. The following controlled substances are included in this schedule:

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- (d) Any material, compound, mixture, or preparation containing limited quantities of any of the following narcotic drugs, or any salts thereof unless specifically exempted or listed in another schedule:
 - 1. Not more than 1.80 grams of codeine per 100 milliliters or not more than 90 milligrams per dosage unit with an equal or greater quantity of an isoquinoline alkaloid of opium.
 - 2. Not more than 1.80 grams of codeine per 100 milliliters or not more than 90 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts.
 - 3. Not more than 300 milligrams of dihydrocodeinone per 100 milliliters or not more than 15 milligrams per dosage unit with a four fold or greater quantity of an isoquinoline alkaloid of opium.
 - 4. Not more than 300 milligrams of dihydrocodeinone per 100 milliliters or not more than 15 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts.
 - 5. Not more than 1.80 grams of dihydrocodeine per 100 milliliters or not more than 90 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts.
 - 6. Not more than 300 milligrams of ethylmorphine per 100 milliliters or not more than 15 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts.
 - 7. Not more than 500 milligrams of opium per 100 milliliters or per 100 grams, or not more than 25 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts.
 - 8. Not more than 50 milligrams of morphine per 100 milliliters or per 100 grams with one or more active, nonnarcotic ingredients in recognized therapeutic amounts.
 - 9. Buprenorphine.

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- (k) Anabolic steroids. The term "anabolic steroid" means any drug or hormonal substance, chemically and pharmacologically related to testosterone (other than estrogens, progestins, and corticosteroids) that promotes muscle growth, including, but not limited to, the following:
 - 1. Methandrostenolone,
 - 2. Stanozolol,
 - 3. Ethylestrenol,
 - 4. Nandrolone phenpropionate,
 - 5. Nandrolone decanoate,
 - 6. Testosterone propionate,
 - 7. Chorionic gonadotropin,
- 8. Boldenone,
 - 8a. Boldione,
 - 9. Chlorotestosterone (4-chlorotestosterone),
- 48 10. Clostebol,
- 49 11. Dehydrochlormethyltestosterone,

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1	<u>11a.</u>	<u>Desoxymethyltesterone</u>		
2		(17[alpha]-methyl-5[alpha]-androst-2-en-17[beta]-ol) (a	also known	as
3		madol),		
4	12.	Dibydrostestosterone (4-dihydrotestosterone),		
5	13.	Drostanolone,		
6	14.	Fluoxymesterone,		
7	15.	Formebulone (formebolone),		
8	16.	Mesterolene,		
9	17.	Methandienone,		
10	18.	Methandranone,		
11	19.	Methandriol,		
12	<u>19a.</u>	Methasterone,		
13	20.	Methenolene,		
14	21.	Methyltestosterone,		
15	22.	Mibolerone,		
16	23.	Nandrolene,		
17	24.	Norethandrolene,		
18	25.	Oxandrolone,		
19	26.	Oxymesterone,		
20	27.	Oxymetholone,		
21	28.	Stanolone,		
22	29.	Testolactone,		
23	30.	Testosterone,		
24	31.	Trenbolone, and		
25	<u>31a.</u>	19-nor-4,9(10)-androstadienedione (estra-4,9(10)-diene-3,	17-dione), and	
26	32.	Any salt, ester, or isomer of a drug or substance describe	ed or listed in t	this
27		subsection, if that salt, ester, or isomer promotes musc	ele growth. Exc	ept
28		such term does not include (i) an anabolic steroid w	which is expres	ssly
29		intended for administration through implants to cattle o	or other nonhum	nan
30		species and which has been approved by the Secretary of	Health and Hun	nan
31		Services for such administration or (ii) chorionic go	onadotropin wł	hen
32		administered by injection for veterinary use by a licensed	veterinarian or	the
33		veterinarian's designated agent. If any person prescrib	bes, dispenses,	or
34		distributes such steroid for human use, such person shall	ll be considered	l to
35		have prescribed, dispensed, or distributed an anabolic		
36		meaning of this subsection.		

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SECTION 6. G.S. 90-92 reads as rewritten:

"§ 90-92. Schedule IV controlled substances.

- (a) This schedule includes the controlled substances listed or to be listed by whatever official name, common or usual name, chemical name, or trade name designated. In determining that a substance comes within this schedule, the Commission shall find: a low potential for abuse relative to the substances listed in Schedule III of this Article; currently accepted medical use in the United States; and limited physical or pyschological dependence relative to the substances listed in Schedule III of this Article. The following controlled substances are included in this schedule:
 - (1) Depressants. Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation which contains any quantity of the following substances, including its salts, isomers, and salts of isomers whenever the existence of such salts, isomers, and salts of isomers is possible within the specific chemical designation:

Temazepam.

Tetrazepam.

Triazolam.

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(b) A murder other than described in subsection (a) of this section or in G.S. 14-23.2 shall be deemed second degree murder. Any person who commits second degree murder shall be punished as a Class B1 felon, except that a person who commits second degree murder shall be punished as a Class B2 felon in either of the following circumstances:

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(2) The murder is one that was proximately caused by the unlawful distribution of opium or any opium, opiate, or opioid; any synthetic or natural salt, compound, derivative, or preparation of opium, or opiate, or opioid; cocaine or other substance described in G.S. 90-90(1)d., or methamphetamine, G.S. 90-90(1)d.; methamphetamine; or a depressant described in G.S. 90-92(a)(1), and the ingestion of such substance caused the death of the user.

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SECTION 10. This act becomes effective December 1, 2017, and applies to offenses committed on or after that date.