

115TH CONGRESS
2D SESSION

H. R. 4922

To amend the Controlled Substances Act to list fentanyl analogues as schedule I controlled substances.

IN THE HOUSE OF REPRESENTATIVES

FEBRUARY 5, 2018

Mr. SENSENBRENNER introduced the following bill; which was referred to the Committee on Energy and Commerce, and in addition to the Committee on the Judiciary, for a period to be subsequently determined by the Speaker, in each case for consideration of such provisions as fall within the jurisdiction of the committee concerned

A BILL

To amend the Controlled Substances Act to list fentanyl analogues as schedule I controlled substances.

1 *Be it enacted by the Senate and House of Representa-*
2 *tives of the United States of America in Congress assembled,*

3 **SECTION 1. SHORT TITLE.**

4 This Act may be cited as the “Stopping Overdoses
5 of Fentanyl Analogues Act”.

6 **SEC. 2. FENTANYL ANALOGUES.**

7 Section 202(c) of the Controlled Substances Act (21
8 U.S.C. 812) is amended—

1 (a) by adding at the end of subsection (b) of Schedule

2 I the following:

3 “(23) Acetyl fentanyl 4-methylphenethyl.

4 “(24) Acrylfentanyl.

5 “(25) 4-fluorobutyrylfentanyl.

6 “(26) 4-fluoroisobutyryl fentanyl.

7 “(27) 3-furanyl fentanyl.

8 “(28) Isobutyryl fentanyl.

9 “(29) Meta-fluorofentanyl.

10 “(30) Methoxyacetyl fentanyl.

11 “(31) 4-methoxybutyrfentanyl.

12 “(32) Ocfentanil.

13 “(33) Ortho-fluorofentanyl.

14 “(34) Tetrahydrofuranfentanyl.

15 “(35) Valeryl fentanyl.”; and

16 (b) by adding at the end of Schedule I the following:

17 “(e)(1) Unless specifically exempted or unless listed

18 in another schedule, any material, compound, mixture, or

19 preparation which contains any quantity of fentanyl ana-

20 logues, or which contains their salts, isomers, and salts

21 of isomers whenever the existence of such salts, isomers,

22 and salts of isomers is possible within the specific chemical

23 designation.

1 “(2) In paragraph (1), the term ‘fentanyl analogues’
2 includes any compound structurally derived from
3 fentanyl—

4 “(A) by replacement of the phenyl portion of
5 the phenethyl group by any monocycle, whether or
6 not further substituted in or on the monocycle;

7 “(B) by substitution in or on the phenethyl
8 group with alkyl, alkenyl, alkoxy, hydroxy, halo,
9 haloalkyl, amino or nitro groups;

10 “(C) by substitution in or on the piperidine ring
11 with alkyl, alkenyl, alkoxy, ester, ether, hydroxy,
12 halo, haloalkyl, amino or nitro groups;

13 “(D) by replacement of the aniline ring with
14 any aromatic monocycle whether or not further sub-
15 stituted in or on the aromatic monocycle; or

16 “(E) by replacement of the N-propionyl group
17 by another acyl group.”.

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