

115TH CONGRESS
1ST SESSION

S. 1553

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

IN THE SENATE OF THE UNITED STATES

JULY 13, 2017

Mr. JOHNSON introduced the following bill; which was read twice and referred to the Committee on the Judiciary

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—

9 (a) by adding at the end of subsection (b) of Schedule
10 I the following:

1 “(23) Acetyl fentanyl 4-methylphenethyl.

2 “(24) Acrylfentanyl.

3 “(25) 4-fluorobutyrylfentanyl.

4 “(26) 4-fluoroisobutyryl fentanyl.

5 “(27) 3-furanyl fentanyl.

6 “(28) Isobutyryl fentanyl.

7 “(29) Meta-fluorofentanyl.

8 “(30) Methoxyacetyl fentanyl.

9 “(31) 4-methoxybutyrfentanyl.

10 “(32) Ocfentanil.

11 “(33) Ortho-fluorofentanyl.

12 “(34) Tetrahydrofuranyl fentanyl.

13 “(35) Valeryl fentanyl.”; and

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

15 “(e)(1) Unless specifically exempted or unless listed
16 in another schedule, any material, compound, mixture, or
17 preparation which contains any quantity of fentanyl ana-
18 logues, or which contains their salts, isomers, and salts
19 of isomers whenever the existence of such salts, isomers,
20 and salts of isomers is possible within the specific chemical
21 designation.

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

1 “(A) by replacement of the phenyl portion of
2 the phenethyl group by any monocycle, whether or
3 not further substituted in or on the monocycle;

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

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

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

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

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