

116TH CONGRESS  
1ST SESSION

**S.** \_\_\_\_\_

To amend the Controlled Substances Act to more effectively regulate selective androgen receptor modulators, and for other purposes.

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IN THE SENATE OF THE UNITED STATES

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Mr. GRASSLEY (for himself and Mr. WHITEHOUSE) introduced the following bill; which was read twice and referred to the Committee on

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**A BILL**

To amend the Controlled Substances Act to more effectively regulate selective androgen receptor modulators, and for other purposes.

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 “Selective Androgen Re-  
5       ceptor Modulators Control Act of 2019” or the “SARMs  
6       Control Act of 2019”.

1 **SEC. 2. AMENDMENTS TO THE CONTROLLED SUBSTANCES**  
2 **ACT.**

3 (a) DEFINITION.—Section 102 of the Controlled Sub-  
4 stances Act (21 U.S.C. 802) is amended—

5 (1) by redesignating paragraph (58) as para-  
6 graph (59);

7 (2) by redesignating the second paragraph des-  
8 ignated as paragraph (57) as paragraph (58);

9 (3) by moving paragraphs (57), (58) (as so re-  
10 designated), and (59) (as so redesignated) 2 ems to  
11 the left; and

12 (4) by adding at the end the following:

13 “(60)(A) The term ‘SARM’—

14 “(i) means any drug or other substance that is  
15 a selective androgen receptor agonist chemically un-  
16 related to testosterone, estrogens, progestins,  
17 corticosteroids, and dehydroepiandrosterone; and

18 “(ii) includes—

19 “(I) (S)-N-(4-cyano-3-  
20 (trifluoromethyl)phenyl)-3-(4-cyanophenoxy)-2-  
21 hydroxy-2-methylpropanamide (commonly  
22 known as ‘ostarine’ or ‘enobosarm’);

23 “(II) 4-((R)-2-((R)-2,2,2-trifluoro-1-hy-  
24 droxyethyl)pyrrolidin-1-yl)-2-  
25 (trifluoromethyl)benzonitrile (commonly known  
26 as ‘LGD-4033’ or ‘ligandrol’);

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1                   “(III) 9-chloro-2-ethyl-1-methyl-3-(2,2,2-  
2 trifluoroethyl)-3,6-dihydro-7H-pyrrolo[3,2-  
3 f]quinolin-7-one (commonly known as ‘LGD-  
4 3303’);

5                   “(IV) isopropyl (S)-(7-cyano-4-(pyridin-2-  
6 ylmethyl)-1,2,3,4-tetrahydrocyclopenta[b]indol-2  
7 -yl)carbamate (commonly known as  
8 ‘LY2452473’ or ‘TT701’);

9                   “(V) 2-chloro-4-(((1R,2S)-1-(5-(4-  
10 cyanophenyl)-1,3,4-oxadiazol-2-yl)-2-  
11 hydroxypropyl)amino)-3-methylbenzonitrile  
12 (commonly known as ‘RAD-140’);

13                   “(VI) (S)-3-(4-acetamidophenoxy)-2-hy-  
14 droxy-2-methyl-N-(4-nitro-3-  
15 (trifluoromethyl)phenyl)propanamide (com-  
16 monly known as ‘andarine’);

17                   “(VII) 2-chloro-4-((7R,7aS)-7-hydroxy-1,3-  
18 dioxotetrahydro-1H-pyrrolo[1,2-c]imidazol-  
19 2(3H)-yl)-3-methylbenzonitrile (commonly  
20 known as ‘BMS-564929’);

21                   “(VIII) 6-ethyl-4-(trifluoromethyl)-6,7,8,9-  
22 tetrahydropyrido[3,2-g]quinolin-2(1H)-one  
23 (commonly known as ‘LG-121071’);

24                   “(IX) (S)-3-(4-chloro-3-fluorophenoxy)-N-  
25 (4-cyano-3-(trifluoromethyl)phenyl)-2-hydroxy-

1           2-methylpropanamide (commonly known as ‘S-  
2           23’); and

3                   “(X) any salt, ester, ether, or substituted  
4           analogue of a drug or other substance described  
5           in subclauses (I) through (IX).

6           “(B) A substance excluded under subparagraph  
7 (A)(i) may at any time be scheduled by the Attorney Gen-  
8 eral in accordance with the authority and requirements  
9 under subsections (a) through (c) of section 201.

10           “(C)(i) A drug or other substance (other than estro-  
11 gens,           progestins,           corticosteroids,           and  
12 dehydroepiandrosterone, unless scheduled under subpara-  
13 graph (B)) that is not listed in subparagraph (A)(ii) and  
14 is derived from, or has a chemical structure substantially  
15 similar to, 1 or more SARMs listed in subparagraph  
16 (A)(ii) shall be considered to be a SARM for purposes of  
17 this title if the drug or other substance—

18                   “(I) has been created or manufactured with the  
19           intent of producing a drug or other substance that—

20                           “(aa) promotes muscle growth; or

21                           “(bb) otherwise causes a pharmacological  
22           effect similar to that of testosterone; or

23                   “(II) has been, or is intended to be, marketed  
24           or otherwise promoted in any manner suggesting  
25           that consuming the drug or other substance will pro-

1       mote muscle growth or any other pharmacological  
2       effect similar to that of testosterone.

3       “(ii) A drug or other substance shall not be consid-  
4       ered to be a SARM for purposes of this subparagraph if  
5       the drug or other substance—

6               “(I) is—

7                       “(aa) an herb or other botanical;

8                       “(bb) a concentrate, metabolite, or extract  
9                       of, or a constituent isolated directly from, an  
10                      herb or other botanical; or

11                     “(cc) a combination of 2 or more sub-  
12                     stances described in item (aa) or (bb);

13               “(II) is a dietary ingredient for purposes of the  
14       Federal Food, Drug, and Cosmetic Act (21 U.S.C.  
15       301 et seq.); and

16               “(III) is not anabolic or androgenic.

17       “(iii) In accordance with section 515(a), any person  
18       claiming the benefit of an exemption or exception under  
19       clause (ii) shall bear the burden of going forward with the  
20       evidence with respect to that exemption or exception.”.

21       (b) AMENDMENT TO SCHEDULE III.—Schedule III in  
22       section 202(c) of the Controlled Substances Act (21  
23       U.S.C. 812(c)) is amended by adding at the end the fol-  
24       lowing:

25               “(f) SARMS.”.

1       (c) TEMPORARY AND PERMANENT SCHEDULING OF  
2 RECENTLY EMERGED SARMS.—Section 201 of the Con-  
3 trolled Substances Act (21 U.S.C. 811) is amended by  
4 adding at the end the following:

5       “(k) TEMPORARY AND PERMANENT SCHEDULING OF  
6 RECENTLY EMERGED SARMS.—

7           “(1) TEMPORARY ORDERS.—

8           “(A) IN GENERAL.—The Attorney General  
9           may issue a temporary order adding a drug or  
10          other substance to the definition of the term  
11          ‘SARM’ under section 102(60) if the Attorney  
12          General finds that—

13               “(i) the drug or other substance satis-  
14               fies the criteria for being considered a  
15               SARM but is not listed in that section or  
16               by regulation of the Attorney General as  
17               being a SARM; and

18               “(ii) adding the drug or other sub-  
19               stance to the definition of the term SARM  
20               will assist in preventing abuse or misuse of  
21               the drug or other substance.

22           “(B) EFFECTIVE DATE; DURATION.—A  
23          temporary order issued under subparagraph  
24          (A)—

1 “(i) shall take effect not earlier than  
2 30 days after the date of publication by  
3 the Attorney General of a notice in the  
4 Federal Register of—

5 “(I) the intention of the Attorney  
6 General to issue the temporary order;  
7 and

8 “(II) the grounds on which the  
9 temporary order is to be issued; and

10 “(ii) shall expire not later than 2  
11 years after the date on which the tem-  
12 porary order becomes effective, except that  
13 the Attorney General may, during the  
14 pendency of proceedings under paragraph  
15 (2), extend the temporary order for not  
16 more than 6 months.

17 “(C) NOTICE TO SECRETARY.—

18 “(i) IN GENERAL.—The Attorney  
19 General shall transmit notice of a tem-  
20 porary order proposed to be issued under  
21 subparagraph (A) to the Secretary.

22 “(ii) CONSIDERATION.—In issuing a  
23 temporary order under subparagraph (A),  
24 the Attorney General shall take into con-  
25 sideration any comments submitted by the

1 Secretary in response to a notice trans-  
2 mitted under this subparagraph.

3 “(D) EFFECT OF PERMANENT SCHED-  
4 ULING.—A temporary order issued under sub-  
5 paragraph (A) shall be vacated upon the  
6 issuance of a permanent order under paragraph  
7 (2).

8 “(E) JUDICIAL REVIEW.—A temporary  
9 order issued under subparagraph (A) shall not  
10 be subject to judicial review.

11 “(2) PERMANENT ORDERS.—

12 “(A) IN GENERAL.—The Attorney General  
13 may by rule issue a permanent order adding a  
14 drug or other substance to the definition of the  
15 term ‘SARM’ under section 102(60) if the drug  
16 or other substance satisfies the criteria for  
17 being considered a SARM under that section.

18 “(B) TIMING.—The Attorney General may  
19 commence a rulemaking under subparagraph  
20 (A) simultaneously with the issuance of a tem-  
21 porary order under paragraph (1).”.

22 (d) LABELING REQUIREMENTS.—

23 (1) IN GENERAL.—Section 305 of the Con-  
24 trolled Substances Act (21 U.S.C. 825) is amended  
25 by adding at the end the following:



1 “(f) FALSE LABELING OF SARMS.—

2 “(1) PROHIBITION.—It shall be unlawful to im-  
3 port, export, manufacture, distribute, dispense, or  
4 possess with intent to manufacture, distribute, or  
5 dispense, a SARM or product containing a SARM,  
6 unless the SARM or product containing the SARM  
7 bears a label clearly identifying the SARM or prod-  
8 uct containing the SARM by the nomenclature used  
9 by the International Union of Pure and Applied  
10 Chemistry.

11 “(2) EXEMPTION.—

12 “(A) IN GENERAL.—A SARM or product  
13 containing a SARM described in subparagraph  
14 (B) shall be exempt from the International  
15 Union of Pure and Applied Chemistry nomen-  
16 clature requirement under paragraph (1) if the  
17 SARM or product containing a SARM is la-  
18 beled in the manner required under the Federal  
19 Food, Drug, and Cosmetic Act (21 U.S.C. 301  
20 et seq.).

21 “(B) EXEMPT PRODUCTS.—A SARM or  
22 product containing a SARM is described in this  
23 subparagraph if the SARM or product con-  
24 taining a SARM—

1 “(i) is the subject of an approved ap-  
2 plication as described in subsection (b) or  
3 (j) of section 505 of the Federal Food,  
4 Drug, and Cosmetic Act (21 U.S.C. 355);  
5 or

6 “(ii) is exempt from the provisions of  
7 section 505 of the Federal Food, Drug,  
8 and Cosmetic Act (21 U.S.C. 355) relating  
9 to new drugs because—

10 “(I) the SARM or product con-  
11 taining a SARM is intended solely for  
12 investigational use as described in  
13 subsection (i) of that section; and

14 “(II) the SARM or product con-  
15 taining a SARM is being used exclu-  
16 sively for purposes of a clinical trial  
17 that is the subject of an effective in-  
18 vestigational new drug application.”.

19 (2) CLARIFICATION REGARDING FELONY DRUG  
20 OFFENSES.—Section 102(44) of the Controlled Sub-  
21 stances Act (21 U.S.C. 802(44)) is amended by in-  
22 serting “SARMs,” after “anabolic steroids,”.

23 (3) CIVIL PENALTIES.—Section 402 of the Con-  
24 trolled Substances Act (21 U.S.C. 842) is amend-  
25 ed—

1 (A) in subsection (a)(16)—

2 (i) by inserting “or (f)” after “sub-  
3 section (e)”; and

4 (ii) by striking “825” and inserting  
5 “305”; and

6 (B) in subsection (c)(1)(D), by inserting  
7 “or a SARM” after “an anabolic steroid”.

8 **SEC. 3. AMENDMENTS TO THE FEDERAL FOOD, DRUG, AND**  
9 **COSMETIC ACT.**

10 Section 413(c) of the Federal Food, Drug, and Cos-  
11 metic Act (21 U.S.C. 350b(c)) is amended—

12 (1) in paragraph (1), by striking “an anabolic  
13 steroid or an analogue of an anabolic steroid” and  
14 inserting “an anabolic steroid, a SARM, an analogue  
15 of an anabolic steroid, or an analogue of a SARM”;  
16 and

17 (2) in paragraph (2)—

18 (A) in subparagraph (A), by striking  
19 “and” at the end;

20 (B) in subparagraph (B), by striking the  
21 period at the end and inserting a semicolon;  
22 and

23 (C) by adding at the end the following:

24 “(C) the term ‘analogue of a SARM’  
25 means a substance that has a chemical struc-

1           ture that is substantially similar to the chemical  
2           structure of a SARM; and  
3           “(D) the term ‘SARM’ has the meaning  
4           given the term in section 102(60) of the Con-  
5           trolled Substances Act (21 U.S.C. 802(60)).”.