

Suspend the Rules and Pass the Bill, H.R. 4981, with An Amendment**(The amendment strikes all after the enacting clause and inserts a new text)**114TH CONGRESS
2^D SESSION**H. R. 4981**

To amend the Controlled Substances Act to improve access to opioid use disorder treatment.

IN THE HOUSE OF REPRESENTATIVES

APRIL 18, 2016

Mr. BUCSHON (for himself and Mr. TONKO) 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 improve access to opioid use disorder treatment.

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 “Opioid Use Disorder
5 Treatment Expansion and Modernization Act”.

1 **SEC. 2. FINDING.**

2 The Congress finds that opioid use disorder has be-
3 come a public health epidemic that must be addressed by
4 increasing awareness and access to all treatment options
5 for opioid use disorder, overdose reversal, and relapse pre-
6 vention.

7 **SEC. 3. OPIOID USE DISORDER TREATMENT MODERNIZA-**
8 **TION.**

9 (a) IN GENERAL.—Section 303(g)(2) of the Con-
10 trolled Substances Act (21 U.S.C. 823(g)(2)) is amend-
11 ed—

12 (1) in subparagraph (B), by striking clauses (i),
13 (ii), and (iii) and inserting the following:

14 “(i) The practitioner is a qualifying practitioner
15 (as defined in subparagraph (G)).

16 “(ii) With respect to patients to whom the prac-
17 titioner will provide such drugs or combinations of
18 drugs, the practitioner has the capacity to provide
19 directly, by referral, or in such other manner as de-
20 termined by the Secretary—

21 “(I) all schedule III, IV, and V drugs, as
22 well as unscheduled medications approved by
23 the Food and Drug Administration, for the
24 treatment of opioid use disorder, including such
25 drugs and medications for maintenance, detoxi-

1 fication, overdose reversal, and relapse preven-
2 tion, as available; and

3 “(II) appropriate counseling and other ap-
4 propriate ancillary services.

5 “(iii)(I) The total number of such patients of
6 the practitioner at any one time will not exceed the
7 applicable number. Except as provided in subclause
8 (II), the applicable number is 30.

9 “(II) The applicable number is 100 if, not soon-
10 er than 1 year after the date on which the practi-
11 tioner submitted the initial notification, the practi-
12 tioner submits a second notification to the Secretary
13 of the need and intent of the practitioner to treat up
14 to 100 patients.

15 “(III) The Secretary may by regulation change
16 such total number.

17 “(IV) The Secretary may exclude from the ap-
18 plicable number patients to whom such drugs or
19 combinations of drugs are directly administered by
20 the qualifying practitioner in the office setting.

21 “(iv) If the Secretary by regulation increases
22 the total number of patients which a qualifying prac-
23 titioner is permitted to treat pursuant to clause
24 (iii)(II), the Secretary shall require such a practi-
25 tioner to obtain a written agreement from each pa-

1 tient, including the patient’s signature, that the pa-
2 tient—

3 “(I) will receive an initial assessment and
4 treatment plan and periodic assessments and
5 treatment plans thereafter;

6 “(II) will be subject to medication adher-
7 ence and substance use monitoring;

8 “(III) understands available treatment op-
9 tions, including all drugs approved by the Food
10 and Drug Administration for the treatment of
11 opioid use disorder, including their potential
12 risks and benefits; and

13 “(IV) understands that receiving regular
14 counseling services is critical to recovery.

15 “(v) The practitioner will comply with the re-
16 porting requirements of subparagraph (D)(i)(IV).”;

17 (2) in subparagraph (D)—

18 (A) in clause (i), by adding at the end the
19 following:

20 “(IV) The practitioner reports to the Secretary,
21 at such times and in such manner as specified by
22 the Secretary, such information and assurances as
23 the Secretary determines necessary to assess wheth-
24 er the practitioner continues to meet the require-
25 ments for a waiver under this paragraph.”;

1 (B) in clause (ii), by striking “Upon re-
2 ceiving a notification under subparagraph (B)”
3 and inserting “Upon receiving a determination
4 from the Secretary under clause (iii) finding
5 that a practitioner meets all requirements for a
6 waiver under subparagraph (B)”;

7 (C) in clause (iii)—

8 (i) by inserting “and shall forward
9 such determination to the Attorney Gen-
10 eral” before the period at the end of the
11 first sentence; and

12 (ii) by striking “physician” and in-
13 serting “practitioner”;

14 (3) in subparagraph (G)—

15 (A) by amending clause (ii)(IV) to read as
16 follows:

17 “(IV) The physician has, with respect to
18 the treatment and management of opiate-de-
19 pendent patients, completed not less than eight
20 hours of training (through classroom situations,
21 seminars at professional society meetings, elec-
22 tronic communications, or otherwise) that is
23 provided by the American Society of Addiction
24 Medicine, the American Academy of Addiction
25 Psychiatry, the American Medical Association,

1 the American Osteopathic Association, the
2 American Psychiatric Association, or any other
3 organization that the Secretary determines is
4 appropriate for purposes of this subclause. Such
5 training shall address—

6 “(aa) opioid maintenance and detoxi-
7 fication;

8 “(bb) appropriate clinical use of all
9 drugs approved by the Food and Drug Ad-
10 ministration for the treatment of opioid
11 use disorder;

12 “(cc) initial and periodic patient as-
13 sessments (including substance use moni-
14 toring);

15 “(dd) individualized treatment plan-
16 ning; overdose reversal; relapse prevention;

17 “(ee) counseling and recovery support
18 services;

19 “(ff) staffing roles and considerations;

20 “(gg) diversion control; and

21 “(hh) other best practices, as identi-
22 fied by the Secretary.”; and

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

24 “(iii) The term ‘qualifying practitioner’
25 means—

1 “(I) a qualifying physician, as defined in
2 clause (ii); or

3 “(II) during the period beginning on the
4 date of the enactment of the Opioid Use Dis-
5 order Treatment Expansion and Modernization
6 Act and ending on the date that is three years
7 after such date of enactment, a qualifying other
8 practitioner, as defined in clause (iv).

9 “(iv) The term ‘qualifying other practitioner’
10 means a nurse practitioner or physician assistant
11 who satisfies each of the following:

12 “(I) The nurse practitioner or physician
13 assistant is licensed under State law to pre-
14 scribe schedule III, IV, or V medications for the
15 treatment of pain.

16 “(II) The nurse practitioner or physician
17 assistant satisfies 1 or more of the following:

18 “(aa) Has completed not fewer than
19 24 hours of initial training addressing each
20 of the topics listed in clause (ii)(IV)
21 (through classroom situations, seminars at
22 professional society meetings, electronic
23 communications, or otherwise) provided by
24 the American Society of Addiction Medi-
25 cine, the American Academy of Addiction

1 Psychiatry, the American Medical Associa-
2 tion, the American Osteopathic Associa-
3 tion, the American Nurses Credentialing
4 Center, the American Psychiatric Associa-
5 tion, the American Association of Nurse
6 Practitioners, the American Academy of
7 Physician Assistants, or any other organi-
8 zation that the Secretary determines is ap-
9 propriate for purposes of this subclause.

10 “(bb) Has such other training or ex-
11 perience as the Secretary determines will
12 demonstrate the ability of the nurse practi-
13 tioner or physician assistant to treat and
14 manage opiate-dependent patients.

15 “(III) The nurse practitioner or physician
16 assistant is supervised by or works in collabora-
17 tion with a qualifying physician, if the nurse
18 practitioner or physician assistant is required
19 by State law to prescribe medications for the
20 treatment of opioid use disorder in collaboration
21 with or under the supervision of a physician.

22 The Secretary may review and update the require-
23 ments for being a qualifying other practitioner under
24 this clause.”; and

25 (4) in subparagraph (H)—

1 (A) in clause (i), by inserting after sub-
2 clause (II) the following:

3 “(III) Such other elements of the requirements
4 under this paragraph as the Secretary determines
5 necessary for purposes of implementing such re-
6 quirements.”; and

7 (B) by amending clause (ii) to read as fol-
8 lows:

9 “(ii) Not later than one year after the date of enact-
10 ment of the Opioid Use Disorder Treatment Expansion
11 and Modernization Act, the Secretary shall update the
12 treatment improvement protocol containing best practice
13 guidelines for the treatment of opioid-dependent patients
14 in office-based settings. The Secretary shall update such
15 protocol in consultation with experts in opioid use disorder
16 research and treatment.”.

17 (b) RECOMMENDATION OF REVOCATION OR SUSPEN-
18 SION OF REGISTRATION IN CASE OF SUBSTANTIAL NON-
19 COMPLIANCE.—The Secretary of Health and Human
20 Services may recommend to the Attorney General that the
21 registration of a practitioner be revoked or suspended if
22 the Secretary determines, according to such criteria as the
23 Secretary establishes by regulation, that a practitioner
24 who is registered under section 303(g)(2) of the Controlled
25 Substances Act (21 U.S.C. 823(g)(2)) is not in substantial

1 compliance with the requirements of such section, as
2 amended by this Act.

3 (c) OPIOID DEFINED.—Section 102(18) of the Con-
4 trolled Substances Act (21 U.S.C. 802(18)) is amended
5 by inserting “or ‘opioid’” after “The term ‘opiate’”.

6 (d) REPORTS TO CONGRESS.—

7 (1) IN GENERAL.—Not later than 2 years after
8 the date of enactment of this Act and not less than
9 over every 5 years thereafter, the Secretary of
10 Health and Human Services, in consultation with
11 the Drug Enforcement Administration and experts
12 in opioid use disorder research and treatment,
13 shall—

14 (A) perform a thorough review of the pro-
15 vision of opioid use disorder treatment services
16 in the United States, including services pro-
17 vided in opioid treatment programs and other
18 specialty and nonspecialty settings; and

19 (B) submit a report to the Congress on the
20 findings and conclusions of such review.

21 (2) CONTENTS.—Each report under paragraph
22 (1) shall include an assessment of—

23 (A) compliance with the requirements of
24 section 303(g)(2) of the Controlled Substances

1 Act (21 U.S.C. 823(g)(2)), as amended by this
2 Act;

3 (B) the measures taken by the Secretary of
4 Health and Human Services to ensure such
5 compliance;

6 (C) whether there is further need to in-
7 crease or decrease the number of patients a
8 waived practitioner is permitted to treat, as
9 provided for by the amendment made by sub-
10 section (a)(1);

11 (D) the extent to which, and proportions
12 with which, the full range of Food and Drug
13 Administration-approved treatments for opioid
14 use disorder are used in routine health care set-
15 tings and specialty substance use disorder treat-
16 ment settings;

17 (E) access to, and use of, counseling and
18 recovery support services, including the percent-
19 age of patients receiving such services;

20 (F) changes in State or local policies and
21 legislation relating to opioid use disorder treat-
22 ment;

23 (G) the use of prescription drug moni-
24 toring programs by practitioners who are per-
25 mitted to dispense narcotic drugs to individuals

1 pursuant to a waiver under section 303(g)(2) of
2 the Controlled Substances Act (21 U.S.C.
3 823(g)(2));

4 (H) the findings resulting from inspections
5 by the Drug Enforcement Administration of
6 practitioners described in subparagraph (G);
7 and

8 (I) the effectiveness of cross-agency col-
9 laboration between Department of Health and
10 Human Services and the Drug Enforcement
11 Administration for expanding effective opioid
12 use disorder treatment.

13 **SEC. 4. SENSE OF CONGRESS.**

14 It is the Sense of Congress that, with respect to the
15 total number of patients that a qualifying physician (as
16 defined in subparagraph (G)(iii) of section 303(g)(2) of
17 the Controlled Substances Act (21 U.S.C. 823(g)(2)) can
18 treat at any one time pursuant to such section, the Sec-
19 retary of Health and Human Services should consider
20 raising such total number to 250 patients following a third
21 notification to the Secretary of the need and intent of the
22 physician to treat up to 250 patients that is submitted
23 to the Secretary not sooner than 1 year after the date
24 on which the physician submitted to the Secretary a sec-
25 ond notification to treat up to 100 patients.

1 **SEC. 5. PARTIAL FILLS OF SCHEDULE II CONTROLLED SUB-**
2 **STANCES.**

3 (a) IN GENERAL.—Section 309 of the Controlled
4 Substances Act (21 U.S.C. 829) is amended by adding at
5 the end the following:

6 “(f) PARTIAL FILLS OF SCHEDULE II CONTROLLED
7 SUBSTANCES.—

8 “(1) PARTIAL FILLS.—

9 “(A) IN GENERAL.—A prescription for a
10 controlled substance in schedule II may be par-
11 tially filled if—

12 “(i) it is not prohibited by State law;

13 “(ii) the prescription is written and
14 filled in accordance with the Controlled
15 Substances Act (21 U.S.C. 801 et seq.),
16 regulations prescribed by the Attorney
17 General, and State law;

18 “(iii) the partial fill is requested by
19 the patient or the practitioner that wrote
20 the prescription; and

21 “(iv) the total quantity dispensed in
22 all partial fillings does not exceed the total
23 quantity prescribed.

24 “(B) OTHER CIRCUMSTANCES.—A pre-
25 scription for a controlled substance in schedule
26 II may be partially filled in accordance with

1 section 1306.13 of title 21, Code of Federal
2 Regulations (as in effect on the date of enact-
3 ment of the Reducing Unused Medications Act
4 of 2016).

5 “(2) REMAINING PORTIONS.—

6 “(A) IN GENERAL.—Except as provided in
7 subparagraph (B), remaining portions of a par-
8 tially filled prescription for a controlled sub-
9 stance in schedule II—

10 “(i) may be filled; and

11 “(ii) shall be filled not later than 30
12 days after the date on which the prescrip-
13 tion is written.

14 “(B) EMERGENCY SITUATIONS.—In emer-
15 gency situations, as described in subsection (a),
16 the remaining portions of a partially filled pre-
17 scription for a controlled substance in schedule
18 II—

19 “(i) may be filled; and

20 “(ii) shall be filled not later than 72
21 hours after the prescription is issued.”.

22 (b) RULE OF CONSTRUCTION.—Nothing in this sec-
23 tion shall be construed to affect the authority of the Attor-
24 ney General to allow a prescription for a controlled sub-
25 stance in schedule III, IV, or V of section 202(c) of the

1 Controlled Substances Act (21 U.S.C. 812(c)) to be par-
2 tially filled.