

AMENDMENT

OFFERED BY M .

Strike section 202 (and update the table of contents accordingly).

Strike section 301 and insert the following new section (and update the table of contents accordingly):

1 SEC. 301. MEDICARE PART D BENEFIT REDESIGN.

2 (a) BENEFIT STRUCTURE REDESIGN.—Section
3 1860D–2(b) of the Social Security Act (42 U.S.C. 1395w–
4 102(b)) is amended—

5 (1) in paragraph (2)—

6 (A) in subparagraph (A)—

7 (i) in the matter preceding clause (i),
8 by inserting “for a year preceding 2022
9 and for costs above the annual deductible
10 specified in paragraph (1) and up to the
11 annual out-of-pocket threshold specified in
12 paragraph (4)(B) for 2022 and each subse-
13 quent year” after “paragraph (3)”; and

14 (ii) in clause (i), by inserting after
15 “25 percent” the following: “(or, for 2022
16 and each subsequent year, 15 percent)”;

1 (B) in subparagraph (C)—

2 (i) in clause (i), in the matter pre-
3 ceding subclause (I), by inserting “for a
4 year preceding 2022,” after “paragraph
5 (4),”; and

6 (ii) in clause (ii)(III), by striking
7 “and each subsequent year” and inserting
8 “and 2021”; and

9 (C) in subparagraph (D)—

10 (i) in clause (i)—

11 (I) in the matter preceding sub-
12 clause (I), by inserting “for a year
13 preceding 2022,” after “paragraph
14 (4),”; and

15 (II) in subclause (I)(bb), by
16 striking “a year after 2018” and in-
17 serting “each of years 2018 through
18 2021”; and

19 (ii) in clause (ii)(V), by striking
20 “2019 and each subsequent year” and in-
21 serting “each of years 2019 through
22 2021”;

23 (2) in paragraph (3)(A)—

1 (A) in the matter preceding clause (i), by
2 inserting “for a year preceding 2022,” after
3 “and (4),”; and

4 (B) in clause (ii), by striking “for a subse-
5 quent year” and inserting “for each of years
6 2007 through 2021”;

7 (3) in paragraph (4)—

8 (A) in subparagraph (A)—

9 (i) in clause (i)—

10 (I) by redesignating subclauses
11 (I) and (II) as items (aa) and (bb),
12 respectively, and indenting appro-
13 priately;

14 (II) in the matter preceding item
15 (aa), as redesignated by subclause (I),
16 by striking “is equal to the greater
17 of—” and inserting “is equal to—

18 “(I) for a year preceding 2022,
19 the greater of—”.

20 (III) by striking the period at the
21 end of item (bb), as redesignated by
22 subclause (I), and inserting “; and”;
23 and

24 (IV) by adding at the end the fol-
25 lowing:

1 “(II) for 2022 and each suc-
2 ceeding year, \$0.”; and

3 (ii) in clause (ii)—

4 (I) by striking “clause (i)(I)” and
5 inserting “clause (i)(I)(aa)”;

6 (II) by adding at the end the fol-
7 lowing new sentence: “The Secretary
8 shall continue to calculate the dollar
9 amounts specified in clause (i)(I)(aa),
10 including with the adjustment under
11 this clause, after 2021 for purposes of
12 section 1860D–14(a)(1)(D)(iii).”;

13 (B) in subparagraph (B)—

14 (i) in clause (i)—

15 (I) in subclause (V), by striking
16 “or” at the end;

17 (II) in subclause (VI)—

18 (aa) by striking “for a sub-
19 sequent year” and inserting “for
20 2021”; and

21 (bb) by striking the period
22 at the end and inserting a semi-
23 colon; and

24 (III) by adding at the end the
25 following new subclauses:

1 “(VII) for 2022, is equal to
2 \$3,100; or

3 “(VIII) for a subsequent year, is
4 equal to the amount specified in this
5 subparagraph for the previous year,
6 increased by the annual percentage in-
7 crease described in paragraph (6) for
8 the year involved.”; and

9 (ii) in clause (ii), by striking “clause
10 (i)(II)” and inserting “clause (i)”;

11 (C) in subparagraph (C)(i), by striking
12 “and for amounts” and inserting “and for a
13 year preceding 2022 for amounts”; and

14 (D) in subparagraph (E), by striking “In
15 applying” and inserting “For each of 2011
16 through 2021, in applying”.

17 (b) DECREASING REINSURANCE PAYMENT
18 AMOUNT.—Section 1860D–15(b)(1) of the Social Security
19 Act (42 U.S.C. 1395w–115(b)(1)) is amended—

20 (1) by striking “equal to 80 percent” and in-
21 serting “equal to—

22 “(A) for a year preceding 2022, 80 per-
23 cent”;

1 (2) in subparagraph (A), as added by para-
2 graph (1), by striking the period at the end and in-
3 serting “; and”; and

4 (3) by adding at the end the following new sub-
5 paragraph:

6 “(B) for 2022 and each subsequent year,
7 the sum of—

8 “(i) an amount equal to 20 percent of
9 the allowable reinsurance costs (as speci-
10 fied in paragraph (2)) attributable to that
11 portion of gross covered prescription drug
12 costs as specified in paragraph (3) in-
13 curred in the coverage year after such indi-
14 vidual has incurred costs that exceed the
15 annual out-of-pocket threshold specified in
16 section 1860D–2(b)(4)(B) with respect to
17 applicable drugs (as defined in section
18 1860D–14B(g)(2)); and

19 “(ii) an amount equal to 30 percent of
20 the allowable reinsurance costs (as speci-
21 fied in paragraph (2)) attributable to that
22 portion of gross covered prescription drug
23 costs as specified in paragraph (3) in-
24 curred in the coverage year after such indi-
25 vidual has incurred costs that exceed the

1 annual out-of-pocket threshold specified in
2 section 1860D–2(b)(4)(B) with respect to
3 generic drugs (as defined in section
4 1860D–14B(g)(5)).”.

5 (c) MANUFACTURER DISCOUNT PROGRAM.—

6 (1) IN GENERAL.—Part D of title XVIII of the
7 Social Security Act is amended by inserting after
8 section 1860D–14A (42 U.S.C. 1495w–114) the fol-
9 lowing new section:

10 **“SEC. 1860D–14B. MANUFACTURER DISCOUNT PROGRAM.**

11 “(a) ESTABLISHMENT.—The Secretary shall estab-
12 lish a manufacturer discount program (in this section re-
13 ferred to as the ‘program’). Under the program, the Sec-
14 retary shall enter into agreements described in subsection
15 (b) with manufacturers and provide for the performance
16 of the duties described in subsection (c). The Secretary
17 shall establish a model agreement for use under the pro-
18 gram by not later than January 1, 2021, in consultation
19 with manufacturers, and allow for comment on such model
20 agreement.

21 “(b) TERMS OF AGREEMENT.—

22 “(1) IN GENERAL.—

23 “(A) AGREEMENT.—An agreement under
24 this section shall require the manufacturer to
25 provide applicable beneficiaries access to—

1 “(i) discounted prices for applicable
2 drugs of the manufacturer that are dis-
3 pensed on or after January 1, 2022; and

4 “(ii) discounted prices for generic
5 drugs of the manufacturer that are dis-
6 pensed on or after January 1, 2022.

7 “(B) PROVISION OF DISCOUNTED PRICES
8 AT THE POINT-OF-SALE.—The discounted prices
9 described in subparagraph (A) shall be provided
10 to the applicable beneficiary at the pharmacy or
11 by the mail order service at the point-of-sale of
12 an applicable drug or a generic drug, as the
13 case may be.

14 “(2) PROVISION OF APPROPRIATE DATA.—Each
15 manufacturer with an agreement in effect under this
16 section shall collect and have available appropriate
17 data, as determined by the Secretary, to ensure that
18 it can demonstrate to the Secretary compliance with
19 the requirements under the program.

20 “(3) COMPLIANCE WITH REQUIREMENTS FOR
21 ADMINISTRATION OF PROGRAM.—Each manufac-
22 turer with an agreement in effect under this section
23 shall comply with requirements imposed by the Sec-
24 retary or a third party with a contract under sub-
25 section (d)(3), as applicable, for purposes of admin-

1 istering the program, including any determination
2 under subparagraph (A) of subsection (c)(1) or pro-
3 cedures established under such subsection (c)(1).

4 “(4) LENGTH OF AGREEMENT.—

5 “(A) IN GENERAL.—An agreement under
6 this section shall be effective for an initial pe-
7 riod of not less than 12 months and shall be
8 automatically renewed for a period of not less
9 than 1 year unless terminated under subpara-
10 graph (B).

11 “(B) TERMINATION.—

12 “(i) BY THE SECRETARY.—The Sec-
13 retary may provide for termination of an
14 agreement under this section for a knowing
15 and willful violation of the requirements of
16 the agreement or other good cause shown.
17 Such termination shall not be effective ear-
18 lier than 30 days after the date of notice
19 to the manufacturer of such termination.
20 The Secretary shall provide, upon request,
21 a manufacturer with a hearing concerning
22 such a termination, and such hearing shall
23 take place prior to the effective date of the
24 termination with sufficient time for such

1 effective date to be repealed if the Sec-
2 retary determines appropriate.

3 “(ii) BY A MANUFACTURER.—A man-
4 ufacturer may terminate an agreement
5 under this section for any reason. Any
6 such termination shall be effective, with re-
7 spect to a plan year—

8 “(I) if the termination occurs be-
9 fore January 30 of a plan year, as of
10 the day after the end of the plan year;
11 and

12 “(II) if the termination occurs on
13 or after January 30 of a plan year, as
14 of the day after the end of the suc-
15 ceeding plan year.

16 “(iii) EFFECTIVENESS OF TERMI-
17 NATION.—Any termination under this sub-
18 paragraph shall not affect discounts for
19 applicable drugs of the manufacturer that
20 are due under the agreement before the ef-
21 fective date of its termination.

22 “(iv) NOTICE TO THIRD PARTY.—The
23 Secretary shall provide notice of such ter-
24 mination to a third party with a contract
25 under subsection (d)(3) within not less

1 than 30 days before the effective date of
2 such termination.

3 “(5) EFFECTIVE DATE OF AGREEMENT.—An
4 agreement under this section shall take effect on a
5 date determined appropriate by the Secretary, which
6 may be at the start of a calendar quarter.

7 “(c) DUTIES DESCRIBED.—The duties described in
8 this subsection are the following:

9 “(1) ADMINISTRATION OF PROGRAM.—Admin-
10 istering the program, including—

11 “(A) the determination of the amount of
12 the discounted price of an applicable drug of a
13 manufacturer and of the discounted price of a
14 generic drug of a manufacturer;

15 “(B) the establishment of procedures
16 under which discounted prices are provided to
17 applicable beneficiaries at pharmacies or by
18 mail order service at the point-of-sale of an ap-
19 plicable drug or a generic drug, as the case may
20 be;

21 “(C) the establishment of procedures to
22 ensure that, not later than the applicable num-
23 ber of calendar days after the dispensing of an
24 applicable drug or a generic drug, as the case
25 may be, by a pharmacy or mail order service,

1 the pharmacy or mail order service is reim-
2 bursed for an amount equal to the difference
3 between—

4 “(i) the negotiated price of the appli-
5 cable drug or generic drug, respectively;
6 and

7 “(ii) the discounted price of the appli-
8 cable drug or generic drug, respectively;

9 “(D) the establishment of procedures to
10 ensure that the discounted price for an applica-
11 ble drug or a generic drug under this section is
12 applied before any coverage or financial assist-
13 ance under other health benefit plans or pro-
14 grams that provide coverage or financial assist-
15 ance for the purchase or provision of prescrip-
16 tion drug coverage on behalf of applicable bene-
17 ficiaries as the Secretary may specify; and

18 “(E) providing a reasonable dispute resolu-
19 tion mechanism to resolve disagreements be-
20 tween manufacturers, applicable beneficiaries,
21 and the third party with a contract under sub-
22 section (d)(3).

23 “(2) MONITORING COMPLIANCE.—

1 “(A) IN GENERAL.—The Secretary shall
2 monitor compliance by a manufacturer with the
3 terms of an agreement under this section.

4 “(B) NOTIFICATION.—If a third party
5 with a contract under subsection (d)(3) deter-
6 mines that the manufacturer is not in compli-
7 ance with such agreement, the third party shall
8 notify the Secretary of such noncompliance for
9 appropriate enforcement under subsection (e).

10 “(3) COLLECTION OF DATA FROM PRESCRIP-
11 TION DRUG PLANS AND MA-PD PLANS.—The Sec-
12 retary may collect appropriate data from prescrip-
13 tion drug plans and MA-PD plans in a timeframe
14 that allows for discounted prices to be provided for
15 applicable drugs and generic drugs under this sec-
16 tion.

17 “(d) ADMINISTRATION.—

18 “(1) IN GENERAL.—Subject to paragraph (2),
19 the Secretary shall provide for the implementation of
20 this section, including the performance of the duties
21 described in subsection (c).

22 “(2) LIMITATION.—In providing for the imple-
23 mentation of this section, the Secretary shall not re-
24 ceive or distribute any funds of a manufacturer
25 under the program.

1 “(3) CONTRACT WITH THIRD PARTIES.—The
2 Secretary shall enter into a contract with 1 or more
3 third parties to administer the requirements estab-
4 lished by the Secretary in order to carry out this
5 section. At a minimum, the contract with a third
6 party under the preceding sentence shall require
7 that the third party—

8 “(A) receive and transmit information be-
9 tween the Secretary, manufacturers, and other
10 individuals or entities the Secretary determines
11 appropriate;

12 “(B) receive, distribute, or facilitate the
13 distribution of funds of manufacturers to ap-
14 propriate individuals or entities in order to
15 meet the obligations of manufacturers under
16 agreements under this section;

17 “(C) provide adequate and timely informa-
18 tion to manufacturers, consistent with the
19 agreement with the manufacturer under this
20 section, as necessary for the manufacturer to
21 fulfill its obligations under this section; and

22 “(D) permit manufacturers to conduct
23 periodic audits, directly or through contracts, of
24 the data and information used by the third
25 party to determine discounts for applicable

1 drugs of the manufacturer and generic drugs of
2 the manufacturer under the program.

3 “(4) PERFORMANCE REQUIREMENTS.—The
4 Secretary shall establish performance requirements
5 for a third party with a contract under paragraph
6 (3) and safeguards to protect the independence and
7 integrity of the activities carried out by the third
8 party under the program under this section.

9 “(5) ADMINISTRATION.—Chapter 35 of title 44,
10 United States Code, shall not apply to the program
11 under this section.

12 “(e) ENFORCEMENT.—

13 “(1) AUDITS.—Each manufacturer with an
14 agreement in effect under this section shall be sub-
15 ject to periodic audit by the Secretary.

16 “(2) CIVIL MONEY PENALTY.—

17 “(A) IN GENERAL.—The Secretary shall
18 impose a civil money penalty on a manufacturer
19 that fails to provide applicable beneficiaries dis-
20 counts for applicable drugs of the manufacturer
21 or generic drugs of the manufacturer in accord-
22 ance with such agreement for each such failure
23 in an amount the Secretary determines is com-
24 mensurate with the sum of—

1 “(i) the amount that the manufac-
2 turer would have paid with respect to such
3 discounts under the agreement, which will
4 then be used to pay the discounts which
5 the manufacturer had failed to provide;
6 and

7 “(ii) 25 percent of such amount.

8 “(B) APPLICATION.—The provisions of
9 section 1128A (other than subsections (a) and
10 (b)) shall apply to a civil money penalty under
11 this paragraph in the same manner as such
12 provisions apply to a penalty or proceeding
13 under section 1128A(a).

14 “(f) CLARIFICATION REGARDING AVAILABILITY OF
15 OTHER COVERED PART D DRUGS.—Nothing in this sec-
16 tion shall prevent an applicable beneficiary from pur-
17 chasing a covered part D drug that is not on the formulary
18 of the prescription drug plan or MA–PD plan that the
19 applicable beneficiary is enrolled in.

20 “(g) DEFINITIONS.—In this section:

21 “(1) APPLICABLE BENEFICIARY.—The term
22 ‘applicable beneficiary’ means an individual who, on
23 the date of dispensing a covered part D drug—

24 “(A) is enrolled in a prescription drug plan
25 or an MA–PD plan;

1 “(B) is not enrolled in a qualified retiree
2 prescription drug plan; and

3 “(C) has incurred costs for covered part D
4 drugs in the year that are equal to or exceed
5 the annual deductible specified in section
6 1860D–2(b)(1) for such year.

7 “(2) APPLICABLE DRUG.—The term ‘applicable
8 drug’ means, with respect to an applicable bene-
9 ficiary, a covered part D drug—

10 “(A) approved under a new drug applica-
11 tion under section 505(c) of the Federal Food,
12 Drug, and Cosmetic Act or, in the case of a bio-
13 logic product, licensed under section 351 of the
14 Public Health Service Act (including a product
15 licensed under subsection (k) of such section);
16 and

17 “(B)(i) if the PDP sponsor of the prescrip-
18 tion drug plan or the MA organization offering
19 the MA–PD plan uses a formulary, which is on
20 the formulary of the prescription drug plan or
21 MA–PD plan that the applicable beneficiary is
22 enrolled in;

23 “(ii) if the PDP sponsor of the prescrip-
24 tion drug plan or the MA organization offering
25 the MA–PD plan does not use a formulary, for

1 which benefits are available under the prescrip-
2 tion drug plan or MA–PD plan that the appli-
3 cable beneficiary is enrolled in; or

4 “(iii) is provided through an exception or
5 appeal.

6 “(3) APPLICABLE NUMBER OF CALENDAR
7 DAYS.—The term ‘applicable number of calendar
8 days’ means—

9 “(A) with respect to claims for reimburse-
10 ment submitted electronically, 14 days; and

11 “(B) with respect to claims for reimburse-
12 ment submitted otherwise, 30 days.

13 “(4) DISCOUNTED PRICE.—

14 “(A) IN GENERAL.—The term ‘discounted
15 price’ means—

16 “(i) with respect to an applicable drug
17 of a manufacturer furnished during a year
18 to an applicable beneficiary—

19 “(I) who has not incurred costs
20 for covered part D drugs in the year
21 that are equal to or exceed the annual
22 out-of-pocket threshold specified in
23 section 1860D–2(b)(4)(B)(i) for the
24 year, 90 percent of the negotiated
25 price of such drug; and

1 “(II) who has incurred such costs
2 in the year that are equal to or exceed
3 such threshold for the year, 90 per-
4 cent of the negotiated price of such
5 drug; and

6 “(ii) with respect to a generic drug of
7 a manufacturer furnished during a year to
8 an applicable beneficiary who has not in-
9 curred costs for covered part D drugs in
10 the year that are equal to or exceed the
11 annual out-of-pocket threshold specified in
12 section 1860D–2(b)(4)(B)(i) for the year,
13 90 percent of the negotiated price of such
14 drug.

15 “(B) CLARIFICATION.—Nothing in this
16 section shall be construed as affecting the re-
17 sponsibility of an applicable beneficiary for pay-
18 ment of a dispensing fee for an applicable drug
19 or a generic drug.

20 “(C) SPECIAL CASE FOR CLAIMS SPANNING
21 DEDUCTIBLE.—In the case where the entire
22 amount of the negotiated price of an individual
23 claim for an applicable drug or a generic drug
24 with respect to an applicable beneficiary does
25 not fall at or above the annual deductible speci-

1 fied in section 1860D–2(b)(1) for the year, the
2 manufacturer of the applicable drug shall pro-
3 vide the discounted price under this section on
4 only the portion of the negotiated price of the
5 applicable drug or generic drug, respectively,
6 that falls at or above such annual deductible.

7 “(5) GENERIC DRUG.—The term ‘generic drug’
8 means, with respect to an applicable beneficiary, a
9 covered part D drug that is not an applicable drug.

10 “(6) MANUFACTURER.—The term ‘manufac-
11 turer’ means any entity which is engaged in the pro-
12 duction, preparation, propagation, compounding,
13 conversion, or processing of prescription drug prod-
14 ucts, either directly or indirectly by extraction from
15 substances of natural origin, or independently by
16 means of chemical synthesis, or by a combination of
17 extraction and chemical synthesis. Such term does
18 not include a wholesale distributor of drugs or a re-
19 tail pharmacy licensed under State law.

20 “(7) NEGOTIATED PRICE.—The term ‘nego-
21 tiated price’ has the meaning given such term in sec-
22 tion 1860D–2(d)(1)(B), except that such negotiated
23 price shall not include any dispensing fee for an ap-
24 plicable drug or a generic drug.

1 “(8) QUALIFIED RETIREE PRESCRIPTION DRUG
2 PLAN.—The term ‘qualified retiree prescription drug
3 plan’ has the meaning given such term in section
4 11860D–22(a)(2).”.

5 (2) SUNSET OF MEDICARE COVERAGE GAP DIS-
6 COUNT PROGRAM.—Section 1860D–14A of the So-
7 cial Security Act (42 U.S.C. 1395–114a) is amend-
8 ed—

9 (A) in subsection (a), in the first sentence,
10 by striking “The Secretary” and inserting
11 “Subject to subsection (h), the Secretary”; and

12 (B) by adding at the end the following new
13 subsection:

14 “(h) SUNSET OF PROGRAM.—

15 “(1) IN GENERAL.—The program shall not
16 apply to applicable drugs dispensed on or after Jan-
17 uary 1, 2022, and, subject to paragraph (2), agree-
18 ments under this section shall be terminated as of
19 such date.

20 “(2) CONTINUED APPLICATION FOR APPLICA-
21 BLE DRUGS DISPENSED PRIOR TO SUNSET.—The
22 provisions of this section (including all responsibil-
23 ities and duties) shall continue to apply after Janu-
24 ary 1, 2022, with respect to applicable drugs dis-
25 pensed prior to such date.”.

1 (3) INCLUSION OF ACTUARIAL VALUE OF MANU-
2 FACTURER DISCOUNTS IN BIDS.—Section 1860D–11
3 of the Social Security Act (42 U.S.C. 1395w–111)
4 is amended—

5 (A) in subsection (b)(2)(C)(iii)—

6 (i) by striking “assumptions regarding
7 the reinsurance” and inserting “assump-
8 tions regarding—

9 “(I) the reinsurance”; and

10 (ii) by adding at the end the fol-
11 lowing:

12 “(II) for 2022 and each subse-
13 quent year, the manufacturer dis-
14 counts provided under section 1860D–
15 14B subtracted from the actuarial
16 value to produce such bid; and”;

17 (B) in subsection (c)(1)(C)—

18 (i) by striking “an actuarial valuation
19 of the reinsurance” and inserting “an ac-
20 tuarial valuation of—

21 “(i) the reinsurance”;

22 (ii) in clause (i), as added by clause
23 (i) of this subparagraph, by adding “and”
24 at the end; and

1 (iii) by adding at the end the fol-
2 lowing:

3 “(ii) for 2022 and each subsequent
4 year, the manufacturer discounts provided
5 under section 1860D–14B;”.

6 (d) DETERMINATION OF ALLOWABLE REINSURANCE
7 COSTS.—Section 1860D–15(b) of the Social Security Act
8 (42 U.S.C. 1395w–115(b)) is amended—

9 (1) in paragraph (2)—

10 (A) by striking “COSTS.—For purposes”
11 and inserting “COSTS.—

12 “(A) IN GENERAL.—Subject to subpara-
13 graph (B), for purposes”.

14 (B) by adding at the end the following new
15 subparagraph:

16 “(B) INCLUSION OF MANUFACTURER DIS-
17 COUNTS ON APPLICABLE DRUGS AND GENERIC
18 DRUGS.—For purposes of applying subpara-
19 graph (A), the term ‘allowable reinsurance
20 costs’ shall include the portion of the negotiated
21 price (as defined in section 1860D–14B(g)(7))
22 of an applicable drug (as defined in section
23 1860D–14(g)(2)) that was paid by a manufac-
24 turer under the manufacturer discount program
25 under section 1860D–14B and the portion of

1 the negotiated price (as so defined) of a generic
2 drug (as defined in section 1860D–14(g)(5))
3 that was paid by a manufacturer under such
4 program.”; and

5 (2) in paragraph (3)—

6 (A) in the first sentence, by striking “For
7 purposes” and inserting “Subject to paragraph
8 (2)(B), for purposes”; and

9 (B) in the second sentence, by inserting
10 “or, in the case of an applicable drug or a ge-
11 neric drug, by a manufacturer” after “by the
12 individual or under the plan”.

13 (e) UPDATING RISK ADJUSTMENT METHODOLOGIES
14 TO ACCOUNT FOR PART D MODERNIZATION REDESIGN.—
15 Section 1860D–15(e) of the Social Security Act (42
16 U.S.C. 1395w–115(e)) is amended by adding at the end
17 the following new paragraph:

18 “(3) UPDATING RISK ADJUSTMENT METH-
19 ODOLOGIES TO ACCOUNT FOR PART D MODERNIZA-
20 TION REDESIGN.—The Secretary shall update the
21 risk adjustment model used to adjust bid amounts
22 pursuant to this subsection as appropriate to take
23 into account changes in benefits under this part pur-
24 suant to the amendments made by section 121 of
25 the Lower Drug Costs Now Act of 2019.”.

1 (f) CONDITIONS FOR COVERAGE OF DRUGS UNDER
2 THIS PART.—Section 1860D–43 of the Social Security
3 Act (42 U.S.C. 1395w–153) is amended—

4 (1) in subsection (a)—

5 (A) in paragraph (2), by striking “and” at
6 the end;

7 (B) in paragraph (3), by striking the pe-
8 riod at the end and inserting a semicolon; and

9 (C) by adding at the end the following new
10 paragraphs:

11 “(4) participate in the manufacturer discount
12 program under section 1860D–14B;

13 “(5) have entered into and have in effect an
14 agreement described in subsection (b) of such sec-
15 tion 1860D–14B with the Secretary; and

16 “(6) have entered into and have in effect, under
17 terms and conditions specified by the Secretary, a
18 contract with a third party that the Secretary has
19 entered into a contract with under subsection (d)(3)
20 of such section 1860D–14B.”;

21 (2) by striking subsection (b) and inserting the
22 following:

23 “(b) EFFECTIVE DATE.—Paragraphs (1) through (3)
24 of subsection (a) shall apply to covered part D drugs dis-
25 pensed under this part on or after January 1, 2011, and

1 before January 1, 2022, and paragraphs (4) through (6)
2 of such subsection shall apply to covered part D drugs
3 dispensed on or after January 1, 2022.”; and

4 (3) in subsection (c), by striking paragraph (2)
5 and inserting the following:

6 “(2) the Secretary determines that in the period
7 beginning on January 1, 2011, and ending on De-
8 cember 31, 2011 (with respect to paragraphs (1)
9 through (3) of subsection (a)) or the period begin-
10 ning on January 1, 2022, and ending December 31,
11 2022 (with respect to paragraphs (4) through (6) of
12 such subsection), there were extenuating cir-
13 cumstances.”.

14 (g) CONFORMING AMENDMENTS.—

15 (1) Section 1860D–2 of the Social Security Act
16 (42 U.S.C. 1395w–102) is amended—

17 (A) in subsection (a)(2)(A)(i)(I), by strik-
18 ing “, or an increase in the initial” and insert-
19 ing “or for a year preceding 2022 an increase
20 in the initial”;

21 (B) in subsection (c)(1)(C)—

22 (i) in the subparagraph heading, by
23 striking “AT INITIAL COVERAGE LIMIT”;
24 and

1 (ii) by inserting “for a year preceding
2 2022 or the annual out-of-pocket threshold
3 specified in subsection (b)(4)(B) for the
4 year for 2022 and each subsequent year”
5 after “subsection (b)(3) for the year” each
6 place it appears; and

7 (C) in subsection (d)(1)(A), by striking “or
8 an initial” and inserting “or for a year pre-
9 ceding 2022, an initial”.

10 (2) Section 1860D–4(a)(4)(B)(i) of the Social
11 Security Act (42 U.S.C. 1395w–104(a)(4)(B)(i)) is
12 amended by striking “the initial” and inserting “for
13 a year preceding 2022, the initial”.

14 (3) Section 1860D–14(a) of the Social Security
15 Act (42 U.S.C. 1395w–114(a)) is amended—

16 (A) in paragraph (1)—

17 (i) in subparagraph (C), by striking
18 “The continuation” and inserting “For a
19 year preceding 2022, the continuation”;

20 (ii) in subparagraph (D)(iii), by strik-
21 ing “1860D–2(b)(4)(A)(i)(I)” and insert-
22 ing “1860D–2(b)(4)(A)(i)(I)(aa)”; and

23 (iii) in subparagraph (E), by striking
24 “The elimination” and inserting “For a
25 year preceding 2022, the elimination”; and

1 (B) in paragraph (2)—

2 (i) in subparagraph (C), by striking
3 “The continuation” and inserting “For a
4 year preceding 2022, the continuation”;
5 and

6 (ii) in subparagraph (E)—

7 (I) by inserting “for a year pre-
8 ceding 2022,” after “subsection (e)”;
9 and

10 (II) by striking “1860D-
11 2(b)(4)(A)(i)(I)” and inserting
12 “1860D-2(b)(4)(A)(i)(I)(aa)”.

13 (4) Section 1860D-21(d)(7) of the Social Secu-
14 rity Act (42 U.S.C. 1395w-131(d)(7)) is amended
15 by striking “section 1860D-2(b)(4)(B)(i)” and in-
16 serting “section 1860D-2(b)(4)(C)(i)”.

17 (5) Section 1860D-22(a)(2)(A) of the Social
18 Security Act (42 U.S.C. 1395w-132(a)(2)(A)) is
19 amended—

20 (A) by striking “the value of any discount”
21 and inserting the following: “the value of—
22 “(i) for years prior to 2022, any dis-
23 count”;

1 (B) in clause (i), as inserted by subpara-
2 graph (A) of this paragraph, by striking the pe-
3 riod at the end and inserting “; and”; and

4 (C) by adding at the end the following new
5 clause:

6 “(ii) for 2022 and each subsequent
7 year, any discount provided pursuant to
8 section 1860D–14B.”.

9 (6) Section 1860D–41(a)(6) of the Social Secu-
10 rity Act (42 U.S.C. 1395w–151(a)(6)) is amended—

11 (A) by inserting “for a year before 2022”
12 after “1860D–2(b)(3)”; and

13 (B) by inserting “for such year” before the
14 period.

15 (h) EFFECTIVE DATE.—The amendments made by
16 this section shall apply to plan year 2022 and subsequent
17 plan years.

Strike section 302 and insert the following new sec-
tion:

1 **SEC. 302. ALLOWING CERTAIN ENROLLEES OF PRESCRIP-**
2 **TION DRUGS PLANS AND MA-PD PLANS**
3 **UNDER MEDICARE PROGRAM TO SPREAD**
4 **OUT COST-SHARING UNDER CERTAIN CIR-**
5 **CUMSTANCES.**

6 (a) STANDARD PRESCRIPTION DRUG COVERAGE.—
7 Section 1860D–2(b)(2) of the Social Security Act (42
8 U.S.C. 1395w–102(b)(2)), as amended by section 121, is
9 further amended—

10 (1) in subparagraph (A), by striking “Subject
11 to subparagraphs (C) and (D)” and inserting “Sub-
12 ject to subparagraphs (C), (D), and (E)”; and

13 (2) by adding at the end the following new sub-
14 paragraph:

15 “(E) ENROLLEE OPTION REGARDING
16 SPREADING COST-SHARING.—

17 “(i) IN GENERAL.—The Secretary
18 shall establish by regulation a process
19 under which, with respect to plan year
20 2022 and subsequent plan years, a pre-
21 scription drug plan or an MA–PD plan
22 shall, in the case of a part D eligible indi-
23 vidual enrolled with such plan for such
24 plan year with respect to whom the plan
25 projects that the dispensing of a covered
26 part D drug to such individual will result

1 in the individual incurring costs within a
2 30-day period that are equal to a signifi-
3 cant percentage (as specified by the Sec-
4 retary pursuant to such regulation) of the
5 annual out-of-pocket threshold specified in
6 paragraph (4)(B) for such plan year, pro-
7 vide such individual with the option to
8 make the coinsurance payment required
9 under subparagraph (A) for such costs in
10 the form of equal monthly installments
11 over the remainder of such plan year.

12 “(ii) SIGNIFICANT PERCENTAGE LIM-
13 TATIONS.—In specifying a significant per-
14 centage pursuant to the regulation estab-
15 lished by the Secretary under clause (i),
16 the Secretary may not specify a percentage
17 that is less than 30 percent or greater
18 than 100 percent.”.

19 (b) ALTERNATIVE PRESCRIPTION DRUG COV-
20 ERAGE.—Section 1860D–2(c) of the Social Security Act
21 (42 U.S.C. 1395w–102(c)) is amended by adding at the
22 end the following new paragraph:

23 “(4) SAME ENROLLEE OPTION REGARDING
24 SPREADING COST-SHARING.—For plan year 2022
25 and subsequent plan years, the coverage provides the

1 enrollee option regarding spreading cost-sharing de-
2 scribed in and required under subsection
3 (b)(2)(E).”.

After section 303, insert the following new section
(and update the table of contents accordingly):

4 **SEC. 304. DRUG DISCOUNTS REQUIRED TO BE PASSED**
5 **THROUGH TO THE PLAN SPONSOR.**

6 (a) IN GENERAL.—Section 1150A of the Social Secu-
7 rity Act (42 U.S.C. 1320b–23) is amended—

8 (1) in the heading, by inserting “**; DRUG DIS-**
9 **COUNTS REQUIRED TO BE PASSED THROUGH**
10 **TO THE PLAN SPONSOR**” before the period at the
11 end; and

12 (2) by adding at the end the following new sub-
13 sections:

14 “(e) **DRUG DISCOUNTS REQUIRED TO BE PASSED**
15 **THROUGH TO THE PLAN SPONSOR.**—

16 “(1) **REQUIREMENT.**—Beginning January 1,
17 2022, a PBM that manages prescription drug cov-
18 erage under a contract with a PDP sponsor or MA
19 organization described in subsection (b)(1) or a
20 qualified health benefits plan described in subsection
21 (b)(2), shall, with respect to the plan sponsor of a
22 health benefits plan, pass through to the plan spon-
23 sor 100 percent of the aggregate amount of the re-

1 bates, discounts, or price concessions (other than
2 bona fide service fees (as defined in subsection (g)))
3 that the PBM negotiates that are attributable to pa-
4 tient utilization under the plan (including any re-
5 bates, discounts, or other price concessions (other
6 than bona fide service fees (as so defined)) that are
7 received by an agent or affiliate of the PBM acting
8 on the PBM’s behalf). Such a PBM may retain bona
9 fide service fees (as so defined), to the extent that
10 such fees are not based on a percentage of the sales
11 for a drug or otherwise linked in any way to the
12 price or formulary position or placement of a drug.

13 “(2) ENFORCEMENT.—A PDP sponsor of a
14 prescription drug plan or an MA organization offer-
15 ing an MA–PD plan under part D of title XVIII
16 may not contract with a PBM that is not in compli-
17 ance with the requirement under paragraph (1).

18 “(f) BONA FIDE SERVICE FEES DEFINED.—The
19 term ‘bona fide service fees’ means, with respect to a
20 PBM, fees paid to such PBM (or an agent or affiliate of
21 such PBM acting on the PBM’s behalf) by a manufac-
22 turer, customer, or client of the PBM that represent the
23 fair market value for a bona fide, itemized service actually
24 performed on behalf of the manufacturer, customer, or cli-
25 ent, that the manufacturer, customer, or client would oth-

1 erwise perform (or contract for) in the absence of the serv-
2 icer arrangement, and that the PBM does not pass on to
3 another party.”.

4 (b) EFFECTIVE DATE.—The amendments made by
5 subsection (a) shall take effect on January 1, 2022.

In section 401, in the matter preceding paragraph
(1), strike “as amended by section 301(d)” and insert
“as amended by section 301(g)”.

In section 404, in the matter preceding paragraph
(1), strike “as amended by sections 301(d)” and insert
“as amended by sections 301(g)”.

