

1 Title: To amend the Federal Food, Drug, and Cosmetic Act to revise and extend the user-fee
2 programs for prescription drugs, medical devices, generic drugs, and biosimilar biological
3 products, and for other purposes.
4
5

6 Be it enacted by the Senate and House of Representatives of the United States of America in
7 Congress assembled,

8 SECTION 1. SHORT TITLE.

9 This Act may be cited as the “FDA Reauthorization Act of 2017”.

10 SEC. 2. TABLE OF CONTENTS.

11 The table of contents for this Act is as follows:

12 Sec.1.Short title.

13 Sec.2.Table of contents.

14 TITLE I—FEES RELATING TO DRUGS

15 Sec.101.Short title; finding.

16 Sec.102.Authority to assess and use drug fees.

17 Sec.103.Reauthorization; reporting requirements.

18 Sec.104.Sunset dates.

19 Sec.105.Effective date.

20 Sec.106.Savings clause.

21 TITLE II—FEES RELATING TO DEVICES

22 Sec.201.Short title; findings.

23 Sec.202.Definitions.

24 Sec.203.Authority to assess and use device fees.

25 Sec.204.Reauthorization; reporting requirements.

26 Sec.205.Conformity assessment pilot program.

27 Sec.206.Reauthorization of review.

28 Sec.207.Electronic format for submissions.

29 Sec.208.Savings clause.

30 Sec.209.Effective date.

31 Sec.210.Sunset clause.

32 TITLE III—FEES RELATING TO GENERIC DRUGS

- 1 Sec.301.Short title; finding.
- 2 Sec.302.Definitions.
- 3 Sec.303.Authority to assess and use human generic drug fees.
- 4 Sec.304.Reauthorization; reporting requirements.
- 5 Sec.305.Sunset dates.
- 6 Sec.306.Effective date.
- 7 Sec.307.Savings clause.

8 TITLE IV—FEES RELATING TO BIOSIMILAR 9 BIOLOGICAL PRODUCTS

- 10 Sec.401.Short title; finding.
- 11 Sec.402.Definitions.
- 12 Sec.403.Authority to assess and use biosimilar fees.
- 13 Sec.404.Reauthorization; reporting requirements.
- 14 Sec.405.Sunset dates.
- 15 Sec.406.Effective date.
- 16 Sec.407.Savings clause.

17 TITLE V—REAUTHORIZATION OF OTHER PROGRAMS

- 18 Sec.501.Reauthorization of provision relating to exclusivity of certain drugs containing single
19 enantiomers.
- 20 Sec.502.Reauthorization of pediatric humanitarian device exceptions.
- 21 Sec.503.Reauthorization of the critical path public-private partnerships.
- 22 Sec.504.Reauthorization of pediatric device consortia.
- 23 Sec.505.Reauthorization of orphan grants program.

24 TITLE I—FEES RELATING TO DRUGS

25 SEC. 101. SHORT TITLE; FINDING.

26 (a) Short Title.—This title may be cited as the “Prescription Drug User Fee Amendments of
27 2017”.

28 (b) Finding.—The Congress finds that the fees authorized by the amendments made in this
29 title will be dedicated toward expediting the drug development process and the process for the
30 review of human drug applications, including postmarket drug safety activities, as set forth in the
31 goals identified for purposes of part 2 of subchapter C of chapter VII of the Federal Food, Drug,
32 and Cosmetic Act, in the letters from the Secretary of Health and Human Services to the
33 Chairman of the Committee on Health, Education, Labor, and Pensions of the Senate and the
34 Chairman of the Committee on Energy and Commerce of the House of Representatives, as set

1 forth in the Congressional Record.

2 **SEC. 102. AUTHORITY TO ASSESS AND USE DRUG**
3 **FEES.**

4 (a) Types of Fees.—

5 (1) IN GENERAL.—Section 736(a) of the Federal Food, Drug, and Cosmetic Act (21
6 U.S.C. 379h(a)) is amended—

7 (A) in the matter preceding paragraph (1), by striking “fiscal year 2013” and
8 inserting “fiscal year 2018”;

9 (B) in the heading of paragraph (1), by striking “AND SUPPLEMENT”;

10 (C) in paragraph (1), by striking “or a supplement” and “or supplement” each place
11 either appears;

12 (D) in paragraph (1)(A)—

13 (i) in clause (i), by striking “(c)(4)” and inserting “(c)(5)”; and

14 (ii) in clause (ii), by striking “A fee established” and all that follows through
15 “are required.” and inserting the following: “A fee established under subsection
16 (c)(5) for a human drug application for which clinical data (other than
17 bioavailability or bioequivalence studies) with respect to safety or effectiveness
18 are not required for approval.”;

19 (E) in the heading of paragraph (1)(C), by striking “OR SUPPLEMENT”;

20 (F) in paragraph (1)(F)—

21 (i) in the heading, by striking “OR INDICATION”; and

22 (ii) by striking the second sentence;

23 (G) by striking paragraph (2) (relating to a prescription drug establishment fee);

24 (H) by redesignating paragraph (3) as paragraph (2);

25 (I) in the heading of paragraph (2), as so redesignated, by striking “PRESCRIPTION
26 DRUG PRODUCT FEE” and inserting “PRESCRIPTION DRUG PROGRAM FEE”;

27 (J) in subparagraph (A) of such paragraph (2), by amending the first sentence to read
28 as follows: “Except as provided in subparagraphs (B) and (C), each person who is
29 named as the applicant in a human drug application, and who, after September 1, 1992,
30 had pending before the Secretary a human drug application or supplement, shall pay
31 the annual prescription drug program fee established for a fiscal year under subsection
32 (c)(5) for each prescription drug product that is identified in such a human drug
33 application approved as of October 1 of such fiscal year.”;

34 (K) in subparagraph (B) of such paragraph (2)—

35 (i) in the heading of subparagraph (B), by inserting after “EXCEPTION” the
36 following: “FOR CERTAIN PRESCRIPTION DRUG PRODUCTS”; and

37 (ii) by striking “A prescription drug product shall not be assessed a fee” and

1 inserting “A prescription drug program fee shall not be assessed for a prescription
2 drug product”; and

3 (L) by adding at the end of such paragraph (2) the following:

4 “(C) LIMITATION.—A person who is named as the applicant in an approved human
5 drug application shall not be assessed more than 5 prescription drug program fees for a
6 fiscal year for prescription drug products identified in such approved human drug
7 application.”.

8 (2) CONFORMING AMENDMENT.—Subparagraph (C) of section 740(a)(3) of the Federal
9 Food, Drug, and Cosmetic Act (21 U.S.C. 379j–12(a)(3)) is amended to read as follows:

10 “(C) LIMITATION.—An establishment shall be assessed only one fee per fiscal year
11 under this section.”.

12 (b) Fee Revenue Amounts.—Subsection (b) of section 736 of the Federal Food, Drug, and
13 Cosmetic Act (21 U.S.C. 379h) is amended to read as follows:

14 “(b) Fee Revenue Amounts.—

15 “(1) IN GENERAL.—For each of the fiscal years 2018 through 2022, fees under subsection
16 (a) shall, except as provided in subsections (c), (d), (f), and (g), be established to generate a
17 total revenue amount under such subsection that is equal to the sum of—

18 “(A) the annual base revenue for the fiscal year (as determined under paragraph (3));

19 “(B) the dollar amount equal to the inflation adjustment for the fiscal year (as
20 determined under subsection (c)(1));

21 “(C) the dollar amount equal to the capacity planning adjustment for the fiscal year
22 (as determined under subsection (c)(2));

23 “(D) the dollar amount equal to the operating reserve adjustment for the fiscal year,
24 if applicable (as determined under subsection (c)(3));

25 “(E) the dollar amount equal to the additional direct cost adjustment for the fiscal
26 year (as determined under subsection (c)(4)); and

27 “(F) additional dollar amounts for each fiscal year as follows:

28 “(i) \$20,077,793 for fiscal year 2018;

29 “(ii) \$21,317,472 for fiscal year 2019;

30 “(iii) \$16,953,329 for fiscal year 2020;

31 “(iv) \$5,426,896 for fiscal year 2021; and

32 “(v) \$2,769,609 for fiscal year 2022.

33 “(2) TYPES OF FEES.—Of the total revenue amount determined for a fiscal year under
34 paragraph (1)—

35 “(A) 20 percent shall be derived from human drug application fees under subsection
36 (a)(1); and

37 “(B) 80 percent shall be derived from prescription drug program fees under

1 subsection (a)(2).

2 “(3) ANNUAL BASE REVENUE.—For purposes of paragraph (1), the dollar amount of the
3 annual base revenue for a fiscal year shall be—

4 “(A) for fiscal year 2018, \$878,590,000; and

5 “(B) for fiscal years 2019 through 2022, the dollar amount of the total revenue
6 amount established under paragraph (1) for the previous fiscal year, not including any
7 adjustments made under subsection (c)(3) or (c)(4).”.

8 (c) Adjustments; Annual Fee Setting.—Subsection (c) of section 736 of the Federal Food,
9 Drug, and Cosmetic Act (21 U.S.C. 379h) is amended to read as follows:

10 “(c) Adjustments; Annual Fee Setting.—

11 “(1) INFLATION ADJUSTMENT.—

12 “(A) IN GENERAL.—For purposes of subsection (b)(1)(B), the dollar amount of the
13 inflation adjustment to the annual base revenue for each fiscal year shall be equal to
14 the product of—

15 “(i) such annual base revenue for the fiscal year under subsection (b)(1)(A);
16 and

17 “(ii) the inflation adjustment percentage under subparagraph (B).

18 “(B) INFLATION ADJUSTMENT PERCENTAGE.—The inflation adjustment percentage
19 under this subparagraph for a fiscal year is equal to the sum of—

20 “(i) the average annual percent change in the cost, per full-time equivalent
21 position of the Food and Drug Administration, of all personnel compensation and
22 benefits paid with respect to such positions for the first 3 years of the preceding 4
23 fiscal years, multiplied by the proportion of personnel compensation and benefits
24 costs to total costs of the process for the review of human drug applications (as
25 defined in section 735(6)) for the first 3 years of the preceding 4 fiscal years; and

26 “(ii) the average annual percent change that occurred in the Consumer Price
27 Index for urban consumers (Washington-Baltimore, DC–MD–VA–WV; Not
28 Seasonally Adjusted; All items; Annual Index) for the first 3 years of the
29 preceding 4 years of available data multiplied by the proportion of all costs other
30 than personnel compensation and benefits costs to total costs of the process for
31 the review of human drug applications (as defined in section 735(6)) for the first 3
32 years of the preceding 4 fiscal years.

33 “(2) CAPACITY PLANNING ADJUSTMENT.—

34 “(A) IN GENERAL.—For each fiscal year, after the annual base revenue established in
35 subsection (b)(1)(A) is adjusted for inflation in accordance with paragraph (1), such
36 revenue shall be adjusted further for such fiscal year, in accordance with this
37 paragraph, to reflect changes in the resource capacity needs of the Secretary for the
38 process for the review of human drug applications.

39 “(B) INTERIM METHODOLOGY.—

40 “(i) IN GENERAL.—Until the capacity planning methodology described in

1 subparagraph (C) is effective, the adjustment under this paragraph for a fiscal year
2 shall be based on the product of—

3 “(I) the annual base revenue for such year, as adjusted for inflation under
4 paragraph (1); and

5 “(II) the adjustment percentage under clause (ii).

6 “(ii) ADJUSTMENT PERCENTAGE.—The adjustment percentage under this clause
7 for a fiscal year is the weighted change in the 3-year average ending in the most
8 recent year for which data are available, over the 3-year average ending in the
9 previous year, for—

10 “(I) the total number of human drug applications, efficacy supplements,
11 and manufacturing supplements submitted to the Secretary;

12 “(II) the total number of active commercial investigational new drug
13 applications; and

14 “(III) the total number of formal meetings scheduled by the Secretary, and
15 written responses issued by the Secretary in lieu of such formal meetings, as
16 identified in section I.H of the letters described in section 101(b) of the
17 Prescription Drug User Fee Amendments of 2017.

18 “(C) CAPACITY PLANNING METHODOLOGY.—

19 “(i) DEVELOPMENT; EVALUATION AND REPORT.—The Secretary shall obtain,
20 through a contract with an independent accounting or consulting firm, a report
21 evaluating options and recommendations for a new methodology to accurately
22 assess changes in the resource and capacity needs of the process for the review of
23 human drug applications. The capacity planning methodological options and
24 recommendations presented in such report shall utilize and be informed by
25 personnel time reporting data as an input. The report shall be published for public
26 comment no later than the end of fiscal year 2020.

27 “(ii) ESTABLISHMENT AND IMPLEMENTATION.—After review of the report
28 described in clause (i) and any public comments thereon, the Secretary shall
29 establish a capacity planning methodology for purposes of this paragraph, which
30 shall—

31 “(I) replace the interim methodology under subparagraph (B);

32 “(II) incorporate such approaches and attributes as the Secretary
33 determines appropriate; and

34 “(III) be effective beginning with the first fiscal year for which fees are set
35 after such capacity planning methodology is established.

36 “(D) LIMITATION.—Under no circumstances shall an adjustment under this
37 paragraph result in fee revenue for a fiscal year that is less than the sum of the amounts
38 under subsections (b)(1)(A) (the annual base revenue for the fiscal year) and (b)(1)(B)
39 (the dollar amount of the inflation adjustment for the fiscal year).

40 “(E) PUBLICATION IN FEDERAL REGISTER.—The Secretary shall publish in the

1 Federal Register notice under paragraph (5) the fee revenue and fees resulting from the
2 adjustment and the methodologies under this paragraph.

3 “(3) OPERATING RESERVE ADJUSTMENT.—

4 “(A) INCREASE.—For fiscal year 2018 and subsequent fiscal years, the Secretary
5 may, in addition to adjustments under paragraphs (1) and (2), further increase the fee
6 revenue and fees if such an adjustment is necessary to provide for not more than 14
7 weeks of operating reserves of carryover user fees for the process for the review of
8 human drug applications.

9 “(B) DECREASE.—If the Secretary has carryover balances for such process in excess
10 of 14 weeks of such operating reserves, the Secretary shall decrease such fee revenue
11 and fees to provide for not more than 14 weeks of such operating reserves.

12 “(C) NOTICE OF RATIONALE.—If an adjustment under subparagraph (A) or (B) is
13 made, the rationale for the amount of the increase or decrease (as applicable) in fee
14 revenue and fees shall be contained in the annual Federal Register notice under
15 paragraph (5) establishing fee revenue and fees for the fiscal year involved.

16 “(4) ADDITIONAL DIRECT COST ADJUSTMENT.—

17 “(A) IN GENERAL.—The Secretary shall, in addition to adjustments under paragraphs
18 (1), (2), and (3), further increase the fee revenue and fees—

19 “(i) for fiscal year 2018, by \$8,730,000; and

20 “(ii) for fiscal year 2019 and subsequent fiscal years, by the amount determined
21 under subparagraph (B).

22 “(B) AMOUNT.—The amount determined under this subparagraph is—

23 “(i) \$8,730,000, multiplied by

24 “(ii) the Consumer Price Index for urban consumers (Washington-Baltimore,
25 DC-MD-VA-WV; Not Seasonally Adjusted; All Items; Annual Index) for the
26 most recent year of available data, divided by such Index for 2016.

27 “(5) ANNUAL FEE SETTING.—The Secretary shall, not later than 60 days before the start of
28 each fiscal year that begins after September 30, 2017—

29 “(A) establish, for the next fiscal year, human drug application fees and prescription
30 drug program fees under subsection (a), based on the revenue amounts established
31 under subsection (b) and the adjustments provided under this subsection; and

32 “(B) publish such fee revenue and fees in the Federal Register.

33 “(6) LIMIT.—The total amount of fees charged, as adjusted under this subsection, for a
34 fiscal year may not exceed the total costs for such fiscal year for the resources allocated for
35 the process for the review of human drug applications.”.

36 (d) Fee Waiver or Reduction.—Section 736(d) of the Federal Food, Drug, and Cosmetic Act
37 (21 U.S.C. 379h(d)) is amended—

38 (1) in paragraph (1)—

39 (A) by inserting “or” at the end of subparagraph (B);

- 1 (B) by striking subparagraph (C); and
2 (C) by redesignating subparagraph (D) as subparagraph (C);
3 (2) by striking paragraph (3) (relating to use of standard costs);
4 (3) by redesignating paragraph (4) as paragraph (3); and
5 (4) in paragraph (3), as so redesignated—

6 (A) in subparagraphs (A) and (B), by striking “paragraph (1)(D)” and inserting
7 “paragraph (1)(C)”; and

8 (B) in subparagraph (B)—

9 (i) by striking clause (ii);

10 (ii) by striking “shall pay” through “(i) application fees” and inserting “shall
11 pay application fees”; and

12 (iii) by striking “; and” at the end and inserting a period.

13 (e) Effect of Failure to Pay Fees.—Section 736(e) of the Federal Food, Drug, and Cosmetic
14 Act (21 U.S.C. 379h(e)) is amended by striking “all fees” and inserting “all such fees”.

15 (f) Limitations.—Section 736(f)(2) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C.
16 379h(f)(2)) is amended by striking “supplements, prescription drug establishments, and
17 prescription drug products” and inserting “prescription drug program fees”.

18 (g) Crediting and Availability of Fees.—Section 736(g) of the Federal Food, Drug, and
19 Cosmetic Act (21 U.S.C. 379h(g)) is amended—

20 (1) in paragraph (3)—

21 (A) by striking “2013 through 2017” and inserting “2018 through 2022”; and

22 (B) by striking “and paragraph (4) of this subsection”; and

23 (2) by striking paragraph (4).

24 (h) Orphan Drugs.—Section 736(k) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C.
25 379h(k)) is amended by striking “product and establishment fees” each place it appears and
26 inserting “prescription drug program fees”.

27 SEC. 103. REAUTHORIZATION; REPORTING 28 REQUIREMENTS.

29 Section 736B of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379h–2) is amended—

30 (1) in subsection (a)(1)—

31 (A) in the matter before subparagraph (A), by striking “2013” and inserting “2018”;
32 and

33 (B) in subparagraph (A), by striking “Prescription Drug User Fee Amendments of
34 2012” and inserting “Prescription Drug User Fee Amendments of 2017”;

35 (2) in subsection (b), by striking “2013” and inserting “2018”; and

1 (3) in subsection (d), by striking “2017” each place it appears and inserting “2022”.

2 **SEC. 104. SUNSET DATES.**

3 (a) Authorization.—Sections 735 and 736 of the Federal Food, Drug, and Cosmetic Act (21
4 U.S.C. 379g; 379h) shall cease to be effective October 1, 2022.

5 (b) Reporting Requirements.—Section 736B of the Federal Food, Drug, and Cosmetic Act (21
6 U.S.C. 379h-2) shall cease to be effective January 31, 2023.

7 (c) Previous Sunset Provision.—Effective October 1, 2017, subsections (a) and (b) of section
8 105 of the Food and Drug Administration Safety and Innovation Act (Public Law 112–144) are
9 repealed.

10 **SEC. 105. EFFECTIVE DATE.**

11 The amendments made by this title shall take effect on October 1, 2017, or the date of the
12 enactment of this Act, whichever is later, except that fees under part 2 of subchapter C of chapter
13 VII of the Federal Food, Drug, and Cosmetic Act shall be assessed for all human drug
14 applications received on or after October 1, 2017, regardless of the date of the enactment of this
15 Act.

16 **SEC. 106. SAVINGS CLAUSE.**

17 Notwithstanding the amendments made by this title, part 2 of subchapter C of chapter VII of
18 the Federal Food, Drug, and Cosmetic Act, as in effect on the day before the date of the
19 enactment of this title, shall continue to be in effect with respect to human drug applications and
20 supplements (as defined in such part as of such day) that on or after October 1, 2012, but before
21 October 1, 2017, were accepted by the Food and Drug Administration for filing with respect to
22 assessing and collecting any fee required by such part for a fiscal year prior to fiscal year 2018.

23 **TITLE II—FEES RELATING TO DEVICES**

24 **SEC. 201. SHORT TITLE; FINDINGS.**

25 (a) Short Title.—This title may be cited as the “Medical Device User Fee Amendments of
26 2017”.

27 (b) Findings.—The Congress finds that the fees authorized under the amendments made by
28 this title will be dedicated toward expediting the process for the review of device applications
29 and for assuring the safety and effectiveness of devices, as set forth in the goals identified for
30 purposes of part 3 of subchapter C of chapter VII of the Federal Food, Drug, and Cosmetic Act
31 in the letters from the Secretary of Health and Human Services to the Chairman of the
32 Committee on Health, Education, Labor, and Pensions of the Senate and the Chairman of the
33 Committee on Energy and Commerce of the House of Representatives, as set forth in the
34 Congressional Record.

35 **SEC. 202. DEFINITIONS.**

36 Section 737 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379i) is amended—

37 (1) by redesignating paragraphs (8) through (13) as paragraphs (9) through (14),

1 respectively;

2 (2) by inserting after paragraph (7) the following new paragraph:

3 “(8) The term ‘de novo classification request’ means a request made under section
4 513(f)(2)(A) with respect to the classification of a device.”;

5 (3) in subparagraph (D) of paragraph (10) (as redesignated by paragraph (1)), by striking
6 “and submissions” and inserting “submissions, and de novo classification requests”; and

7 (4) in paragraph (11) (as redesignated by paragraph (1)), by striking “2011” and inserting
8 “2016”.

9 SEC. 203. AUTHORITY TO ASSESS AND USE DEVICE 10 FEES.

11 (a) Types of Fees.—Section 738(a) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C.
12 379j(a)) is amended—

13 (1) in paragraph (1), by striking “fiscal year 2013” and inserting “fiscal year 2018”; and

14 (2) in paragraph (2)—

15 (A) in subparagraph (A)—

16 (i) in the matter preceding clause (i), by striking “October 1, 2012” and
17 inserting “October 1, 2017”;

18 (ii) in clause (viii), by striking “2” and inserting “3.4”; and

19 (iii) by adding at the end the following new clause:

20 “(xi) For a de novo classification request, a fee equal to 30 percent of the fee
21 that applies under clause (i).”; and

22 (B) in subparagraph (B)(v)(I), by striking “or premarket notification submission”
23 and inserting “premarket notification submission, or de novo classification request”.

24 (b) Fee Amounts.—Section 738(b) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C.
25 379j(b)) is amended to read as follows:

26 “(b) Fee Amounts.—

27 “(1) IN GENERAL.—Subject to subsections (c), (d), (e), and (h), for each of fiscal years
28 2018 through 2022, fees under subsection (a) shall be derived from the base fee amounts
29 specified in paragraph (2), to generate the total revenue amounts specified in paragraph (3).

30 “(2) BASE FEE AMOUNTS SPECIFIED.—For purposes of paragraph (1), the base fee amounts
31 specified in this paragraph are as follows:6,L2(4,4,4,4,0,0),tp0,p8,8/8,s60,8,8,8,8,8

32 1“Fee Type1Fiscal Year 20181Fiscal Year 20191Fiscal Year 20201Fiscal Year 20211Fiscal
33 Year 2022

34 Premarket Application\$294,000\$300,000\$310,000\$328,000\$329,000

35 Establishment Registration\$4,375\$4,548\$4,760\$4,975\$4,978

36 “(3) TOTAL REVENUE AMOUNTS SPECIFIED.—For purposes of paragraph (1), the total

1 revenue amounts specified in this paragraph are as follows:

2 “(A) \$183,280,756 for fiscal year 2018.

3 “(B) \$190,654,875 for fiscal year 2019.

4 “(C) \$200,132,014 for fiscal year 2020.

5 “(D) \$211,748,789 for fiscal year 2021.

6 “(E) \$213,687,660 for fiscal year 2022.”.

7 (c) Annual Fee Setting; Adjustments.—Section 738(c) of the Federal Food, Drug, and
8 Cosmetic Act (21 U.S.C. 379j(c)) is amended—

9 (1) in paragraph (1), by striking “2012” and inserting “2017”;

10 (2) in paragraph (2)—

11 (A) in subparagraph (A), by striking “2014” and inserting “2018”;

12 (B) by striking subparagraph (B) and inserting the following new subparagraph:

13 “(B) APPLICABLE INFLATION ADJUSTMENT.—The applicable inflation adjustment for
14 fiscal year 2018 and each subsequent fiscal year is the product of—

15 “(i) the base inflation adjustment under subparagraph (C) for such fiscal year;
16 and

17 “(ii) the product of the base inflation adjustment under subparagraph (C) for
18 each of the fiscal years preceding such fiscal year, beginning with fiscal year
19 2016.”;

20 (C) in subparagraph (C), in the heading, by striking “TO TOTAL REVENUE AMOUNTS”;
21 and

22 (D) by amending subparagraph (D) to read as follows:

23 “(D) ADJUSTMENT TO BASE FEE AMOUNTS.—For each of fiscal years 2018 through
24 2022, the Secretary shall—

25 “(i) adjust the base fee amounts specified in subsection (b)(2) for such fiscal
26 year by multiplying such amounts by the applicable inflation adjustment under
27 subparagraph (B) for such year; and

28 “(ii) if the Secretary determines necessary, increase (in addition to the
29 adjustment under clause (i)) such base fee amounts, on a uniform proportionate
30 basis, to generate the total revenue amounts under subsection (b)(3), as adjusted
31 for inflation under subparagraph (A).”; and

32 (3) in paragraph (3)—

33 (A) by striking “2014 through 2017” and inserting “2018 through 2022”; and

34 (B) by striking “further adjusted” and inserting “increased”.

35 (d) Small Businesses; Fee Waiver and Fee Reduction Regarding Premarket Approval Fees.—
36 Section 738(d) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379j(d)) is amended—

1 (1) in paragraph (1), by striking “specified in clauses (i) through (v) and clauses (vii),
2 (ix), and (x)” and inserting “specified in clauses (i) through (vii) and clauses (ix), (x), and
3 (xi)”; and

4 (2) in paragraph (2)(C)—

5 (A) by striking “supplement, or” and inserting “supplement,”; and

6 (B) by inserting “, or a de novo classification request” after “class III device”.

7 (e) Small Businesses; Fee Reduction Regarding Premarket Notification Submissions.—
8 Section 738(e)(2)(C) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379j(e)(2)(C)) is
9 amended by striking “50” and inserting “25”.

10 (f) Fee Waiver or Reduction.—

11 (1) REPEAL.—Section 738 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379j)
12 is amended by striking subsection (f).

13 (2) CONFORMING CHANGES.—

14 (A) Section 515(c)(4)(A) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C.
15 360e(c)(4)(A)) is amended by striking “738(h)” and inserting “738(g)”.

16 (B) Section 738 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379j), as
17 amended by paragraph (1), is further amended—

18 (i) by redesignating subsections (g) through (l) as subsections (f) through (k);

19 (ii) in subsection (a)(2)(A), by striking “(d), (e), and (f)” and inserting “(d) and
20 (e)”; and

21 (iii) in subsection (a)(3)(A), by striking “and subsection (f)”.

22 (g) Effect of Failure to Pay Fees.—Subsection (f)(1), as redesignated, of section 738 of the
23 Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379j) is amended—

24 (1) by striking “or periodic reporting concerning a class III device” and inserting
25 “periodic reporting concerning a class III device, or de novo classification request”; and

26 (2) by striking “all fees” and inserting “all such fees”.

27 (h) Conditions.—Subsection (g)(1)(A), as redesignated, of section 738 of the Federal Food,
28 Drug, and Cosmetic Act (21 U.S.C. 379j) is amended by striking “\$280,587,000” and inserting
29 “\$320,825,000”.

30 (i) Crediting and Availability of Fees.—Subsection (h), as redesignated, of section 738 of the
31 Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379j) is amended—

32 (1) in paragraph (3)—

33 (A) by striking “2013 through 2017” and inserting “2018 through 2022”; and

34 (B) by striking “subsection (c)” and all that follows through the period at the end
35 and inserting “subsection (c).”; and

36 (2) by striking paragraph (4).

1 SEC. 204. REAUTHORIZATION; REPORTING 2 REQUIREMENTS.

3 (a) Performance Reports.—Section 738A(a) of the Federal Food, Drug, and Cosmetic Act (21
4 U.S.C. 379j-1(a)) is amended—

5 (1) in paragraph (1)—

6 (A) in subparagraph (A)—

7 (i) by striking “2013” and inserting “2018”; and

8 (ii) by striking “the Medical Device User Fee Amendments of 2012” and
9 inserting “Medical Device User Fee Amendments of 2017”; and

10 (B) in subparagraph (B), by striking “the Medical Device User Fee Amendments of
11 2012” and inserting “Medical Device User Fee Amendments of 2017”; and

12 (2) in paragraph (2), by striking “2013 through 2017” and inserting “2018 through 2022”.

13 (b) Reauthorization.—Section 738A(b) of the Federal Food, Drug, and Cosmetic Act (21
14 U.S.C. 379j-1(b)) is amended—

15 (1) in paragraph (1), by striking “2017” and inserting “2022”; and

16 (2) in paragraph (5), by striking “2017” and inserting “2022”.

17 SEC. 205. CONFORMITY ASSESSMENT PILOT 18 PROGRAM.

19 (a) In General.—Section 514 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360d) is
20 amended by adding at the end the following:

21 “(d) Pilot Accreditation Scheme for Conformity Assessment.—

22 “(1) IN GENERAL.—The Secretary shall establish a pilot program under which—

23 “(A) testing laboratories may be accredited, by accreditation bodies meeting criteria
24 specified by the Secretary, to assess the conformance of a device with certain standards
25 recognized under this section; and

26 “(B) subject to paragraph (2), determinations by testing laboratories so accredited
27 that a device conforms with such standard or standards shall be accepted by the
28 Secretary for purposes of demonstrating such conformity under this section unless the
29 Secretary finds that a particular such determination shall not be so accepted.

30 “(2) SECRETARIAL REVIEW OF ACCREDITED LABORATORY DETERMINATIONS.—The
31 Secretary may—

32 “(A) review determinations by testing laboratories accredited pursuant to this
33 subsection, including by conducting periodic audits of such determinations or
34 processes of accredited bodies or testing laboratories and, following such review,
35 taking additional measures under this Act, such as suspension or withdrawal of
36 accreditation of such testing laboratory under paragraph (1)(A) or requesting additional
37 information with respect to such device, as the Secretary determines appropriate; and

1 “(B) if the Secretary becomes aware of information materially bearing on safety or
2 effectiveness of a device assessed for conformity by a testing laboratory so accredited,
3 take such additional measures under this Act as the Secretary determines appropriate,
4 such as suspension or withdrawal of accreditation of such testing laboratory under
5 paragraph (1)(A), or requesting additional information with regard to such device.

6 “(3) IMPLEMENTATION AND REPORTING.—

7 “(A) PUBLIC MEETING.—The Secretary shall publish in the Federal Register a notice
8 of a public meeting to be held no later than September 30, 2018, to discuss and obtain
9 input and recommendations from stakeholders regarding the goals and scope of, and a
10 suitable framework and procedures and requirements for, the pilot program under this
11 subsection.

12 “(B) PILOT PROGRAM GUIDANCE.—The Secretary shall—

13 “(i) not later than September 30, 2019, issue draft guidance regarding the goals
14 and implementation of the pilot program under this subsection; and

15 “(ii) not later than September 30, 2021, issue final guidance with respect to the
16 implementation of such program.

17 “(C) PILOT PROGRAM INITIATION.—Not later than September 30, 2020, the Secretary
18 shall initiate the pilot program under this subsection.

19 “(D) REPORT.—The Secretary shall make available on the website of the Food and
20 Drug Administration an annual report on the progress of the pilot program under this
21 subsection.

22 “(4) SUNSET.—As of October 1, 2022—

23 “(A) the authority for accreditation bodies to accredit testing laboratories pursuant to
24 paragraph (1)(A) shall cease to have force or effect;

25 “(B) the Secretary—

26 “(i) may not accept a determination pursuant to paragraph (1)(B) made by a
27 testing laboratory after such date; and

28 “(ii) may accept such a determination made prior to such date;

29 “(C) except for purposes of accepting a determination described in subparagraph
30 (B)(ii), the Secretary shall not continue to recognize the accreditation of testing
31 laboratories accredited under paragraph (1)(A); and

32 “(D) the Secretary may take actions in accordance with paragraph (2) with respect to
33 the determinations made prior to such date and recognition of the accreditation of
34 testing laboratories pursuant to determinations made prior to such date.”.

35 SEC. 206. REAUTHORIZATION OF REVIEW.

36 Section 523 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360m) is amended—

37 (1) in subsection (a)(3)—

38 (A) in subparagraph (A), by striking clauses (ii) and (iii) and inserting the following:

1 “(ii) a device classified under section 513(f)(2) or designated under section
2 515C(d); or

3 “(iii) a device that is of a type, or subset of a type, listed as not eligible for
4 review under subparagraph (B)(iii).”;

5 (B) by striking subparagraph (B) and inserting the following:

6 “(B) DESIGNATION FOR REVIEW.—The Secretary shall—

7 “(i) issue draft guidance on the factors the Secretary will use in determining
8 whether a class I or class II device type, or subset of such device types, is eligible
9 for review by an accredited person, including—

10 “(I) the risk of the device type, or subset of such device type; and

11 “(II) whether the device type, or subset of such device type, is
12 permanently implantable, life sustaining, or life supporting;

13 “(ii) not later than 24 months after the date on which the Secretary issues such
14 draft guidance, finalize such guidance; and

15 “(iii) beginning on the date such guidance is finalized, designate and post on
16 the Internet website of the Food and Drug Administration, an updated list of class
17 I and class II device types, or subsets of such device types, and the Secretary’s
18 determination with respect to whether each such device type, or subset of a device
19 type, is eligible or not eligible for review by an accredited person under this
20 section based on the factors described in clause (i).”; and

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

22 “(C) INTERIM RULE.—Until the date on which the updated list is designated and
23 posted in accordance with subparagraph (B)(iii), the list in effect on the date of
24 enactment the Medical Device User Fee Amendments of 2017 shall be in effect.”;

25 (2) in subsection (b)—

26 (A) in paragraph (2)—

27 (i) by striking subparagraph (D); and

28 (ii) by redesignating subparagraph (E) as subparagraph (D); and

29 (B) in paragraph (3)—

30 (i) by redesignating subparagraph (E) as subparagraph (F);

31 (ii) in subparagraph (F) (as so redesignated), by striking “The operations of”
32 and all that follows through “it will—” and inserting “Such person shall agree, at
33 a minimum, to include in its request for accreditation a commitment to, at the time
34 of accreditation, and at any time it is performing any review pursuant to this
35 section—”; and

36 (iii) by inserting after subparagraph (D) the following new subparagraph:

37 “(E) The operations of such person shall be in accordance with generally accepted
38 professional and ethical business practices.”; and

1 (3) in subsection (c), by striking “2017” and inserting “2022”.

2 **SEC. 207. ELECTRONIC FORMAT FOR SUBMISSIONS.**

3 Section 745A(b) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379k–1(b)) is
4 amended by adding at the end the following new paragraph:

5 “(3) PRESUBMISSIONS AND SUBMISSIONS SOLELY IN ELECTRONIC FORMAT.—

6 “(A) IN GENERAL.—Beginning on October 1, 2021 (or such later date as may be
7 specified by the Secretary under subparagraph (B)), presubmissions and submissions
8 for devices described in paragraph (1) (and any appeals of action taken by the
9 Secretary with respect to such presubmissions or submissions) shall be submitted
10 solely in such electronic format as specified by the Secretary in guidance issued under
11 subparagraph (C).

12 “(B) EXTENSION.—The Secretary may, if the Secretary determines an extension of
13 the date specified in subparagraph (A) is necessary for the development and adoption
14 of the electronic format referred to in such paragraph, extend such date until such later
15 date as the Secretary may specify, but in no event later than April 1, 2023.

16 “(C) GUIDANCE.—The Secretary shall, not later than January 1, 2021, or such later
17 date as may be specified by the Secretary under subparagraph (B), issue guidance
18 providing for—

19 “(i) any further standards for the submission by electronic format required
20 under subparagraph (A);

21 “(ii) a timetable for the establishment by the Secretary of such further
22 standards; and

23 “(iii) set forth criteria for waivers of and exemptions from the requirements of
24 this subsection.”.

25 **SEC. 208. SAVINGS CLAUSE.**

26 Notwithstanding the amendments made by this title, part 3 of subchapter C of chapter VII of
27 the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379i et seq.), as in effect on the day before
28 the date of the enactment of this title, shall continue to be in effect with respect to the
29 submissions listed in section 738(a)(2)(A) of such Act (as defined in such part as of such day)
30 that on or after October 1, 2012, but before October 1, 2017, were accepted by the Food and
31 Drug Administration for filing with respect to assessing and collecting any fee required by such
32 part for a fiscal year prior to fiscal year 2018.

33 **SEC. 209. EFFECTIVE DATE.**

34 The amendments made by this title shall take effect on October 1, 2017, or the date of the
35 enactment of this Act, whichever is later, except that fees under part 3 of subchapter C of chapter
36 VII of the Federal Food, Drug, and Cosmetic Act shall be assessed for all submissions listed in
37 section 738(a)(2)(A) of such Act received on or after October 1, 2017, regardless of the date of
38 the enactment of this Act.

1 SEC. 210. SUNSET CLAUSE.

2 (a) Authorization.—Sections 737 and 738 of the Federal Food, Drug, and Cosmetic Act (21
3 U.S.C. 739i; 739j) shall cease to be effective October 1, 2022.

4 (b) Reporting Requirements.—Section 738A (21 U.S.C. 739j–1) of the Federal Food, Drug,
5 and Cosmetic Act (regarding reauthorization and reporting requirements) shall cease to be
6 effective January 31, 2023.

7 (c) Previous Sunset Provision.—

8 (1) IN GENERAL.—Effective October 1, 2017, section 207(a) of the Medical Device User
9 Fee Amendments of 2012 (Public Law 112–144) is repealed.

10 (2) CONFORMING AMENDMENT.—The Food and Drug Administration Safety and
11 Innovation Act (Public Law 112–144) is amended in the table of contents in section 2 by
12 striking the item relating to section 207.

13 TITLE III—FEES RELATING TO GENERIC DRUGS

14 SEC. 301. SHORT TITLE; FINDING.

15 (a) Short Title.—This title may be cited as the “Generic Drug User Fee Amendments of
16 2017”.

17 (b) Finding.—The Congress finds that the fees authorized by the amendments made in this
18 title will be dedicated to human generic drug activities, as set forth in the goals identified for
19 purposes of part 7 of subchapter C of chapter VII of the Federal Food, Drug, and Cosmetic Act,
20 in the letters from the Secretary of Health and Human Services to the Chairman of the
21 Committee on Health, Education, Labor, and Pensions of the Senate and the Chairman of the
22 Committee on Energy and Commerce of the House of Representatives, as set forth in the
23 Congressional Record.

24 SEC. 302. DEFINITIONS.

25 Section 744A of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379j–41) is amended—

26 (1) in paragraph (1)(B), by striking “application for a positron emission tomography
27 drug.” and inserting “application—

28 “(i) for a positron emission tomography drug; or

29 “(ii) submitted by a State or Federal governmental entity for a drug that is not
30 distributed commercially.”; and

31 (2) by redesignating paragraphs (5) through (12) as paragraphs (6) through (13),
32 respectively; and

33 (3) by inserting after paragraph (4) the following:

34 “(5) The term ‘contract manufacturing organization facility’ means a manufacturing
35 facility of a finished dosage form of a drug approved pursuant to an abbreviated new drug
36 application, where such manufacturing facility is not identified in an approved abbreviated
37 new drug application held by the owner of such facility or an affiliate of such owner or

1 facility.”.

2 **SEC. 303. AUTHORITY TO ASSESS AND USE HUMAN**
3 **GENERIC DRUG FEES.**

4 (a) Types of Fees.—Section 744B(a) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C.
5 379j-42(a)) is amended—

6 (1) in the matter preceding paragraph (1), by striking “fiscal year 2013” and inserting
7 “fiscal year 2018”;

8 (2) in paragraph (1), by adding at the end the following:

9 “(E) SUNSET.—This paragraph shall cease to be effective October 1, 2022.”.

10 (3) in paragraph (2)—

11 (A) by amending subparagraph (C) to read as follows:

12 “(C) NOTICE.—Not later than 60 days before the start of each of fiscal years 2018
13 through 2022, the Secretary shall publish in the Federal Register the amount of the
14 drug master file fee established by this paragraph for such fiscal year.”; and

15 (B) in subparagraph (E)—

16 (i) in clause (i)—

17 (I) by striking “no later than the date” and inserting “on the earlier of—

18 “(I) the date”;

19 (II) by striking the period and inserting “; or”; and

20 (III) by adding at the end the following:

21 “(II) the date on which the drug master file holder requests the initial
22 completeness assessment.”; and

23 (ii) in clause (ii), by striking “notice provided for in clause (i) or (ii) of
24 subparagraph (C), as applicable” and inserting “notice provided for in
25 subparagraph (C)”;

26 (4) in paragraph (3)—

27 (A) in the heading, by striking “AND PRIOR APPROVAL SUPPLEMENT”;

28 (B) in subparagraph (A), by striking “or a prior approval supplement to an
29 abbreviated new drug application”;

30 (C) by amending subparagraphs (B) and (C) to read as follows:

31 “(B) NOTICE.—Not later than 60 days before the start of each of fiscal years 2018
32 through 2022, the Secretary shall publish in the Federal Register the amount of the fees
33 under subparagraph (A) for such fiscal year.

34 “(C) FEE DUE DATE.—The fees required by subparagraphs (A) and (F) shall be due
35 no later than the date of submission of the abbreviated new drug application or prior
36 approval supplement for which such fee applies.”;

1 (D) in subparagraph (D)—

2 (i) in the heading, by inserting “, IS WITHDRAWN PRIOR TO BEING RECEIVED, OR
3 IS NO LONGER RECEIVED” after “RECEIVED”;

4 (ii) by striking “The Secretary shall” and all that follows through the period and
5 inserting the following:

6 “(i) APPLICATIONS NOT CONSIDERED TO HAVE BEEN RECEIVED AND
7 APPLICATIONS WITHDRAWN PRIOR TO BEING RECEIVED.—The Secretary shall
8 refund 75 percent of the fee paid under subparagraph (A) for any abbreviated new
9 drug application that the Secretary considers not to have been received within the
10 meaning of section 505(j)(5)(A) for a cause other than failure to pay fees, or that
11 has been withdrawn prior to being received within the meaning of section
12 505(j)(5)(A).

13 “(ii) APPLICATIONS NO LONGER RECEIVED.—The Secretary shall refund 100
14 percent of the fee paid under subparagraph (A) for any abbreviated new drug
15 application if the Secretary initially receives the application under section
16 505(j)(5)(A) and subsequently determines that an exclusivity period for a listed
17 drug should have prevented the Secretary from receiving such application, such
18 that the abbreviated new drug application is no longer received within the
19 meaning of section 505(j)(5)(A).”;

20 (E) in subparagraph (E), by striking “or prior approval supplement”; and

21 (F) in the matter preceding clause (i) of subparagraph (F)—

22 (i) by striking “2012” and inserting “2017”; and

23 (ii) by striking “subsection (d)(3)” and inserting “subsection (d)(2)”;

24 (5) in paragraph (4)—

25 (A) in subparagraph (A)—

26 (i) in the matter preceding clause (i) and in clause (iii), by striking “, or
27 intended to be identified, in at least one generic drug submission that is pending
28 or” and inserting “in at least one generic drug submission that is”;

29 (ii) in clause (i), by striking “or intended to be identified in at least one generic
30 drug submission that is pending or” and inserting “in at least one generic drug
31 submission that is”;

32 (iii) in clause (ii), by striking “produces,” and all that follows through “such a”
33 and inserting “is identified in at least one generic drug submission in which the
34 facility is approved to produce one or more active pharmaceutical ingredients or
35 in a Type II active pharmaceutical ingredient drug master file referenced in at
36 least one such”; and

37 (iv) in clause (iii), by striking “to fees under both such clauses” and inserting
38 “only to the fee attributable to the manufacture of the finished dosage forms”; and

39 (B) by amending subparagraphs (C) and (D) to read as follows:

40 “(C) NOTICE.—Within the timeframe specified in subsection (d)(1), the Secretary

1 shall publish in the Federal Register the amount of the fees under subparagraph (A) for
2 such fiscal year.”.

3 “(D) FEE DUE DATE.—For each of fiscal years 2018 through 2022, the fees under
4 subparagraph (A) for such fiscal year shall be due on the later of—

5 “(i) the first business day on or after October 1 of each such year; or

6 “(ii) the first business day after the enactment of an appropriations Act
7 providing for the collection and obligation of fees for such year under this section
8 for such year.”;

9 (6) by redesignating paragraph (5) as paragraph (6); and

10 (7) by inserting after paragraph (4) the following:

11 “(5) GENERIC DRUG APPLICANT PROGRAM FEE.—

12 “(A) IN GENERAL.—A generic drug applicant program fee shall be assessed annually
13 as described in subsection (b)(2)(E).

14 “(B) AMOUNT.—The amount of fees established under subparagraph (A) shall be
15 established under subsection (d).

16 “(C) NOTICE.—Within the timeframe specified in subsection (d)(1), the Secretary
17 shall publish in the Federal Register the amount of the fees under subparagraph (A) for
18 such fiscal year.

19 “(D) FEE DUE DATE.—For each of fiscal years 2018 through 2022, the fees under
20 subparagraph (A) for such fiscal year shall be due on the later of—

21 “(i) the first business day on or after October 1 of each such fiscal year; or

22 “(ii) the first business day after the date of enactment of an appropriations Act
23 providing for the collection and obligation of fees for such fiscal year under this
24 section for such fiscal year.”.

25 (b) Fee Revenue Amounts.—Section 744B(b) of the Federal Food, Drug, and Cosmetic Act
26 (21 U.S.C. 379j-42(b)) is amended—

27 (1) in paragraph (1)—

28 (A) in subparagraph (A)—

29 (i) in the heading, by striking “2013” and inserting “2018”;

30 (ii) by striking “2013” and inserting “2018”;

31 (iii) by striking “\$299,000,000” and inserting “\$493,600,000”; and

32 (iv) by striking “Of that amount” and all that follows through the end of clause
33 (ii); and

34 (B) in subparagraph (B)—

35 (i) in the heading, by striking “2014 THROUGH 2017” and inserting “2019
36 THROUGH 2022”;

37 (ii) by striking “2014 through 2017” and inserting “2019 through 2022”;

1 (iii) by striking “paragraphs (2) through (4)” and inserting “paragraphs (2)
2 through (5)”; and

3 (iv) by striking “\$299,000,000” and inserting “\$493,600,000”; and

4 (2) in paragraph (2)—

5 (A) in the matter preceding subparagraph (A)—

6 (i) by striking “paragraph (1)(A)(ii) for fiscal year 2013 and paragraph (1)(B)
7 for each of fiscal years 2014 through 2017” and inserting “such paragraph for a
8 fiscal year”; and

9 (ii) by striking “through (4)” and inserting “through (5)”;

10 (B) in subparagraph (A), by striking “Six percent” and inserting “Five percent”;

11 (C) by amending subparagraphs (B) and (C) to read as follows:

12 “(B) Thirty-three percent shall be derived from fees under subsection (a)(3) (relating
13 to abbreviated new drug applications).

14 “(C) Twenty percent shall be derived from fees under subsection (a)(4)(A)(i)
15 (relating to generic drug facilities). The amount of the fee for a contract manufacturing
16 organization facility shall be equal to one-third the amount of the fee for a facility that
17 is not a contract manufacturing organization facility. The amount of the fee for a
18 facility located outside the United States and its territories and possessions shall be
19 \$15,000 higher than the amount of the fee for a facility located in the United States and
20 its territories and possessions.”;

21 (D) in subparagraph (D)—

22 (i) by striking “Fourteen percent” and inserting “Seven percent”;

23 (ii) by striking “not less than \$15,000 and not more than \$30,000” and inserting
24 “\$15,000”; and

25 (iii) by striking “, as determined” and all that follows through the period at the
26 end and inserting a period; and

27 (E) by adding at the end the following:

28 “(E)(i) Thirty-five percent shall be derived from fees under subsection (a)(5)
29 (relating to generic drug applicant program fees). For purposes of this subparagraph, if
30 a person has affiliates, a single program fee shall be assessed with respect to that
31 person, including its affiliates, and may be paid by that person or any one of its
32 affiliates. The Secretary shall determine the fees as follows:

33 “(I) If a person (including its affiliates) owns at least one but not more than 5
34 approved abbreviated new drug applications on the due date for the fee under this
35 subsection, the person (including its affiliates) shall be assessed a small business
36 generic drug applicant program fee equal to one-tenth of the large size operation
37 generic drug applicant program fee.

38 “(II) If a person (including its affiliates) owns at least 6 but not more than 19
39 approved abbreviated new drug applications on the due date for the fee under this

1 subsection, the person (including its affiliates) shall be assessed a medium size
2 operation generic drug applicant program fee equal to two-fifths of the large size
3 operation generic drug applicant program fee.

4 “(III) If a person (including its affiliates) owns 20 or more approved
5 abbreviated new drug applications on the due date for the fee under this
6 subsection, the person (including its affiliates) shall be assessed a large size
7 operation generic drug applicant program fee.

8 “(ii) For purposes of this subparagraph, an abbreviated new drug application shall be
9 deemed not to be approved if the applicant has submitted a written request for
10 withdrawal of approval of such abbreviated new drug application by April 1 of the
11 previous fiscal year.”.

12 (c) Adjustments.—Section 744B(c) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C.
13 379j-42(c)) is amended—

14 (1) in paragraph (1)—

15 (A) by striking “2014” and inserting “2019”;

16 (B) by inserting “to equal the product of the total revenues established in such notice
17 for the prior fiscal year multiplied” after “a fiscal year,”; and

18 (C) by striking the flush text following subparagraph (C); and

19 (2) in paragraph (2)—

20 (A) by striking “2017” each place it appears and inserting “2022”; and

21 (B) by striking “2018” and inserting “2023”.

22 (d) Annual Fee Setting.—Section 744B of the Federal Food, Drug, and Cosmetic Act (21
23 U.S.C. 379j-42) is amended—

24 (1) in subsection (c)(2), by striking “Such fees may only be used in fiscal year 2018.”;

25 (2) in subsection (d)—

26 (A) by striking paragraphs (1) and (2) and inserting the following:

27 “(1) FISCAL YEARS 2018 THROUGH 2022.—Not more than 60 days before the first day of
28 each of fiscal years 2018 through 2022, the Secretary shall establish the fees described in
29 paragraphs (2) through (5) of subsection (a), based on the revenue amounts established
30 under subsection (b) and the adjustments provided under subsection (c).”;

31 (B) by redesignating paragraph (3) as paragraph (2); and

32 (C) in paragraph (2) (as so redesignated), in the matter preceding subparagraph (A),
33 by striking “fees under paragraphs (1) and (2)” and inserting “fee under paragraph (1)”.

34 (e) Identification of Facilities.—Section 744B(f) of the Federal Food, Drug, and Cosmetic Act
35 (21 U.S.C. 379j-42(f)) is amended—

36 (1) by striking paragraph (1);

37 (2) by redesignating paragraphs (2) through (4) as paragraphs (1) through (3),
38 respectively;

- 1 (3) in paragraph (1) (as so redesignated)—
2 (A) by striking “paragraph (4)” and inserting “paragraph (3)”; and
3 (B) by striking “Such information shall” and all that follows through the end of
4 subparagraph (B) and inserting “Such information shall, for each fiscal year, be
5 submitted, updated, or reconfirmed on or before June 1 of the previous fiscal year.”;
6 and
7 (4) in paragraph (2), as so redesignated—
8 (A) in the heading, by striking “CONTENTS OF NOTICE” and inserting “INFORMATION
9 REQUIRED TO BE SUBMITTED”;
10 (B) in the matter preceding subparagraph (A), by striking “paragraph (2)” and
11 inserting “paragraph (1)”;
12 (C) in subparagraph (A), by striking “or intended to be identified”;
13 (D) in subparagraph (D), by striking “and” at the end;
14 (E) in subparagraph (E), by striking the period and inserting “; and”; and
15 (F) by adding at the end the following:
16 “(F) whether the facility is a contract manufacturing organization facility.”.
17 (f) Effect of Failure to Pay Fees.—Section 744B(g) of the Federal Food, Drug, and Cosmetic
18 Act (21 U.S.C. 379–42(g)) is amended—
19 (1) in paragraph (1), by adding at the end the following: “This paragraph shall cease to be
20 effective on October 1, 2022.”.
21 (2) in paragraph (2)(C)(ii), by striking “of 505(j)(5)(A)” and inserting “of section
22 505(j)(5)(A)”;
23 (3) by adding at the end the following:
24 “(5) GENERIC DRUG APPLICANT PROGRAM FEE.—
25 “(A) IN GENERAL.—A person who fails to pay a fee as required under subsection
26 (a)(5) by the date that is 20 calendar days after the due date, as specified in
27 subparagraph (D) of such subsection, shall be subject to the following:
28 “(i) The Secretary shall place the person on a publicly available arrears list.
29 “(ii) Any abbreviated new drug application submitted by the generic drug
30 applicant or an affiliate of such applicant shall not be received, within the
31 meaning of section 505(j)(5)(A).
32 “(iii) All drugs marketed pursuant to any abbreviated new drug application held
33 by such applicant or an affiliate of such applicant shall be deemed misbranded
34 under section 502(aa).
35 “(B) APPLICATION OF PENALTIES.—The penalties under subparagraph (A) shall
36 apply until the fee required under subsection (a)(5) is paid.”.
37 (g) Limitations.—Section 744B(h)(2) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C.

1 379–42(h)(2)) is amended by striking “for Type II active pharmaceutical ingredient drug master
2 files, abbreviated new drug applications and prior approval supplements, and generic drug
3 facilities and active pharmaceutical ingredient facilities”.

4 (h) Crediting and Availability of Fees.—Section 744B(i) of the Federal Food, Drug, and
5 Cosmetic Act (21 U.S.C. 379–42(i)) is amended—

6 (1) in paragraph (2)—

7 (A) by striking subparagraph (C) (relating to fee collection during first program
8 year);

9 (B) in subparagraph (D)—

10 (i) in the heading, by striking “IN SUBSEQUENT YEARS”; and

11 (ii) by striking “(after fiscal year 2013)”; and

12 (C) by redesignating subparagraph (D) as subparagraph (C); and

13 (2) in paragraph (3), by striking “fiscal years 2013 through 2017” and inserting “fiscal
14 years 2018 through 2022”.

15 (i) Information on Abbreviated New Drug Applications Held by Applicants and Their
16 Affiliates.—Section 744B of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379–42) is
17 amended by adding at the end the following:

18 “(o) Information on Abbreviated New Drug Applications Owned by Applicants and Their
19 Affiliates.—

20 “(1) IN GENERAL.—By April 1 of each year, each person that owns an abbreviated new
21 drug application, or any affiliate of such person, shall submit to the Secretary a list of —

22 “(A) all approved abbreviated new drug applications owned by such person; and

23 “(B) if any affiliate of such person also owns an abbreviated new drug application,
24 all approved abbreviated new drug applications owned by any such affiliate.

25 “(2) FORMAT AND METHOD.—The Secretary shall specify in guidance the format and
26 method for submission of lists under this subsection.”.

27 SEC. 304. REAUTHORIZATION; REPORTING 28 REQUIREMENTS.

29 Section 744C of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379j–43) is amended—

30 (1) in subsection (a)—

31 (A) by striking “2013” and inserting “2018”; and

32 (B) by striking “Generic Drug User Fee Amendments of 2012” and inserting
33 “Generic Drug User Fee Amendments of 2017”;

34 (2) in subsection (b), by striking “2013” and inserting “2018”; and

35 (3) in subsection (d), by striking “2017” each place it appears and inserting “2022”.

1 **SEC. 305. SUNSET DATES.**

2 (a) Authorization.—Sections 744A and 744B of the Federal Food, Drug, and Cosmetic Act
3 (21 U.S.C. 379j–41; 379j–42) shall cease to be effective October 1, 2022.

4 (b) Reporting Requirements.—Section 744C of the Federal Food, Drug, and Cosmetic Act (21
5 U.S.C. 379j–43) shall cease to be effective January 31, 2023.

6 (c) Previous Sunset Provision.—Effective October 1, 2017, subsections (a) and (b) of section
7 304 of the Food and Drug Administration Safety and Innovation Act (Public Law 112–144) are
8 repealed.

9 **SEC. 306. EFFECTIVE DATE.**

10 The amendments made by this title shall take effect on October 1, 2017, or the date of the
11 enactment of this Act, whichever is later, except that fees under part 7 of subchapter C of chapter
12 VII of the Federal Food, Drug, and Cosmetic Act shall be assessed for all abbreviated new drug
13 applications received on or after October 1, 2017, regardless of the date of the enactment of this
14 Act.

15 **SEC. 307. SAVINGS CLAUSE.**

16 Notwithstanding the amendments made by this title, part 7 of subchapter C of chapter VII of
17 the Federal Food, Drug, and Cosmetic Act, as in effect on the day before the date of the
18 enactment of this title, shall continue to be in effect with respect to abbreviated new drug
19 applications (as defined in such part as of such day) that on or after October 1, 2012, but before
20 October 1, 2017, were received by the Food and Drug Administration within the meaning of
21 505(j)(5)(A) of such Act (21 U.S.C. 355(j)(5)(A)), prior approval supplements that were
22 submitted, and drug master files for Type II active pharmaceutical ingredients that were first
23 referenced with respect to assessing and collecting any fee required by such part for a fiscal year
24 prior to fiscal year 2018.

25 **TITLE IV—FEES RELATING TO BIOSIMILAR**
26 **BIOLOGICAL PRODUCTS**

27 **SEC. 401. SHORT TITLE; FINDING.**

28 (a) Short Title.—This title may be cited as the “Biosimilar User Fee Amendments of 2017”.

29 (b) Finding.—The Congress finds that the fees authorized by the amendments made in this
30 title will be dedicated to expediting the process for the review of biosimilar biological product
31 applications, including postmarket safety activities, as set forth in the goals identified for
32 purposes of part 8 of subchapter C of chapter VII of the Federal Food, Drug, and Cosmetic Act,
33 in the letters from the Secretary of Health and Human Services to the Chairman of the
34 Committee on Health, Education, Labor, and Pensions of the Senate and the Chairman of the
35 Committee on Energy and Commerce of the House of Representatives, as set forth in the
36 Congressional Record.

37 **SEC. 402. DEFINITIONS.**

1 (a) Adjustment Factor.—Section 744G(1) of the Federal Food, Drug, and Cosmetic Act (21
2 U.S.C. 379j–51(1)) is amended to read as follows:

3 “(1) The term ‘adjustment factor’ applicable to a fiscal year is the Consumer Price Index
4 for all urban consumers (all items; United States city average) for October of the preceding
5 fiscal year divided by such Index for October 2011.”.

6 (b) Biosimilar Biological Product.—Section 744G(3) of the Federal Food, Drug, and Cosmetic
7 Act (21 U.S.C. 379j–51(3)) is amended by striking “means a product” and inserting “means a
8 specific strength of a biological product in final dosage form”.

9 SEC. 403. AUTHORITY TO ASSESS AND USE 10 BIOSIMILAR FEES.

11 (a) Types of Fees.—Section 744H(a) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C.
12 379j–52(a)) is amended—

13 (1) in the matter preceding paragraph (1), by striking “fiscal year 2013” and inserting
14 “fiscal year 2018”;

15 (2) in the heading of paragraph (1), by striking “BIOSIMILAR” and inserting “BIOSIMILAR
16 BIOLOGICAL PRODUCT”;

17 (3) in paragraph (1)(A)(i), by striking “(b)(1)(A)” and inserting “(c)(5)”;

18 (4) in paragraph (1)(B)(i), by striking “(b)(1)(B) for biosimilar biological product
19 development” and inserting “(c)(5) for the biosimilar biological product development
20 program”;

21 (5) in paragraph (1)(B)(ii), by striking “annual biosimilar biological product development
22 program fee” and inserting “annual biosimilar biological product development fee”;

23 (6) in paragraph (1)(B)(iii), by striking “annual biosimilar development program fee” and
24 inserting “annual biosimilar biological product development fee”;

25 (7) in paragraph (1)(B), by adding at the end the following:

26 “(iv) REFUND.—If a person submits a marketing application for a biosimilar
27 biological product before October 1 of a fiscal year and such application is
28 accepted for filing on or after October 1 of such fiscal year, the person may
29 request a refund equal to the annual biosimilar development fee paid by the
30 person for the product for such fiscal year. To qualify for consideration for a
31 refund under this clause, a person shall submit to the Secretary a written request
32 for such refund not later than 180 days after the marketing application is accepted
33 for filing.”;

34 (8) in paragraph (1)(C), by striking “for a product effective October 1 of a fiscal year by,”
35 and inserting “for a product, effective October 1 of a fiscal year, by,”;

36 (9) in paragraph (1)(D)—

37 (A) in clause (i) in the matter preceding subclause (I), by inserting “, if the person
38 seeks to resume participation in such program,” before “pay a fee”;

39 (B) in clause (i)(I), by inserting after “grants a request” the following: “by such

- 1 person”; and
- 2 (C) in clause (i)(II), by inserting after “discontinued” the following: “by such
3 person”;
- 4 (10) in the heading of paragraph (1)(E), by striking “BIOSIMILAR DEVELOPMENT
5 PROGRAM”;
- 6 (11) in the heading of subparagraph (F) of paragraph (1), by striking “BIOSIMILAR
7 DEVELOPMENT PROGRAM FEES” and inserting “BIOSIMILAR BIOLOGICAL PRODUCT
8 DEVELOPMENT FEES”;
- 9 (12) in paragraph (1)(F)—
- 10 (A) in the heading of subparagraph (F), by striking “BIOSIMILAR DEVELOPMENT
11 PROGRAM” before “FEES”; and
- 12 (B) by amending clause (i) to read as follows:
- 13 “(i) REFUNDS.—Except as provided in subparagraph (B)(iv), the Secretary shall
14 not refund any initial or annual biosimilar biological product development fee
15 paid under subparagraph (A) or (B), or any reactivation fee paid under
16 subparagraph (D).”;
- 17 (13) in paragraph (2)—
- 18 (A) in the heading of paragraph (2), by striking “AND SUPPLEMENT”;
- 19 (B) by amending subparagraphs (A) and (B) to read as follows:
- 20 “(A) IN GENERAL.—Each person that submits, on or after October 1, 2017, a
21 biosimilar biological product application shall be subject to the following fees:
- 22 “(i) A fee established under subsection (c)(5) for a biosimilar biological
23 product application for which clinical data (other than comparative bioavailability
24 studies) with respect to safety or effectiveness are required for approval.
- 25 “(ii) A fee established under subsection (c)(5) for a biosimilar biological
26 product application for which clinical data (other than comparative bioavailability
27 studies) with respect to safety or effectiveness are not required for approval. Such
28 fee shall be equal to half of the amount of the fee described in clause (i).
- 29 “(B) RULE OF APPLICABILITY; TREATMENT OF CERTAIN PREVIOUSLY PAID FEES.—Any
30 person who pays a fee under subparagraph (A), (B), or (D) of paragraph (1) for a
31 product before October 1, 2017, but submits a biosimilar biological product application
32 for that product after such date, shall—
- 33 “(i) be subject to any biosimilar biological product application fees that may be
34 assessed at the time when such biosimilar biological product application is
35 submitted; and
- 36 “(ii) be entitled to no reduction of such application fees based on the amount of
37 fees paid for that product before October 1, 2017 under such subparagraphs (A),
38 (B), or (D).”;
- 39 (C) in the heading of subparagraph (D), by striking “OR SUPPLEMENT”; and

1 (D) in subparagraphs (C) through (F)—

2 (i) by striking “or supplement” each place it appears; and

3 (ii) in subparagraph (D), by striking “or a supplement”;

4 (14) by amending paragraph (3) to read as follows:

5 “(3) BIOSIMILAR BIOLOGICAL PRODUCT PROGRAM FEE.—

6 “(A) IN GENERAL.—Each person who is named as the applicant in a biosimilar
7 biological product application shall pay the annual biosimilar biological product
8 program fee established for a fiscal year under subsection (c)(5) for each biosimilar
9 biological product that—

10 “(i) is identified in such a biosimilar biological product application approved as
11 of October 1 of such fiscal year; and

12 “(ii) as of October 1 of such fiscal year, does not appear on a list, developed
13 and maintained by the Secretary, of discontinued biosimilar biological products.

14 “(B) DUE DATE.—The biosimilar biological product program fee for a fiscal year
15 shall be due on the later of—

16 “(i) the first business day on or after October 1 of each such year; or

17 “(ii) the first business day after the enactment of an appropriations Act
18 providing for the collection and obligation of fees for such year under this section.

19 “(C) ONE FEE PER PRODUCT PER YEAR.—The biosimilar biological product program
20 fee shall be paid only once for each product for each fiscal year.

21 “(D) LIMITATION.—A person who is named as the applicant in a biosimilar
22 biological product application shall not be assessed more than 5 biosimilar biological
23 product program fees for a fiscal year for biosimilar biological products identified in
24 such biosimilar biological product application.”.

25 (b) Fee Revenue Amounts.—Subsection (b) of section 744H of the Federal Food, Drug, and
26 Cosmetic Act (21 U.S.C. 379j–52) is amended to read as follows:

27 “(b) Fee Revenue Amounts.—

28 “(1) FISCAL YEAR 2018.—For fiscal year 2018, fees under subsection (a) shall be
29 established to generate a total revenue amount equal to the sum of—

30 “(A) \$45,000,000; and

31 “(B) the dollar amount equal to the fiscal year 2018 adjustment (as determined under
32 subsection (c)(4)).

33 “(2) SUBSEQUENT FISCAL YEARS.—For each of the fiscal years 2019 through 2022, fees
34 under subsection (a) shall, except as provided in subsection (c), be established to generate a
35 total revenue amount equal to the sum of—

36 “(A) the annual base revenue for the fiscal year (as determined under paragraph (4));

37 “(B) the dollar amount equal to the inflation adjustment for the fiscal year (as
38 determined under subsection (c)(1));

1 “(C) the dollar amount equal to the capacity planning adjustment for the fiscal year
2 (as determined under subsection (c)(2)); and

3 “(D) the dollar amount equal to the operating reserve adjustment for the fiscal year,
4 if applicable (as determined under subsection (c)(3)).

5 “(3) ALLOCATION OF REVENUE AMOUNT AMONG FEES; LIMITATIONS ON FEE AMOUNTS.—

6 “(A) ALLOCATION.—The Secretary shall determine the percentage of the total
7 revenue amount for a fiscal year to be derived from, respectively—

8 “(i) initial and annual biosimilar development fees and reactivation fees under
9 subsection (a)(1);

10 “(ii) biosimilar biological product application fees under subsection (a)(2); and

11 “(iii) Biosimilar biological product program fees under subsection (a)(3).

12 “(B) LIMITATIONS ON FEE AMOUNTS.—Until the first fiscal year for which the
13 capacity planning adjustment under subsection (c)(2) is effective, the amount of any
14 fee under subsection (a) for a fiscal year after fiscal year 2018 shall not exceed 125
15 percent of the amount of such fee for fiscal year 2018.

16 “(C) BIOSIMILAR BIOLOGICAL PRODUCT DEVELOPMENT FEES.—The initial biosimilar
17 biological product development fee under subsection (a)(1)(A) for a fiscal year shall be
18 equal to the annual biosimilar biological product development fee under subsection
19 (a)(1)(B) for that fiscal year.

20 “(D) REACTIVATION FEE.—The reactivation fee under subsection (a)(1)(D) for a
21 fiscal year shall be equal to twice the amount of the annual biosimilar biological
22 product development fee under subsection (a)(1)(B) for that fiscal year.

23 “(4) ANNUAL BASE REVENUE.—For purposes of paragraph (2), the dollar amount of the
24 annual base revenue for a fiscal year shall be the dollar amount of the total revenue amount
25 for the previous fiscal year, excluding any adjustments to such revenue amount under
26 subsection (c)(3).”.

27 (c) Adjustments; Annual Fee Setting.—Section 744H of the Federal Food, Drug, and
28 Cosmetic Act (21 U.S.C. 379j–52) is amended—

29 (1) by redesignating subsections (c) through (h) as subsections (d) through (i),
30 respectively;

31 (2) in subsections (a)(2)(F) and (g), by striking “subsection (c)” and inserting “subsection
32 (d)”;

33 (3) in subsection (a)(4)(A), by striking “subsection (b)(1)(F)” and inserting “subsection
34 (c)(5)”;

35 (4) by inserting after subsection (b) the following:

36 “(c) Adjustments; Annual Fee Setting.—

37 “(1) INFLATION ADJUSTMENT.—

38 “(A) IN GENERAL.—For purposes of subsection (b)(2)(B), the dollar amount of the
39 inflation adjustment to the annual base revenue for each fiscal year shall be equal to

1 the product of—

2 “(i) such annual base revenue for the fiscal year under subsection (b); and

3 “(ii) the inflation adjustment percentage under subparagraph (B).

4 “(B) INFLATION ADJUSTMENT PERCENTAGE.—The inflation adjustment percentage
5 under this subparagraph for a fiscal year is equal to the sum of—

6 “(i) the average annual percent change in the cost, per full-time equivalent
7 position of the Food and Drug Administration, of all personnel compensation and
8 benefits paid with respect to such positions for the first 3 years of the preceding 4
9 fiscal years, multiplied by the proportion of personnel compensation and benefits
10 costs to total costs of the process for the review of biosimilar biological product
11 applications (as defined in section 744G(13)) for the first 3 years of the preceding
12 4 fiscal years; and

13 “(ii) the average annual percent change that occurred in the Consumer Price
14 Index for urban consumers (Washington-Baltimore, DC-MD-VA-WV; Not
15 Seasonally Adjusted; All items; Annual Index) for the first 3 years of the
16 preceding 4 years of available data multiplied by the proportion of all costs other
17 than personnel compensation and benefits costs to total costs of the process for
18 the review of biosimilar biological product applications (as defined in section
19 744G(13)) for the first 3 years of the preceding 4 fiscal years.

20 “(2) CAPACITY PLANNING ADJUSTMENT.—

21 “(A) IN GENERAL.—Beginning with the fiscal year described in subparagraph
22 (B)(ii)(II), the Secretary shall, in addition to the adjustment under paragraph (1),
23 further increase the fee revenue and fees under this section for a fiscal year to reflect
24 changes in the resource capacity needs of the Secretary for the process for the review
25 of biosimilar biological product applications.

26 “(B) CAPACITY PLANNING METHODOLOGY.—

27 “(i) DEVELOPMENT; EVALUATION AND REPORT.—The Secretary shall obtain,
28 through a contract with an independent accounting or consulting firm, a report
29 evaluating options and recommendations for a new methodology to accurately
30 assess changes in the resource and capacity needs of the process for the review of
31 biosimilar biological product applications. The capacity planning methodological
32 options and recommendations presented in such report shall utilize and be
33 informed by personnel time reporting data as an input. The report shall be
34 published for public comment not later than September 30, 2020.

35 “(ii) ESTABLISHMENT AND IMPLEMENTATION.—After review of the report
36 described in clause (i) and receipt and review of public comments thereon, the
37 Secretary shall establish a capacity planning methodology for purposes of this
38 paragraph, which shall—

39 “(I) incorporate such approaches and attributes as the Secretary determines
40 appropriate; and

41 “(II) be effective beginning with the first fiscal year for which fees are set

1 after such capacity planning methodology is established.

2 “(C) LIMITATION.—Under no circumstances shall an adjustment under this
3 paragraph result in fee revenue for a fiscal year that is less than the sum of the amounts
4 under subsections (b)(2)(A) (the annual base revenue for the fiscal year) and (b)(2)(B)
5 (the dollar amount of the inflation adjustment for the fiscal year).

6 “(D) PUBLICATION IN FEDERAL REGISTER.—The Secretary shall publish in the
7 Federal Register notice under paragraph (5) the fee revenue and fees resulting from the
8 adjustment and the methodologies under this paragraph.

9 “(3) OPERATING RESERVE ADJUSTMENT.—

10 “(A) INTERIM APPLICATION; FEE REDUCTION.—Until the first fiscal year for which
11 the capacity planning adjustment under paragraph (2) is effective, the Secretary may,
12 in addition to the adjustment under paragraph (1), reduce the fee revenue and fees
13 under this section for a fiscal year as the Secretary determines appropriate for long-
14 term financial planning purposes.

15 “(B) GENERAL APPLICATION AND METHODOLOGY.—Beginning with the first fiscal
16 year for which the capacity planning adjustment under paragraph (2) is effective, the
17 Secretary may, in addition to the adjustments under paragraphs (1) and (2)—

18 “(i) reduce the fee revenue and fees under this section as the Secretary
19 determines appropriate for long-term financial planning purposes; or

20 “(ii) increase the fee revenue and fees under this section if such an adjustment
21 is necessary to provide for not more than 21 weeks of operating reserves of
22 carryover user fees for the process for the review of biosimilar biological product
23 applications.

24 “(C) FEDERAL REGISTER NOTICE.—If an adjustment under subparagraph (A) or (B) is
25 made, the rationale for the amount of the increase or decrease (as applicable) in fee
26 revenue and fees shall be contained in the annual Federal Register notice under
27 paragraph (5) establishing fee revenue and fees for the fiscal year involved.

28 “(4) FISCAL YEAR 2018 ADJUSTMENT.—

29 “(A) IN GENERAL.—For fiscal year 2018, the Secretary shall adjust the fee revenue
30 and fees under this section in such amount (if any) as needed to reflect an updated
31 assessment of the workload for the process for the review of biosimilar biological
32 product applications.

33 “(B) METHODOLOGY.—The Secretary shall publish under paragraph (5) a
34 description of the methodology used to calculate the fiscal year 2018 adjustment under
35 this paragraph in the Federal Register notice establishing fee revenue and fees for
36 fiscal year 2018.

37 “(C) LIMITATION.—No adjustment under this paragraph shall result in an increase in
38 fee revenue and fees under this section in excess of \$9,000,000.

39 “(5) ANNUAL FEE SETTING.—For fiscal year 2018 and each subsequent fiscal year, the
40 Secretary shall, not later than 60 days before the start of each such fiscal year—

1 “(A) establish, for the fiscal year, initial and annual biosimilar biological product
2 development fees and reactivation fees under subsection (a)(1), biosimilar biological
3 product application fees under subsection (a)(2), and biosimilar biological product
4 program fees under subsection (a)(3), based on the revenue amounts established under
5 subsection (b) and the adjustments provided under this subsection; and

6 “(B) publish such fee revenue and fees in the Federal Register.

7 “(6) LIMIT.—The total amount of fees assessed for a fiscal year under this section may
8 not exceed the total costs for such fiscal year for the resources allocated for the process for
9 the review of biosimilar biological product applications.”.

10 (d) Application Fee Waiver for Small Business.—Subsection (d)(1) of section 744H of the
11 Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379j–52), as redesignated by subsection
12 (c)(1), is amended—

13 (1) by striking subparagraph (B);

14 (2) by striking “shall pay—” and all that follows through “application fees” and inserting
15 “shall pay application fees”; and

16 (3) by striking “; and” at the end and inserting a period.

17 (e) Effect of Failure to Pay Fees.—Subsection (e) of section 744H of the Federal Food, Drug,
18 and Cosmetic Act (21 U.S.C. 379j–52), as redesignated by subsection (c)(1), is amended by
19 striking “all fees” and inserting “all such fees”.

20 (f) Crediting and Availability of Fees.—Subsection (f) of section 744H of the Federal Food,
21 Drug, and Cosmetic Act (21 U.S.C. 379j–52), as redesignated by subsection (c)(1), is
22 amended—

23 (1) in paragraph (2)—

24 (A) by striking subparagraph (C) (relating to fee collection during first program
25 year) and inserting the following:

26 “(C) COMPLIANCE.—The Secretary shall be considered to have met the requirements
27 of subparagraph (B) in any fiscal year if the costs described in such subparagraph are
28 not more than 15 percent below the level specified in such subparagraph.”; and

29 (B) in subparagraph (D)—

30 (i) in the heading, by striking “IN SUBSEQUENT YEARS”; and

31 (ii) by striking “(after fiscal year 2013)”; and

32 (2) in paragraph (3), by striking “2013 through 2017” and inserting “2018 through 2022”.

33 **SEC. 404. REAUTHORIZATION; REPORTING** 34 **REQUIREMENTS.**

35 Section 744I of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379j–53) is amended—

36 (1) in subsection (a)—

37 (A) by striking “2013” and inserting “2018”; and

1 (B) by striking “Biosimilar User Fee Act of 2012” and inserting “Biosimilar User
2 Fee Amendments of 2017”;
3 (2) in subsection (b), by striking “2013” and inserting “2018”;
4 (3) by striking subsection (d);
5 (4) by redesignating subsection (e) as subsection (d); and
6 (5) in subsection (d), as so redesignated, by striking “2017” each place it appears and
7 inserting “2022”.

8 **SEC. 405. SUNSET DATES.**

9 (a) Authorization.—Sections 744G and 744H of the Federal Food, Drug, and Cosmetic Act, as
10 amended by section 403 of this Act, shall cease to be effective October 1, 2022.

11 (b) Reporting Requirements.—Section 744I of the Federal Food, Drug, and Cosmetic Act, as
12 amended by section 404 of this Act, shall cease to be effective January 31, 2023.

13 (c) Previous Sunset Provision.—

14 (1) IN GENERAL.—Effective October 1, 2017, section 404 of the Food and Drug
15 Administration Safety and Innovation Act (Public Law 112–144) is repealed.

16 (2) CONFORMING AMENDMENT.—The Food and Drug Administration Safety and
17 Innovation Act (Public Law 112–144) is amended in the table of contents in section 2 by
18 striking the item relating to section 404.

19 **SEC. 406. EFFECTIVE DATE.**

20 The amendments made by this title shall take effect on October 1, 2017, or the date of the
21 enactment of this Act, whichever is later, except that fees under part 8 of subchapter C of chapter
22 VII of the Federal Food, Drug, and Cosmetic Act shall be assessed for all biosimilar biological
23 product applications received on or after October 1, 2017, regardless of the date of the enactment
24 of this Act.

25 **SEC. 407. SAVINGS CLAUSE.**

26 Notwithstanding the amendments made by this title, part 8 of subchapter C of chapter VII of
27 the Federal Food, Drug, and Cosmetic Act, as in effect on the day before the date of the
28 enactment of this title, shall continue to be in effect with respect to biosimilar biological product
29 applications and supplements (as defined in such part as of such day) that were accepted by the
30 Food and Drug Administration for filing on or after October 1, 2012, but before October 1, 2017,
31 with respect to assessing and collecting any fee required by such part for a fiscal year prior to
32 fiscal year 2018.

33 **TITLE V—REAUTHORIZATION OF OTHER PROGRAMS**

34 **SEC. 501. REAUTHORIZATION OF PROVISION** 35 **RELATING TO EXCLUSIVITY OF CERTAIN DRUGS** 36 **CONTAINING SINGLE ENANTIOMERS.**

1 Section 505(u)(4) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(u)(4)) is
2 amended by striking “2017” and inserting “2022”.

3 **SEC. 502. REAUTHORIZATION OF PEDIATRIC**
4 **HUMANITARIAN DEVICE EXCEPTIONS.**

5 Section 520(m)(6)(A)(iv) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C.
6 360j(m)(6)(A)(iv)) is amended by striking “2017” and inserting “2022”.

7 **SEC. 503. REAUTHORIZATION OF THE CRITICAL PATH**
8 **PUBLIC-PRIVATE PARTNERSHIPS.**

9 Section 566(f) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360bbb-5(f)) is
10 amended by striking “2013 through 2017” and inserting “2018 through 2022”.

11 **SEC. 504. REAUTHORIZATION OF PEDIATRIC DEVICE**
12 **CONSORTIA.**

13 Section 305(e) of Pediatric Medical Device Safety and Improvement Act of 2007 (Public Law
14 110-85; 42 U.S.C. 282 note)) is amended by striking “2013 through 2017” and inserting “2018
15 through 2022”.

16 **SEC. 505. REAUTHORIZATION OF ORPHAN GRANTS**
17 **PROGRAM.**

18 Section 5(c) of the Orphan Drug Act (21 U.S.C. 360ee(c)) is amended by striking “2013
19 through 2017” and inserting “2018 through 2022”.