

In the House of Representatives, U. S.,

February 25, 2019.

Resolved, That the bill from the Senate (S. 483) entitled “An Act to enact into law a bill by reference.”, do pass with the following

AMENDMENT:

Strike out all after the enacting clause and insert:

1 ***SECTION 1. SHORT TITLE; TABLE OF CONTENTS.***

2 (a) *SHORT TITLE.*—*This Act may be cited as the “Pesticide Registration Improvement Extension Act of 2018”.*

3 (b) *TABLE OF CONTENTS.*—*The table of contents for*
4 *this Act is as follows:*

Sec. 1. Short title; table of contents.

Sec. 2. Extension and modification of maintenance fee authority.

Sec. 3. Reregistration and Expedited Processing Fund.

Sec. 4. Experimental use permits for pesticides.

Sec. 5. Pesticide registration service fees.

Sec. 6. Revision of tables regarding covered pesticide registration applications and other covered actions and their corresponding registration service fees.

Sec. 7. Agricultural worker protection standard; certification of pesticide applicators.

6 ***SEC. 2. EXTENSION AND MODIFICATION OF MAINTENANCE***

7 ***FEE AUTHORITY.***

8 (a) *MAINTENANCE FEE.*—*Section 4(i)(1) of the Fed-*
9 *eral Insecticide, Fungicide, and Rodenticide Act (7 U.S.C.*

10 *136a–1(i)(1)) is amended—*

1 (1) *in subparagraph (C), by striking “an aggregate amount of \$27,800,000 for each of fiscal years*
2 *2013 through 2017” and inserting “an average*
3 *amount of \$31,000,000 for each of fiscal years 2019*
4 *through 2023”;*

5
6 (2) *in subparagraph (D)—*

7 (A) *in clause (i), by striking “\$115,500 for*
8 *each of fiscal years 2013 through 2017” and in-*
9 *serting “\$129,400 for each of fiscal years 2019*
10 *through 2023”;* and

11 (B) *in clause (ii), by striking “\$184,800 for*
12 *each of fiscal years 2013 through 2017” and in-*
13 *serting “\$207,000 for each of fiscal years 2019*
14 *through 2023”;*

15 (3) *in subparagraph (E)(i)—*

16 (A) *in subclause (I), by striking “\$70,600*
17 *for each of fiscal years 2013 through 2017” and*
18 *inserting “\$79,100 for each of fiscal years 2019*
19 *through 2023”;* and

20 (B) *in subclause (II), by striking “\$122,100*
21 *for each of fiscal years 2013 through 2017” and*
22 *inserting “\$136,800 for each of fiscal years 2019*
23 *through 2023”;* and

24 (4) *in subparagraph (I), by striking “2017..”*
25 *and inserting “2023.”.*

1 (b) *PROHIBITION ON OTHER FEES.*—Section 4(i)(2) of
2 *the Federal Insecticide, Fungicide, and Rodenticide Act* (7
3 *U.S.C. 136a–1(i)(2)*) is amended—

4 (1) by striking “the date of enactment of this sec-
5 tion and ending on September 30, 2019” and insert-
6 ing “the effective date of the *Pesticide Registration*
7 *Improvement Extension Act of 2018* and ending on
8 *September 30, 2025*”; and

9 (2) by inserting after “registration of a pesticide
10 under this Act” the following: “or any other action
11 covered under a table specified in section 33(b)(3),”.

12 (c) *EXTENSION OF PROHIBITION ON TOLERANCE*
13 *FEES.*—Section 408(m)(3) of the *Federal Food, Drug, and*
14 *Cosmetic Act* (21 *U.S.C. 346a(m)(3)*) is amended by strik-
15 ing “2017” and inserting “2023”.

16 **SEC. 3. REREGISTRATION AND EXPEDITED PROCESSING**
17 **FUND.**

18 (a) *AUTHORIZED USE OF FUND.*—Section 4(k)(2)(A)
19 *of the Federal Insecticide, Fungicide, and Rodenticide Act*
20 *(7 U.S.C. 136a–1(k)(2)(A))* is amended—

21 (1) in the first sentence, by striking “the fund”
22 and inserting “the *Reregistration and Expedited*
23 *Processing Fund*”;

24 (2) by striking “paragraph (3),” in the first sen-
25 tence and all that follows through the period at the

1 *end of the second sentence and inserting the following:*
2 *“paragraph (3), to offset the costs of registration re-*
3 *view under section 3(g), including the costs associated*
4 *with any review under the Endangered Species Act of*
5 *1973 (16 U.S.C. 1531 et seq.) required as part of the*
6 *registration review, to offset the costs associated with*
7 *tracking and implementing registration review deci-*
8 *sions, including registration review decisions designed*
9 *to reduce risk, for the purposes specified in para-*
10 *graphs (4) and (5), and to enhance the information*
11 *systems capabilities to improve the tracking of pes-*
12 *ticide registration decisions.”;*

13 *(3) in clause (i), by striking “are allocated sole-*
14 *ly” and all that follows through “3(g);” and inserting*
15 *the following: “are allocated solely for the purposes*
16 *specified in the first sentence of this subparagraph;”;*
17 *and*

18 *(4) in clause (ii), by striking “necessary to*
19 *achieve” and all that follows through “3(g);” and in-*
20 *serting the following: “necessary to achieve the pur-*
21 *poses specified in the first sentence of this subpara-*
22 *graph;”.*

23 *(b) SET-ASIDE FOR REVIEW OF INERT INGREDIENTS*
24 *AND EXPEDITED PROCESSING OF SIMILAR APPLICA-*
25 *TIONS.—Section 4(k)(3)(A) of the Federal Insecticide, Fun-*

1 *gicide, and Rodenticide Act (7 U.S.C. 136a-1(k)(3)(A)) is*
 2 *amended, in the matter preceding clause (i), by striking*
 3 *“The Administrator shall use” and all that follows through*
 4 *“personnel and resources—” and inserting the following:*
 5 *“For each of fiscal years 2018 through 2023, the Adminis-*
 6 *trator shall use between $\frac{1}{9}$ and $\frac{1}{8}$ of the maintenance fees*
 7 *collected in such fiscal year to obtain sufficient personnel*
 8 *and resources—”.*

9 (c) *SET-ASIDE FOR EXPEDITED RULEMAKING AND*
 10 *GUIDANCE DEVELOPMENT FOR CERTAIN PURPOSES.—*
 11 *Paragraph (4) of section 4(k) of the Federal Insecticide,*
 12 *Fungicide, and Rodenticide Act (7 U.S.C. 136a-1(k)) is*
 13 *amended to read as follows:*

14 “(4) *EXPEDITED RULEMAKING AND GUIDANCE*
 15 *DEVELOPMENT FOR CERTAIN PRODUCT PERFORMANCE*
 16 *DATA REQUIREMENTS.—*

17 “(A) *SET-ASIDE.—For each of fiscal years*
 18 *2018 through 2023, the Administrator shall use*
 19 *not more than \$500,000 of the amounts made*
 20 *available to the Administrator in the Reregistra-*
 21 *tion and Expedited Processing Fund for the ac-*
 22 *tivities described in subparagraph (B).*

23 “(B) *PRODUCTS CLAIMING EFFICACY*
 24 *AGAINST INVERTEBRATE PESTS OF SIGNIFICANT*
 25 *PUBLIC HEALTH OR ECONOMIC IMPORTANCE.—*

1 *The Administrator shall use amounts made*
2 *available under subparagraph (A) to develop, re-*
3 *ceive comments with respect to, finalize, and im-*
4 *plement the necessary rulemaking and guidance*
5 *for product performance data requirements to*
6 *evaluate products claiming efficacy against the*
7 *following invertebrate pests of significant public*
8 *health or economic importance (in order of im-*
9 *portance):*

10 *“(i) Bed bugs.*

11 *“(ii) Premise (including crawling in-*
12 *sects, flying insects, and baits).*

13 *“(iii) Pests of pets (including pet pests*
14 *controlled by spot-ons, collars, shampoos,*
15 *powders, or dips).*

16 *“(iv) Fire ants.*

17 *“(C) DEADLINES FOR GUIDANCE.—The Ad-*
18 *ministrator shall develop, and publish guidance*
19 *required by subparagraph (B), with respect to*
20 *claims of efficacy against pests described in such*
21 *subparagraph as follows:*

22 *“(i) With respect to bed bugs, issue*
23 *final guidance not later than 30 days after*
24 *the effective date of the Pesticide Registra-*
25 *tion Improvement Extension Act of 2018.*

1 “(ii) *With respect to pests specified in*
2 *clause (ii) of such subparagraph—*

3 “(I) *submit draft guidance to the*
4 *Scientific Advisory Panel and for pub-*
5 *lic comment not later than June 30,*
6 *2018; and*

7 “(II) *complete any response to*
8 *comments received with respect to such*
9 *draft guidance and finalize the guid-*
10 *ance not later than September 30,*
11 *2019.*

12 “(iii) *With respect to pests specified in*
13 *clauses (iii) and (iv) of such subpara-*
14 *graph—*

15 “(I) *submit draft guidance to the*
16 *Scientific Advisory Panel and for pub-*
17 *lic comment not later than June 30,*
18 *2019; and*

19 “(II) *complete any response to*
20 *comments received with respect to such*
21 *draft guidance and finalize the guid-*
22 *ance not later than March 31, 2021.*

23 “(D) *REVISION.—The Administrator shall*
24 *revise the guidance required by subparagraph*
25 *(B) from time to time, but shall permit appli-*

1 *cants and registrants sufficient time to obtain*
2 *data that meet the requirements specified in such*
3 *revised guidance.*

4 “(E) *DEADLINE FOR PRODUCT PERFORM-*
5 *ANCE DATA REQUIREMENTS.—The Administrator*
6 *shall, not later than September 30, 2021, issue*
7 *regulations prescribing product performance*
8 *data requirements for any pesticide intended for*
9 *preventing, destroying, repelling, or mitigating*
10 *any invertebrate pest of significant public health*
11 *or economic importance specified in clauses (i)*
12 *through (iv) of subparagraph (B).”.*

13 (d) *SET-ASIDE FOR GOOD LABORATORY PRACTICES*
14 *INSPECTIONS.—Section 4(k) of the Federal Insecticide,*
15 *Fungicide, and Rodenticide Act (7 U.S.C. 136a–1(k)) is*
16 *amended—*

17 (1) *by redesignating paragraphs (5) and (6) as*
18 *paragraphs (6) and (7), respectively;*

19 (2) *by inserting after paragraph (4) the fol-*
20 *lowing new paragraph:*

21 “(5) *GOOD LABORATORY PRACTICES INSPEC-*
22 *TIONS.—*

23 “(A) *SET-ASIDE.—For each of fiscal years*
24 *2018 through 2023, the Administrator shall use*
25 *not more than \$500,000 of the amounts made*

1 *available to the Administrator in the Reregistra-*
2 *tion and Expedited Processing Fund for the ac-*
3 *tivities described in subparagraph (B).*

4 “(B) *ACTIVITIES.*—*The Administrator shall*
5 *use amounts made available under subparagraph*
6 *(A) for enhancements to the good laboratory*
7 *practices standards compliance monitoring pro-*
8 *gram established under part 160 of title 40 of the*
9 *Code of Federal Regulations (or successor regula-*
10 *tions), with respect to laboratory inspections and*
11 *data audits conducted in support of pesticide*
12 *product registrations under this Act. As part of*
13 *such monitoring program, the Administrator*
14 *shall make available to each laboratory inspected*
15 *under such program in support of such registra-*
16 *tions a preliminary summary of inspection ob-*
17 *servations not later than 60 days after the date*
18 *on which such an inspection is completed.”; and*
19 *(3) in paragraph (7), as so redesignated, by*
20 *striking “paragraphs (2), (3), and (4)” and inserting*
21 *“paragraphs (2), (3), (4), and (5)”.*

22 **SEC. 4. EXPERIMENTAL USE PERMITS FOR PESTICIDES.**

23 *Section 5(a) of the Federal Insecticide, Fungicide, and*
24 *Rodenticide Act (7 U.S.C. 136c(a)) is amended—*

1 (1) by striking “permit for a pesticide.” and in-
2 serting “permit for a pesticide. An application for an
3 experimental use permit for a covered application
4 under section 33(b) shall conform with the require-
5 ments of that section.”; and

6 (2) by inserting “(or in the case of an applica-
7 tion for an experimental use permit for a covered ap-
8 plication under section 33(b), not later than the last
9 day of the applicable timeframe for such application
10 specified in such section)” after “all required sup-
11 porting data”.

12 **SEC. 5. PESTICIDE REGISTRATION SERVICE FEES.**

13 (a) *EXTENSION AND MODIFICATION OF FEE AUTHOR-*
14 *ITY.—Section 33(b) of the Federal Insecticide, Fungicide,*
15 *and Rodenticide Act (7 U.S.C. 136w–8(b)) is amended—*

16 (1) *in paragraph (2)—*

17 (A) *in the heading, by striking “PESTICIDE*
18 *REGISTRATION”;* and

19 (B) *in subparagraph (A), by inserting “or*
20 *for any other action covered by a table specified*
21 *in paragraph (3)” after “covered by this Act that*
22 *is received by the Administrator on or after the*
23 *effective date of the Pesticide Registration Im-*
24 *provement Act of 2003”;*

25 (2) *in paragraph (5)—*

1 (A) in the heading, by striking “PESTICIDE
2 REGISTRATION APPLICATIONS” and inserting
3 “COVERED APPLICATIONS”; and

4 (B) by striking “pesticide registration ap-
5 plication” both places it appears and inserting
6 “covered application”;

7 (3) in paragraph (6)—

8 (A) in subparagraph (A)—

9 (i) by striking “pesticide registration”;

10 and

11 (ii) by striking “October 1, 2013, and
12 ending on September 30, 2015” and insert-
13 ing “October 1, 2019, and ending on Sep-
14 tember 30, 2021”;

15 (B) in subparagraph (B)—

16 (i) by striking “pesticide registration”;

17 and

18 (ii) by striking “2015” each place it
19 appears and inserting “2021”; and

20 (C) in subparagraph (C), by striking “re-
21 vised registration service fee schedules” and in-
22 serting “service fee schedules revised pursuant to
23 this paragraph”;

24 (4) in paragraph (7)—

25 (A) in subparagraph (A)—

1 (i) by striking “covered pesticide reg-
2 istration” and inserting “covered applica-
3 tion”; and

4 (ii) by inserting before the period at
5 the end the following: “, except that no
6 waiver or fee reduction shall be provided in
7 connection with a request for a letter of cer-
8 tification (commonly referred to as a Gold
9 Seal letter)”; and

10 (B) in subparagraph (F)(i), by striking
11 “pesticide registration”; and
12 (5) in paragraph (8)—

13 (A) in subparagraph (A), by striking “pes-
14 ticide registration”;

15 (B) in subparagraph (B)(i), by striking
16 “pesticide registration”; and

17 (C) in subparagraph (C)—

18 (i) in clause (i), by striking “pesticide
19 registration” and inserting “covered”; and

20 (ii) in clause (ii)(I), by striking “pes-
21 ticide registration” and inserting “covered”.

22 (b) *PESTICIDE REGISTRATION FUND SET-ASIDES FOR*
23 *WORKER PROTECTION, PARTNERSHIP GRANTS, AND PES-*
24 *TICIDE SAFETY EDUCATION.*—Section 33(c)(3)(B) of the

1 *Federal Insecticide, Fungicide, and Rodenticide Act* (7
2 *U.S.C. 136w–8(c)(3)(B)*) is amended—

3 (1) in the heading, by inserting “, PARTNERSHIP
4 GRANTS, AND PESTICIDE SAFETY EDUCATION” after
5 “WORKER PROTECTION”;

6 (2) in clause (i)—

7 (A) by striking “2017” and inserting
8 “2023”; and

9 (B) by inserting before the period at the end
10 the following: “, with an emphasis on field-worker
11 populations in the United States”;

12 (3) in clause (ii), by striking “2017” and insert-
13 ing “2023”; and

14 (4) in clause (iii), by striking “2017” and in-
15 serting “2023”.

16 (c) *REFORMS TO REDUCE DECISION TIME REVIEW*
17 *PERIODS*.—Section 33(e) of the *Federal Insecticide, Fun-*
18 *gicide, and Rodenticide Act* (7 *U.S.C. 136w–8(e)*) is amend-
19 ed—

20 (1) by striking “Pesticide Registration Improve-
21 ment Extension Act of 2012” and inserting “Pesticide
22 Registration Improvement Extension Act of 2018”;
23 and

24 (2) by inserting at the end the following new sen-
25 tence: “Such reforms shall include identifying oppor-

1 *tunities for streamlining review processes for applica-*
2 *tions for a new active ingredient or a new use and*
3 *providing prompt feedback to applicants during such*
4 *review process.”.*

5 *(d) DECISION TIME REVIEW PERIODS.—Section 33(f)*
6 *of the Federal Insecticide, Fungicide, and Rodenticide Act*
7 *(7 U.S.C. 136w–8(f)) is amended—*

8 *(1) in paragraph (1)—*

9 *(A) by striking “Pesticide Registration Im-*
10 *provement Extension Act of 2012” and inserting*
11 *“Pesticide Registration Improvement Extension*
12 *Act of 2018”; and*

13 *(B) by inserting after “covered pesticide*
14 *registration actions” the following: “or for any*
15 *other action covered by a table specified in sub-*
16 *section (b)(3)”;*

17 *(2) in paragraph (3), by striking subparagraph*
18 *(C) and inserting the following new subparagraph:*

19 *“(C) applications for any other action cov-*
20 *ered by a table specified in subsection (b)(3).”;*
21 *and*

22 *(3) in paragraph (4)(A)—*

23 *(A) by striking “a pesticide registration ap-*
24 *plication” and inserting “a covered applica-*
25 *tion”;* *and*

1 (B) by striking “covered pesticide registra-
2 tion application” and inserting “covered appli-
3 cation”.

4 (e) *REPORTING REQUIREMENTS.*—Section 33(k) of the
5 *Federal Insecticide, Fungicide, and Rodenticide Act* (7
6 *U.S.C. 136w–8(k)*) is amended—

7 (1) in paragraph (1) by striking “2017” and in-
8 serting “2023”; and

9 (2) in paragraph (2)—

10 (A) in subparagraph (D), by striking clause
11 (i) and inserting the following new clause:

12 “(i) the number of pesticides or pes-
13 ticide cases reviewed and the number of reg-
14 istration review decisions completed, includ-
15 ing—

16 “(I) the number of cases cancelled;

17 “(II) the number of cases requir-
18 ing risk mitigation measures;

19 “(III) the number of cases remov-
20 ing risk mitigation measures;

21 “(IV) the number of cases with no
22 risk mitigation needed; and

23 “(V) the number of cases in which
24 risk mitigation has been fully imple-
25 mented;”;

1 (B) in subparagraph (G)—

2 (i) in clause (i)—

3 (I) by striking “section 4(k)(4)”
4 and inserting “paragraphs (4) and (5)
5 of section 4(k)”; and

6 (II) by striking “that section”
7 and inserting “such paragraphs”;

8 (ii) by striking clauses (ii), (iii), (iv),
9 (v), and (vi);

10 (iii) by inserting after clause (i) the
11 following new clause:

12 “(ii) implementing enhancements to—

13 (I) the electronic tracking of cov-
14 ered applications;

15 (II) the electronic tracking of
16 conditional registrations;

17 (III) the endangered species
18 database;

19 (IV) the electronic review of la-
20 bels submitted with covered applica-
21 tions; and

22 (V) the electronic review and as-
23 sessment of confidential statements of
24 formula submitted with covered appli-
25 cations; and”;

1 (iv) by redesignating clause (vii) as
2 clause (iii);

3 (C) in subparagraph (I), by striking “and”
4 at the end;

5 (D) in subparagraph (J), by striking the
6 period at the end and inserting a semicolon; and

7 (E) by adding at the end the following new
8 subparagraphs:

9 “(K) a review of the progress made in devel-
10 oping, updating, and implementing product per-
11 formance test guidelines for pesticide products
12 that are intended to control invertebrate pests of
13 significant public health importance and, by reg-
14 ulation, prescribing product performance data
15 requirements for such pesticide products reg-
16 istered under section 3;

17 “(L) a review of the progress made in the
18 priority review and approval of new pesticides
19 to control invertebrate public health pests that
20 may transmit vector-borne disease for use in the
21 United States, including each territory or posses-
22 sion of the United States, and United States
23 military installations globally;

24 “(M) a review of the progress made in im-
25 plementing enhancements to the good laboratory

1 *practices standards compliance monitoring pro-*
2 *gram established under part 160 of title 40 of the*
3 *Code of Federal Regulations (or successor regula-*
4 *tions);*

5 *“(N) the number of approvals for active in-*
6 *gredients, new uses, and pesticide end use prod-*
7 *ucts granted in connection with the Design for*
8 *the Environment program (or any successor pro-*
9 *gram) of the Environmental Protection Agency;*
10 *and*

11 *“(O) with respect to funds in the Pesticide*
12 *Registration Fund reserved under subsection*
13 *(c)(3), a review that includes—*

14 *“(i) a description of the amount and*
15 *use of such funds—*

16 *“(I) to carry out activities relat-*
17 *ing to worker protection under clause*
18 *(i) of subsection (c)(3)(B);*

19 *“(II) to award partnership grants*
20 *under clause (ii) of such subsection;*
21 *and*

22 *“(III) to carry out the pesticide*
23 *safety education program under clause*
24 *(iii) of such subsection;*

1 “(ii) an evaluation of the appropriate-
2 ness and effectiveness of the activities,
3 grants, and program described in clause (i);

4 “(iii) a description of how stakeholders
5 are engaged in the decision to fund such ac-
6 tivities, grants, and program; and

7 “(iv) with respect to activities relating
8 to worker protection carried out under sub-
9 paragraph (B)(i) of such subsection, a sum-
10 mary of the analyses from stakeholders, in-
11 cluding from worker community-based orga-
12 nizations, on the appropriateness and effec-
13 tiveness of such activities.”.

14 (f) *TERMINATION OF EFFECTIVENESS.*—Section 33(m)
15 of the Federal Insecticide, Fungicide, and Rodenticide Act
16 (7 U.S.C. 136w–8(m)) is amended—

17 (1) in paragraph (1), by striking “2017” and in-
18 serting “2023”; and

19 (2) in paragraph (2)—

20 (A) in subparagraph (A)—

21 (i) by striking “FISCAL YEAR 2018.—
22 During fiscal year 2018” and inserting
23 “FISCAL YEAR 2024.—During fiscal year
24 2024”; and

1 (ii) by striking “2017” and inserting
2 “2023”;

3 (B) in subparagraph (B)—

4 (i) by striking “FISCAL YEAR 2019.—
5 During fiscal year 2019” and inserting
6 “FISCAL YEAR 2025.—During fiscal year
7 2025”; and

8 (ii) by striking “2017” and inserting
9 “2023”;

10 (C) in subparagraph (C), by striking “SEP-
11 TEMBER 30, 2019.—Effective September 30, 2019”
12 and inserting “SEPTEMBER 30, 2025.—Effective
13 September 30, 2025”; and

14 (D) in subparagraph (D), by striking
15 “2017” both places it appears and inserting
16 “2023”.

17 **SEC. 6. REVISION OF TABLES REGARDING COVERED PES-**
18 **TICIDE REGISTRATION APPLICATIONS AND**
19 **OTHER COVERED ACTIONS AND THEIR COR-**
20 **RESPONDING REGISTRATION SERVICE FEES.**

21 Paragraph (3) of section 33(b) of the Federal Insecti-
22 cide, Fungicide, and Rodenticide Act (7 U.S.C. 136w–8(b))
23 is amended to read as follows:

24 “(3) SCHEDULE OF COVERED APPLICATIONS AND
25 OTHER ACTIONS AND THEIR REGISTRATION SERVICE

1 *FEES.—Subject to paragraph (6), the schedule of reg-*
 2 *istration applications and other covered actions and*
 3 *their corresponding registration service fees shall be*
 4 *as follows:*

“TABLE 1. — REGISTRATION DIVISION — NEW ACTIVE INGREDIENTS

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
<i>R010</i>	<i>1</i>	<i>New Active Ingredient, Food use. (2)(3)</i>	<i>24</i>	<i>753,082</i>
<i>R020</i>	<i>2</i>	<i>New Active Ingredient, Food use; reduced risk. (2)(3)</i>	<i>18</i>	<i>627,568</i>
<i>R040</i>	<i>3</i>	<i>New Active Ingredient, Food use; Experimental Use Permit application; establish temporary tolerance; submitted before application for registration; credit 45% of fee toward new active ingredient application that follows. (3)</i>	<i>18</i>	<i>462,502</i>
<i>R060</i>	<i>4</i>	<i>New Active Ingredient, Non-food use; outdoor. (2)(3)</i>	<i>21</i>	<i>523,205</i>
<i>R070</i>	<i>5</i>	<i>New Active Ingredient, Non-food use; outdoor; reduced risk. (2)(3)</i>	<i>16</i>	<i>436,004</i>

“TABLE 1. — REGISTRATION DIVISION — NEW ACTIVE INGREDIENTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
R090	6	<i>New Active Ingredient, Non-food use; outdoor; Experimental Use Permit application; submitted before application for registration; credit 45% of fee toward new active ingredient application that follows. (3)</i>	16	323,690
R110	7	<i>New Active Ingredient, Non-food use; indoor. (2)(3)</i>	20	290,994
R120	8	<i>New Active Ingredient, Non-food use; indoor; reduced risk. (2)(3)</i>	14	242,495
R121	9	<i>New Active Ingredient, Non-food use; indoor; Experimental Use Permit application; submitted before application for registration; credit 45% of fee toward new active ingredient application that follows. (3)</i>	18	182,327
R122	10	<i>Enriched isomer(s) of registered mixed-isomer active ingredient. (2)(3)</i>	18	317,128

“TABLE 1. — REGISTRATION DIVISION — NEW ACTIVE INGREDIENTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
<i>R123</i>	<i>11</i>	<i>New Active Ingredient, Seed treatment only; includes agricultural and non-agricultural seeds; residues not expected in raw agricultural commodities. (2)(3)</i>	<i>18</i>	<i>471,861</i>
<i>R125</i>	<i>12</i>	<i>New Active Ingredient, Seed treatment; Experimental Use Permit application; submitted before application for registration; credit 45% of fee toward new active ingredient application that follows. (3)</i>	<i>16</i>	<i>323,690</i>

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) All requests for new uses (food and/or nonfood) contained in any application for a new active ingredient or a first food use are covered by the base fee for that new active ingredient or first food use application and retain the same decision time review period as the new active ingredient or first food use application. The application must be received by the agency in one package. The base fee for the category covers a maximum of five new products. Each application for an additional new product registration and new inert approval that is submitted in the new active ingredient application package or first food use application package is subject to the registration service fee for a new product or a new inert approval. All such associated applications that are submitted together will be subject to the new active ingredient or first food use decision review time. In the case of a new active ingredient application, until that new active ingredient is approved, any subsequent application for another new product containing the same active ingredient or an amendment to the proposed labeling will be deemed a new active ingredient application, subject to the registration service fee and decision review time for a new active ingredient. In the case of a first food use application, until that first food use is approved, any subsequent application for an additional new food use or uses will be subject to the registration service fee and decision review time for a first food use. Any information that (a) was neither requested nor required by the Agency, and (b) is submitted by the applicant at the applicant's initiative to support the application after completion of the technical deficiency screening, and (c) is not itself a covered registration application, must be assessed 25% of the full registration service fee for the new active ingredient or first food use application.

(3) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

“TABLE 2. — REGISTRATION DIVISION — NEW USES

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R130	13	First food use; indoor; food/food handling. (2) (3)	21	191,444
R140	14	Additional food use; Indoor; food/food handling. (3) (4)	15	44,672
R150	15	First food use. (2)(3)	21	317,104

“TABLE 2. — REGISTRATION DIVISION — NEW USES—
Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
<i>R155</i>	<i>16 (new)</i>	<i>First food use, Experimental Use Permit application; a.i. registered for non-food outdoor use. (3)(4)</i>	<i>21</i>	<i>264,253</i>
<i>R160</i>	<i>17</i>	<i>First food use; reduced risk. (2)(3)</i>	<i>16</i>	<i>264,253</i>
<i>R170</i>	<i>18</i>	<i>Additional food use. (3) (4)</i>	<i>15</i>	<i>79,349</i>
<i>R175</i>	<i>19</i>	<i>Additional food uses covered within a crop group resulting from the conversion of existing approved crop group(s) to one or more revised crop groups. (3)(4)</i>	<i>10</i>	<i>66,124</i>
<i>R180</i>	<i>20</i>	<i>Additional food use; reduced risk. (3)(4)</i>	<i>10</i>	<i>66,124</i>
<i>R190</i>	<i>21</i>	<i>Additional food uses; 6 or more submitted in one application. (3)(4)</i>	<i>15</i>	<i>476,090</i>
<i>R200</i>	<i>22</i>	<i>Additional Food Use; 6 or more submitted in one application; Reduced Risk. (3)(4)</i>	<i>10</i>	<i>396,742</i>

“TABLE 2. — REGISTRATION DIVISION — NEW USES—
Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
R210	23	<i>Additional food use; Experimental Use Permit application; establish temporary tolerance; no credit toward new use registration. (3)(4)</i>	12	48,986
R220	24	<i>Additional food use; Experimental Use Permit application; crop destruct basis; no credit toward new use registration. (3)(4)</i>	6	19,838
R230	25	<i>Additional use; non-food; outdoor. (3) (4)</i>	15	31,713
R240	26	<i>Additional use; non-food; outdoor; reduced risk. (3)(4)</i>	10	26,427
R250	27	<i>Additional use; non-food; outdoor; Experimental Use Permit application; no credit toward new use registration. (3)(4)</i>	6	19,838
R251	28	<i>Experimental Use Permit application which requires no changes to the tolerance(s); non-crop destruct basis. (3)</i>	8	19,838

“TABLE 2. — REGISTRATION DIVISION — NEW USES—
Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
R260	29	<i>New use; non-food; indoor. (3) (4)</i>	12	15,317
R270	30	<i>New use; non-food; indoor; reduced risk. (3)(4)</i>	9	12,764
R271	31	<i>New use; non-food; indoor; Experimental Use Permit application; no credit toward new use registration. (3)(4)</i>	6	9,725
R273	32	<i>Additional use; seed treatment; limited uptake into Raw Agricultural Commodities; includes crops with established tolerances (e.g., for soil or foliar application); includes food and/or non-food uses. (3)(4)</i>	12	50,445

“TABLE 2. — REGISTRATION DIVISION — NEW USES—
Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R274	33	Additional uses; seed treatment only; 6 or more submitted in one application; limited uptake into raw agricultural commodities; includes crops with established tolerances (e.g., for soil or foliar application); includes food and/or non-food uses. (3)(4)	12	302,663

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) All requests for new uses (food and/or nonfood) contained in any application for a new active ingredient or a first food use are covered by the base fee for that new active ingredient or first food use application and retain the same decision time review period as the new active ingredient or first food use application. The application must be received by the agency in one package. The base fee for the category covers a maximum of five new products. Each application for an additional new product registration and new inert approval that is submitted in the new active ingredient application package or first food use application package is subject to the registration service fee for a new product or a new inert approval. All such associated applications that are submitted together will be subject to the new active ingredient or first food use decision review time. In the case of a new active ingredient application, until that new active ingredient is approved, any subsequent application for another new product containing the same active ingredient or an amendment to the proposed labeling will be deemed a new active ingredient application, subject to the registration service fee and decision review time for a new active ingredient. In the case of a first food use application, until that first food use is approved, any subsequent application for an additional new food use or uses will be subject to the registration service fee and decision review time for a first food use. Any information that (a) was neither requested nor required by the Agency, and (b) is submitted by the applicant at the applicant's initiative to support the application after completion of the technical deficiency screening, and (c) is not itself a covered registration application, must be assessed 25% of the full registration service fee for the new active ingredient or first food use application.

(3) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

(4) Amendment applications to add the new use(s) to registered product labels are covered by the base fee for the new use(s). All items in the covered application must be submitted together in one package. Each application for an additional new product registration and new inert approval(s) that is submitted in the new use application package is subject to the registration service fee for a new product or a new inert approval. However, if a new use application only proposes to register the new use for a new product and there are no amendments in the application, then review of one new product application is covered by the new use fee. All such associated applications that are submitted together will be subject to the new use decision review time. Any application for a new product or an amendment to the proposed labeling (a) submitted subsequent to submission of the new use application and (b) prior to conclusion of its decision review time and (c) containing the same new uses, will be deemed a separate new-use application, subject to a separate registration service fee and new decision review time for a new use. If the new-use application includes non-food (indoor and/or outdoor), and food (outdoor and/or indoor) uses, the appropriate fee is due for each type of new use and the longest decision review time applies to all of the new uses requested in the application. Any information that (a) was neither requested nor required by the Agency, and (b) is submitted by the applicant at the applicant's initiative to support the application after completion of the technical deficiency screen, and (c) is not itself a covered registration application, must be assessed 25% of the full registration service fee for the new use application.

“TABLE 3. — REGISTRATION DIVISION — IMPORT AND OTHER TOLERANCES

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
R280	34	Establish import tolerance; new active ingredient or first food use. (2)	21	319,072
R290	35	Establish Import tolerance; Additional new food use.	15	63,816

“TABLE 3. — REGISTRATION DIVISION — IMPORT AND OTHER TOLERANCES—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
R291	36	<i>Establish import tolerances; additional food uses; 6 or more crops submitted in one petition.</i>	15	382,886
R292	37	<i>Amend an established tolerance (e.g., decrease or increase) and/or harmonize established tolerances with Codex MRLs; domestic or import; applicant-initiated.</i>	11	45,341
R293	38	<i>Establish tolerance(s) for inadvertent residues in one crop; applicant-initiated.</i>	12	53,483
R294	39	<i>Establish tolerances for inadvertent residues; 6 or more crops submitted in one application; applicant-initiated.</i>	12	320,894

“TABLE 3. — REGISTRATION DIVISION — IMPORT AND OTHER TOLERANCES—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R295	40	<i>Establish tolerance(s) for residues in one rotational crop in response to a specific rotational crop application; submission of corresponding label amendments which specify the necessary plant-back restrictions; applicant-initiated. (3) (4)</i>	15	66,124
R296	41	<i>Establish tolerances for residues in rotational crops in response to a specific rotational crop petition; 6 or more crops submitted in one application; submission of corresponding label amendments which specify the necessary plant-back restrictions; applicant-initiated. (3) (4)</i>	15	396,742
R297	42	<i>Amend 6 or more established tolerances (e.g., decrease or increase) in one petition; domestic or import; applicant-initiated.</i>	11	272,037

“TABLE 3. — REGISTRATION DIVISION — IMPORT AND OTHER TOLERANCES—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R298	43	Amend an established tolerance (e.g., decrease or increase); domestic or import; submission of corresponding amended labels (requiring science review). (3) (4)	13	58,565
R299	44	Amend 6 or more established tolerances (e.g., decrease or increase); domestic or import; submission of corresponding amended labels (requiring science review). (3) (4)	13	285,261

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) All requests for new uses (food and/or nonfood) contained in any application for a new active ingredient or a first food use are covered by the base fee for that new active ingredient or first food use application and retain the same decision time review period as the new active ingredient or first food use application. The application must be received by the agency in one package. The base fee for the category covers a maximum of five new products. Each application for an additional new product registration and new inert approval that is submitted in the new active ingredient application package or first food use application package is subject to the registration service fee for a new product or a new inert approval. All such associated applications that are submitted together will be subject to the new active ingredient or first food use decision review time. In the case of a new active ingredient application, until that new active ingredient is approved, any subsequent application for another new product containing the same active ingredient or an amendment to the proposed labeling will be deemed a new active ingredient application, subject to the registration service fee and decision review time for a new active ingredient. In the case of a first food use application, until that first food use is approved, any subsequent application for an additional new food use or uses will be subject to the registration service fee and decision review time for a first food use. Any information that (a) was neither requested nor required by the Agency, and (b) is submitted by the applicant at the applicant's initiative to support the application after completion of the technical deficiency screening, and (c) is not itself a covered registration application, must be assessed 25% of the full registration service fee for the new active ingredient or first food use application.

(3) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

(4) Amendment applications to add the revised use pattern(s) to registered product labels are covered by the base fee for the category. All items in the covered application must be submitted together in one package. Each application for an additional new product registration and new inert approval(s) that is submitted in the amendment application package is subject to the registration service fee for a new product or a new inert approval. However, if an amendment application only proposes to register the amendment for a new product and there are no amendments in the application, then review of one new product application is covered by the base fee. All such associated applications that are submitted together will be subject to the category decision review time.

“TABLE 4. — REGISTRATION DIVISION — NEW PRODUCTS

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R300	45	<i>New product; or similar combination product (already registered) to an identical or substantially similar in composition and use to a registered product; registered source of active ingredient; no data review on acute toxicity, efficacy or CRP — only product chemistry data; cite-all data citation, or selective data citation where applicant owns all required data, or applicant submits specific authorization letter from data owner. Category also includes 100% repackage of registered end-use or manufacturing-use product that requires no data submission nor data matrix. (2)(3)</i>	4	1,582

“TABLE 4. — REGISTRATION DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R301	46	<i>New product; or similar combination product (already registered) to an identical or substantially similar in composition and use to a registered product; registered source of active ingredient; selective data citation only for data on product chemistry and/or acute toxicity and/or public health pest efficacy (identical data citation and claims to cited product(s)), where applicant does not own all required data and does not have a specific authorization letter from data owner. (2)(3)</i>	4	1,897

“TABLE 4. — REGISTRATION DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R310	47	<p><i>New end-use or manufacturing-use product with registered source(s) of active ingredient(s); includes products containing two or more registered active ingredients previously combined in other registered products; excludes products requiring or citing an animal safety study; requires review of data package within RD only; includes data and/or waivers of data for only:</i></p> <ul style="list-style-type: none"> ● <i>product chemistry and/or</i> ● <i>acute toxicity and/or</i> ● <i>child resistant packaging and/or</i> ● <i>pest(s) requiring efficacy (4) - for up to 3 target pests. (2)(3)</i> 	7	7,301

“TABLE 4. — REGISTRATION DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R314	48	<p><i>New end use product containing up to three registered active ingredients never before registered as this combination in a formulated product; new product label is identical or substantially similar to the labels of currently registered products which separately contain the respective component active ingredients; excludes products requiring or citing an animal safety study; requires review of data package within RD only; includes data and/or waivers of data for only:</i></p> <ul style="list-style-type: none"> ● <i>product chemistry and/or</i> ● <i>acute toxicity and/or</i> ● <i>child resistant packaging and/or</i> ● <i>pest(s) requiring efficacy (4) - for up to 3 target pests. (2)(3)</i> 	8	8,626

“TABLE 4. — REGISTRATION DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R319	49	<p><i>New end use product containing up to three registered active ingredients never before registered as this combination in a formulated product; new product label is identical or substantially similar to the labels of currently registered products which separately contain the respective component active ingredients; excludes products requiring or citing an animal safety study; requires review of data package within RD only; includes data and/or waivers of data for only:</i></p> <ul style="list-style-type: none"> ● <i>product chemistry and/or</i> ● <i>acute toxicity and/or</i> ● <i>child resistant packaging and/or</i> ● <i>pest(s) requiring efficacy (4) - for 4 to 7 target pests. (2)(3)</i> 	10	12,626

“TABLE 4. — REGISTRATION DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R318	50 (new)	<p><i>New end use product containing four or more registered active ingredients never before registered as this combination in a formulated product; new product label is identical or substantially similar to the labels of currently registered products which separately contain the respective component active ingredients; excludes products requiring or citing an animal safety study; requires review of data package within RD only; includes data and/or waivers of data for only:</i></p> <ul style="list-style-type: none"> ● <i>product chemistry and/or</i> ● <i>acute toxicity and/or</i> ● <i>child resistant packaging and/or</i> ● <i>pest(s) requiring efficacy (4) - for up to 3 target pests. (2)(3)</i> 	9	13,252

“TABLE 4. — REGISTRATION DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R321	51 (new)	<p><i>New end use product containing four or more registered active ingredients never before registered as this combination in a formulated product; new product label is identical or substantially similar to the labels of currently registered products which separately contain the respective component active ingredients; excludes products requiring or citing an animal safety study; requires review of data package within RD only; includes data and/or waivers of data for only:</i></p> <ul style="list-style-type: none"> ● <i>product chemistry and/or</i> ● <i>acute toxicity and/or</i> ● <i>child resistant packaging and/or</i> ● <i>pest(s) requiring efficacy (4) - for 4 to 7 target pests. (2)(3)</i> 	11	17,252

“TABLE 4. — REGISTRATION DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R315	52	<p><i>New end-use, on-animal product, registered source of active ingredient(s), with the submission of data and/or waivers for only:</i></p> <ul style="list-style-type: none"> ● <i>animal safety and</i> ● <i>pest(s) requiring efficacy (4) and/or</i> ● <i>product chemistry and/or</i> ● <i>acute toxicity and/or</i> ● <i>child resistant packaging. (2)</i> <p><i>(3)</i></p>	9	9,820

“TABLE 4. — REGISTRATION DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R316	53 (new)	<p><i>New end-use or manufacturing product with registered source(s) of active ingredient(s) including products containing two or more registered active ingredients previously combined in other registered products; excludes products requiring or citing an animal safety study; and requires review of data and/or waivers for only:</i></p> <ul style="list-style-type: none"> ● <i>product chemistry and/or</i> ● <i>acute toxicity and/or</i> ● <i>child resistant packaging and/or</i> ● <i>pest(s) requiring efficacy (4) - for greater than 3 and up to 7 target pests. (2)(3)</i> 	9	11,301

“TABLE 4. — REGISTRATION DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R317	54 (new)	<p>New end-use or manufacturing product with registered source(s) of active ingredient(s) including products containing 2 or more registered active ingredients previously combined in other registered products; excludes products requiring or citing an animal safety study; and requires review of data and/or waivers for only:</p> <ul style="list-style-type: none"> ● product chemistry and/or ● acute toxicity and/or ● child resistant packaging and/or ● pest(s) requiring efficacy (4) - for greater than 7 target pests. <p>(2)(3)</p>	10	15,301
R320	55	<p>New product; new physical form; requires data review in science divisions. (2)(3)</p>	12	13,226

“TABLE 4. — REGISTRATION DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R331	56	<i>New product; re-pack of identical registered end-use product as a manufacturing-use product, or identical registered manufacturing-use product as an end use product; same registered uses only. (2)(3)</i>	3	2,530
R332	57	<i>New manufacturing-use product; registered active ingredient; unregistered source of active ingredient; submission of completely new generic data package; registered uses only; requires review in RD and science divisions. (2)(3)</i>	24	283,215
R333	58	<i>New product; MUP or End use product with unregistered source of active ingredient; requires science data review; new physical form; etc. Cite-all or selective data citation where applicant owns all required data. (2)(3)</i>	10	19,838

“TABLE 4. — REGISTRATION DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R334	59	New product; MUP or End use product with unregistered source of the active ingredient; requires science data review; new physical form; etc. Selective data citation. (2)(3)	11	23,100

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) An application for a new end-use product using a source of active ingredient that (a) is not yet registered but (b) has an application pending with the Agency for review, will be considered an application for a new product with an unregistered source of active ingredient.

(3) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant’s written or electronic confirmation of agreement to the Agency.

(4) For the purposes of classifying proposed registration actions into PRIA categories, “pest(s) requiring efficacy” are: public health pests listed in PR Notice 2002-1, livestock pests (e.g. Horn flies, Stable flies), wood-destroying pests (e.g. termites, carpenter ants, wood-boring beetles) and certain invasive species (e.g. Asian Longhorned beetle, Emerald Ashborer). This list may be updated/refined as invasive pest needs arise. To determine the number of pests for the PRIA categories, pests have been placed into groups (general; e.g., cockroaches) and pest specific (specifically a test species). If seeking a label claim against a pest group (general), use the group listing below and each group will count as 1. The general pests groups are: mites, dust mites, chiggers, ticks, hard ticks, soft ticks, cattle ticks, scorpions, spiders, centipedes, lice, fleas, cockroaches, keds, bot flies, screwworms, filth flies, blow flies, house flies, flesh flies, mosquitoes, biting flies, horse flies, stable flies, deer flies, sand flies, biting midges, black flies, true bugs, bed bugs, stinging bees, wasps, yellow jackets, hornets, ants (excluding carpenter ants), fire and harvester ants, wood destroying beetles, carpenter ants, termites, subterranean termites, dry wood termites, arboreal termites, damp wood termites and invasive species. If seeking a claim against a specific pest without a general claim then each specific pest will count as 1.

“TABLE 5. — REGISTRATION DIVISION — AMENDMENTS

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R340	60	<i>Amendment requiring data review within RD (e.g., changes to precautionary label statements); includes adding/modifying pest(s) claims for up to 2 target pests, excludes products requiring or citing an animal safety study. (2)(3)(4)</i>	4	4,988
R341	61 (New)	<i>Amendment requiring data review within RD (e.g., changes to precautionary label statements), includes adding/modifying pest(s) claims for greater than 2 target pests, excludes products requiring or citing an animal safety study. (2)(3)(4)</i>	6	5,988
R345	62	<i>Amending on-animal products previously registered, with the submission of data and/or waivers for only:</i> <ul style="list-style-type: none"> ● <i>animal safety and</i> ● <i>pest(s) requiring efficacy (4) and/or</i> ● <i>product chemistry and/or</i> ● <i>acute toxicity and/or</i> ● <i>child resistant packaging. (2)(3)</i> 	7	8,820

“TABLE 5. — REGISTRATION DIVISION — AMENDMENTS—
Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R350	63	Amendment requiring data review in science divisions (e.g., changes to REI, or PPE, or PHI, or use rate, or number of applications; or add aerial application; or modify GW/SW advisory statement). (2)(3)	9	13,226
R351	64	Amendment adding a new unregistered source of active ingredient. (2)(3)	8	13,226
R352	65	Amendment adding already approved uses; selective method of support; does not apply if the applicant owns all cited data. (2) (3)	8	13,226
R371	66	Amendment to Experimental Use Permit; (does not include extending a permit's time period). (3)	6	10,090

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) (a) EPA-initiated amendments shall not be charged registration service fees. (b) Registrant-initiated fast-track amendments are to be completed within the timelines specified in FIFRA Section 3(c)(3)(B) and are not subject to registration service fees. (c) Registrant-initiated fast-track amendments handled by the Antimicrobials Division are to be completed within the timelines specified in FIFRA Section 3(h) and are not subject to registration service fees. (d) Registrant initiated amendments submitted by notification under PR Notices, such as PR Notice 98-10, continue under PR Notice timelines and are not subject to registration service fees. (e) Submissions with data and requiring data review are subject to registration service fees.

(3) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

(4) For the purposes of classifying proposed registration actions into PRIA categories, "pest(s) requiring efficacy" are: public health pests listed in PR Notice 2002-1, livestock pests (e.g. Horn flies, Stable flies), wood-destroying pests (e.g. termites, carpenter ants, wood-boring beetles) and certain invasive species (e.g. Asian Longhorned beetle, Emerald Ashborer). This list may be updated/refined as invasive pest needs arise. To determine the number of pests for the PRIA categories, pests have been placed into groups (general; e.g., cockroaches) and pest specific (specifically a test species). If seeking a label claim against a pest group (general), use the group listing below and each group will count as 1. The general pests groups are: mites, dust mites, chiggers, ticks, hard ticks, soft ticks, cattle ticks, scorpions, spiders, centipedes, lice, fleas, cockroaches, keds, bot flies, screwworms, filth flies, blow flies, house flies, flesh flies, mosquitoes, biting flies, horse flies, stable flies, deer flies, sand flies, biting midges, black flies, true bugs, bed bugs, stinging bees, wasps, yellow jackets, hornets, ants (excluding carpenter ants), fire and harvester ants, wood destroying beetles, carpenter ants, termites, subterranean termites, dry wood termites, arboreal termites, damp wood termites and invasive species. If seeking a claim against a specific pest without a general claim then each specific pest will count as 1.

"TABLE 6. — REGISTRATION DIVISION — OTHER ACTIONS

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
R124	67	Conditional Ruling on Pre-application Study Waivers; applicant-initiated.	6	2,530

“TABLE 6. — REGISTRATION DIVISION — OTHER ACTIONS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
R272	68	Review of Study Protocol applicant-initiated; excludes DART, pre-registration conference, Rapid Response review, DNT protocol review, protocol needing HSRB review.	3	2,530
R275	69	Rebuttal of agency reviewed protocol, applicant initiated.	3	2,530
R370	70	Cancer reassessment; applicant-initiated.	18	198,250

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

“TABLE 7. — ANTIMICROBIALS DIVISION — NEW ACTIVE INGREDIENTS

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
A380	71	New Active Ingredient; Indirect Food use; establish tolerance or tolerance exemption if required. (2)(3)	24	137,841
A390	72	New Active Ingredient; Direct Food use; establish tolerance or tolerance exemption if required. (2)(3)	24	229,733

“TABLE 7. — ANTIMICROBIALS DIVISION — NEW ACTIVE INGREDIENTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
A410	73	New Active Ingredient Non-food use.(2)(3)	21	229,733
A431	74	New Active Ingredient, Non-food use; low-risk. (2)(3)	12	80,225

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) All requests for new uses (food and/or nonfood) contained in any application for a new active ingredient or a first food use are covered by the base fee for that new active ingredient or first food use application and retain the same decision time review period as the new active ingredient or first food use application. The application must be received by the agency in one package. The base fee for the category covers a maximum of five new products. Each application for an additional new product registration and new inert approval that is submitted in the new active ingredient application package or first food use application package is subject to the registration service fee for a new product or a new inert approval. All such associated applications that are submitted together will be subject to the new active ingredient or first food use decision review time. In the case of a new active ingredient application, until that new active ingredient is approved, any subsequent application for another new product containing the same active ingredient or an amendment to the proposed labeling will be deemed a new active ingredient application, subject to the registration service fee and decision review time for a new active ingredient. In the case of a first food use application, until that first food use is approved, any subsequent application for an additional new food use or uses will be subject to the registration service fee and decision review time for a first food use. Any information that (a) was neither requested nor required by the Agency, and (b) is submitted by the applicant at the applicant's initiative to support the application after completion of the technical deficiency screening, and (c) is not itself a covered registration application, must be assessed 25% of the full registration service fee for the new active ingredient or first food use application.

(3) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

“TABLE 8. — ANTIMICROBIALS DIVISION — NEW USES

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
A440	75	<i>New Use, Indirect Food Use, establish tolerance or tolerance exemption. (2)(3)(4)</i>	21	31,910
A441	76	<i>Additional Indirect food uses; establish tolerances or tolerance exemptions if required; 6 or more submitted in one application. (3)(4)(5)</i>	21	114,870
A450	77	<i>New use, Direct food use, establish tolerance or tolerance exemption. (2)(3)(4)</i>	21	95,724
A451	78	<i>Additional Direct food uses; establish tolerances or tolerance exemptions if required; 6 or more submitted in one application. (3)(4)(5)</i>	21	182,335
A500	79	<i>New use, non-food. (4)(5)</i>	12	31,910

“TABLE 8. — ANTIMICROBIALS DIVISION — NEW USES—
Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
A501	80	New use, non-food; 6 or more submitted in one application. (4)(5)	15	76,583

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) All requests for new uses (food and/or nonfood) contained in any application for a new active ingredient or a first food use are covered by the base fee for that new active ingredient or first food use application and retain the same decision time review period as the new active ingredient or first food use application. The application must be received by the agency in one package. The base fee for the category covers a maximum of five new products. Each application for an additional new product registration and new inert approval that is submitted in the new active ingredient application package or first food use application package is subject to the registration service fee for a new product or a new inert approval. All such associated applications that are submitted together will be subject to the new active ingredient or first food use decision review time. In the case of a new active ingredient application, until that new active ingredient is approved, any subsequent application for another new product containing the same active ingredient or an amendment to the proposed labeling will be deemed a new active ingredient application, subject to the registration service fee and decision review time for a new active ingredient. In the case of a first food use application, until that first food use is approved, any subsequent application for an additional new food use or uses will be subject to the registration service fee and decision review time for a first food use. Any information that (a) was neither requested nor required by the Agency, and (b) is submitted by the applicant at the applicant's initiative to support the application after completion of the technical deficiency screening, and (c) is not itself a covered registration application, must be assessed 25% of the full registration service fee for the new active ingredient or first food use application.

(3) If EPA data rules are amended to newly require clearance under section 408 of the FFDCA for an ingredient of an antimicrobial product where such ingredient was not previously subject to such a clearance, then review of the data for such clearance of such product is not subject to a registration service fee for the tolerance action for two years from the effective date of the rule.

(4) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

(5) *Amendment applications to add the new use(s) to registered product labels are covered by the base fee for the new use(s). All items in the covered application must be submitted together in one package. Each application for an additional new product registration and new inert approval(s) that is submitted in the new use application package is subject to the registration service fee for a new product or a new inert approval. However, if a new use application only proposes to register the new use for a new product and there are no amendments in the application, then review of one new product application is covered by the new use fee. All such associated applications that are submitted together will be subject to the new use decision review time. Any application for a new product or an amendment to the proposed labeling (a) submitted subsequent to submission of the new use application and (b) prior to conclusion of its decision review time and (c) containing the same new uses, will be deemed a separate new-use application, subject to a separate registration service fee and new decision review time for a new use. If the new-use application includes non-food (indoor and/or outdoor), and food (outdoor and/or indoor) uses, the appropriate fee is due for each type of new use and the longest decision review time applies to all of the new uses requested in the application. Any information that (a) was neither requested nor required by the Agency, and (b) is submitted by the applicant at the applicant's initiative to support the application after completion of the technical deficiency screen, and (c) is not itself a covered registration application, must be assessed 25% of the full registration service fee for the new use application.*

“TABLE 9. — ANTIMICROBIALS DIVISION — NEW PRODUCTS AND AMENDMENTS

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
A530	81	<i>New product, identical or substantially similar in composition and use to a registered product; no data review or only product chemistry data; cite all data citation or selective data citation where applicant owns all required data; or applicant submits specific authorization letter from data owner. Category also includes 100% re-package of registered end-use or manufacturing use product that requires no data submission nor data matrix. (2)(3)</i>	4	1,278

“TABLE 9. — ANTIMICROBIALS DIVISION — NEW
PRODUCTS AND AMENDMENTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registra- tion Service Fee (\$)
A531	82	<i>New product; identical or substantially similar in composition and use to a registered product; registered source of active ingredient; selective data citation only for data on product chemistry and/or acute toxicity and/or public health pest efficacy, where applicant does not own all required data and does not have a specific authorization letter from data owner. (2)(3)</i>	4	1,824
A532	83	<i>New product; identical or substantially similar in composition and use to a registered product; registered active ingredient; unregistered source of active ingredient; cite-all data citation except for product chemistry; product chemistry data submitted. (2)(3)</i>	5	5,107

“TABLE 9. — ANTIMICROBIALS DIVISION — NEW
PRODUCTS AND AMENDMENTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registra- tion Service Fee (\$)
A540	84	New end use prod- uct; FIFRA §2(mm) uses only; up to 25 public health or- ganisms. (2)(3)(5)(6)	5	5,107
A541	85 (new)	New end use prod- uct; FIFRA §2(mm) uses only; 26-50 pub- lic health orga- nisms. (2)(3)(5)(6)	7	8,500
A542	86 (new)	New end use prod- uct; FIFRA §2(mm) uses only; ≥ 51 pub- lic health orga- nisms. (2)(3)(5)	10	15,000
A550	87	New end-use prod- uct; uses other than FIFRA §2(mm); non- FQPA product. (2)(3)(5)	9	13,226
A560	88	New manufac- turing use prod- uct; registered active ingre- dient; selective data citation. (2)(3)	6	12,596

“TABLE 9. — ANTIMICROBIALS DIVISION — NEW PRODUCTS AND AMENDMENTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
A565	89 (new)	New manufacturing-use product; registered active ingredient; unregistered source of active ingredient; submission of new generic data package; registered uses only; requires science review. (2)(3)	12	18,234
A570	90	Label amendment requiring data review; up to 25 public health organisms. (3)(4)(5)(6)	4	3,831
A573	91 (new)	Label amendment requiring data review; 26-50 public health organisms. (2)(3)(5)(7)	6	6,350
A574	92 (new)	Label amendment requiring data review; ≥ 51 public health organisms. (2)(3)(5)(7)	9	11,000

“TABLE 9. — ANTIMICROBIALS DIVISION — NEW
PRODUCTS AND AMENDMENTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
A572	93	New Product or amendment requiring data review for risk assessment by Science Branch (e.g., changes to REI, or PPE, or use rate). (2)(3)(4)	9	13,226

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) An application for a new end-use product using a source of active ingredient that (a) is not yet registered but (b) has an application pending with the Agency for review, will be considered an application for a new product with an unregistered source of active ingredient.

(3) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant’s written or electronic confirmation of agreement to the Agency.

(4)(a) EPA-initiated amendments shall not be charged registration service fees. (b) Registrant-initiated fast-track amendments are to be completed within the timelines specified in FIFRA Section 3(c)(3)(B) and are not subject to registration service fees. (c) Registrant-initiated fast-track amendments handled by the Antimicrobials Division are to be completed within the timelines specified in FIFRA Section 3(h) and are not subject to registration service fees. (d) Registrant initiated amendments submitted by notification under PR Notices, such as PR Notice 98–10, continue under PR Notice timelines and are not subject to registration service fees. (e) Submissions with data and requiring data review are subject to registration service fees.

(5) The applicant must identify the substantially similar product if opting to use cite-all or the selective method to support acute toxicity data requirements.

(6) Once a submission for a new product with public health organisms has been submitted and classified in either A540 or A541, additional organisms submitted for the same product before expiration of the first submission’s original decision review time period will result in reclassification of both the original and subsequent submission into the appropriate new category based on the sum of the number of organisms in both submissions. A reclassification would result in a new PRLA start date and require additional fees to meet the fee of the new category.

(7) Once a submission for a label amendment with public health organisms has been submitted and classified in either A570 or A573, additional organisms submitted for the same product before expiration of the first submission's original decision review time period will result in reclassification of both the original and subsequent submission into the appropriate new category based on the sum of the number of organisms in both submissions. A reclassification would result in a new PRIA start date and require additional fees to meet the fee of the new category.

**“TABLE 10. — ANTIMICROBIALS DIVISION —
EXPERIMENTAL USE PERMITS AND OTHER ACTIONS**

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
A520	94	<i>Experimental Use Permit application, non-food use. (2)</i>	9	6,383
A521	95	<i>Review of public health efficacy study protocol within AD, per AD Internal Guidance for the Efficacy Protocol Review Process; Code will also include review of public health efficacy study protocol and data review for devices making pesticidal claims; applicant-initiated; Tier 1.</i>	4	4,726

“TABLE 10. — ANTIMICROBIALS DIVISION — EXPERIMENTAL USE PERMITS AND OTHER ACTIONS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
A522	96	<i>Review of public health efficacy study protocol outside AD by members of AD Efficacy Protocol Review Expert Panel; Code will also include review of public health efficacy study protocol and data review for devices making pesticidal claims; applicant-initiated; Tier 2.</i>	12	12,156
A537	97 (new)	<i>New Active Ingredient/New Use, Experimental Use Permit application; Direct food use; Establish tolerance or tolerance exemption if required. Credit 45% of fee toward new active ingredient/new use application that follows.</i>	18	153,156

“TABLE 10. — ANTIMICROBIALS DIVISION — EXPERIMENTAL USE PERMITS AND OTHER ACTIONS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
A538	98 (new)	<i>New Active Ingredient/New Use, Experimental Use Permit application; Indirect food use; Establish tolerance or tolerance exemption if required Credit 45% of fee toward new active ingredient/new use application that follows.</i>	18	95,724
A539	99 (new)	<i>New Active Ingredient/New Use, Experimental Use Permit application; Nonfood use. Credit 45% of fee toward new active ingredient/new use application that follows.</i>	15	92,163
A529	100	<i>Amendment to Experimental Use Permit; requires data review or risk assessment. (2)</i>	9	11,429
A523	101	<i>Review of protocol other than a public health efficacy study (i.e., Toxicology or Exposure Protocols).</i>	9	12,156

“TABLE 10. — ANTIMICROBIALS DIVISION — EXPERIMENTAL USE PERMITS AND OTHER ACTIONS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
A571	102	Science reassessment: Cancer risk, refined ecological risk, and/or endangered species; applicant-initiated.	18	95,724
A533	103 (new)	Exemption from the requirement of an Experimental Use Permit. (2)	4	2,482
A534	104 (new)	Rebuttal of agency reviewed protocol, applicant initiated.	4	4,726
A535	105 (new)	Conditional Ruling on Pre-application Study Waiver or Data Bridging Argument; applicant-initiated.	6	2,409
A536	106 (new)	Conditional Ruling on Pre-application Direct Food, Indirect Food, Nonfood use determination; applicant-initiated.	4	2,482

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

“TABLE 11. — BIOPESTICIDES DIVISION — NEW ACTIVE INGREDIENTS

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
B580	107	New active ingredient; food use; petition to establish a tolerance. (2)(3)	20	51,053
B590	108	New active ingredient; food use; petition to establish a tolerance exemption. (2)(3)	18	31,910
B600	109	New active ingredient; non-food use. (2)(3)	13	19,146
B610	110	New active ingredient; Experimental Use Permit application; petition to establish a temporary tolerance or temporary tolerance exemption. (3)	10	12,764

“TABLE 11. — BIOPESTICIDES DIVISION — NEW ACTIVE INGREDIENTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
<i>B611</i>	<i>111</i>	<i>New active ingredient; Experimental Use Permit application; petition to establish permanent tolerance exemption. (3)</i>	<i>12</i>	<i>12,764</i>
<i>B612</i>	<i>112</i>	<i>New active ingredient; no change to a permanent tolerance exemption. (2)(3)</i>	<i>10</i>	<i>17,550</i>
<i>B613</i>	<i>113</i>	<i>New active ingredient; petition to convert a temporary tolerance or a temporary tolerance exemption to a permanent tolerance or tolerance exemption. (2)(3)</i>	<i>11</i>	<i>17,550</i>
<i>B620</i>	<i>114</i>	<i>New active ingredient; Experimental Use Permit application; non-food use including crop destruct. (3)</i>	<i>7</i>	<i>6,383</i>

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) All requests for new uses (food and/or nonfood) contained in any application for a new active ingredient or a first food use are covered by the base fee for that new active ingredient or first food use application and retain the same decision time review period as the new active ingredient or first food use application. The application must be received by the agency in one package. The base fee for the category covers a maximum of five new products. Each application for an additional new product registration and new inert approval that is submitted in the new active ingredient application package or first food use application package is subject to the registration service fee for a new product or a new inert approval. All such associated applications that are submitted together will be subject to the new active ingredient or first food use decision review time. In the case of a new active ingredient application, until that new active ingredient is approved, any subsequent application for another new product containing the same active ingredient or an amendment to the proposed labeling will be deemed a new active ingredient application, subject to the registration service fee and decision review time for a new active ingredient. In the case of a first food use application, until that first food use is approved, any subsequent application for an additional new food use or uses will be subject to the registration service fee and decision review time for a first food use. Any information that (a) was neither requested nor required by the Agency, and (b) is submitted by the applicant at the applicant's initiative to support the application after completion of the technical deficiency screening, and (c) is not itself a covered registration application, must be assessed 25% of the full registration service fee for the new active ingredient or first food use application.

(3) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

“TABLE 12. — BIOPESTICIDES DIVISION — NEW USES

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
B630	115	First food use; petition to establish a tolerance exemption. (2)(4)	13	12,764
B631	116	New food use; petition to amend an established tolerance. (3)(4)	12	12,764

“TABLE 12. — BIOPESTICIDES DIVISION — NEW USES—
Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
B640	117	First food use; petition to establish a tolerance. (2)(4)	19	19,146
B643	118	New Food use; petition to amend an established tolerance exemption. (3)(4)	10	12,764
B642	119	First food use; indoor; food/food handling. (2)(4)	12	31,910
B644	120	New use, no change to an established tolerance or tolerance exemption. (3)(4)	8	12,764
B650	121	New use; non-food. (3)(4)	7	6,383
B645	122 (new)	New food use; Experimental Use Permit application; petition to amend or add a tolerance exemption. (4)	12	12,764
B646	123 (new)	New use; non-food use including crop destruct; Experimental Use Permit application. (4)	7	6,383

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) *All requests for new uses (food and/or nonfood) contained in any application for a new active ingredient or a first food use are covered by the base fee for that new active ingredient or first food use application and retain the same decision time review period as the new active ingredient or first food use application. The application must be received by the agency in one package. The base fee for the category covers a maximum of five new products. Each application for an additional new product registration and new inert approval that is submitted in the new active ingredient application package or first food use application package is subject to the registration service fee for a new product or a new inert approval. All such associated applications that are submitted together will be subject to the new active ingredient or first food use decision review time. In the case of a new active ingredient application, until that new active ingredient is approved, any subsequent application for another new product containing the same active ingredient or an amendment to the proposed labeling will be deemed a new active ingredient application, subject to the registration service fee and decision review time for a new active ingredient. In the case of a first food use application, until that first food use is approved, any subsequent application for an additional new food use or uses will be subject to the registration service fee and decision review time for a first food use. Any information that (a) was neither requested nor required by the Agency, and (b) is submitted by the applicant at the applicant's initiative to support the application after completion of the technical deficiency screening, and (c) is not itself a covered registration application, must be assessed 25% of the full registration service fee for the new active ingredient or first food use application.*

(3) *Amendment applications to add the new use(s) to registered product labels are covered by the base fee for the new use(s). All items in the covered application must be submitted together in one package. Each application for an additional new product registration and new inert approval(s) that is submitted in the new use application package is subject to the registration service fee for a new product or a new inert approval. However, if a new use application only proposes to register the new use for a new product and there are no amendments in the application, then review of one new product application is covered by the new use fee. All such associated applications that are submitted together will be subject to the new use decision review time. Any application for a new product or an amendment to the proposed labeling (a) submitted subsequent to submission of the new use application and (b) prior to conclusion of its decision review time and (c) containing the same new uses, will be deemed a separate new-use application, subject to a separate registration service fee and new decision review time for a new use. If the new-use application includes non-food (indoor and/or outdoor), and food (outdoor and/or indoor) uses, the appropriate fee is due for each type of new use and the longest decision review time applies to all of the new uses requested in the application. Any information that (a) was neither requested nor required by the Agency, and (b) is submitted by the applicant at the applicant's initiative to support the application after completion of the technical deficiency screen, and (c) is not itself a covered registration application, must be assessed 25% of the full registration service fee for the new use application.*

(4) *Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.*

“TABLE 13. — BIOPESTICIDES DIVISION — NEW PRODUCTS

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B652	124	<i>New product; registered source of active ingredient; requires petition to amend established tolerance or tolerance exemption; requires 1) submission of product specific data; or 2) citation of previously reviewed and accepted data; or 3) submission or citation of data generated at government expense; or 4) submission or citation of scientifically-sound rationale based on publicly available literature or other relevant information that addresses the data requirement; or 5) submission of a request for a data requirement to be waived supported by a scientifically-sound rationale explaining why the data requirement does not apply. (2)(3)</i>	13	12,764

“TABLE 13. — BIOPESTICIDES DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B660	125	<p><i>New product; registered source of active ingredient(s); identical or substantially similar in composition and use to a registered product. No data review, or only product chemistry data; cite-all data citation, or selective data citation where applicant owns all required data or authorization from data owner is demonstrated. Category includes 100% re-package of registered end-use or manufacturing-use product that requires no data submission or data matrix. For microbial pesticides, the active ingredient(s) must not be re-isolated.</i></p> <p><i>(2)(3)</i></p>	4	1,278

“TABLE 13. — BIOPESTICIDES DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B670	126	New product; registered source of active ingredient(s); requires: 1) submission of product specific data; or 2) citation of previously reviewed and accepted data; or 3) submission or citation of data generated at government expense; or 4) submission or citation of a scientifically-sound rationale based on publicly available literature or other relevant information that addresses the data requirement; or 5) submission of a request for a data requirement to be waived supported by a scientifically-sound rationale explaining why the data requirement does not apply. (2)(3)	7	5,107

“TABLE 13. — BIOPESTICIDES DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B671	127	New product; unregistered source of active ingredient(s); requires a petition to amend an established tolerance or tolerance exemption; requires: 1) submission of product specific data; or 2) citation of previously reviewed and accepted data; or 3) submission or citation of data generated at government expense; or 4) submission or citation of a scientifically-sound rationale based on publicly available literature or other relevant information that addresses the data requirement; or 5) submission of a request for a data requirement to be waived supported by a scientifically-sound rationale explaining why the data requirement does not apply. (2)(3)	17	12,764

“TABLE 13. — BIOPESTICIDES DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B672	128	<i>New product; unregistered source of active ingredient(s); non-food use or food use requires: 1) submission of product specific data; or 2) citation of previously reviewed and accepted data; or 3) submission or citation of data generated at government expense; or 4) submission or citation of a scientifically-sound rationale based on publicly available literature or other relevant information that addresses the data requirement; or 5) submission of a request for a data requirement to be waived supported by a scientifically-sound rationale explaining why the data requirement does not apply. (2)(3)</i>	13	9,118

“TABLE 13. — BIOPESTICIDES DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B673	129	<i>New product MUP/EP; unregistered source of active ingredient(s); citation of Technical Grade Active Ingredient (TGAI) data previously reviewed and accepted by the Agency. Requires an Agency determination that the cited data supports the new product. (2)(3)</i>	10	5,107
B674	130	<i>New product MUP; Repack of identical registered end-use product as a manufacturing-use product; same registered uses only. (2)(3)</i>	4	1,278
B675	131	<i>New Product MUP; registered source of active ingredient; submission of completely new generic data package; registered uses only. (2)(3)</i>	10	9,118

“TABLE 13. — BIOPESTICIDES DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B676	132	<p><i>New product; more than one active ingredient where one active ingredient is an unregistered source; product chemistry data must be submitted; requires: 1) submission of product specific data, and 2) citation of previously reviewed and accepted data; or 3) submission or citation of data generated at government expense; or 4) submission or citation of a scientifically-sound rationale based on publicly available literature or other relevant information that addresses the data requirement; or 5) submission of a request for a data requirement to be waived supported by a scientifically-sound rationale explaining why the data requirement does not apply. (2)(3)</i></p>	13	9,118

“TABLE 13. — BIOPESTICIDES DIVISION — NEW PRODUCTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B677	133	<p><i>New end-use non-food animal product with submission of two or more target animal safety studies; includes data and/or waivers of data for only:</i></p> <ul style="list-style-type: none"> ● <i>product chemistry and/or</i> ● <i>acute toxicity and/or</i> ● <i>public health pest efficacy and/or</i> ● <i>animal safety studies and/or</i> ● <i>child resistant packaging.</i> (2)(3) 	10	8,820

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) An application for a new end-use product using a source of active ingredient that (a) is not yet registered but (b) has an application pending with the Agency for review, will be considered an application for a new product with an unregistered source of active ingredient.

(3) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant’s written or electronic confirmation of agreement to the Agency.

“TABLE 14. — BIOPESTICIDES DIVISION — AMENDMENTS

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Reg-istration Service Fee (\$)
B621	134	Amendment; Experimental Use Permit; no change to an established temporary tolerance or tolerance exemption. (3)	7	5,107
B622	135	Amendment; Experimental Use Permit; petition to amend an established or temporary tolerance or tolerance exemption. (3)	11	12,764
B641	136	Amendment of an established tolerance or tolerance exemption.	13	12,764
B680	137	Amendment; registered sources of active ingredient(s); no new use(s); no changes to an established tolerance or tolerance exemption. Requires data submission. (2)(3)	5	5,107
B681	138	Amendment; unregistered source of active ingredient(s). Requires data submission. (2)(3)	7	6,079
B683	139	Label amendment; requires review/update of previous risk assessment(s) without data submission (e.g., labeling changes to REI, PPE, PHI). (2)(3)	6	5,107
B684	140	Amending non-food animal product that includes submission of target animal safety data; previously registered. (2)(3)	8	8,820
B685	141 (new)	Amendment; add a new biochemical unregistered source of active ingredient or a new microbial production site. Requires submission of analysis of samples data and source/production site-specific manufacturing process description. (3)	5	5,107

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) (a) EPA-initiated amendments shall not be charged registration service fees. (b) Registrant-initiated fast-track amendments are to be completed within the timelines specified in FIFRA Section 3(c)(3)(B) and are not subject to registration service fees. (c) Registrant-initiated fast-track amendments handled by the Antimicrobials Division are to be completed within the timelines specified in FIFRA Section 3(h) and are not subject to registration service fees. (d) Registrant initiated amendments submitted by notification under PR Notices, such as PR Notice 98-10, continue under PR Notice timelines and are not subject to registration service fees. (e) Submissions with data and requiring data review are subject to registration service fees.

(3) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

“TABLE 15. — BIOPESTICIDES DIVISION — SCLP

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
<i>B690</i>	<i>142</i>	<i>New active ingredient; food or non-food use. (2)(6)</i>	<i>7</i>	<i>2,554</i>
<i>B700</i>	<i>143</i>	<i>Experimental Use Permit application; new active ingredient or new use. (6)</i>	<i>7</i>	<i>1,278</i>
<i>B701</i>	<i>144</i>	<i>Extend or amend Experimental Use Permit. (6)</i>	<i>4</i>	<i>1,278</i>

“TABLE 15. — BIOPESTICIDES DIVISION — SCLP—
Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registra- tion Service Fee (\$)
B710	145	<p><i>New product; reg- istered source of active ingre- dient(s); iden- tical or substan- tially similar in composition and use to a reg- istered product; no change in an established toler- ance or tolerance exemption. No data review, or only product chemistry data; cite-all data ci- tation, or selec- tive data cita- tion where ap- plicant owns all required data or authorization from data owner is demonstrated. Category in- cludes 100% re- package of reg- istered end-use or manufac- turing-use prod- uct that requires no data submis- sion or data ma- trix. (3)(6)</i></p>	4	1,278

“TABLE 15. — BIOPESTICIDES DIVISION — SCLP—
Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registra- tion Service Fee (\$)
<i>B720</i>	<i>146</i>	<i>New product; reg- istered source of active ingre- dient(s); re- quires: 1) sub- mission of prod- uct specific data; or 2) citation of previously re- viewed and ac- cepted data; or 3) submission or citation of data generated at gov- ernment expense; or 4) submission or citation of a scientifically- sound rationale based on pub- licly available literature or other relevant information that addresses the data require- ment; or 5) sub- mission of a re- quest for a data requirement to be waived sup- ported by a sci- entifically-sound rationale ex- plaining why the data require- ment does not apply. (3)(6)</i>	<i>5</i>	<i>1,278</i>
<i>B721</i>	<i>147</i>	<i>New product; un- registered source of active ingre- dient. (3)(6)</i>	<i>7</i>	<i>2,676</i>

“TABLE 15. — BIOPESTICIDES DIVISION — SCLP—
Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
B722	148	New use and/or amendment; petition to establish a tolerance or tolerance exemption. (4)(5)(6)	7	2,477
B730	149	Label amendment requiring data submission. (4)(6)	5	1,278

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) All requests for new uses (food and/or nonfood) contained in any application for a new active ingredient or a first food use are covered by the base fee for that new active ingredient or first food use application and retain the same decision time review period as the new active ingredient or first food use application. The application must be received by the agency in one package. The base fee for the category covers a maximum of five new products. Each application for an additional new product registration and new inert approval that is submitted in the new active ingredient application package or first food use application package is subject to the registration service fee for a new product or a new inert approval. All such associated applications that are submitted together will be subject to the new active ingredient or first food use decision review time. In the case of a new active ingredient application, until that new active ingredient is approved, any subsequent application for another new product containing the same active ingredient or an amendment to the proposed labeling will be deemed a new active ingredient application, subject to the registration service fee and decision review time for a new active ingredient. In the case of a first food use application, until that first food use is approved, any subsequent application for an additional new food use or uses will be subject to the registration service fee and decision review time for a first food use. Any information that (a) was neither requested nor required by the Agency, and (b) is submitted by the applicant at the applicant's initiative to support the application after completion of the technical deficiency screening, and (c) is not itself a covered registration application, must be assessed 25% of the full registration service fee for the new active ingredient or first food use application.

(3) An application for a new end-use product using a source of active ingredient that (a) is not yet registered but (b) has an application pending with the Agency for review, will be considered an application for a new product with an unregistered source of active ingredient.

(4) (a) EPA-initiated amendments shall not be charged registration service fees. (b) Registrant-initiated fast-track amendments are to be completed within the timelines specified in FIFRA Section 3(c)(3)(B) and are not subject to registration service fees. (c) Registrant-initiated fast-track amendments handled by the Antimicrobials Division are to be completed within the timelines specified in FIFRA Section 3(h) and are not subject to registration service fees. (d) Registrant initiated amendments submitted by notification under PR Notices, such as PR Notice 98-10, continue under PR Notice timelines and are not subject to registration service fees. (e) Submissions with data and requiring data review are subject to registration service fees.

(5) Amendment applications to add the new use(s) to registered product labels are covered by the base fee for the new use(s). All items in the covered application must be submitted together in one package. Each application for an additional new product registration and new inert approval(s) that is submitted in the new use application package is subject to the registration service fee for a new product or a new inert approval. However, if a new use application only proposes to register the new use for a new product and there are no amendments in the application, then review of one new product application is covered by the new use fee. All such associated applications that are submitted together will be subject to the new use decision review time. Any application for a new product or an amendment to the proposed labeling (a) submitted subsequent to submission of the new use application and (b) prior to conclusion of its decision review time and (c) containing the same new uses, will be deemed a separate new-use application, subject to a separate registration service fee and new decision review time for a new use. If the new-use application includes non-food (indoor and/or outdoor), and food (outdoor and/or indoor) uses, the appropriate fee is due for each type of new use and the longest decision review time applies to all of the new uses requested in the application. Any information that (a) was neither requested nor required by the Agency, and (b) is submitted by the applicant at the applicant's initiative to support the application after completion of the technical deficiency screen, and (c) is not itself a covered registration application, must be assessed 25% of the full registration service fee for the new use application.

(6) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

“TABLE 16. — BIOPESTICIDES DIVISION — OTHER ACTIONS

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
B614	150	Pre-application; Conditional Ruling on rationales for addressing a data requirement in lieu of data; applicant-initiated; applies to one rationale at a time.	3	2,530

“TABLE 16. — BIOPESTICIDES DIVISION — OTHER ACTIONS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B615	151	Rebuttal of agency reviewed protocol, applicant initiated.	3	2,530
B682	152	Protocol review; applicant initiated; excludes time for HSRB review.	3	2,432

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

“TABLE 17. — BIOPESTICIDES DIVISION — PIP

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B740	153	<p>Experimental Use Permit application; no petition for tolerance/tolerance exemption. Includes:</p> <ol style="list-style-type: none"> 1. non-food/feed use(s) for a new (2) or registered (3) PIP (12); 2. food/feed use(s) for a new or registered PIP with crop destruct (12); 3. food/feed use(s) for a new or registered PIP in which an established tolerance/tolerance exemption exists for the intended use(s). <p>(4)(12)</p>	6	95,724

“TABLE 17. — BIOPESTICIDES DIVISION — PIP—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B741	154 (new)	<p><i>Experimental Use Permit application; no petition for tolerance/tolerance exemption. Includes:</i></p> <ol style="list-style-type: none"> <i>1. non-food/feed use(s) for a new (2) or registered (3) PIP;</i> <i>2. food/feed use(s) for a new or registered PIP with crop destruct;</i> <i>3. food/feed use(s) for a new or registered PIP in which an established tolerance/tolerance exemption exists for the intended use(s);</i> <p><i>SAP Review. (12)</i></p>	12	159,538
B750	155	<p><i>Experimental Use Permit application; with a petition to establish a temporary or permanent tolerance/tolerance exemption for the active ingredient. Includes new food/feed use for a registered (3) PIP.</i></p> <p><i>(4)(12)</i></p>	9	127,630

“TABLE 17. — BIOPESTICIDES DIVISION — PIP—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B770	156	<i>Experimental Use Permit application; new (2) PIP; with petition to establish a temporary tolerance/tolerance exemption for the active ingredient; credit 75% of B771 fee toward registration application for a new active ingredient that follows; SAP review. (5)(12)</i>	15	191,444
B771	157	<i>Experimental Use Permit application; new (2) PIP; with petition to establish a temporary tolerance/tolerance exemption for the active ingredient; credit 75% of B771 fee toward registration application for a new active ingredient that follows. (12)</i>	10	127,630
B772	158	<i>Application to amend or extend an Experimental Use Permit; no petition since the established tolerance/tolerance exemption for the active ingredient is unaffected. (12)</i>	3	12,764

“TABLE 17. — BIOPESTICIDES DIVISION — PIP—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B773	159	<i>Application to amend or extend an Experimental Use Permit; with petition to extend a temporary tolerance/tolerance exemption for the active ingredient. (12)</i>	5	31,910
B780	160	<i>Registration application; new (2) PIP; non-food/feed. (12)</i>	12	159,537
B790	161	<i>Registration application; new (2) PIP; non-food/feed; SAP review. (5)(12)</i>	18	223,351
B800	162	<i>Registration application; new (2) PIP; with petition to establish permanent tolerance/tolerance exemption for the active ingredient based on an existing temporary tolerance/tolerance exemption. (12)</i>	13	172,300
B810	163	<i>Registration application; new (2) PIP; with petition to establish permanent tolerance/tolerance exemption for the active ingredient based on an existing temporary tolerance/tolerance exemption. SAP review. (5)(12)</i>	19	236,114

"TABLE 17. — BIOPESTICIDES DIVISION — PIP—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B820	164	Registration application; new (2) PIP; with petition to establish or amend a permanent tolerance/tolerance exemption of an active ingredient. (12)	15	204,208
B840	165	Registration application; new (2) PIP; with petition to establish or amend a permanent tolerance/tolerance exemption of an active ingredient. SAP review. (5)(12)	21	268,022
B851	166	Registration application; new event of a previously registered PIP active ingredient(s); no petition since permanent tolerance/tolerance exemption is already established for the active ingredient(s). (12)	9	127,630
B870	167	Registration application; registered (3) PIP; new product; new use; no petition since a permanent tolerance/tolerance exemption is already established for the active ingredient(s). (4) (12)	9	38,290

“TABLE 17. — BIOPESTICIDES DIVISION — PIP—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B880	168	Registration application; registered (3) PIP; new product or new terms of registration; additional data submitted; no petition since a permanent tolerance/tolerance exemption is already established for the active ingredient(s). (6) (7) (12)	9	31,910
B881	169	Registration application; registered (3) PIP; new product or new terms of registration; additional data submitted; no petition since a permanent tolerance/tolerance exemption is already established for the active ingredient(s). SAP review. (5)(6)(7)(12)	15	95,724
B882	170 (new)	Registration application; new (2) PIP, seed increase with negotiated acreage cap and time-limited registration; with petition to establish a permanent tolerance/tolerance exemption for the active ingredient based on an existing temporary tolerance/tolerance exemption; SAP Review. (8)(12)	15	191,444

“TABLE 17. — BIOPESTICIDES DIVISION — PIP—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B883	171	<i>Registration application; new (2) PIP, seed increase with negotiated acreage cap and time-limited registration; with petition to establish a permanent tolerance/tolerance exemption for the active ingredient based on an existing temporary tolerance/tolerance exemption. (8) (12)</i>	9	127,630
B884	172	<i>Registration application; new (2) PIP, seed increase with negotiated acreage cap and time-limited registration; with petition to establish a permanent tolerance/tolerance exemption for the active ingredient. (8)(12)</i>	12	159,537
B885	173	<i>Registration application; registered (3) PIP, seed increase; breeding stack of previously approved PIPs, same crop; no petition since a permanent tolerance/tolerance exemption is already established for the active ingredient(s). (9)(12)</i>	6	31,910

“TABLE 17. — BIOPESTICIDES DIVISION — PIP—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B886	174 (new)	Registration application; new (2) PIP, seed increase with negotiated acreage cap and time-limited registration; with petition to establish a permanent tolerance/tolerance exemption for the active ingredient. SAP Review. (8) (12)	18	223,351
B890	175	Application to amend a seed increase registration; converts registration to commercial registration; no petition since permanent tolerance/tolerance exemption is already established for the active ingredient(s). (12)	9	63,816
B891	176	Application to amend a seed increase registration; converts registration to a commercial registration; no petition since a permanent tolerance/tolerance exemption already established for the active ingredient(s); SAP review. (5)(12)	15	127,630

“TABLE 17. — BIOPESTICIDES DIVISION — PIP—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B900	177	Application to amend a registration, including actions such as extending an expiration date, modifying an IRM plan, or adding an insect to be controlled. (10)(11)(12)	6	12,764
B901	178	Application to amend a registration, including actions such as extending an expiration date, modifying an IRM plan, or adding an insect to be controlled. SAP review. (10) (11) (12)	12	76,578
B902	179	PIP Protocol review.	3	6,383
B903	180	Inert ingredient tolerance exemption; e.g., a marker such as NPT II; reviewed in BPPD.	6	63,816
B904	181	Import tolerance or tolerance exemption; processed commodities/food only (inert or active ingredient).	9	127,630
B905	182 (new)	SAP Review.	6	63,816
B906	183 (new)	Petition to establish a temporary tolerance/tolerance exemption for one or more active ingredients.	3	31,907

“TABLE 17. — BIOPESTICIDES DIVISION — PIP—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
B907	184 (new)	<i>Petition to establish a temporary tolerance/tolerance exemption for one or more active ingredients based on an existing temporary tolerance/tolerance exemption.</i>	3	12,764
B908	185 (new)	<i>Petition to establish a temporary tolerance/tolerance exemption for one or more active ingredients or inert ingredients.</i>	3	44,671

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) New PIP = a PIP with an active ingredient that has not been registered.

(3) Registered PIP = a PIP with an active ingredient that is currently registered.

(4) Transfer registered PIP through conventional breeding for new food/feed use, such as from field corn to sweet corn.

(5) The scientific data involved in this category are complex. EPA often seeks technical advice from the Scientific Advisory Panel on risks that pesticides pose to wildlife, farm workers, pesticide applicators, non-target species, as well as insect resistance, and novel scientific issues surrounding new technologies. The scientists of the SAP neither make nor recommend policy decisions. They provide advice on the science used to make these decisions. Their advice is invaluable to the EPA as it strives to protect humans and the environment from risks posed by pesticides. Due to the time it takes to schedule and prepare for meetings with the SAP, additional time and costs are needed.

(6) Registered PIPs stacked through conventional breeding.

(7) Deployment of a registered PIP with a different IRM plan (e.g., seed blend).

(8) The negotiated acreage cap will depend upon EPA’s determination of the potential environmental exposure, risk(s) to non-target organisms, and the risk of targeted pest developing resistance to the pesticidal substance. The uncertainty of these risks may reduce the allowable acreage, based upon the quantity and type of non-target organism data submitted and the lack of insect resistance management data, which is usually not required for seed-increase registrations. Registrants are encouraged to consult with EPA prior to submission of a registration application in this category.

(9) Application can be submitted prior to or concurrently with an application for commercial registration.

(10) For example, IRM plan modifications that are applicant-initiated.

(11) EPA-initiated amendments shall not be charged fees.

(12) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

“TABLE 18. — INERT INGREDIENTS

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
I001	186	Approval of new food use inert ingredient. (2)(3)	13	27,000
I002	187	Amend currently approved inert ingredient tolerance or exemption from tolerance; new data. (2)	11	7,500
I003	188	Amend currently approved inert ingredient tolerance or exemption from tolerance; no new data. (2)	9	3,308
I004	189	Approval of new non-food use inert ingredient. (2)	6	11,025
I005	190	Amend currently approved non-food use inert ingredient with new use pattern; new data. (2)	6	5,513

“TABLE 18. — INERT INGREDIENTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
I006	191	<i>Amend currently approved non-food use inert ingredient with new use pattern; no new data. (2)</i>	3	3,308
I007	192	<i>Approval of substantially similar non-food use inert ingredients when original inert is compositionally similar with similar use pattern. (2)</i>	4	1,654
I008	193	<i>Approval of new or amended polymer inert ingredient, food use. (2)</i>	5	3,749
I009	194	<i>Approval of new or amended polymer inert ingredient, non-food use. (2)</i>	4	3,087
I010	195	<i>Petition to amend a single tolerance exemption descriptor, or single non-food use descriptor, to add ≤ 10 CASRNs; no new data. (2)</i>	6	1,654
I011	196 <i>(new)</i>	<i>Approval of new food use safener with tolerance or exemption from tolerance. (2)(8)</i>	24	597,683
I012	197 <i>(new)</i>	<i>Approval of new non-food use safener. (2)(8)</i>	21	415,241

“TABLE 18. — INERT INGREDIENTS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
I013	198 (new)	Approval of additional food use for previously approved safener with tolerance or exemption from tolerance. (2)	15	62,975
I014	199 (new)	Approval of additional non-food use for previously approved safener. (2)	15	25,168
I015	200 (new)	Approval of new generic data for previously approved food use safener. (2)	24	269,728
I016	201 (new)	Approval of amendment(s) to tolerance and label for previously approved safener. (2)	13	55,776

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) If another covered application is submitted that depends upon an application to approve an inert ingredient, each application will be subject to its respective registration service fee. The decision review time line for both submissions will be the longest of the associated applications. If the application covers multiple ingredients grouped by EPA into one chemical class, a single registration service fee will be assessed for approval of those ingredients.

(3) If EPA data rules are amended to newly require clearance under section 408 of the FFDCA for an ingredient of an antimicrobial product where such ingredient was not previously subject to such a clearance, then review of the data for such clearance of such product is not subject to a registration service fee for the tolerance action for two years from the effective date of the rule.

(4) Any other covered application that is associated with and dependent on the HSRB review will be subject to its separate registration service fee. The decision review times for the associated actions run concurrently, but will end at the date of the latest review time.

(5) Any other covered application that is associated with and dependent on the SAP review will be subject to its separate registration service fee. The decision review time for the associated action will be extended by the decision review time for the SAP review.

(6) An application for a new end-use product using a source of active ingredient that (a) is not yet registered but (b) has an application pending with the Agency for review, will be considered an application for a new product with an unregistered source of active ingredient.

(7) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

(8) If a new safener is submitted in the same package as a new active ingredient, and that new active ingredient is determined to be reduced risk, then the safener would get the same reduced timeframe as the new active ingredient.

“TABLE 19. — EXTERNAL REVIEW AND MISCELLANEOUS ACTIONS

EPA No.	New CR No.	Action	Decision Review Time (Months)⁽¹⁾	Registration Service Fee (\$)
M001	202	Study protocol requiring Human Studies Review Board review as defined in 40 CFR Part 26 in support of an active ingredient. (4)	9	7,938
M002	203	Completed study requiring Human Studies Review Board review as defined in 40 CFR Part 26 in support of an active ingredient. (4)	9	7,938

“TABLE 19. — EXTERNAL REVIEW AND MISCELLANEOUS ACTIONS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
M003	204	<i>External technical peer review of new active ingredient, product, or amendment (e.g., consultation with FIFRA Scientific Advisory Panel) for an action with a decision time-frame of less than 12 months. Applicant initiated request based on a requirement of the Administrator, as defined by FIFRA § 25(d), in support of a novel active ingredient, or unique use pattern or application technology. Excludes PIP active ingredients. (5)</i>	12	63,945

“TABLE 19. — EXTERNAL REVIEW AND MISCELLANEOUS ACTIONS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
M004	205	<i>External technical peer review of new active ingredient, product, or amendment (e.g., consultation with FIFRA Scientific Advisory Panel) for an action with a decision time-frame of greater than 12 months. Applicant initiated request based on a requirement of the Administrator, as defined by FIFRA § 25(d), in support of a novel active ingredient, or unique use pattern or application technology. Excludes PIP active ingredients. (5)</i>	18	63,945

“TABLE 19. — EXTERNAL REVIEW AND MISCELLANEOUS ACTIONS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
M005	206	<i>New Product: Combination, Contains a combination of active ingredients from a registered and/or unregistered source; conventional, anti-microbial and/or biopesticide. Requires coordination with other regulatory divisions to conduct review of data, label and/or verify the validity of existing data as cited. Only existing uses for each active ingredient in the combination product. (6)(7)</i>	9	22,050
M006	207	<i>Request for up to 5 letters of certification (Gold Seal) for one actively registered product (excludes distributor products). (8)</i>	1	277
M007	208	<i>Request to extend Exclusive Use of data as provided by FIFRA Section 3(c)(1)(F)(ii).</i>	12	5,513

“TABLE 19. — EXTERNAL REVIEW AND MISCELLANEOUS ACTIONS—Continued

EPA No.	New CR No.	Action	Decision Review Time (Months)₍₁₎	Registration Service Fee (\$)
M008	209	Request to grant Exclusive Use of data as provided by FIFRA Section 3(c)(1)(F)(vi) for a minor use, when a FIFRA Section 2(l)(2) determination is required.	15	1,654
M009	210 (new)	Non-FIFRA Regulated Determination: Applicant initiated, per product.	4	2,363
M010	211 (new)	Conditional ruling on pre-application, product substantial similarity.	4	2,363
M011	212 (new)	Label amendment to add the DfE logo; requires data review; no other label changes. (9)	4	3,648

(1) A decision review time that would otherwise end on a Saturday, Sunday, or federal holiday, will be extended to end on the next business day.

(2) If another covered application is submitted that depends upon an application to approve an inert ingredient, each application will be subject to its respective registration service fee. The decision review time line for both submissions will be the longest of the associated applications. If the application covers multiple ingredients grouped by EPA into one chemical class, a single registration service fee will be assessed for approval of those ingredients.

(3) If EPA data rules are amended to newly require clearance under section 408 of the FFDCA for an ingredient of an antimicrobial product where such ingredient was not previously subject to such a clearance, then review of the data for such clearance of such product is not subject to a registration service fee for the tolerance action for two years from the effective date of the rule.

(4) Any other covered application that is associated with and dependent on the HSRB review will be subject to its separate registration service fee. The decision review times for the associated actions run concurrently, but will end at the date of the latest review time.

(5) Any other covered application that is associated with and dependent on the SAP review will be subject to its separate registration service fee. The decision review time for the associated action will be extended by the decision review time for the SAP review.

(6) An application for a new end-use product using a source of active ingredient that (a) is not yet registered but (b) has an application pending with the Agency for review, will be considered an application for a new product with an unregistered source of active ingredient.

(7) Where the action involves approval of a new or amended label, on or before the end date of the decision review time, the Agency shall provide to the applicant a draft accepted label, including any changes made by the Agency that differ from the applicant-submitted label and relevant supporting data reviewed by the Agency. The applicant will notify the Agency that the applicant either (a) agrees to all of the terms associated with the draft accepted label as amended by the Agency and requests that it be issued as the accepted final Agency-stamped label; or (b) does not agree to one or more of the terms of the draft accepted label as amended by the Agency and requests additional time to resolve the difference(s); or (c) withdraws the application without prejudice for subsequent resubmission, but forfeits the associated registration service fee. For cases described in (b), the applicant shall have up to 30 calendar days to reach agreement with the Agency on the final terms of the Agency-accepted label. If the applicant agrees to all of the terms of the accepted label as in (a), including upon resolution of differences in (b), the Agency shall provide an accepted final Agency-stamped label to the registrant within 2 business days following the registrant's written or electronic confirmation of agreement to the Agency.

(8) Due to low fee and short time frame this category is not eligible for small business waivers. Gold seal applies to one registered product.

(9) This category includes amendments the sole purpose of which is to add DfE (or equivalent terms that do not use "safe" or derivatives of "safe") logos to a label. DfE is a voluntary program. A label bearing a DfE logo is not considered an Agency endorsement because the ingredients in the qualifying product must meet objective, scientific criteria established and widely publicized by EPA."

1 **SEC. 7. AGRICULTURAL WORKER PROTECTION STANDARD;**

2 **CERTIFICATION OF PESTICIDE APPLICATORS.**

3 (a) *IN GENERAL.*—*Except as provided in subsection*
 4 *(b), during the period beginning on the date of enactment*
 5 *of this Act and ending not earlier than October 1, 2021,*
 6 *the Administrator of the Environmental Protection Agency*
 7 *(referred to in this section as the "Administrator")—*

8 (1) *shall carry out—*

9 (A) *the final rule of the Administrator enti-*
 10 *tled "Pesticides; Agricultural Worker Protection*
 11 *Standard Revisions" (80 Fed. Reg. 67496 (No-*
 12 *vember 2, 2015)); and*

1 (B) the final rule of the Administrator enti-
2 tled “Pesticides; Certification of Pesticide Appli-
3 cators” (82 Fed. Reg. 952 (January 4, 2017));
4 and

5 (2) shall not revise or develop revisions to the
6 rules described in subparagraphs (A) and (B) of
7 paragraph (1).

8 (b) *EXCEPTIONS.*—Prior to October 1, 2021, the Ad-
9 ministrators may propose, and after a notice and public
10 comment period of not less than 90 days, promulgate revi-
11 sions to the final rule described in subsection (a)(1)(A) ad-
12 dressing application exclusion zones under part 170 of title
13 40, Code of Federal Regulations, consistent with the Federal
14 Insecticide, Fungicide, and Rodenticide Act (7 U.S.C. 136
15 et seq.).

16 (c) *GAO REPORT.*—The Comptroller General of the
17 United States shall—

18 (1) conduct a study on the use of the designated
19 representative, including the effect of that use on the
20 availability of pesticide application and hazard infor-
21 mation and worker health and safety; and

22 (2) not later than October 1, 2021, make pub-
23 lically available a report describing the study under
24 paragraph (1), including any recommendations to

- 1 *prevent the misuse of pesticide application and haz-*
- 2 *ard information, if that misuse is identified.*

Attest:

Clerk.

116TH CONGRESS
1ST SESSION

S. 483

AMENDMENT