Attachment J

Pesticide Registration Improvement Renewal Act (PRIA 2)
Tables - FY 2009/FY 2010 Fee Schedule for Registration Applications

The following PRIA fee schedule tables were published in the Federal Register of August 5, 2008 (Volume 73, Number 151) Page 45438-45450 and apply to pesticide registration applications received by the Agency on October 1, 2008 through September 30, 2010.

How to Read the PRIA Fee Tables

1. Each table consists of the following columns:

    - The column entitled "EPA No." assigns an EPA identifier to each fee category. There are 140 categories spread across the 3 Divisions. There are 58 RD categories, 27 AD categories, and 55 BPPD categories. For tracking purposes, OPP has assigned a 3-digit identifier to each category, beginning with RD categories, followed by AD and BPPD categories. The categories are prefaced with a letter designation indicating which Division of OPP is responsible for applications in that category (R= Registration Division, A=Antimicrobials Division, B=Biopesticides and Pollution Prevention Division). **A detailed explanation of the type of application that falls under a fee category is linked to the fee category number. When accessed, the link will go to the appropriate page of the Fee Determination Decision Tree with the fee category interpretation.**

    - The column entitled "CR No." cross-references the current Congressional Record category number for convenience. However, EPA will be using the categories as numbered in the "EPA No." column in its tracking systems.

    - The column entitled "Action" describes the categories of action. In establishing the expanded fee schedule categories, Congress eliminated some of the more confusing terminology of the original categories. For example, instead of the term "fast-track," the schedule in the Congressional Record uses the regulatory phrase "identical or substantially similar in composition and use to a registered product." Further information may be obtained by accessing the link provided with the fee category in the first column.

    - The column entitled "Decision Time" lists the decision times in months for each type of action for Fiscal Years 2009 and 2010. The 2010 decision times apply to 2011 and 2012. The decision review periods in the tables are based upon EPA fiscal years (FY), which run from October 1 through September 30.

    - The column entitled "FY 09/FY 10 Registration Service Fee ($)" lists the registration service fee for the action for fiscal year 2009 (October 1, 2008 through September 30, 2009) and fiscal year 2010 (October 1, 2009 through September 30, 2010).

2. The following acronyms are used in some of the tables:

    - DART - Dose Adequacy Response Team
    - DNT - Developmental Neurotoxicity
    - HSRB - Human Studies Review Board
    - GW/SW - Ground Water/Surface Water
    - PHI - Pre-Harvest Interval
    - PPE - Personal Protective Equipment
    - REI - Restricted Entry Interval
    - SAP - FIFRA Scientific Advisory Panel

Fee Schedule Tables - Effective October 1, 2008

- A. Registration Division
- B. Antimicrobials Division
- C. Biopesticides and Pollution Prevention Division

A. Registration Division (RD)

The Registration Division of OPP is responsible for the processing of pesticide applications and associated tolerance petitions for pesticides that are termed "conventional chemicals," excluding pesticides intended for antimicrobial uses. The term "conventional chemical" is a term of art that is intended to distinguish synthetic chemicals from those that are of naturally occurring or non-synthetic origin, synthetic chemicals that are identical to naturally-occurring chemicals and microbial pesticides. Tables 1 through 5 cover RD actions.

TABLE 1. REGISTRATION DIVISION–NEW ACTIVE INGREDIENTS

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R010	1	Food use[1]	24	24	542,115
R020	2	Food use; reduced risk[1]	18	18	542,115
R030	3	Food use; Experimental Use Permit application submitted simultaneously with application for registration; decision time for Experimental Use Permit and temporary tolerance same as #R040[1]	24	24	599,235
R040	4	Food use; Experimental Use Permit application; establish temporary tolerance; submitted before application for registration; credit $326,025 toward new active ingredient application that follows	18	18	399,525
R050	5	Food use; application submitted after Experimental Use Permit application; decision time begins after Experimental Use Permit and temporary tolerance are granted[1]	14	14	199,815
R060	6	Non-food use; outdoor[1]	21	21	376,635
R070	7	Non-food use; outdoor; reduced risk[1]	16	16	376,635

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R080	8	Non-food use; outdoor; Experimental Use Permit application submitted simultaneously with application for registration; decision time for Experimental Use Permit same as #R090[1]	21	21	416,640
R090	9	Non-food use; outdoor; Experimental Use Permit application submitted before application for registration; credit $228,225 toward new active ingredient application that follows	16	16	279,615
R100	10	Non-food use; outdoor; submitted after Experimental Use Permit application; decision time begins after Experimental Use Permit is granted [1]	12	12	137,025
R110	11	Non-food use; indoor[1]	20	20	209,475
R120	12	Non-food use; indoor; reduced risk[1]	14	14	209,475
R121	13	Non-food use; indoor; Experimental Use Permit application submitted before application for registration; credit $100,000 toward new active ingredient application that follows	18	18	157,500
R122	14	Enriched isomer(s) of registered mixed-isomer active ingredient[1]	18	18	273,945
R123	15	Seed treatment only; includes non-food and food uses; limited uptake into Raw Agricultural Commodities[1]	18	18	407,610
R124	16	Conditional Ruling on Preapplication Study Waivers; applicant-initiated	6	6	2,184

[1]All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

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TABLE 2. REGISTRATION DIVISION–NEW USES

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R130	17	First food use; indoor; food/food handling[1]	21	21	165,375
R140	18	Additional food use; Indoor; food/food handling	15	15	38,588
R150	19	First food use[1]	21	21	228,270

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R160	20	First food use; reduced risk[1]	16	16	228,270
R170	21	Additional food use	15	15	57,120
R180	22	Additional food use; reduced risk	10	10	57,120
R190	23	Additional food uses; 6 or more submitted in one application	15	15	342,720
R200	24	Additional food uses; 6 or more submitted in one application; reduced risk	10	10	342,720
R210	25	Additional food use; Experimental Use Permit application; establish temporary tolerance; no credit toward new use registration	12	12	42,315
R220	26	Additional food use; Experimental Use Permit application; crop destruct basis; no credit toward new use registration	6	6	17,136
R230	27	Additional use; non-food; outdoor	15	15	22,827
R240	28	Additional use; non-food; outdoor; reduced risk	10	10	22,827
R250	29	Additional use; non-food; outdoor; Experimental Use Permit application; no credit toward new use registration	6	6	17,136
R260	30	New use; non-food; indoor	12	12	11,025
R270	31	New use; non-food; indoor; reduced risk	9	9	11,025
R271	32	New use; non-food; indoor; Experimental Use Permit application; no credit toward new use registration	6	6	8,400
R272	33	Review of Study Protocol; applicant-initiated; excludes DART, pre-registration conferences, Rapid Response review, DNT protocol review, protocols needing HSRB review	3	3	2,184
R273	34	Additional use; seed treatment; limited uptake into Raw Agricultural Commodities; includes crops with established tolerances (e.g., for soil or foliar application); includes food or non-food uses	12	12	43,575
R274	35	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	12	12	261,450

[1] All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

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TABLE 3. REGISTRATION DIVISION–IMPORT AND OTHER TOLERANCES

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R280	36	Establish import tolerance; new active ingredient or first food use[1]	21	21	275,625
R290	37	Establish import tolerance; additional food use	15	15	55,125
R291	38	Establish import tolerances; additional food uses; 6 or more crops submitted in one petition	15	15	330,750
R292	39	Amend an established tolerance (e.g., decrease or increase); domestic or import; applicant-initiated	10	10	39,165
R293	40	Establish tolerance(s) for inadvertent residues in one crop; applicant-initiated	12	12	46,200
R294	41	Establish tolerances for inadvertent residues; 6 or more crops submitted in one application; applicant-initiated	12	12	277,200
R295	42	Establish tolerance(s) for residues in one rotational crop in response to a specific rotational crop application; applicant-initiated	15	15	57,120
R296	43	Establish tolerances for residues in rotational crops in response to a specific rotational crop petition; 6 or more crops submitted in one application; applicant-initiated	15	15	342,720

[1] All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

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TABLE 4. REGISTRATION DIVISION–NEW PRODUCTS

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R300	44	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	3	3	1,365

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
		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.			
R301	45	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.	4	4	1,638
R310	46	New end-use or manufacturing-use product; requires review of data package within RD; includes reviews and/or waivers of data for only: product chemistry and/or acute toxicity and/or public health pest efficacy	6	6	4,578
R311	49	New product; requires approval of new food-use inert; applicant-initiated; excludes approval of safeners	12	12	16,317
R312	50	New product; requires approval of new non-food-use inert; applicant-initiated	6	6	8,715
R313	51	New product; requires amendment to existing inert tolerance exemption (e.g., adding post-harvest use); applicant-initiated	10	10	11,991
R320	47	New product; new physical form; requires data review in science divisions	12	12	11,424
R330	48	New manufacturing-use product; registered active ingredient; selective data citation	12	12	17,136
R331	52	New product; repack of identical registered end-use product as a manufacturing-use product; same registered uses only	3	3	2,184
R332	53	New manufacturing-use product; registered active ingredient; unregistered source of active ingredient; submission of completely new generic data package; registered uses only	24	24	244,650

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TABLE 5. REGISTRATION DIVISION–AMENDMENTS TO REGISTRATION

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
R340	54	Amendment requiring data review within RD (e.g., changes to precautionary label statements, or source changes to an unregistered source of active ingredient)[1]	4	4	3,444
R350	55	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)[1]	8	8	11,424
R370	56	Cancer reassessment; applicant-initiated	18	18	171,255
R371	57	Amendment to Experimental Use Permit; requires data review / risk assessment	6	6	8,715
R372	58	Refined ecological and/or endangered species assessment; applicant-initiated	18	12	163,065

[1]EPA-initiated amendments shall not be charged fees. Fast-track amendments handled by the Antimicrobials Division are to be completed within the FIFRA stated timelines listed in Section 3(h) and are not subject to PRIA fees. Label amendments submitted by notification under PR Notices, such as PR Notice 95–2 and PR Notice 98–10, continue under PR Notice timelines and are not subject to PRIA fees.

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B. Antimicrobials Division (AD)

The Antimicrobials Division of OPP is responsible for the processing of pesticide applications and associated tolerances for conventional chemicals intended for antimicrobial uses, that is, uses that are defined under FIFRA section 2(mm)(1)(A), including products for use against bacteria, protozoa, non-agricultural fungi, and viruses. AD is also responsible for a selected set of conventional chemicals intended for other uses, including most wood preservatives and antifoulants. Tables 6 through 8 cover AD actions.

TABLE 6. ANTIMICROBIALS DIVISION–NEW ACTIVE INGREDIENTS

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
A380	59	Food use; establish tolerance exemption[1]	24	24	99,225
A390	60	Food use; establish tolerance[1]	24	24	165,375
A400	61	Non-food use; outdoor; FIFRA §2(mm) uses[1]	18	18	82,688

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
A410	62	Non-food use; outdoor; uses other than FIFRA §2(mm)[1]	21	21	165,375
A420	63	Non-food use; indoor; FIFRA §2(mm) uses[1]	18	18	55,125
A430	64	Non-food use; indoor; uses other than FIFRA §2(mm)[1]	20	20	82,688
A431	65	Non-food use; indoor; low-risk and low-toxicity food-grade active ingredient(s); efficacy testing for public health claims required under GLP and following DIS/TSS or AD-approved study protocol	12	12	57,750

[1]All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

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TABLE 7. ANTIMICROBIALS DIVISION–NEW USES

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
A440	66	First food use; establish tolerance exemption[1]	21	21	27,563
A450	67	First food use; establish tolerance [1]	21	21	82,688
A460	68	Additional food use; establish tolerance exemption	15	15	11,025
A470	69	Additional food use; establish tolerance	15	15	27,563
A480	70	Additional use; non-food; outdoor; FIFRA §2(mm) uses	9	9	16,538
A490	71	Additional use; non-food; outdoor; uses other than FIFRA §2(mm)	15	15	27,563
A500	72	Additional use; non-food; indoor; FIFRA §2(mm) uses	9	9	11,025
A510	73	Additional use; non-food; indoor; uses other than FIFRA §2(mm)	12	12	11,025
A520	74	Experimental Use Permit application	9	9	5,513
A521	75	Review of public health efficacy study protocol within AD; per AD Internal Guidance for the Efficacy Protocol Review Process; applicant-initiated; Tier 1	4	3	2,100
A522	76	Review of public health efficacy study protocol	15	12	10,500

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
		outside AD by members of AD Efficacy Protocol Review Expert Panel; applicant-initiated; Tier 2			

[1]All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

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TABLE 8. ANTIMICROBIALS DIVISION–NEW PRODUCTS AND AMENDMENTS

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
A530	77	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.	3	3	1,103
A531	78	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.	4	4	1,575
A532	85	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	4	4	4,410
A540	79	New end use product; FIFRA §2(mm) uses only	4	4	4,410
A550	80	New end-use product; uses other than FIFRA §2(mm); non-FQPA product	6	6	4,410
A560	81	New manufacturing-use product; registered active	12	12	16,538

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
		ingredient; selective data citation			
A570	82	Label amendment requiring data submission [1]	4	4	3,308
A571	83	Cancer reassessment; applicant-initiated	18	18	82,688
A572	84	Refined ecological risk and/or endangered species assessment; applicant-initiated	18	12	78,750

[1]EPA-initiated amendments shall not be charged fees. Fast-track amendments handled by the Antimicrobials Division are to be completed within the FIFRA stated timelines listed in Section 3(h) and are not subject to PRIA fees. Label amendments submitted by notification under PR Notices, such as PR Notice 95–2 and PR Notice 98–10, continue under PR Notice timelines and are not subject to PRIA fees.

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C. Biopesticides and Pollution Prevention Division (BPPD)

The Biopesticides and Pollution Prevention Division of OPP is responsible for the processing of pesticide applications for biochemical pesticides, microbial pesticides, and plant-incorporated protectants (PIPs).

The fee tables for BPPD tables are presented by type of pesticide rather than by type of action: Microbial and biochemical pesticides, straight chain lepidopteran pheromones (SCLPs), and PIPs. Within each table, the types of application are the same as those in other divisions and use the same terminology as in Unit III. Tables 9 through 11 cover BPPD actions.

TABLE 9. BIOPESTICIDES AND POLLUTION PREVENTION DIVISION–MICROBIAL AND BIOCHEMICAL PESTICIDES; NEW PRODUCTS AND AMENDMENTS

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
B580	86	New active ingredient; food use; establish tolerance[1]	18	18	44,100
B590	87	New active ingredient; food use; establish tolerance exemption [1]	16	16	27,563
B600	88	New active ingredient; non-food use[1]	12	12	16,538
B610	89	Food use; Experimental Use Permit application; establish temporary tolerance exemption	9	9	11,025
B620	90	Non-food use; Experimental Use Permit application	6	6	5,513
B621	91	Extend or amend Experimental Use Permit	6	6	4,410

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
B630	92	First food use; establish tolerance exemption	12	12	11,025
B631	93	Amend established tolerance exemption	9	9	11,025
B640	94	First food use; establish tolerance [1]	18	18	16,538
B641	95	Amend established tolerance (e.g., decrease or increase)	12	12	11,025
B650	96	New use; non-food	6	6	5,513
B660	97	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.	3	3	1,103
B670	98	New product; registered source of active ingredient; all Tier I data for product chemistry, toxicology, non-target organisms, and product performance must be addressed with product specific data or with request for data waivers supported by scientific rationales	6	6	4,410
B671	99	New product; food use; unregistered source of active ingredient; requires amendment of established tolerance or tolerance exemption; all Tier I data requirements for product chemistry, toxicology, non-target organisms, and product performance must be addressed with product-specific data or with request for data waivers supported by scientific rationales	16	16	11,025
B672	100	New product; non-food use or food use having established tolerance or tolerance exemption; unregistered source of active ingredient; no data compensation issues; all Tier I data requirements for product chemistry, toxicology, non-target organisms, and product performance must be addressed with product-specific data or with request for data waivers supported by scientific rationales	12	12	7,875
B680	101	Label amendment requiring data submission[2]	4	4	4,410
B681	102	Label amendment; unregistered source of active	6	6	5,250

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
		ingredient; supporting data require scientific review			
B682	103	Protocol review; applicant-initiated; excludes time for HSRB review (pre application)	3	3	2,100

[1]All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

[2]EPA-initiated amendments shall not be charged fees. Fast-track amendments handled by the Antimicrobials Division are to be completed within the FIFRA stated timelines listed in Section 3(h) and are not subject to PRIA fees. Label amendments submitted by notification under PR Notices, such as PR Notice 95–2 and PR Notice 98–10, continue under PR Notice timelines and are not subject to PRIA fees.

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TABLE 10. BIOPESTICIDES AND POLLUTION PREVENTION DIVISION–STRAIGHT CHAIN LEPIDOPTERAN PHEROMONES(SCLPS)

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
B690	104	New active ingredient; food or non-food use [1]	6	6	2,205
B700	105	Experimental Use Permit application; new active ingredient or new use	6	6	1,103
B701	106	Extend or amend Experimental Use Permit	3	3	1,103
B710	107	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.	3	3	1,103
B720	108	New product; registered source of active ingredient; all Tier I data for product chemistry, toxicology, non-target organisms, and product performance must be addressed with product specific data or with request for data waivers supported by scientific rationales	4	4	1,103

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
B721	109	New product; unregistered source of active ingredient	6	6	2,310
B722	110	New use and/or amendment to tolerance or tolerance exemption	6	6	2,310
B730	111	Label amendment requiring data submission [2]	4	4	1,103

[1]All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.

[2]EPA-initiated amendments shall not be charged fees. Fast-track amendments handled by the Antimicrobials Division are to be completed within the FIFRA stated timelines listed in Section 3(h) and are not subject to PRIA fees. Label amendments submitted by notification under PR Notices, such as PR Notice 95–2 and PR Notice 98–10, continue under PR Notice timelines and are not subject to PRIA fees.

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TABLE 11. BIOPESTICIDE AND POLLUTION PREVENTION DIVISION–PLANT INCORPORATED PROTECTANTS (PIPS)

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
B740	112	Experimental Use Permit application; registered active ingredient; non-food/feed or crop destruct basis; no SAP review required [1]	6	6	82,688
B750	113	Experimental Use Permit application; registered active ingredient; establish temporary tolerance or tolerance exemption; no SAP review required [1]	9	9	110,250
B760	114	Experimental Use Permit application; new active ingredient; non-food/feed or crop destruct basis; SAP review required; credit $78,750 toward new active ingredient application that follows	12	12	137,813
B761	115	Experimental Use Permit application; new active ingredient; non-food/feed or crop destruct; no SAP review required; credit $78,750 toward new active ingredient application that follows	7	7	82,688
B770	116	Experimental Use Permit application; new active ingredient; establish temporary tolerance or tolerance exemption; SAP review required; credit $105,000	15	15	165,375

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
		toward new active ingredient application that follows			
B771	117	Experimental Use Permit application; new active ingredient; establish temporary tolerance or tolerance exemption; no SAP review required; credit $105,000 toward new active ingredient application that follows	10	10	110,250
B772	118	Amend or extend Experimental Use Permit; minor changes to experimental design; established temporary tolerance or tolerance exemption is unaffected	3	3	11,025
B773	119	Amend or extend existing Experimental Use Permit; minor changes to experimental design; extend established temporary tolerance or tolerance exemption	5	5	27,563
B860	120	Amend Experimental Use Permit; first food use or major revision of experimental design	6	6	11,025
B780	121	New active ingredient; non-food/feed; no SAP review required [2]	12	12	137,813
B790	122	New active ingredient; Non-food/feed; SAP review required [2]	18	18	192,938
B800	123	New active ingredient; establish permanent tolerance or tolerance exemption based on temporary tolerance or tolerance exemption; no SAP review required[2]	12	12	220,500
B810	124	New active ingredient; establish permanent tolerance or tolerance exemption based on temporary tolerance or tolerance exemption; SAP review required[2]	18	18	275,625
B820	125	New active ingredient; establish tolerance or tolerance exemption; no SAP review required [2]	15	15	275,625
B840	126	New active ingredient; establish tolerance or tolerance exemption; SAP review required[2]	21	21	330,750
B830	127	New active ingredient; Experimental Use Permit application submitted simultaneously; establish tolerance or tolerance exemption; no SAP review required [2]	15	15	330,750
B850	128	New active ingredient; Experimental Use Permit requested simultaneously; establish tolerance or tolerance exemption; SAP review required[2]	21	21	385,875
B851	129	New active ingredient; different genetic event of a	9	9	110,250

EPA No.	CR No.	Action	Decision time (months)		FY 09/FY 10 Registration Service Fee ($)
			FY 09	FY 10	
		previously approved active ingredient; same crop; no tolerance action required; no SAP review required			
B852	130	New active ingredient; different genetic event of a previously approved active ingredient; same crop; no tolerance action required; SAP review required	9	9	165,375
B870	131	New use [1]	9	9	33,075
B880	132	New product; no SAP review required[3]	9	9	27,563
B881	133	New product; SAP review required[3]	15	15	82,688
B890	134	Amendment; seed production to commercial registration; no SAP review required	9	9	55,125
B891	135	Amendment; seed production to commercial registration; SAP review required	15	15	110,250
B900	136	Amendment (except #B890); No SAP review required; (e.g., new IRM requirements that are applicant initiated; or amending a conditional registration to extend the registration expiration date with additional data submitted)[4]	6	6	11,025
B901	137	Amendment (except #B890); SAP review required[4]	12	12	66,150
B902	138	PIP Protocol review	3	3	5,513
B903	139	Inert ingredient tolerance exemption; e.g., a marker such as NPT II; reviewed in BPPD	6	6	55,125
B904	140	Import tolerance or tolerance exemption; processed commodities/food only	9	9	110,250

[1]Example: Transfer existing PIP trait by traditional breeding, such as from field corn to sweet corn.
[2]May be either a registration for seed increase or a full commercial registration. If a seed increase registration is granted first, full commercial registration is obtained using B890.
[3]Example: Stacking PIP traits within a crop using traditional breeding techniques.
[4]EPA-initiated amendments shall not be charged fees. Fast-track amendments handled by the Antimicrobials Division are to be completed within the FIFRA stated timelines listed in Section 3(h) and are not subject to PRIA fees. Label amendments submitted by notification under PR Notices, such as PR Notice 95–2 and PR Notice 98–10, continue under PR Notice timelines and are not subject to PRIA fees.