Table I

Number of PRIA Actions Completed in FY 2023 and Average Exceedance in Days, by Category

Key to the Table:

- R - Conventional Pesticides
- A - Antimicrobial Pesticides
- B - Biopesticides
- EUP - Experimental Use Permit
- I – Inert Ingredient
- M – Miscellaneous
- PIP - Plant-Incorporated Protectants
- SAP - FIFRA Scientific Advisory Panel
- SCLP - Straight Chain Lepidopteran Pheromones

PRIA Category	Description of Category	Number of Completed Decisions	Average Exceedance (Days)
R010	New active ingredient, food use	3	633
R020	New active ingredient, food use, reduced risk	2	870
R124	Conditional ruling on pre-application study waivers; applicant-initiated	9	-14
R170	New use, additional food use	61	378
R175	New use, additional food uses covered within a crop grouping/conversion	22	359
R180	New use, additional food use; reduced risk	4	1086
R190	New use, additional food uses; 6 or more submitted in one application	23	273
R230	New use, additional use; non-food; outdoor	5	310
R250	New use, additional use; non-food; outdoor; EUP; no credit toward new use registration	1	253
R260	New use; non-food; indoor	1	390
R272	Review of study protocol; applicant-initiated; excludes DART, pre-registration conferences, rapid response review, DNT protocol review, protocols needing HSRB review	20	7

PRIA Category	Description of Category	Number of Completed Decisions	Average Exceedance (Days)
R273	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	3	183
R274	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 nonfood uses	2	286
R275	Rebuttal of agency reviewed protocol; applicant-initiated	1	82
R280	Establish import tolerance; new active ingredient or first food use	1	2
R290	Establish import tolerance; additional food use	13	132
R292	Amend an established tolerance (e.g., decrease or increase) and/or harmonize established tolerances with Codex MRLs; domestic or import; applicant-initiated	1	(-287)
R298	Amend an established tolerance (e.g., decrease or increase); domestic or import; submission of corresponding amended labels (requiring science review)	21	169
R300	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	106	82
R301	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 owne	81	152

PRIA Category	Description of Category	Number of Completed Decisions	Average Exceedance (Days)
R310	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: • product chemistry and/or • acute toxicity and/or • child resistant packaging and/ or • pest(s) requiring efficacy- for up to 3 target pests	67	135
R314	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: • product chemistry and/or • acute toxicity and/or • child resistant packaging and/ or • pest(s) requiring efficacy- for up to 3 target pests	22	130
R317	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: • product chemistry and/or • acute toxicity and/or • child resistant packaging and/ or • pest(s) requiring efficacy- for greater than 7 target pests	1	304
R318	New end use product containing four or more registered active ingredients never before registered as this combination in a formulated product; new	14	208

PRIA Category	Description of Category	Number of Completed Decisions	Average Exceedance (Days)
	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: • product chemistry and/or • acute toxicity and/or • child resistant packaging and/ or • pest(s) requiring efficacy- for up to 3 target pests		
R320	New product; new physical form; requires data review in science divisions	21	129
R333	New product with unregistered source of a.i..; cite-all or selective data citation where applicant owns all required data	37	141
R334	New product with unregistered source of a.i.; selective data citation	92	117
R340	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	72	104
R341	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	1	107
R350	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)	41	229
R351	Amendment adding a new unregistered source of active ingredient.	118	10
R352	Amendment adding already approved uses; selective method of support; does not apply if the applicant owns all cited data	35	26
R370	Cancer reassessment; applicant-initiated	1	380
R371	Amendment to Experimental Use Permit; (does not include extending a permit's time period)	1	50
A410	New Active Ingredient Non-food use	4	378
A431	New Active Ingredient, Non-food use; low-risk	1	477
A450	New use, Direct food use, establish tolerance or tolerance exemption	1	1

PRIA Category	Description of Category	Number of Completed Decisions	Average Exceedance (Days)
A460	Additional food use; establish tolerance exemption	2	(-59)
A470	Label amendment requiring data review; 0 to 10 public health organisms	10	(-26)
A500	New use, non-food	4	167
A521	Review of public health efficacy study protocol within AD; per AD Internal Guidance for the Efficacy Protocol Review Process; applicant-initiated; Tier 1	7	43
A523	Protocol review; other than public health efficacy	1	-4
A530	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 when applicant owns all required data, or applicant submits specific authorization letter for data owner. Category also includes 100% re-package of registered end-use or manufacturing-use product that requires no data submission nor data matrix	27	41
A531	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	24	26
A532	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	11	61
A534	Rebuttal of Agency reviewed protocol, applicant initiated	1	102
A535	Conditional ruling on pre-application study waiver or data bridging argument; applicant-initiated	8	184
A540	New end use product; FIFRA §2(mm) uses only; up to 25 public health organisms	50	96
A541	New end use product; FIFRA §2(mm) uses only; 26–50 public health organisms	7	110
A542	New end use product; FIFRA §2(mm) uses only; ≥ 51 public health organisms.	6	167
A550	New end-use product; uses other than FIFRA §2(mm); non-FQPA product	3	85

PRIA Category	Description of Category	Number of Completed Decisions	Average Exceedance (Days)
A560	New manufacturing use product; registered active ingredient; selective data citation	11	80
A565	New manufacturing-use product; registered active ingredient; unregistered source of active ingredient; submission of new generic data package; registered uses only; requires science review	1	630
A570	Approval of amendment(s) to tolerance and label for previously approved safener	146	45
A571	Science reassessment: refined ecological risk, and/or endangered species; applicant-initiated	1	0
A572	New Product or amendment requiring data review for risk assessment by Science Branch (e.g., changes to Restricted Entry Interval, or Personal Protective Equipment, or use rate	4	100
A573	Label amendment requiring data review; 26–50 public health organisms	1	223
A574	Label amendment requiring data review; ≥ 51 public health organisms	1	0
B590	New active ingredient; food use; petition to establish a tolerance exemption	27	99
B600	New active ingredient; nonfood use	7	121
B611	New active ingredient; Experimental Use Permit application; petition to establish permanent tolerance exemption	2	92
B612	New active ingredient; no change to a permanent tolerance exemption	1	147
B614	Pre-application; Conditional Ruling on rationales for addressing a data requirement in lieu of data; applicant-initiated; applies to one rationale at a time	14	114
B621	Amendment; Experimental Use Permit; no change to an established temporary tolerance or tolerance exemption	1	(-159)
B630	First food use; petition to establish a tolerance exemption	2	325
B643	New Food use; petition to amend an established tolerance exemption	2	57
B644	New use, no change to an established tolerance or tolerance exemption	2	7
B650	New use; nonfood	6	72
B660	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	8	75

PRIA Category	Description of Category	Number of Completed Decisions	Average Exceedance (Days)
	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		
B670	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	26	100
B672	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	15	103
B673	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	8	111
B680	Amendment; registered sources of active ingredient(s); no new use(s); no changes to an established tolerance or tolerance exemption. Requires data submission	5	73
B681	Amendment; unregistered source of active ingredient(s). Requires data submission	11	121
B685	Amendment; add a new biochemical unregistered source of active ingredient or a new microbial production site. Requires submission of	14	72

PRIA Category	Description of Category	Number of Completed Decisions	Average Exceedance (Days)
	analysis of samples data and source/production site specific manufacturing process description		
B690	SCLP; New active ingredient; food or non-food use	1	16
B700	SCLP; Experimental Use Permit application; new active ingredient or new use	2	49
B710	SCLP; New product; registered source of active ingredient(s); identical or substantially similar in composition and use to a registered product; no change in an established tolerance or tolerance exemption. 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	1	(-35)
B720	SCLP; 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	1	1
B721	SCLP; New product; unregistered source of active ingredient	1	91
B730	SCLP; Label amendment requiring data submission	3	9
B771	PIP; 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	2	778
B890	PIP; 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)	1	205

PRIA Category	Description of Category	Number of Completed Decisions	Average Exceedance (Days)
B900	PIP; Application to amend a registration, including actions such as extending an expiration date, modifying an IRM plan, or adding an insect to be controlled	1	(-15)
B903	PIP; Inert ingredient tolerance exemption; e.g., a marker such as NPT II; reviewed in BPPD	1	898
B910	PIP; Biotechnology Notification for small scale field testing of genetically engineered microbes	1	(-9)
I001	New food use inert ingredient	17	419
I002	Amend currently approved inert ingredient tolerance or exemption from tolerance; new data	5	468
I003	Amend currently approved inert ingredient tolerance or exemption from tolerance; no new data	1	745
I004	New non-food use inert ingredient	9	126
I007	Substantially similar non-food use inert ingredients when original inert is compositionally similar with similar use pattern	1	(-6)
I008	New or amended polymer inert ingredient, food use.	8	143
I009	New or amended polymer inert ingredient, non-food use	3	(-63)
I010	Petition to amend a single tolerance exemption descriptor, or single non-food use descriptor, to add ≤ 10 CASRNs; no new data	1	663
I018	Petition to add one approved inert ingredient (CASRN) to the Commodity Inert Ingredient List; no data	14	48
M001	Human Studies protocol requiring HSRB review	2	25
M002	Completed human study requiring HSRB review	3	186
M005	New product, combination of AIs across divisions	2	215
M006	Gold Seal letter request	260	56
M007	Extend exclusive use of data as provided by FIFRA Section 3(c)(1)(F)(ii)	2	61
M009	Grant exclusive use of data for a minor use as provided by FIFRA Section 3(c)(1)(F)(vi)	51	68
M010	Conditional ruling on pre-application, product substantial similarity	2	52
M012	Request for up to 5 letters of certification (Certificate of Establishment) for one actively registered product or one product produced for export (excludes distributor products)	3	