Table II

Number of PRIA Actions Pending as of the End of FY 2023

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 Pending Decisions
R010	New active ingredient, food use	20
R020	New active ingredient, food use, reduced risk	39
R060	New Active Ingredient, Non-food use; outdoor	7
R122	Enriched isomer(s) of registered mixed-isomer active ingredient	11
R124	Conditional ruling on pre-application study waivers; applicant-initiated	9
R140	Additional food use; Indoor; food/food handling	6
R150	First food use	10
R17	New Use, each additional food use	1
R170	New use, additional food use	158
R175	New use, additional food uses covered within a crop grouping/conversion	31
R180	New use, additional food use; reduced risk	6
R190	New use, additional food uses; 6 or more submitted in one application	22
R230	New use, additional use; non-food; outdoor	28
R240	Additional use; non-food; outdoor; reduced risk	10

PRIA Category	Description of Category	Number of Pending Decisions
R260	New use; non-food; indoor	7
R272	Review of study protocol; applicant-initiated; excludes DART, pre-registration conferences, rapid response review, DNT protocol review, protocols needing HSRB review	6
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	7
R281	Establish tolerances for residues in imported commodities; additional new food use; submission of residue chemistry data review conducted by Codex or other competent national regulatory authority	1
R290	Establish import tolerance; additional food use	16
R292	Amend an established tolerance (e.g., decrease or increase) and/or harmonize established tolerances with Codex MRLs; domestic or import; applicant-initiated	6
R295	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	13
R296	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	1
R298	Amend an established tolerance (e.g., decrease or increase); domestic or import; submission of corresponding amended labels (requiring science review)	13
R299	Amend 6 or more established tolerances (e.g., decrease or increase); domestic or import; submission of corresponding amended labels (requiring science review)	6
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,	50

PRIA Category	Description of Category	Number of Pending Decisions
	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.	
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 owner.	100
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	91
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	31

PRIA Category	Description of Category	Number of Pending Decisions
R315	New end-use on-animal product, registered source of active ingredient(s) with submission of data and/or waivers for only: • animal safety and • pest(s) requiring efficacy and/or • product chemistry and/or • acute toxicity and/or • child resistant packaging	6
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
R318	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: • product chemistry and/or • acute toxicity and/or • child resistant packaging and/ or • pest(s) requiring efficacy- for up to 3 target pests	18
R319	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	4

PRIA Category	Description of Category	Number of Pending Decisions
	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 4 to 7 target pests	
R320	New product; new physical form; requires data review in science divisions	22
R321	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: • product chemistry and/or • acute toxicity and/or • child resistant packaging and/or • pest(s) requiring efficacy - for 4 to 7 target pests	1
R333	New product with unregistered source of a.i..; cite-all or selective data citation where applicant owns all required data	86
R334	New product with unregistered source of a.i.; selective data citation	101
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	28
R345	Amending on-animal products previously registered, with the submission of data and/or waivers for only: • animal safety and • pest(s) requiring efficacy and/or • product chemistry and/or	1

PRIA Category	Description of Category	Number of Pending Decisions
	• acute toxicity and/or • child resistant packaging	
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)	54
R351	Amendment adding a new unregistered source of active ingredient.	73
R352	Amendment adding already approved uses; selective method of support; does not apply if the applicant owns all cited data.	15
R363	New product; repack of identical registered manufacturing-use product as an end-use product; same registered uses only, with no additional data	1
R370	Cancer reassessment; applicant-initiated	4
A380	New Active Ingredient; Indirect Food use; establish tolerance or tolerance exemption if required	1
A410	New Active Ingredient Non-food use	6
A431	New Active Ingredient, Non-food use; low-risk	3
A451	Additional Direct food uses; establish tolerances or tolerance exemptions if required; 6 or more submitted in one application	2
A460	Additional food use; establish tolerance exemption	19
A461	New end-use product; FIFRA §2(mm) uses only; 11 to 20 public health organisms	3
A462	New end-use product; FIFRA §2(mm) uses only; 21 to 30 public health organisms	3
A463	New end-use product; FIFRA §2(mm) uses only; 31 to 40 public health organisms	1
A470	Label amendment requiring data review; 0 to 10 public health organisms	43
A471	Label amendment requiring data review; 11 to 20 public health organisms	3
A472	Label amendment requiring data review; 21 to 30 public health organisms	1
A500	New use, non-food	5
A521	Review of public health efficacy study protocol within AD; per AD Internal Guidance for the Efficacy Protocol Review Process; applicant-initiated; Tier 1	3

PRIA Category	Description of Category	Number of Pending Decisions
A522	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 study protocol; applicant-initiated; Tier 2	1
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	21
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	8
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	8
A535	Conditional ruling on pre-application study waiver or data bridging argument; applicant-initiated	5
A540	New end use product; FIFRA §2(mm) uses only; up to 25 public health organisms	32
A541	New end use product; FIFRA §2(mm) uses only; 26–50 public health organisms	6
A550	New end-use product; uses other than FIFRA §2(mm); non-FQPA product	5
A560	New manufacturing use product; registered active ingredient; selective data citation	5
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

PRIA Category	Description of Category	Number of Pending Decisions
A570	Approval of amendment(s) to tolerance and label for previously approved safener	24
A571	Science reassessment: refined ecological risk, and/or endangered species; applicant-initiated	1
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	6
A573	Label amendment requiring data review; 26–50 public health organisms	6
B590	New active ingredient; food use; petition to establish a tolerance exemption	96
B600	New active ingredient; nonfood use	11
B612	New active ingredient; no change to a permanent tolerance exemption	4
B613	New active ingredient; petition to convert a temporary tolerance or a temporary tolerance exemption to a permanent tolerance or tolerance exemption	2
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	3
B617	Pre-application; biochemical classification determination	1
B630	First food use; petition to establish a tolerance exemption	6
B641	Amendment of an established tolerance or tolerance exemption	2
B643	New Food use; petition to amend an established tolerance exemption	2
B644	New use, no change to an established tolerance or tolerance exemption	1
B650	New use; nonfood	1
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 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	6

PRIA Category	Description of Category	Number of Pending Decisions
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	17
B671	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
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	6
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	4

PRIA Category	Description of Category	Number of Pending Decisions
B680	Amendment; registered sources of active ingredient(s); no new use(s); no changes to an established tolerance or tolerance exemption. Requires data submission	8
B681	Amendment; unregistered source of active ingredient(s). Requires data submission	10
B683	Amendment; no change to an established tolerance or tolerance exemption; requires review/update of previous risk assessment(s) without data submission (e.g., labeling changes to Restricted Entry Interval, Personal Protective Equipment, Preharvest Interval)	2
B685	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	4
B721	SCLP; New product; unregistered source of active ingredient	1
B750	PIP; 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 PIP	2
B773	PIP; Application to amend or extend a PIP Experimental Use Permit; with petition to extend a temporary tolerance/tolerance exemption for the active ingredient	1
B780	PIP; Registration application; new PIP; non-food/feed or food/feed without tolerance petition based on an existing permanent tolerance exemption	1
B820	PIP; Registration application; new PIP; with petition to establish or amend a permanent tolerance/tolerance exemption of an active ingredient	2
B880	PIP; Registration application; registered 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)	1
B884	PIP; Registration application; new PIP, seed increase with negotiated acreage cap and time-limited registration; with petition to establish a permanent tolerance/tolerance exemption for the active ingredient	12

PRIA Category	Description of Category	Number of Pending Decisions
B885	PIP; Registration application; registered 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)	1
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	5
B903	PIP; Inert ingredient tolerance exemption; e.g., a marker such as NPT II; reviewed in BPPD	1
B906	PIP; Petition to establish a temporary tolerance/tolerance exemption for one or more active ingredients	1
B909	PIP; tolerance exemption determination; applicant-initiated; request to determine if an existing tolerance exemption applies to a PIP	1
I001	New food use inert ingredient	16
I002	Amend currently approved inert ingredient tolerance or exemption from tolerance; new data	1
I003	Amend currently approved inert ingredient tolerance or exemption from tolerance; no new data	2
I004	New non-food use inert ingredient	8
I007	Substantially similar non-food use inert ingredients when original inert is compositionally similar with similar use pattern	1
I008	New or amended polymer inert ingredient, food use.	2
I009	New or amended polymer inert ingredient, non-food use	2
I016	Approval of amendment(s) to tolerance and label for previously approved safener	1
I017	Petition to add one approved inert ingredient (CASRN) to the Commodity Inert Ingredient List; no data.	2
M001	Human Studies protocol requiring HSRB review	1
M002	Completed human study requiring HSRB review	1
M005	New product, combination of AIs across divisions	2
M006	Gold Seal letter request	112
M007	Extend exclusive use of data as provided by FIFRA Section 3(c)(1)(F)(ii)	3

PRIA Category	Description of Category	Number of Pending Decisions
M009	Grant exclusive use of data for a minor use as provided by FIFRA Section 3(c)(1)(F)(vi)	26
M010	Conditional ruling on pre-application, product substantial similarity	1
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)	4
M014	Pre-application nano-particle determination	1