PRIA 4 Interpretations

TABLE 2. REGISTRATION DIVISION - NEW USES

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
R130	13	First Food Use; Indoor; Food/Food Handling (2) (3)	An application that proposes the first indoor food use. First food use includes a proposed use of any U. S. registered active ingredient for which there is no registered "food use". The use requires the establishment of, or the exemption from the requirement of a tolerance under section 408 of the FFDCA. The application submission must contain a petition to establish tolerances or exemption(s) from tolerance for all food/feed commodities covered by the pending registration application(s). Indoor means that the proposed use is for use inside of manmade structures. All indoor food uses included in any original application or petition for the first food use are covered by the base fee for the application in this category if submitted within the original application. Some examples of indoor food uses include use in a food handling and/or processing establishment, use on food crops in a greenhouse, aquatic uses involving potable water, irrigation, or requiring tolerances for fish, or shellfish, use in home gardens, and uses involving livestock, such as livestock housing, and livestock dips.	21	201,017

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			All of the inerts used in the product must be either approved, pending with the Agency, or a new inert is submitted within the package for the applicable uses. A maximum of five new products are covered by the base fee. After the first five new products, each application for an additional new product or new inert ingredient approval that is submitted within this first food use application package is subject to the registration service fee for a new product or a new inert ingredient approval. All such associated applications that are submitted together will be subject to the first food use decision review time. Until the first food use is approved, any subsequent application for another new use(s) containing the same active ingredient will be charged a first food use service fee and decision review timeframe. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration		

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			application, the applicant will be charged an additional 25% of the full registration service fee for the first food use application. The Agency will provide the applicant with a pre-decisional determination 4 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested first food use registration. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice.		
R140	14	Additional food use;	An application that proposes an additional indoor food use. This category includes a proposed indoor food use of any U. S. registered active ingredient for which there	15	46,906

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		Indoor, Food/Food Handling (3) (4)	currently is a registered food use. The use requires the establishment of the exemption from the requirement of a tolerance under section 408 of the FFDCA. If residues are reasonably foreseeable or likely to occur in food or feed or around food, the application submission must contain a petition to establish tolerances or exemption(s) from tolerance for all food/feed commodities covered by the pending registration application(s). Increases in exposure such as a dosage rate increase or different method of application will result in the application being treated as a new use. Indoor means that the proposed use is for use inside of manmade structures. Some examples of indoor food uses include: use in a food handling and/or processing establishment and use on food crops in a greenhouse. The fee applies to each additional food use requested (i.e. the fee for this category is multiplied by 4 if 4 uses are proposed). If a crop group or subgroup is requested, the fee is based on the number of representative crops in that group or subgroup that are not currently registered. If all of the representative crops have been registered, then requesting the crop group will count as one additional use. All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses.		

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			The Agency will provide the applicant with a pre-decisional determination 4 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested additional food use registration. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice. Amendment applications to add new use(s) to registered product labels are covered by the base fee for this category as long as they are all submitted in the same package. Each application for a new product and/or new inert approval submitted in this package, however, is subject to its own registration service fee. The only exception would be if the new use(s) were to be added only to a new product (no		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			amendments to registered product labels in the application package) in which case the review of the one new product application would be covered by the base fee for the new uses. Any new product or amendment to the proposed labeling, which contains the same new use(s), that is submitted subsequent to the submission of the new use application but prior to its decision review time expiration date, will be deemed a separate new use application subject to a separate fee and new decision review time. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the new use application. Finally, if the new use(s) application include 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.		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
R150	15	First Food Use (2) (3)	An application that proposes the first food use. First food use includes a proposed use for any U. S. registered active ingredient for which there is no registered food use. The use requires the establishment of, or the exemption from the requirement of a tolerance under section 408 of the FFDCA. The application submission must contain a petition to establish tolerances or exemption(s) from tolerance for all food/feed commodities covered by the pending registration application(s). All uses included in any original application or petition for the first food use are covered by the base fee for the application in this category if submitted within the original application. Examples of food uses include: use on foods, for example, corn or apples; aquatic uses involving potable water, irrigation, or requiring tolerances for fish, or shellfish; uses on areas where food may be grown or raised such as pasture, rangeland, home garden, beehive, and uses involving livestock, such as livestock housing, livestock dips, and livestock ear tags. All of the inerts used in the product must be either approved, pending with the Agency, or a new inert is submitted within the package for the applicable uses. A maximum of five new products are covered by the base fee. After the first five	21	332,960

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			new products, each application for an additional new product or new inert ingredient approval that is submitted within this first food use package is subject to the registration service fee for a new product or a new inert ingredient approval. All such associated applications that are submitted together will be subject to the first food use decision review time. Until the first food use is approved, any subsequent application for another new use(s) containing the same active ingredient will be charged a first food use service fee and decision review timeframe. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the first food use application. The Agency will provide the applicant with a pre-decisional determination 4 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested first food use registration. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice.		
R155	16 new	First food use, Experimental Use Permit application; a.i. registered for non-food outdoor use (3)(4)	An Experimental Use Permit (EUP) application for the first food use(s) for any U. S. registered active ingredient that is currently registered for non-food outdoor use(s). The first food use(s) requires the establishment of or the exemption from the requirement of a tolerance under section 408 of the FFDCA. The application submission must contain a petition to establish tolerances or exemption(s) from tolerance for all food/feed commodities covered by the pending registration application. All uses included in any original application or petition for the first food use EUP are covered by the base fee for the application in this category if submitted within the original application. Examples of food uses include: use on foods, for example, corn or apples; aquatic uses involving potable water, irrigation, or	21	277,466

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			requiring tolerances for fish, or shellfish; uses on areas where food may be grown or raised such as pasture, rangeland, home garden, beehive, and uses involving livestock, such as livestock housing, livestock dips, and livestock ear tags. The Agency will not accept a certification for crop destruct once the review clock has started (the "clock" or decision review timeframe starts 21 days after the Agency receives the application and the required fees or approves a fee waiver or fee exemption). A change to a crop destruct application would require the applicant to withdraw their application and submit a new application. All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 4 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested experimental use permit. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time		

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			the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice.		
R160	17	First Food Use; Reduced Risk (2) (3)	An application that proposes the first food use. First food use includes a proposed use for any U. S. registered active ingredient for which there is no registered food use. The use requires the establishment of, or the exemption from the requirement of a tolerance under section 408 of the FFDCA. The application submission must contain a petition to establish tolerances or exemption(s) from tolerance for all food/feed commodities covered by the pending registration application(s). All food uses included in any original application or petition for the first food use are covered by the base fee for the application in this category if submitted within the original application. Examples of food uses include: use on foods, for example, corn or apples; aquatic uses involving potable water, irrigation, or requiring tolerances for fish, or shellfish; uses on areas where food may be grown or raised such as pasture, rangeland, home garden, beehive, and uses involving livestock, such as livestock housing, livestock	16	277,466

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			dips, and livestock ear tags. A "reduced risk" (https://www.epa.gov/pesticide-registration/conventional-reduced-risk-pesticide-program) submission must accompany the application for registration. The Agency's Reduced Risk Committee will evaluate the submission and make the determination, based on criteria and guidance listed in PR Notice 97-3 and in FIFRA 3(10) (B) (- iv), whether the requested use(s) qualify as "reduced risk" when compared to currently registered pesticides for the same use(s). The reduced risk status of any use of a chemical is an initial assessment. Should information warrant, or should the Agency determine at any time that the data base for the chemical is unacceptable or upon a more thorough review found to be insufficient to demonstrate that the use/application is reduced risk, the Agency may reject reduced risk status. In the event that any uses do not qualify as "reduced risk" by decision of the Reduced Risk Committee, the application will not receive the reduced risk decision timeframes. The fee category will be changed to the category R150 and the action will receive the R150 decision review timeframe. All of the inerts used in the product must be either approved, pending with the Agency, or a new inert is submitted within the package for the applicable uses.		

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			A maximum of five new products are covered by the base fee. After the first five new products, each application for an additional new product or new inert ingredient approval that is submitted within this first food use package is subject to the registration service fee for a new product or a new inert ingredient approval. All such associated applications that are submitted together will be subject to the first food use decision review time. Until the first food use is approved, any subsequent application for another new use(s) containing the same active ingredient will be charged a first food use service fee and decision review timeframe. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the first food use application.		

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			The Agency will provide the applicant with a pre-decisional determination 4 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested first food use registration. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice.		
R170	18	Additional Food Use (3) (4)	An application that proposes an additional food use. Additional food use includes a proposed food use for any U. S. registered active ingredient for which there currently is an approved food use. The use requires the establishment of, or the exemption from the requirement of a tolerance under section 408 of the FFDCA. The application submission must contain a petition to establish tolerances or exemption(s) from	15	83,317

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			tolerance for all food/feed commodities covered by the pending registration application(s). A different pattern of use that significantly changes or increases exposure such as a dosage rate increase or different method of application will result in the application being treated as a new use. Examples of food uses include: use on foods, for example, corn or apples; aquatic uses involving potable water, irrigation, or requiring tolerances for fish, or shellfish; uses on areas where food may be grown or raised such as pasture, rangeland, home garden, beehive, and uses involving livestock, such as livestock housing, livestock dips, and livestock ear tags. The fee applies to each additional food use requested up to 5 uses (i.e. the fee for this category is multiplied by 4 if 4 uses are proposed). If six or more additional food uses are requested in the application, fee category R190 applies. If a crop group or subgroup is requested, the fee is based on the number of representative crops in that group or subgroup that are not currently registered. If all of the representative crops have been registered, then requesting the crop group will count as one additional use. Some unusual examples of outdoor uses are livestock uses, (i.e. ear tags), livestock dips, and feed through treatments of livestock. All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses.		

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			The Agency will provide the applicant with a pre-decisional determination 4 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested additional food use registration. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice. Amendment applications to add new use(s) to registered product labels are covered by the base fee for this category as long as they are all submitted in the same package. Each application for a new product submitted in this package and/or new inert approval,		

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			however, is subject to its own registration service fee. The only exception would be if the new use(s) were to be added only to a new product (no amendments to registered product labels in the application package) in which case the review of the one new product application would be covered by the base fee for the new uses. Any new product or amendment to the proposed labeling, which contains the same new use(s), that is submitted subsequent to the submission of the new use application but prior to its decision review time expiration date, will be deemed a separate new use application subject to a separate fee and new decision review time. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the new use application. Finally, if the new use(s) application include non-food (indoor and/or outdoor) and food (outdoor and/or indoor) uses, the appropriate fee is due for each type of new use,		

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			and the longest decision review time applies to all of the new uses requested in the application.		
R175	19	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)	An application that proposes conversion of one or more crop groups (or subgroups) for a currently registered active ingredient resulting from a published federal register definition for a revised crop group (or subgroup). The application may reflect the conversion of multiple crop group revisions under the base fee for this category. A request to add uses through the establishment of a crop group or subgroup tolerance where a crop group or subgroup tolerance does not already exist does not fall into this category. The appropriate category will be one of the food use categories (see R170 interpretation). A request to add uses associated with a crop group or subgroup update but before that new crop group definition has been formally established in the Federal Register does not fall into this category. The application will not contain new data for review in this category. If conversion of a crop group or subgroup requires submission of new data, the action does not belong in this category. The application requires a petition for the establishment of, or the exemption from the requirement of a tolerance under Section 408 of the FFDCA as well as a new or amended product label which incorporates the new crop group or subgroup term.	10	69,431

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			The application submission must contain a petition to establish tolerances or exemption(s) from tolerance for all food/feed commodities covered by the pending registration application(s). All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 2 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested additional food use registration. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the		

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			Agency has 2 business days to review; or (c) withdraw the application without prejudice. Amendment applications to add new use(s) to registered product labels are covered by the base fee for this category as long as they are all submitted in the same package. Each application for a new product submitted in this package and/or new inert approval, however, is subject to its own registration service fee. The only exception would be if the new use(s) were to be added only to a new product (no amendments to registered product labels in the application package) in which case the review of the one new product application would be covered by the base fee for the new uses. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the new use application.		
R180	20	Additional Food Use;	An application that proposes an additional food use. Additional food use includes a proposed food use for any U. S. registered active ingredient for which there currently is	10	69,431

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
		Reduced Risk (3) (4)	a registered food use. The use requires the establishment of, or the exemption from the requirement of a tolerance under section 408 of the FFDCA. If residues are reasonably foreseeable or likely to occur in food or feed or around food, the application submission must contain a petition to establish tolerances or exemption(s) from tolerance for all food/feed commodities covered by the pending registration application(s). A different pattern of use that significantly changes or increases exposure such as a dosage rate increase or different method of application will result in the application being treated as a new use. Examples of food uses include: use on foods, for example, corn or apples; aquatic uses involving potable water, irrigation, or requiring tolerances for fish, or shellfish; uses on areas where food may be grown or raised such as pasture, rangeland, home garden, beehive, and uses involving livestock, such as livestock housing, livestock dips, and livestock ear tags. The fee applies to each additional food use requested up to 5 uses (i.e. the fee for this category is multiplied by 4 if 4 uses are proposed). If six or more additional food uses are requested in the application, fee category R200 applies. If a crop group or subgroup is requested, the fee is based on the number of representative crops in that group or subgroup that are not currently registered. If all of the representative crops have been registered, then requesting the crop group will count as one additional use. Some unusual examples of outdoor uses are livestock uses, (i.e. ear tags), livestock dips, and feed through treatments of livestock.		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			A "reduced risk" (https://www.epa.gov/pesticide-registration/conventional-reduced-risk-pesticide-program) submission must accompany the application for registration. The Agency's Reduced Risk Committee will evaluate the submission and make the determination, based on criteria and guidance listed in PR Notice 97-3 and in FIFRA 3 (c)(10) (B) (-iv), whether the requested use(s) qualify as "reduced risk" when compared to currently registered pesticides for the same use(s). In the event that any uses do not qualify as reduced risk, the application will not receive the reduced risk decision timeframes. The reduced risk status of any use of a chemical is an initial assessment. Should information warrant, or should the Agency determine at any time that the data base for the chemical is unacceptable or upon a more thorough review found to be insufficient to demonstrate that the use/application is reduced risk, the Agency may reject reduced risk status. In the event that any uses do not qualify as "reduced risk" by decision of the Reduced Risk Committee, the application will not receive the reduced risk decision timeframes. The fee category will be changed to the category R170, and the action will receive the R170 decision review timeframe. All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses.		

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			The Agency will provide the applicant with a pre-decisional determination 2 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested additional food use registration. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice. Amendment applications to add new use(s) to registered product labels are covered by the base fee for this category as long as they are all submitted in the same package. Each application for a new product and/or new inert approval submitted in this package, however, is subject to its own registration service fee. The only exception would be if		

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			the new use(s) were to be added only to a new product (no amendments to registered product labels in the application package) in which case the review of the one new product application would be covered by the base fee for the new uses. Any new product or amendment to the proposed labeling, which contains the same new use(s), that is submitted subsequent to the submission of the new use application but prior to its decision review time expiration date, will be deemed a separate new use application subject to a separate fee and new decision review time. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the new use application. Finally, if the new use(s) application include 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.		

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R190	21	Additional Food Uses, 6 or more submitted in one application (3) (4)	An application that proposes additional food uses. Additional food use includes a proposed food use for any U. S. registered active ingredient for which there currently is a registered food use. The use requires the establishment of, or the exemption from the requirement of a tolerance under section 408 of the FFDCA. The application submission must contain a petition to establish tolerances or exemption(s) from tolerance for all food/feed commodities covered by the pending registration application(s). A different pattern of use that significantly changes or increases exposure such as a dosage rate increase or different method of application will result in the application being treated as a new use. Examples of food uses include: use on foods, for example, corn or apples; aquatic uses involving potable water, irrigation, or requiring tolerances for fish, or shellfish; uses on areas where food may be grown or raised such as pasture, rangeland, home garden, beehive, and uses involving livestock, such as livestock housing, livestock dips, and livestock ear tags. The application must propose at least (6) specific additional food or feed crops or 6 or more additional representative commodities for crop subgroups or crop groups. If a crop group or subgroup is requested, the fee is based on the number of representative crops in that group or subgroup that are not currently registered. If all of the representative crops have been registered, then requesting the crop group will count as one additional use.	15	499,895

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			All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 4 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested additional food use registrations. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice.		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			Amendment applications to add new use(s) to registered product labels are covered by the base fee for this category as long as they are all submitted in the same package. Each application for a new product submitted in this package and/or new inert approval, however, is subject to its own registration service fee. The only exception would be if the new use(s) were to be added only to a new product (no amendments to registered product labels in the application package) in which case the review of the one new product application would be covered by the base fee for the new uses. Any new product or amendment to the proposed labeling, which contains the same new use(s), that is submitted subsequent to the submission of the new use application but prior to its decision review time expiration date, will be deemed a separate new use application subject to a separate fee and new decision review time. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the new use application.		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			Finally, if the new use(s) application include 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.		
R200	22	Additional Food Use; 6 or more submitted in one application; Reduced Risk (3) (4)	An application that proposes additional food uses. Additional food use includes a proposed food use for any U. S. registered active ingredient for which there currently is a registered food use. The use requires the establishment of, or the exemption from the requirement of a tolerance under section 408 of the FFDCA. The application submission must contain a petition to establish tolerances or exemption(s) from tolerance for all food/feed commodities covered by the pending registration application(s). A different pattern of use that significantly changes or increases exposure such as a dosage rate increase or different method of application will result in the application being treated as a new use. Examples of food uses include: use on foods, for example, corn or apples; aquatic uses involving potable water, irrigation, or requiring tolerances for fish, or shellfish; uses on areas where food may be grown or raised such as pasture, rangeland, home garden, beehive, and uses involving livestock, such as livestock housing, livestock dips, and livestock ear tags. The application must propose at least (6) specific additional food or feed crops or 6 or more additional	10	416,580

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			representative commodities for crop subgroups or crop groups. If a crop group or subgroup is requested, the fee is based on the number of representative crops in that group or subgroup that are not currently registered. If all of the representative crops have been registered, then requesting the crop group will count as one additional use. A "reduced risk" (https://www.epa.gov/pesticide-registration/conventional-reduced-risk-pesticide-program) submission must accompany the application for registration. The Agency's Reduced Risk Committee will evaluate the submission and make the determination, based on criteria and guidance listed in PR Notice 97-3 and in FIFRA 3 (c)(10)(B)(iv), whether the requested use(s) qualify as "reduced risk" when compared to currently registered pesticides for the same use(s). In the event that any uses do not qualify as reduced risk, the application will not receive the reduced risk decision timeframes. The reduced risk status of any use of a chemical is an initial assessment. Should information warrant, or should the Agency determine at any time that the data base for the chemical is unacceptable or upon a more thorough review found to be insufficient to demonstrate that the use/application is reduced risk, the Agency may reject reduced risk status. In the event that any uses do not qualify as "reduced risk" by decision of the Reduced Risk Committee, the application will not receive the reduced		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			risk decision timeframes. The fee category will be changed to the category R190 and the action will receive the R190 timeframe. All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 2 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested additional food use registrations. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			prejudice. Amendment applications to add new use(s) to registered product labels are covered by the base fee for this category as long as they are all submitted in the same package. Each application for a new product submitted in this package and/or new inert approval, however, is subject to its own registration service fee. The only exception would be if the new use(s) were to be added only to a new product (no amendments to registered product labels in the application package) in which case the review of the one new product application would be covered by the base fee for the new uses. Any new product or amendment to the proposed labeling, which contains the same new use(s), that is submitted subsequent to the submission of the new use application but prior to its decision review time expiration date, will be deemed a separate new use application subject to a separate fee and new decision review time. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			applicant will be charged an additional 25% of the full registration service fee for the new uses application.		
R210	23	Additional food use; Experimental Use Permit application; establish temporary tolerance; no credit toward new use registration (3) (4)	An Experimental Use Permit (EUP) application for a new food use(s) that includes a proposed additional food use for any U. S. registered active ingredient that is currently not registered for the proposed use. The use requires the establishment of or the exemption from the requirement of a tolerance under section 408 of the FFDCA. The application submission must contain a petition to establish tolerances or exemption(s) from tolerance for all food/feed commodities covered by the pending registration application(s). Increases in exposure such as a dosage rate increase or different method of application that will result in a temporary tolerance increase belong to this category. Examples of food uses include: use on foods, for example, corn or apples; aquatic uses involving potable water, irrigation, or requiring tolerances for fish, or shellfish; uses on areas where food may be grown or raised such as pasture, rangeland, home garden, beehive, and uses involving livestock, such as livestock housing, livestock dips, and livestock ear tags. The Agency will not accept a certification for crop destruct once the review clock has started (the "clock" or decision review timeframe starts 21 days after the Agency receives the application and the required fees or approves a fee waiver or fee exemption). A change to a crop destruct application would require the applicant to	12	51,436

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			withdraw their application and start the process application again. All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 2 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested experimental use permit. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice.		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the experimental use permit application.		
R220	24	Additional food use; Experimental Use Permit application; Crop Destruct Basis; no credit toward new use registration (3) (4)	An Experimental Use Permit (EUP) application for a new food use(s) includes a proposed food for any U. S. registered active ingredient that is currently not registered for the proposed use. Food/feed commodities covered by the pending application(s) must have a certification that all food/feed treated under the EUP will be destroyed or fed to experimental animals for testing purposes only. Examples of food uses include: use on foods, for example, corn or apples; aquatic uses involving potable water, irrigation, or requiring tolerances for fish, or shellfish; uses on areas where food may be grown or raised such as pasture, rangeland, home garden, beehive, and uses involving livestock, such as livestock housing, livestock dips, and livestock ear tags. All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 2 weeks prior	6	20,830

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested experimental use permit. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the experimental use permit application.		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
R230	25	Additional use; Non-food; Outdoor (3) (4)	An application that proposes a new non-food use. A non-food use includes a proposed use that is not a food use as described in the food use categories. A different pattern of use that significantly changes or increases exposure such as a dosage rate increase or different method of application will result in the application being treated as a new use. Outdoor use means any use that is not indoor as described in the indoor category. Non-food outdoor uses could include treatment of ornamentals in a shade house, termiticide use around the perimeter of a house and turf uses. All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 4 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested additional use registration. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time	15	33,299

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice. Amendment applications to add new use(s) to registered product labels are covered by the base fee for this category as long as they are all submitted in the same package. Each application for a new product submitted in this package and/or new inert approval, however, is subject to its own registration service fee. The only exception would be if the new use(s) were to be added only to a new product (no amendments to registered product labels in the application package) in which case the review of the one new product application would be covered by the base fee for the new uses. Any new product or amendment to the proposed labeling, which contains the same new use(s), that is submitted subsequent to the submission of the new use application but prior to its decision review time expiration date, will be deemed a separate new use application subject to a separate fee and new decision review time.		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the new use application. Finally, if the new use(s) application include 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.		
R240	26	Additional use; Non-food, Outdoor, Reduced Risk (3) (4)	An application that proposes a new non-food use. A non-food use includes a proposed use that is not a food use as described in the food use categories. A different pattern in a non-food outdoor use that significantly changes or increases exposure such as a dosage rate increase or different method of application will result in the application that belongs in this category. Outdoor use means any use that is not indoor as described in the indoor category. Examples of non-food outdoor uses are treatment of ornamentals in a shade house, termiticide use around the perimeter of a house, and	10	27,749

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			turf uses. A "reduced risk" (https://www.epa.gov/pesticide-registration/conventional-reduced-risk-pesticide-program) submission must accompany the application for registration. The Agency's Reduced Risk Committee will evaluate the submission and make the determination, based on criteria and guidance listed in PR Notice 97-3 and in FIFRA 3(c)(10) (B) (-iv), whether the requested use(s) qualify as "reduced risk" when compared to currently registered pesticides for the same use(s). In the event that any uses do not qualify as reduced risk, the application will not receive the reduced risk decision timeframes. The reduced risk status of any use of a chemical is an initial assessment. Should information warrant, or should the Agency determine at any time that the data base for the chemical is unacceptable or upon a more thorough review found to be insufficient to demonstrate that the use/application is reduced risk, the Agency may reject reduced risk status. In the event that any uses do not qualify as "reduced risk" by decision of the Reduced Risk Committee, the application will not receive the reduced risk decision timeframes. The fee category will be changed to the non-reduced risk category and the action will receive the longer timeframes (e.g. from an R240 New Use, Non- Food Use, "reduced risk" to an R230 New Use, Non-Food Use).		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 2 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested additional use registration. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice.		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			Amendment applications to add new use(s) to registered product labels are covered by the base fee for this category as long as they are all submitted in the same package. Each application for a new product submitted in this package and/or new inert approval, however, is subject to its own registration service fee. The only exception would be if the new use(s) were to be added only to a new product (no amendments to registered product labels in the application package) in which case the review of the one new product application would be covered by the base fee for the new uses. Any new product or amendment to the proposed labeling, which contains the same new use(s), that is submitted subsequent to the submission of the new use application but prior to its decision review time expiration date, will be deemed a separate new use application subject to a separate fee and new decision review time. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the new use application.		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			Finally, if the new use(s) application include 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.		
R250	27	Additional Use; Non-food; Outdoor; Experimental Use Permit Application; no credit toward new Use registration (3) (4)	An Experimental Use Permit (EUP) application that proposes a new non-food use for any U.S. registered active ingredient that is currently not registered for the proposed use. A non-food use includes a proposed use that is not a food use as described in the food use categories. A different pattern in a non-food outdoor use that significantly changes or increases exposure such as a dosage rate increase or different method of application will result in the application that belongs in this category. Outdoor use means any use that is not indoor as described in the indoor category. Fees will not cover any subsequent application for registration of the new use. Non-food outdoor uses could include treatment of ornamentals in a shade house, and turf uses. All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses.	6	20,830

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			An Experimental Use Permit (EUP) application that proposes a new non-food use for any U.S. registered active ingredient that is currently not registered for the proposed use. A non-food use includes a proposed use that is not a food use as described in the food use categories. A different pattern in a non-food outdoor use that significantly changes or increases exposure such as a dosage rate increase or different method of application will result in the application that belongs in this category. Outdoor use means any use that is not indoor as described in the indoor category. Fees will not cover any subsequent application for registration of the new use. Non-food outdoor uses could include treatment of ornamentals in a shade house, and turf uses. All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 2 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested experimental use permit. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the experimental use permit application.		
R251	28	Experimental Use Permit application which requires	An Experimental Use Permit (EUP) application for food use which requires no changes to the existing tolerance(s) and the crop is not destroyed. Any U.S. registered active ingredient that currently has approved tolerance(s) for the proposed use. Due to the extended registration process in certain states, this category provides the ability	8	20,830

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
		no changes to the tolerance(s); non- crop destruct basis (3)	to conduct an EUP without the need for crop destruct or for establishing temporary tolerance(s) while the state registration is under review. This category would allow the conduct of research in States for a new application method on a crop for which tolerance(s) were already federally approved. For example, in order to get a California (CA) EUP, CA requires a Federal EUP to do testing. Testing may be required by CA for an aerial application when only the ground application method is approved in the state. Examples of food uses include: use on foods, for example, corn or apples; aquatic uses involving potable water, irrigation, or requiring tolerances for fish, or shellfish; uses on areas where food may be grown or raised such as pasture, rangeland, home garden, beehive, and uses involving livestock, such as livestock housing, livestock dips, and livestock ear tags. All of the inerts used in the product must be approved for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 2 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested experimental use permit. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice.		
R260	29	New Use, Non-food, Indoor (3) (4)	An application that proposes a new non-food use. A non-food use includes a proposed use that is not a food use as described in the food use categories. A different pattern in a non- food indoor use that significantly changes or increases exposure such as a dosage rate increase or different method of application will result in the application that belongs in this category. The proposed use is for use inside of manmade structures and is not a food use. Some examples of indoor uses are termiticides and indoor residential treatments (i.e. cockroach treatments). Treatment of ornamentals in a shade house is classified as outdoor uses and is not covered in this category. All of the inerts used in the product must be either approved or pending with the	12	16,083

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			Agency for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 2 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested non-food use registration. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice. Amendment applications to add new use(s) to registered product labels are covered by the base fee for this category as long as they are all submitted in the same package. Each application for a new product submitted in this package and/or new inert approval,		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			however, is subject to its own registration service fee. The only exception would be if the new use(s) were to be added only to a new product (no amendments to registered product labels in the application package) in which case the review of the one new product application would be covered by the base fee for the new uses. Any new product or amendment to the proposed labeling, which contains the same new use(s), that is submitted subsequent to the submission of the new use application but prior to its decision review time expiration date, will be deemed a separate new use application subject to a separate fee and new decision review time. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the new use application. Finally, if the new use(s) application include 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.		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
R270	30	New Use, Non-food, Indoor, Reduced Risk (3) (4)	An application that proposes a new non-food use. A non-food use includes a proposed use that is not a food use as described in the food use categories. A different pattern in a non- food indoor use that significantly changes or increases exposure such as a dosage rate increase or different method of application will result in the application that belongs in this category. The proposed use is for use inside of manmade structures and is not a food use. Some examples of indoor uses are termiticides and indoor residential treatments (i.e. cockroach treatments). Treatment of ornamentals in a shade house is classified as outdoor uses and is not covered in this category. A "reduced risk" (https://www.epa.gov/pesticide-registration/conventional-reduced-risk-pesticide-program) submission must accompany the application for registration. The Agency's Reduced Risk Committee will evaluate the submission and make the determination, based on criteria and guidance listed in PR Notice 97-3 and in FIFRA 3 (c (10) (B) (-iv), whether the requested use(s) qualify as "reduced risk" when compared to currently registered pesticides for the same use(s). In the event that any uses do not qualify as reduced risk, the application will not receive the reduced risk decision timeframes. The reduced risk status of any use of a chemical is an initial assessment. Should information warrant, or should the Agency determine at any time that the data	9	13,403

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			base for the chemical is unacceptable or upon a more thorough review found to be insufficient to demonstrate that the use/application is reduced risk, the Agency may reject reduced risk status. In the event that any uses do not qualify as "reduced risk" by decision of the Reduced Risk Committee, the application will not receive the reduced risk decision timeframes. The fee category will be changed to the non-reduced risk category and the action will receive the longer timeframes (e.g. from an R270 New Use, Non-Food Use "reduced risk" to an R260 New Use, Non-Food Use). All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 2 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested non-food use registration. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time		

EPA No.	CR No.	Action	Interpretation	Decision Review Time (Months)	FY'20-FY'21 Registration Service Fee ($)
			the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice. Amendment applications to add new use(s) to registered product labels are covered by the base fee for this category as long as they are all submitted in the same package. Each application for a new product submitted in this package and/or new inert approval, however, is subject to its own registration service fee. The only exception would be if the new use(s) were to be added only to a new product (no amendments to registered product labels in the application package) in which case the review of the one new product application would be covered by the base fee for the new uses. Any new product or amendment to the proposed labeling, which contains the same new use(s), that is submitted subsequent to the submission of the new use application but		

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			prior to its decision review time expiration date, will be deemed a separate new use application subject to a separate fee and new decision review time. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the new use application. Finally, if the new use(s) application include 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.		
R271	31	New use; non-food; indoor; Experimental Use Permit application; no credit toward	An Experimental Use Permit (EUP) application for a new non-food use(s) includes a proposed non-food use for any U. S. registered active ingredient that is currently not registered for the proposed use. A non-food use includes a proposed use that is not a food use as described in the food use categories. Increases in exposure such as a dosage rate increase or different method of application will result in the application being treated as a new use. The proposed use is for use inside of manmade structures	6	10,212

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		new use registration (3) (4)	and is not a food use. Some examples of indoor uses are termiticide structural protection and indoor residential treatments (i.e. cockroach treatments). Treatment of ornamentals in a shade house is classified as outdoor uses and is not covered in this category. All of the inerts used in the product must be either approved or pending with the Agency for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 2 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested experimental use permit. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit		

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			a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the experimental use permit application.		
R273	32	Additional use; seed treatment; limited uptake into Raw Agricultural Commodities; includes crops with established tolerances (e.g., for soil or foliar	An application that proposes an additional seed treatment use only for any U.S. registered active ingredient for food use or non-food use seed treatment that is not expected to result in residues above already set tolerance levels in raw agricultural commodities. In order for a seed treatment to be considered in this category when proposed for seed treatment use on a food crop, data from a radiotracer study must be available showing limited uptake of residues (radioactivity) from treated seed into the aerial portion of the growing crop. Guidance is available at (https://www.epa.gov/test-guidelines-pesticides-and-toxic-substances/series-860-residue-chemistry-test-guidelines)	12	52,968

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		application); includes food and/or non-food uses (3) (4)	If residues occur in the aerial portion of the plant that require the establishment of a new tolerance or modification of an existing tolerance, or if there is no data available to make this determination, the seed treatment falls into a different category. Examples of food uses are corn, soybean, and wheat. If a seed treatment use is proposed on ornamental seed or other non-food use seed treatments, then the application would be in this category because it is known, without consideration of any data, that a tolerance is not required. The fee applies to each seed treatment use requested up to 5 uses (i.e. the fee for this category is multiplied by 4 if 4 seed uses are proposed). If a crop group or subgroup is requested, the fee is based on the number of representative crops in that group or subgroup that are not currently registered. If all of the representative crops have been registered, then requesting the crop group will count as one additional use. If a numerical tolerance needs to be established, the application does not belong in this category. If six or more seed treatment uses are being proposed, this is not the correct category (see R274). All of the inerts used in the product must be approved for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 2 weeks		

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			prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested additional use registration. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice. Amendment applications to add new use(s) to registered product labels are covered by the base fee for this category as long as they are all submitted in the same package. Each application for a new product submitted in this package and/or new inert approval, however, is subject to its own registration service fee. The only exception would be if the new use(s) were to be added only to a new product (no amendments to registered product labels in the application package) in which case the review of the one new		

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			product application would be covered by the base fee for the new uses. Any new product or amendment to the proposed labeling, which contains the same new use(s), that is submitted subsequent to the submission of the new use application but prior to its decision review time expiration date, will be deemed a separate new use application subject to a separate fee and new decision review time. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the new use application. Finally, if the new use(s) application include 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.		
R274	33	Additional uses; seed treatment	An application that proposes additional seed treatment uses only for any U.S. registered active ingredient that is not expected to result in residues above already set tolerance	12	317,797

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		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)	levels. The application must propose at least (6) specific seed treatment uses or 6 or more representative seeds for crop subgroups or crop groups. In order for a seed treatment to be considered in this category, data from a radiotracer study must be available showing limited uptake of residues (radioactivity) from treated seed into the aerial portion of the growing crop. Guidance is available at https://www.epa.gov/test-guidelines-pesticides-and-toxic-substances/series-860-residue-chemistry-test-guidelines If residues occur in the aerial portion of the plant that require the establishment of a new tolerance or modification of an existing tolerance, or if there is no data available to make this determination, the seed treatments fall into a different category. Examples of food uses are corn, soybean, and wheat. If a seed treatment use is proposed on ornamental seed or other non-food use seed treatment uses, then the application would be in this category because it is known, without consideration of any data, that a tolerance is not required. If a crop group or subgroup is requested, the fee is based on the number of representative crops in that group or subgroup that are not currently registered. If all of the representative crops have been registered, then requesting the crop group will count as one additional use. If a numerical tolerance needs to be established, the application does not belong in this category.		

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			All of the inerts used in the product must be approved for the applicable uses. The Agency will provide the applicant with a pre-decisional determination 2 weeks prior to the PRIA decision review time due date which specifies any label changes that have to be made in order to grant the requested additional use registrations. If the label issues cannot be resolved prior to the PRIA decision review time due date and if a PRIA due date time extension has not been agreed upon, then the Agency will issue to the applicant its regulatory decision with the specific label changes and supporting documentation on or just before the PRIA decision review time due date. At that time the applicant must either (a) agree to all of the label changes and submit a revised label that incorporates all of these label changes; or (b) does not agree with one or more of the label changes and request up to 30 days to reach agreement with the Agency and submit a revised label that incorporates all of the agreed upon label changes, which the Agency has 2 business days to review; or (c) withdraw the application without prejudice. Amendment applications to add new use(s) to registered product labels are covered by		

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			the base fee for this category as long as they are all submitted in the same package. Each application for a new product submitted in this package and/or new inert approval, however, is subject to its own registration service fee. The only exception would be if the new use(s) were to be added only to a new product (no amendments to registered product labels in the application package) in which case the review of the one new product application would be covered by the base fee for the new uses. Any new product or amendment to the proposed labeling, which contains the same new use(s), that is submitted subsequent to the submission of the new use application but prior to its decision review time expiration date, will be deemed a separate new use application subject to a separate fee and new decision review time. If the applicant on his own initiative submits any additional information that was neither requested nor required by the Agency, after completion of the technical deficiency screening, and which does not itself constitute a covered registration application, the applicant will be charged an additional 25% of the full registration service fee for the new uses application. Finally, if the new use(s) application include 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		

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			longest decision review time applies to all of the new uses requested in the application.		