

EU PESTICIDE RENEWAL MONITOR

August 31, 2020¹

In the European Union, all active ingredients undergo periodic reevaluation, as laid out in Regulation (EC) No. 1107/2009². The reevaluation process typically follows the scheme indicated in the figure below.

This document provides an overview of active ingredients that are currently undergoing or are scheduled for the periodic active ingredient renewal in the European Union and it is prepared based on publicly available information. It lists active ingredients that expired or are expected to expire, as of **August 31, 2020**. Please note that the renewal process relates to the approval of active ingredients used in the EU market. It is lengthy and unpredictable. Only final, published Regulations are definitive. Pesticide Renewal is a separate process to the EU MRL and Import Tolerance setting, which is governed by specific legislation. MRL change or removal may happen after non-renewal has been finalized.

For additional information on the registration status and expiry dates for active ingredients that are not listed in this document, please refer to the [EU pesticide database](#)³.

For information on specific active ingredients, please consult the [EFSA website](#)⁴ (then click on the “Pesticide Dossier” tab) or contact the companies that are supporting the active ingredient through the renewal process by contacting croplife@croplife.org.

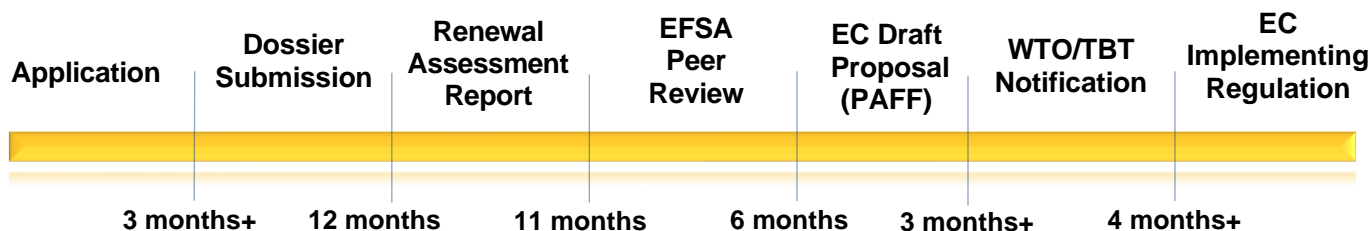


Figure 1: Schematic overview of the EU reevaluation process for non-renewals or restricted renewals. Timelines are indicative as outlined in Regulation (EC) No. 1107/2009. Steps can take longer than indicated.

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¹ This EU Pesticide Renewal Monitor issue reflects substances listed as proposed non-renewal or restricted renewal on the PAFF agenda up to the July 16-17, 2020 meeting.

² <https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32009R1107&from=EN>

³ <https://ec.europa.eu/food/plant/pesticides/eu-pesticides-database/public/?event=homepage&language=EN>

⁴ <http://registerofquestions.efsa.europa.eu/roqFrontend/ListOfQuestionsNoLogin>

APPLICATION FOR RENEWAL - EXPIRED (September 2019 - August 2020)

Chemical companies must submit an application for renewal of approval of their active ingredient to EU authorities. If they do not, the active ingredient will automatically expire in the EU on a set date. Active ingredients below **have already expired** due to **no submission of application for renewal** or **withdrawal of application for renewal**. This list includes substances that have expired since September 2019.

Quizalofop-P	30/11/2019	Phlebiopsis gigantea	
Teflubenzuron	30/11/2019	multiple strains	30/04/2020
Difenacoum	30/12/2019	Epoconazole	30/04/2020
Chlorsulfuron	31/12/2019	Triflumizole	30/06/2020
Cyromazine	31/12/2019	Spirodiclofen	31/07/2020
Lufenuron	31/12/2019		

APPLICATION FOR RENEWAL - EXPECTED TO EXPIRE (Up to August 2021)

Applications for **renewal of approval were not submitted** or **applications have been withdrawn** for the active ingredients below. The expiration date is outlined. Please note that the expiration of active ingredients in the EU is not notified to the World Trade Organization (WTO).

FEN 560	31/10/2020	Carboxin	31/05/2021
Triflumuron	31/03/2021	Diethofencarb	31/05/2021
Fenbuconazole	30/04/2021	Etridiazole	31/05/2021
Metosulam	30/04/2021	Fenoxycarb	31/05/2021
Trichoderma artroviride		Myclobutanil	31/05/2021
strain IMI 206040	30/04/2021	Oryzalin	31/05/2021
Bromadiolone	31/05/2021	Profoxydim	31/07/2021
Carbetamide	31/05/2021		

UP NEXT FOR REVIEW (Up to January 2021)

Under the EU pesticide review program, the active ingredients listed below have **upcoming deadlines** for the **submission of the application for renewal**. The list shows all active ingredients that are scheduled to go through the periodic renewal process. This does not imply that these active ingredients will have issues with renewal in the EU.

Bacillus firmus I01582	30/09/2020	Eugenol	30/11/2020
Bixafen	30/09/2020	Geraniol	30/11/2020
Candida oleophila strain O	30/09/2020	Aureobasidium pullulans (strains DSM 14940 and DSM 14941)	31/01/2021
Halosulfuron methyl	30/09/2020	Disodium phosphonate	31/01/2021
Maltodextrin	30/09/2020	Fluopyram	31/01/2021
Paecilomyces fumosoroseus strain Fe9901	30/09/2020	Penflufen	31/01/2021
Potassium phosphonates	30/09/2020	Pseudomonas sp. DSMZ 13134	31/01/2021
Spiromesifen	30/09/2020	Pyriofenone	31/01/2021
Thymol	30/11/2020	Sedaxane	31/01/2021

STANDING COMMITTEE ON PLANTS, ANIMALS, FOOD AND FEED (PAFF)

Active ingredients below have been under discussion for **non-renewal** or **restricted renewal** and are now **under consideration** by the Standing Committee on Plants, Animals, Food and Feed (PAFF) as available in the May 18-19 summary report and July 16-17, 2020 meeting agenda. A final decision may have not yet been made.

- **Azadirachtin**
- **Benalaxyl**
- **Benfluralin**
- **Bifenazate**
- **Bromoxynil**
- **Carbon dioxide**
- **Ethametsulfuron-methyl**
- **Etoxazole**
- **Fenamiphos**
- **Fenpyrazamine**
- **Indoxacarb**
- **Mancozeb**
- **Thiophanate-methyl**

WTO NOTIFICATION (September 2019 – August 2020)

Active ingredients below have been notified to the **WTO as proposed for non-renewal or restricted renewal**. Active ingredients that are renewed do not need to be notified to the WTO. These WTO Technical Barrier to Trade (TBT) notifications advise that the active ingredient is subject to a proposed non-renewal for use in the EU. This is not necessarily indicative of the final EU decision. Further Committee work is done by the Commission and EU Member States after notification, which can change the proposal. Definitive non-renewal regulation may take a significant period of time after the WTO consultation has completed. These notifications do not advise what will happen with EU MRLs. This information is provided in notifications via WTO Sanitary and Phytosanitary (SPS), typically some years later. For full explanation on the justification for proposed restricted or non-renewal of approval, please refer to the draft implementing regulation. Notification date is outlined in parentheses ().

Fenamiphos: proposed non-renewal based on incomplete data and acute risk for consumers was identified for all the representative uses concerning fruiting vegetables. (04/03/2020)

Bromoxynil: proposed non-renewal based on suggested classification as toxic for reproduction category 1B. (17/04/2020)

Mancozeb: proposed non-renewal based on classification as toxic for reproduction category 1B. (17/04/2020)

Benfluralin: proposed non-renewal based on long-term risk to birds and mammals including the risk from secondary poisoning of earthworm eating birds and mammals, as well as the genotoxic potential of an impurity could not be excluded. (06/05/2020)

Benalaxyl: proposed non-renewal based on potential groundwater contamination, long term risk to birds and earthworm-eating birds from secondary poisoning as well as lack of data to assess the endocrine disrupting potential. (14/05/2020)

Proposed restricted renewal

Etoxazole: proposed renewal as a candidate for substitution and restricted to use on ornamental plants in permanent greenhouses. (17/08/2020)

COMMISSION IMPLEMENTING REGULATION (September 2019 - August 2020)

The Commission has published the final decision on non-renewal or restricted renewal in the EU for the active ingredients below. For full explanation on the justification for restricted or non-renewal of approval, please refer to the published implementing regulation. Publication date is outlined in parentheses ().

Methiocarb: non-renewal based on an unacceptable risk to workers, even when considering the use of personal protective equipment and a high risk to birds, mammals and earthworms. (30/09/2019)

Chlorpyrifos: non-renewal based on genotoxic potential, developmental neurotoxicity, and classification of the substance as toxic for reproduction, category 1B. (13/01/2020)

Chlorpyrifos-methyl: non-renewal based on genotoxic potential, developmental neurotoxicity, and classification of the substance as toxic for reproduction, category 1B. (13/01/2020)

COMMISSION IMPLEMENTING REGULATION (continued)

Thiacloprid: non-renewal based on a critical concern in relation to the contamination of groundwater by metabolites containing carcinogenic properties. There is also an undetermined risk to aquatic organisms, bees, and non-target plants, as well as concerns on the impact on reproductive toxicity. (14/01/2020)

Beta-cyfluthrin: non-renewal based on unacceptable risk to workers, high risk to residents, to non-target arthropods and to aquatic organisms. (30/06/2020)

Restricted renewal

Alpha-Cypermethrin: restricted renewal as a candidate for substitution until October 31, 2026. (09/10/2019)

Metalaxyl-M: restricted renewal limits seeds treated with metalaxyl-M to only be grown in greenhouses. (05/05/2020)

MRL CHANGES (September 2019 - August 2020)

As a **result of non-renewal** or **expiration** of approval, restrictive MRLs have either been proposed (WTO notification) or implemented (Commission Regulation) for the substances below. Notifications of MRL changes will appear as WTO SPS notifications. Please note that not all MRL changes are included here as MRLs may change even if the active ingredient is reaproved.

WTO SPS Notification

- **Azinphos-methyl:** WTO notification G/SPS/N/EU/371. (28/02/2020)
- **Flufenoxuron:** WTO notification G/SPS/N/EU/371. (28/02/2020)
- **Oxadiazon:** WTO notification G/SPS/N/EU/371. (28/02/2020)
- **Phosalone:** WTO notification G/SPS/N/EU/371. (28/02/2020)
- **Tall oil pitch:** WTO notification G/SPS/N/EU/371. (28/02/2020)
- **Tall oil crude:** WTO notification G/SPS/N/EU/371. (28/02/2020)
- **Chlorothalonil:** WTO notification G/SPS/N/EU/394. (15/07/2020)
- **Chlorpropham:** WTO notification G/SPS/N/EU/394. (15/07/2020)
- **Dimethoate:** WTO notification G/SPS/N/EU/394. (15/07/2020)
- **Ethoprophos:** WTO notification G/SPS/N/EU/394. (15/07/2020)
- **Fenamidone:** WTO notification G/SPS/N/EU/394. (15/07/2020)
- **Methiocarb:** WTO notification G/SPS/N/EU/394. (15/07/2020)
- **Propiconazole:** WTO notification G/SPS/N/EU/394. (15/07/2020)
- **Pymetrozine:** WTO notification G/SPS/N/EU/394. (15/07/2020)

MRL CHANGES (*continued*)Implementing Regulation

- **Amitrole:** Commission Regulation 2019/1792 on 17/10/2019. *Effective date: 18/05/2020*
- **Flupyrulfuron-methyl:** Commission Regulation 2019/1792 on 17/10/2019. *Effective date: 18/05/2020*
- **Isoproturon:** Commission Regulation 2019/1792 on 17/10/2019. *Effective date: 18/05/2020*
- **Triasulfuron:** Commission Regulation 2019/1792 on 17/10/2019. *Effective date: 18/05/2020*
- **Fipronil:** Commission Regulation 2019/1792 on 17/10/2019. *Effective date: 18/05/2020*
- **Imazosulfuron:** Commission Regulation 2019/1792 on 17/10/2019. *Effective date: 18/05/2020*
- **Orthosulfamuron:** Commission Regulation 2019/1792 on 17/10/2019. *Effective date: 18/05/2020*
- **Chlorpyrifos:** Commission Regulation 2020/1085 on 24/07/2020. *Effective date: 13/11/2020*
- **Chlorpyrifos-methyl:** Commission Regulation 2020/1085 on 24/07/2020. *Effective date: 13/11/2020*

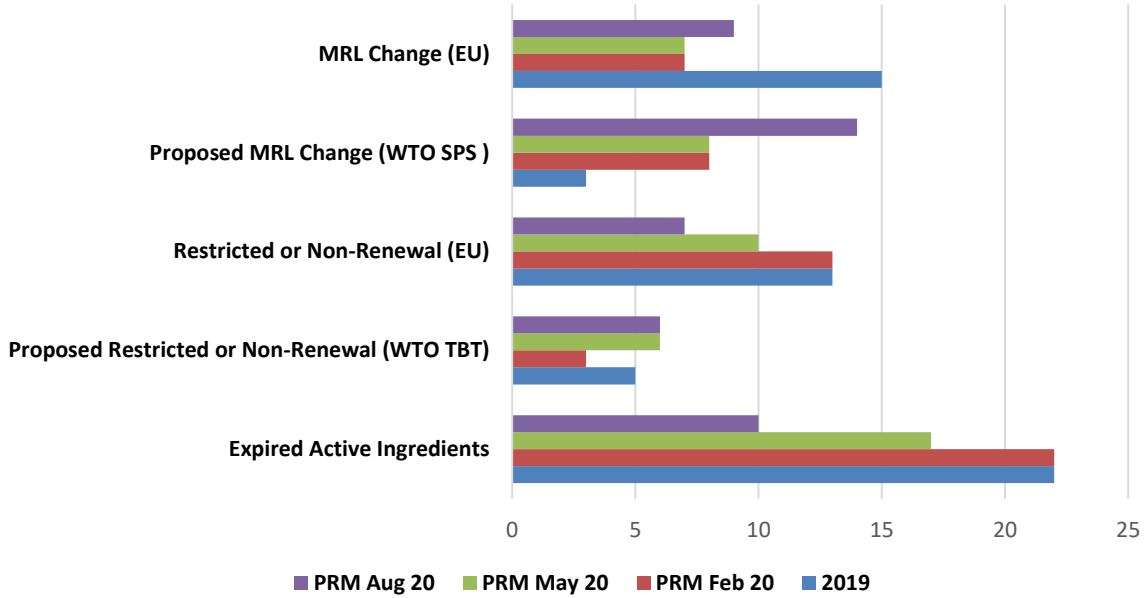
Table I. EU Pesticide Renewal Monitor in Numbers

Time scope / Review step	2019 (January-December)	PRM February 2020 (February 2019- February 2020)	PRM May 2020 (June 2019- May 2020)	PRM August 2020 (September 2019- August 2020)
Expired Active Ingredients	22	22	17	10
	Fuberidazole, Bacillus thuringiensis subsp. Tenebrionis strain NB 176 (TM 14 1), Fenpropimorph, Tralkoxydim, Trichoderma polysporum strain IMI 206039, Ammonium acetate, Limestone, Methomyl, Pepper, Putrescine, Sea-algae extract, Sodium aluminium silicate, Sodium hypochlorite, Triadimenol, Trimethylamine hydrochloride, Quinalofop-P, Teflubenzuron, Difenacoum, Chlorsulfuron, Cyromazine, Lufenuron, Propanil	Propanil, Fuberidazole, Fenpropimorph, Tralkoxydim, Trichoderma polysporum strain IMI 206039, Bacillus thuringiensis subsp. Tenebrionis strain NB 176 (TM 14 1), Sea-algae extract, Ammonium acetate, Limestone, Methomyl, Pepper, Putrescine, Sodium hypochlorite, Triadimenol, Trimethylamine hydrochloride, Sodium aluminium silicate, Quinalofop-P, Teflubenzuron, Difenacoum, Chlorsulfuron, Cyromazine, Lufenuron	Sea-algae extract, Ammonium acetate, Limestone, Methomyl, Pepper, Putrescine, Sodium hypochlorite, Triadimenol, Trimethylamine hydrochloride, Sodium aluminium silicate, Quinalofop-P, Teflubenzuron, Difenacoum, Chlorsulfuron, Cyromazine, Lufenuron, Phlebobiosis gigantea multiple strains	Quinalofop-P, Teflubenzuron, Difenacoum, Chlorsulfuron, Cyromazine, Lufenuron, Phlebobiosis gigantea multiple strains, Epoxiconazole, Triflumizole, Spirodiclofen
Proposed Restricted or Non-Renewal of Approval (WTO TBT)	5	3	6	6
	Thiacloprid, Metalaxyl-M, Chlorpyrifos, Chlorpyrifos-methyl, Thiophanate-methyl	Thiophanate-methyl, Beta-cyfluthrin, Metalaxyl-M	Beta-cyfluthrin, Fenamiphos, Bromoxynil, Mancozeb, Benfluralin, Benalaxyl	Fenamiphos, Bromoxynil, Mancozeb, Benfluralin, Benalaxyl, Etoxazole

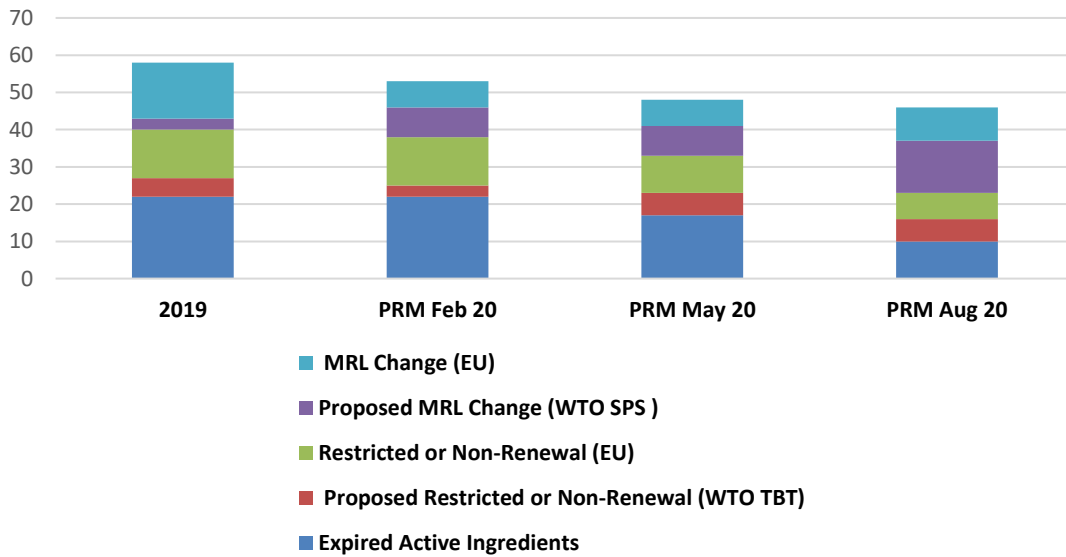
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Time scope / Review step	2019 (January-December)	PRM February 2020 (February 2019- February 2020)	PRM May 2020 (June 2019- May 2020)	PRM August 2020 (September 2019- August 2020)
	13	13	10	7
Restricted or Non-Renewal of Approval (Commission Regulation)	Ethoprophos, Chlorothalonil, Chlorpropham, Dimethoate, Desmedipham, Methiocarb, Chlorpyrifos, Chlorpyrifos-methyl, Thiachlopid, Methoxyfenozide, Cyflumetofen, Tolclofos-methyl, Alpha-Cypermethrin	Ethoprophos, Chlorothalonil, Chlorpropham, Dimethoate, Desmedipham, Methiocarb, Chlorpyrifos, Chlorpyrifos-methyl, Thiachlopid, Methoxyfenozide, Cyflumetofen, Tolclofos-methyl, Alpha-Cypermethrin	Chlorpropham, Dimethoate, Desmedipham, Methiocarb, Chlorpyrifos, Chlorpyrifos-methyl, Thiachlopid, Tolclofos-methyl, Alpha-Cypermethrin, Metalaxyl-M	Methiocarb, Chlorpyrifos, Chlorpyrifos-methyl, Thiachlopid, Beta-cyfluthrin, Alpha-Cypermethrin, Metalaxyl-M
	3	8	8	14
Active Ingredients with Proposed MRL Changes due to Non-Approval (WTO SPS)	Flufenoxuron, Chlorpyrifos, Chlorpyrifos-methyl	Azinphos-methyl, Flufenoxuron, Oxadiazon, Phosalone, Tall oil pitch, Tall oil crude, Chlorpyrifos, Chlorpyrifos-methyl	Azinphos-methyl, Flufenoxuron, Oxadiazon, Phosalone, Tall oil pitch, Tall oil crude, Chlorpyrifos, Chlorpyrifos-methyl	Azinphos-methyl, Flufenoxuron, Oxadiazon, Phosalone, Tall oil pitch, Tall oil crude, Chlorothalonil, Chlorpropham, Dimethoate, Ethoprophos, Fenamidone, Methiocarb, Propiconazole, Pymetrozine
	15	7	7	9
Active Ingredients with MRL Changes due to Non-Approval (Commission Regulation)	Iprodione, Linuron Buprofezin, Diflubenzuron, Picoxystrobin, Tepraloxydim, Ethoxysulfuron, Ioxynil, Amitrole, Flupyrsulfuron-methyl, Isoproturon, Triasulfuron, Fipronil, Imazosulfuron, Orthosulfamuron	Amitrole, Flupyrsulfuron-methyl, Isoproturon, Triasulfuron, Fipronil, Imazosulfuron, Orthosulfamuron	Amitrole, Flupyrsulfuron-methyl, Isoproturon, Triasulfuron, Fipronil, Imazosulfuron, Orthosulfamuron	Amitrole, Flupyrsulfuron-methyl, Isoproturon, Triasulfuron, Fipronil, Imazosulfuron, Orthosulfamuron, Chlorpyrifos, Chlorpyrifos-methyl
Total Number of Active Ingredients	58	53	48	46

Active Ingredients per Review Stage



Total Active Ingredients per EU Pesticide Renewal Monitor



EU PESTICIDE RENEWAL MONITOR Q&A

What is the EU Pesticide Renewal Monitor report?

- Provides an overview on the active ingredients that are scheduled for the periodic review or are currently undergoing the periodic review. The periodic review is a normal and regular procedure for all pesticides approved in the EU, according to Regulation 1107/2009 that regulates market access for pesticides in the EU.
- Active ingredients that have not been renewed, have a restricted renewal or are expired (no renewal dossier submitted) are also listed.

Why was this report created?

- CropLife International commissioned the Pesticide Renewal Monitor report to support governments, farmers and the agri-food value chain to better understand which active ingredients are currently undergoing the pesticide renewal process in the European Union.
- The EU pesticide renewal procedure can be complex, challenging to track and the status of products in the review process may be difficult to interpret.
- Information on review status of active ingredients is publicly available but spread over multiple websites and reports. Most information is not available in one place.
- The EU needs to notify WTO member states when an active ingredient approval is changed or not renewed (TBT notification). WTO notifications are only made late in the EU renewal process, consequently stakeholders are informed late in the process. A subsequent decision to lower or delete MRLs (potentially including ITs and MRLs corresponding to Codex), which is done through a SPS notification, may be too late for stakeholders to adapt their supply and/or to engage in the discussion.
- The US Department of Agriculture (USDA) has commissioned a similar report for US relevant active ingredients which has been valuable for US stakeholders.

What is the difference between the USDA Early Alert and the CropLife International EU Pesticide Monitor?

- The USDA Early Alert covers active ingredients undergoing the EU renewal that have MRLs established in the United States.
- The CropLife International report covers all active ingredients undergoing the renewal in the EU. i.e. the CropLife International report covers more substances than the USDA report.

Why was the report developed?

- To inform interested stakeholders on the process and timing of active ingredient renewal in the EU, to create awareness for non-EU stakeholders
- In case of restrictive or non-renewal: alert stakeholders that/when MRLs may be lowered to the Limit of Quantification (LOQ).
- Provide timely information so stakeholders could engage in support of adequate transition periods or maintenance of MRLs that are important for trade with the EU.

Who is responsible for submitting a dossier for active ingredient renewal in the European Union?

- All stakeholders are invited to provide information relevant for the renewal of the active ingredients.

- Data requirements for active ingredient approval are in place (Regulation EU 283/2013).
- Normally, industry (individual companies, several companies jointly or task forces) generates the data and compiles and submits the dossier. A list of dossier submitters is available on the EFSA website: <http://registerofquestions.efsa.europa.eu/roqFrontend/ListOfQuestionsNoLogin> (then click on the “Pesticide Dossier” tab)

What is the relationship between active ingredient renewal and MRL / IT?

- When an active ingredient is not renewed in the EU or if the conditions of use were amended, MRLs may be deleted or lowered.
- Deletion of MRLs following the revocation of pesticide authorizations in the EU does not apply to MRLs based on Codex MRLs and Import Tolerances established to facilitate global trade, provided that they are deemed acceptable with regard to consumer safety.

Whom can I contact for questions on a specific active ingredient?

- Companies that submitted a dossier for EU renewal can be retrieved through the EFSA website: <http://registerofquestions.efsa.europa.eu/roqFrontend/ListOfQuestionsNoLogin> (then click on the “Pesticide Dossier” tab)
- Individual companies will have the relevant information on the active ingredient, products they sell to farmers and the range of crops that the product is approved for use.
- In case of issues in identifying the relevant authorization holder for a specific active ingredient, please feel free to contact CropLife International.

Is this a complete list of substances registered in the EU?

- No. Only active ingredients are listed that will undergo or currently undergo the renewal process. Also, substances that expired, were not renewed or had a restricted renewal are mentioned.
- For more information and for information on the approval status and expiry dates for active ingredient that are not listed in this document, we refer to the EU pesticide database: <https://ec.europa.eu/food/plant/pesticides/eu-pesticides-database/public/?event=homepage&language=EN>

When will this report be updated?

- This report will be updated on a quarterly basis or at the discretion of CropLife International.