EUROPEAN COMMISSION
HEALTH & FOOD SAFETY DIRECTORATE-GENERAL

Safety of the food chain
Pesticides and Biocides

SANCO/2012/11284 –rev. 22

December 2018

DRAFT WORKING DOCUMENT

AIR III RENEWAL PROGRAMME

SUMMARY

This document provides an overview of indicative submission dates for supplementary dossiers for the renewal of substances expiring between 1 January 2013 and 31 December 2018. It is foreseen that the document will be updated regularly. **It is recommended to refer to the dates provided in the relevant acts adopted as published in the Official Journal (see http://eur-lex.europa.eu/homepage.html).**

Commission Implementing Regulation (EU) No 844/2012 *setting out the provisions necessary for the implementation of the renewal procedure for active substances, as provided for in Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market* has been published on 19 September 2012.

Commission Implementing Regulation (EU) No 686/2012 *allocating to Member States, for the purposes of the renewal procedure, the evaluation of the active substances whose approval expires by 31 December 2018 at the latest* has been published on 27 July 2012.

Regarding the substances covered by **Group 1** a Regulation to set new expiry dates has been published on 15 September 2012 (Commission Regulation (EU) No 823/2012). The submission dates for supplementary dossiers for this Group (batch 1, 2 and 3) are now fixed by Regulations (EU) No 823/2012 and Regulation (EU) No 844/2012.

Applications for batch 1 of Group 1 had to be submitted by 31 July 2013. For the substances for which no application has been received the expiry date has been set at the earliest date possible after the original date of expiry (Regulation (EU) No 186/2014).

Applications for batch 2 of Group 1 had to be submitted by 31 October 2013. For the substance for which no application has been received the expiry date will be set at the earliest date possible after the original date of expiry (Regulation (EU) No 460/2014).

All applications for batch 3 of Group 1 have been received before the deadline of 31 January 2014.

Regarding the substances covered by **Group 2** a Regulation to set new expiry dates has been published on 14 December 2012 (Commission Regulation (EU) No 1197/2012). As a consequence now also the submission dates for supplementary dossiers for this Group (batch 4 and 5) are fixed.

Regarding the substances covered by **Group 3** no Regulation will be prepared to set already all the submission dates for the supplementary dossiers. When applications have been received new expiry dates will be set for batches of substances.

Regarding the first batch of substances covered by Group 3 –which had an expiry date of 28 February 2016 and 31 March 2016- Regulation (EU) No 533/2013 has been published to set for these substances a new expiry date of 31 October 2017.

Regarding the second batch of substances covered by Group 3 –which had an expiry date of 30 April 2016 and 30 June 2016- Regulation (EU) No 762/2013 has been published to set for these substances a new expiry date of 31 October 2017 respectively 31 January 2018.

Regarding the third batch of substances covered by Group 3 –which had an expiry date of 31 July 2016 and 30 September 2016- Regulation (EU) No 1136/2013 has been published to set for these substances a new expiry date of 31 January 2018.

Regarding the fourth batch of substances covered by Group 3 –which had an expiry date of 30 November 2016- Regulation (EU) No 85/2014 has been published to set for these substances a new expiry date of 31 January 2018.

Regarding the fifth batch of substances covered by Group 3 –which had an expiry date of 31 January 2017- Regulation (EU) No 487/2014 has been published to set for these substances a new expiry date of 30 April 2018.

Regarding the sixth batch of substances covered by Group 3 –which had an expiry date of 30 April 2017 and 31 May 2017- Regulation (EU) No 678/2014 has been published to set for these substances a new expiry date of 30 April 2018.

Regarding the seventh batch of substances covered by Group 3 –which had an expiry date of 31 May 2017- Regulation (EU) No 878/2014 has been published to set for these substances a new expiry date of 30 April 2018.

Applications for the eight batch of substances covered by Group 3 have been received and a draft Regulation to extend the approval periods for these substances, with a current expiry date of 31 July 2017, has been put on the agenda of the Standing Committee meeting in October 2014 for an opinion/vote.

Regarding the eight batch of substances covered by Group 3 –which had an expiry date of 31 July 2017- Regulation (EU) 2015/415 has been published to set for these substances a new expiry date of 30 April 2018.

Regarding the ninth batch of substances covered by Group 3 –which had an expiry date of 30 September 2017 and 30 November 2017- Regulation (EU) 2015/404has been published to set for these substances a new expiry date of 31 July 2018.

Regarding the substances covered by **Group 4** no extension is expected to be necessary routinely. However extensions still can be possible ‘where for reasons beyond the control of the applicant it appears that the approval is likely to expire before a decision has been taken on renewal’.

GROUP 1: 40 substances with previous expiry date before 14 June 2014

For all the substances which approvals expired before 14 June 2014 applicants were not in a position to apply at the latest three years before the approval period was due to lapse. Therefore, it was necessary to extend the period of approval of these active substances to enable the applicants to prepare their applications and to enable the Commission to examine such applications (batch 1, 2 and 3). Regulation (EU) No 823/2012 to set new expiry dates has been published on 15 September 2012.

Batch 1: 15 active substances

Applications for batch 1 of Group 1 had to be submitted by 31 July 2013. No application for renewal of the approval of the active substances ethoxysulfuron, oxadiargyl and warfarin has been submitted. Now the expiry date will be set at the earliest date possible after the original date of expiry (Regulation (EU) No 186/2014). Regulation (EU) 2017/841 extended the approval periods for some active substances in batch 1. Regulation (EU) 2018/917 extended the approval periods for some active substances in batch 2.

Active substance	**Previous Expiry date**		**New Expiry date**	**Application** (Previous E-date – 3 y)	**Dossier submission** (New E-date – 2.5 y)
Ethofumesate	28 February 2013		Renewal	31 July 2013	31 January 2014
Cyazofamid	30 June 2013		31 July 2019	31 July 2013	31 January 2014
Ethoxysulfuron*	Exp. 31 March 2014				
Foramsulfuron	30 June 2013		31 July 2019	31 July 2013	31 January 2014
Imazamox	30 June 2013		Renewal	31 July 2013	31 January 2014
Oxadiargyl*	Exp. 31 March 2014				
Oxasulfuron	30 June 2013		Non-renewal	31 July 2013	31 January 2014
Carfentrazone ethyl	30 September 2013		Renewal	31 July 2013	31 January 2014
Fenamidone	30 September 2013		Non-renewal	31 July 2013	31 January 2014
Isoxaflutole	30 September 2013		31 July 2019	31 July 2013	31 January 2014
Mesotrione	30 September 2013		Renewal	31 July 2013	31 January 2014
Trifloxystrobin	30 September 2013		Renewal	31 July 2013	31 January 2014
Warfarin*	Exp. 31 March 2014				
Linuron	31 December 2013		Non-renewal	31 July 2013	31 January 2014
Pendimethalin	31 December 2013		Renewal	31 July 2013	31 January 2014

***** *No application for renewal of approval has been submitted.*

Batch 2: 14 active substances

Applications for batch 2 of Group 1 had to be submitted by 31 October 2013. No application for renewal of the approval of the active substance cyfluthrin has been submitted. Now the expiry date will be set at the earliest date possible after the original date of expiry (Regulation (EU) No 460/2014). Regulation (EU) 2017/1511 and Regulation (EU) 2018/1262 extended the approval periods for some active substances in batch 2.

Active substance	**Previous Expiry date**		**New Expiry date**	**Application** **(Previous E-date – 3 y)**	**Dossier submission** **(New E-date – 2.5 y)**
Deltamethrin	31 October 2013		31 October 2019	31 October 2013	30 April 2014
2,4-DB	31 December 2013		Renewal	31 October 2013	30 April 2014
Beta-cyfluthrin	31 December 2013		31 October 2019	31 October 2013	30 April 2014
Coniothyrium minitans Strain CON/M/91-08 (DSM 9660)	31 December 2013		Renewal	31 October 2013	30 April 2014
Cyfluthrin*	Exp. 30 April 2014				
Dimethenamid-P	31 December 2013		31 October 2019	31 October 2013	30 April 2014
Flufenacet	31 December 2013		31 October 2019	31 October 2013	30 April 2014
Flurtamone	31 December 2013		Non-renewal	31 October 2013	30 April 2014
Fosthiazate	31 December 2013		31 October 2019	31 October 2013	30 April 2014
Iodosulfuron	31 December 2013		Renewal	31 October 2013	30 April 2014
Iprodione	31 December 2013		Non-renewal	31 October 2013	30 April 2014
Maleic hydrazide	31 December 2013		Renewal	31 October 2013	30 April 2014
Picoxystrobin	31 December 2013		Non-renewal	31 October 2013	30 April 2014
Silthiofam	31 December 2013		Renewal	31 October 2013	30 April 2014

***** *No application for renewal of approval has been submitted.*

Batch 3: 11 active substances

All applications for batch 3 of Group 1 have been received before the deadline of 31 January 2014. Regulation (EU) 2018/84 extended the approval periods for some active substances in batch 3.

Active substance	**Previous Expiry date**		**New Expiry date**	**Application** (Previous E-date – 3 y)	**Dossier submission** (New E-date – 2.5 y)
Mesosulfuron	31 March 2014		Renewal	31 January 2014	31 July 2014
Propineb	31 March 2014		Non-renewal	31 January 2014	31 July 2014
Propoxycarbazone	31 March 2014		Renewal	31 January 2014	31 July 2014
Propyzamide	31 March 2014		Renewal	31 January 2014	31 July 2014
Zoxamide	31 March 2014		Renewal	31 January 2014	31 July 2014
Benzoic acid	31 May 2014		Renewal	31 January 2014	31 July 2014
Flazasulfuron	31 May 2014		Renewal	31 January 2014	31 July 2014
Mecoprop**	31 May 2014		Exp. 31 January 2017	31 January 2014	
Mecoprop-P	31 May 2014		31 January 2019	31 January 2014	31 July 2014
Propiconazole	31 May 2014		Non-renewal	31 January 2014	31 July 2014
Pyraclostrobin	31 May 2014		31 January 2019	31 January 2014	31 July 2014

*** No dossier submitted*

GROUP 2: 23 substances with current expiry dates of 31/7/2014 – 30/11/2015

Second group of 23 substances with expiry date between: 31/7/2014 and 30/112015. Extension only if application/letter had been received no later than 3 years before the previous expiry date. This because applicants were not in a position to submit an application according to the format and provisions of Regulation (EU) No 844/2012 *setting out the provisions necessary for the implementation of the renewal procedure for active substances, as provided for in Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market* (batch 4 and 5). Regulation (EU) No 1197/2012 to set new expiry dates has been published on 14 December 2012. Regulation (EU) 2018/524 extended the approval periods for some active substances in batch 4.

Batch 4: 7 active substances

All applications for batch 4 of Group 2 have been received before the deadline of 30 April 2014.

Active substance	**Previous Expiry date**		**New Expiry date**	**Application** (Previous E-date – 3 y)	**Dossier submission** (New E-date – 2.5 y)
Molinate*	Exp. 31 July 2014				
Thiram	31 July 2014		Non-renewal	30 April 2014	31 October 2014
Ziram	31 July 2014		30 April 2019	30 April 2014	31 October 2014
Quinoxyfen	31 August 2014		Non-renewal	30 April 2014	31 October 2014
Mepanipyrim	30 September 2014		30 April 2019	30 April 2014	31 October 2014
Pseudomonas chlororaphis Strain: MA 342	30 September 2014		Renewal	30 April 2014	31 October 2014
Carbendazim*	Exp. 30 Nov 2014				
Acetamiprid	31 December 2014		Renewal	30 April 2014	31 October 2014
Thiacloprid	31 December 2014		30 April 2019	30 April 2014	31 October 2014

***** *No application/letter for renewal of approval has been submitted.*

Batch 5: 16 active substances

Regulation (EU) 2018/917 extended the approval periods for some active substances in batch 5.

Active substance	**Previous Expiry date**		**New Expiry date**	**Application** **(Previous E-date – 3 y)**	**Dossier submission** **(New E-date – 2.5 y)**
Chlorpropham	31 January 2015		31 July 2019	31 July 2014	31 January 2015
Alpha-cypermethrin	28 February 2015		31 July 2019	31 July 2014	31 January 2015
Benalaxyl	28 February 2015		31 July 2019	31 July 2014	31 January 2015
Bromoxynil	28 February 2015		31 July 2019	31 July 2014	31 January 2015
Desmedipham	28 February 2015		31 July 2019	31 July 2014	31 January 2015
Ioxynil*	Exp. 28 Feb 2015				
Phenmedipham	28 February 2015		31 July 2019	31 July 2014	31 January 2015
Ampelomyces quisqualis Strain: AQ 10	31 March 2015		Renewal	31 July 2014	31 January 2015
Gliocladium catenulatum Strain: J1446	31 March 2015		31 July 2019	31 July 2014	31 January 2015
Imazosulfuron†	Exp. 31 July 2017				
Laminarin	31 March 2015		Renewal	31 July 2014	31 January 2015
Methoxyfenozide	31 March 2015		31 July 2019	31 July 2014	31 January 2015
S-metolachlor	31 March 2015		31 July 2019	31 July 2014	31 January 2015
Etoxazole	31 May 2015		31 July 2019	31 July 2014	31 January 2015
Tepraloxydim†	Exp. 31 May 2015				
Bitertanol*	Exp. 29 Aug 2013				
Bifenazate	30 November 2015		31 July 2019	31 July 2014	31 January 2015
Milbemectin	30 November 2015		31 July 2019	31 July 2014	31 January 2015

***** *No application/letter for renewal of approval has been submitted.*
† *Active substance is no longer defended at European level.*

GROUP 3: 55 substances with current expiry dates of 28/2/2016 – 30/11/2017

Third group of **55 substances** with expiry date between: 28/2/2016 and 30/11/2017. Extension only if an application according to the format and provisions of Regulation (EU) No 844/2012 *setting out the provisions necessary for the implementation of the renewal procedure for active substances, as provided for in Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market* has been received no later than 3 years before the current expiry date. Extension is necessary because it is the intention to receive all dossiers according to the new data requirements which will be applicable from 1 January 2014 (based on the current expiry dates the dossier submission for this group of substances should be done before 1 January 2014). Besides EFSA can only deal with a workload of around 60 substances a year and for the first two groups of substances we can already expect up to a maximum of 47 dossiers for 2014. Group 3 covers batches 6, 7, 8 and 9. Regarding the first batches of substances –which had expiry dates from 28 February 2016 till 31 July May 2017- Regulations (EU) No 533/2013, No 762/2013, No 1136/2013, No 85/2014, No 487/2014, No 678/2014, No 878/2014, 2015/404 and 2015/415 have been published to set for these substances a new expiry date of 31 October 2017 respectively 31 July 2018. Additional draft Regulations to set new expiry dates will be submitted to the meeting of the Standing Committee on Plants, Animals, Food and Feed in 2014 for an opinion/vote.

Batch 6: 11 active substances

Voted in Standing Committee meeting of 16-17 May 2013 (Regulation (EU) No 533/2013)

Implementing Regulation (EU) 2017/1511 and Implementing Regulation (EU) 2018/1262 extended the approval periods for some active substances in batch 6.

Active substance	**Previous Expiry date**		**New Expiry date**	**Application** (Previous E-date – 3 y)	**Dossier submission** (New E-date – 2.5 y)
Chlorothalonil	28 February 2016		31 October 2019	28 February 2013	30 April 2015
Chlorotoluron	28 February 2016		31 October 2019	28 February 2013	30 April 2015
Cypermethrin	28 February 2016		31 October 2019	28 February 2013	30 April 2015
Daminozide	28 February 2016		31 October 2019	28 February 2013	30 April 2015
Thiophanate-methyl	28 February 2016		31 October 2019	28 February 2013	30 April 2015
Tribenuron	28 February 2016		Renewal	28 February 2013	30 April 2015
1-methyl-cyclopropene	31 March 2016		31 October 2019	31 March 2013	30 April 2015
Forchlorfenuron	31 March 2016		Renewal	31 March 2013	30 April 2015
Indoxacarb	31 March 2016		31 October 2019	31 March 2013	30 April 2015

Batch 6: 11 active substances (cont'd)

Voted in Standing Committee meeting of 15-16 July 2013 (Regulation (EU) No 762/2013).

Active substance	Previous Expiry date		New Expiry date	Application (Previous E-date – 3 y)	Dossier submission (New E-date – 2.5 y)
MCPA	30 April 2016		31 October 2019	30 April 2013	30 April 2015
MCPB	30 April 2016		31 October 2019	30 April 2013	30 April 2015

Batch 7: 10 active substances

Regulation (EU) 2018/84 and Implementing Regulation (EU) 2018/1796 extended the approval periods for some active substances in batch 7.

Active substance	Previous Expiry date		New Expiry date	Application (Previous E-date – 3 y)	Dossier submission (New E-date – 2.5 y)
Chlorpyrifos	30 June 2016		31 January 2020	30 June 2013	31 July 2015
Chlorpyrifos-methyl	30 June 2016		31 January 2020	30 June 2013	31 July 2015
Mancozeb	30 June 2016		31 January 2020	30 June 2013	31 July 2015
Maneb**	Exp. 31 January 2017			30 June 2013	
Metiram	30 June 2016		31 January 2020	30 June 2013	31 July 2015

*** No dossier submitted*

Batch 7: 10 active substances (cont'd)

Voted in Standing Committee meeting of 2-3 October 2013 (Regulation (EU) No 1136/2013)

Active substance	Previous Expiry date		New Expiry date	Application (Previous E-date – 3 y)	Dossier submission (New E-date – 2.5 y)
Clothianidin†	31 July 2016		31 January 2019	31 July 2013	31 July 2015
Oxamyl	31 July 2016		31 January 2020	31 July 2013	31 July 2015
Pethoxamid	31 July 2016		Renewal	31 July 2013	31 July 2015
Dimoxystrobin	30 September 2016		31 January 2020	30 September 2013	31 July 2015

† *Active substance is no longer defended at European level.*

Batch 7: 10 active substances (cont'd)

Voted in Standing Committee meeting of 13 December 2013 (Regulation (EU) No 85/2014)

Active substance	Previous Expiry date		New Expiry date	Application (Previous E-date – 3 y)	Dossier submission (New E-date – 2.5 y)
Copper compounds: Copper hydroxide Copper oxychloride Copper oxide Bordeaux mixture Tribasic copper sulphate	30 November 2016		Renewal	30 November 2013	31 July 2015

Batch 8: 17 active substances

Voted in Standing Committee meeting of 19-20 March 2014 (Regulation (EU) No 487/2014)

Regulation (EU) 2018/524 extended the approval periods for the active substances in batch 8.

Active substance	Previous Expiry date		New Expiry date	Application (Previous E-date – 3 y)	Dossier submission (New E-date – 2.5 y)
Bacillus subtilis (Cohn 1872) Strain QST 713, identical with strain AQ 713	31 January 2017		30 April 2019	31 January 2014	31 October 2015
Clodinafop	31 January 2017		30 April 2019	31 January 2014	31 October 2015
Metrafenone	31 January 2017		30 April 2019	31 January 2014	31 October 2015
Pirimicarb	31 January 2017		30 April 2019	31 January 2014	31 October 2015
Rimsulfuron	31 January 2017		30 April 2019	31 January 2014	31 October 2015
Spinosad	31 January 2017		30 April 2019	31 January 2014	31 October 2015
Thiamethoxam †	31 January 2017		30 April 2019	31 January 2014	31 October 2015
Tolclofos-methyl	31 January 2017		30 April 2019	31 January 2014	31 October 2015
Triticonazole	31 January 2017		30 April 2019	31 January 2014	31 October 2015

† *Active substance is no longer defended at European level.*

Batch 8: 17 active substances (cont'd)

Voted in Standing Committee meeting of 15-16 May 2014 (Regulation (EU) No 678/2014)

Active substance	Previous Expiry date		New Expiry date	Application (Previous E-date – 3 y)	Dossier submission (New E-date – 2.5 y)
Clopyralid	30 April 2017		30 April 2019	30 April 2014	31 October 2015
Cyprodinil	30 April 2017		30 April 2019	30 April 2014	31 October 2015
Fosetyl	30 April 2017		30 April 2019	30 April 2014	31 October 2015
Trinexapac	30 April 2017		30 April 2019	30 April 2014	31 October 2015
Pyrimethanil	31 May 2017		30 April 2019	31 May 2014	31 October 2015

Batch 8: 17 active substances (cont'd)

Voted in Standing Committee meeting of 10-11 July 2014 (Regulation (EU) No 878/2014)

Active substance	Previous Expiry date		New Expiry date	Application (Previous E-date – 3 y)	Dossier submission (New E-date – 2.5 y)
Dichlorprop-P	31 May 2017		30 April 2019	31 May 2014	31 October 2015
Metconazole	31 May 2017		30 April 2019	31 May 2014	31 October 2015
Triclopyr	31 May 2017		30 April 2019	31 May 2014	31 October 2015

Batch 9: 17 active substances

Voted in Standing Committee meeting of 9-10 October 2014 (Regulation (EU) 2015/415)

Regulation (EU) 2018/917 extended the approval periods for some active substances in batch 9.

Active substance	Previous Expiry date		New Expiry date	Application (Previous E-date – 3 y)	Dossier submission (New E-date – 2.5 y)
Ethephon	31 July 2017		31 July 2019	31 July 2014	31 January 2016
Fenamiphos	31 July 2017		31 July 2019	31 July 2014	31 January 2016

Batch 9: 17 active substances (cont'd)

Voted in Standing Committee meeting of 11-12 December 2014 (Regulation (EU) 2015/404)

Active substance	Previous Expiry date		New Expiry date	Application (Previous E-date – 3 y)	Dossier submission (New E-date – 2.5 y)
Captan	30 September 2017		31 July 2019	30 September 2014	31 January 2016
Dimethoate	30 September 2017		31 July 2019	30 September 2014	31 January 2016
Dimethomorph	30 September 2017		31 July 2019	30 September 2014	31 January 2016
Ethoprophos	30 September 2017		31 July 2019	30 September 2014	31 January 2016
Fipronil**	Exp. 30 September 2017			30 September 2014	
Folpet	30 September 2017		31 July 2019	30 September 2014	31 January 2016
Formetanate	30 September 2017		31 July 2019	30 September 2014	31 January 2016
Glufosinate†	30 September 2017		31 July 2018	30 September 2014	31 January 2016
Methiocarb	30 September 2017		31 July 2019	30 September 2014	31 January 2016
Metribuzin	30 September 2017		31 July 2019	30 September 2014	31 January 2016
Phosmet	30 September 2017		31 July 2019	30 September 2014	31 January 2016
Pirimiphos-methyl	30 September 2017		31 July 2019	30 September 2014	31 January 2016
Propamocarb	30 September 2017		31 July 2019	30 September 2014	31 January 2016

*** No dossier submitted*

† Active substance is no longer defended at European level.

Batch 9: 17 active substances (cont'd)

Voted in Standing Committee meeting of 11-12 December 2014 (Regulation (EU) 2015/404)

Active substance	Previous Expiry date		New Expiry date	Application (Previous E-date – 3 y)	Dossier submission (New E-date – 2.5 y)
Beflubutamid	30 November 2017		31 July 2019	30 November 2014	31 January 2016
Spodoptera exigua nuclear polyhedrosis virus*	Exp. 30 November 2017				

*** No application for renewal of approval has been submitted.*

GROUP 4: substances with current expiry date of 31/7/2018 or later

Fourth group of **28 substances** (the number is related to the original 150 substances from 'AIR-3') with expiry date from: 31/7/2018. From this date onwards applications and dossiers will be dealt with according to the provisions of the "Regulation setting out the provisions necessary for the implementation of the renewal procedure for active substances, as provided for in Regulation (EC) No 1107/2009" (Regulation (EU) No 844/2012). No extension will be necessary routinely however extensions still can be possible 'where for reasons beyond the control of the applicant it appears that the approval is likely to expire before a decision has been taken on renewal'. Regulation (EU) 2018/917 extended the approval periods for some active substances in Group 4. Implementing Regulation (EU) 2018/1262 and Implementing Regulation (EU) 2018/1796 extended the approval periods for some active substances in Group 4.

Active substance	**Previous Expiry date**		**New Expiry date**	**Application** (Previous E-date – 3 y)	**Dossier submission** (New E-date – 2.5 y)
Benthiavalicarb	31 July 2018		31 July 2019	31 July 2015	31 January 2016
Boscalid	31 July 2018		31 July 2019	31 July 2015	31 January 2016
Carvone	31 July 2018		31 July 2019	31 July 2015	31 January 2016
Fluoxastrobin	31 July 2018		31 July 2019	31 July 2015	31 January 2016
Paecilomyces lilacinus strain 251	31 July 2018		31 July 2019	31 July 2015	31 January 2016
Prothioconazole	31 July 2018		31 July 2019	31 July 2015	31 January 2016
Diuron	30 September 2018		30 September 2019	30 September 2015	31 March 2016
Clomazone	31 October 2018		31 October 2019	31 October 2015	30 April 2016
Fludioxonil	31 October 2018		31 October 2019	31 October 2015	30 April 2016
Prosulfocarb	31 October 2018		31 October 2019	31 October 2015	30 April 2016
Tritosulfuron	30 November 2018		30 November 2019	30 November 2015	31 May 2016
Amidosulfuron	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Bifenox	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Chloridazon*	31 December 2018				
Clofentezine	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Dicamba	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Difenoconazole	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Diflubenzuron	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Diflufenican	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Fenoxaprop-P	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Fenpropidin	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Imazaquin*	31 December 2018				
Lenacil	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Nicosulfuron	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Oxadiazon*	31 December 2018				

Active substance	**Previous Expiry date**		**New Expiry date**	**Application** **(Previous E-date – 3 y)**	**Dossier submission** **(New E-date – 2.5 y)**
Picloram	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Pyriproxyfen	31 December 2018		31 December 2019	31 December 2015	30 June 2016
Quinoclamine †	31 December 2018			31 December 2015	30 June 2016

** No application received, † Active substance is no longer defended at European level.*