Commission Implementing Regulation (EU) 2016/183 of 11 February 2016 amending 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 (Text with EEA relevance)

COMMISSION IMPLEMENTING REGULATION (EU) 2016/183

of 11 February 2016

amending 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

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EC) No 1107/2009 of the European Parliament and of the Council of 21 October 2009 concerning the placing of plant protection products on the market and repealing Council Directives 79/117/EEC and 91/414/EEC[(1)], and in particular Article 19, thereof,

Whereas:

(1) Commission Implementing Regulation (EU) No 686/2012[(2)] allocates, for the purposes of the renewal procedures, the evaluation of active substances whose approval expires by 31 December 2018 at the latest to the Member States, naming for each active substance a rapporteur and a co-rapporteur. It is appropriate to allocate the evaluation of active substances whose approval expires by 31 December 2021 at the latest.

(2) That allocation should be made in such a way that a balance is achieved as regards the distribution of the responsibilities and the work between Member States.

(3) Implementing Regulation (EU) No 686/2012 should therefore be amended accordingly.

(4) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

HAS ADOPTED THIS REGULATION:

Article 1

Implementing Regulation (EU) No 686/2012 is amended as follows:

(1) The title is replaced by the following:
Commission Implementing Regulation (EU) No 686/2012 of 26 July 2012 allocating to Member States, for the purposes of the renewal procedure, the evaluation of active substances.

(2) The Annex to Implementing Regulation (EU) No 686/2012 is amended in accordance with the Annex to this Regulation.

***Changes to legislation:** There are outstanding changes not yet made to Commission Implementing Regulation (EU) 2016/183. Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes*

Article 2

This Regulation shall enter into force on the twentieth day following that of its publication in the *Official Journal of the European Union*.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

Done at Brussels, 11 February 2016.

For the Commission

The President

Jean-Claude JUNCKER

***Changes to legislation:** There are outstanding changes not yet made to Commission Implementing Regulation (EU) 2016/183. Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes*

ANNEX

The Annex to Implementing Regulation (EU) No 686/2012 is amended as follows:

(1) The word ‘ANNEX’ is replaced by the following:

ANNEX PART AAllocation of the evaluation of active substances whose approval expires by 31 December 2018 at the latest

(2) The following Part B is added:

PART B

Allocation of the evaluation of active substances whose approval expires after 31 December 2018 and not later than 31 December 2021

Active substance	Rapporteur Member State	Co-rapporteur Member State
1-Decanol	PL	IT
1-Naphthylacetamide (1-NAD)	HU	FR
1-Naphthylacetic acid (1-NAA)	HU	FR
2,5-Dichlorobenzoic acid methylester	FR	AT
2-Phenylphenol (incl. sodium salt orthophenyl phenol)	ES	EL
6-Benzyladenine	SE	NL
8-Hydroxyquinoline incl. oxyquinoleine	ES	NL
Abamectin (aka avermectin)	AT	MT
Acetic acid	AT	NL
Aclonifen	NL	NO
Acrinathrin	FR	ES
Aluminium ammonium sulphate	IE	UK
Aluminium phosphide	AT	EE
Aluminium silicate (aka kaolin)	EL	FR
Aluminium sulphate	NL	CZ
Azimsulfuron	EL	FR

Changes to legislation: *There are outstanding changes not yet made to Commission Implementing Regulation (EU) 2016/183. Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes*

Azoxystrobin	UK	NO
Bacillus thuringiensis subsp. *Aizawai* strain ABTS-1857	NL	DE
Bacillus thuringiensis subsp. *Aizawai* strain GC-91	NL	DE
Bacillus thuringiensis subsp. *Israeliensis* (serotype H-14) strain AM65-52	SE	ES
Bacillus thuringiensis subsp. *Kurstaki* strain ABTS 351	DK	NL
Bacillus thuringiensis subsp. *Kurstaki* strain EG 2348	DK	NL
Bacillus thuringiensis subsp. *Kurstaki* strain PB 54	DK	NL
Bacillus thuringiensis subsp. *Kurstaki* strain SA 11	DK	NL
Bacillus thuringiensis subsp. *Kurstaki* strain SA12	DK	NL
Bacillus thuringiensis subsp. *Tenebrionis* strain NB 176 (TM 14 1)	IT	DE
Beauveria bassiana strain ATCC 74040	DE	NL
Beauveria bassiana strain GHA	DE	NL
Benfluralin	NO	NL
Bensulfuron	IT	ES
Bifenthrin	BE	HU
Bispyribac	IT	PT
Blood meal	AT	LT
Bromadiolone	IT	RO
Bromuconazole	BE	CZ
Bupirimate	NL	UK

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Buprofezin	IT	AT
Calcium carbide	EE	CZ
Calcium carbonate	ES	HU
Calcium phosphide	AT	DE
Carbetamide	UK	FR
Carbon dioxide	FR	AT
Carboxin	HR	LV
Chlormequat	UK	IT
Chlorsulfuron	PL	EL
Clethodim	SE	LT
Cycloxydim	NL	DK
Cydia pomonella Granulovirus (CpGV)	DE	NL
Cyflufenamid	DE	AT
Cymoxanil	LT	FI
Cyproconazole	IE	EE
Cyromazine	ES	IT
Dazomet	BG	NL
Denathonium benzoate	IT	PL
Diclofop	PT	FR
Diethofencarb	BE	ES
Difenacoum	IT	PT
Dimethachlor	HR	AT
Dithianon	AT	EL
Dodemorph	NL	IT
Dodine	ES	DE
Epoxiconazole	UK	PL
Ethylene	NL	UK
Etofenprox	SK	IT
Etridiazole	NL	ES
Extract from tea tree	PL	BG
Fat distilation residues	CZ	FR
Fatty acids C7 to C20 (Pelargonic acid (CAS 112-05-0))	EL	AT