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 (Text with EEA relevance)

[[F1]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**]**

(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) For active substances whose approval expires by 31 December 2018 at the latest, it is appropriate to allocate the evaluation for the purposes of the renewal procedures to the Member States, naming for each active substance a rapporteur and a co-rapporteur. 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.

(2) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS REGULATION:

Textual Amendments

F1 Substituted by 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).

[*F2**Article 1*

For the purposes of the renewal procedure, the evaluation of each active substance set out in the first column of the Annex is allocated either to a rapporteur Member State, as set out in the second column of that Annex, and to a co-rapporteur Member State, as set out in the third column of that Annex, or to a group of Member States acting jointly as rapporteur Member State, as set out in the fourth column of that Annex, where applicable. In the latter case, no co-rapporteur is appointed.**]**

Changes to legislation: *There are outstanding changes not yet made to Commission Implementing Regulation (EU) No 686/2012. 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*

Textual Amendments

F2 Substituted by Commission Implementing Regulation (EU) 2019/724 of 10 May 2019 amending Implementing Regulation (EU) No 686/2012 as regards the nomination of rapporteur Member States and co-rapporteur Member States for the active substances glyphosate, lambda-cyhalothrin, imazamox and pendimethalin and amending Implementing Regulation (EU) No 844/2012 as regards the possibility that a group of Member States assumes jointly the role of the rapporteur Member State (Text with EEA relevance).

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.

***Changes to legislation:** There are outstanding changes not yet made to Commission Implementing Regulation (EU) No 686/2012. 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*

[F1ANNEX

PART A

Allocation of the evaluation of active substances whose approval expires by 31 December 2018 at the latest]

Active substance	**Rapporteur Member State**	**Co-rapporteur Member State**
1-methyl-cyclopropene	UK	PT
2,4-DB	BE	EL
Acetamiprid	NL	ES
Alpha-cypermethrin	BE	EL
Amidosulfuron	FI	HR
Ampelomyces quisqualis Strain: AQ 10	FR	DE
Bacillus subtilis (Cohn 1872) Strain QST 713, identical with strain AQ 713	DE	DK
Beflubutamid	DE	LT
Benalaxyl	RO	PT
Benthiavalicarb	PL	FR
Benzoic acid	HU	NL
Beta-cyfluthrin	DE	HU
Bifenazate	SE	IT
Bifenox	PL	BE
Bitertanol	SE	CZ
Boscalid	SK	FR
Bromoxynil	FR	DE
Captan	AT	IT
Carbendazim	DE	SI
Carfentrazone ethyl	BE	FR
Carvone	NL	SE
Chloridazon	DE	PL
Chlorothalonil	NL	BE
Chlorotoluron	BG	FR
Chlorpropham	NL	ES

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Chlorpyrifos	ES	PL
Chlorpyrifos-methyl	ES	PL
Clodinafop	EL	DE
Clofentezine	ES	NL
Clomazone	DK	DE
Clopyralid	FI	PL
Clothianidin	DE	ES
Coniothyrium minitans Strain CON/M/91-08 (DSM 9660)	NL	EE
Copper compounds	FR	DE
Cyazofamid	FR	LV
Cyfluthrin	DE	HU
Cypermethrin	BE	DE
Cyprodinil	FR	BG
Daminozide	CZ	HU
[F3Deltamethrin	AT	]
Desmedipham	FI	DK
Dicamba	DK	RO
Dichlorprop-P	IE	PL
Difenoconazole	ES	UK
Diflubenzuron	EL	SK
[F3Diflufenican	CZ	]
Dimethenamid-P	DE	BG
Dimethoate	IT	BG
[F4Dimethomorph	NL	DE]
Dimoxystrobin	HU	IE
Diuron	DE	DK
Ethephon	NL	PL
Ethofumesate	AT	DK
Ethoprophos	IT	IE
Ethoxysulfuron	IT	AT
Etoxazole	EL	UK
Fenamidone	CZ	FR
Fenamiphos	EL	CY

***Changes to legislation:** There are outstanding changes not yet made to Commission Implementing Regulation (EU) No 686/2012. 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*

Fenoxaprop-P	AT	FI
Fenpropidin	CZ	DE
Fipronil	AT	NL
Flazasulfuron	ES	FR
Fludioxonil	FR	ES
Flufenacet	PL	FR
⌊[F3]Fluoxastrobin	DE	⌋
Flurtamone	CZ	IE
Folpet	AT	IT
Foramsulfuron	FI	SK
Forchlorfenuron	ES	EL
Formetanate	ES	EL
Fosetyl	FR	EE
Fosthiazate	DE	EL
Gliocladium catenulatum Strain: J1446	HU	NL
Glufosinate	DE	FR
Imazamox	FR	IT
Imazaquin	BE	IE
Imazosulfuron	SI	FI
Indoxacarb	FR	ES
Iodosulfuron	SE	FI
Ioxynil	FR	AT
Iprodione	FR	BE
Isoxaflutole	IT	SI
Laminarin	NL	FR
Lenacil	BE	AT
Linuron	IT	DE
Maleic hydrazide	DK	BE
Mancozeb	UK	EL
Maneb	IT	UK
MCPA	PL	NL
MCPB	PL	NL
Mecoprop	PL	IE