Commission Implementing Regulation (EU) 2015/2047 of 16 November 2015 renewing the approval of the active substance esfenvalerate, as a candidate for substitution, in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market, and amending the Annex to Commission Implementing Regulation (EU) No 540/2011 (Text with EEA relevance)

COMMISSION IMPLEMENTING REGULATION (EU) 2015/2047

of 16 November 2015

renewing the approval of the active substance esfenvalerate, as a candidate for substitution, in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market, and amending the Annex to Commission Implementing Regulation (EU) No 540/2011

(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 24 in conjunction with Article 20(1) thereof,

Whereas:

(1) The approval of the active substance esfenvalerate, as set out in Part A of the Annex to Commission Implementing Regulation (EU) No 540/2011[(2)], expires on 30 June 2016.

(2) An application for the renewal of the inclusion of esfenvalerate in Annex I to Council Directive 91/414/EEC[(3)] was submitted in accordance with Article 4 of Commission Regulation (EU) No 1141/2010[(4)] within the time period provided for in that Article.

(3) The applicant submitted the supplementary dossiers required in accordance with Article 9 of Regulation (EU) No 1141/2010. The application was found to be complete by the rapporteur Member State.

(4) The rapporteur Member State prepared a renewal assessment report in consultation with the co-rapporteur Member State and submitted it to the European Food Safety Authority (hereinafter 'the Authority') and the Commission on 30 July 2013.

(5) The Authority communicated the renewal assessment report to the applicant and to the Member States for comments and forwarded the comments received to the Commission. The Authority also made the supplementary summary dossier available to the public.

(6) On 22 October 2014 the Authority communicated to the Commission its conclusion[(5)] on whether esfenvalerate can be expected to meet the approval criteria provided for

Changes to legislation: *There are outstanding changes not yet made to Commission Implementing Regulation (EU) 2015/2047. 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*

in Article 4 of Regulation (EC) No 1107/2009. The Commission presented the draft review report for esfenvalerate to the Standing Committee on Plants, Animals, Food and Feed on 20 March 2015.

(7) It has been established with respect to one or more representative uses of at least one plant protection product containing the active substance that the approval criteria provided for in Article 4 are satisfied. Those approval criteria are therefore deemed to be satisfied.

(8) The risk assessment for the renewal of the approval of esfenvalerate is based on a limited number of representative uses, which however do not restrict the uses for which plant protection products containing esfenvalerate may be authorised. It is therefore appropriate not to maintain the restriction to uses as an insecticide.

(9) The Commission however considers that esfenvalerate is a candidate for substitution pursuant to Article 24 of Regulation (EC) No 1107/2009. Esfenvalerate is a toxic substance in accordance with points 3.7.2.2 and 3.7.2.3 respectively, of Annex II to Regulation (EC) No 1107/2009, given that bioconcentration factor is greater than 2 000 and the long-term no-observed effect concentration for freshwater organisms is less than 0,01 mg/L. Esfenvalerate therefore fulfils the condition set in the second indent of point 4 of Annex II to Regulation (EC) No 1107/2009.

(10) It is therefore appropriate to renew the approval of esfenvalerate as a candidate for substitution.

(11) In accordance with Article 20(3) of Regulation (EC) No 1107/2009, in conjunction with Article 13(4) thereof, the Annex to Implementing Regulation (EU) No 540/2011 should be amended accordingly.

(12) Commission Implementing Regulation (EU) 2015/1885[(6)] extended the expiry date of esfenvalerate to allow the renewal process to be completed before the expiry of the substance. However, given that a decision on renewal has been taken ahead of the original expiry date, this Regulation should apply from the day after the original date of expiry.

(13) 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

Renewal of the approval of the active substance as a candidate for substitution

The approval of the active substance esfenvalerate, as a candidate for substitution, is renewed as set out in Annex I.

***Changes to legislation:** There are outstanding changes not yet made to Commission Implementing Regulation (EU) 2015/2047. 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

Amendments to Implementing Regulation (EU) No 540/2011

The Annex to Implementing Regulation (EU) No 540/2011 is amended in accordance with Annex II to this Regulation.

Article 3

Entry into force and date of application

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

It shall apply from 1 January 2016.

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

Done at Brussels, 16 November 2015.

For the Commission

The President

Jean-Claude JUNCKER

Changes to legislation: *There are outstanding changes not yet made to Commission Implementing Regulation (EU) 2015/2047. 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 I

Common Name, Identification Numbers	**IUPAC Name**	**Purity**[a]	**Date of approval**	**Expiration of approval**	**Specific provisions**
Esfenvalerate CAS No: 66230-04-4 CIPAC No: 481	(αS)-α-cyano-3-phenoxybenzyl (2S)-2-(4-chlorophenyl)-3-methylbutyrate	830 g/kg The impurity toluene shall not exceed 310 g/kg in the technical material.	1 January 2016	31 December 2022	For the implementation of the uniform principles, as referred to in Article 29(6) of Regulation (EC) No 1107/2009, the conclusions of the review report on esfenvalerate, and in particular Appendices I and II thereof, shall be taken into account. In this overall assessment Member States shall pay particular attention to: — the risk from esfenvalerate and the 2SαR-isomer of fenvalerate to aquatic organisms including the risk for bio-

a Further details on identity and specification of active substance are provided in the review report.