

COMMISSION DIRECTIVE 2010/56/EU**of 20 August 2010****amending Annex I to Council Directive 91/414/EEC to renew the inclusion of prohexadione as active substance****(Text with EEA relevance)**

THE EUROPEAN COMMISSION,

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

Having regard to Council Directive 91/414/EEC of 15 July 1991 concerning the placing of plant protection products on the market ⁽¹⁾, and in particular Article 6(1) thereof,

Whereas:

- (1) The inclusion of prohexadione (formerly prohexadione-calcium) in Annex I to Directive 91/414/EEC expires on 31 December 2011. A notification was submitted in accordance with Article 4 of Commission Regulation (EC) No 737/2007 of 27 June 2007 on laying down the procedure of the renewal of the inclusion of a first group of active substances in Annex I to Council Directive 91/414/EEC and establishing the list of those substances ⁽²⁾ for the renewal of the inclusion of prohexadione to as active substance in Annex I to Directive 91/414/EEC within the time period provided for in that Article.
- (2) That notification was found to be admissible by Commission Decision 2008/656/EC of 28 July 2008 on the admissibility of the notifications concerning the renewal of the inclusion in Annex I to Council Directive 91/414/EEC of the active substances azimsulfuron, azoxystrobin, fluroxypyr, imazalil, kresoxim-methyl, prohexadione-calcium and spiroxamin, and establishing the list of the notifiers concerned ⁽³⁾.
- (3) Within the time period provided for in Article 6 of Regulation (EC) No 737/2007, the notifier submitted the data required in accordance with Article 6 of Regulation (EC) No 737/2007 together with an explanation as regards the relevance of each new study submitted.
- (4) The rapporteur Member State prepared an 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 22 May 2009. In addition to the assessment of the substance, that report includes a list of the studies the rapporteur Member State relied on for its assessment.
- (5) The Authority communicated the assessment report to the notifier and to all the Member States and

forwarded the comments received to the Commission. The Authority also made the assessment report available to the public.

- (6) At the request of the Commission, the assessment report was peer reviewed by the Member States and the Authority, the Authority presented its conclusion on the peer review of the risk assessment of prohexadione (considered variant prohexadione-calcium) ⁽⁴⁾ to the Commission on 6 April 2010. The assessment report and the conclusion from the Authority were reviewed by the Member States and the Commission within the Standing Committee on the Food Chain and Animal Health and finalised on 9 July 2010 in the format of the Commission review report for prohexadione.
- (7) It has appeared from the various examinations made that plant protection products containing prohexadione may be expected to continue to satisfy, in general, the requirements laid down in Article 5(1)(a) and (b) of Directive 91/414/EEC, in particular as regards the uses which were examined and detailed in the Commission review report. It is therefore appropriate to renew the inclusion of prohexadione in Annex I to Directive 91/414/EEC, in order to ensure that plant protection products containing this active substance may continue to be authorised where they comply with that Directive.
- (8) A reasonable period should be allowed to elapse before the inclusion of an active substance in Annex I to Directive 91/414/EEC is renewed in order to permit Member States and the interested parties to prepare themselves to meet the new requirements resulting from the renewal.
- (9) Without prejudice to the obligations defined by Directive 91/414/EEC as a consequence of renewing the inclusion of an active substance in Annex I thereof, Member States should be allowed a period of six months after renewal to review authorisations of plant protection products containing prohexadione to make sure that the requirements laid down in Directive 91/414/EEC, in particular in its Article 13, and the relevant conditions set out in Annex I to that Directive, continue to be satisfied. As appropriate, Member States should renew, where appropriate with modifications, or refuse to renew authorisations. By way of derogation from that deadline, a longer period should be provided for the submission and assessment of the update of the complete Annex III dossier of each plant protection product for each intended use in accordance with the uniform principles laid down in Directive 91/414/EEC.

⁽¹⁾ OJ L 230, 19.8.1991, p. 1.

⁽²⁾ OJ L 169, 29.6.2007, p. 10.

⁽³⁾ OJ L 214, 9.8.2008, p. 70.

⁽⁴⁾ European Food Safety Authority; Conclusion on the peer review of the pesticide risk assessment of the active substance prohexadione – on request from the European Commission EFSA Journal 2010; 8(3):1555.

- (10) It is therefore appropriate to amend Directive 91/414/EEC accordingly.
- (11) The measures provided for in this Directive are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS DIRECTIVE:

Article 1

Annex I to Directive 91/414/EEC is amended in accordance with the Annex to this Directive.

Article 2

Member States shall adopt and publish by 31 January 2012 at the latest the laws, regulations and administrative provisions necessary to comply with this Directive. They shall forthwith communicate to the Commission the text of those provisions and a correlation table between those provisions and this Directive.

They shall apply those provisions from 1 February 2012.

When Member States adopt those provisions, they shall contain a reference to this Directive or be accompanied by such a reference on the occasion of their official publication. Member States shall determine how such reference is to be made.

Article 3

1. Member States shall in accordance with Directive 91/414/EEC, where necessary, amend or withdraw existing authorisations for plant protection products containing prohexadione as an active substance by 31 January 2012.

By that date they shall in particular verify that the conditions in Annex I to that Directive relating to prohexadione are met, with the exception of those identified in part B of the entry concerning that active substance, and that the holder of the authorisation has, or has access to, a dossier satisfying the requirements of Annex II to that Directive in accordance with the conditions of Article 13 of that Directive.

2. By way of derogation from paragraph 1, for each authorised plant protection product containing prohexadione as either the only active substance or as one of several active substances, all of which were listed in Annex I to Directive 91/414/EEC by 31 July 2011 at the latest, Member States shall, where necessary, re-evaluate the products, to take into

account developments occurred in the scientific and technical knowledge and in accordance with the uniform principles provided for in Annex VI to Directive 91/414/EEC, on the basis of a dossier satisfying the requirements of Annex III to that Directive and taking into account part B of the entry in Annex I to that Directive concerning prohexadione. On the basis of that evaluation, they shall determine whether the product still satisfies the conditions set out in Article 4(1)(b), (c), (d) and (e) of Directive 91/414/EEC. Following that determination Member States shall, where necessary, amend or withdraw the authorisation by 31 July 2015.

3. By way of derogation from paragraphs 1 and 2, for each authorised plant protection product containing prohexadione as one of several active substances, all of which were listed in Annex I to Directive 91/414/EEC by 31 July 2011 at the latest, and at least one of which was included in Annex I to Directive 91/414/EEC between 1 January 2009 and 31 July 2011, Member States shall re-evaluate the product in accordance with the uniform principles provided for in Annex VI to Directive 91/414/EEC, on the basis of a dossier satisfying the requirements of Annex III to that Directive and taking into account part B of the entry in Annex I to that Directive concerning prohexadione. On the basis of that evaluation, they shall determine whether the product satisfies the conditions set out in Article 4(1)(b), (c), (d) and (e) of Directive 91/414/EEC.

Following that determination Member States shall, where necessary, amend or withdraw the authorisation by 31 July 2015 or by the date fixed for such an amendment or withdrawal in the respective Directive or Directives which added the relevant substance or substances to Annex I to Directive 91/414/EEC, whichever is the latest.

Article 4

This Directive shall enter into force on 1 August 2011.

Article 5

This Directive is addressed to the Member States.

Done at Brussels, 20 August 2010.

For the Commission

The President

José Manuel BARROSO

ANNEX

In Annex I to Directive 91/414/EEC, row No 8 is replaced by the following:

No	Common Name, Identification Numbers	IUPAC Name	Purity ⁽¹⁾	Entry into force	Expiration of inclusion	Specific provisions
'8	prohexadione CAS No: 127277-53-6 (<i>prohexadione-calcium</i>) CIPAC No: 567 (<i>prohexadione</i>) No: 567.020 (<i>prohexadione-calcium</i>)	3,5-dioxo-4-propionylcyclohexanecarboxylic acid	≥ 890 g/kg (expressed as prohexadione-calcium)	1 August 2011	31 July 2021	PART A Only uses as plant growth regulator may be authorised. PART B For the implementation of the uniform principles of Annex VI, the conclusions of the review report on prohexadione and in particular Appendices I and II thereof, as finalised in the Standing Committee on the Food Chain and Animal Health on 9 July 2010 shall be taken into account.'

⁽¹⁾ Further details on identity and specification of active substance are provided in the review report.