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SANCO/03874/2007 rev.1

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COMMISSION OF THE EUROPEAN COMMUNITIES

Brussels, December 2007

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DRAFT*

COMMISSION DECISION

of [...]

**concerning the non-inclusion of azocyclotin, cyhexatin and thidiazuron in Annex I to
Council Directive 91/414/EEC and the withdrawal of authorisations for plant protection
products containing those active substances
(Text with EEA relevance)**

COMMISSION DECISION

of [...]

**concerning the non-inclusion of azocyclotin, cyhexatin and thidiazuron in Annex I to Council Directive 91/414/EEC and the withdrawal of authorisations for plant protection products containing those active substances
(Text with EEA relevance)**

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Council Directive 91/414/EEC of 15 July 1991 concerning the placing of plant protection products on the market¹, and in particular the fourth subparagraph of Article 8(2) thereof,

Whereas:

- (1) Article 8(2) of Directive 91/414/EEC provides that a Member State may, during a period of twelve years following the notification of that Directive, authorise the placing on the market of plant protection products containing active substances not listed in Annex I to that Directive that are already on the market two years after the date of notification, while those substances are gradually being examined within the framework of a programme of work.
- (2) Commission Regulations (EC) No 451/2000² and (EC) No 1490/2002³ lay down the detailed rules for the implementation of the second and third stage of the programme of work referred to in Article 8(2) of Directive 91/414/EEC.
- (3) Azocyclotin, cyhexatin and thidiazuron are substances designated in the third stage programme.
- (4) The sole notifiers for azocyclotin, cyhexatin and thidiazuron informed the Commission on 25 January 2007, 24 January 2007 and 7 February 2007 respectively, that they no longer wished to participate in the programme of work for these active substances, and therefore further information will not be submitted. As a consequence, these active substances should not be included in Annex I to Directive 91/414/EEC.
- (5) Measures should be taken to ensure that existing authorisations for plant protection products containing azocyclotin, cyhexatin or thidiazuron are withdrawn within a

¹ OJ L 230, 19.8.1991, p. 1. Directive as last amended by [INSERT LAST AMENDMENT].

² OJ L 55, 29.2.2000, p. 25. Regulation as last amended by Regulation (EC) No 1044/2003 (OJ L 151, 19.6.2003, p.32).

³ OJ L 224, 21.8.2002, p. 23. Regulation as last amended by Regulation (EC) No 1095/2007 (OJ L 246, 21.9.2007, p. 19).

prescribed period and are not renewed and that no authorisations for such products are granted.

- (6) For these active substances for which there is only a short period of advance notice for the withdrawal of plant protection products containing such substances, it is reasonable to provide for a period of grace for disposal, storage, placing on the market and use of existing stocks for a period no longer than twelve months to allow existing stocks to be used in no more than one further growing. In cases where a longer advance notice period is provided, such period can be shortened to expire at the end of the growing season.
- (7) This Decision does not prejudice the submission of an application for azocyclotin, cyhexatin or thidiazuron according to the provisions of Article 6(2) of Directive 91/414/EEC in view of a possible inclusion in its Annex I.
- (8) The measures provided for in this Decision are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS DECISION:

Article 1

Azocyclotin, cyhexatin and thidiazuron shall not be included in Annex I to Directive 91/414/EEC.

Article 2

Member States shall ensure that:

- (a) authorisations for plant protection products containing azocyclotin, cyhexatin and thidiazuron are withdrawn by ...*;
- (b) from ... ** no authorisations for plant protection products containing azocyclotin, cyhexatin and thidiazuron are granted or renewed under the derogation provided for in Article 8(2) of Directive 91/414/EEC

Article 3

Any period of grace granted by Member States in accordance with Article 4(6) of Directive 91/414/EEC, shall be as short as possible and shall expire not later than ***.

* 6 months from the date of adoption of this Decision

** Date of publication of this Decision

*** 18 months from the date of adoption of this Decision

Article 4

This Decision is addressed to the Member States.

Done at Brussels, [...]

For the Commission
Markos Kyprianou
Member of the Commission