

ANNEX

In Annex I to Directive 91/414/EEC, the following rows are added at the end of the table:

No	Common Name, Identification Numbers	IUPAC Name	Purity ^a	Entry into force	Expiration of inclusion	Specific provisions
'129	Dimoxystrobin CAS No 149961-52-4 CIPAC No 739	(E)-o-(2,5-dimethylphenoxy)methyl-methoxyimino-N-methylphenylacetamide	≥ 980 g/kg	1 October 2006	30 September 2016	<p>PART A Only uses as fungicide may be authorised.</p> <p>PART B In assessing applications to authorise plant protection products containing dimoxystrobin for indoor uses, Member States shall pay particular attention to the criteria in Article 4(1) (b), and shall ensure that any necessary data and information</p>

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

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is provided before such an authorisation is granted.

For the implementation of the uniform principles of Annex VI, the conclusions of the review report on dimoxystrobin, and in particular Appendices I and II thereof, as finalised in the Standing Committee on the Food Chain and Animal Health on 27 January 2006 shall be taken into account. In this overall assessment Member States

— must pay particular attention to the protection of groundwater,

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						<p>when the active substance is applied in a situation with a low crop interception factor, or in regions with vulnerable soil and/or climate conditions; must pay particular attention to the protection of aquatic organisms.</p> <p>Conditions of use shall include risk mitigation measures, where appropriate. The concerned Member States shall request the submission of</p> <p>— a refined risk</p>
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						<p>assessment for birds and mammals considering the formulated active substance; a comprehensive aquatic risk assessment considering the high chronic risk to fish and the effectiveness of potential risk mitigation measures, particularly taking into account run-off and drainage.</p> <p>They shall ensure that the notifiers at whose request dimoxystrobin has been included in this Annex provide such studies to the Commission</p>
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						within two years from the entry into force of this Directive.’
a	Further details on identity and specification of active substances are provided in the review report.					