## ANNEX

The following entry shall be added at the end of the table in Annex I to Directive 91/414/EEC:

'No	Common name, identificatio numbers	IUPAC name on	Purity <sup>a</sup>	Entry into force	Expiration of inclusion	Specific provisio	ons
125	identification	on (R)-2- [4-(5-	≥ 950 g/kg (expressed as clodinafop- propargyl)	1 February 2007		PART A PART B	Only uses as herbicide may be authorised.
							thereof, as finalised in the
							Standing Committee on the Food

							Chain and Animal Health on 27 January 2006 shall be taken into account.
126	Pirimicarb CAS No 23103-98-2 CIPAC No 231	2- dimethylami dimethylpyri yl dimethylcart	imidin-4-	1 February 2007	31 January 2017	PART A	uses as insecticide may be authorised.
						PART B	the implementation of the uniform principles of Annex VI, the conclusions of the review report on pirimicarb, and in particular Appendices I and II thereof, as
<b>a</b> Further deta	ails on identity and	specification of ac	tive substance are	provided in the rev	view report.'		finalised in the

Standing Committee on the Food Chain and Animal Health on 27 January 2006 shall be taken into account. Member States must pay particular attention to the safety of operators and ensure that conditions of use prescribe the application of adequate personal protective equipment. Member States must pay particular attention to the protection of aquatic organisms and must ensure that the conditions of authorisation include risk

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

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mitigation measures, where appropriate, such as buffer zones. The on, ly

						The
						concerned
						Member
						States shall
						request the
						submission
						of further
						studies to
						confirm
						the long
						term risk
						assessment
						for birds
						and for
						potential
						groundwater
						contamination,
						in
						particular
						concerning
						metabolite
						R35140.
						They shall
						ensure that
						the notifiers
						at whose
						request pirimicarb
						has been
						included in
						this Annex
						provide
						such
						studies
						to the
						Commission
						within two
						years from
						the entry
						into force
						of this
						Directive.
127	Rimsulfuron	1-(4-6	≥960 g/kg	1 February	31 January	PART A Only
·		dimethoxypy		2007	2017	uses
a Further deta	ils on identity and		` <b>A</b>			

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

a

CAS No 122931-48-0	yl)-3-(3- ethylsulfony	as I <b>12</b> msulfuron)			as herbicide
(rimsulfuron)	pyridylsulfo	nvl)			may
CIPAC No	urea				be
716	ureu				authorised.
/10					
				PART B	
					the
					implement
					of
					the
					uniform
					principles
					of
					Annex
					VI,
					the
					conclusion
					of
					the
					review
					report
					on
					rimsulfuro
					and
					in
					particular
					Appendice
					I
					and
					II
					thereof,
					as
					finalised
					in
					the
					Standing
					Committee
					on
					the
					Food
					Chain
					and
					Animal
					Health
					on
					27
					January
					2006
					shall
					be
					taken

						Member States must pay particula attention to the protection of non target plants an groundw in vulnerab situation Condition of authorisa should include r mitigatic measures where appropria	r n d ater le s. ns ution isk m 5,
128 a Further det	Tolclofos- methyl CAS No 57018-04-9 CIPAC No 479	O-2,6- dichloro-p- tolyl O,O- dimethyl phosphorothi O-2,6- dichloro-4- methylpheny O,O- dimethyl phosphorothi	l	1 February 2007	31 January 2017	PART A	uses as fungicide may be authorised.

> (seed) treatment in potato and soil treatment in lettuce within greenhouses, Member States shall pay particular attention to the criteria in Article 4(1) (b), and shall ensure that any necessary data and information 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

<b>a</b> Further details on identity and specification of active substance are provided in the review rep
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					tolclofos methyl, and in particula Appendid I and II thereof, a finalised in the Standing Committ on the Fo Chain an Animal Health on 27 Janua 2006 shall be taken inta	r ces as ee bod d n ry o
129	Triticonazole CAS No 131983-72-7 CIPAC No 652	e(±)- (E)-5-(4- chlorobenzyl dimethyl-1- (1 <i>H</i> -1,2,4- triazol-1- ylmethyl)cyd	1 February 2007	31 January 2017	PART A	uses as fungicide may be authorised.

> the criteria in Article 4(1) (b), and shall ensure that any necessary data and information 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 triticonazole, 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

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

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account. In this overall assessment Member States: must \_\_\_\_ pay particular attention to the operator safety. Conditions of authorisation should include protective measures, where appropriate, must pay particular attention to the potential for groundwater contamination, in particular from the highly persistent active substance and its metabolite RPA 406341, in vulnerable zones, must pay particular attention

<b>a</b> Further details on identity and specification of active substance are provided in the review report	a	Further details on identity an	l specification of active substance are	provided in the review report.
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> to the protection of granivorous birds (long term risk). Conditions of authorisation should include risk mitigation measures, where appropriate. The concerned Member States shall request the submission of further studies to confirm the risk assessment for granivorous birds. They shall ensure that the notifier at whose request triticonazole has been included in this Annex provide such studies to the Commission within two years from the entry into force

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