

ANNEX

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

No	Common name, identification numbers	IUPAC name	Purity <sup>a</sup>	Entry into force	Expiration of inclusion	Specific provisions
'324	Myclobutanil CAS No: 88671-89-0 CIPAC No: 442	(RS)-2-(4-chlorophenyl)-1H-1,2,4-triazol-1-ylmethyl)hexan-2-one	≥ 925 g/kg The impurity 1-methylpyrrolidin-2-one shall not exceed 1g/kg in the technical material	1 June 2011	31 May 2021	PART A Only uses as fungicide may be authorised.  PART B For the implementation of the uniform principles of Annex VI, the conclusions of the review report on myclobutanil, and in particular Appendices I and II thereof, as finalised in the Standing Committee on the Food

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

---

*Status: EU Directives are being published on this site to aid cross referencing from UK legislation. After IP completion day (31 December 2020 11pm) no further amendments will be applied to this version.*

---

Chain and Animal Health on 23 November 2010 shall be taken into account.

In this overall assessment Member States shall pay particular attention to the operator safety and ensure that conditions of use prescribe the application of adequate personal protective equipment where appropriate. Conditions of use shall include risk mitigation measures, where appropriate. The Member States concerned shall request the submission of confirmatory information

---

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

---

---

**Status:** EU Directives are being published on this site to aid cross referencing from UK legislation. After IP completion day (31 December 2020 11pm) no further amendments will be applied to this version.

---

					on the residues of myclobutanil and its metabolites in following growing seasons and information confirming that the available residue data cover all compounds of the residue definition. The Member States concerned shall ensure that the applicant submits such confirmatory information to the Commission by 31 January 2013.'
--	--	--	--	--	---

---

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

---