Commission Regulation (EC) No 1490/2002 of 14 August 2002 laying down further detailed rules for the implementation of the third stage of the programme of work referred to in Article 8(2) of Council Directive 91/414/EEC and amending Regulation (EC) No 451/2000 (Text with EEA relevance) (revoked)

Article 1	Subject matter and scope
Article 2	Definitions
Article 3	Member State authority
Article 4	Measures in case of imbalances
Article 5	Withdrawal or replacement of notifier
Article 6	Submission and checking of data lists
Article 7	Submission of dossiers
Article 8	Submission of information by third parties
Article 9	Completeness check of dossiers
Article 10	Evaluation by the rapporteur Member State
Article 11	Receipt of and access to the draft assessment report
Article 11a	Examination of the draft assessment report
Article 11b	Active substance with clear indications that they do not have any
	harmful effects
Article 11c	Consultation of the EFSA
Article 11d	Submission of additional information after the draft assessment
	report has been submitted to the EFSA
Article 11e	Withdrawal by notifier
Article 11f	Active substance for which there are clear indications of harmful
	effects
Article 12	Presentation of a draft directive or draft decision
Article 12a	View by the EFSA
Article 13	Finalised review report
Article 14	Suspension of time limits
Article 15	Measures taken by Member States
Article 16	Interim progress report
Article 17	Fees
Article 18	Other charges, taxes, levies or fees
Article 19	Temporary measures
Article 20	Amendment to Regulation (EC) No 451/2000
Article 21	Entry into force
	Signature

ANNEX I

List of active substances (column A), rapporteur Member States (column B) and notifying producers (code identification) (column C)

PART A

PART B

ANNEX II

List of notifiers' code identification, names and addresses

ANNEX III

Coordinating authority in the Member States (more details are available at the following website: http://www.europa.eu.int/comm/food/fs/ph_ps/pro/index_en.htm)

	AUSTRIA
	BELGIUM
	DENMARK
	GERMANY
	GREECE
	SPAIN
	FINLAND
	FRANCE
	IRELAND
	ITALY
	LUXEMBOURG
	NETHERLANDS
	PORTUGAL
	SWEDEN
	UNITED KINGDOM
	ANNEX IV
Organisations in the Member States to be contacted concerning further details on the payment of the fees referred to in Article 17 and to which such fees have to be paid	

AUSTRIA BELGIUM DENMARK GERMANY Commission Regulation (EC) No 1490/2002. (See end of Document for details)

Changes to legislation: There are currently no known outstanding effects for the

GREECE

SPAIN

FINLAND

FRANCE

IRELAND

ITALY

LUXEMBOURG

THE NETHERLANDS

PORTUGAL

SWEDEN

UNITED KINGDOM

ANNEX V

Criteria for clear indications of no harmful effects

An active substance shall be considered as fulfilling the requirement,...

- 1. The active substance satisfies the following criteria:
- 2. At least one supported representative use of the active substance...

ANNEX VI

Criteria for clear indications of harmful effects

An active substance shall be considered as fulfilling the requirement,...

- 1. As regards the active substance, the existing evidence is not...
- 2. As regards each supported representative use, at least one of...

Changes to legislation:

There are currently no known outstanding effects for the Commission Regulation (EC) No 1490/2002.