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COMMISSION DECISION

of 16 October 2000

adopting Community import decisions for certain chemicals pursuant to Council Regulation (EEC) No 2455/92 concerning the export and import of certain dangerous chemicals

(notified under document number C(2000) 2685)

(Text with EEA relevance)

(2000/657/EC)

(OJ L 275, 27.10.2000, p. 44)

Amended by:

		Official Journal		
		No	page	date
►**M1**	Commission Decision 2001/852/EC of 19 November 2001	L 318	28	4.12.2001

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COMMISSION DECISION

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adopting Community import decisions for certain chemicals pursuant to Council Regulation (EEC) No 2455/92 concerning the export and import of certain dangerous chemicals

(notified under document number C(2000) 2685)

(Text with EEA relevance)

(2000/657/EC)

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Council Regulation (EEC) No 2455/92 of 23 July 1992 concerning the export and import of certain dangerous chemicals [1], as last amended by Regulation (EC) No 2247/98 [2], and in particular Article 5(2) thereof,

Whereas:

(1) Regulation (EEC) No 2455/92 provides that the Commission is to decide for each chemical subject to the Prior Informed Consent (PIC) procedure whether or not the Community consents, possibly subject to specified conditions, to its import.

(2) The United Nations Environment Programme (UNEP) and the Food and Agriculture Organisation (FAO) have been appointed to provide secretariat services for the operation of the interim PIC procedure established by the Final Act of the Conference of Plenipotentiaries on the Convention on the Prior Informed Consent (PIC) Procedure for Certain Hazardous Chemicals and Pesticides in International Trade, signed at Rotterdam on 10 September 1998, in particular the Resolution on interim arrangements thereof.

(3) Additional chemicals have been added to the interim PIC procedure, as pesticide or pesticide formulation, for which the Commission has received information from the interim Secretariat in the form of decision guidance documents.

(4) The Commission, acting as common designated authority, is required to forward decisions on chemicals to the Secretariat of the interim PIC procedure, on behalf of the Community and its Member States.

(5) The interim Secretariat has requested that the participants in the PIC procedure use the specific form for importing country response to report on their import decisions.

(6) The Commission is required, whenever practicable, to make use of existing Community procedures and to ensure that the responses do not conflict with existing Community legislation. However, the Commission also needs to take into consideration Member States' bans or severe restrictions, pending a Community decision.

[1] OJ L 251, 29.8.1992, p. 13.
[2] OJ L 282, 20.10.1998, p. 12.

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(7) The substances binapacryl, captafol, hexachlorobenzene, pentachlorophenol and toxaphene are banned or severely restricted at Community level, in particular by Council Directive 79/117/EEC of 21 December 1978 prohibiting the placing on the market and use of plant protection products containing certain active substances ([1]), as last amended by Directive 91/188/EEC ([2]), or by Council Directive 76/769/EEC of 27 July 1976 on the approximation of the laws, regulations and administrative provisions of the Member States relating to restrictions on the marketing and use of certain dangerous substances and preparations ([3]), as last amended by Directive 1999/77/EC ([4]). Accordingly, a final decision on import should be taken for these substances.

(8) The substances 2,4,5-T, chlorobenzilate, lindane, methamidophos, methylparathion, monochrotophos, parathion and phosphamidon are subject to Community legislation, and in particular to Council Directive 91/414/EEC of 15 July 1991 concerning the placing of plant protection products on the market ([5]), as last amended by Directive 2000/50/EC ([6]), or Directive 98/8/EC of the European Parliament and of the Council of 16 February 1998 concerning the placing of biocidal products on the market ([7]), both providing for a transitional period during which the Member States are allowed to take national decisions on substances and products falling within their scopes, pending a Community decision. Accordingly, an interim decision on import should be taken for these substances.

(9) The measures provided for in this Decision are in accordance with the opinion of the Committee set up pursuant to Article 29 of Council Directive 67/548/EEC ([8]),

HAS DECIDED AS FOLLOWS:

Sole Article

The decisions on import concerning the chemical substances 2,4,5-T, binapacryl, captafol, chlorobenzilate, hexachlorobenzene, lindane, methamidophos, methylparathion, monochrotophos, parathion, pentachlorophenol, phosphamidon and toxaphene are hereby adopted as reported on the importing country response forms contained in the Annex.

([1]) OJ L 33, 8.2.1979, p. 36.
([2]) OJ L 92, 13.4.1991, p. 42.
([3]) OJ L 262, 27.9.1976, p. 201.
([4]) OJ L 207, 6.8.1999, p. 18.
([5]) OJ L 230, 19.8.1991, p. 1.
([6]) OJ L 198, 4.8.2000, p. 39.
([7]) OJ L 123, 24.4.1998, p. 1.
([8]) OJ 196, 16.8.1967, p. 1.

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ANNEX

RESPONSE FORMS FOR IMPORTING COUNTRY

The European Community (Member States: Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Netherlands, Portugal, Spain, Sweden, United Kingdom and the members of the EEA Agreement: Iceland, Liechtenstein and Norway)

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SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	2,4,5-T
1.2.	CAS number	93-76-5
1.3.	Type of formulation and content of active ingredient	
SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES		
	X Pesticide ❑ Industrial ❑ Severely hazardous pesticide formulation	
SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY		
3.1.	**X** This is a first time import response for this chemical in the country.	
3.2.	❑ This is a modification of a previous response. The previous response was a final decision. ❑ Yes ❑ No The previous response was an interim response. ❑ Yes ❑ No Date of issue of the previous response: _____	
SECTION 4. RESPONSE REGARDING FUTURE IMPORT		
❑ **Final decision** ***(fill in section 5)*** **OR** **X Interim response** ***(fill in section 6)***		
SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES		
5.1.	❑ **No consent to import** Is the import of the chemical from all sources simultaneously prohibited? ❑ Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? ❑ Yes ❑ No	
5.2.	❑ **Consent to import**	
5.3.	❑ **Consent to import only subject to specified conditions** Are the conditions for import of the chemical the same for all sources of import? ❑ Yes ❑ No Are the conditions for domestic production of the chemical for domestic use the same as for all imports? ❑ Yes ❑ No	
5.4.	**National legislative or administrative measure upon which the final decision is based**	
	Description of the national legislative or administrative measure: The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure:	

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5.5.	**Remarks** See under points 5.3. and 5.4.			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			
SECTION 6. INTERIM RESPONSE				
6.1.	❑ **No consent to import**			
	Is the import of the chemical from all sources simultaneously prohibited?		❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?		❑ Yes	❑ No
6.2.	❑ **Consent to import**			
6.3.	**X Consent to import only subject to specified conditions** The specified conditions are: Member States that do not consent to import: Austria, Belgium, Denmark, Finland, Germany, Ireland, Italy, Luxembourg, the Netherlands, Spain, Sweden and the members of the EEA Agreement: Iceland, Liechtenstein and Norway. Member States that consent to import (for import prior written authorisation is required): France, Greece, Portugal and the United Kingdom.			
	Are the conditions for import of the chemical the same for all sources of import?		**X** Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?		**X** Yes	❑ No
6.4.	**Indication of active consideration in order to reach a final decision**			
	Is a final decision under active consideration?		**X Yes**	❑ **No**
	The following administrative action is being undertaken during the period in which a final decision is being considered: — 2,4,5-T is included in the Community programme for evaluation of existing active substances under Council Directive 91/414/EEC of 15 July 1991 concerning the placing of plant protection products on the market (OJ L 230 of 19.8.1991, p. 1), — the chemical is already prohibited in the following Member States: Austria, Belgium, Denmark, Finland, Germany, Ireland, Italy, Luxembourg, the Netherlands, Spain, Sweden and the members of the EEA Agreement: Iceland, Liechtenstein and Norway, in their national legislation. Approximate time needed before a final decision can be reached: by 2003 The complete name and address of the responsible institution/authority actively considering a final decision: European Community and its Member States (see address in section 8)			

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6.5.	**Information or assistance requested in order to reach a final decision**			
	The following additional information is requested from the Secretariat: The following additional information is requested from the country that notified the final regulatory action: The following assistance is requested from the Secretariat in evaluating the chemical:			
6.6.	**Remarks**			
	Has there ever been a request for registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			

SECTION 7. RELEVANT ADDITIONAL INFORMATION
2,4,5-T is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ 196, 16.8.1967, p. 1) as: Xn; R 22 (harmful; harmful if swallowed) — Xi; R 36/37/38 (irritant; irritating to eyes, respiratory system and skin) — N; R 50/53 (dangerous to the environment; very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment)

SECTION 8. DESIGNATED NATIONAL AUTHORITY	
Institution	European Commission, Directorate-General for the Environment
Address	Rue de la Loi/Wetstraat 200, B-1049 Brussels

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SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Binapacryl
1.2.	CAS number	485-31-4
1.3.	Type of formulation and content of active ingredient	
SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES		
	X Pesticide ❑ Industrial ❑ Severely hazardous pesticide formulation	
SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY		
3.1.	**X** This is a first time import response for this chemical in the country.	
3.2.	❑ This is a modification of a previous response. The previous response was a final decision. ❑ Yes ❑ No The previous response was an interim response. ❑ Yes ❑ No Date of issue of the previous response: _____	
SECTION 4. RESPONSE REGARDING FUTURE IMPORT		
X Final decision *(fill in section 5)* **OR** **❑ Interim response *(fill in section 6)***		
SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES		
5.1.	**X No consent to import** Is the import of the chemical from all sources simultaneously prohibited? **X** Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? **X** Yes ❑ No	
5.2.	**❑ Consent to import**	
5.3.	**❑ Consent to import only subject to specified conditions** Are the conditions for import of the chemical the same for all sources of import? ❑ Yes ❑ No Are the conditions for domestic production of the chemical for domestic use the same as for all imports? ❑ Yes ❑ No	
5.4.	**National legislative or administrative measure upon which the final decision is based**	
	Description of the national legislative or administrative measure: Binapacryl is listed in Annex I to Council Regulation (EEC) No 2455/92 of 23 July 1992 concerning the export and import of certain dangerous chemicals (OJ L 251, 29.8.1992, p. 13) as banned for use as a plant protection product. It is prohibited to use or place on the market all plant protection products containing binapacryl as an active ingredient according to Council Directive 79/117/EEC of 21 December 1978 prohibiting the placing on the market and use of plant protection products containing certain active substances (OJ L 33, 8.2.1979, p. 36), as amended by Directive 90/533/EEC of 15 October 1990 (OJ L 296, 27.10.1990, p. 63). The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure: European Community and its Member States (see address in section 8)	

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5.5.	**Remarks** See under points 5.3. and 5.4.			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			
SECTION 6. INTERIM RESPONSE				
6.1.	❑ **No consent to import**			
	Is the import of the chemical from all sources simultaneously prohibited?		❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?		❑ Yes	❑ No
6.2.	❑ **Consent to import**			
6.3.	❑ **Consent to import only subject to specified conditions**			
	The specified conditions are:			
	Are the conditions for import of the chemical the same for all sources of import?		❑ Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?		❑ Yes	❑ No
6.4.	**Indication of active consideration in order to reach a final decision**			
	Is a final decision under active consideration?		❑ **Yes**	❑ **No**
	The following administrative action is being undertaken during the period in which a final decision is being considered:			
	Approximate time needed before a final decision can be reached: _____			
	The complete name and address of the responsible institution/authority actively considering a final decision:			

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6.5.	**Information or assistance requested in order to reach a final decision**			
	The following additional information is requested from the Secretariat: The following additional information is requested from the country that notified the final regulatory action: The following assistance is requested from the Secretariat in evaluating the chemical:			
6.6.	**Remarks**			
	Has there ever been a request for registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			
SECTION 7. RELEVANT ADDITIONAL INFORMATION				
Binapacryl is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ L 196, 16.8.1967, p. 1) as: Repr. Cat. 2; R 61 (reproductive toxicity in category 2; may cause harm to the unborn child) — Xn; R 21/22 (harmful in contact with skin and if swallowed).				
SECTION 8. DESIGNATED NATIONAL AUTHORITY				
Institution	European Commission, Directorate-General for the Environment			
Address	Rue de la Loi/Wetstraat 200, B-1049 Brussels			

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SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Captafol
1.2.	CAS number	2425-06-1
1.3.	Type of formulation and content of active ingredient	
SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES		
	X Pesticide ❑ Industrial ❑ Severely hazardous pesticide formulation	
SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY		
3.1.	**X** This is a first time import response for this chemical in the country.	
3.2.	❑ This is a modification of a previous response. The previous response was a final decision. ❑ Yes ❑ No The previous response was an interim response. ❑ Yes ❑ No Date of issue of the previous response: _____	
SECTION 4. RESPONSE REGARDING FUTURE IMPORT		
X Final decision *(fill in section 5)* **OR** **❑ Interim response *(fill in section 6)***		
SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES		
5.1.	**X No consent to import** Is the import of the chemical from all sources simultaneously prohibited? **X** Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? **X** Yes ❑ No	
5.2.	**❑ Consent to import**	
5.3.	**❑ Consent to import only subject to specified conditions** Are the conditions for import of the chemical the same for all sources of import? ❑ Yes ❑ No Are the conditions for domestic production of the chemical for domestic use the same as for all imports? ❑ Yes ❑ No	
5.4.	**National legislative or administrative measure upon which the final decision is based**	
	Description of the national legislative or administrative measure: Captafol is listed in Annex I to Council Regulation (EEC) No 2455/92 of 23 July 1992 concerning the export and import of certain dangerous chemicals as banned for use as a plant protection product (OJ L 251, 29.8.1992, p. 13). It is prohibited to use or place on the market all plant protection products containing captafol as an active ingredient according to Directive 79/117/EEC of 21 December 1978 (OJ L 33, 8.2.1979, p. 36), as amended by Directive 90/533/EEC of 15 October 1990 (OJ L 296, 27.10.1990, p. 63). The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure: European Community and its Member States (see address in section 8)	

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5.5.	**Remarks** See under points 5.3. and 5.4.			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			
SECTION 6. INTERIM RESPONSE				
6.1.	❑ **No consent to import**			
	Is the import of the chemical from all sources simultaneously prohibited?		❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?		❑ Yes	❑ No
6.2.	❑ **Consent to import**			
6.3.	❑ **Consent to import only subject to specified conditions**			
	The specified conditions are:			
	Are the conditions for import of the chemical the same for all sources of import?		❑ Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?		❑ Yes	❑ No
6.4.	**Indication of active consideration in order to reach a final decision**			
	Is a final decision under active consideration?		❑ **Yes**	❑ **No**
	The following administrative action is being undertaken during the period in which a final decision is being considered:			
	Approximate time needed before a final decision can be reached: _____			
	The complete name and address of the responsible institution/authority actively considering a final decision:			

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6.5.	**Information or assistance requested in order to reach a final decision**			
	The following additional information is requested from the Secretariat: The following additional information is requested from the country that notified the final regulatory action: The following assistance is requested from the Secretariat in evaluating the chemical:			
6.6.	**Remarks**			
	Has there ever been a request for registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			

SECTION 7. RELEVANT ADDITIONAL INFORMATION
Captafol is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ 196, 16.8.1967, p. 1) as: Carc. Cat. 2; R 45 (carcinogen in category 2; may cause cancer) — R 43 (may cause sensitisation by skin contact) — N; R 50/53 (dangerous to the environment; very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment)

SECTION 8. DESIGNATED NATIONAL AUTHORITY	
Institution	European Commission, Directorate-General for the Environment
Address	Rue de la Loi/Wetstraat 200, B-1049 Brussels

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SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Chlorobenzilate
1.2.	CAS number	510-15-6
1.3.	Type of formulation and content of active ingredient	
SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES		
	X Pesticide ❑ Industrial ❑ Severely hazardous pesticide formulation	
SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY		
3.1.	**X** This is a first time import response for this chemical in the country.	
3.2.	❑ This is a modification of a previous response. The previous response was a final decision. ❑ Yes ❑ No The previous response was an interim response. ❑ Yes ❑ No Date of issue of the previous response: _____	
SECTION 4. RESPONSE REGARDING FUTURE IMPORT		
❑ **Final decision** ***(fill in section 5)*** **OR** **X Interim response** ***(fill in section 6)***		
SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES		
5.1.	❑ **No consent to import** Is the import of the chemical from all sources simultaneously prohibited? ❑ Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? ❑ Yes ❑ No	
5.2.	❑ **Consent to import**	
5.3.	❑ **Consent to import only subject to specified conditions** Are the conditions for import of the chemical the same for all sources of import? ❑ Yes ❑ No Are the conditions for domestic production of the chemical for domestic use the same as for all imports? ❑ Yes ❑ No	
5.4.	**National legislative or administrative measure upon which the final decision is based**	
	Description of the national legislative or administrative measure: The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure:	

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5.5.	**Remarks** See under points 5.3. and 5.4.			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			
SECTION 6. INTERIM RESPONSE				
6.1.	❑ **No consent to import**			
	Is the import of the chemical from all sources simultaneously prohibited?		❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?		❑ Yes	❑ No
6.2.	❑ **Consent to import**			
6.3.	**X Consent to import only subject to specified conditions** The specified conditions are: Member States that do not consent to import: Belgium, Denmark, Finland, Greece, Ireland, Italy, Luxembourg, the Netherlands, Spain, Sweden and the members of the EEA Agreement: Iceland, Liechtenstein and Norway. Member States that consent to import (for import prior written authorisation is required): Austria, France, Germany, Portugal and the United Kingdom.			
	Are the conditions for import of the chemical the same for all sources of import?		**X** Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?		**X** Yes	❑ No
6.4.	**Indication of active consideration in order to reach a final decision**			
	Is a final decision under active consideration?		**X Yes**	❑ **No**
	The following administrative action is being undertaken during the period in which a final decision is being considered: — chlorobenzilate is included in the Community programme for evaluation of existing active substances under Council Directive 91/414/EEC of 15 July 1991 concerning the placing of plant protection products on the market (OJ L 230, 19.8.1991, p. 1), — the chemical is already prohibited in the following Member States: Belgium, Denmark, Finland, Greece, Ireland, Italy, Luxembourg, the Netherlands, Spain, Sweden and the members of the EEA Agreement: Iceland, Liechtenstein and Norway, in their national legislation. Approximate time needed before a final decision can be reached: by 2003 The complete name and address of the responsible institution/authority actively considering a final decision: European Community and its Member States (see address in section 8)			

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6.5.	**Information or assistance requested in order to reach a final decision**			
	The following additional information is requested from the Secretariat: The following additional information is requested from the country that notified the final regulatory action: The following assistance is requested from the Secretariat in evaluating the chemical:			
6.6.	**Remarks**			
	Has there ever been a request for registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			

SECTION 7. RELEVANT ADDITIONAL INFORMATION
Chlorobenzilate is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ 196, 16.8.1967, p. 1) as: Xn; R 22 (harmful; harmful if swallowed) — N; R 50/53 (dangerous to the environment; very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment).

SECTION 8. DESIGNATED NATIONAL AUTHORITY	
Institution	European Commission, Directorate-General for the Environment
Address	Rue de la Loi/Wetstraat 200, B-1049 Brussels

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SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Hexachlorobenzene
1.2.	CAS number	118-74-1
1.3.	Type of formulation and content of active ingredient	
SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES		
	X Pesticide ❑ Industrial ❑ Severely hazardous pesticide formulation	
SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY		
3.1.	**X** This is a first time import response for this chemical in the country.	
3.2.	❑ This is a modification of a previous response. The previous response was a final decision. ❑ Yes ❑ No The previous response was an interim response. ❑ Yes ❑ No Date of issue of the previous response: _____	
SECTION 4. RESPONSE REGARDING FUTURE IMPORT		
X Final decision *(fill in section 5)* **OR** **❑ Interim response *(fill in section 6)***		
SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES		
5.1.	**X No consent to import** Is the import of the chemical from all sources simultaneously prohibited? **X** Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? **X** Yes ❑ No	
5.2.	**❑ Consent to import**	
5.3.	**❑ Consent to import only subject to specified conditions** Are the conditions for import of the chemical the same for all sources of import? ❑ Yes ❑ No Are the conditions for domestic production of the chemical for domestic use the same as for all imports? ❑ Yes ❑ No	
5.4.	**National legislative or administrative measure upon which the final decision is based**	
	Description of the national legislative or administrative measure: Hexachlorobenzene is listed in Annex I to Council Regulation (EEC) No 2455/92 of 23 July 1992 concerning the export and import of certain dangerous chemicals (OJ L 251, 29.8.1992, p. 13) as banned for use as a plant protection product. It is prohibited to place on the market or use plant protection products containing hexachlorobenzene as an active ingredient in accordance with Directive 79/117/EEC of 21 December 1978 prohibiting the placing on the market and use of plant protection products containing certain active substances (OJ L 33, 8.2.1979, p. 36). The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure: European Community and its Member States (see address in section 8)	

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5.5.	**Remarks** See under points 5.3. and 5.4.			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			
SECTION 6. INTERIM RESPONSE				
6.1.	**❑ No consent to import**			
	Is the import of the chemical from all sources simultaneously prohibited?		❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?		❑ Yes	❑ No
6.2.	**❑ Consent to import**			
6.3.	**❑ Consent to import only subject to specified conditions**			
	The specified conditions are:			
	Are the conditions for import of the chemical the same for all sources of import?		❑ Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?		❑ Yes	❑ No
6.4.	**Indication of active consideration in order to reach a final decision**			
	Is a final decision under active consideration?		**❑ Yes**	**❑ No**
	The following administrative action is being undertaken during the period in which a final decision is being considered:			
	Approximate time needed before a final decision can be reached: _____			
	The complete name and address of the responsible institution/authority actively considering a final decision:			

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6.5.	**Information or assistance requested in order to reach a final decision**			
	The following additional information is requested from the Secretariat: The following additional information is requested from the country that notified the final regulatory action: The following assistance is requested from the Secretariat in evaluating the chemical:			
6.6.	**Remarks**			
	Has there ever been a request for registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			

SECTION 7. RELEVANT ADDITIONAL INFORMATION
Hexachlorobenzene is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ 196 of 16.8.1967, p. 1) as: Carc. Cat. 2; R 45 (carcinogen in category 2; may cause cancer) — T; R 48/25 (toxic; toxic: danger of serious damage to health by prolonged exposure if swallowed) — N; R 50/53 (dangerous to the environment; very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment).

SECTION 8. DESIGNATED NATIONAL AUTHORITY	
Institution	European Commission, Directorate-General for the Environment
Address	Rue de la Loi/Wetstraat 200, B-1049 Brussels

▼**M1**

Interim Secretariat for the Rotterdam Convention on the Prior Informed Consent Procedure for Certain Hazardous Chemicals and Pesticides in International Trade

UNEP

FORM
FOR IMPORTING COUNTRY RESPONSE

IMPORTANT: See instructions before filling in the form

COUNTRY: **European Community (Member States: Belgium, Denmark, Germany, Greece, Spain, France, Ireland, Italy, Luxembourg, Netherlands, Austria, Portugal, Finland, Sweden, United Kingdom)**

SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Ethylene dichloride (1,2-dichloroethane)
1.2.	CAS number	107-06-2
1.3.	Type of formulation and content of active ingredient	Liquid

SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES

X Pesticide

❑ Industrial

❑ Severely hazardous pesticide formulation

SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY			
3.1.	X This is a first time import response for this chemical in the country.		
3.2.	❑ This is a modification of a previous response.		
	The previous response was a final decision.	❑ Yes	❑ No
	The previous response was an interim response.	❑ Yes	❑ No
	Date of issue of the previous response: _____		

SECTION 4. RESPONSE REGARDING FUTURE IMPORT

X **Final decision** ***(Fill in Section 5, page 2)*** OR ❑ **Interim response** ***(Fill in Section 6, pages 3 and 4)***

SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES			
5.1.	X **No consent to import**		
	Is the import of the chemical from all sources simultaneously prohibited?	X Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?	X Yes	❑ No
5.2.	❑ **Consent to import**		

▼**M1**

5.3.	❑ **Consent to import only subject to specified conditions**		
	The specified conditions are:		
	Are the conditions for import of the chemical the same for all sources of import?	❑ Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?	❑ Yes	❑ No
5.4.	**National legislative or administrative measure upon which the final decision is based**		
	Description of the national legislative or administrative measure: Ethylene dichloride (1,2-dichloroethane) is listed in Annex I to Regulation (EEC) No 2455/92 of 23 July 1992 concerning the export and import of certain dangerous chemicals (OJ L 251, 29.8.1992, p. 13), as amended by Regulation (EEC) No 3135/94 (OJ L 332, 22.12.1994, p. 1), as banned for use as a plant protection product. It is prohibited to use or place on the market all plant protection products containing 1,2-dichloroethane as an active ingredient according to Council Directive 79/117/EEC of 21 December 1978 prohibiting the placing on the market and use of plant protection products containing certain active substances (OJ L 33, 8.2.1979, p. 36), as amended by Directive 87/181/EEC (OJ L 71, 14.3.1987, p. 33). The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure: European Community and its Member States (see address in Section 8)		
5.5.	**Remarks**		
	Has there ever been a request of registration of this chemical in the country?	❑ Yes	❑ No
	Is this chemical currently registered in the country?	❑ Yes	❑ No
	Is this chemical manufactured in the country?	❑ Yes	❑ No
	Is this chemical formulated in the country?	❑ Yes	❑ No
	If yes to either one of these last two questions: — Is this intended for domestic use?	❑ Yes	❑ No
	If yes to either one of these last two questions: — Is this intended for export?	❑ Yes	❑ No
	Other remarks:		

SECTION 6. INTERIM RESPONSE			
6.1.	❑ **No consent to import**		
	Is the import of the chemical from all sources simultaneously prohibited?	❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?	❑ Yes	❑ No
6.2.	❑ **Consent to import**		
6.3.	❑ **Consent to import only subject to specified conditions**		
	The specified conditions are:		
	Are the conditions for import of the chemical the same for all sources of import?	❑ Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?	❑ Yes	❑ No

▼**M1**

6.4.	**Indication of active consideration in order to reach a final decision**			
	Is a final decision under active consideration?		❑ **Yes**	❑ **No**
	The following administrative action is being undertaken during the period a final decision is being considered: Approximate time needed before a final decision can be reached: _____ The complete name and address of the responsible institution/authority actively considering a final Decision:			
6.5.	**Information or assistance requested in order to reach a final decision**			
	The following additional information is requested from the Secretariat: The following additional information is requested from the country that notified the final regulatory action: The following assistance is requested from the Secretariat in evaluating the chemical:			
6.6.	**Remarks**			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If yes to either one of these last two questions:	Is this intended for domestic use?	❑ Yes	❑ No
		Is this intended for export?	❑ Yes	❑ No
	Other remarks:			

SECTION 7. RELEVANT ADDITIONAL INFORMATION
Ethylene dichloride (1,2-dichloroethane) is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ L 196, 16.8.1967, p. 1) as F; R 11 - Carc.Cat.2; R 45 - Xn; R 22 - Xi; R 36/37/38. R 45: May cause cancer. R 11: Highly flammable. R 22: Harmful if swallowed. R 36/37/38: Irritating to eyes, respiratory system and skin. Ethylene dichloride (1,2-dichloroethane) has been classified by the EC as a category 2 carcinogen (probably carcinogenic to humans).

SECTION 8. DESIGNATED NATIONAL AUTHORITY	
Institution	European Commission, Directorate-General for the Environment
Address	Rue de la Loi/Wetstraat 200 B-1049 Brussels

▼**M1**

Interim Secretariat for the Rotterdam Convention on the Prior Informed Consent Procedure for Certain Hazardous Chemicals and Pesticides in International Trade

UNEP

FORM
FOR IMPORTING COUNTRY RESPONSE

IMPORTANT: See instructions before filling in the form

COUNTRY: **European Community (Member States: Belgium, Denmark, Germany, Greece, Spain, France, Ireland, Italy, Luxembourg, Netherlands, Austria, Portugal, Finland, Sweden, United Kingdom)**

SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Ethylene oxide
1.2.	CAS number	75-21-8
1.3.	Type of formulation and content of active ingredient	Liquefied gas.

SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES

X Pesticide

❑ Industrial

❑ Severely hazardous pesticide formulation

SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY			
3.1.	X This is a first time import response for this chemical in the country.		
3.2.	❑ This is a modification of a previous response.		
	The previous response was a final decision.	❑ Yes	❑ No
	The previous response was an interim response.	❑ Yes	❑ No
	Date of issue of the previous response: _____		

SECTION 4. RESPONSE REGARDING FUTURE IMPORT

X **Final decision** ***(Fill in Section 5, page 2)*** OR ❑ **Interim response** ***(Fill in Section 6, pages 3 and 4)***

SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES			
5.1.	X **No consent to import**		
	Is the import of the chemical from all sources simultaneously prohibited?	X Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?	X Yes	❑ No
5.2.	❑ **Consent to import**		

▼**M1**

5.3.	❑ **Consent to import only subject to specified conditions** The specified conditions are: Are the conditions for import of the chemical the same for all sources of import? ❑ Yes ❑ No Are the conditions for domestic production of the chemical for domestic use the same as for all imports? ❑ Yes ❑ No		
5.4.	**National legislative or administrative measure upon which the final decision is based**		
	Description of the national legislative or administrative measure: Ethylene oxide is listed in Annex I to Council Regulation (EEC) No 2455/92 of 23 July 1992 concerning the export and import of certain dangerous chemicals (OJ L 251, 29.8.1992, p. 13), as amended by Regulation (EEC) No 3135/94 (OJ L 332, 22.12.1994, p. 1) as banned for use as a plant protection product. It is prohibited to use or place on the market all plant protection products containing ethylene oxide as an active ingredient according to Council Directive 79/117/EEC of 21 December 1978 prohibiting the placing on the market and use of plant protection products containing certain active substances (OJ L 33, 8.2.1979, p. 36), as amended by Directive 86/355/EEC (OJ L 212, 2.8.1986, p. 33). The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure: European Community and its Member States (see address in Section 8).		
5.5.	**Remarks see under points 5.3 and 5.4**		
	Has there ever been a request of registration of this chemical in the country?		❑ Yes ❑ No
	Is this chemical currently registered in the country?		❑ Yes ❑ No
	Is this chemical manufactured in the country?		❑ Yes ❑ No
	Is this chemical formulated in the country?		❑ Yes ❑ No
	If yes to either one of these last two questions:	Is this intended for domestic use?	❑ Yes ❑ No
		Is this intended for export?	❑ Yes ❑ No
	Other remarks:		

SECTION 6. INTERIM RESPONSE			
6.1.	❑ **No consent to import** Is the import of the chemical from all sources simultaneously prohibited? ❑ Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? ❑ Yes ❑ No		
6.2.	❑ **Consent to import**		
6.3.	❑ **Consent to import only subject to specified conditions** The specified conditions are: Are the conditions for import of the chemical the same for all sources of import? ❑ Yes ❑ No Are the conditions for domestic production of the chemical for domestic use the same as for all imports? ❑ Yes ❑ No		

▼**M1**

6.4.	**Indication of active consideration in order to reach a final decision**		
	Is a final decision under active consideration?	❑ **Yes**	❑ **No**
	The following administrative action is being undertaken during the period a final decision is being considered:		
	Approximate time needed before a final decision can be reached: _____		
	The complete name and address of the responsible institution/authority actively considering a final decision:		
6.5.	**Information or assistance requested in order to reach a final decision**		
	The following additional information is requested from the Secretariat:		
	The following additional information is requested from the country that notified the final regulatory action:		
	The following assistance is requested from the Secretariat in evaluating the chemical:		
6.6.	**Remarks**		
	Has there ever been a request of registration of this chemical in the country?	❑ Yes	❑ No
	Is this chemical currently registered in the country?	❑ Yes	❑ No
	Is this chemical manufactured in the country?	❑ Yes	❑ No
	Is this chemical formulated in the country?	❑ Yes	❑ No
	If yes to either one of these last two questions: — Is this intended for domestic use?	❑ Yes	❑ No
	If yes to either one of these last two questions: — Is this intended for export?	❑ Yes	❑ No
	Other remarks:		

SECTION 7. RELEVANT ADDITIONAL INFORMATION
Ethylene oxide is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ L 196, 16.8.1967, p. 1), as F+; R 12 - Carc.Cat.2; R 45 - Muta.Cat.2; R 46 - T; R 23 - Xi; R 36/37/38 R 45: May cause cancer. R 46: May cause heritable genetic damage. R 12: Extremely flammable. R 23: Toxic by inhalation. R 36/37/38: Irritating to eyes, respiratory system and skin. Ethylene oxide has been classified by the EC as a category 2 carcinogen (probably carcinogenic to humans). Ethylene oxide has also been classified by the EC as a category 2 mutagen (probably mutagenic to humans).

SECTION 8. DESIGNATED NATIONAL AUTHORITY	
Institution	European Commission, Directorate-General for the Environment
Address	Rue de la Loi/Wetstraat 200 B-1049 Brussels

▼B

SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Methamidophos
1.2.	CAS number	10265-92-6
1.3.	Type of formulation and content of active ingredient	
SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES		
	❑ Pesticide ❑ Industrial **X** Severely hazardous pesticide formulation	
SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY		
3.1.	**X** This is a first time import response for this chemical in the country.	
3.2.	❑ This is a modification of a previous response. The previous response was a final decision. ❑ Yes ❑ No The previous response was an interim response. ❑ Yes ❑ No Date of issue of the previous response: _____	
SECTION 4. RESPONSE REGARDING FUTURE IMPORT		
❑ **Final decision** ***(fill in section 5)*** **OR** **X Interim response** ***(fill in section 6)***		
SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES		
5.1.	❑ **No consent to import** Is the import of the chemical from all sources simultaneously prohibited? ❑ Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? ❑ Yes ❑ No	
5.2.	❑ **Consent to import**	
5.3.	❑ **Consent to import only subject to specified conditions** Are the conditions for import of the chemical the same for all sources of import? ❑ Yes ❑ No Are the conditions for domestic production of the chemical for domestic use the same as for all imports? ❑ Yes ❑ No	
5.4.	**National legislative or administrative measure upon which the final decision is based**	
	Description of the national legislative or administrative measure: The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure:	

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5.5.	**Remarks** See under points 5.3. and 5.4.			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			
SECTION 6. INTERIM RESPONSE				
6.1.	❑ **No consent to import**			
	Is the import of the chemical from all sources simultaneously prohibited?		❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?		❑ Yes	❑ No
6.2.	❑ **Consent to import**			
6.3.	**X Consent to import only subject to specified conditions** The specified conditions are: Member States that do not consent to import: Denmark, Ireland, Sweden and the members of the EEA Agreement: Iceland, Liechtenstein and Norway. Member States that consent to import (for import prior written authorisation is required): Austria, Belgium, Finland, France, Germany, Greece, Italy, Luxembourg, the Netherlands, Spain, Portugal and the United Kingdom.			
	Are the conditions for import of the chemical the same for all sources of import?		**X** Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?		**X** Yes	❑ No
6.4.	**Indication of active consideration in order to reach a final decision**			
	Is a final decision under active consideration?		**X Yes**	❑ **No**
	The following administrative action is being undertaken during the period in which a final decision is being considered: — methamidophos is included in the Community programme for evaluation of existing active substances under Council Directive 91/414/EEC of 15 July 1991 concerning the placing of plant protection products on the market (OJ L 230, 19.8.1991, p. 1), — the chemical is already prohibited in the following Member States: Denmark, Ireland, Sweden and the members of the EEA Agreement Iceland, Liechtenstein and Norway, in their national legislation. Approximate time needed before a final decision can be reached: by 2003 The complete name and address of the responsible institution/authority actively considering a final decision: European Community and its Member States (see address in section 8)			

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6.5.	**Information or assistance requested in order to reach a final decision**			
	The following additional information is requested from the Secretariat: The following additional information is requested from the country that notified the final regulatory action: The following assistance is requested from the Secretariat in evaluating the chemical:			
6.6.	**Remarks**			
	Has there ever been a request for registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			

SECTION 7. RELEVANT ADDITIONAL INFORMATION

Methamidophos is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ L 196 of 16.8.1967, p. 1) as: T +; R 28 (very toxic; very toxic if swallowed) — T; R 24 (toxic; toxic in contact with skin) — Xi; R 36 (irritant; irritating to eyes) — N; R 50 (dangerous to the environment; very toxic to aquatic organisms).

SECTION 8. DESIGNATED NATIONAL AUTHORITY

Institution	European Commission, Directorate-General for the Environment
Address	Rue de la Loi/Wetstraat 200, B-1049 Brussels

▼B

SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Methyl parathion
1.2.	CAS number	298-00-0
1.3.	Type of formulation and content of active ingredient	
SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES		
	❑ Pesticide ❑ Industrial **X** Severely hazardous pesticide formulation	
SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY		
3.1.	**X** This is a first time import response for this chemical in the country.	
3.2.	❑ This is a modification of a previous response. The previous response was a final decision. ❑ Yes ❑ No The previous response was an interim response. ❑ Yes ❑ No Date of issue of the previous response: _____	
SECTION 4. RESPONSE REGARDING FUTURE IMPORT		
❑ **Final decision** ***(fill in section 5)*** **OR** **X Interim response** ***(fill in section 6)***		
SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES		
5.1.	❑ **No consent to import** Is the import of the chemical from all sources simultaneously prohibited? ❑ Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? ❑ Yes ❑ No	
5.2.	❑ **Consent to import**	
5.3.	❑ **Consent to import only subject to specified conditions** Are the conditions for import of the chemical the same for all sources of import? ❑ Yes ❑ No Are the conditions for domestic production of the chemical for domestic use the same as for all imports? ❑ Yes ❑ No	
5.4.	**National legislative or administrative measure upon which the final decision is based**	
	Description of the national legislative or administrative measure: The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure:	

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5.5.	**Remarks** See under points 5.3. and 5.4.			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			
SECTION 6. INTERIM RESPONSE				
6.1.	❑ **No consent to import**			
	Is the import of the chemical from all sources simultaneously prohibited?		❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?		❑ Yes	❑ No
6.2.	❑ **Consent to import**			
6.3.	**X Consent to import only subject to specified conditions** The specified conditions are: Member States that do not consent to import: Belgium, Denmark, Finland, Ireland, Sweden and the members of the EEA Agreement: Iceland, Liechtenstein and Norway. Member States that consent to import (for import prior written authorisation is required): Austria, France, Germany, Greece, Italy, Luxembourg, the Netherlands, Spain, Portugal and the United Kingdom.			
	Are the conditions for import of the chemical the same for all sources of import?		**X** Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?		**X** Yes	❑ No
6.4.	**Indication of active consideration in order to reach a final decision**			
	Is a final decision under active consideration?		**X Yes**	❑ **No**
	The following administrative action is being undertaken during the period in which a final decision is being considered: — methyl parathion is included in the Community programme for evaluation of existing active substances under Council Directive 91/414/EEC of 15 July 1991 concerning the placing of plant protection products on the market (OJ L 230, 19.8.1991, p. 1) and Directive 98/8/EC of the European Parliament and of the Council of 16 February 1998 concerning the placing of biocidal products on the market (OJ L 123, 24.4.1998, p. 1), — the chemical is already prohibited in the following Member States: Belgium, Denmark, Finland, Ireland, Sweden and the members of the EEA Agreement: Iceland, Liechtenstein and Norway, in their national legislation. Approximate time needed before a final decision can be reached: by 2003 (as PPP) and by 2008 (as biocides) The complete name and address of the responsible institution/authority actively considering a final decision: European Community and its Member States (see address in section 8)			

▼B

6.5.	**Information or assistance requested in order to reach a final decision**			
	The following additional information is requested from the Secretariat: The following additional information is requested from the country that notified the final regulatory action: The following assistance is requested from the Secretariat in evaluating the chemical:			
6.6.	**Remarks**			
	Has there ever been a request for registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			

SECTION 7. RELEVANT ADDITIONAL INFORMATION

Methyl parathion is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ 196, 16.8.1967, p. 1) as: T+; R 28 (very toxic; very toxic if swallowed) — T; R 24 (toxic; toxic in contact with skin)

SECTION 8. DESIGNATED NATIONAL AUTHORITY

Institution	European Commission, Directorate-General for the Environment
Address	Rue de la Loi/Wetstraat 200, B-1049 Brussels

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SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Monocrotophos
1.2.	CAS number	6923-22-4
1.3.	Type of formulation and content of active ingredient	
SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES		
	❑ Pesticide ❑ Industrial **X** Severely hazardous pesticide formulation	
SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY		
3.1.	**X** This is a first time import response for this chemical in the country.	
3.2.	❑ This is a modification of a previous response. The previous response was a final decision. ❑ Yes ❑ No The previous response was an interim response. ❑ Yes ❑ No Date of issue of the previous response: _____	
SECTION 4. RESPONSE REGARDING FUTURE IMPORT		
❑ **Final decision** ***(fill in section 5)*** **OR** **X Interim response** ***(fill in section 6)***		
SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES		
5.1.	❑ **No consent to import** Is the import of the chemical from all sources simultaneously prohibited? ❑ Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? ❑ Yes ❑ No	
5.2.	❑ **Consent to import**	
5.3.	❑ **Consent to import only subject to specified conditions** Are the conditions for import of the chemical the same for all sources of import? ❑ Yes ❑ No Are the conditions for domestic production of the chemical for domestic use the same as for all imports? ❑ Yes ❑ No	
5.4.	**National legislative or administrative measure upon which the final decision is based**	
	Description of the national legislative or administrative measure: The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure:	

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5.5.	**Remarks** See under points 5.3. and 5.4.			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			
SECTION 6. INTERIM RESPONSE				
6.1.	❑ **No consent to import**			
	Is the import of the chemical from all sources simultaneously prohibited?		❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?		❑ Yes	❑ No
6.2.	❑ **Consent to import**			
6.3.	**X Consent to import only subject to specified conditions** The specified conditions are: Member States that do not consent to import: Belgium, Denmark, Ireland, Luxembourg, the Netherlands, Sweden and the members of the EEA Agreement: Iceland, Liechtenstein and Norway. Member States that consent to import (for import prior written authorisation is required): Austria, Finland, France, Germany, Greece, Italy, Spain, Portugal and the United Kingdom.			
	Are the conditions for import of the chemical the same for all sources of import?		**X** Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?		**X** Yes	❑ No
6.4.	**Indication of active consideration in order to reach a final decision**			
	Is a final decision under active consideration?		**X Yes**	❑ **No**
	The following administrative action is being undertaken during the period in which a final decision is being considered: — monochrotophos is included in the Community programme for evaluation of existing active substances under Council Directive 91/414/EEC of 15 July 1991 concerning the placing of plant protection products on the market (OJ L 230, 19.8.1991, p. 1), — the chemical is already prohibited in the following Member States: Belgium, Denmark, Ireland, Luxembourg, the Netherlands, Sweden and the members of the EEA Agreement: Iceland, Liechtenstein and Norway, in their national legislation. Approximate time needed before a final decision can be reached: by 2003 The complete name and address of the responsible institution/authority actively considering a final decision: European Community and its Member States (see address in section 8)			

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6.5.	**Information or assistance requested in order to reach a final decision**			
	The following additional information is requested from the Secretariat: The following additional information is requested from the country that notified the final regulatory action: The following assistance is requested from the Secretariat in evaluating the chemical:			
6.6.	**Remarks**			
	Has there ever been a request for registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			

SECTION 7. RELEVANT ADDITIONAL INFORMATION
Monochrotophos is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ 196, 16.8.1967, p. 1) as: Muta. Cat. 3; R 40 (mutagenic category 3: possible risks of irreversible effects) — T+; R 26/28 (very toxic; very toxic by inhalation and if swallowed) — T; R 24 (toxic; toxic in contact with skin) — N; R 50/53 (dangerous to the environment; very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment)

SECTION 8. DESIGNATED NATIONAL AUTHORITY	
Institution	European Commission, Directorate-General for the Environment
Address	Rue de la Loi/Wetstraat 200, B-1049 Brussels

▼**M1**

Interim Secretariat for the Rotterdam Convention on the Prior Informed Consent Procedure for Certain Hazardous Chemicals and Pesticides in International Trade

UNEP

FORM
FOR IMPORTING COUNTRY RESPONSE

IMPORTANT: See instructions before filling in the form

COUNTRY: European Community (Member States: Belgium, Denmark, Germany, Greece, Spain, France, Ireland, Italy, Luxembourg, Netherlands, Austria, Portugal, Finland, Sweden, United Kingdom

SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Lindane
1.2.	CAS number	58-89-9
1.3.	Type of formulation and content of active ingredient	

SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES

X Pesticide

❑ Industrial

❑ Severely hazardous pesticide formulation

SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY	
3.1.	❑ This is a first time import response for this chemical in the country.
3.2.	X This is a modification of a previous response. The previous response was a final decision. ❑ Yes X No The previous response was an interim response. X Yes ❑ No Date of issue of the previous response: 27.10.2000 _____

SECTION 4. RESPONSE REGARDING FUTURE IMPORT

❑ **Final decision** *(Fill in Section 5, page 2)* OR X **Interim response** *(Fill in Section 6, pages 3 and 4)*

SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES	
5.1.	❑ **No consent to import** Is the import of the chemical from all sources simultaneously prohibited? ❑ Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? ❑ Yes ❑ No
5.2.	❑ **Consent to import**

▼**M1**

5.3.	❑ **Consent to import only subject to specified conditions**			
	The specified conditions are:		❑ Yes	❑ No
	Are the conditions for import of the chemical the same for all sources of import?			
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?		❑ Yes	❑ No
5.4.	**National legislative or administrative measure upon which the final decision is based**			
	Description of the national legislative or administrative measure:			
	The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure:			
5.5.	**Remarks**			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If yes to either one of these last two questions:	Is this intended for domestic use?	❑ Yes	❑ No
		Is this intended for export?	❑ Yes	❑ No
	Other remarks:			

SECTION 6. INTERIM RESPONSE			
6.1.	❑ **No consent to import**		
	Is the import of the chemical from all sources simultaneously prohibited?	❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?	❑ Yes	❑ No
6.2.	❑ **Consent to import**		
6.3.	X **Consent to import only subject to specified conditions**		
	The specified conditions are:		
	For plant protection products		
	It is prohibited to use or place on the market all plant protection products containing lindane. (Commission Decision 2000/801/EC of 20 December 2000 (OJ L 324, 21.12.2000, p. 42) concerning the non-inclusion of lindane in Annex I to Council Directive 91/414/EEC and the withdrawal of authorisations for plant protection products containing this active substance).		
	For biocidal products		
	Member States that consent to import (for import prior written authorisation is required): Belgium, Germany, Greece, Spain, France, Luxembourg, Austria, Portugal and the United Kingdom.		
	Member States that do not consent to import: Denmark, Ireland, Italy, the Netherlands, Finland, Sweden.		
	Are the conditions for import of the chemical the same for all sources of import?	X Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?	X Yes	❑ No

▼M1

6.4.	**Indication of active consideration in order to reach a final decision**
	Is a final decision under active consideration? **X Yes** **❑ No** The following administrative action is being undertaken during the period a final decision is being considered: Lindane was excluded from Annex I to Council Directive 91/414/EEC and thus authorisations for plant-protection products containing this active substance have been withdrawn. (Commission Decision 2000/801/EC (OJ L 324, 21.12.2000, p. 42)). However it is also included in the Community Programme for evaluation of existing active substances under Directive 98/8/EC of the European Parliament and of the Council of 16 February 1998 (OJ L 123, 24.4.1998, p. 1) concerning the placing of biocidal products on the market. Approximate time needed before a final decision can be reached: by 2008, when evaluation for biocidal use will be completed The complete name and address of the responsible institution/authority actively considering a final decision: European Community and its Member States (see address in Section 8)
6.5.	**Information or assistance requested in order to reach a final decision**
	The following additional information is requested from the Secretariat: The following additional information is requested from the country that notified the final regulatory action: The following assistance is requested from the Secretariat in evaluating the chemical:
6.6.	**Remarks**

Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
Is this chemical currently registered in the country?		❑ Yes	❑ No
Is this chemical manufactured in the country?		❑ Yes	❑ No
Is this chemical formulated in the country?		❑ Yes	❑ No
If yes to either one of these last two questions:	Is this intended for domestic use?	❑ Yes	❑ No
	Is this intended for export?	❑ Yes	❑ No
Other remarks:			

SECTION 7. Relevant additional information
Lindane is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ L 196, 16.8.1967, p. 1) as: T; R 23/24/25-R 36/38-N; R 50/53 (R 23/24/25 Toxic by inhalation, in contact with skin and if swallowed. R 36/38 Irritating to eyes and skin. R 50/53 Very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment)

SECTION 8. Designated national authority	
Institution	European Commission, Directorate-General for the Environnement
Address	Rue de la Loi/Wetstraat 200, B-1049 Brussels

▼**M1**

Interim Secretariat for the Rotterdam Convention on the Prior Informed Consent Procedure for Certain Hazardous Chemicals and Pesticides in International Trade

UNEP

FORM
FOR IMPORTING COUNTRY RESPONSE

IMPORTANT: See instructions before filling in the form

COUNTRY: European Community (Member States: Belgium, Denmark, Germany, Greece, Spain, France, Ireland, Italy, Luxembourg, Netherlands, Austria, Portugal, Sweden, United Kingdom)

SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Parathion
1.2.	CAS number	56-38-2
1.3.	Type of formulation and content of active ingredient	

SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES

❑ Pesticide

❑ Industrial

X Severely hazardous pesticide formulation

SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY	
3.1.	❑ This is a first time import response for this chemical in the country.
3.2.	X This is a modification of a previous response. The previous response was a final decision. ❑ Yes X No The previous response was an interim response. X Yes ❑ No Date of issue of the previous response: 27.10.2000 _____

SECTION 4. RESPONSE REGARDING FUTURE IMPORT

❑ **Final decision** ***(Fill in Section 5, page 2)*** OR X **Interim response** ***(Fill in Section 6, pages 3 and 4)***

SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES	
5.1.	❑ **No consent to import** Is the import of the chemical from all sources simultaneously prohibited? ❑ Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? ❑ Yes ❑ No
5.2.	❑ **Consent to import**

▼**M1**

5.3.	❑ **Consent to import only subject to specified conditions**			
	Are the conditions for import of the chemical the same for all sources of import?		❑ Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?		❑ Yes	❑ No
5.4.	**National legislative or administrative measure upon which the final decision is based**			
	Description of the national legislative or administrative measure:			
	The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure:			
5.5.	**Remarks** See under point 5.3 and 5.4			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If yes to either one of these last two questions:	Is this intended for domestic use?	❑ Yes	❑ No
		Is this intended for export?	❑ Yes	❑ No
	Other remarks:			

SECTION 6. INTERIM RESPONSE			
6.1.	❑ **No consent to import**		
	Is the import of the chemical from all sources simultaneously prohibited?	❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?	❑ Yes	❑ No
6.2.	❑ **Consent to import**		
6.3.	X **Consent to import only subject to specified conditions**		
	The specified conditions are:		
	For plant protection products		
	It is prohibited to use or place on the market all plant protection products containing parathion. (Commission Decision 2001/520/EC of 9 July 2001 (OJ L 187, 10.7.2001, p. 47) concerning the non-inclusion of parathion in Annex I to Council Directive 91/414/EEC and the withdrawal of authorisations for plant protection products containing this active substance).		
	For biocidal products		
	Member States that consent to import (for import prior written authorisation is required): Belgium, Germany, Greece, Spain, France, Italy, Luxembourg, Austria, Portugal and the United Kingdom.		
	Member States that do not consent to import: Denmark, Ireland, the Netherlands, Sweden.		
	Are the conditions for import of the chemical the same for all sources of import?	X Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?	X Yes	❑ No

▼**M1**

6.4.	**Indication of active consideration in order to reach a final decision**			
	Is a final decision under active consideration?		**X Yes**	**❑ No**
	Parathion was excluded from Annex I to Council Directive 91/414/EEC and thus authorisations for plant-protection products containing this active sustance have been withdrawn. Commission Decision 2001/520/EC (OJ L 187, 10.7.2001, p. 47).			
	However it is also included in the Community Programme for evaluation of existing active substances under Directive 98/8/EC of the European Parliament and of the Council of 16 February 1998 (OJ L 123, 24.4.1998, p. 1) concerning the placing of biocidal products on the market.			
	Approximate time needed before a final decision can be reached: by 20/08, when evaluation for biocidal use will be completed			
	The complete name and address of the responsible institution/authority actively considering a final decision: European Community and its Member States (see address in Section 8).			
6.5.	**Information or assistance requested in order to reach a final decision**			
	The following additional information is requested from the Secretariat:			
	The following additional information is requested from the country that notified the final regulatory action:			
	The following assistance is requested from the Secretariat in evaluating the chemical:			
6.6.	**Remarks**			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If yes to either one of these last two questions:	Is this intended for domestic use?	❑ Yes	❑ No
		Is this intended for export?	❑ Yes	❑ No
	Other remarks:			

SECTION 7. RELEVANT ADDITIONAL INFORMATION
Parathion is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ L 196, 16.8.1967, p.1) as: T+; R 27/28 (Very Toxic; Very toxic in contact with skin and if swallowed.) — N; R50-53 (Dangerous to the environment; Very Toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment).

SECTION 8. DESIGNATED NATIONAL AUTHORITY	
Institution	European Commission — Directorate General for the Environment
Address	Rue de la Loi/Wetstraat 200 B-1049 Brussels Belgium

▼B

SECTION 1. IDENTIY OF CHEMICAL		
1.1.	Common name	Pentachlorophenol
1.2.	CAS number	87-86-5
1.3.	Type of formulation and content of active ingredient	
SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES		
	X Pesticide ❑ Industrial ❑ Severely hazardous pesticide formulation	
SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY		
3.1.	**X** This is a first time import response for this chemical in the country.	
3.2.	❑ This is a modification of a previous response. The previous response was a final decision. ❑ Yes ❑ No The previous response was an interim response. ❑ Yes ❑ No Date of issue of the previous response: _____	
SECTION 4. RESPONSE REGARDING FUTURE IMPORT		
X Final decision *(fill in section 5)* **OR** ❑ **Interim response *(fill in section 6)***		
SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES		
5.1.	❑ **No consent to import** Is the import of the chemical from all sources simultaneously prohibited? ❑ Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? ❑ Yes ❑ No	
5.2.	❑ **Consent to import**	

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5.3.	**X Consent to import only subject to specified conditions** Member States that do not consent to import: Austria, Belgium, Denmark, Finland, Germany, Greece, Italy, Luxembourg, the Netherlands, Sweden and the members of the EEA Agreement: Iceland, Liechtenstein and Norway. Member States that consent to import for restricted uses by way of derogation until 31 December 2008: France, Ireland, Portugal and the United Kingdom. Member State that consents to import for restricted uses by way of derogation until 1 January 2004: Spain. The following conditions apply: Substances and preparations containing PCP, its salts or esters may be placed on the market for use in industrial installations not permitting the emission and/or discharge of PCP in quantities greater than those prescribed by existing legislation: (a) in the treatment of wood. Treated wood may not be used inside buildings or for the manufacture or retreatment of containers intended for growing purposes, packaging that may come into contact with raw materials, intermediate or finished products destined for human and/or animal consumption; (b) in the impregnation of fibres and heavy-duty textiles not intended in any case for clothing or for decorative furnishings; (c) for special exceptions authorised on a case-by-case basis. In any case, PCP used alone or as a component of preparations employed within the framework of the above exceptions must have a total hexachlorodibenzoparadioxin (HCDD) content of less than 2 ppm, may not be placed on the market in packages of less than 20 litres and may not be sold to the general public. Without prejudice to other labelling requirements, the packaging of such preparations should be marked clearly and indelibly: 'Reserved for industrial and professional use'.		
	Are the conditions for import of the chemical the same for all sources of import?	**X** Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?	**X** Yes	❑ No
5.4.	**National legislative or administrative measure upon which the final decision is based**		
	Description of the national legislative or administrative measure: Pentachlorophenol is listed in Annex I to Council Regulation (EEC) No 2455/92 of 23 July 1992 concerning the export and import of certain dangerous chemicals as a severely restricted chemical (OJ L 251, 29.8.1992, p. 13). The placing on the market and use of products containing pentachlorophenol, its salts and esters, are prohibited by Directive 76/769/EEC of 27 July 1976 on the approximation of the laws, regulations and administrative provisions of the Member States relating to restrictions on the marketing and use of certain dangerous substances and preparations (OJ L 262, 27.9.76, p. 201), as amended by Directive 91/173/EEC of 21 March 1991 (OJ L 85, 5.4.1991, p. 34) and Directive 1999/51/EC (OJ L 142, 5.6.1999, p. 22). The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure: European Community and its Member States (see address in section 8)		
5.5.	**Remarks** See under points 5.3 and 5.4		
	Has there ever been a request of registration of this chemical in the country?	❑ Yes	❑ No
	Is this chemical currently registered in the country?	❑ Yes	❑ No
	Is this chemical manufactured in the country?	❑ Yes	**X** No
	Is this chemical formulated in the country?	❑ Yes	**X** No
	If the answer is 'yes' to either of these last two questions: — Is it intended for domestic use?	❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions: — Is it intended for export?	❑ Yes	❑ No
	Other remarks		

▼B

SECTION 6. INTERIM RESPONSE			
6.1.	❑ **No consent to import**		
	Is the import of the chemical from all sources simultaneously prohibited?	❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?	❑ Yes	❑ No
6.2.	❑ **Consent to import**		
6.3.	❑ **Consent to import only subject to specified conditions**		
	The specified conditions are:		
	Are the conditions for import of the chemical the same for all sources of import?	❑ Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?	❑ Yes	❑ No
6.4.	**Indication of active consideration in order to reach a final decision**		
	Is a final decision under active consideration?	❑ **Yes**	❑ **No**
	The following administrative action is being undertaken during the period in which a final decision is being considered:		
	Approximate time needed before a final decision can be reached: _____		
	The complete name and address of the responsible institution/authority actively considering a final decision:		
6.5.	**Information or assistance requested in order to reach a final decision**		
	The following additional information is requested from the Secretariat:		
	The following additional information is requested from the country that notified the final regulatory action:		
	The following assistance is requested from the Secretariat in evaluating the chemical:		

6.6.	**Remarks**			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either one of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			

SECTION 7. RELEVANT ADDITIONAL INFORMATION
Pentachlorophenol is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ 196, 16.8.1967, p. 1) as: Carc. Cat. 3; R 40 (carcinogen in category 3; possible risks of irreversible effects) — T+; R 26 (very toxic; very toxic by inhalation) — T; R 24/25 (toxic; toxic in contact with skin and if swallowed) — Xi; R 36/37/38 (irritant; irritating to eyes, respiratory system and skin) — N; R 50/53 (dangerous to the environment; very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment).

SECTION 8. DESIGNATED NATIONAL AUTHORITY	
Institution	European Commission, Directorate-General for the Environment
Address	Rue de la Loi/Wetstraat 200, B-1049 Brussels

▼B

SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Phosphamidon
1.2.	CAS number	13171-21-6/23783-98-4/297-99-4
1.3.	Type of formulation and content of active ingredient	
SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES		
	❑ Pesticide ❑ Industrial **X** Severely hazardous pesticide formulation	
SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY		
3.1.	**X** This is a first time import response for this chemical in the country.	
3.2.	❑ This is a modification of a previous response. The previous response was a final decision. ❑ Yes ❑ No The previous response was an interim response. ❑ Yes ❑ No Date of issue of the previous response: _____	
SECTION 4. RESPONSE REGARDING FUTURE IMPORT		
❑ **Final decision** ***(fill in section 5)*** **OR** **X Interim response** ***(fill in section 6)***		
SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES		
5.1.	❑ **No consent to import** Is the import of the chemical from all sources simultaneously prohibited? ❑ Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? ❑ Yes ❑ No	
5.2.	❑ **Consent to import**	
5.3.	❑ **Consent to import only subject to specified conditions** Are the conditions for import of the chemical the same for all sources of import? ❑ Yes ❑ No Are the conditions for domestic production of the chemical for domestic use the same as for all imports? ❑ Yes ❑ No	
5.4.	**National legislative or administrative measure upon which the final decision is based**	
	Description of the national legislative or administrative measure: The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure:	

▼B

5.5.	**Remarks** See under points 5.3. and 5.4.			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			
SECTION 6. INTERIM RESPONSE				
6.1.	❑ **No consent to import**			
	Is the import of the chemical from all sources simultaneously prohibited?		❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?		❑ Yes	❑ No
6.2.	❑ **Consent to import**			
6.3.	**X Consent to import only subject to specified conditions** The specified conditions are: Member States that do not consent to import: Belgium, Denmark, Ireland, Luxembourg, the Netherlands, and the members of the EEA Agreement: Iceland, Liechtenstein and Norway. Member States that consent to import (for import prior written authorisation is required): Austria, Finland, France, Germany, Greece, Italy, Spain, Portugal, Sweden and the United Kingdom.			
	Are the conditions for import of the chemical the same for all sources of import?		**X** Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?		**X** Yes	❑ No
6.4.	**Indication of active consideration in order to reach a final decision**			
	Is a final decision under active consideration?		**X Yes**	❑ **No**
	The following administrative action is being undertaken during the period in which a final decision is being considered: — phosphamidon is included in the Community programme for evaluation of existing active substances under Council Directive 91/414/EEC of 15 July 1991 concerning the placing of plant protection products on the market (OJ L 230, 19.8.1991, p. 1), — the chemical is already prohibited in the following Member States: Belgium, Denmark, Ireland, Luxembourg, the Netherlands and the members of the EEA Agreement: Iceland, Liechtenstein and Norway, in their national legislation. Approximate time needed before a final decision can be reached: by 2003 The complete name and address of the responsible institution/authority actively considering a final decision: European Community and its Member States (see address in section 8)			

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6.5.	**Information or assistance requested in order to reach a final decision**			
	The following additional information is requested from the Secretariat: The following additional information is requested from the country that notified the final regulatory action: The following assistance is requested from the Secretariat in evaluating the chemical:			
6.6.	**Remarks**			
	Has there ever been a request for registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			

SECTION 7. RELEVANT ADDITIONAL INFORMATION

Phosphamidon is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ 196, 16.8.1967, p. 1) as: Muta. Cat. 3; R 40 (mutagenic category 3: possible risks of irreversible effects — T+; R 28 (very toxic; very toxic if swallowed) — T; R 24 (toxic: toxic in contact with skin) — N; R 50/53 (dangerous to the environment; very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment)

SECTION 8. DESIGNATED NATIONAL AUTHORITY

Institution	European Commission, Directorate-General for the Environment
Address	Rue de la Loi/Wetstraat 200, B-1049 Brussels

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SECTION 1. IDENTITY OF CHEMICAL		
1.1.	Common name	Toxaphene
1.2.	CAS number	8001-35-2
1.3.	Type of formulation and content of active ingredient	
SECTION 2. THE IMPORT RESPONSE PROVIDED IN THIS FORM APPLIES TO THE FOLLOWING CATEGORY OR CATEGORIES		
	X Pesticide ❑ Industrial ❑ Severely hazardous pesticide formulation	
SECTION 3. INDICATION REGARDING PREVIOUS RESPONSE, IF ANY		
3.1.	**X** This is a first time import response for this chemical in the country.	
3.2.	❑ This is a modification of a previous response. The previous response was a final decision. ❑ Yes ❑ No The previous response was an interim response. ❑ Yes ❑ No Date of issue of the previous response: _____	
SECTION 4. RESPONSE REGARDING FUTURE IMPORT		
X Final decision *(fill in section 5)* **OR** ❑ **Interim response *(fill in section 6)***		
SECTION 5. FINAL DECISION, PURSUANT TO NATIONAL LEGISLATIVE OR ADMINISTRATIVE MEASURES		
5.1.	**X No consent to import** Is the import of the chemical from all sources simultaneously prohibited? **X** Yes ❑ No Is domestic production of the chemical for domestic use simultaneously prohibited? **X** Yes ❑ No	
5.2.	❑ **Consent to import**	
5.3.	❑ **Consent to import only subject to specified conditions** Are the conditions for import of the chemical the same for all sources of import? ❑ Yes ❑ No Are the conditions for domestic production of the chemical for domestic use the same as for all imports? ❑ Yes ❑ No	
5.4.	**National legislative or administrative measure upon which the final decision is based**	
	Description of the national legislative or administrative measure: Toxaphene is listed in Annex I to Council Regulation (EEC) No 2455/92 of 23 July 1992 concerning the export and import of certain dangerous chemicals as banned for use as a plant protection product (OJ L 251, 29.8.1992, p. 13). It is prohibited to use or place on the market all plant protection products containing toxaphene as an active ingredient according to Directive 79/117/EEC of 21 December 1978 prohibiting the placing on the market and use of plant protection products containing active substances (OJ L 33, 8.2.1979, p. 36), as amended by Directive 83/131/EEC of 14 March 1983 (OJ L 91, 9.4.1983, p. 35). The complete name and address of the institution/authority responsible for issuing this national legislative or administrative measure: European Community and its Member States (see address in section 8)	

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5.5.	**Remarks** See under points 5.3. and 5.4.			
	Has there ever been a request of registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			
SECTION 6. INTERIM RESPONSE				
6.1.	❑ **No consent to import**			
	Is the import of the chemical from all sources simultaneously prohibited?		❑ Yes	❑ No
	Is domestic production of the chemical for domestic use simultaneously prohibited?		❑ Yes	❑ No
6.2.	❑ **Consent to import**			
6.3.	❑ **Consent to import only subject to specified conditions**			
	The specified conditions are:			
	Are the conditions for import of the chemical the same for all sources of import?		❑ Yes	❑ No
	Are the conditions for domestic production of the chemical for domestic use the same as for all imports?		❑ Yes	❑ No
6.4.	**Indication of active consideration in order to reach a final decision**			
	Is a final decision under active consideration?		❑ **Yes**	❑ **No**
	The following administrative action is being undertaken during the period in which a final decision is being considered:			
	Approximate time needed before a final decision can be reached: _____			
	The complete name and address of the responsible institution/authority actively considering a final decision:			

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6.5.	**Information or assistance requested in order to reach a final decision**			
	The following additional information is requested from the Secretariat: The following additional information is requested from the country that notified the final regulatory action: The following assistance is requested from the Secretariat in evaluating the chemical:			
6.6.	**Remarks**			
	Has there ever been a request for registration of this chemical in the country?		❑ Yes	❑ No
	Is this chemical currently registered in the country?		❑ Yes	❑ No
	Is this chemical manufactured in the country?		❑ Yes	❑ No
	Is this chemical formulated in the country?		❑ Yes	❑ No
	If the answer is 'yes' to either of these last two questions:	Is it intended for domestic use?	❑ Yes	❑ No
		Is it intended for export?	❑ Yes	❑ No
	Other remarks			

SECTION 7. RELEVANT ADDITIONAL INFORMATION
Toxaphene is classified under Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances (OJ 196, 16.8.1967, p. 1) as: Carc. Cat. 3; R 40 (carcinogen in category 3; possible risks of irreversible effects) — T; R 25 (toxic; toxic if swallowed) — Xn; R 21 (harmful; harmful in contact with skin) — Xi; R 37/38 (irritant; irritating to respiratory system and skin) — N; R 50/53 (dangerous to the environment; very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment)

SECTION 8. DESIGNATED NATIONAL AUTHORITY	
Institution	European Commission, Directorate-General for the Environment
Address	Rue de la Loi/Wetstraat 200, B-1049 Brussels