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**Stockholm Convention on Persistent Organic Pollutants Persistent Organic Pollutants Revi ew Committee First meeting** Geneva, 7–11 November 2005 Item 4 of the provisional agenda\*

**Operational procedures** 

# Verification process by the Secretariat\*\*

## Note by the Secretariat

1. Pursuant to paragraph 1 of Article 8 of the Stockholm Convention on Persistent Organic Pollutants, a Party which submits a proposal to the Secretariat for listing a chemical in Annex A, B or C of the Convention must provide in the proposal the information specified in Annex D of the Convention (Information Requirements and Screening Criteria).

2. Paragraph 2 of the same Article reads as follows:

"The Secretariat shall verify whether the proposal contains the information specified in Annex D. If the Secretariat is satisfied that the proposal contains the information so specified it shall forward the proposal to the Persistent Organic Pollutants Review Committee."

3. Pursuant to the above, the Secretariat has examined the five proposals which were submitted before 15 August 2005 using the verification process described below. It is important to keep in mind that the verification process is not an evaluation of the rigour or strength of the scientific information provided.

4. The verification process undertaken by the Secretariat involves an examination of the information which is provided in proposals:

(a) *Chemical identity*. Is there a clear identification of the chemical(s) proposed, as described in Annex D, subparagraphs 1 (a) (i) and (ii)?

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<sup>\*</sup> UNEP/POPS/POPRC.1/1.

<sup>\*\*</sup> Stockholm Convention, paragraph 8 and Annex D.

(b) *Persistence*. Is evidence provided on the half-life of the chemical consistent with the criteria listed in Annex D, subparagraph 1 (b) (i) or is some other evidence provided in relation to persistence, as described in Annex D, subparagraph 1 (b) (ii)?

(c) *Bioaccumulation*. Is evidence provided on bioconcentration, bioaccumulation or the logarithmic octanol-water partition coefficient consistent with the criteria set forth in Annex D, subparagraph 1 (c) (i) or is other evidence provided to show that a chemical presents other reasons for concern as described in Annex D, subparagraph 1 (c) (ii), or are monitoring data in biota provided as described in Annex D, subparagraph 1 (c) (iii)?

(d) *Potential for long-range environmental transport*. Are there reports of measured levels of the chemical in locations distant from the source of its release which are of potential concern, as described in Annex D, subparagraph 1 (d) (i), or are monitoring data provided relating to long-range transport, as described in Annex D, subparagraph 1 (d) (ii), or is information given on environmental fate properties or model predictions, as described in Annex D, subparagraph 1 (d) (ii)?

(e) *Adverse effects*. Is evidence provided of direct or potential adverse effects on human health or the environment, as described in Annex D, subparagraphs 1 (e) (i) and (ii) respectively?

(f) *Reasons for concern*. Does the proposal contain a statement of the reasons for concern and the need for global control as described in Annex D, paragraph 2?

(g) *Additional information*. Is additional information provided to support the evaluation of the proposal as described in Annex D, paragraph 3?

5. Where information or the source of information was considered ambiguous or inaccessible, the Secretariat sought clarification from the proponent.

6. The Secretariat assembled for each proposal a verification dossier containing a conclusion as to whether or not the proposal provided the information specified in Annex D. Those dossiers are reproduced in the annex to the present note.

## Annex

Pentabromodiphenyl ether (pentaBDE) proposal (Norway)

1 (a) Chamical idantita	(i) Nomes CAS	The CAS sharring a new second as sisters
1 (a) Chemical identity	(i) Names, CAS number, etc.	The CAS chemical names and registry
		numbers are provided for all major
		components of the commercial mixtures.
		Common and trade names are provided for
		"commercial" pentaBDE.
	(ii) Structure, isomers, etc.	The CAS chemical name, registry number,
		molecular weight and the structural formula
		for the major component of commercial
		pentaBDE (BDE-99) are provided.
		Chemical names and molecular formulae
		are provided for all other components, but
		not the molecular weights.
1 (b) Persistence	(i) Evidence of half-life greater	Persistence in aerobic sediment and water is
	than or	greater than the 6-month and 2-month
		criteria (respectively) for both the major
		components of pentaBDE. Persistence in
		soil is close to the 6-month criterion.
	(ii) Evidence that it is otherwise	Not required (however, evidence is
	sufficiently persistent.	provided that pentaBDE residues have
		persisted decades in marine sediments).
1 (c) Bioaccumulation	(i) Evidence of BCF/BAF	Evidence is provided that the BCF value for
	greater than or	commercial pentaBDE in carp is 27,400,
	6	well above the criterion. No BAF is
		provided. The logKow for all components
		of the commercial mixture is above 5.
	(ii) Evidence of other reasons for	Evidence is provided that tetrabrominated
	concern or	and pentabrominated diphenyl ethers have
		very high bioaccumulation potential and
		that they biomagnify.
	(iii) Monitoring data indicating	Evidence is provided of biomagnification in
	bioaccumulation potential.	Baltic, Atlantic and Arctic species.
1 (d) Potential for long-	(i) Measured levels of concern in	Evidence is provided that increasing
range environmental	distant locations <b>or</b>	concentrations of tetrabromo - and
transport	distant locations of	pentabromodiphenyl ethers have been found
transport		in Arctic whales.
	(ii) Monitoring data showing	Evidence is provided of components of
	transfer may have occurred <b>or</b>	commercial pentaBDE in air samples in
	transfer may have becarred of	remote locations (Arctic).
	(iii) Environmental fate	Vapour pressures for components of
	properties/models demonstrating	commercial pentaBDE range from
	the potential for transport.	$4.7 \times 10^{-5}$ to $9.6 \times 10^{-8}$ Pa.
	the potential for transport.	
		Half-life values in air range from 10 to 20 days.
1 (e) Adverse effects	(i) Evidence of adverse effects	No human or ecosystem evidence is
	or	provided.
	(ii) Toxicity or ecotoxicity data	Evidence of laboratory animal toxicity is
	which indicate potential for	provided (liver toxicity and developmental
	damage.	neurotoxicity, and also immunotoxicity).
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## 2. Statement of concern

A statement of the reasons for concern is provided (see document UNEP/POPS/POPRC.1/5).

#### 3. Additional information

- Environmental Health Criteria (EHC) 162: Brominated Diphenyl Ethers. IPCS, 1994.
- Risk Assessment Report for Diphenyl Ether, Pentabromo Derivative (Pentabromodiphenyl ether), Final Report of August 2000, European Commission, 2000.
- Brominated Flame Retardants. Report 5065 (author, C.A. de Wit), Swedish Environmental Protection Agency, 2000.

Supporting documentation for the pentaBDE proposal also included an extensive 50 pages of "additional information" citing over 100 references.

#### Secretariat evaluation

The proposal identifies the chemical as required under Annex D, subparagraph 1 (a) and provides information on the chemical relating to the screening criteria set forth in Annex D, subparagraphs 1 (b) to (e). It includes a statement of the reasons for concern and the need for global control. Additional information, in the form of a review paper developed for the proposal, was provided. The Secretariat is satisfied that the proposal contains the information specified in Annex D.

1 (a) Chemical identity	(i) Names, CAS number, etc.	The CAS chemical name and registry
		number, synonyms, trade names and
		European Commission registration number
		are provided. The structural formula, molecular formula
	(ii) Structure, isomers, etc.	,
1 (b) Persistence	(i) Evidence of half-life greater	and molecular weight are provided. Persistence in soil is greater than the
1 (b) Persistence	C C	÷
	than or	6-month criterion.
	(ii) Evidence it is otherwise	Not required (however, evidence is provided
	sufficiently persistent.	that it is not expected to
		hydrolyse/biodegrade and that
		photodegradation is not significant).
1 (c) Bioaccumulation	(i) Evidence of BCF/BAF	Evidence that BCF values in three aquatic
	greater than or	species exceed the 5,000 criterion. The
		logKow is between 4.5 and 6.0.
	(ii) Evidence of other reasons for	Not provided.
	concern or	
	(iii) Monitoring data indicating	Not provided.
	bio-accumulation potential.	
1 (d) Potential for long-	(i) Measured levels of concern in	Not provided.
range environmental	distant locations or	
transport	(ii) Monitoring data showing	Atmospheric transport of Chlordecone has
	transfer may have occurred or	been reported.
	(iii) Environmental fate	A vapour pressure of under $3 \times 10^{-5}$ Pa,
	properties/models demonstrating	insignificant photodegradation and a
	the potential for transport	half-life in air of up to 50 years are given as
		evidence that the potential for long-range
		transport is significant.
1 (e) Adverse effects	(i) Evidence of adverse effects	Evidence is provided of high toxicity to
	or	aquatic organisms and some evidence of
		reproductive effects in terrestrial vertebrates.
	(ii) Toxicity or ecotoxicity data	Evidence is provided of laboratory-animal
	which indicate potential for	toxicity following long-term exposure
	damage.	(tremors/neurological signs and liver
		hypertrophy). Evidence is provided of
		carcinogenicity in male and female rats and
		mice.

### Chlordecone proposal (European Union)

#### 2. Statement of concern

A statement of the reasons for concern is provided (see document UNEP/POPS/POPRC.1/6).

### 3. Additional information

Supporting documentation includes two extensive reviews for chlordecone:

- Toxicology profile for mirex and chlordecone, US DHHS, 1995.
- Environmental Health Criteria 43: Chlordecone. IPCS, 1990.

#### Secretariat evaluation

The proposal identifies the chemical as required under Annex D, subparagraph 1 (a) and provides information on the chemical relating to the screening criteria set out in Annex D, subparagraphs 1 (b) to (e). It includes a statement of the reasons for concern and the need for global control. Additional information in the form of a national and an international review was provided. The Secretariat is satisfied that the proposal contains the information specified in Annex D.

1 (a) Chemical identity	(i) Names, CAS number, etc.	The CAS chemical name and registry
		number and trade name for HBB are
		provided.
	(ii) Structure, isomers, etc.	The structural formula, chemical formula
		and molecular weight for HBB are
		provided.
1 (b) Persistence	(i) Evidence of half-life greater	Not provided.
	than <b>or</b>	L
	(ii) Evidence it is otherwise	Evidence is provided that polybrominated
	sufficiently persistent.	biphenyls (PBBs) in general are persistent
	sufficiently persistent.	under field conditions. No significant
		decline in PBB levels were reported in river
		sediment following termination of PBB
		production. Evidence is provided that PBBs
		are resistant to microbial degradation.
1 (c) Bioaccumulation	(i) Evidence of BCF/BAF	Evidence is provided that the BCF value in
	greater than or	minnows exceeds the 5,000 criterion. The
		logKow is between 6.4 and 7.0.
	(ii) Evidence of other reasons for	Not provided.
	concern or	
	(iii) Monitoring data indicating	Not provided.
	bioaccumulation potential	
1 (d) Potential for long-	(i) Measured levels of concern in	Evidence is provided that PBBs have been
range environmental	distant locations or	reported in Arctic mammals.
transport	(ii) Monitoring data showing	Not provided.
-	transfer may have occurred or	*
	(iii) Environmental fate	HBB has a vapour pressure of $6.9 \times 10^{-9}$ Pa.
	properties/models demonstrating	Specific information on its
	the potential for transport.	photodecomposition in air is not reported.
		Evidence of PBB in the Arctic is suggestive
		of transport and persistence of HBB.
1 (a) Advance offects	(i) Evidence of advance offects	
1 (e) Adverse effects	(i) Evidence of adverse effects	No evidence is provided of adverse effects
	or	on human health or on the environment.
		IARC considers HBB a possible human
		carcinogen based on laboratory animal
		studies.
	(ii) Toxicity or ecotoxicity data	Evidence of laboratory animal toxicity and
	which indicate potential for	carcinogenicity following long-term
	damage.	exposure to PBBs.

### Hexabromobiphenyl (HBB) proposal (European Union)

#### 2. Statement of concern

A statement of the reasons for concern is provided (see document UNEP/POPS/POPRC.1/7)

#### 3. Additional information

Supporting documentation includes one extensive review for polybrominated biphenyls:
Environmental Health Criteria 152: Polybrominated biphenyls, IPCS, 1994

#### Secretariat evaluation

The proposal identifies the chemical as required under Annex D, subparagraph 1 (a) and provides information on the chemical relating to the screening criteria set forth in Annex D, subparagraphs 1 (b) to (e). It includes a statement of the reasons for concern and the need for global control. Additional information, in the form of an international review, was provided. The Secretariat is satisfied that the proposal contains the information specified in Annex D.

1 (a) Chemical identity	(i) Names, CAS number, etc.	The CAS chemical name, CAS registry number and common/trade names are
		provided.
	(ii) Structure, isomers, etc.	The structural formula for
		hexachlorocyclohexane and its gamma
		isomer (Lindane), molecular formula and
		molecular weight are provided.
1 (b) Persistence	(i) Evidence of half-life greater	Persistence in water is greater than the
	than <b>or</b>	2-month criterion and persistence in soil is greater than the 6-month criterion.
	(ii) Evidence it is otherwise	Not required (however, evidence is provided
	sufficiently persistent.	that it is not readily degraded by light and
		degraded very slowly by microbial action).
1 (c) Bioaccumulation	(i) Evidence of BCF/BAF	Evidence is provided that the log BCF value
	greater than or	can be as high as 3.85, the log BAF is
		4.1 and the logKow is 3.5. Those values do
		not meet the criteria set in Annex D,
		subparagraph 1 (c) (i).
	(ii) Evidence of other reasons for	Evidence is provided that Lindane is
	concern or	considered highly toxic to some aquatic
		species. Its beta isomer accumulates more
		rapidly and bioconcentrates to higher levels
		in the environment.
	(iii) Monitoring data indicating	Evidence is provided that Lindane has been
	bio-accumulation potential.	found accumulating in aquatic species and
		marine mammals far from its site of
		manufacture and use.
1 (d) Potential for long-	(i) Measured levels of concern in	Evidence is provided that Lindane has been
range environmental	distant locations or	found accumulating in aquatic species and marine mammals far from its site of
transport		manufacture and use.
	(ii) Monitoring data showing	Evidence is provided that Lindane is
	transfer may have occurred <b>or</b>	routinely measurable in Arctic air, ice pack,
	transfer may have becarred of	sea water and freshwater.
	(iii) Environmental fate	A vapour pressure of $3.8 \times 10^{-3}$ Pa,
	properties/models demonstrating	insignificant photodegradation and a
	the potential for transport.	half-life in air of 2.3 to 13 days are given as
	I CONTRACTOR	evidence that the potential for long-range
		transport is significant. Estimated annual
		airborne deposition in the Arctic is
		13,000 kg.
1 (e) Adverse effects	(i) Evidence of adverse effects	Evidence is provided of high toxicity to
	or	some aquatic organisms.
	(ii) Toxicity or ecotoxicity data	Evidence is provided of laboratory-animal
	which indicate potential for	toxicity following acute exposures (central
	damage.	nervous system signs, convulsions,
		respiratory failure, pulmonary oedema and
		dermatitis) and long-term exposure
		(convulsions, effects on liver, kidney and
		reproductive organs, impaired immune
		system). Evidence is provided of
		carcinogenicity in mice.

Lindane (gamma hexachlorocyclohexane) pro	oposal (Mexico)
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#### 2. Statement of concern

A statement of the reasons for concern is provided (see document UNEP/POPS/POPRC.1/8).

#### 3. Additional information

Supporting documentation includes three reviews for Lindane:

- National Diagnostic Report on Lindane, Mexican National Institute of Ecology (INE), 2004. (www.ine.gob.mx/dgicurg/download/Proyectos-2003/EL\_LINDANO\_EN\_MEXICO.pdf)
- Technical Review Report on Lindane, UNECE, 2004 (www.unece.org/env/popsxg/docs/2004/Dossier\_Lindane.pdf)
- (Draft) Decision Document on Lindane, Commission for Environmental Cooperation, 2000. (www.cec.org/files/pdf/POLLUTANTS/linddd\_en.pdf)

An additional review:

Toxicological Profiles for Hexachlorocyclohexanes (HCH), US-ATSDR, 2003 (www.atsdr.cdc.gov/toxprofiles/tp43.html)

#### Secretariat evaluation

The proposal identifies the chemical as required by Annex D, subparagraph 1 (a) and provides information on the chemical relating to the screening criteria set out in Annex D, subparagraphs 1 (b) to (e). It includes a statement of the reasons for concern and the need for global control. Additional information, in the form of national and international reviews, was provided. Other international and domestic reviews are available. The Secretariat is satisfied that the proposal contains the information specified in Annex D.

1 (a) Chamicalide 4'	Namaa CAC 1	
1 (a) Chemical identity (i)	Names, CAS number, etc.	The CAS chemical name, synonyms and trade names for PEOS are provided. As a
		trade names for PFOS are provided. As a fully fluoring and anion PEOS does not have
		fully fluorinated anion, PFOS does not have a CAS number. The proposal covers a list of
		96 PFOS-related substances for which CAS
		numbers are provided along with their
		chemical names.
(ii)	) Structure, isomers, etc.	The structural formula for the PFOS anion
(II)	structure, isomers, etc.	is provided, and also for the related family
		of perfluoroalkyl sulfonates (PFAS).
<b>1 (b) Persistence</b> (i)	Evidence of half-life greater	Persistence measured as half-life in water
	an or	and under light was 41 years and 3.7 years
		respectively. No biodegradation was found
		in anaerobic and aerobic tests.
(ii)	) Evidence it is otherwise	Evidence is provided of high levels of
	fficiently persistent.	PFOS in many species, including several in
		locations remote from the site(s) of
		manufacture and use.
<b>1 (c) Bioaccumulation</b> (i)	Evidence of BCF/BAF	PFOS is both lipophobic and hydrophobic
gre	eater than or	and accumulates bound to protein, not fat.
-		Its BCF in sunfish (whole body) is
		2,796 and in trout (plasma) 3,100: neither
		value exceeds the Annex D criteria. The
		logKow for PFOS is not measurable.
(ii)	) Evidence of other reasons for	Evidence is provided that the
CO	ncern <b>or</b>	biomagnification potential as measured in
		top-of-the-food-chain species is very large
		(e.g., from seals to polar bears, the BMF is
		greater than 160).
	i) Monitoring data indicating	Evidence is provided of extensive
bio	paccumulation potential.	monitoring in a wide range of species.
		Levels in a variety of aquatic species and
		birds are reported to reach mg/kg
	M 11 1 C '	bodyweight levels.
. , , , , , , , , , , , , , , , , , , ,	Measured levels of concern in stant locations <b>or</b>	Evidence is provided that there is
transport	stant locations or	widespread contamination of species remote from the sites of manufacture or use.
-	) Monitoring data showing	Not provided. Assumed to be transported
	nsfer may have occurred or	over long ranges bound to particles.
	) Environmental fate	Vapour pressure for the potassium salt of
	operties/models demonstrating	PFOS is $3.31 \times 10^{-4}$ Pa. Evidence is
	e potential for transport.	provided that other PFOS-containing
	potential for transport.	substances have considerably higher vapour
		pressures. Half-life values in air probably
		exceed 2 days.
1 (e) Adverse effects (i)	Evidence of adverse effects	No human or ecosystem evidence is
or		provided.
	) Toxicity or ecotoxicity data	Evidence of laboratory-animal toxicity in
	nich indicate potential for	monkeys and rats (gastrointestinal lesions
	mage.	and organ weight loss, infant mortality,
	-	inhibition of lung maturation). No-effect
		minoriton of fung maturation). No-effect
		levels were in the submilligram range.
		levels were in the submilligram range.

## Perfluorooctane sulfonate (PFOS) proposal (Sweden)

#### 2. Statement of concern

A statement of the reasons for concern is provided (see document UNEP/POPS/POPRC.1/9).

#### 3. Additional information

Supporting documentation for PFOS and compounds which contain PFOS is provided in a report submitted together with the proposal. The report provides information on recent environmental levels, PFOS-containing compounds, production/use/emissions data, socio-economic factors, etc. The supporting documentation mentioned includes :

- Hazard Assessment of Perfluorooctane Sulfonate and its Salts, OECD, 2002.
- Perfluorooctane Sulfonate: Risk Assessment Strategy and Analysis of Advantages and Drawbacks, United Kingdom, 2004.
- Environmental Risk Evaluation Report: Perfluorooctane sulfonate (PFOS), UK, 2004.

#### Secretariat evaluation

The proposal identifies the chemical and related compounds as required under Annex D, subparagraph 1 (a) and provides information relating to the screening criteria set forth in Annex D, subparagraphs 1 (b) to (e). It includes a statement of the reasons for concern and the need for global control. Additional information, in the form of a review paper developed for the proposal and other national and international supporting documents, was provided. The Secretariat is satisfied that the proposal contains the information specified in Annex D.

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