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**Conference of the Parties to the Basel Convention
on the Control of Transboundary Movements of
Hazardous Wastes and Their Disposal
Thirteenth meeting**

Geneva, 24 April–5 May 2017

Agenda item 4 (b) (i)

**Matters related to the implementation of the Convention:
scientific and technical matters: technical guidelines**

Technical guidelines

Addendum

**Technical guidelines on the environmentally sound management of wastes consisting of, containing or contaminated with polychlorinated biphenyls, polychlorinated terphenyls, polychlorinated naphthalenes or polybrominated biphenyls including hexabromobiphenyl**

 Note by the Secretariat

 At its thirteenth meeting, the Conference of the Parties to the Basel Convention on the Control of Transboundary Movements of Hazardous Wastes and Their Disposal adopted, in decision BC‑13/4 on technical guidelines on the environmentally sound management of wastes consisting of, containing or contaminated with persistent organic pollutants, the technical guidelines on the environmentally sound management of wastes consisting of, containing or contaminated with polychlorinated biphenyls, polychlorinated terphenyls, polychlorinated naphthalenes or polybrominated biphenyls including hexabromobiphenyl, on the basis of the draft technical guidelines contained in document UNEP/CHW.13/6/Add.4. The technical guidelines referred to above were prepared by Japan, in consultation with the small intersessional working group on the development of technical guidelines on persistent organic pollutants wastes and taking into account comments received from Parties and others and comments provided at the tenth meeting of the Open‑ended Working Group of the Basel Convention. The technical guidelines were further revised on 1 March 2017 taking into account the outcome of the face-to-face meeting of the small intersessional working group on the development of technical guidelines on persistent organic pollutants wastes held from 20 to 22 February 2017 in Bonn, Germany (see document UNEP/CHW.13/INF/63). The text of the final version of the technical guidelines, as adopted, is set out in the annex to the present note. The present note, including its annex, has not been formally edited.

Annex

Technical guidelines on the environmentally sound management of wastes consisting of, containing or contaminated with polychlorinated biphenyls, polychlorinated terphenyls, polychlorinated naphthalenes or polybrominated biphenyls including hexabromobiphenyl

 **Revised final version (5 May 2017)**

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# **Abbreviations and acronyms**

|  |  |
| --- | --- |
| ABS | acrylonitrile-butadiene-styrene copolymers (plastics) |
| ABNT | Associação Brasileira de Normas Técnicas (Brazilian National Standards Organization) |
| AMAP | Arctic Monitoring and Assessment Programme |
| AOACASTM | Association of Official Agricultural Chemists (United States of America)American Society for Testing and Materials |
| ATSDRCASDIN | Agency for Toxic Substances and Disease Registry (United States of America)Chemical Abstracts ServiceDeutsches Institut für Normung e.V. (German Institute for Standardization) |
| ENEPA | European StandardsEnvironmental Protection Agency (United States of America) |
| ESM HBB | environmentally sound managementhexabromobiphenyl |
| HCB  | hexachlorobenzene  |
| IARC | International Agency for Research on Cancer |
| IPCS | International Programme on Chemical Safety (of WHO) |
| ISO | International Organization for Standardization |
| JIS | Japanese Industrial Standards |
| NEN | The Netherlands Standardization Institute |
| NVN | Dutch standards |
| OEWG | Open-ended Working Group of the Basel Convention |
| PBB  | polybrominated biphenyl |
| PBDD | polybrominated dibenzo-*p*-dioxin |
| PBDF | polybrominated dibenzofuran |
| PCB  | polychlorinated biphenyl |
| PCDD | polychlorinated dibenzo-*p*-dioxin |
| PCDF | polychlorinated dibenzofuran |
| PCN  | polychlorinated naphthalene |
| PCT  | polychlorinated terphenyl |
| PeCB | Pentachlorobenzene |
| POP  | persistent organic pollutant |
| POPRC | Persistent Organic Pollutants Review Committee |
| TEF  | toxic equivalency factor  |
| UNECE | United Nations Economic Commission for Europe |
| UNEP | United Nations Environment Programme |
| WHO | World Health Organization |

# **Units of measurement**

|  |  |
| --- | --- |
| mg  | Milligram |
| kg  | Kilogram |
| Mg | megagram (1,000 kg or 1 tonne)  |
| mg/kg  | milligram (s) per kilogram. Corresponds to parts per million (ppm) by mass. |

# **I. Introduction**

## A. Scope

1. The present document supersedes the *Updated technical guidelines for the environmentally sound management of wastes consisting of, containing or contaminated with polychlorinated biphenyls (PCBs), polychlorinated terphenyls (PCTs) or polybrominated biphenyls (PBBs)* *including* *hexabromobiphenyl (HBB)* of May 2015.
2. The present technical guidelines provide guidance on the environmentally sound management (ESM) of wastes consisting of, containing or contaminated with polychlorinated biphenyls (PCBs), polychlorinated terphenyls (PCTs), polychlorinated naphthalenes (PCNs) or polybrominated biphenyls (PBBs) including hexabromobiphenyl (HBB), pursuant to several decisions of two multilateral environmental agreements on chemicals and wastes.[[1]](#footnote-2) PCB was listed in Annex A (elimination) to the Stockholm Convention at the time of its adoption. HBB was listed in Annex A to the Stockholm Convention in 2009, through the adoption of an amendment that entered into force in 2010. PCNs, including dichlorinated naphthalenes (di-CNs), trichlorinated naphthalenes (tri-CNs), tetrachlorinated naphthalenes (tetra-CNs), pentachlorinated naphthalenes (penta-CNs), hexachlorinated naphthalenes (hexa-CNs), heptachlorinated naphthalenes (hepta-CNs) and octachlorinated naphthalene (octa-CN), were listed in Annex A of the Stockholm Convention in 2015 through the adoption of an amendment that entered into force in 2016.
3. The present technical guidelines address PCBs, PCNs and HBB together with PCTs and PBBs other than HBB as a class or category of substances, owing to similarities in the physico-chemical and toxicological properties of all of these substances. Among other topics, the guidelines address all activities pertaining to waste management. It should be noted that PCTs and PBBs other than HBB are not currently subject to the Stockholm Convention. It should also be noted that monochlorinated naphthalenes (mono-CNs) are not subject to the Stockholm Convention.
4. Unintentionally produced PCBs and PCNs listed in Annex C to the Stockholm Convention (unitentional production) are not covered by the present technical guidelines. They are covered instead by the *Technical guidelines on the environmentally sound management of wastes containing or contaminated with unintentionally produced polychlorinated dibenzo-p-dioxins (PCDDs), polychlorinated dibenzofurans (PCDFs), hexachlorobenzene (HCB), polychlorinated biphenyls (PCBs), pentachlorobenzene (PeCB) or polychlorinated naphthalenes (PCNs)* (Unintentional POPs technical guidelines) (UNEP, 2017a).
5. The present document should be used in conjunction with the *General technical guidelines on the environmentally sound management of wastes consisting of, containing or contaminated with persistent organic pollutants”* (UNEP, 2017b) (hereinafter referred to as “General technical guidelines”). The General technical guidelines are intended to serve as an umbrella guide for the ESM of wastes consisting of, containing or contaminated with persistent organic pollutants (POPs).

## B. Description, production, use and wastes

### 1. Description

#### (a) PCBs

1. PCBs are synthetic aromatic compounds formed in such a manner that the hydrogen atoms on the biphenyl molecule (two benzene rings bonded together by a single carbon-carbon bond) may be replaced by up to 10 chlorine atoms. The basic chemical structure of PCBs is shown in figure 1 below; the general molecular formula for PCBs is C12H10-nCln, where n=1-10 (CAS No. 1336-36-3). In theory there are 209 congeners, although only about 130 congeners have actually been found in commercial chemical formulations (Holoubek, 2000). Typically, four to six of the 10 possible substitution sites are occupied by a chlorine atom (Environment Canada, 1988). In case of dielectric fluids, PCB mixtures mainly containing either tri-, tetra-, or pentachlorinated homologues are used. The physical properties are, for instance, regarding trade named Aroclor 1254 which is one of the popular commercial PCB products consisting most of pentachlorobiphenyls, the boiling point is of 365°C -390°C, the specific gravity (at 25°C) is 1.54 g/cm,3 the vapour pressure (at 25°C) is 0.010 Pa, the water solubility (at 24°C) is 0.057 mg/L, and it is viscous liquid at an ordinary temperature (ATSDR, 2000). The more highly chlorinated PCB congeners are virtually insoluble in water and highly resistant to degradation.

**Figure 1:** Chemical structure of PCBs



1. Since PCBs are stable to heat and biodegradation, once released into the environment they are persistent and accumulate in the organic components of soils, sediments, biological tissues and organic carbons dissolved in aquatic systems, thereby entering the ecological food chain. PCBs especially accumulate in fish and marine mammals, reaching levels that may be many thousands of times higher than in water. The general population may be exposed to PCBs by ingesting contaminated food and by inhaling contaminated air. PCBs are transported from soil and sediment to the atmosphere and can easily cycle between air, water and soil and enter the air by evaporation from both soil and water. In air, PCBs can be carried long distances and have been found in snow and seawater in areas far away from where they were released, such as the Arctic (ATSDR, 2000).
2. PCBs include 12 congeners to which the World Health Organization (WHO) has assigned toxicity equivalency factors (TEFs) because they exhibit dioxin-like toxicity (Van den Berg et al, 2006).
3. PCBs, including the abovementioned 12 dioxin-like congeners, have been classified as carcinogenic to humans (Group 1) by the International Agency for Research on Cancer (IARC, 2014).

#### (b) PCTs

1. PCTs also constitute a group of halogenated hydrocarbons. They are very similar to PCBs in terms of chemical structure, except that they contain three phenyl rings instead of two and therefore can have up to 14 chlorine atoms attached. The number of possible PCT congeners is large; however, only a few occur in commercial chemical formulations. PCTs and PCBs have similar chemical and physical properties. PCTs are virtually insoluble in water and highly resistant to degradation. One difference between PCTs and PCBs is that PCTs are generally less volatile. The basic chemical structure of PCTs is shown in figure 2 below; PCTs have the general molecular formula C18H14-nCln, where n=1-14 (CAS No: 61788-33-8).

**Figure 2:** Chemical structure of PCTs



#### (c) PCNs

1. PCNs are a group of compounds based on the naphthalene ring system, but where one or more hydrogen atoms have been replaced by chlorine. The basic structure of the PCNs is shown in figure 3 below and has the molecular formula C10H8-nCln, where n=1-8 (CAS no: 70776-03-3). PCNs comprise of 75 possible congeners in eight homologue groups with one to eight chlorine atoms substituted around the planar aromatic naphthalene molecule. Almost all congeners have actually been found in commercial formulations (Noma et al. 2004). All 75 PCN congeners and 8 homologues have different CAS numbers. The system of nomenclature for PCN is similar to that of PCB. The homologue groups are mono-CNs, di-CNs, tri-CNs, tetra-CNs, penta-CNs, hexa-CNs, hepta-CNs and octa-CN. However, mono-CNs are not subject to the Stockholm Convention. Physical-chemical properties vary considerably due to the degree of chlorine substitution. The physical state ranges from thin liquids to hard waxes. Tri- through octa-CNs are very lipophilic with high log Kow (>5) and their water solubility and vapour pressure decrease with the degree of chlorination. Mono-CNs and di-CNs are slightly soluble in water while the higher CNs have a water solubility of a few µg/L. For example, Halowax 1001 (the trade name of PCN mixture) consisting most of tetra-CNs and penta-CNs has a boiling point of 308 °C, a melting point of 98 °C and physical form of flakes, while Halowax 1000 consisting most of mono-CNs and di-CNs has a boiling point of 250 °C, a melting point of 33 °C and physical form of liquid (US EPA, 1975; Brinkman et al. 1976; Crookes et al. 1993; IPCS, 2001).

**Figure 3:** The structural formula of PCNs



\* For PCNs listed as POPs (x + y) varies from 2 to 8

1. Although two publications on the WHO TEF schemes recommend to include certain PCN, no TEFs have been proposed through the WHO experts so far (van den Berg et al., 2006; van den Berg et al., 2013).

####  (d) PBBs

1. PBBs are the bromine analogues of PCBs and thus there are 209 possible PBB congeners. Only a few, however, occur in commercial chemical formulations (IPCS, 1994). They are solids or waxy substances at room temperature. They are virtually insoluble in water and highly resistant to degradation. The basic chemical structure of PBBs is shown in figure 4 below; PBBs have the general molecular formula C12H10-nBrn, where n=1-10.

**Figure 4:** Structural formula of PBBs



1. HBB belongs to a wider group of PBBs. Hexabrominated congeners possibly exist in 42 isomeric forms, which are listed under a variety of CAS numbers, e.g., CAS No. 36355-01-8 for all HBB isomers and CAS No. 59080-40-9 for 2,2’,4,4’,5,5’-HBB. HBB is white and solid at normal temperatures and has a vapour pressure of 6.9x10-6 Pa and a melting point of 72°C (ATSDR, 2004).
2. Some PBBs have been assigned toxicity equivalency factors similar to those of PCBs (Van den Berg et al., 2013).
3. PBBs have been classified by IARC as probably carcinogenic to humans (Group 2A) (IARC, 2014).

### 2. Production

#### (a) PCBs

1. PCBs have excellent dielectric properties, longevity, non-flammability and resistance to thermal and chemical degradation. For this reason, prior to national bans they were manufactured for use in electrical equipment, heat exchangers, hydraulic systems and several other specialized applications.
2. The main period of manufacture of PCBs was from 1930 to late 1977 in the United States of America, up to 1983 in China, up to the mid-1980s in Europe, up to 1993 in the Russian Federation and from 1954 to 1972 in Japan.[[2]](#footnote-3)
3. Chlorination of PCBs was conducted continuously until a certain target percentage of chlorine based on weight was achieved. Manufactured PCBs were used as insulating oils and as heat media. Electric equipment can contain notably high concentrations of PCBs. For example, capacitors may be filled with up to 100 per cent PCBs and transformers with approximately 60-90 per cent PCBs (Ivanov et al., 1992; Kukharchyk et al., 2008). In addition, PCBs were added in small quantities to inks, plastics, paints, sealants, adhesives and dye solvents for carbonless paper. At room temperature, most of the PCBs added to these products were oily liquids or waxy solids.
4. Prominent trade names of PCB products include those listed below (see annex I to the present guidelines for a more detailed list of PCB trade names and synonyms, and section IV.D for a discussion of trade names in inventory identification):

(a) Apirolio (Italy);

(b) Aroclor (United States of America and United Kingdom);

(c) Askarel (United States and United Kingdom);

(d) Clophen (Germany);

(e) Delor (Former Czechoslovakia);

(f) Elaol (Germany);

(g) Fenchlor (Italy);

(h) Inerteen (United States);

(i) Kanechlor (Japan);

(j) Phenoclor (France);

(k) Pyralene (France);

(l) Pyranol (United States);

(m) Pyroclor (United States and United Kingdom);

(n) Santotherm (Japan);

(o) Sovol (former Union of Soviet Socialist Republics (USSR));

(p) Sovtol (former USSR);

(q) Trichlorodiphenyls (former USSR).

1. In the Aroclor series, a four-digit number follows the word Aroclor. The first two digits of the number are either 10 or 12. The number 12 indicates a normal Aroclor while the number 10 indicates a distillation product of an Aroclor. The second two digits of the four-digit code indicate the percentage of chlorine in the mixture by weight. Therefore, Aroclor 1254 contains approximately 54 per cent chlorine by weight.
2. Commercial PCB products and articles were sold for their industrial properties rather than for their chemical composition (IPCS, 1992). They contained a number of impurities and were often mixed with solvents, such as tri- and tetrachlorobenzenes. PCBs mixed with tri- and tetrachlorobenzenes were called Askarel. Contaminants in commercial mixtures include PCDFs and PCNs. Studies have found from 0.8 mg/kg to 40 mg/kg of PCDFs in PCB commercial mixtures (IPCS, 1992). PCBs are also formed unintentionally in some thermal and chemical processes.
3. The cumulative worldwide production of PCBs has been estimated to be 1-1.5 million tonnes.

#### (b) PCTs

1. PCTs were manufactured in much smaller quantities than PCBs and were given the same or similar trade names. They were used for the same sorts of applications as PCBs, although most were used in waxes, plastics, hydraulic fluids, paints and adhesives (Jensen and Jørgensen, 1983).
2. Prominent trade names of PCT products are Aroclor (United States) and Kanechlor KC-C (Japan). Aroclor series PCTs were identified by the digits 54 in the first two spaces of the four-digit code, e.g., Aroclor 5432, 5442 and 5460 (IPCS, 1992) (see annex I to the present guidelines for a more detailed list of PCT trade names and synonyms, and section IV.D for a discussion of trade names in inventory identification).
3. PCTs were produced in the United States, France, Germany, Italy and Japan until the early 1980s, when all production is thought to have ceased. The cumulative world production is estimated to have been 60,000 tonnes between 1955 and 1980 (UNECE, 2002).

#### (c) PCNs

1. Parties to the Stockholm Convention must prohibit and/or eliminate the production of PCNs. The specific exemption for production of PCNs is intermediates in production of polyfluorinated naphthalenes, including octafluoronaphthalene (see paragraph 65).
2. PCNs were produced for technical use in the first decade of the 20th century, e.g. as dielectrics, for flame proofing or for the protection of paper and fabrics – including for paper inlays for gas masks in World War I. After the Second World War, their production began to decline with the advent of plastic substances for insulation and the use of PCBs for dielectrics in transformers, partly due to severe occupational exposure problems. Still, PCNs remained high volume chemicals until the 1970s. Their production decreased significantly since 1977. (US EPA, 1975; Brinkman et al. 1976; Crookes et al., 1993; Falandysz, 1998; Hayward, 1998; Plassche et al., 2003; AMAP, 2004).
3. The production data are limited, and the original data on the quantum of PCNs produced worldwide are not available.
4. Estimates for the total global PCNs production so far vary between 200,000–400,000 tonnes (AMAP, 2004) and 150,000 tonnes (about one tenth of the PCBs ever produced (Falandysz, 1998). In the United States of America, approximately a total of between 50,000 and 150,000 tonnes of PCNs have been produced from 1910 until 1960. The production of PCNs in the United States of America stopped in 1980. The production of technical PCNs by Koppers Company, main producer of Halowax, ceased in 1977, and the last producer of the United States of America of PCNs (Chemispheres Company) stopped manufacture in 1980 (US EPA, 1975; US EPA, 1983). In Germany, the production of PCNs started around 1910 and ceased in the mid-1980s. A cumulative total of around 75,000 tonnes of PCNs were manufactured up to 1972 and around 300 tonnes were produced in 1984. Bayer produced PCNs (Nibren waxes) in a range of 100 to 200 tonnes per year between 1980 and 1983 and ceased PCNs production in 1983 (IPCS, 2001; Plassche et al., 2003). In France, small amounts of PCNs were produced by Prodelec (Brinkman et al., 1976). In the United Kingdom, PCNs were produced from around 1919 until the mid-1960s. No data as to the amounts produced appear to be available (Crookes et al., 1993). In Japan, the production of PCNs from 1940 to 1976 was of approximately 4,000 tonnes, and manufacture was banned from 1979 (Japan Ministry of Economy, Trade and Industry, 1979). In China, there is no information available about the production (Pan et al., 2011), however a small quantity production (not specified) of octa-CN for scientific purposes was reported in Jiangsu province (UNEP, 2012). In the former USSR, PCNs or Halowax was produced by Chimprom enterprises, while volumes were not assessed or reported. To date, intentional production of PCNs is assumed to have ended. PCNs are unintentionally generated in some thermal and chemical processes (UNEP, 2012).
5. Commercial PCBs also contained traces of PCNs (0.01–0.09%) (Falandysz, 1998; Kannan et al., 2000; Yamashita et al., 2000). Studies have found the impurity PCDDs (1.5 to 370 ng/g), PCDFs (250 to 16,000 ng/g), and PCBs (220 to 640,000 ng/g) in all Halowax formulations investigated (Noma et al., 2005, 2005a).
6. Prominent trade names of PCNs products include those listed below (see annex I for a more detailed list of PCNs trade names and synonyms and section IV.D for a discussion of trade names in inventory identification) (Nomura, 1951; US EPA, 1975; Brinkman et al., 1976; Crookes et al., 1993; Plassche et al., 2003; Falandysz, 2008):

(a) Basileum (Germany);

(b) Cerifal (Italy);

(c) Chlonacire wax (France);

(d) Halowax (United States of America)[[3]](#footnote-4);

(e) Hodogaya Amber wax (Japan);

(f) Nankai wax (Japan);

(g) Nibren wax (Germany);

(h) N-Oil, N-Wax (United States of America);

(i) Perna wax (Germany);

(j) Seekay wax (United Kingdom);

(k) Tokyo Ohka wax (Japan);

(l) Woskol: Zakady Azotowe (Poland).

#### (d) PBBs

1. PBBs exhibit unusual chemical stability and are stable in acids, bases, heat, and reducing and oxidizing agents. However, in chemical reactions, bromine is better than chlorine as a leaving group (IPCS, 1994). For this reason, PBBs were manufactured mainly for use as flame retardants.
2. It is estimated that at least 11,000 tonnes of PBBs were produced worldwide, but the amounts of production of some PBB-producing countries are not available (IPCS, 1994). In the United States, the commercial production of PBBs began in 1970, and about 6,000 tonnes were produced from 1970 to 1976. The first PBB compound produced in the United States was HBB, but its production was discontinued in 1975. HBB was commercially called FireMaster in the United States and its production constituted about 88 per cent of total PBB production (ATSDR, 2004). PBBs were also manufactured in the United Kingdom until 1977, and in Germany until the mid-1980s. Although PBBs were never produced in Japan, they were imported into that country until 1978. It has been reported that PBB production ended worldwide with the cessation of decabromobiphenyl production in France in 2000 (UNEP, 2006).
3. PBBs produced for commercial uses include mixtures of various brominated biphenyls, mainly containing HBB and octa-, nona- and decabromobiphenyls, as well as other PBB congeners (IPCS, 1994). All commercial PBB mixtures were relatively highly brominated, with bromine contents ranging from about 76 per cent HBB to 81-85 per cent octa- to decabromobiphenyl mixtures (IPCS, 1994; IARC, 2014).
4. Prominent trade names of PBB products include those listed in table 1 below (see annex I to the present guidelines for a more detailed list of PBB trade names and synonyms, and section IV.D for a discussion of trade names in inventory identification).

**Table 1:** Main constituents, trade names and country of origin[[4]](#footnote-5)

| **Main PBB congener** | **Trade name** | **Country in which the chemical has been produced** |
| --- | --- | --- |
| Hexabromobiphenyls | FireMaster FF-1 | United States  |
| FireMaster BP-6 | United States  |
| Octabromobiphenyls | BB-8 |  |
| Bromkal 80 | Germany |
| Bromkal 80–9D | Germany |
| Octabromobiphenyl FR 250 13A | United States  |
| Technical octabromobiphenyl | United States  |
| Decabromobiphenyl | Adine 0102 | France |
| Berkflam B-10 | United Kingdom  |
| Flammex B-10 | United Kingdom  |
| HFO 101 | United Kingdom  |
| Technical decabromobiphenyl | United States  |

### 3. Use

#### (a) PCBs

1. PCBs were used in a very wide variety of industrial and consumer applications. Such applications have been categorized by WHO as completely closed, nominally closed, and open-ended (IPCS, 1992), and included the following:

(a) Completely closed systems:

(i) Electrical transformers;

(ii) Electrical capacitors (including lamp ballasts);

(iii) Electrical switches, relays, breakers, reclosers and others;

(iv) Electrical cables;

(v) Electrical bushings;

(vi) Electrical reactors;

(vii) Electrical regulators;

(viii) Electric motors and magnets (very small amounts);

(b) Nominally closed systems:

(i) Hydraulic systems;

(ii) Heat transfer systems (heaters, heat exchangers);

(iii) Vacuum pumps;

(iv) Vapour diffusion pumps;

(c) Open-ended systems:

(i) Plasticizer in polyvinyl chloride, neoprene and other artificial rubbers;

(ii) Ingredient in paint and other coatings;

(iii) Ingredient in ink and carbonless copy paper;

(iv) Ingredient in adhesives;

(v) Pesticide extender;

(vi) Ingredient in sealants and caulking material;

(vii) Fire retardant in fabrics, carpets, polyurethane foam, etc.;

(viii) Lubricants (microscope oils, brake linings, cutting oils, bridge bearings, other lubricants).

1. Although electrical transformers containing PCBs are defined as a “completely closed” application, industrial practices caused these PCBs to be transferred to other types of equipment, thus creating additional points of contact with the environment. A common practice was to top up or recharge non‑PCB (mineral oil) transformers with PCBs when no other fluid was available.
2. PCB oils were also added to or disposed of with non-PCB fluids such as heating or cooling fluids, hydraulic fluids, brake fluids, engine oils and off-specification fuels. There are numerous anecdotal reports of employees in electrical utilities using PCB fluids to wash their hands and taking PCB fluids home for use as lubricants in home heaters, hydraulic systems and motors. Since most of the fluorescent lamp ballasts that were made before PCBs were banned contained PCBs, many homes and businesses that installed fluorescent lamps unknowingly acquired PCBs.

#### (b) PCTs

1. PCTs were used in almost exactly the same applications as PCBs but in much smaller amounts. Little is known, however, about remaining quantities of PCTs because PCT inventories have not been developed (UNECE, 2002). It is known that very small amounts of PCTs were used in electrical equipment (Jensen and Jørgensen, 1983).

#### (c) PCNs

1. Parties to the Stockholm Convention must prohibit and/or eliminate the use of PCNs. The specific exemption for use of PCNs is production of polyfluorinated naphthalenes, including octafluoronaphthalene (see paragraph 65).
2. PCNs have been used mainly for their chemical inertness, including low flammability, their electrically insulating properties and recalcitrance including resistance to biodegradation and biocidal function; they share these properties and their scope of application with the PCBs (Hayward, 1998).
3. PCNs were used for the same applications as PCBs with a wide variety because of their structural similarity. The uses were industrial and consumer applications in completely closed, nominally closed and open-ended systems similar to PCBs. PCNs were gradually substituted by PCBs in many applications. Major uses are shown below (US EPA, 1975; Brinkman et al., 1976; US EPA, 1983; Crookes et al., 1993; IPCS, 2001; UNEP, 2012).
4. Mono-CNs and mixtures of mono- and di-CNs have been used for chemical-resistant gauge fluids and instrument seals, as heat exchange fluids, as high boiling specialty solvents, for colour dispersions, as engine crankcase additives, and as ingredients in motor tune-up compounds. Mono-CNs have also been used as a raw material for dyes and as a wood preservative with fungicidal and insecticidal properties.
5. The tri- and higher chlorinated naphthalene products have been used as impregnants for condensers and capacitors and dipping encapsulating compounds in electronic and automotive applications, as temporary binders in paper coating and impregnation, binders for ceramic components, in precision casting of alloys, in electroplating stop-off compounds, as additives in gear oils and cutting compounds, in flame-proofing and insulation of electrical cable and conductors, as moisture-proof sealants and caulking material, as separators in batteries, in refractive index testing oils, as masking compounds in electroplating, in grinding wheel lubricants, and ingredient in paint and other coatings.
6. The most important uses, in terms of volume, have been cable insulation and flame-proofing, wood preservation, engine and gear oil additives, electroplating masking compounds, feedstock for dye production, dye carriers, dielectric impregnates for capacitors/condensers, and refractive index testing oils. The use of PCNs as wood preservatives was popular in the 1940s and 1950s, but they are no longer used for this purpose in the United States of America. The US Environmental Protection Agency stated that only very small amounts of PCNs (about 15 tons/year in 1981) were still being used in the United States of America, mainly as refractive index testing oils and as capacitor dielectrics. It did note that the most likely possible new uses for PCNs would be as intermediates for polymers and as flame retardants in plastics. (Crookes et al., 1993).

#### (d) PBBs

1. The principal use of PBBs was as flame retardants. PBBs are an additive type flame retardant. Mixed with a dry solid or liquid polymeric material, PBB provides a filter-type, flame retardant action by chemically releasing hydrogen bromide when ignited. Other uses of PBBs are: as color activators in light sensitive compositions; as relative molecular mass control agents for polybutadiene; as wood preservatives; as voltage stabilizing agents in electrical insulation; and as functional fluids, such as dielectric media (IPCS, 1994).
2. In the United States and Canada, FireMaster was used as a flame retardant in three main commercial products: acrylonitrile-butadienestyrene (ABS) thermoplastics (10 per cent PBBs) for housing business machines, industrial equipment (e.g., motor housing) and electronic products (e.g., radio and TV parts); as a flame retardant in coatings and lacquers; and in polyurethane foams for auto upholstery. Of the estimated 2,200 tonnes of HBB produced in 1974, about 900 tonnes were used in ABS plastic products and an even larger amount was used in cable coatings. Decabromobiphenyl Adine 0102 was used as a flame retardant in thermoplastics and thermosets (e.g., in polyesters, epoxy resins, polystyrene, ABS, polyolefin and PVC), elastomers (e.g., in PU-elastomers and India rubber) and cellulosic (e.g., in chip-boards), as well as in paints and varnishes (IPCS, 1994).
3. More recently, PBBs of predominantly low bromine content were found in electronic waste such as cable coatings, stuffing powder for electronic components and circuit boards, suggesting that it was used in such equipment (Zhao et al., 2008; IARC, 2014).

### 4. Wastes

1. Wastes consisting of, containing or contaminated with PCBs, PCTs, PCNs or PBBs (hereinafter referred to as “PCB, PCT, PCN or PBB wastes”) can be found in:

(a) Equipment containing or contaminated with PCBs, PCTs or PCNs (capacitors, circuit breakers, electrical cables, electric motors, electromagnets, heat transfer equipment, hydraulic equipment, switches, transformers, vacuum pumps, voltage regulators);

(b) Solvents contaminated with PCBs, PCTs or PCNs;

(c) Waste vehicles and shredder light fraction (fluff) containing or contaminated with PCBs or PCNs;

(d) Demolition wastes containing or contaminated with PCBs or PCNs (painted materials, resin‑based floorings, sealants, sealed glazing units);

(e) Oils consisting of, containing or contaminated with PCBs, PCTs or PCNs (dielectric fluids, heat transfer fluids, hydraulic fluids, motor oil);

(f) Electrical cables isolated by polymers containing or contaminated with PCBs, PCNs or PBBs;

(g) Soils and sediments, rock and aggregates (e.g., excavated bedrock, gravel, rubble) contaminated with PCBs, PCTs, PCNs or PBBs;

(h) Sludge contaminated with PCBs, PCTs, PCNs or PBBs;

(i) Plastics containing or contaminated with PCNs or PBBs and equipment containing such materials;

(j) Fire suppression equipment containing or contaminated with PBBs;

(k) Containers and absorbent materials contaminated through the handling, packaging, transportation or storage of PCB, PCT, PCN or PBB wastes;

(l) Wood contaminated with PCNs.

1. It should be noted that the abovementioned categories apply mainly to PCBs, which were produced in much larger quantities than PCTs, PCNs and PBBs and are stored as wastes awaiting disposal. It should also be noted that PCNs may be contained in traces in waste consisting of, containing or contaminated with PCBs (see paragraph 31). Since PBBs were used in electrotechnical products and automotive parts, it is possible that such products, if manufactured prior to 2000, contain PBBs. PBBs may also be present in shredder residue generated during the recycling process of waste vehicles and waste electrical and electronic equipment (WEEE).

# **II. Relevant provisions of the Basel and Stockholm conventions**

## A. Basel Convention

1. Article 1 (“Scope of Convention”) defines the types of waste that are subject to the Basel Convention. Subparagraph 1 (a) of that Article sets forth a two-step process for determining whether a “waste” is a “hazardous waste” subject to the Convention: first, the waste must belong to any category contained in Annex I to the Convention (“Categories of wastes to be controlled”), and second, the waste must possess at least one of the characteristics listed in Annex III to the Convention (“List of hazardous characteristics”).
2. Annex I to the Convention lists some of the wastes that may consist of, contain or be contaminated with PCBs,PCTs or PCNs. These include:

(a) Y5: Wastes from the manufacture, formulation and use of wood preserving chemicals;

(b) Y6: Wastes from the production, formulation and use of organic solvents;

(c) Y8: Waste mineral oils unfit for their originally intended use;

(d) Y9: Waste oils/water, hydrocarbons/water mixtures, emulsions;

(e) Y10: Waste substances and articles containing or contaminated with polychlorinated biphenyls (PCBs) and/or polychlorinated terphenyls (PCTs) and/or polybrominated biphenyls (PBBs);

(f) Y11: Waste tarry residues arising from refining, distillation and any pyrolytic treatment;

(g) Y12: Wastes from production, formulation and use of inks, dyes, pigments, paints, lacquers, varnish;

(h) Y13: Wastes from production, formulation and use of resins, latex, plasticizers, glues/adhesives;

(i) Y14: Waste chemical substances arising from research and development or teaching activities which are not identified and/or are new and whose effects on man and/or the environment are not known;

(j) Y18: Residues arising from industrial waste disposal operations;

(k) Y39: Phenols; phenol compounds including chlorophenols;

(l) Y41: Halogenated organic solvents;

(m) Y42: Organic solvents excluding halogenated solvents;

(n) Y45: Organohalogen compounds other than substances referred to in this Annex (e.g. Y39, Y41, Y42, Y43, Y44).

1. Annex I to the Convention lists some of the wastes that may consist of, contain or be contaminated with PBBs. These include:

(a) Y10: Waste substances and articles containing or contaminated with polychlorinated biphenyls (PCBs) and/or polychlorinated terphenyls (PCTs) and/or polybrominated biphenyls (PBBs);

(b) Y12: Wastes from production, formulation and use of inks, dyes, pigments, paints, lacquers, varnish;

(c) Y13: Wastes from production, formulation and use of resins, latex, plasticizers, glues/adhesives;

(d) Y14: Waste chemical substances arising from research and development or teaching activities which are not identified and/or are new and whose effects on man and/or the environment are not known;

(e) Y18: Residues arising from industrial waste disposal operations;

(f) Y41: Halogenated organic solvents;

(g) Y42: Organic solvents excluding halogenated solvents;

(h) Y45: Organohalogen compounds other than substances referred to in this Annex (e.g. Y39, Y41, Y42, Y43, Y44).

1. Annex I wastes are presumed to exhibit one or more Annex III hazardous characteristics, which may include H4.1 “Flammable solids”, H6.1 “Poisonous (Acute)”, H11 “Toxic (Delayed or chronic)”, H12 “Ecotoxic”, or H13 “Capable after disposal of yielding another hazardous material” unless, through “national tests”, they can be shown not to exhibit such characteristics. National tests may be useful for identifying a particular hazardous characteristic listed in Annex III until such time as the hazardous characteristic is fully defined. Guidance papers for Annex III hazardous characteristics H11, H12 and H13 were adopted on an interim basis by the Conference of the Parties to the Basel Convention at its sixth and seventh meetings.
2. List A of Annex VIII describes wastes that are “characterized as hazardous under Article 1, paragraph 1 (a) of this Convention” although “their designation on this Annex does not preclude the use of Annex III [hazard characteristics] to demonstrate that a waste is not hazardous” (Annex I, paragraph (b)). The following Annex VIII waste categories in particular are applicable to PCBs, PCTs, PCNs or PBBs:

(a) A1180: Waste electrical and electronic assemblies or scrap[[5]](#footnote-6) containing components such as accumulators and other batteries included on list A, mercury-switches, glass from cathode-ray tubes and other activated glass and PCB-capacitors, or contaminated with Annex I constituents (e.g., cadmium, mercury, lead, polychlorinated biphenyl) to an extent that they possess any of the characteristics contained in Annex III (note the related entry on list B B1110)[[6]](#footnote-7);

(b) A1190: Waste metal cables coated or insulated with plastics containing or contaminated with coal tar, PCB, lead, cadmium, other organohalogen compounds or other Annex I constituents to an extent that they exhibit Annex III characteristics;

(c) A3180: Wastes, substances and articles containing, consisting of or contaminated with polychlorinated biphenyl (PCB), polychlorinated terphenyl (PCT), polychlorinated naphthalene (PCN) or polybrominated biphenyl (PBB), or any other polybrominated analogues of these compounds, at a concentration level of 50 mg/kg or more[[7]](#footnote-8).

1. List A of Annex VIII includes a number of wastes or waste categories that have the potential to contain or be contaminated with PCBs, PCTs or PCNs, including:

(a) A1090: Ashes from the incineration of insulated copper wire;

(b) A1100: Dusts and residues from gas cleaning systems of copper smelters;

(c) A2040: Waste gypsum arising from chemical industry processes, when containing Annex I constituents to the extent that it exhibits an Annex III hazardous characteristic (note the related entry on list B B2080);

(d) A2060: Coal-fired power plant fly-ash containing Annex I substances in concentrations sufficient to exhibit Annex III characteristics (note the related entry on list B B2050);

(e) A3020: Waste mineral oils unfit for their originally intended use;

(f) A3040: Waste thermal (heat transfer) fluids;

(g) A3050: Wastes from production, formulation and use of resins, latex, plasticizers, glues/adhesives excluding such wastes specified on list B (note the related entry on list B B4020);

(h) A3070: Waste phenols, phenol compounds including chlorophenol in the form of liquids or sludges;

(i) A3120: Fluff - light fraction from shredding;

(j) A3150: Waste halogenated organic solvents;

(k) A3160: Waste halogenated or unhalogenated non-aqueous distillation residues arising from organic solvent recovery operations;

(l) A4040: Wastes from the manufacture, formulation and use of wood-preserving chemicals[[8]](#footnote-9);

(m) A4070: Wastes from the production, formulation and use of inks, dyes, pigments, paints, lacquers, varnish excluding any such waste specified on list B (note the related entry on list B B4010);

(n) A4100: Wastes from industrial pollution control devices for cleaning of industrial off-gases but excluding such wastes specified on list B;

(o) A4130: Waste packages and containers containing Annex I substances in concentrations sufficient to exhibit Annex III hazard characteristics;

(p) A4140: Wastes consisting of or containing off specification or outdated[[9]](#footnote-10) chemicals corresponding to Annex I categories and exhibiting Annex III hazard characteristics;

(q) A4150: Waste chemical substances arising from research and development or teaching activities which are not identified and/or are new and whose effects on human health and/or the environment are not known;

(r) A4160: Spent activated carbon not included on list B (note the related entry on list B B2060).

1. List A of Annex VIII includes a number of wastes or waste categories that have the potential to contain or be contaminated with PBBs, including:

(a) A3050: Wastes from production, formulation and use of resins, latex, plasticizers, glues/adhesives excluding such wastes specified on list B (note the related entry on list B B4020);

(b) A3150: Waste halogenated organic solvents;

(c) A3160: Waste halogenated or unhalogenated non-aqueous distillation residues arising from organic solvent recovery operations;

(d) A4070: Wastes from the production, formulation and use of inks, dyes, pigments, paints, lacquers, varnish excluding any such waste specified on list B (note the related entry on list B B4010);

(e) A4100: Wastes from industrial pollution control devices for cleaning of industrial off-gases but excluding such wastes specified on list B;

(f) A4130: Waste packages and containers containing Annex I substances in concentrations sufficient to exhibit Annex III hazard characteristics;

(g) A4140: Wastes consisting of or containing off specification or outdated12 chemicals corresponding to Annex I categories and exhibiting Annex III hazard characteristics;

(h) A4150: Waste chemical substances arising from research and development or teaching activities which are not identified and/or are new and whose effects on human health and/or the environment are not known;

(i) A4160: Spent activated carbon not included on list B (note the related entry on list B B2060).

1. List B of Annex IX to the Convention lists wastes that “will not be wastes covered by Article 1, paragraph 1 (a), of this Convention unless they contain Annex I material to an extent causing them to exhibit an Annex III characteristic.”
2. List B of Annex IX includes a number of wastes or waste categories that have the potential to contain or be contaminated with PCBs,PCTs or PCNs and its related substances, including: B1100: Metal-bearing wastes arising from melting, smelting and refining of metals.[[10]](#footnote-11)
3. List B of Annex IX includes a number of wastes or waste categories that have the potential to contain or be contaminated with PBBs, including:

(a) B3010: Cured waste resins or condensation products and fluorinated polymer wastes;[[11]](#footnote-12)

(b) B3030: Textile wastes.[[12]](#footnote-13)

1. For further information, see section II.A of the General technical guidelines.

## B. Stockholm Convention[[13]](#footnote-14)

1. The present technical guidelines cover intentionally produced PCBs, PCNs, including di- through octa-CNs, and HBB whose production and use are to be eliminated in accordance with Article 3 of, and Annex A to, the Stockholm Convention.
2. Annex A, part I, to the Convention does not include any exemption for the production or use of HBB.
3. Annex A, part I for PCNs, including di- through octa-CNs, allow specific exemptions for production of those chemicals as intermediates in production of polyfluorinated naphthalenes, including octafluoronaphthalene, and the use of those chemicals for the production of polyfluorinated naphthalenes, including octafluoronaphthalene.
4. Annex A, part II (“Polychlorinated biphenyls”) outlines specific requirements for PCBs, as follows:

“Each Party shall

(a) With regard to the elimination of the use of polychlorinated biphenyls in equipment (e.g. transformers, capacitors or other receptacles containing liquid stocks) by 2025, subject to review by the Conference of the Parties, take action in accordance with the following priorities:

(i) Make determined efforts to identify, label and remove from use equipment containing greater than 10 per cent polychlorinated biphenyls and volumes greater than 5 litres;

(ii) Make determined efforts to identify, label and remove from use equipment containing greater than 0.05 per cent polychlorinated biphenyls and volumes greater than 5 litres;

(iii) Endeavour to identify and remove from use equipment containing greater than 0.005 per cent polychlorinated biphenyls and volumes greater than 0.05 litres;

(b) Consistent with the priorities in subparagraph (a), promote the following measures to reduce exposures and risk to control the use of polychlorinated biphenyls:

(i) Use only in intact and non-leaking equipment and only in areas where the risk from environmental release can be minimised and quickly remedied;

(ii) Not use in equipment in areas associated with the production or processing of food or feed;

(iii) When used in populated areas, including schools and hospitals, take all reasonable measures to protect from electrical failure which could result in a fire, and regular inspection of equipment for leaks;

(c) Notwithstanding paragraph 2 of Article 3, ensure that equipment containing polychlorinated biphenyls, as described in subparagraph (a), shall not be exported or imported except for the purpose of environmentally sound waste management;

(d) Except for maintenance and servicing operations, not allow recovery for the purpose of reuse in other equipment of liquids with polychlorinated biphenyls content above 0.005 per cent;

(e) Make determined efforts designed to lead to environmentally sound waste management of liquids containing polychlorinated biphenyls and equipment contaminated with polychlorinated biphenyls having a polychlorinated biphenyls content above 0.005 per cent, in accordance with paragraph 1 of Article 6, as soon as possible but no later than 2028, subject to review by the Conference of the Parties;

(f) In lieu of note (ii) in Part I of this Annex, endeavour to identify other articles containing more than 0.005 per cent polychlorinated biphenyls (e.g. cable‑sheaths, cured caulk and painted objects) and manage them in accordance with paragraph 1 of Article 6;

(g) Provide a report every five years on progress in eliminating polychlorinated biphenyls and submit it to the Conference of the Parties pursuant to Article 15”;

(h) The reports described in subparagraph (g) shall, as appropriate, be considered by the Conference of the Parties in its reviews relating to polychlorinated biphenyls. The Conference of the Parties shall review progress towards elimination of polychlorinated biphenyls at five year intervals or other period, as appropriate, taking into account such reports.

1. For further information, see section II.B of the General technical guidelines.

# **III. Issues under the Stockholm Convention to be addressed cooperatively with the Basel Convention**[[14]](#footnote-15)

## A. Low POP content

1. The following provisional definitions of low POP content should be applied:
	1. PCBs: 50 mg/kg;[[15]](#footnote-16)
	2. HBB: 50 mg/kg;[[16]](#footnote-17)
	3. PCNs: 10 mg/kg.[[17]](#footnote-18)
2. The low POP content described in the Stockholm Convention is independent from the provisions on hazardous waste under the Basel Convention.
3. Wastes with a content of PCBs, HBB, or PCNs above the values specified in paragraph 68 must be disposed of in such a way that the POP content is destroyed or irreversibly transformed in accordance with the methods described in section IV.G.2. Otherwise, they may be disposed of in an environmentally sound manner when destruction or irreversible transformation does not represent the environmentally preferable option in accordance with the methods described in section IV.G.3.
4. Wastes with a content of PCBs, HBB, or PCNs at or below the values specified in paragraph 68 should be disposed of in accordance with the methods referred to in subsection IV.G.4 outlining disposal methods when POP content is low and sections IV.I.1 and IV.I.2 addressing pertinent higher and lower risk situations.
5. For further information on low POP content, refer to section III.A of the General technical guidelines.

## B. Levels of destruction and irreversible transformation

1. For the provisional definition of levels of destruction and irreversible transformation, see section III.B of the General technical guidelines.

## C. Methods that constitute environmentally sound disposal

1. See section IV.G below and section IV.G of the General technical guidelines*.*

# **IV. Guidance on environmentally sound management (ESM)**

## A. General considerations

1. For further information, see section IV.A of the General technical guidelines.

## B. Legislative and regulatory framework

1. Parties to the Basel and Stockholm conventions should examine their national strategies, policies, controls,[[18]](#footnote-19) standards and procedures to ensure that they are in agreement with the two conventions and their obligations under them, including those that pertain to ESM of wastes consisting of, containing or contaminated with PCBs, PCNs and HBB.
2. Elements of a regulatory framework applicable to PCBs, PCTs, PCNs and PBBs should include measures to prevent the generation of wastes and to ensure the environmentally sound management of generated wastes. Such elements could include the following:

(a) Environmental protection legislation establishing a regulatory regime, setting release limits and establishing environmental quality criteria;

(b) Prohibitions on the production, sale, use, import and export of PCBs, PCTs, PCNs and PBBs;

(c) Phase-out dates for PCBs that remain in service, inventory or storage;

(d) Transportation requirements for hazardous materials and waste;

(e) Specifications for containers, equipment, bulk containers and storage sites;

(f) Specification of acceptable analytical and sampling methods for PCBs, PCTs, PCNs and PBBs;

(g) Requirements for waste management and disposal facilities;

(h) Definitions of hazardous waste and conditions and criteria for the identification and classification of PCB, PCT, PCN and PBB wastes as hazardous wastes;

(i) A general requirement for public notification and review of proposed government waste-related regulations, policy, certificates of approval, licences, inventory information and national emissions data;

(j) Requirements for identification, assessment and remediation of contaminated sites;

(k) Requirements concerning the health and safety of workers;

(l) Other legislative measures on, e.g., waste prevention and minimization, inventory development and emergency response.

1. The timing of the phase-out of PCBs (and, to a lesser extent, of PCTs, PCNs or PBBs) will probably be the most critical legislative concern for most countries, given that most of them already have some form of legislative framework dealing with PCBs.
2. For further information, see section IV.B of the General technical guidelines.

## C. Waste prevention and minimization

1. Both the Basel and Stockholm conventions advocate waste prevention and minimization, and PCB compounds, PCNs and HBB are targeted for a complete phase-out in the Stockholm Convention. PCBs, PCTs, PCNs and PBBs should be taken out of service and disposed of in an environmentally sound manner.
2. Quantities of waste containing these compounds should be minimized through isolation and source separation to prevent mixing and contamination of other waste streams. For example, electrical equipment, painted materials, resin-based floorings, sealants and sealed glazing units containing PCBs can contaminate large amounts of demolition waste and should be separated where practicable prior to demolition.
3. The mixing and blending of wastes with a content of PCBs, HBB, or PCNs above the values specified in paragraph 68 with other materials solely for the purpose of generating a mixture with a content of PCBs, HBB, or PCNs at or below the values specified in paragraph 68 is not environmentally sound. Nevertheless, the mixing or blending of materials before waste treatment may be necessary in order to enable treatment or to optimize treatment efficiency.
4. To facilitate the reuse of electrical equipment containing insulating oil contaminated with PCBs, such as transformers, a procedure called retrofilling may be implemented in which the equipment is emptied of the insulating oil contaminated with PCBs and refilled with PCB-free insulating oil, such as mineral oil. In retrofilling procedures, care should be taken to avoid cross-contamination of retrofilled oils with any PCBs that may have penetrated the porous parts of the equipment, such as wood, cardboard, insulating paper and resins, and may gradually leach into retrofilled oils. As a preventive measure, some countries have enacted regulations whereby, in the absence of analyses to determine the presence or absence of PCBs in electrical equipment oils, such oils are presumed to contain PCBs until proven otherwise.[[19]](#footnote-20) Decontamination methodsshould be planned carefully to reduce the number of times that retrofilling procedures are conducted by requiring that initial PCB levels be considered and that every effort be made to empty equipment entirely. Retrofilled equipment should be periodically tested for PCBs and, when PCB levels exceed the low POP content, retrofilling should be performed again.
5. For further information, see section IV.C of the General technical guidelines.

## D. Identification of wastes

1. Article 6, paragraph 1 (a), of the Stockholm Convention requires each Party to, *inter alia,* develop appropriate strategies for the identification of products and articles in use and wastes consisting of, containing or contaminated with POPs. The identification of POPs wastes is the starting point for their effective ESM.
2. For general information on identification of waste, see section IV.D of the General technical guidelines.

### 1. Identification

1. PCBs, PCTs and PCNs have historically been found in a number of locations, as follows:

(a) In completely closed or nominally closed systems, including:

(i) Electrical utilities: transformers, capacitors, switches, voltage regulators, circuit breakers, light ballasts, and waste electrical and electronic equipment (WEEE) containing small capacitors and cables;

(ii) Industrial facilities: transformers, capacitors, voltage regulators, circuit breakers, light ballasts, heat transfer fluids and hydraulic fluids;

(iii) Railroad systems: transformers, capacitors, voltage regulators and circuit breakers;

(iv) Mining operations: hydraulic fluids, earthing coils, transformers and capacitors;

(v) Military installations: transformers, capacitors, voltage regulators and hydraulic fluids;

(vi) Residential/commercial buildings: capacitors, circuit breakers and light ballasts;

(vii) Research laboratories: vacuum pumps, light ballasts, capacitors and circuit breakers;

(viii) Electronics manufacturing plants: vacuum pumps, light ballasts, transformers, capacitors and circuit breakers;

(ix) Wastewater discharge facilities: capacitors, vacuum pumps and well motors;

(x) Communal and housing services (water pump stations, water distribution): capacitors;

(xi) Automotive service stations: reused oil;

(b) In open-ended systems, including:

(i) Residential/commercial buildings: elastic joints and fillers, sealants,[[20]](#footnote-21) paints, concrete, plaster, carbonless copy paper, fabrics, polyurethane foam, lubricants, synthetic rubber, adhesives and wood preservatives;

(ii) Steel structures such as bridges, tanks, ships or laying pipes: paints and coatings.

1. When identifying wastes containing or contaminated with PCBs, PCTs and PCNs, Parties may use an approach developed for PCBs inventorying and find it is useful to refer to *the Guidelines for the identification of PCBs and materials containing PCBs* (UNEP, 1999) and t*he PEN magazine, Issue “Inventories of PCBs – The Place to Start”* (UNEP, 2010).
2. For closed electrical equipment such as transformers and capacitors, generally it is possible to identify whether such equipment contains PCBs, PCTs or PCNs by inspecting the type designation on the equipment name plates and the product labels or literature issued by the manufacturer, and by referring to the date of production of the equipment. However, it should be noted that the information on PCB content indicated in the name plates may be incorrect in some cases. Through retrofilling or maintenance work, cross-contamination of insulating oils that result in PCB content above the values specified in paragraph 68 can occur even with equipment known as PCB-free and recently manufactured equipment. The insulating oils in all closed electrical equipment should therefore be analysed for possible PCB, PCT or PCN content.
3. In the case of light ballasts and WEEE equipped with small capacitors, it is difficult to determine whether they contain PCBs, PCTs or PCNs as dielectric fluids. The PCB, PCT or PCN content of such equipment should be carefully determined by referring to equipment type designations and production dates.
4. For open-ended materials such as joint sealants or paints separated from demolition waste, it is impossible to judge whether these contain PCBs or PCNs solely from their appearance. Therefore, the time when such materials were applied should be verified and, if the materials were produced during the time when PCBs or PCNs were used as plasticizers, a test for the presence of PCBs or PCNs in the waste should be performed.
5. It is difficult even for experienced technicians to identify the characteristics of effluents, substances, containers or equipment solely from their external appearance or labels. With respect to electrical equipment such as transformers and capacitors, it is possible to identify the brand of the equipment and thus confirm the year and country where it was manufactured, as well as its manufacturer. By referring to available information or by contacting the manufacturer, it is possible to determine whether the equipment contains PCBs, PCTs or PCNs. If PCB, PCT or PCN-containing equipment does not have any label pertaining to its insulating oil, experienced investigators can obtain information on the original contents and other information through labels of similar equipment, by referring to relevant guidance manuals, such as the Guidelines for the identification of PCBs and materials containing PCBs (UNEP, 1999), or by contacting the manufacturer.
6. PBBs have historically been found in a number of consumer products where they were used as flame retardants, including a variety of plastic products such as computer monitors, televisions, textiles and plastic foams (including those in WEEE and shredder residue generated during the recycling process of waste vehicles).
7. Parties may find the information on production, use and waste types provided in section I.B of the present guidelines useful in identifying PCBs, PCTs, PCNs and PBBs.

### 2. Inventories

1. Inventories are an important tool for identifying, quantifying and characterizing wastes. A step-by-step approach for the development of national inventories of PCBs, PCTs, PCNs and PBBs generally includes the following steps:

(a) Step 1: planning (i.e., identifying relevant sectors of use and production of PCBs, PCTs, PCNs and PBBs;

(b) Step 2: choosing data collection methodologies using a tiered approach;

(c) Step 3: collecting and compiling data from national statistics on the production, use, import and export of PCBs, PCT, PCNs and PBBs;

(d) Step 4: managing and evaluating the data obtained in step 3 using an estimation method;

(e) Step 5: preparing an inventory report;

(f) Step 6: periodically updating the inventory report.

1. For further information, refer to the *Guidelines for the identification of PCBs and materials containing PCBs* (UNEP, 1999).

## E. Sampling, analysis and monitoring

1. For general information on sampling, analysis and monitoring, see section IV.E of the General technical guidelines.

### 1. Sampling

1. It is difficult to extract dielectric fluid samples from sealed waste electrical equipment such as capacitors. To obtain such samples, a small hole should carefully be drilled on the top of the equipment. After taking the sample, the hole should be plugged and repaired.
2. During sampling of shredder residue, efforts should be made to ensure sample homogeneity.
3. The types of matrices that are of special interest for analysis of PCBs, PCTs, PCNs and PBBs include:

(a) Industrial synthetic PCB, PCT and PCN-containing oils from transformers or other equipment or in bulk storage;

(b) Mineral oils from retrofilled transformers contaminated with PCBs or in bulk storage;

(c) Waste motor oils and other waste oils, fuels and organic liquids;

(d) Elastic joints and fillers, sealants and paints;

(e) Fire suppressants and retardants (PBBs).

### 2. Analysis

1. Analysis refers to the extraction, purification, separation, identification, quantification and reporting of POP concentrations in the matrix of interest. The development and dissemination of reliable analysis methods and the accumulation of high-quality analytical data are important to understand the environmental impact of hazardous chemicals, including POPs.
2. As with all polybrominated flame retardants, samples should not be exposed to sunlight prior to analysis, since PBBs are unstable when exposed to ultraviolet radiation (IARC, 2014).
3. Methods of analysing the various matrices for POPs have been developed by ISO, the European Committee for Standardization (CEN – EN standards), ABNT, AOAC, ASTM, DIN, EPA, JIS, NEN and NVN. Some examples of analytical methods for PCBs include the following:

(a) Methods for oils or insulating liquids:

 (i) EN 12766-1 (2000): Petroleum products and used oils - Determination of PCBs and related products - Part 1: Separation and determination of selected PCB congeners by gas chromatography (GC) using an electron capture detector (ECD);

 (ii) EN 12766-2 (2002): Petroleum products and used oils - Determination of PCBs and related products - Part 2: Calculation of polychlorinated biphenyl (PCB) content;

 (iii) EN 61619 (1997): Insulating liquids - Contamination by polychlorinated biphenyls (PCBs) - Method of determination by capillary column gas chromatography;

 (iv) EPA Method 4020: Screening for polychlorinated biphenyls by immunoassay (www.epa.gov/solidwaste/hazard/testmethods/sw846/pdfs/4020.pdf);

 (v) EPA Method 8082A: Polychlorinated biphenyls (PCBs) by gas chromatography (www.epa.gov/osw/hazard/testmethods/sw846/pdfs/8082a.pdf);

 (vi) EPA Method 9079: Screening test method for polychlorinated biphenyls in transformer oil (www.epa.gov/solidwaste/hazard/testmethods/sw846//pdfs/9079.pdf);

 (vii) ABNT NBR 13882:2005: Electrical Insulating Liquids - Determination of PCB contents;

(b) Methods for solid materials:

 (i) EN 15308 (2008):Characterization of waste - Determination of selected polychlorinated biphenyls (PCBs) in solid waste by using capillary gas chromatography with electron capture or mass spectrometric detection;

 (ii) EPA Method 8080: Organochlorine Pesticides and PCBs;

 (iii) Methods for Examining Standards of General Wastes under Special Control and Industrial Waste under Special Control, Notice 192 of the Japan Ministry of Welfare and Labour, 3 July 1992;

(c) Methods for water, sludge, gases and others:

 (i) DIN 38414-20 (1996): German standard methods for the examination of water, waste water and sludge - Sludge and sediments (group S) - Part 20: Determination of 6 polychlorinated biphenyls (PCBs) (P 20);

 (ii) EN 1948 (2006): Stationary source emissions - determination of the mass concentration of PCDDs/PCDFs and dioxin-like PCBs. Part 1 Sampling, Part 2: Extraction and clean-up of PCDDs/PCDFs, Part 3: Identification and quantification of PCDDs/PCDFs;

 (iii) EPA Method 1668, Revision A: Chlorinated Biphenyl Congeners in Water, Soil, Sediment, and Tissue by HRGC/HRMS, United States Office of Water, EPA No. EPA 821-R-00-002, Environmental Protection Agency (4303), December 1999;

 (iv) EPA Method 8275A: Semivolatile organic compounds (PAHs and PCBs) in soils/sludges and solid wastes using thermal extraction/gas chromatography/mass spectrometry (TE/GC/MS), EPA analytical chemistry guidance SW-846;

 (v) EPA Method 9078: Screening test method for polychlorinated biphenyls in soil (www.epa.gov/epaoswer/hazwaste/test/pdfs/9078.pdf);

 (vi) ISO 6468 (1996): Water quality - Determination of certain organochlorine insecticides, polychlorinated biphenyls and chlorobenzenes - Gas chromatographic method after liquid-liquid extraction;

 (vii) ISO 10382 (2002): Soil quality - Determination of organochlorine pesticides and polychlorinated biphenyls - Gas-chromatographic method with electron capture detection;

 (viii) JIS K 0093 (2006): Testing method for polychlorinated biphenyls in industrial water and wastewater;

 (ix) NEN 7374 (2004): Leaching characteristics - Column test for the determination of the leaching of PAH, PCB, OCP and EOX, phenol and creosols from granular materials - Solid earthy and stony materials;

 (x) Norwegian Institute for Water Research method no. H 3-2: Determination of organochlorine compounds in sediments, water and biological material by gas chromatography;

 (xi) NVN 7350 (1997): Leaching characteristics of solid earthy and stony building and waste materials - Leaching tests - Determination of the leaching of PAH, PCB and EOX from granular materials with the cascade test;

 (xii) NVN 7376 (2004): Leaching characteristics - Determination of the leaching of PAH, PCB, OCP and EOX, phenol and creosols from building and monolithic waste materials with a diffusion test - Solid earthy and stony materials.

1. A method of analysing water for PCNs has been developed by the International Organization for Standardization (ISO), which is as follows: ISO/TS 16780 (2015) Water quality– Determination of polychlorinated naphthalenes (PCN)—Method using gas chromatography (GC) and mass spectrometry (MS).
2. PCNs analysis is commonly based on fractionation and carbon clean-up followed by GC/MS. It should be noted that there are some problems in PCNs analysis. A problem is the limited commercial availability of authentic standards and quite limited commercially available 13C-labeled PCN congeners as internal standards. Another problem is the co-elution of some PCN congeners in capillary gas chromatographic separation. Further problem is that PCBs could co-elute with PCNs and interfere with their detection, if gas chromatography alone was used. However, useful knowledge can be obtained from the following literature regarding the method of analysis of various matrices for PCNs:

(a) Abad E, et al., 1999. “Dioxin like compounds from municipal waste incinerator emissions: assessment of the presence of polychlorinated naphthalenes”, *Chemosphere*, vol. 38, pp. 109-120;

(b) Falandysz J., et al., 2006. “HRGC/HRMS analysis of chloronaphthalenes in several batches of Halowax 1000, 1001, 1013, 1014 and 1099”, *Journal of Environmental Science & Health, Part A*. vol. 41, pp. 2237-2255;

(c) Helm P.A., 1999. “Complete separation of isomeric penta- and hexachloronaphthalenes by capillary gas chromatography”, *Journal of High Resolution Chromatography*, vol. 22, pp. 639-643;

(d) Järnberg U., et al., 1994. “Gas chromatographic retention behaviour of Polychlorinated naphthalenes on non-polar, polarizable, polar and smectic capitally columns”, *Journal of Chromatography A*, vol. 783, pp. 385-396;

(e) Liu G., et al., 2014. “Sources of unintentionally produced polychlorinated naphthalenes”, *Chemosphere*, vol. 94, pp. 1-12;

(f) Noma Y., et al., 2006. “Behavior of PCNs, PCDDs, PCDFs, and Co-PCBs in the thermal destruction of wastes containing PCNs”, *Chemosphere*, vol. 62, pp. 1183-1195;

(g) Taniyasu S., et al., 2003. “Isomer-specific analysis of chlorinated biphenyls, naphthalenes and dibenzofurans in Delor: polychlorinated biphenyl preparations from the former Czechoslovakia”, *Environmental Pollution*, vol.126, pp. 169-178.

1. A method of analysing electrotechnical products for PBBs have been developed by the International Electrotechnical Commission (IEC), which is as follows: IEC 62321 (2008): Electrotechnical products – Determination of levels of six regulated substances (lead, mercury, cadmium, hexavalent chromium, polybrominated biphenyls, polybrominated diphenyl ethers).
2. Furthermore, useful knowledge can be obtained from the following literature regarding the method of analysis of various matrices for PBBs:

(a) US Agency for Toxic Substances and Disease Registry, 2004. *Toxicological profile for polybrominated biphenyls and polybrominated diphenyl ethers*;

(b) Kemmlein, S. et al., 2009. “Brominated flame retardants in the European chemicals policy of REACH-Regulation and determination in materials”, *Journal of Chromatography A*, vol. 1216 No. 3, pp. 320-333;

(c) Clarke, B. et al., 2008. “Polybrominated diphenyl ethers and polybrominated biphenyls in Australian sewage sludge”, *Chemosphere*, vol. 73, pp. 980-989;

(d) Covaci, A. et al., 2003. “Determination of brominated flame retardants, with emphasis on polybrominated diphenyl ethers (PBDEs) in environmental and human samples: A review”, *Environment International*, vol. 29, pp. 735-756;

(e) Hanari, N. et al., 2006. “Occurrence of polybrominated biphenyls, polybrominated dibenzo-p-dioxins, and polybrominated dibenzofurans as impurities in commercial polybrominated diphenyl ether mixtures”, *Environmental Science & Technology*, vol. 40, pp. 4400-4405.

1. To determine dioxin-like PCBs and PBBs, which could be of special interest to Parties, internationally accepted methods such as those for analysing PCDDs/PCDFs should be applied.
2. For screening purposes, test kits are available for the quantification of PCBs in oils and soils (based on immunoassays or chlorine determinations). If the result is negative, a confirmatory PCB analysis is not necessary. If the result is positive, confirmatory chemical analysis should be performed, or the waste may be regarded as waste containing or contaminated with PCBs.

### 3. Monitoring

1. Monitoring and surveillance serve as elements for identifying and tracking environmental concerns and human health risks. Information collected from monitoring programs feeds into science-based decision-making processes and is used for the evaluation of the effectiveness of risk management measures, including regulations.
2. Monitoring programmes should be implemented in facilities managing PCB, PCT, PCN or PBB wastes.

## F. Handling, collection, packaging, labelling, transportation and storage

1. For general information on handling, collection, packaging, labelling, transportation and storage, see section IV.F of the General technical guidelines.

### 1. Handling

1. Special attention should be paid to possible PCB, PCTs or PCN leakages due to corrosion or defects of PCB, PCT or PCN-containing electrical equipment such as transformers and capacitors, as such equipment generally has a lifetime of several decades. Special attention should also be paid to damages that may result from moving such equipment. Heavy electrical machineries require caution in handling bushings, as these are prone to breakage under heavy loads. When dealing with highly concentrated PCBs, PCTs or PCNs, operators should use masks and rubber gloves to avoid inhalation of volatilized PCBs or skin contact with PCBs, PCTs or PCNs.
2. When conducting repairs in or renovation or demolition of older buildings, renovators and contractors should pay attention to the possibility of PCBs or PCNs being contained in building joints, window or door sealants or fillers, and paint coatings on steel bridges or structures. Should these materials contain PCBs or PCNs, they should be carefully removed and isolated to prevent PCB or PCN-containing dust from spreading to surrounding areas. The work should be conducted wearing appropriate protective equipment such as suitable gloves, disposable coveralls, protective goggles and respiratory protection masks that meet international standards.

### 2. Collection

1. A significant fraction of total national inventories of PCBs, PCTs, PCNs and PBBs may be held in small quantities by small business owners or homeowners (for example, in PCB fluorescent light ballasts; small electrical devices, heat exchangers and heaters containing PCB, PCT or PCN fluids; fire suppression systems containing PBBs; and small containers and small stockpiles of those substances).It is difficult for those who own small quantities of PCBs, PCTs, PCNs or PBBs to dispose of such materials. For example, regulations may require them to be registered as waste generators, logistical considerations may prevent or discourage pick-up (e.g., no industrial waste pick-up is allowed or available in their neighbourhood), and disposal costs may be prohibitive. National, regional and municipal governments should consider establishing collection stations for those small quantities so that each small-quantity owner does not have to make individual transport and disposal arrangements.
2. In cases where PCBs, PCTs and PCNs are found when conducting repairs in, or renovation or demolition of, older buildings (e.g., elastic joints and fillers, sealants, paints, concrete and plaster containing PCBs, PCTs and PCNs), the safety of workers should be ensured and wastes should be carefully removed and collected separately to prevent dust containing PCBs, PCTs and PCNs from spreading to surrounding areas.
3. Collection arrangements and collection depots for PCB, PCT, PCN and PBB wastes should provide for the separation of those wastes from other wastes.
4. It is imperative that collection depots do not become long-term storage facilities for PCB, PCT, PCN or PBB wastes. Large amounts of wastes, even if properly stored, pose a higher risk to the environment and human health than small quantities scattered over large areas.

### 3. Packaging

1. PCB, PCT, PCN and PBB wastes should be properly packaged before storage for ease of transport and as a safety measure to reduce the risk of leaks and spills:
2. With regard to transformers whose insulating oils have been drawn out, drained oils and carcasses should be packed separately. The risk of leakage during transport to a treatment facility may be lowered by separating insulating oils from transformers. Such separation should ideally be taken into account when evaluating packaging methods. Separation procedures should be performed by professional operators using special tools;
3. Liquid wastes should be placed in double-bung steel drums or other approved containers;
4. Solid wastes such as sealants and paints should be placed in steel drums or other approved containers lined with plastic bags;
5. Regulations governing the transport of hazardous materials often require the use of containers that meet certain specifications (e.g., 16‑gauge, made of steel, internally coated with epoxy). Containers used for storage should meet such specifications given that they may be transported in the future;
6. Large drained equipment may be stored as is or may be placed inside large containers (overpack drums) or heavy plastic wraps if leakage is a concern;
7. Small pieces of equipment, whether drained or not, should be placed in drums with an absorbent material, where appropriate, to prevent excessive movement of container contents and enable any free liquids and spills to be absorbed. Numerous small pieces of equipment may be placed in the same drum so long as an adequate amount of absorbent material is present in the drum. Loose absorbents may be purchased from safety suppliers;
8. Drums and equipment may be placed on pallets for movement by forklift truck and for storage. Drums and equipment should be strapped to the pallets before they are moved.

### 4. Labelling

1. Every container and piece of equipment containing or contaminated with PCBs, PCTs, PCNs or PBBs should be clearly labelled with a hazard-warning label and a label giving details of the equipment or container. Such details should include the contents of the container or equipment (e.g., exact counts of equipment, volume of liquid, type of waste carried), the name of the site from which the container or equipment originated so as to allow its traceability and, if applicable, the date of repackaging and the name and telephone number of the person responsible for the repackaging operation.

### 5. Transportation

1. Since PCBs are transported mostly in liquid form, measures should be taken to prevent leakage during transport. Transformers and capacitors, for example, should be secured in metallic containers to reduce the risk of breakage of equipment bushings from impact during transport, and include absorbent materials in their packaging.

### 6. Storage

1. While many countries have adopted storage regulations or developed storage guidelines concerning PCBs, most do not have specific storage regulations or guidance concerning PCTs, PCNs and PBBs. Although PCBs, PCTs, PCNs and PBBs have similar toxicity, PCBs are liquid at room temperature, while PCTs, PCNs other than mono-CNs and PBBs are solid at room temperature and have a lower vapour pressure. The storage conditions required for PCTs, PCNs and PBBs may therefore differ from those required for PCBs.
2. An oil pan (steel tray) should be laid underneath stored equipment. Storage sites should be maintained and inspected to verify whether there have been any releases of PCBs, PCTs, PCNs or PBBs into the environment.
3. In order to prevent the spillage of PCBs, PCTs or PCNs from equipment that has tumbled down during disasters such as earthquakes, tornadoes and heavy rains, or from leakage due to equipment corrosion, storage sites should have structures to prevent underground leakage. Moreover, it should be taken into account that PCBs, PCTs or PCNs might be released into the environment through vaporization during storage. It should be noted that PCNs are usually more volatile than PCBs because of the higher vapour pressure.

## G. Environmentally sound disposal

### 1. Pre-treatment

1. Cutting and milling of capacitors or disassembling of external parts such as radiators, conservators and bushings of transformers for the purpose of size reduction should be carried out before destruction in dedicated facilities. Caution should be taken during disassembling and dismantling operations, since such operations increase the risk of operator exposure to PCBs, PCTs or PCNs and of PCBs, PCTs or PCNs releases into the environment.
2. When destroying PCBs in waste oils or liquid wastes through alkali metal reduction, dewatering or oil-water separation should be done as a pre-treatment procedure in order to avoid the violent reaction of water with, and the excessive consumption of, alkali metals.
3. Since wastes containing PCBs from open-ended systems such as joint sealants or paints are generally bulky, crushing or shredding should be done as a pre-treatment to reduce them into smaller pieces and, when necessary, thermal desorption or vacuum thermal desorption should be implemented to effectively treat the PCBs in the wastes.
4. For further information on pre-treatment, see subsection IV.G.1 of the General technical guidelines.

### 2. Destruction and irreversible transformation methods

1. For information on destruction and irreversible transformation methods relating to PCBs, PCTs, PCNs and HBB see subsection IV.G.2 of the General technical guidelines.
2. It should be noted that PCDDs/PCDFs can be generated from the combustion and incineration of PCB, PCT or PCN wastes, while the combustion and incineration of PBB wastes can generate PBDDs/PBDFs.

### 3. Other disposal methods when neither destruction nor irreversible transformation is the environmentally preferable option

1. For information, see subsection IV.G.3 of the General technical guidelines.

### 4. Other disposal methods when the POP content is low

1. When cleansing or decomposing PCB-contaminated transformer insulating oils on site the spilling or leakage of effluents should be prevented even if the level of PCBs in the oils is relatively low.
2. For further information, see subsection IV.G.4 of the General technical guidelines.

## H. Remediation of contaminated sites

1. For information, see subsection IV.H of the General technical guidelines.

## I. Health and safety

1. For information, including on the distinction between higher- and lower-risk situations, see section IV.I of the General technical guidelines.

### 1. Higher-risk situations

1. For information on higher-risk situations, see subsection IV.I.1 of the General technical guidelines. Potential higher-risk situations specific to PCBs, PCTs, PCNs or PBBs may include:

(a) In electrical rooms with large or multiple PCB transformers, circuit breakers or capacitors;

(b) At sites at which PCB-containing transformers, circuit breakers, hydraulic equipment or vacuum pumps have been used or maintained;

(c) At sites where PCBs are separated from equipment and transferred to another container or where pre-handling measures such as disassembling of equipment are carried out. Caution should be taken at those sites, since they present increased risk of exposure for operators;

(d) In buildings in which PCBs have been used in elastic joints and fillers, paints or sealants.

### 2. Lower-risk situations

1. For information on lower risk situations, see subsection IV.I.2 of the General technical guidelines. Lower-risk situations specific to PCBs, PCTs, PCNs or PBBs may include:

(a) Those that involve only products or articles that contain or are contaminated with PCBs in small quantities or at low concentrations (e.g., some electrical and electronic equipment and waste equipment);

(b) Those that involve electrical transformers or other equipment with low-level PCB-contaminated mineral oils.

## J. Emergency response

1. Emergency response plans should be in place for PCBs, PCTs, PCNs and PBBs that are in service, in storage, in transport and at disposal sites. Further information on emergency response plans is given in section IV.J of the General technical guidelinesand in *Preparation of a national environmentally sound plan for PCBs and PCB-contaminated equipment: training manual* (UNEP, 2003).

## K. Public participation

1. Parties to the Basel or Stockholm Convention should have open public participation processes.
2. For further information see section IV.K of the General technical guidelines.

# **Annex I to the technical guidelines**

#  **Synonyms and trade names for PCBs, PCTs,** **PCNs, PBBs other than HBB,** **and HBB**

|  |  |
| --- | --- |
| **Chemical** | **Some synonyms and trade names**[[21]](#footnote-22)1 |
| PCBs | Abestol, Aceclor, Adkarel, ALC, Apirolio (Italy), Apirorlio, Areclor, Arochlor, Arochlors, Aroclor/Arochlor(s) (USA), Arubren, Asbestol (USA), Ask/Askarel/Askael, Auxol, Bakola, Biclor, Blacol (Germany), Biphenyl, Clophen (Germany), Cloresil, Chlophen, Chloretol, Chlorextol (USA), Chlorfin, Chlorinal/Chlorinol, Chlorinated biphenyl, Chlorinated diphenyl, Chlorobiphenyl, Chlorodiphenyl, Chlorofen (Poland), Chlorphen, Chorextol, Chorinol, Clophen/Clophenharz (Germany), Cloresil, Clorinal, Clorphen, Crophene (Germany), Decachlorodiphenyl, Delofet O-2, Delor (former Czechoslovakia), Delor/Del (former Czechoslovakia), Delorene, Delorit, Delotherm DK/DH (former Czechoslovakia), Diaclor (USA), Diarol, Dicolor, Diconal, Disconon, DK (Italy), Ducanol, Duconal, Duconol, Dykanol (USA), Dyknol, Educarel, EEC-18, Elaol (Germany), Electrophenyl, Elemex (USA), Elinol, Eucarel, Euracel, Fenchlor (Italy), Fenclor (Italy), Fenocloro, Gilotherm, Hexol, Hivar, Hydelor, Hydol, Hydrol, Hyrol, Hyvol (USA), Inclor, Inerteen (USA), Inertenn, Kanechlor (Japan), Kaneclor, Kennechlor (Japan) , Kenneclor, Leromoll, Magvar, MCS 1489, Montar, Monter, Nepoli, Nepolin, Niren, NoFlamol, No-Flamol (USA), Nitrosovol (former USSR), Non-Flamol, Olex-sf-d, Orophene, Pheaoclor, Pheneclor, Phenochlor, Phenoclor (France), Plastivar, Polychlorinated diphenyl, Polychlorinated diphenyls, Polychlorobiphenyl, Polychlorodiphenyl, Prodelec, Pydraul, Pyraclor, Pyralene (France), Pyranol (USA), Pyroclor (USA), Pyrochlor, Pyronol, Safe-T-Kuhl, Saft-Kuhl, Saf-T-Kohl, Saf-T-Kuhl (USA), Santosol, Santotherm (Japan), Santothern, Santovac, Sat-T-America, Siclonyl, Solvol, Sorol, Soval, Sovol (former USSR), Sovtol, Tarnol (Poland), Terphenychlore, Therminal, Therminol, Trichlorodiphenyls (former USSR),Turbinol |
| PCTs | Aroclor (USA), Clophen Harz (Germany), Cloresil A, B, and 100 (Italy), Electrophenyl T-50 and T60 (France), Kanechlor KC-C (Japan), Leromoll (Germany), Phenoclor (France), Pydraul (USA) |
| PCNs | Basileum SP‑70 (Germany), Cerifal (Italy), Chlonacire wax 90, 115 and 130 (France), Halowax 1000, 1001, 1013, 1014, 1031, 1051, 1099, 1099B, 2141 and 2148 (USA), Halowax (former USSR), Hodogaya Amber wax (Japan), Nankai wax (Japan), Nibren wax D88, D116N and D130 (Germany), N-Oil (USA), N-Wax (USA), Perna wax (Germany), Seekay wax R68, R93, R123, R700, RC93 and RC123 (United Kingdom), Tokyo Ohka wax (Japan), Woskol (Poland) |
| PBBsother thanHBB | Adine 0102 (France), Berkflam B10 (United Kingdom), Bromkal 80 (Germany), Bromkal 80-9D (Germany), Octabromobiphenyl FR250 13A (USA), Flammex B-10 (United Kingdom), HFO 101 (United Kingdom), BB-8, BB-9, OBB, Technical octabromobiphenyl (USA), DBB, Technical decabromobiphenyl (USA) |
| HBB | FireMaster BP-6 (USA), FireMaster FF-1 (USA) |

# **Annex II to the technical guidelines**

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1. Decisions V/8, VI/23, VII/13 and VIII/16, BC-10/9, BC-11/3, BC-12/3 and BC-13/4 of the Conference of the Parties to the Basel Convention on the Control of Transboundary Movements of Hazardous Wastes and Their Disposal; decisions OEWG‑I/4, OEWG-II/10 OEWG-III/8, OEWG‑IV/11, OEWG-V/12, OEWG-8/5, OEWG-9/3 and OEWG-10/4 of the Open-ended Working Group (OEWG) of the Basel Convention; resolution 5 of the Conference of Plenipotentiaries of the Stockholm Convention on Persistent Organic Pollutants; decisions INC-6/5 and INC-7/6 of the Intergovernmental Negotiating Committee for an International Legally Binding Instrument for Implementing International Action on Certain Persistent Organic Pollutants; and decisions SC‑1/21, SC-2/6, SC‑4/13 and SC-7/14 of the Conference of the Parties to the Stockholm Convention on Persistent Organic Pollutants. [↑](#footnote-ref-2)
2. The estimated production amount and the period of manufacture of PCBs are summarized in table 1 of UNEP/POPS/COP.7/INF/9. [↑](#footnote-ref-3)
3. In the Halowax series, a four-digit number follows the word Halowax. They exhibit a wide range from nearly pure mono-CNs (Halowax 1031) to nearly pure octa-CN (Halowax 1051) and from 22% to 70% of the chlorine content, respectively. [↑](#footnote-ref-4)
4. IPCS, 1994 and IARC, 2014. [↑](#footnote-ref-5)
5. This entry does not include scrap assemblies from electric power generation. [↑](#footnote-ref-6)
6. PCBs are at a concentration level of 50 mg/kg or more. [↑](#footnote-ref-7)
7. The 50 mg/kg level is considered to be an internationally practical level for all wastes. However, many individual countries have established lower regulatory levels (e.g., 20 mg/kg) for specific wastes. [↑](#footnote-ref-8)
8. This entry does not include wood treated with wood preserving chemicals. [↑](#footnote-ref-9)
9. “Outdated” means unused within the period recommended by the manufacturer. [↑](#footnote-ref-10)
10. Refer to Annex IX to the Basel Convention for a full description of this entry. [↑](#footnote-ref-11)
11. *Ibid* 10. [↑](#footnote-ref-12)
12. *Ibid* 10. [↑](#footnote-ref-13)
13. This section does not apply to PCTs, to mono-CNs and to PBBs other than HBB. [↑](#footnote-ref-14)
14. This section does not apply to PCTs, to mono-CNs and to PBBs other than HBB. [↑](#footnote-ref-15)
15. Determined in accordance with national or international methods and standards. [↑](#footnote-ref-16)
16. *Ibid* 15. [↑](#footnote-ref-17)
17. *Ibid* 15. [↑](#footnote-ref-18)
18. In these guidelines, national legislation and control measures include sub-national and other applicable forms of governance.  [↑](#footnote-ref-19)
19. For example, see Argentina’s 2002 *Law 25.670*. [↑](#footnote-ref-20)
20. Buildings built mainly between 1950 and 1980 may contain PCBs in joint sealants. [↑](#footnote-ref-21)
21. 1 The list of trade names provided in annex I is not intended to be exhaustive. [↑](#footnote-ref-22)