

**FOREWORD**

**INTRODUCTION**

**PHOSPHORUS TRICHLORIDE**

**CAS N°: 7719-12-2**

## SIDS Initial Assessment Report

For

### SIAM 19

Berlin, Germany, 19-22 October 2004

- 1. Chemical Name:** Phosphorus trichloride
- 2. CAS Number:** 7719-12-2
- 3. Sponsor Country:** Germany  
Contact Point:  
BMU (Bundesministerium für Umwelt, Naturschutz und  
Reaktorsicherheit)  
Contact person:  
Prof. Dr. Ulrich Schlottmann  
Postfach 12 06 29  
D- 53048 Bonn
- 4. Shared Partnership with:**
- 5. Roles/Responsibilities of the Partners:**
  - Name of industry sponsor /consortium Bayer AG, Germany  
Contact person:  
Dr. Burkhardt Stock  
D-51368 Leverkusen  
Gebäude 9115
  - Process used The BUA Peer Review Process : see next page
- 6. Sponsorship History**
  - How was the chemical or category brought into the OECD HPV Chemicals Programme? by ICCA-Initiative
- 7. Review Process Prior to the SIAM:** last literature search (update):  
8 April 2004 (Human Health): databases medline, toxline; search profile CAS-No. and special search terms  
5 March 2004 (Ecotoxicology): databases CA, biosis; search profile CAS-No. and special search terms OECD/ICCA
- 8. Quality check process:** As basis for the SIDS-Dossier the IUCLID was used. All data have been checked and validated by BUA. A final evaluation of the human health part has been performed by the Federal Institute for Risk Assessment (BfR) and of the ecotoxicological part by the Federal Environment Agency (UBA).
- 9. Date of Submission:** Deadline for circulation: 23 July 2004
- 10. Date of last Update:** Last literature search: IUCLID Chapters 1-4: 2002-07-26,  
Chapter 5: 2003-05-01

**11. Comments:****OECD/ICCA - The BUA\* Peer Review Process**

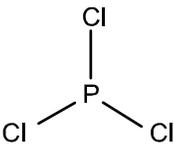
Qualified BUA personnel (toxicologists, ecotoxicologists) perform a quality control on the full SIDS dossier submitted by industry. This quality control process follows internal BUA guidelines/instructions for the OECD/ICCA peer review process and includes:

- a full (or update) literature search to verify completeness of data provided by industry in the IUCLID/HEDSET
- Review of data and assessment of the quality of data
- Review of data evaluation
- Check of adequacy of selection process for key studies for OECD endpoints, and, where relevant, for non-OECD endpoints by checking original reports/publications
- Review of key study description according to robust summary requirements; completeness and correctness is checked against original reports/publications (if original reports are missing: reliability (4), i.e. reliability not assignable)
- Review of validity of structure-activity relationships
- Review of full SIDS dossier (including SIAR, SIAP and proposal for conclusion and recommendation for further work)
- In case of data gaps, review of testing plan or rationale for not testing

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\* BUA (GDCh-Beratergremium für Altstoffe): Advisory Committee on Existing Chemicals of the Association of German Chemists (GDCh)

**SIDS INITIAL ASSESSMENT PROFILE**

<b>CAS No.</b>	7719-12-2
<b>Chemical Name</b>	Phosphorus trichloride
<b>Structural Formula</b>	
<b>SUMMARY CONCLUSIONS OF THE SIAR</b>	
<p><b>Human Health</b></p> <p>Phosphorus trichloride is quickly hydrolysed at first contact with water. It is, therefore, very unlikely that phosphorus trichloride will reach tissues distant from the portal of entry and become systemically available. The products of hydrolysis, hydrochloric acid and phosphorous acid, also act at the portal of entry.</p> <p>The acute toxicity of phosphorus trichloride is high. It is characterised by immediate irritation/corrosion at the portal of entry in experimental animals and humans due to the irritant/corrosive properties of the products of hydrolysis. After inhalation (4h in some studies, unspecified in others) the LC<sub>50</sub> was determined in rats as 226 to &gt;500 mg/m<sup>3</sup>. The oral LD<sub>50</sub> presumably by gavage in corn oil or vegetable oil (data for mode of application and for vehicle sometimes not given) was 18 to 550 mg/kg bw. in rats showing a very steep dose/mortality-curve in individual studies. The dermal LD<sub>low</sub> in rabbits is 500 mg/kg bw. The most relevant route of exposure is inhalation. Therefore, the primary target tissues are the mucous membranes of mouth, eyes and respiratory tract. After oral exposure stomach ulceration is to be expected.</p> <p>Phosphorus trichloride is corrosive to the skin and mucous membranes of the eyes and the respiratory tract. Data on sensitisation for phosphorus trichloride have not been identified. The hydrolysis product hydrochloric acid gave no indication for a sensitising potential in humans and experimental animals. Data on phosphorous acid, the second hydrolysis product, are not available, but no specific effects are expected due to its structure.</p> <p>After 4-weeks of whole-body inhalation exposure to 0.5, 3, or 10 ppm (2.8, 17.1 or 56.8 mg/m<sup>3</sup>) phosphorus trichloride (6h/day, 5d/wk for 4 weeks), irritation of the eyes and the respiratory tract (suppurative inflammation, and inflammation and squamous metaplasia of respiratory epithelium) was observed in rats. There were no other symptoms not related to the irritant properties of phosphorus trichloride. The NOAEC in rats was 3 ppm (17 mg/m<sup>3</sup>). Chronic bronchitis can develop in humans. Repeated dose toxicity studies employing other routes were not identified in the literature. Because phosphorus trichloride as well as its hydrolysis products are toxicants acting at the portal of entry, and because phosphorus trichloride is unlikely to reach tissues distant from the portal of entry due to rapid hydrolysis, direct systemic toxicity is not likely to occur following exposure to phosphorus trichloride by any route.</p> <p>Phosphorus trichloride did not show mutagenic activity in a bacterial mutagenicity assay. Neither micronuclei nor chromosomal aberrations were induced in mouse bone marrow and human blood cells in vivo.</p> <p>As phosphorus trichloride decomposes to acid within seconds in aqueous media the resulting acidity of the hydrolysis products may cause unspecific effects of low pH in in-vitro tests. The change in pH may induce chromosomal aberrations and other DNA damage.</p> <p>In vivo, reduced pH levels could lead to chromosomal changes and DNA damage at the portal-of-entry of phosphorus trichloride. However, it is unlikely that systemic changes in pH would occur after exposure to phosphorus trichloride that are sufficient in magnitude to induce this effect in distant tissues or organs. The excess phosphate produced by hydrolysis of phosphorus trichloride may play a role in the development of effects on kidney, bone and calcium levels. Also by other routes (oral, dermal) phosphorous trichloride is expected to produce effects at the site of first contact (irritation, corrosion). The long term effects observed in humans (chronic bronchitis) are considered as sequelae of the irritation in the lungs which after prolonged periods may lead to an impairment of lung function (i.e.</p>	

oxygen availability).

No carcinogenicity studies with phosphorus trichloride were identified. The hydrolysis product hydrochloric acid (rats and mice) gave no indications for an increased tumour incidence after life-time exposure by inhalation. The other product of hydrolysis and subsequent partial neutralisation of phosphorus trichloride, mono sodium phosphite, gave also no indication of a carcinogenic potential after long term oral exposure. At low concentrations the hydrolysis products, phosphoric and hydrochloric acid, will be neutralized immediately in the physiologic medium at the portal of entry. Nevertheless prolonged irritation could give rise to a constant stimulus to local cell proliferation.

The repeated treatment of male animals with phosphorus trichloride via gavage or inhalation did not induce sperm morphology aberrations in rats and mice. There were no significant effects on intra-uterine development in rats. No malformations were detected. Skeletal development in treated fetuses was retarded but without a dose effect relation. The NOAEL was 19.3 mg/kg bw/day.

Due to the rapid hydrolysis it is unlikely that  $\text{PCl}_3$  could reach the reproductive organs or the embryo/fetus. At high concentrations major toxic effects (severe irritation and/or corrosion) on the parents are expected that could influence reproductive success. Specific toxicity to reproduction or developmental toxicity in mammals are not likely to occur following exposure to phosphorus trichloride by any route.

As, due to the corrosive nature of the substance, exposure is limited to the technically feasible extent in industrial settings, no consumer exposure is anticipated and it is unlikely that  $\text{PCl}_3$  could reach the reproductive organs, reproductive toxicity studies in animals are not warranted. No recommendation will result for further testing within the context of the SIDS program.

### Environment

Phosphorus trichloride is a moisture/water sensitive fluid with a melting point of  $-93.6\text{ }^\circ\text{C}$ , a boiling point of  $76.1\text{ }^\circ\text{C}$ , and a density of  $1.575\text{ g/cm}^3$  at  $20\text{ }^\circ\text{C}$ . The vapour pressure of the substance is  $129.7\text{ hPa}$  at  $20\text{ }^\circ\text{C}$ . The log  $K_{ow}$ , the water solubility and several other parameters cannot be determined due to hydrolysis. Phosphorus trichloride hydrolyzes completely in water with a  $t_{1/2}$  of less than 10 seconds at  $20\text{ }^\circ\text{C}$ , forming phosphonic acid and hydrochloric acid. In the atmosphere,  $\text{PCl}_3$  is oxidised by several photooxidants. Any emission into water, air, or the terrestrial compartment would be affected by humidity and also results in the formation of the hydrolysis products. Hydrochloric acid dissociates readily in water causing a pH shift which determines the impact of phosphoryl trichloride on aquatic life. The tolerance of water organisms towards pH is diverse. Recommended pH values for test species listed in OECD guidelines are between 6 and 9. Phosphonic acid ( $\text{pK}_a = 2.0$ )/ phosphoric acid ( $\text{pK}_a = 2.1$ ) is of medium acidity and partly dissociates in water causing a pH shift. Phosphoric acid and phosphates may affect aquatic life due to eutrophication.

Several aquatic toxicity tests have been undertaken in non-buffered solution. The observed toxicity effects in these studies can be attributed to the acidity of the degradation products and are not used for the hazard assessment. Acute toxicity of phosphorus trichloride (buffered) to fish (*Danio rerio*) tested according to the German guideline proposal "Lethal effects on Brachydanio rerio", was  $\geq 1000\text{ mg/l}$  (96 h-LC<sub>0</sub>, nominal concentration), which equals a LC<sub>0</sub> of  $\geq 597\text{ mg/l}$  of (buffered) phosphonic acid. With *Daphnia magna* an EC<sub>50</sub> (48 h) of  $> 100\text{ mg/l}$  in buffered solution was determined (92/69/EEC, method C.2). Algal toxicity was determined in a growth inhibition test with *Desmodesmus subspicatus*(92/69/EEC, method C.3). In buffered solution no effect was observed at  $100\text{ mg/l}$  (nominal). There are no results available on chronic toxicity. With activated sludge a 3 h-EC<sub>50</sub> of  $9450\text{ mg/l}$  (nominal) and an EC<sub>0</sub> of  $3520\text{ mg/l}$  (nominal) were measured according to the ISO 8192 (pH not reported).

There are test results available for acute testing from three trophic levels (all in buffered media). Using the lowest acute test result, a 48 h-EC<sub>50</sub> of  $\geq 100\text{ mg/l}$  (*Daphnia magna*, nominal concentration of buffered solution), and an assessment factor of 1000, a PNEC<sub>aqua</sub>  $\geq 0.1\text{ mg/l}$  was obtained.

### Exposure

In 2002, the global production capacity of phosphorus trichloride is estimated to be 0.8 million tonnes by about 20 producers. The global distribution of this capacity was approximately 0.5 million tonnes/year in OECD countries and 0.3 million tonnes /year in non-member countries.

Phosphorus trichloride is used as an intermediate for the manufacturing of wide range of chemicals (percentages

reported for the USA 2001):

- Pesticide intermediate (70 %)
- Phosphorus oxychloride (12 %)
- Surfactants and sequestrants, including phosphorus acid, used primarily for water treatment chemicals (11 %)
- Plastics additives, including flame retardants, plasticizers, phosphite antioxidants, and stabilizers (5 %)
- Miscellaneous, including lube oil and paint additives (2 %)

At one company in the Sponsor country phosphorus trichloride is manufactured and processed in closed systems. The exhausts from manufacturing and processing (including filling) of phosphorus trichloride are connected to air washing units. Thus, at this company, during production and processing virtually no phosphorus trichloride is emitted into the atmosphere. Due to water-free production, processing, and rapid hydrolysis phosphorus trichloride is not detectable in the wastewater.

In this company, the exposure of workers is well below the maximum admissible concentration of phosphorus trichloride in the workplace air (MAK) of 2.8 mg/m<sup>3</sup> (0.5 ppm). The exposure of workers to the hydrolysis product hydrochloric acid is also well below the MAK value of 8 mg/m<sup>3</sup> (5 ppm) for hydrogen chloride. Immunoglobulines against phosphorus trichloride have been detected.

No direct use of phosphorus trichloride is known. There is no exposure of consumers. Phosphorus trichloride is not listed in the Finnish, Norwegian and Swiss product registers. In the Swedish product register it is listed as an industrial chemical intermediate. Product register entry is confidential for Denmark. Phosphorus trichloride can be converted by chemical synthesis to nerve gases. Therefore the production and export of phosphorus trichloride is stringently controlled under the International Chemical Weapons Convention.

### RECOMMENDATION AND RATIONALE FOR THE RECOMMENDATION AND NATURE OF FURTHER WORK RECOMMENDED

**Human Health:** The chemical possesses properties indicating a hazard for human health (acute toxicity, corrosiveness). Based on data presented by the Sponsor country (relating to production by one producer which accounts for 1-6% of global production and relating to the use pattern in several OECD countries), exposure is limited to the technically feasible extent in occupational settings in the sponsor country. There is no exposure of consumers. No recommendation for further testing within the context of the SIDS program is therefore warranted. Although there are no valid data regarding reproductive effects, due to the fast hydrolysis it is unlikely that PCl<sub>3</sub> could reach organs and tissues distant from the site of first contact, therefore, and due to the corrosive properties studies in animals are not warranted. The chemical is currently of low priority for further work.

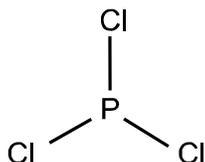
**Environment:** The chemical is currently of low priority for further work due to its low hazard profile. One of the degradation products, hydrochloric acid, has already been assessed within the OECD SIDS-Program.

## SIDS Initial Assessment Report

### 1 IDENTITY

#### 1.1 Identification of the Substance

CAS Number: 7719-12-2  
IUPAC Name: Phosphorus trichloride  
Molecular Formula:  $\text{PCl}_3$   
Structural Formula:



Molecular Weight: 137.33  
Synonyms: Chloride of phosphorus  
Phosphorus chloride  
Phosphorus trichloride  
Phosphorus(III)chloride  
Trichlorophosphine

#### 1.2 Purity/Impurities/Additives

Technical phosphorus trichloride has a purity of > 99.7 % w/w (Buechel, Moretto and Woditsch, 2000; Bayer AG, 2002). The following impurities have been reported:

- Phosphorus oxychloride  $\leq 0.3$  % w/w (Bayer AG, 2002)
- Distillation residue  $\leq 0.1$  % w/w (Bayer AG, 2002)
- Iron  $\leq 0.0001$  % w/w (Bayer AG, 2002)
- Arsenic 0.000002 % w/w (Bayer AG, 2002)

### 1.3 Physico-Chemical properties

**Table 1** Summary of physico-chemical properties

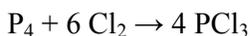
Property	Value	Reference	IUCLID
Substance type	Inorganic compound	Buechel, Moretto and Woditsch (2000)	1.1.1
Physical state	Colourless liquid, pungent odour	Bayer AG (2002)	1.1.1
Melting point	-93.6 °C	Riess (2002)	2.1
Boiling point at 1013 hPa	76.1 °C	Riess (2002)	2.2
Density at 20 °C	1.575 g/cm <sup>3</sup>	Riess (2002)	2.3
Vapour pressure at 20 °C	129.7 hPa	Riess (2002)	2.4
Conversion factors at 25 °C (calculated)	1 ppm = 5.6 mg/m <sup>3</sup> 1 mg/m <sup>3</sup> = 0.18 ppm	HSDB (2002)	2.14
Octanol/water partition coefficient (log Kow)	Not applicable*		2.5
Water solubility	Not stable in water due to hydrolysis*		2.6.1
pH value at 25 °C	Approximately 1 (at 5 g/l)*	Bayer Chemicals (2003)	2.14
Vapour density in relation to air	4.75	Sax (1979)	2.14

\*Rapid hydrolysis, cf. Chapter 2.2.3

## 2 GENERAL INFORMATION ON EXPOSURE

### 2.1 Production Volumes and Use Pattern

Phosphorus trichloride is manufactured by the strongly exothermic reaction of white phosphorus with chlorine



Phosphorus trichloride is manufactured in several ways:

- Chlorine is led into a suspension of phosphorus in phosphorus trichloride. Due to the heat released during the reaction, phosphorus trichloride evaporates. Phosphorus trichloride is condensed in reflux condensers and is partly returned into the phosphorus suspension.
- Phosphorus is burnt with stoichiometric amounts of chlorine.

Raw phosphorus trichloride is purified by fractional distillation (Buechel, Moretto and Woditsch, 2000).

In 1995 the phosphorus trichloride manufacturing capacities were about 0.2 million tonnes in the USA, 0.2 million tonnes in Western Europe, and 0.02 million tonnes in Japan. The phosphorus

trichloride consumption of the USA increased from 73 000 tonnes in 1983 to about 142 500 tonnes in 1994 (Buechel, Moretto and Woditsch, 2000). The US production capacity was reported to be about 315 000 tonnes and the US consumption to be about 277 000 tonnes in 2001 (TIG, 2004).

The global production capacity was estimated to be 0.8 million tonnes for about 20 producers in 2002. Approximately 0.5 million tonnes/year of the manufacturing capacity are in OECD countries and 0.3 million tonnes/year in non-member countries. In Western Europe there are 5 producers of phosphorus trichloride. Three of them have production plants in Germany. In 2003, Bayer manufactured 10 000 - 50 000 tonnes of phosphorus trichloride in the Bayer Leverkusen industrial park (Bayer Chemicals, 2004a).

Phosphorus trichloride is a basic chemical which is used industrially as an intermediate. Because of its reactivity phosphorus trichloride participates in a large number of chemical reactions (Greenwood and Earnshaw, 1988), e.g.

- Addition of sulfur
- Reaction with carbonic acids or acetic acid anhydrid
- Oxidation with oxygen or chlorine, to yield  $\text{POCl}_3$  or  $\text{PCl}_5$
- Substitutions with e.g. ammonia, amines, alcohols, thiols, esters
- Hydrolysis with water to produce phosphorus acid ( $\text{H}_3\text{PO}_3$ )
- Halogen ligand exchange with  $\text{ZnF}_2$ ,  $\text{PBr}_3$ , HI
- Friedel-Crafts-alkylations
- Grignard-reactions
- Formation of adducts e.g. with  $\text{BBr}_3$  or  $\text{Ni}(\text{CO})_4$ .

Due to these properties phosphorus trichloride is used as an intermediate for the manufacturing of wide range of chemicals (percentages reported for the USA 2001; TIG, 2004):

- Pesticide intermediate (70 %)
- Phosphorus oxychloride (12 %)
- Surfactants and sequestrants, including phosphorus acid, used primarily for water treatment chemicals (11 %)
- Plastics additives, including flame retardants, plasticizers, phosphite antioxidants, and stabilizers (5 %)
- Miscellaneous, including lube oil and paint additives (2 %)

No direct use is known (Bayer Chemicals, 2004a). Phosphorus trichloride is not listed in the Finnish, Norwegian (SPIN, 2004) and Swiss product registers (Swiss Product Register, 2003). In the Swedish product register, 5 products are mentioned to be used as industrial products for the manufacture of chemicals and chemical products in closed systems (SPIN, 2004). According to SPIN (2004) entries in the Danish Product Register are confidential.

Phosphorus trichloride can be converted by multistage chemical synthesis to nerve gases. Therefore the production and export of phosphorus trichloride is stringently controlled under the International Chemical Weapons Convention (1993). The Chemical Weapons Convention lists phosphorus trichloride as precursor to chemical weapons.

## 2.2 Environmental Exposure and Fate

### 2.2.1 Sources of Environmental Exposure

Information on exposure from manufacturing and processing of the chemical is available for the Bayer production plant at Leverkusen, Germany.

Phosphorus trichloride is manufactured, processed and filled in closed, waterfree systems (e.g. transport via pipeline, sampling without dead volume, gas shuttle pipe for filling processes). There is no direct wastewater in connection with the phosphorus trichloride production process itself. Cleaning of the reactors takes place only in the case of maintenance (Bayer Chemicals, 2004a).

The exhaust from manufacturing and processing of phosphorus trichloride is connected to a central gas washing unit. Water from the air washing unit is led to the industrial biological wastewater treatment plant. There is no detectable emission of phosphorus trichloride into the atmosphere. For this reason, phosphorus trichloride is not listed in the official Emission Declaration of Bayer in 2000 (Bayer Chemicals, 2004a).

Waste from the manufacturing and processing of phosphorus trichloride is incinerated in a incinerator for hazardous wastes equipped with an exhaust air cleaning device (Bayer Chemicals, 2004a).

The wastewater from the Bayer production plant is led to the Leverkusen industrial and municipal wastewater treatment plant. During the wastewater treatment at neutral pH (hydraulic retention time about 3 d) a rapid hydrolysis of phosphorus trichloride (half-life < 10 s, *cf.* Chapter 2.2.3) occurs. The hydrolysis products are neutralized (Bayer Chemicals, 2004a). Additionally, phosphorus acid is slowly oxidised by oxygen (air) to phosphoric acid (Merck, 2001). The pH value of the outlet is monitored continuously and so possible reduction of pH caused by phosphorus trichloride can be detected (Bayer Chemicals, 2004a).

The concentrated sewage sludge of the wastewater treatment plant is incinerated in a hazardous waste incinerator especially constructed for this sludge (Bayer Chemicals, 2004a).

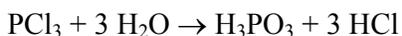
There is no information available on environmental exposure from production and use as synthesis intermediate at other manufacturing and processing sites. Because of the hydrolytic properties a relevant entry of phosphorus trichloride into the environment is unrealistic.

### 2.2.2 Photodegradation

Estimation of the photodegradation of phosphorus trichloride is not applicable by current assessment models due to the inorganic character of the substance. Due to its reactivity, it is expected that phosphorus trichloride will be oxidised by several agents in air (e.g. ozone, oxygen radicals). In aerosol phosphorus trichloride will be affected by air humidity, which leads to hydrolysis of the substance (*cf.* Chapter 2.2.3).

### 2.2.3 Stability in Water

In water, phosphorus trichloride hydrolyzes within seconds to phosphonic acid and hydrochloric acid (Melhem and Reid, 1998; Riess 2002). This reaction is the basis of the industrial production of phosphonic acid (Buechel, Moretto and Woditsch, 2000).



Phosphonic acid and phosphorus acid are tautomeric molecules (Roempp, 1999). Phosphorus acid is slowly oxidised by oxygen (air) to phosphoric acid (Merck, 2001).

Since HCl, which is formed in a ratio of 3:1 with regard to  $\text{H}_3\text{PO}_3$ , is a much stronger acid than  $\text{H}_3\text{PO}_3$  ( $\text{pK}_a$  -7.0 versus 2.0) all effects on pH shift are mainly caused by HCl.

The reaction of phosphorus trichloride and water was studied by adding small amounts of neat phosphorus trichloride (purity > 99 %) into an excess of well stirred water, and following the generation of acidic reaction products using a pH electrode. With this experimental set up it is not possible to distinguish the apparent reaction rate from, e.g., the mixing delay or the inertia of the measuring system. However, the half-life of phosphorus trichloride in water was estimated to be less than 10 seconds at 23 °C (Bayer Chemicals, 2004b).

This result is in line with several other studies. Mitchell (1925) observed that phosphorus trichloride released most of its acidity (98 %) within about 15 seconds. He suggests that intermediate stages of hydrolysis are  $\text{P}(\text{OH})_2\text{Cl}$  and  $\text{P}(\text{OH})_3$ .  $\text{P}(\text{OH})_2\text{Cl}$  reacts within seconds to  $\text{P}(\text{OH})_3$ . The tautomerisation of  $\text{P}(\text{OH})_3$  to  $\text{HP}(\text{O})(\text{OH})_2$  may take minutes to hours (Mitchell, 1925). Motorova and Noseko (1989) measured the rate of hydrolysis in dilute aqueous-organic solvent (dioxan) using calorimetric techniques. The half-life of phosphorus trichloride was found to be 1 - 13 seconds, for  $\text{PCl}_3$  concentrations between 0.61 and 1.8 mmol/l dioxan and water concentrations in the range 4 – 350 mmol/l dioxan.

In general, adding small amounts of phosphorus trichloride to water (molar ratio of water to phosphorus trichloride > 3), the hydrolysis products of phosphorus trichloride are phosphonic acid and hydrochloric acid (Melhem and Reid, 1998). If the molar ratio of water to phosphorus trichloride is between 2.5 and 3, a mixture of phosphorus and pyrophosphonic acid is formed. A molar ratio below 2.5 results in a product of indefinite composition, called lower oxides of phosphorus (Kirk-Othmer 1996). When the molar ratio of water to phosphorus trichloride is low, traces of phosphine may be formed in hot water (Geuther, 1872).

Thus, an environmental impact of phosphorus trichloride itself is not likely, but its impact is determined by the pH effect of the hydrolysis products. Phosphonic acid ( $\text{pK}_a = 2.0$ ) / phosphoric acid ( $\text{pK}_a = 2.1$ ) is of medium acidity and partly dissociates in water causing a pH shift. Since hydrochloric acid is a much stronger acid than phosphonic acid/phosphorus acid and their oxidation product phosphoric acid, the environmental effects of the hydrolysis products are due to hydrochloric acid. However, phosphoric acid and phosphates may affect aquatic life due to eutrophication.

For assessment of the environmental impact of the hydrolysis product hydrochloric acid, it is referred to the validated results of the hazard assessments on hydrochloric acid within the OECD SIDS-Program:

#### **Hydrochloric acid (CAS-No. 7647-01-0):**

Hydrochloric acid is a strong mineral acid, that dissociates readily in water to chloride ions and hydrated protons, and it is miscible with water. Dilute hydrochloric acid is nearly totally dissociated. This total ionisation also implies that hydrochloric acid will not adsorb on particulate matters or surfaces and will not accumulate in living tissues. For assessment of the environmental impact of hydrochloric acid it is referred to the validated results of the hazard assessments within the OECD SIDS-Program OECD, 2002).

### 2.2.4 Transport between Environmental Compartments

Since phosphorus trichloride hydrolyzes rapidly in water (*cf.* Chapter 2.2.3), no transfer coefficients can be measured.

### 2.2.5 Biodegradation

Since phosphorus trichloride hydrolyzes rapidly in water (*cf.* Chapter 2.2.3), no biodegradation can be measured. Hydrochloric acid is an inorganic degradation product which is not further degradable. Phosphonic acid is oxidized to phosphoric acid.

### 2.2.6 Bioaccumulation

Since phosphorus trichloride hydrolyzes rapidly in water (*cf.* Chapter 2.2.3), no BCF can be measured. Bioaccumulation of hydrochloric acid is not expected (OECD, 2002). Phosphonic acid is oxidized to phosphoric acid. Phosphoric acid is used by organisms as an essential nutrient.

### 2.2.7 Environmental Monitoring

No monitoring data available.

## 2.3 Human Exposure

Experience with human exposure is described in Chapter 5.10 of the IUCLID and Chapter 3.1.2 of this SIAR.

### 2.3.1 Occupational Exposure

#### Workplaces

During manufacturing and processing of phosphorus trichloride workers may be exposed, with the dermal and inhalational routes being the primary routes of exposure. In accordance with the principles of Responsible Care and Sustainable Development, at Bayer Chemicals the exposure of workers is reduced to the lowest technically practicable level (Bayer Chemicals, 2004a). Exposure information on other producing and/or processing sites in Germany is not available.

At the Bayer manufacturing site, workplaces where phosphorus trichloride is manufactured or processed in closed systems (Bayer Chemicals, 2004a), include

- Manufacturing processes: Conversion of P<sub>4</sub> with Cl<sub>2</sub> to phosphorus trichloride, distillation.
- Processing: on site in chemical synthesis, e.g. production of phosphorus oxichloride and phosphonates.

In the Bayer industrial park in Leverkusen most of the phosphorus trichloride is transported via pipeline. For transports to customers outside this industrial park, mostly railcars and ISO-containers (20 feet-containers) are used. A minor amount (less than 5 %) of phosphorus trichloride is transported in steel barrels (Bayer Chemicals, 2004a).

### Precautionary measures at the workplace

Surveys of the Bayer workplaces have been performed according to German Technical Guidance TRGS 402 (1997). This includes regular checks in the working area for any possible exposure to phosphorus trichloride and appropriate control measures (Bayer Chemicals, 2004a).

To protect workers several precautionary and protective measures are taken. These measures include technical equipment like suction devices at filling and sampling stations as well as appropriate personal protection equipment as prescribed in detail for different work situations e.g. during sampling, maintenance, and repair work. During sampling, for instance, gas filter masks, goggles, and rubber gloves have to be worn. Depending on the work to be done during maintenance, gas filter masks (classification ABEK) or a respirator with independent air supply have to be used as well as protective clothing (Bayer Chemicals, 2004a).

Down stream users of phosphorus trichloride are informed by way of a material safety data sheet on the recommended safety measures (see above, Bayer Chemicals, 2004a).

### Potential exposure at the workplace

The maximum admissible concentration of phosphorus trichloride in the workplace air (MAK) is 2.8 mg/m<sup>3</sup> (0.5 ppm) in Germany. Workplace air measurements of phosphorus trichloride were performed in the Bayer Chemicals manufacturing and processing units. In the manufacturing unit, 13 total shift measurements were done in the relevant areas between 1987 and 1993. 6 values of these (0.009 - 0.7 mg/m<sup>3</sup>) were above the detection limit (0.004 - 0.09 mg/m<sup>3</sup> depending on sampling conditions). Phosphorus trichloride was not detected in the other 7 samples. Phosphorus trichloride was also measured in 2 phosphorus trichloride processing units. 8 total shift values were measured between 1986 and 1996. 2 of these values (0.08 and 0.36 mg/m<sup>3</sup>) were above the limit of detection, the other 6 samples were below the limit of detection (0.04 - 0.6 mg/m<sup>3</sup> depending on sampling conditions). All results were below one third of the MAK value (Bayer Chemicals, 2004a).

Since there was no relevant exposure to phosphorus trichloride, the monitoring program was modified to include all compounds which release hydrochloric acid upon hydrolysis, e.g. phosphorus oxychloride. The MAK value of hydrochloric acid is 8 mg/m<sup>3</sup> (5 ppm). Between 1999 and 2003, eight hydrochloric acid measurements were performed in the manufacturing unit, and one in a processing unit. All results were below the limit of detection (0.8 mg/m<sup>3</sup>) (Bayer Chemicals, 2004a).

In general, the exposure of workers to phosphorus trichloride and to the hydrolysis product hydrochloric acid is negligible.

### Biological monitoring

In the framework of the Bayer occupational health surveillance program, the level of an immunoglobulin E (IgE) specific for phosphorus trichloride (and phosphorus oxychloride) was determined in the last 5 years (1999 - 2003) in about 900 workers routinely handling these substances. This specific IgE would indicate a possible sensitising effect of phosphorus chloride (and phosphorus oxychloride). With a detection limit of 0.35 kU, in the previous five years no specific IgE against phosphorus trichloride (and phosphorus oxychloride) was seen neither in the occupational surveillance program nor in any case of product contact. Phosphorus trichloride (and phosphorus oxychloride) appears to have no sensitisation potential (Bayer Industry Services, 2004).

### 2.3.2 Consumer Exposure

Phosphorus trichloride is exclusively used as an intermediate for chemical synthesis (*cf* Chapter 2.1). No direct use is known (Bayer Chemicals, 2004a). Phosphorus trichloride is not listed in the Finnish, Norwegian, (SPIN, 2004), and Swiss product registers (Swiss Product Register, 2003). For the Danish Product Register the entry is confidential (SPIN, 2004). The Swedish product register lists 5 products (industrial intermediates use in closed systems) with a total quantity of 950 t in 2000 (SPIN, 2004; Swedish Product Register, 2003).

In several products of the Sponsor company no phosphorus trichloride could be detected. To cover all chlorine containing compounds in products manufactured from phosphorus chloride, two phosphonates were analysed for chloride with a determination limit of about 1 mg/kg. No chloride could be quantified (Bayer Chemicals, 2004a).

Phosphorus trichloride can be converted by multistage chemical synthesis to nerve gases. Therefore the production and export of phosphorus trichloride is stringently controlled under the International Chemical Weapons Convention (1993). The Chemical Weapons Convention lists phosphorus trichloride as precursor to chemical weapons.

Thus, an exposure of consumers to phosphorus trichloride is unlikely to occur.

## 3 HUMAN HEALTH HAZARDS

### 3.1 Effects on Human Health

#### 3.1.1 Toxicokinetics, Metabolism and Distribution

Hydrolysis of phosphorus trichloride is mostly complete within 4-6 seconds in excess water. Products are hydrochloric acid (HCl) and phosphorous acid ( $H_3PO_3$ ). The rate of pH-change upon dissolution of phosphorus trichloride in water is comparable to the addition of concentrated hydrochloric acid. The change of pH of water was recorded during addition of phosphorus trichloride at room temperature. (Bayer Chemicals, 2004b)

At low concentrations the free acids resulting from the hydrolysis of phosphorus trichloride will be neutralised quickly by body fluids. The resulting chloride ions are natural components of food and ubiquitously found in living tissues, and are therefore not expected to pose a hazard. At high concentrations, which exceed the buffer capacity of body fluids the acids will damage the tissue at the portal of entry dependent upon concentration and duration of exposure. An availability of phosphorus trichloride or free acids in tissues distant from the portal of entry is hence not expected.

#### Studies in Animals and Humans

No studies in animals or humans were available in the literature

#### Conclusion

Phosphorus trichloride is quickly hydrolysed at first contact with water. It is, therefore, very unlikely that phosphorus trichloride will reach tissues distant from the portal of entry and become systemically available. The products of hydrolysis, hydrochloric acid and phosphorous acid, are also acting at the portal of entry.

### 3.1.2 Acute Toxicity

#### Studies in Animals

##### *Inhalation*

Toxicity after inhalation was determined in rats and guinea pigs by Weeks et al. (1964).

In rats the LC<sub>50</sub> of phosphorus trichloride was 104.3 ppm (592 mg/m<sup>3</sup>). Twenty female rats per group were exposed by whole body exposure. Animals were observed and deaths were recorded up to 14 days post exposure. Hydrolysis of phosphorus trichloride in the chamber atmosphere was about 40 percent. Animals showed signs of irritation (agitation, restlessness, irregular breathing, eyes closed, chromodakryorhea) during exposure to phosphorus trichloride. All deaths occurred within 10 days. Histopathology revealed necrosis in the epithelium and its supporting structures in the nostrils. Pulmonary damage was negligible. The kidney showed nephrosis of the tubules of the cortico-medullary region (Weeks et al., 1964).

In guinea pigs the LC<sub>50</sub> of phosphorus trichloride was 50.1 ppm (285 mg/m<sup>3</sup>) under similar conditions. Animals showed the same signs of irritation during exposure to phosphorus trichloride as described for rats (Weeks et al., 1964).

Monsanto (1983) reported a mean lethal concentration of 118 ppm (measured concentration as inorganic phosphorus corresponding to 673 mg/m<sup>3</sup>, corresponding to a nominal concentration of 453 ppm) after 4 hours “nose only” or “whole body” exposure in rats. Chamber analysis was based on inorganic phosphorus. Higher values were reported, if analysis was based on chloride or gravimetry. The mass median aerodynamic diameter was < 0.65 microns at the high doses and 1.75 microns at the lowest dose. Four groups of 5 male and 5 female Sprague-Dawley rats were exposed for 4 h to phosphorus trichloride by whole body as well as nose only exposure. Afterwards they were observed for 14 days. The LC<sub>50</sub> (4 h) was above 453 ppm (nominal). Three males and two female animals died. Clinical signs of intoxication were reduction of body weight, wheezing, laboured respiration, localised sores, swollen nose, blocked nostrils, and red crusts around the eyes. (Randall and Robinson, 1990; Monsanto, 1983a). Similar findings were also reported by Hoechst AG (1977) in an inhalation hazard test in rats.

Molodkina gave an LC<sub>50</sub> of 226 mg/m<sup>3</sup> for rats with similar symptoms as described by other authors like reduction of body weight, wheezing, laboured respiration, localised sores, swollen nose, blocked nostrils, agitation, pawing of nose, sedation, coordination disturbances, lateral position, fibrilar twitching, convulsions. Additionally, lacrimation, and corneal opacity were reported (Molodkina, 1974; Roshchin and Molodkina, 1977).

In an early report Butjagin (1904) described the effects of phosphorus trichloride on rabbits and cats.

Two to three rabbits per concentration range were exposed to phosphorus trichloride at concentrations of 4 - 30, 40 - 330, or 370 - 3870 ppm and observed for clinical signs for 6, 6 - 10, or 3 - 4 hours, respectively (whole body exposure). Low concentrations caused sedation and reduced respiratory frequency. The medium concentrations produced agitation, nasal discharge, rhinitis, sedation, reduced respiratory frequency, irregular respiration, corneal corrosion, dyspnea, tremor, lacrimation, and salivation. After the high concentration animals exhibited sneezing, agitation, closed eyes, white secretion from eyes, corneal opacity (starting at 118 min), weight loss and death. At necropsy animals of the medium dose group showed hyperemia of the trachea, slight lung oedema, and partial infiltration of right lung. Rabbits treated with the high concentration had severe lung oedema, emphysema, severe catarrh, and corrosion of the tongue and the cornea.

Although cats proved to be more sensitive the type of effects was identical to the findings in rabbits. The concentration range was 4 - 1080 ppm (23 - 6145 mg/m<sup>3</sup>) and animals were exposed for 6 hours. Salivation, nasal discharge, reduced, irregular respiratory frequency, dyspnea, cough, breathing through open mouth, lacrimation, and sedation were recorded. At higher doses immediate agitation, sneezing, rhinitis, conjunctivitis, loss of weight, and death occurred. Necropsy revealed discoloration and corrosion, of nasal cavity, injection and foam in the trachea, lung oedema, ecchymosis, corneal opacity, conjunctivitis, oedema of epiglottis, and severe emphysema (Butjagin, 1904).

#### *Dermal*

Dermal toxicity of phosphorus trichloride was determined in New Zealand White rabbits. The dose range was 1000 to 2000 mg/kg bw. No mortality was seen at 1000 mg/kg. Due to the low animal number an LD<sub>50</sub> could not be derived. The LD<sub>low</sub> was 1260 mg/kg bw. Animals showed reduced appetite and activity (3 to 15 days in survivors), increasing weakness, collapse, and death. At necropsy the lungs were hyperaemic, the livers discoloured, and the gall bladder enlarged. Kidneys and spleen were darkened and the gastrointestinal tract was slightly inflamed (Monsanto, 1977; Randall and Robinson, 1990).

A second study in rabbits afforded corrosion in animals treated with 250 mg/kg bw and death (LD<sub>low</sub>) in the one rabbit treated with 500 mg/kg bw (Mobil International 1977).

#### *Oral*

The oral LD<sub>50</sub> was determined as 550 mg/kg in a study using 6 rats per dose group. Signs of intoxication were nausea, disturbance of movement coordination, fatigue, weakness, chromodakryorrhea. The respiratory frequency was reduced to 50-80 per minute. Twenty to 40 minutes after application of the LD<sub>50</sub> cyanosis, weakness, convulsions, and dyspnea were observed. At necropsy the lungs of deceased animals were intensely red discoloured. The livers were dark-grey, the stomachs were distended and haemorrhagic (Molodkina, 1974; Roshchin and Molodkina, 1977).

Monsanto found a LD<sub>50</sub> of 18 mg/kg bw in Sprague Dawley rats. The dose range was 12.6 to 25.1 mg/kg bw. Doses of 12.6, 15.8, 20.0, or 25.1 mg/kg bw produced 0, 2, 3, 5 deaths, resp., in groups of 5 rats. Symptoms of toxicity were reduced appetite and activity (3 to 10 days in survivors), increasing weakness, collapse, and death. At necropsy the lungs were congested and the liver discoloured. Acute gastrointestinal inflammation with ulceration of the stomach was detected also in some survivors (Monsanto, 1977; Randall and Robinson, 1990).

A third study by Mobil employed male and female rats. Animals were fasted for 24 hours before administration of the test substance. The test material was administered orally, by intubation, as a 10 % solution in corn oil at doses of 100, 200, 300, 400, or 500 mg/kg bw. Animals were observed at 1, 3, 6, 24, 48, and 72 hours, then daily up to 14 days. The oral LD<sub>50</sub> was calculated as 200 ± 29 mg/kg bw. Signs of toxicity were decreased locomotor activity, piloerection, ptosis, wet underside and death. Normal body activity returned within 9 days in all surviving animals. All surviving animals were killed and autopsied. Necropsy revealed rugation of the pyloric mucosa, enlarged and darkened spleen, irregular thickening of the pyloric mucosa, stomach fused to liver or liver and pancreas, chronic pulmonary disease (Mobil International 1977a).

#### Studies in Humans

No studies but a number of case reports are available describing the effects after phosphorus trichloride exposure in humans.

### *Inhalation*

Workers employed in the production of phosphorous trichloride were exposed to concentrations of 10 - 20 mg/m<sup>3</sup> under normal condition and 80 - 150 mg/m<sup>3</sup> at times when the plant was out of order. In acute poisoning after 2 - 6 hours burning sensation in eyes and throat, photophobia, feeling of chest oppression, dry cough, and irritation of mucous membranes were reported. In subacute poisoning the symptoms occurred after 1 - 8 weeks with signs of irritation and asthmatic bronchitis (Sassi, 1952).

Patients accidentally exposed to phosphorus trichloride after a railroad accident reported of burning eyes (85 %), shortness of breath (59 %), throat irritation (59 %) and lacrimation (59 %). Headache and nausea (48 %), burning skin (44 %), increased sputum production (41 %), generalised or pleuric chest pain (33 %) and rash/itch (33 %) were also observed. Additionally, wheezing (26 %), blurred vision (22 %), vomiting (15 %) and abdominal pain (15 %) occurred. Lactic dehydrogenase was increased in 22 % of patients and recovered within four weeks after exposure. Pulmonary function tests showed some abnormalities in some patients (Wason et al., 1982; Wason et al., 1984).

Phosphorus trichloride caused irritation of eyes and mucous membranes in workers. Symptoms appeared immediately or delayed for up to 1 day. One patient died after several days due to an asthmatic fit caused by the exposure to phosphorus trichloride in concentrations less than 1 mg/l air. Rhinitis and conjunctivitis were accompanied by pain in nose and throat. At higher concentrations dyspnea and death followed (Reinl, 1956).

### Conclusion

The acute toxicity of phosphorus trichloride is high. It is characterised by immediate irritation/corrosion at the portal of entry in experimental animals and humans due to the irritant/corrosive properties of the products of hydrolysis. After inhalation (4h in some studies, unspecified in others) the LC<sub>50</sub> was determined in rats as 226 to > 500 mg/m<sup>3</sup>. The oral LD<sub>50</sub> presumably by gavage in corn oil or vegetable oil (data for mode of application and for vehicle sometimes not given) was 18 to 550 mg/kg bw. in rats showing a very steep dose/mortality-curve in individual studies. The dermal LD<sub>low</sub> was 500 mg/kg bw. The most relevant route of exposure is inhalation. Therefore, the primary target tissues are the mucous membranes of mouth, eyes and respiratory tract. After oral exposure stomach ulceration is to be expected.

### **3.1.3 Irritation**

#### Skin Irritation

##### *Studies in Animals*

To determine the skin irritation potential of phosphorus trichloride 100 µl of undiluted test substance were applied to the shaved skin of rabbits. The treated area was washed 60 seconds thereafter with water, Lutrol, or paraffin oil. Nevertheless, this exposure caused skin corrosion (Bayer, 1984).

Other authors similarly reported severely irritating or corrosive effects (Molodkina, 1974; Randall and Robinson, 1990; Radionova and Ivanov 1979).

Additionally signs of severe irritation at the site of first contact were reported in almost all studies aimed at systemic toxicity in animals.

## Eye Irritation

### *Studies in Animals*

Undiluted phosphorus trichloride was applied to the conjunctival sac of rabbits. The dose was 0.1 ml per eye. Phosphorus trichloride caused corrosion (Randall and Robinson, 1990). In an other study necrosis of the eye causing irreversible loss of vision is reported (Molodkina 1974).

Additionally signs of severe irritation at the site of first contact were reported in most studies aimed at systemic toxicity in animals.

### *Studies in Humans*

Phosphorus trichloride caused irritation of eyes and mucous membranes in workers. Symptoms appeared immediately or delayed for up to 1 day (Reinl, 1956, Weichardt 1957, Sassi, 1952, Wason et al., 1982; Wason et al., 1984).

## Respiratory Tract Irritation

### *Studies in Animals*

Symptoms of respiratory tract irritation are reported in almost all studies. The findings are described in the respective chapters(see chapters 3.1.2, 3.1.5)

### *Studies in Humans*

Phosphorus trichloride caused irritation of mucous membranes in workers. Symptoms appeared immediately or delayed for up to 1 day. Rhinitis and conjunctivitis were accompanied by pain in nose and throat. At higher concentrations dyspnea and death followed (Reinl 1956; Weichardt 1957; Zamakovskaya 1940; Sassi 1952; Wason et al. 1982; Wason et al. 1984).

## Conclusion

Phosphorus trichloride is corrosive to the skin and mucous membranes of the eyes and the respiratory tract.

### **3.1.4 Sensitisation**

No data on sensitisation potential for phosphorus trichloride were identified in the available literature. The hydrolysis product hydrochloric acid was tested in a Guinea Pig Maximisation Test (concentration of 1 %) and also in a Mouse Ear Swelling Test (concentrations of up to 5 %). Both tests gave no indication for a sensitising potential (Gad et al., 1986).

## Conclusion

Data on sensitisation potential for phosphorus trichloride were not identified. The hydrolysis product hydrochloric acid gave no indication for a sensitising potential in humans and experimental animals. Data on phosphorous acid, the second hydrolysis product, are not available, but no specific effects are expected due to its structure.

### **3.1.5 Repeated Dose Toxicity**

#### Studies in Animals

No studies employing oral or dermal exposure were identified.

### *Inhalation*

Fifteen male and 15 female Sprague-Dawley rats per group were exposed to phosphorus trichloride for 4 weeks (6h\*5 days\*4 weeks, whole body). The concentration range was 0 (air control), 0.5, 3.0, and 10.0 ppm (2,85 – 57 mg/m<sup>3</sup>). While the low concentrations did not produce any effects, the histopathological examination of the nasal cavity revealed squamous metaplasia of respiratory epithelium and focal suppurative inflammation of the anterior nasal region in the group exposed to 10 ppm. No other effects were recorded in clinical, clinical pathologic (haematology; clinical chemistry; urinalysis), and ophtalmoscopical examinations. Organ weights (absolute; relative to brain and body weight) and gross pathology findings were determined at necropsy. The NOAEC was 3 ppm (17 mg/m<sup>3</sup>) (Monsanto, 1983b).

A study conducted by CIIT employed F344/CrlBr and Sprague-Dawley rats at an age of 6 - 7 weeks. Thirtyone males and 21 females per group were exposed to hydrogen chloride whole body; (6h\*5d/w; 0, 10; 20 or 50 ml/m<sup>3</sup>) for 90 days. The parameters examined included: Clinical signs (daily), body weight (weekly), food consumption (weekly), urinalysis, hematology, clinical chemistry, necropsy, organ weights, histopathology [nasal turbinates, trachea, lung, brain, heart, kidney, liver, testis, adrenal, duodenum, eyes and optic nerve, mesenteric lymph nodes, aorta, sternum bone, ear canal, bone marrow, colon, epididymis, jejunum, mandibular lymph nodes, oviducts, ovaries, prostate, skin, pituitary glands, spinal cord, sciatic nerve, peripheral nerve, salivary gland, spleen, thyroid glands, urinary bladder, uterus, thymus, fore and glandular stomach, pancreas, parathyroid, skeletal muscle, seminal vesicle, tongue, femur bone, cecum, esophagus, ileum, lacrimal gland, mammary gland, larynx. NOAEC = 20 ppm (disregarding irritation). The treatment resulted in no mortality. Body weight (50 ppm) and food consumption (20 and 50 ppm) were reduced. Clinical pathology showed no effects in hematology, clin. chemistry, and urinalysis. At the histopathology examination rhinitis in the nasal cavity was detected in all treatment groups. The finding was accompanied with occasional hyperkeratosis (CIIT 1984).

### Studies in Humans

#### *Inhalation*

Workers employed in the production of phosphorus trichloride were exposed to concentrations of 10 - 20 mg/m<sup>3</sup> under normal condition and 80-150 mg/m<sup>3</sup> in cases of technical failures. After 1 - 8 weeks with signs of irritation and asthmatic bronchitis were reported (Sassi, 1952).

Symptoms reported by exposed workers in Russia were respiratory tract and eye irritation, cough, asthma, loss of voice (Zamakovhskaya, 1940).

Phosphorus trichloride is mentioned as a cause of occupational asthma. Forty eight of 170 (28.2 %) workers exposed to phosphorus chlorides (PCl<sub>3</sub>, PCl<sub>5</sub>, POCl<sub>3</sub>) are reported to have developed asthma (Buess and Lerner, 1956).

### Conclusion

After 4-weeks of whole-body inhalation exposure to 0.5, 3, or 10 ppm (2.8, 17.1 or 56.8 mg/m<sup>3</sup>) phosphorus trichloride (6h/day, 5d/wk for 4 weeks), irritation of the eyes and the respiratory tract (suppurative inflammation, and inflammation and squamous metaplasia of respiratory epithelium) was observed in rats.. Symptoms not related to the irritant properties of phosphorus trichloride were not detected. The NOAEC in rats was 3 ppm (17 mg/m<sup>3</sup>). Studies employing other routes were not identified in the literature. Chronic bronchitis may develop in humans. Repeated dose toxicity studies employing other routes were not identified in the literature. Because phosphorus trichloride as well as its hydrolysis products are toxicants acting at the portal-of-entry, and because phosphorus

trichloride is unlikely to reach tissues distant from the portal of entry due to rapid hydrolysis, direct systemic toxicity is not likely to occur following exposure to phosphorus trichloride by any route.

The excess phosphate produced by hydrolysis of phosphorus trichloride may play a role in the development of effects on kidney, bone and calcium levels. Also by other routes (oral, dermal) phosphorous trichloride is expected to produce effects at the site of first contact (irritation, corrosion). The long term effects observed in humans (chronic bronchitis) are considered as sequelae of the irritation in the lungs which after prolonged periods may lead to an impairment of lung function (i.e. oxygen availability).

### 3.1.6 Mutagenicity

#### *In vitro Studies*

Only two studies in bacteria could be identified in the available literature.

In a modified Ames-Test using *S. typhimurium* (TA1535, TA100, TA1537, D3052, TA1538, TA98, C3076, G46) and *E. coli* tester strains (WP2, WP2uvrA-) phosphorus trichloride had no mutagenic effects with and without metabolic activation by rat-S-9 mix (McMahon et al., 1979). *In vitro* studies with mammalian cells are not available.

*S. typhimurium* (TA1535, TA100, TA1537, TA1538, TA98) and *S. cerevisiae* (D4) were treated in a second study by Mobil with 0.001 to 5.0 µl of phosphorus trichloride per plate in the presence and absence of a metabolic activation system. No mutagenic effects were observed (Mobil International 1977b).

#### *In vivo Studies*

Male mice were injected 5 times intraperitoneally with phosphorus trichloride at doses of 10.94, 21.88, or 43.75 mg/kg and killed on day 6. The bone marrow was prepared for the determination of chromosomal aberrations and micronuclei. For chromosomal aberrations 100 cells were scored and for micro-nuclei 1000 polychromatic erythrocytes were examined. Phosphorus trichloride did not induce chromosomal aberrations or increases of the rate of micronuclei in mouse bone marrow cells (He et al., 1989).

#### Study in humans

Twenty-four workers exposed to phosphorus trichloride and 10 non-exposed were examined for chromosomal aberrations in peripheral lymphocytes. Phosphorus trichloride did not increase the incidence of chromosome aberrations (He et al., 1989).

#### Conclusion

Phosphorus trichloride did not show mutagenic activity in a bacterial mutagenicity assay. Neither micronuclei nor chromosomal aberrations were induced in mouse bone marrow and human blood cells *in vivo*.

As phosphorus trichloride decomposes to acid within seconds in aqueous media the resulting acidity of the hydrolysis products may cause unspecific effects of low pH in *in-vitro* tests. The change in pH may induce chromosomal aberrations and other DNA damage.

*In vivo*, reduced pH levels could lead to chromosomal changes and DNA damage at the portal-of-entry of phosphorus trichloride. However, it is unlikely that systemic changes in pH would occur after exposure to phosphorus trichloride, that are sufficient in magnitude to induce this effect in distant tissues or organs. The excess phosphate produced by hydrolysis of phosphorus trichloride

may play a role in the development of effects on kidney, bone and calcium levels. Also by other routes (oral, dermal) phosphorous trichloride is expected to produce effects at the site of first contact (irritation, corrosion). The long term effects observed in humans (chronic bronchitis) are considered as sequelae of the irritation in the lungs which after prolonged periods may lead to an impairment of lung function (i.e. oxygen availability).

### 3.1.7 Carcinogenicity

No carcinogenicity studies with phosphorus trichloride in experimental animals were identified in the available literature. As phosphorus trichloride hydrolyses quickly to form hydrochloric and phosphorous acids, chronic effects are expected mostly from exposure to these degradation products. Data are available only regarding hydrochloric acid/hydrogen chloride.

#### *Inhalation*

Albert et al. (1982) reported data from a chronic whole body inhalation exposure study with HCl in rats, discussed in detail by Sellakumar et al. (1985). One hundred male Sprague-Dawley rats were exposed to 10 ppm hydrogen chloride (HCl) for 6 hours/day, 5 days/week (duration-adjusted concentration = 2.5 mg/m<sup>3</sup>) for their lifetimes. All animals were observed daily, weighed monthly, and allowed to die naturally or killed when moribund. Complete necropsy was performed on all animals, with particular attention given to the respiratory tract. Histological sections were prepared from the nasal cavity (one lateral section from each side of the head), lung (one section from each lobe), trachea, larynx, liver, kidneys, testes, and other organs where gross pathological signs were present. However, Sellakumar et al. (1985) did not discuss histopathological events in organs other than the respiratory tract. HCl-exposed animals showed no differences in body weights or survival when compared with air controls. The data indicated 62/99 exposed animals with epithelial or squamous hyperplasia in the nasal mucosa (location not specified) vs. 51/99 in the concurrent control group. Incidence of squamous metaplasia was 9 and 5 in the exposed and control rats, respectively. There was increased hyperplasia of laryngeal-tracheal segments in HCl-exposed rats (larynx 22/99, trachea 26/99) vs. the controls (larynx 2/99, trachea 6/99). The authors did not make any comments concerning the severity of these changes. The tumour incidence in organs other than the respirator tract was similar in the treated and control groups. The total incidences of tumours at various sites being 19/99, 25/99 and 24/99 in treated, air control and colony control animals, respectively.

#### *Oral*

The repeated oral application of hydrochloric acid in mice gave no indication for an increased tumour incidence and also did not promote the activity of a known carcinogen. However, possibly only the gastro-intestinal tract was examined (Dyer, Kelly and Dunn, 1946).

The product of hydrolysis and subsequent partial neutralisation of phosphorus trichloride, mono sodium phosphite, was studied in rats. The US EPA IRIS data base reported a study with dietary treatment of rats with for 27 months, which gave no indication of a carcinogenic potential even at doses of 32000 ppm in the diet (US EPA IRIS, 2003; US EPA, 1986).

#### Studies in Humans

No data regarding carcinogenicity in humans were identified in the available literature.

#### Conclusion

No carcinogenicity studies with phosphorus trichloride were identified. The hydrolysis product hydrochloric acid (rats and mice) gave no clear indications for an increased tumour incidence in

the respiratory tract after life-time exposure by inhalation. The other product of hydrolysis and subsequent partial neutralisation of phosphorus trichloride, mono sodium phosphite, gave also no indication of a carcinogenic potential after long-term oral exposure. At low concentrations the hydrolysis products, phosphoric and hydrochloric acid, will be neutralized immediately in the physiologic medium at the portal of entry. Nevertheless prolonged irritation could give rise to a constant stimulus to local cell proliferation.

### 3.1.8 Toxicity for Reproduction

#### Studies on Fertility

##### *Studies in Animals*

He et al. describe studies on the effects of phosphorus trichloride on sperm morphology in rats and mice. Male mice were treated by gavage with phosphorus trichloride in vegetable oil for 5 days at doses of 45 - 178 mg/kg bw/day and the morphology of sperm cells was evaluated 4 weeks later. Rats were treated for 45 or 60 days at concentrations of 2, 33.5, and 97.76 mg/m<sup>3</sup>. No abnormalities of sperm cells were seen in rats and mice treated with phosphorus trichloride. (He et al., 1989)

#### Studies on Developmental toxicity

##### *Studies in Animals*

He et al. also treated pregnant rats with oral doses of 6.44, 9.7 and 19.3 mg/kg bw/day phosphorus trichloride on days 6 - 15 of pregnancy. The study also included a positive control group (substance described as: Dikushuang, the chinese name of a pesticide. The dose was 7 mg/kg bw). Body weights and reproductive indices (number of fetuses; corpora lutea, corpora lutea per rat, number of pups/litter, dead fetuses, % dead fetuses, resorptions, % resorptions, litters with resorptions, % litters with resorptions) were determined. Regarding the offspring the number of pups, size, tail length, weight of placenta and the size of the frontal fontanella were determined.

There were no significant differences between treated and negative control animals. No malformations were detected. Skeletal development in treated fetuses was retarded but without a dose effect relation. The NOAEL was 19.3 mg/kg bw. (He et al., 1989).

There were no studies identified for the hydrolysis products hydrochloric acid and phosphorous acid.

#### Conclusion

The repeated treatment of male animals with phosphorus trichloride via gavage or by inhalation did not induce sperm morphology aberrations in rats and mice.

There were no significant effects on intra-uterine development in rats. No malformations were detected. Skeletal development in treated fetuses was retarded but without a dose effect relation. The NOAEL was 19.3 mg/kg bw.

Due to the rapid hydrolysis it is unlikely that PCl<sub>3</sub> could reach the reproductive organs or the embryo/fetus. At high concentrations major toxic effects (severe irritation and/or corrosion) on the parents are expected that could influence reproductive success. Specific toxicity to reproduction or developmental toxicity in mammals are not likely to occur following exposure to phosphorus trichloride by any route.

As, due to the corrosive nature of the substance, exposure is limited to the technically feasible extent in industrial settings, no consumer exposure is anticipated and it is unlikely that PCl<sub>3</sub> could reach the reproductive organs, reproductive toxicity studies in animals are not warranted.

### 3.2 Initial Assessment for Human Health

Phosphorus trichloride is quickly hydrolysed at first contact with water. It is, therefore, very unlikely that phosphorus trichloride will reach tissues distant from the portal of entry and become systemically available. The products of hydrolysis, hydrochloric acid and phosphorous acid, are also acting at the portal of entry.

The acute toxicity of phosphorus trichloride is high. It is characterised by immediate irritation/corrosion at the portal of entry in experimental animals and humans due to the irritant/corrosive properties of the products of hydrolysis. After 4h-inhalation the LC<sub>50</sub> was determined in rats as 226 to > 500 mg/m<sup>3</sup>. The oral LD<sub>50</sub> presumably by gavage in corn oil or vegetable oil (data for mode of application and for vehicle sometimes not given) was 18 to 550 mg/kg bw. in rats showing a very steep dose/mortality-curve in individual studies. The dermal LD<sub>low</sub> was 1260 mg/kg bw. The most relevant route of exposure is inhalation. Therefore, the primary target tissues are the mucous membranes of mouth, eyes and respiratory tract. After oral exposure stomach ulceration is to be expected.

Phosphorus trichloride is corrosive to the skin and mucous membranes of the eyes and the respiratory tract.

Data on sensitisation potential for phosphorus trichloride were not identified. The hydrolysis product hydrochloric acid gave no indication for a sensitising potential in humans and experimental animals. Data on phosphorous acid, the second hydrolysis product, are not available, but no specific effects are expected due to its structure

After 4-weeks of whole-body inhalation exposure to 0.5, 3, or 10 ppm (2.8, 17.1 or 56.8 mg/m<sup>3</sup>) phosphorus trichloride (6h/day, 5d/wk for 4 weeks), irritation of the eyes and the respiratory tract (suppurative inflammation, and inflammation and squamous metaplasia of respiratory epithelium) was observed in rats.. Symptoms not related to the irritant properties of phosphorus trichloride were not detected. The NOAEC in rats was 3 ppm (17 mg/m<sup>3</sup>). Chronic bronchitis can develop in humans.

Repeated dose toxicity studies employing other routes were not identified in the literature. Because phosphorus trichloride as well as its hydrolysis products are toxicants acting at the portal of entry, and because phosphorus trichloride is unlikely to reach tissues distant from the portal of entry due to hydrolysis, direct systemic toxicity is not likely to occur following exposure to phosphorus trichloride by any route.

The excess phosphate produced by hydrolysis of phosphorus trichloride may play a role in the development of effects on kidney, bone and calcium levels. Also by other routes (oral, dermal) phosphorous trichloride is expected to produce effects at the site of first contact (irritation, corrosion). The long term effects observed in humans (chronic bronchitis) are considered as sequelae of the irritation in the lungs which after prolonged periods may lead to an impairment of lung function (i.e. oxygen availability).

Phosphorus trichloride did not show mutagenic activity in a bacterial mutagenicity assay. Neither micronuclei nor chromosomal aberrations were induced in mouse bone marrow and human blood cells in vivo.

As phosphorus trichloride decomposes to acid within seconds in aqueous media the resulting acidity of the hydrolysis products may cause unspecific effects of low pH in in-vitro tests. The change in pH may induce chromosomal aberrations and other DNA damage.

In vivo, reduced pH-levels could lead to chromosomal changes and DNA damage at the portal-of-entry of phosphorus trichloride. However, it is unlikely that systemic changes in pH would occur

after exposure to phosphorus trichloride, that are sufficient in magnitude to induce this effect in distant tissues or organs.

No carcinogenicity studies with phosphorus trichloride were identified. The hydrolysis product hydrochloric acid (rats and mice) gave no clear indications for an increased tumour incidence after life-time exposure by inhalation. The other product of hydrolysis and subsequent partial neutralisation of phosphorus trichloride, mono sodium phosphite, gave also no indication of a carcinogenic potential after long term oral exposure.

At low concentrations the hydrolysis products, phosphoric and hydrochloric acid, will be neutralized immediately in the physiologic medium at the portal of entry. Nevertheless prolonged irritation could give rise to a constant stimulus to local cell proliferation.

The repeated treatment of male animals with phosphorus trichloride via gavage or by inhalation did not induce sperm morphology aberrations in rats and mice.

There were no significant effects on intra-uterine development in rats. No malformations were detected. Skeletal development in treated fetuses was retarded but without a dose effect relation. The NOAEL was 19.3 mg/kg bw

Phosphorus trichloride is a toxicant acting at the portal-of-entry. It is quickly hydrolysed in aqueous media (half-life < 10 seconds) and the resulting free acids will be neutralised immediately at low concentrations. At high concentrations the acids will damage the tissue at the site of first contact. Therefore it is unlikely that phosphorus trichloride could reach the reproductive organs or the embryo/fetus. At high concentrations major toxic effects (severe irritation and/or corrosion) on the parents are expected that could influence reproductive success. Specific toxicity to reproduction or developmental toxicity in mammals are not likely to occur following exposure to phosphorus trichloride by any route.

As, due to the corrosive nature of the substance, exposure is limited to the technically feasible extent in industrial settings, no consumer exposure is anticipated and it is unlikely that  $\text{PCl}_3$  could reach the reproductive organs, reproductive toxicity studies in animals are not warranted.

## **4 HAZARDS TO THE ENVIRONMENT**

### **4.1 Aquatic Effects**

In water, phosphorus trichloride hydrolyzes to phosphonic acid and hydrogen chloride. The experimentally determined half-life is < 10 seconds (*cf.* Chapter 2.2.3). Right from the start of the test, ecotoxicological measurements as described in chapter 4.1 will cover the effects of the degradation products phosphonic acid and hydrogen chloride.

The hydrolysis product hydrochloric acid is tested with aquatic species (OECD, 2002). Hydrochloric acid causes a pH shift in water (Table 2). It is the resulting pH that determines the impact of hydrogen chloride on aquatic life as shown with buffered test substance solution. Thus toxic effects are not due to substance inherent properties but caused by the low pH (OECD, 2002).

Some experiments with phosphorus trichloride were performed in the presence of buffer to avoid the pH effects of the acids formed by hydrolysis of phosphorus trichloride. Comparison of experiments in the presence and absence of buffer (with and without neutralisation) confirmed the conclusions drawn from the OECD-SIDS Hydrochloric Acid (2002; see below). Regarding natural systems, the impact of dissociated acids depends on the buffer capacity of the system. Buffer

function is attributed to humic substances, alkaline earth carbonates, clay minerals, silicates, as well as amphoteric oxides.

**Table 2 Theoretical pH-values of hydrochloric acid in non-buffered water**

Hydrochloric acid concentration (mg/l)	Corresponding phosphorus trichloride concentration (mg/l)	pH
0.036	0.046	6
0.36	0.46	5
3.6	4.7	4
36	46	3

The tolerance of water organisms towards pH is diverse. pH-values recommended in OECD guidelines for testing issues are compiled in Table 3.

**Table 3 pH values recommended in OECD guidelines for testing issues**

Group (Trophic level)	Recommendation
Fish	pH 6.0 to pH 8.5 is preferable
<i>Daphnia</i>	within the range of pH 6 to pH 9
Algae	approximately pH 8

### Acute Toxicity Test Results

Short term tests on aquatic toxicity are available for each trophic level (Table 4).

Acute toxicity to fish (*Danio rerio*) was tested in a static test system according to the method proposal of the German Environmental Protection Agency “Lethal effects on *Brachydanio rerio*”. Phosphorus trichloride was not monitored because it hydrolyzes. A limit test was conducted with an adjusted pH (pH ca. 7.5) value at 1000 mg/l (nominal concentration). During 96 h no effects were observed at the tested concentration level, and a LC<sub>0</sub> of ≥ 1000 mg/l was determined for phosphorus trichloride, which equals a LC<sub>0</sub> of ≥ 597 mg/l of (neutralized) phosphonic acid as the hydrolysis product (Bayer AG, 1991).

In contrast, acute toxicity was found in a study without adjustment of pH. This test was not conducted according to any guideline (Gurova, Krasnov and Mazmanidi, 1970). In the test media (dechlorinated tap water and Wolga water) pH values are assumed to vary from 3.3 to 7 (information not given for the acute study but for the long-term study of the same authors performed with the same concentration range as described below.) (*cf.* Table 3). The 3 d-NOEC (= LC<sub>0</sub>) was found to be about 60 mg/l (LC<sub>100</sub>: 75 mg/l) for sturgeon eggs (*Acipenser stellatus*). The hatching success of the fish larvae was reduced by about 10 % at 60 and 70 mg/l. Growth of hatchlings was not tested at 70 mg/l, as all hatchlings showed abnormalities at this concentration. With regard to length of the hatchlings after 5 days, a NOEC of 20 mg/l was observed. A small reduction (5 %) of fish larvae weight was observed at the lowest concentration tested (20 mg/l). In an insufficiently described experiment, dace (*Leuciscus leuciscus*) were more sensitive to phosphorus trichloride and its degradation products, respectively, and a 10 d-LC<sub>100</sub> of 25 mg/l was observed. However, from this study, no EC<sub>50</sub> can be derived (Gurova, Krasnov and Mazmanidi, 1970). A test on prolonged toxicity of phosphorus trichloride to 3 fish species was performed by

Gurova, Krasnov, and Mazmanidi (1970) with the same conditions as above. In 30 d tests with the 3 fish species *Carassius carassius*, *Perca fluviatilis*, and *Esox lucius*, NOEC values of 40-50 mg/l were found (Gurova, Krasnov and Mazmanidi, 1970) in non-buffered media.

With the invertebrate *Daphnia magna* one acute test according to the European guideline 92/69/EEC, method C.2, is available. In non-buffered test solution the pH decreased from pH 7.9 in the controls to about pH 6.7 at 12.5 mg/l, pH 6.1 at 25 mg/l, pH 3.6 at 50 mg/l, and pH 2.9 at 100 mg/l. For a test period of 24 h an EC<sub>0</sub> (immobilisation) of 25 mg/l, an EC<sub>50</sub> of 35.4 mg/l, and an EC<sub>100</sub> of 50 mg/l were obtained. The same effect concentrations were reported after a test period of 48 hours. In buffered test solution no effect was observed at the highest tested concentration of 100 mg/l of phosphorus trichloride, suggesting that the effects in the non-buffered solutions were solely due to the pH decrease (Bayer AG, 2003a).

Algal toxicity was determined by a test with *Desmodesmus subspicatus* in the presence of phosphorus trichloride and its hydrolysis products. In a growth inhibition test according the European guideline 92/69/EEC, method C.3, equivalent to OECD TG 201, in non-buffered solution, the pH depended on the nominal phosphorus trichloride concentration and was pH 8.2 in the controls and pH 2.9 at 100 mg/l (nominal concentration at the start of the incubation period). In non-buffered solution, a 72 h-E<sub>r</sub>C<sub>50</sub> of 33 mg/l was determined for growth rate (population density) and a 72 h-E<sub>b</sub>C<sub>50</sub> of 30 mg/l for growth (integral of biomass). The 72h-NOEC was 12.5 mg/l for both growth rate and biomass. In buffered solution no effect was observed at the highest phosphorus trichloride concentration tested (nominal 100 mg/l). Therefore, it can be concluded that the effects found in this study are caused by pH effects (Bayer AG, 2003b).

#### Chronic Toxicity Test Results

There are no results available on chronic toxicity.

#### Toxicity to Microorganisms

A test with activated sludge with a duration of 3 h was conducted according to the ISO 8192 (Test for the Inhibition of Oxygen Consumption by Activated Sludge). The inoculum contained 6 g of dry matter per litre (the pH was not reported). An EC<sub>50</sub> of 9450 mg/l and an EC<sub>0</sub> of 3520 mg/l were determined (Bayer AG, 1991).

**Table 4 Aquatic toxicity of phosphorus trichloride and its hydrolysis products**

Species	Endpoint	Eposure regime	Duration	Effect concentration	Reference	IUCLID
Fish						
<i>Danio rerio</i>	Mortality	Static	96 h-LC <sub>50</sub>	>= 1000 mg/l (n)*	Bayer AG, 1991	4.1
<i>Acipenser stellatus</i> , eggs	Mortality (hatching success)	Semistatic	3 d-NOEC (= LC <sub>0</sub> ) 3 d-LC <sub>10</sub> 3 d-LC <sub>100</sub>	60 mg/l (n) >60-70 mg/l (n) 75 mg/l (n)	Gurova, Krasnov and Mazmanidi 1970	4.1
<i>Acipenser stellatus</i> , hatchlings	Growth (weight, length)	Semistatic	EC <sub>10</sub> NOEC	60 mg/l (n) 20 mg/l (n)	Gurova, Krasnov and Mazmanidi 1970	4.1
<i>Leuciscus leuciscus</i>	Mortality	Semistatic	10 d-LC <sub>100</sub>	25 mg/l (n)	Gurova, Krasnov, and Mazmanidi 1970	4.1
<i>Carassius carassius</i> , <i>Perca fluviatilis</i> , and <i>Esox lucius</i>	Mortality and Growth (weight)	Semistatic	30 d-NOEC	40-50 mg/l (n)	Gurova, Krasnov, and Mazmanidi 1970	4.5.1
Invertebrates						
<i>Daphnia magna</i>	Immobility	Static	48 h-EC <sub>0</sub> 48 hEC <sub>50</sub> 48 h-EC <sub>100</sub> 48 h-EC <sub>50</sub>	25 mg/l (n) 35.4 mg/l (n) 50 mg/l (n) >= 100 mg/l (n)*	Bayer AG 2003a	4.2
Algae						
<i>Desmodesmus subspicatus</i>	Growth	Static	72 h-E <sub>r</sub> C <sub>50</sub> 72 h-E <sub>b</sub> C <sub>50</sub> 72 h-EC <sub>50</sub>  72 h-NOEC (for both population growth and biomass) 72 h-NOEC	33 mg/l (n) 30 mg/l (n) >= 100 mg/l (n)*  12.5 mg/l (n)  >= 100 mg/l (n)*	Bayer AG 2003b	4.3
Activated Sludge	Respiration inhibition	Static	3 h-EC <sub>50</sub>	9450 mg/l (n)	Bayer AG 1991	4.4

(n) = nominal concentration; \*buffered test medium

#### Determination of PNEC<sub>aqua</sub>

For phosphorus trichloride and its hydrolysis products, the lowest acute effect concentration was found for the dace (*Leuciscus leuciscus*) with a 10 d-EC<sub>100</sub> = 25 mg/l in a non-guideline study (Gurova, Krasnov and Mazmanidi, 1970). In this study, non-buffered media were used and the pH was presumably 3.3. As has been shown by several other studies (OECD,2002), this effect was due to the pH which was outside the pH range tolerated by fish (*cf.* Table 2 and 3). Other tests in non-buffered media showed similarly that effects depended on the pH (Bayer AG 1991, Bayer AG 2003a, b). However, there are acute test results available from test with buffered test media at three

trophic levels. In this series of experiments, the lowest acute effect concentration is a 48 h-EC<sub>50</sub> of  $\geq 100$  mg/l (n) of *Daphnia magna*. Applying an assessment factor of 1000 according to the EU Technical Guidance Document results in a

$$\text{PNEC}_{\text{aqua}} > 0.1 \text{ mg/l.}$$

Due to the fast hydrolysis this PNEC<sub>aqua</sub> covers also the hydrolysis products hydrochloric acid and phosphonic acid.

#### 4.2 Terrestrial Effects

No data available.

#### 4.3 Other Environmental Effects

No data available.

#### 4.4 Initial Assessment for the Environment

Phosphorus trichloride is a moisture/water sensitive fluid with a melting point of -93.6 °C, a boiling point of 76.1 °C, and a density of 1.575 g/cm<sup>3</sup> at 20 °C. The vapour pressure of the substance is 129.7 hPa at 20 °C. The log K<sub>ow</sub>, the water solubility and several other parameters cannot be determined due to hydrolysis. Phosphorus trichloride hydrolyzes completely in water with a t<sub>1/2</sub> of less than 10 seconds at 20 °C, forming phosphonic acid and hydrochloric acid. In the atmosphere, PCl<sub>3</sub> is oxidised by several photooxidants. Any emission into water, air, or the terrestrial compartment would be affected by humidity and also results in the formation of the hydrolysis products. Hydrochloric acid dissociates readily in water causing a pH shift which determines the impact of phosphoryl trichloride on aquatic life.. The tolerance of water organisms towards pH is diverse. Recommended pH values for test species listed in OECD guidelines are between 6 and 9. Phosphonic acid (pK<sub>a</sub> = 2.0)/ phosphoric acid (pK<sub>a</sub> = 2.1) is of medium acidity and partly dissociates in water causing a pH shift. Phosphoric acid and phosphates may affect aquatic life due to eutrophication.

Several aquatic toxicity tests have been undertaken in non-buffered solution. The observed toxicity effects in these studies can be attributed to the acidity of the degradation products and are not used for the hazard assessment. Acute toxicity of phosphorus trichloride (buffered) to fish (*Danio rerio*) tested according to the German guideline proposal “Lethal effects on Brachydanio rerio”, was  $\geq 1000$  mg/l (96 h-LC<sub>0</sub>, nominal concentration), which equals an LC<sub>0</sub> of  $\geq 597$  mg/l of (buffered) phosphonic acid. With *Daphnia magna* an EC<sub>50</sub> (48 h) of  $> 100$  mg/l in buffered solution was determined (92/69/EEC, method C.2). Algal toxicity was determined in a growth inhibition test with *Desmodesmus subspicatus* (92/69/EEC, method C.3). In buffered solution no effect was observed at 100 mg/l (nominal). There are no results available on chronic toxicity. With activated sludge a 3 h-EC<sub>50</sub> of 9450 mg/l (nominal) and an EC<sub>0</sub> of 3520 mg/l (nominal) were measured according to the ISO 8192 (pH not reported).

There are test results available for acute testing from three trophic levels (all in buffered media). Using the lowest acute test result, a 48 h-EC<sub>50</sub> of  $\geq 100$  mg/l (*Daphnia magna*, nominal concentration of buffered solution), and an assessment factor of 1000, a PNEC<sub>aqua</sub>  $> 0.1$  mg/l was obtained.

## 5 RECOMMENDATIONS

The chemical is currently of low priority for further work.

### **Human Health:**

The chemical possesses properties indicating a hazard for human health (acute toxicity, corrosiveness). Based on data presented by the Sponsor country (relating to production by one producer which accounts for 1-6% of global production and relating to the use pattern in several OECD countries), exposure is limited to the technically feasible extent in occupational settings in the sponsor country. There is no exposure of consumers. No recommendation for further testing within the context of the SIDS program is therefore warranted. Although there are no valid data regarding reproductive effects, due to the fast hydrolysis it is unlikely that  $\text{PCl}_3$  could reach organs and tissues distant from the site of first contact, therefore, and due to the corrosive properties further studies in animals are not warranted. The chemical is currently of low priority for further work.

### **Environment:**

The chemical is currently of low priority for further work due to its low hazard profile. One of the degradation products, hydrochloric acid, has already been assessed within the OECD SIDS-Program.

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# I U C L I D

## Data Set

**Existing Chemical** : ID: 7719-12-2  
**CAS No.** : 7719-12-2  
**EINECS Name** : phosphorus trichloride  
**EC No.** : 231-749-3  
**TSCA Name** : Phosphorous trichloride  
**Molecular Formula** : Cl<sub>3</sub>P

**Producer related part**

**Company** : Bayer AG  
**Creation date** : 22.09.1992

**Substance related part**

**Company** : Bayer AG  
**Creation date** : 22.09.1992

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**Flags (profile)** : Flags: without flag, non confidential, WGK (DE), TA-Luft (DE), Material Safety Dataset, Risk Assessment, Directive 67/548/EEC, SIDS

**1.0.1 APPLICANT AND COMPANY INFORMATION****1.0.2 LOCATION OF PRODUCTION SITE, IMPORTER OR FORMULATOR****1.0.3 IDENTITY OF RECIPIENTS****1.0.4 DETAILS ON CATEGORY/TEMPLATE****1.1.0 SUBSTANCE IDENTIFICATION**

**IUPAC Name** : Phosphorus trichloride  
**Smiles Code** : ClP(Cl)Cl  
**Molecular formula** : Cl<sub>3</sub>P  
**Molecular weight** : 137.33  
**Petrol class** :

**Flag** : Critical study for SIDS endpoint  
 17.10.2003 (1)

**1.1.1 GENERAL SUBSTANCE INFORMATION**

**Purity type** : typical for marketed substance  
**Substance type** : inorganic  
**Physical status** : liquid  
**Purity** : > 99.7 % w/w  
**Colour** :  
**Odour** :

**Flag** : Critical study for SIDS endpoint  
 09.02.2004 (2)

**Purity type** : typical for marketed substance  
**Substance type** : inorganic  
**Physical status** : liquid  
**Purity** : = 99.7 % w/w  
**Colour** : colourless  
**Odour** : pungent

**Method** : Purity: Method CH-P ELS 31.91 GC-WLD  
**Flag** : Critical study for SIDS endpoint  
 27.01.2004 (3)

**Purity type** : typical for marketed substance  
**Substance type** : inorganic  
**Physical status** : liquid  
**Purity** : ca. 99 % w/w  
**Colour** : colourless  
**Odour** : pungent

09.02.2004 (4) (5)

**1.1.2 SPECTRA****1.2 SYNONYMS AND TRADENAMES****Chloride of phosphorus**

17.10.2003 (6) (7)

**Phosphorus chloride**

24.09.2003 (6) (7)

**Phosphorus trichloride**

31.07.2003 (6) (1) (7)

**Phosphorus(III)chloride**

31.07.2003 (7)

**Trichlorophosphine**

24.09.2003 (6)

**1.3 IMPURITIES**

**Purity** : typical for marketed substance  
**CAS-No** : 10025-87-3  
**EC-No** : 233-046-7  
**EINECS-Name** : phosphoryl trichloride  
**Molecular formula** : POCl3  
**Value** : < 1 % w/w

**Flag** : Critical study for SIDS endpoint  
 09.02.2004 (5)

**Purity** : typical for marketed substance  
**CAS-No** : 10025-87-3  
**EC-No** : 233-046-7  
**EINECS-Name** : phosphoryl trichloride  
**Molecular formula** : POCl3  
**Value** : <= .3 % w/w

**Flag** : Critical study for SIDS endpoint  
 09.02.2004 (3)

**Purity** : typical for marketed substance  
**CAS-No** :  
**EC-No** :  
**EINECS-Name** : Distillation residue  
**Molecular formula** :  
**Value** : <= .1 % w/w

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<b>Flag</b> 09.02.2004	:	Critical study for SIDS endpoint	(3)
<b>Purity</b>	:	typical for marketed substance	
<b>CAS-No</b>	:	7439-89-6	
<b>EC-No</b>	:	231-096-4	
<b>EINECS-Name</b>	:	iron	
<b>Molecular formula</b>	:		
<b>Value</b>	:	<= .001 g/kg	
<b>Remark</b>	:	Value given as <= 1 mg/kg	
<b>Flag</b> 09.02.2004	:	Critical study for SIDS endpoint	(3)
<b>Purity</b>	:	typical for marketed substance	
<b>CAS-No</b>	:	7440-38-2	
<b>EC-No</b>	:	231-148-6	
<b>EINECS-Name</b>	:	arsenic	
<b>Molecular formula</b>	:	As	
<b>Value</b>	:	= 0 % w/w	
<b>Remark</b>	:	Value given as = 0.02 mg/kg	
<b>Flag</b> 09.02.2004	:	Critical study for SIDS endpoint	(3)

## 1.4 ADDITIVES

## 1.5 TOTAL QUANTITY

<b>Quantity</b>	:	- tonnes in 1995	
<b>Remark</b>	:	In 1995, the phosphorus trichloride manufacturing capacities were about 0.2 million tonnes in the USA, 0.2 million tonnes in Western Europe, and 0.02 million tonnes in Japan. The phosphorus trichloride consumption of the USA increased from 73,000 tonnes in 1983 to about 142,500 tonnes in 1994	
<b>Flag</b> 14.07.2005	:	Critical study for SIDS endpoint	(2)
<b>Quantity</b>	:	- tonnes in 2001	
<b>Result</b>	:	The US production capacity is reported to be about 315,000 tonnes and the US consumption to be about 277,000 tonnes in 2001	
<b>Flag</b> 14.07.2005	:	Critical study for SIDS endpoint	(8)
<b>Quantity</b>	:	- tonnes in 2002	
<b>Result</b>	:	The global production capacity was estimated to be 0.8 million tonnes for about 20 producers in 2002. Approximately 0.5 million tonnes/a of the manufacturing capacity are in OECD countries and 0.3 million tonnes/a in non-member countries. In Western Europe there are 5 producers of phosphorus trichloride. Three of them have production plants in Germany in 2002	
<b>Flag</b> 14.07.2005	:	Critical study for SIDS endpoint	(9)

**1.6.1 LABELLING**

<b>Labelling</b>	:	as in Directive 67/548/EEC	
<b>Specific limits</b>	:	no	
<b>Symbols</b>	:	T+, C, ,	
<b>Nota</b>	:	, ,	
<b>R-Phrases</b>	:	(14) Reacts violently with water (26/28) Very toxic by inhalation and if swallowed (35) Causes severe burns (48/20) Harmful: danger of serious damage to health by prolonged exposure through inhalation	
<b>S-Phrases</b>	:	(1/2) Keep locked up and out of reach of children (7/8) Keep container tightly closed and dry (26) In case of contact with eyes, rinse immediately with plenty of water and seek medical advice (36/37/39) Wear suitable protective clothing, gloves and eye/face protection (45) In case of accident or if you feel unwell, seek medical advice immediately (show the label where possible)	
<b>Remark</b>	:	Index-No. 015-007-00-4 EG-No. 231-749-3 Substance is dealt with in the 25th ATP. As an editorial mistake R29 is missing in the labelling. Will be corrected in the 29th ATP which is not yet published.	
25.03.2004			(10)

**1.6.2 CLASSIFICATION**

<b>Classified</b>	:	as in Directive 67/548/EEC	
<b>Class of danger</b>	:		
<b>R-Phrases</b>	:	(14) Reacts violently with water	
<b>Specific limits</b>	:		
17.08.2004			(10)
<b>Classified</b>	:	as in Directive 67/548/EEC	
<b>Class of danger</b>	:		
<b>R-Phrases</b>	:	(29) Contact with water liberates toxic gas	
<b>Specific limits</b>	:		
18.12.2003			(10)
<b>Classified</b>	:	as in Directive 67/548/EEC	
<b>Class of danger</b>	:	very toxic	
<b>R-Phrases</b>	:	(26/28) Very toxic by inhalation and if swallowed	
<b>Specific limits</b>	:		
18.12.2003			(10)
<b>Classified</b>	:	as in Directive 67/548/EEC	
<b>Class of danger</b>	:	harmful	

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**R-Phrases** : (48/20) Harmful: danger of serious damage to health by prolonged exposure through inhalation

**Specific limits** :

18.12.2003 (10)

**Classified** : as in Directive 67/548/EEC

**Class of danger** : corrosive

**R-Phrases** : (35) Causes severe burns

**Specific limits** :

18.12.2003 (10)

**1.6.3 PACKAGING****1.7 USE PATTERN**

**Type of use** : type  
**Category** : Use in closed system

09.02.2004 (5)

**Type of use** : industrial  
**Category** : Chemical industry: used in synthesis

17.12.2003 (5)

**Type of use** : use  
**Category** : Intermediates

17.12.2003 (5)

**1.7.1 DETAILED USE PATTERN****1.7.2 METHODS OF MANUFACTURE****1.8 REGULATORY MEASURES****1.8.1 OCCUPATIONAL EXPOSURE LIMIT VALUES**

**Type of limit** : MAK (DE)  
**Limit value** : 2.8 mg/m<sup>3</sup>

**Remark** : 1 mg/m<sup>3</sup> = 0.175 ml/m<sup>3</sup> (ppm)  
Ceiling limit: Category I (should not be exceeded)  
MAK: 0.5 ppm

25.03.2004 (11)

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**Type of limit** : TLV (US)  
**Limit value** : .2 other: ppm  
**Short term exposure limit value**  
**Limit value** : .5 other: ppm  
**Time schedule** : 15 minute(s)  
**Frequency** : times

25.03.2004

(12)

**1.8.2 ACCEPTABLE RESIDUES LEVELS****1.8.3 WATER POLLUTION**

**Classified by** : other: VwVwS  
**Labelled by** :  
**Class of danger** : 1 (weakly water polluting)

**Remark** : Official German Classification with identification number (Kenn-Nr.) 1245  
26.03.2004

**1.8.4 MAJOR ACCIDENT HAZARDS**

**Legislation** : Stoerfallverordnung (DE)  
**Substance listed** : yes  
**No. in Seveso directive** :

**Remark** : Major accidents regulations: no. 1 (highly toxic)  
25.03.2004

(13)

**1.8.5 AIR POLLUTION****1.8.6 LISTINGS E.G. CHEMICAL INVENTORIES****1.9.1 DEGRADATION/TRANSFORMATION PRODUCTS****1.9.2 COMPONENTS****1.10 SOURCE OF EXPOSURE****1.11 ADDITIONAL REMARKS****1.12 LAST LITERATURE SEARCH**

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ID: 7719-12-2

DATE: 13.02.2006

**Type of search** : Internal and External  
**Chapters covered** : 1  
**Date of search** : 26.07.2002

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**Type of search** : Internal and External  
**Chapters covered** : 2  
**Date of search** : 26.07.2002

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**Type of search** : Internal and External  
**Chapters covered** : 3, 4  
**Date of search** : 26.07.2002

10.02.2004

**Type of search** : Internal and External  
**Chapters covered** : 5  
**Date of search** : 01.05.2003

10.02.2004

**1.13 REVIEWS**

**2.1 MELTING POINT**

<b>Value</b>	:	-93.6 °C	
<b>Sublimation</b>	:		
<b>Method</b>	:	other: no data	
<b>Year</b>	:	2003	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Remark</b>	:	Data selected as critical data because carefully peer-reviewed. MAK commission gave similar data. The melting point of -111.85 °C, found by mistake from S. v. Wroblewski, K. Olzewski (Ann. Phys. 3, 20 81983) 243-257) was cited by Landolt-Börnstein (1905) and International Critical Tables 1 (1926) p. 106.	
<b>Reliability</b>	:	(2) valid with restrictions Data from handbook or collection of data	
<b>Flag</b>	:	Critical study for SIDS endpoint	
22.09.2004			(5)
<b>Value</b>	:	-93.6 °C	
<b>Sublimation</b>	:		
<b>Method</b>	:	other: no data	
<b>Year</b>	:	2003	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Reliability</b>	:	(2) valid with restrictions Data from handbook or collection of data	
22.09.2004			(14)
<b>Value</b>	:	-93.6 °C	
<b>Sublimation</b>	:		
<b>Method</b>	:	other: no data	
<b>Year</b>	:	1988	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Reliability</b>	:	(4) not assignable Data from handbook or collection of data, not peer-reviewed	
22.09.2004			(15)
<b>Value</b>	:	-92 °C	
<b>Sublimation</b>	:		
<b>Method</b>	:	other: no data	
<b>Year</b>	:	1984	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Reliability</b>	:	(2) valid with restrictions Data from handbook or collection of data	
22.09.2004			(16)
<b>Value</b>	:	-111.8 °C	
<b>Sublimation</b>	:		
<b>Method</b>	:	other: no data	
<b>Year</b>	:	1979	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	

**Remark** : The melting point of -111.85 °C, found by mistake from S. v. Wroblewski, K. Olzewski (Ann. Phys. 3, 20 81983) 243-257) was cited by Landolt-Börnstein (1905) and International Critical Tables 1 (1926) p. 106.

**Reliability** : (2) valid with restrictions  
Data from handbook or collection of data

22.09.2004 (17)

**Value** : -112 °C  
**Sublimation** :  
**Method** : other: no data  
**Year** : 2003  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Remark** : The melting point of -111.85 °C, found by mistake from S. v. Wroblewski, K. Olzewski (Ann. Phys. 3, 20 81983) 243-257) was cited by Landolt-Börnstein (1905) and International Critical Tables 1 (1926) p. 106.

**Reliability** : (2) valid with restrictions  
Data from handbook or collection of data

22.09.2004 (18)

**Value** : -112 °C  
**Sublimation** :  
**Method** : other: no data  
**Year** : 1991  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Remark** : The melting point of -111.85 °C, found by mistake from S. v. Wroblewski, K. Olzewski (Ann. Phys. 3, 20 81983) 243-257) was cited by Landolt-Börnstein (1905) and International Critical Tables 1 (1926) p. 106.

**Reliability** : (2) valid with restrictions  
Data from handbook or collection of data

22.09.2004 (19)

**Value** : -112 °C  
**Sublimation** :  
**Method** : other: no data  
**Year** : 2001  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Remark** : The melting point of -111.85 °C, found by mistake from S. v. Wroblewski, K. Olzewski (Ann. Phys. 3, 20 81983) 243-257) was cited by Landolt-Börnstein (1905) and International Critical Tables 1 (1926) p. 106.

**Reliability** : (2) valid with restrictions  
Data from handbook or collection of data

22.09.2004 (20)

## 2.2 BOILING POINT

**Value** : 76.1 °C at 1013 hPa  
**Decomposition** :  
**Method** : other: no data  
**Year** : 2003  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

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<b>Reliability</b>	:	(2) valid with restrictions Data from handbook or collection of data	
<b>Flag</b>	:	Critical study for SIDS endpoint	
22.09.2004			(5)
<b>Value</b>	:	76.1 °C at 1013 hPa	
<b>Decomposition</b>	:		
<b>Method</b>	:	other: no data	
<b>Year</b>	:	2003	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Reliability</b>	:	(2) valid with restrictions Data from handbook or collection of data	
22.09.2004			(14)
<b>Value</b>	:	76.1 °C at 1013 hPa	
<b>Decomposition</b>	:		
<b>Method</b>	:	other: no data	
<b>Year</b>	:	1988	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Reliability</b>	:	(4) not assignable Data from handbook or collection of data, not peer-reviewed	
22.09.2004			(15)
<b>Value</b>	:	76 °C at 1013 hPa	
<b>Decomposition</b>	:		
<b>Method</b>	:	other: no data	
<b>Year</b>	:	2003	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Reliability</b>	:	(2) valid with restrictions Data from handbook or collection of data	
22.09.2004			(18)
<b>Value</b>	:	76 °C at 1013 hPa	
<b>Decomposition</b>	:		
<b>Method</b>	:	other: no data	
<b>Year</b>	:	2001	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Reliability</b>	:	(2) valid with restrictions Data from handbook or collection of data	
22.09.2004			(20)
<b>Value</b>	:	75.5 °C at 998.6 hPa	
<b>Decomposition</b>	:		
<b>Method</b>	:	other: no data	
<b>Year</b>	:	1991	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Reliability</b>	:	(2) valid with restrictions Data from handbook or collection of data	

22.09.2004 (19)

**Value** : 74.2 °C at  
**Decomposition** :  
**Method** : other: no data  
**Year** : 1979  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Reliability** : (2) valid with restrictions  
 Data from handbook or collection of data

22.09.2004 (17)

**Value** : 75 - 76 °C at 1020 hPa  
**Decomposition** :  
**Method** : other: not specified  
**Year** : 1896  
**GLP** : no  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Reliability** : (4) not assignable  
 Documentation insufficient for assessment

22.09.2004 (21)

**Value** : 74 °C at  
**Decomposition** :  
**Method** : other: no data  
**Year** : 1984  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Reliability** : (2) valid with restrictions  
 Data from handbook or collection of data

22.09.2004 (16)

### 2.3 DENSITY

**Type** : density  
**Value** : 1.575 g/cm<sup>3</sup> at 20 °C  
**Method** : other: no data  
**Year** : 2003  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Reliability** : (2) valid with restrictions  
 Data from handbook or collection of data

**Flag** : Critical study for SIDS endpoint

22.09.2004 (5)

**Type** : relative density  
**Value** : 1.574 at 21 °C  
**Method** : other: no data  
**Year** : 2001  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Result** : Value relative to the density of water at 4°C.

**Reliability** : (2) valid with restrictions

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	Data from handbook or collection of data	
22.09.2004		(20)
<b>Type</b>	: density	
<b>Value</b>	: 1.574 g/cm <sup>3</sup> at 21 °C	
<b>Method</b>	: other: no data	
<b>Year</b>	: 1979	
<b>GLP</b>	: no data	
<b>Test substance</b>	: other TS: Phosphorus trichloride, purity not given	
<b>Reliability</b>	: (2) valid with restrictions	
	Data from handbook or collection of data	
22.09.2004		(17)
<b>Type</b>	: relative density	
<b>Value</b>	: 1.6 at °C	
<b>Method</b>	: other: no data	
<b>Year</b>	: 2003	
<b>GLP</b>	: no data	
<b>Test substance</b>	: other TS: Phosphorus trichloride, purity not given	
<b>Result</b>	: Value relative to the density of water at ambient temperature.	
<b>Reliability</b>	: (2) valid with restrictions	
	Data from handbook or collection of data	
22.09.2004		(18)
<b>Type</b>	: relative density	
<b>Value</b>	: 1.575 at 20 °C	
<b>Method</b>	: other: no data	
<b>Year</b>	: 2003	
<b>GLP</b>	: no data	
<b>Test substance</b>	: other TS: Phosphorus trichloride, purity not given	
<b>Result</b>	: Value relative to the density of water at 4°C.	
<b>Reliability</b>	: (2) valid with restrictions	
	Data from handbook or collection of data	
22.09.2004		(14)
<b>Type</b>	: relative density	
<b>Value</b>	: 1.573 at 20 °C	
<b>Method</b>	: other: no data	
<b>Year</b>	: 1988	
<b>GLP</b>	: no data	
<b>Test substance</b>	: other TS: Phosphorus trichloride, purity not given	
<b>Result</b>	: Value relative to the density of water at 4°C.	
<b>Reliability</b>	: (4) not assignable	
	Data from handbook or collection of data, not peer-reviewed	
22.09.2004		(15)
<b>Type</b>	: density	
<b>Value</b>	: 1.57 g/cm <sup>3</sup> at 20 °C	
<b>Method</b>	: other: no data	
<b>Year</b>	: 2003	
<b>GLP</b>	: no data	
<b>Test substance</b>	: other TS: Phosphorus trichloride, purity not given	
<b>Reliability</b>	: (4) not assignable	
	Manufacturer data without proof	

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22.09.2004 (22)

**Type** : relative density  
**Value** : 1.5941 at 11 °C  
**Method** : other: not specified  
**Year** : 1896  
**GLP** : no  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Reliability** : (4) not assignable  
 Documentation insufficient for assessment

22.09.2004 (21)

**Type** : relative density  
**Value** : 1.574 at 21 °C  
**Method** : other: no data  
**Year** : 1991  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Result** : Value relative to the density of water at 4°C.  
**Reliability** : (2) valid with restrictions  
 Data from handbook or collection of data

22.09.2004 (19)

**Type** : density  
**Value** : 1.858 at -95.1 °C  
**Method** : other: not specified  
**Year** : 1965  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Reliability** : (2) valid with restrictions  
 Data from handbook or collection of data

22.09.2004 (23)

**Type** : density  
**Value** : 2.015 at -183 °C  
**Method** : other: not specified  
**Year** : 1965  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Reliability** : (2) valid with restrictions  
 Data from handbook or collection of data

22.09.2004 (23)

**Type** : density  
**Value** : at °C  
**Method** : other: not specified  
**Year** : 1965  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Result** : t\* -70 -20.5 0 20.8 35.2  
 d\*\* 1.744 1.653 1.613 1.574 1.547  
  
 t\* 50.3 64.8 75.1  
 d\*\* 1.518 1.492 1.475

\* t=temperature  
\*\* d=density

t*	0	15	25	35	50
d**	1.6114	1.5844	1.5659	1.5470	1.5181
t*	d**	t*	d**		
0	1.609	20.8	1.574		
0	1.6094	21	1.5696		
0	1.612	25	1.5659		
0	1.61275	30	1.5567		
10	1.597	33.2	1.5505		
16.4	1.582	43.5	1.5332		
17.9	1.5792	46.2	1.527		
18	1.579	59.7	1.5000		
18	1.585	61.5	1.5008		
18	1.576	76	1.471		
20	1.5778				
20	1.5761				

**Reliability** : (2) valid with restrictions  
Data from handbook or collection of data

22.09.2004

(23)

### 2.3.1 GRANULOMETRY

### 2.4 VAPOUR PRESSURE

**Value** : 129.7 hPa at 20 °C  
**Decomposition** :  
**Method** : other (measured): description of the method is not given  
**Year** : 2003  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Result** : Vapour pressure at 70°C: 857.5 hPa  
**Reliability** : (2) valid with restrictions  
Data from handbook or collection of data

**Flag** : Critical study for SIDS endpoint

22.09.2004

(5)

**Value** : 133 hPa at 21 °C  
**Decomposition** :  
**Method** : other (measured): description of the method is not given  
**Year** : 2001  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Reliability** : (2) valid with restrictions  
Data from handbook or collection of data

22.09.2004

(20)

**Value** : 133 hPa at 21 °C  
**Decomposition** :  
**Method** :  
**Year** : 1979  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

<b>Reliability</b>	:	(2) valid with restrictions Data from handbook or collection of data	
22.09.2004			(18) (17)
<b>Value</b>	:	127 hPa at 20 °C	
<b>Decomposition</b>	:		
<b>Method</b>	:	other (measured): description of the method is not given	
<b>Year</b>	:	1988	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Result</b>	:	Further, following values are reported: vapour pressure at 30°C 194 hPa vapour pressure at 50°C 415 hPa	
<b>Reliability</b>	:	(4) not assignable Data from handbook or collection of data, not peer-reviewed	
22.09.2004			(15)

## 2.5 PARTITION COEFFICIENT

<b>Partition coefficient</b>	:	octanol-water	
<b>Log pow</b>	:	at °C	
<b>pH value</b>	:		
<b>Method</b>	:	other (calculated): Expert judgement	
<b>Year</b>	:	2003	
<b>GLP</b>	:	no	
<b>Test substance</b>	:	other TS: Phosphorus trichloride	
<b>Result</b>	:	"Endpoint Partition Coefficient" is not applicable because the substance is not stable in water due to hydrolysis	
<b>Reliability</b>	:	(2) valid with restrictions Basic data given	
<b>Flag</b>	:	Critical study for SIDS endpoint	
22.09.2004			(24)

### 2.6.1 SOLUBILITY IN DIFFERENT MEDIA

<b>Solubility in</b>	:	Water	
<b>Value</b>	:	at °C	
<b>pH value</b>	:		
<b>concentration</b>	:	at °C	
<b>Temperature effects</b>	:		
<b>Examine different pol.</b>	:		
<b>pKa</b>	:	at 25 °C	
<b>Description</b>	:		
<b>Stable</b>	:		
<b>Deg. product</b>	:		
<b>Method</b>	:	other: Expert judgement	
<b>Year</b>	:	1990	
<b>GLP</b>	:	no	
<b>Test substance</b>	:	other TS: Phosphorus trichloride	
<b>Remark</b>	:	Not stable in water due to hydrolysis, see chapter 3.1.2	
<b>Reliability</b>	:	(2) valid with restrictions Data from handbook or collection of data	
<b>Flag</b>	:	Critical study for SIDS endpoint	

## 2. PHYSICO-CHEMICAL DATA

ID: 7719-12-2

DATE: 13.02.2006

22.09.2004 (25)

**Solubility in Value** : Organic Solvents  
 : at °C  
**pH value concentration** :  
 : at °C  
**Temperature effects** :  
**Examine different pol.** :  
**pKa** : at 25 °C  
**Description** :  
**Stable** :  
**Deg. product** :  
**Method** : other: no data  
**Year** : 1991  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Result** : Soluble in ether, benzene, chloroform, carbon disulfide and carbon tetrachloride.

**Reliability** : (2) valid with restrictions  
Data from handbook or collection of data

**Flag** : Critical study for SIDS endpoint

22.09.2004 (19)

**Solubility in Value** : Organic Solvents  
 : at °C  
**pH value concentration** :  
 : at °C  
**Temperature effects** :  
**Examine different pol.** :  
**pKa** : at 25 °C  
**Description** :  
**Stable** :  
**Deg. product** :  
**Method** : other: no data  
**Year** : 2003  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Result** : Soluble in benzene, chloroform, carbon tetrachloride, and diethyl ether. Miscible with phosphoryl chloride.

**Reliability** : (2) valid with restrictions  
Data from handbook or collection of data

**Flag** : Critical study for SIDS endpoint

22.09.2004 (5)

**Solubility in Value** : Organic Solvents  
 : at °C  
**pH value concentration** :  
 : at °C  
**Temperature effects** :  
**Examine different pol.** :  
**pKa** : at 25 °C  
**Description** :  
**Stable** :  
**Deg. product** :  
**Method** : other: no data  
**Year** : 2001  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Result** : Soluble in benzene, chloroform, ether, and carbon disulfide.  
**Reliability** : (2) valid with restrictions  
 Data from handbook or collection of data

22.09.2004

(20)

**2.6.2 SURFACE TENSION****2.7 FLASH POINT**

**Method** :  
**Year** :  
**GLP** :  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Remark** : not applicable  
**Flag** : Critical study for SIDS endpoint

22.09.2004

**2.8 AUTO FLAMMABILITY****2.9 FLAMMABILITY****2.10 EXPLOSIVE PROPERTIES****2.11 OXIDIZING PROPERTIES****2.12 DISSOCIATION CONSTANT****2.13 VISCOSITY**

**Value** : .65 - mPa s (dynamic) at 0 °C  
**Result** :  
**Method** : other: no data  
**Year** : 2003  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Result** : Further, a viscosity of 0.438 mPa\*s at 50°C is reported.  
**Reliability** : (2) valid with restrictions  
 Data from handbook or collection of data

22.09.2004

(5)

**Test type** : other: no data  
**Test procedure** :  
**Method** : other: no data  
**Year** : 1965  
**GLP** : no data

## 2. PHYSICO-CHEMICAL DATA

ID: 7719-12-2

DATE: 13.02.2006

<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Result</b>	:	t*    -15    0    10    20    30 h**   0.874 0.755 0.690 0.636 0.600	
		t*    40    50    60    70 h**   0.571 0.548 0.526 0.508	
		New measurements reveal lower values: t*    0    15    25    35    50 h**   0.656 0.560 0.501 0.479 0.453	
		* t=temperature ** h=viscosity	
<b>Reliability</b>	:	(2) valid with restrictions Data from handbook or collection of data	
22.09.2004			(23)
<b>Value Result</b>	:	.438 - mPa s (dynamic) at 50 °C	
<b>Method</b>	:	other: no data	
<b>Year</b>	:	1999	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Reliability</b>	:	(4) not assignable Manufacturer data without proof	
22.09.2004			(26)
<b>Value Result</b>	:	.536 - mPa s (dynamic) at 20 °C	
<b>Method</b>	:	other: DIN 53 015	
<b>Year</b>	:	2002	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Reliability</b>	:	(4) not assignable Manufacturer data without proof	
22.09.2004			(3)

## 2.14 ADDITIONAL REMARKS

<b>Memo</b>	:	Conversion factors	
<b>Result</b>	:	1 ml/m <sup>3</sup> (ppm) = 5.69 mg/m <sup>3</sup> 1 mg/m <sup>3</sup> = 0.175 ml/m <sup>3</sup> (ppm)	
<b>Flag</b>	:	Critical study for SIDS endpoint	
03.02.2004			(16)
<b>Memo</b>	:	Conversion factors at 20°C and 1.013 bar	
<b>Result</b>	:	1 ppm = 5.71 mg/m <sup>3</sup> 1 mg/m <sup>3</sup> = 0.175 ppm	
<b>Reliability</b>	:	(4) not assignable Data from handbook or collection of data, not peer-reviewed	
02.02.2004			(15)

<b>Memo</b>	:	Hazards identification and decomposition products							
<b>Result</b>	:	The substance decomposes on heating producing toxic and corrosive fumes including hydrogen chloride and phosphorus oxides. Reacts violently with water producing heat and decomposition products including hydrochloric acid and phosphorus acid, causing fire and explosion hazard. [Note: Hydrolysis in water to form hydrochloric acid and phosphoric acid.]							
<b>Reliability</b>	:	(2) valid with restrictions							
18.11.2003		Data from handbook or collection of data						(18)	
<b>Memo</b>	:	Refraction index							
<b>Result</b>	:	I*	263	274	288	298	325	340	
		n**	1.666	1.634	1.610	1.597	1.573	1.564	
		I*	346	361	467	480	508	537	
		n**	1.561	1.555	1.529	1.525	1.524	1.520	
		* I=wavelength in mμ							
		** n=refraction index							
<b>Reliability</b>	:	(2) valid with restrictions							
02.02.2004								(23)	
<b>Memo</b>	:	Relative vapour density							
<b>Result</b>	:	4.74 (air = 1)							
<b>Reliability</b>	:	(4) not assignable							
		Data from handbook or collection of data, not peer-reviewed							
02.02.2004								(15)	
<b>Memo</b>	:	Relative vapour density							
<b>Result</b>	:	4.75 (air = 1)							
<b>Reliability</b>	:	(2) valid with restrictions							
		Data from handbook or collection of data							
<b>Flag</b>	:	Critical study for SIDS endpoint							
18.11.2003								(18) (17)	
<b>Memo</b>	:	pH value							
<b>Result</b>	:	ca. pH 1 at 5 g/l water							
<b>Reliability</b>	:	(4) not assignable							
		Manufacturer data without proof							
<b>Flag</b>	:	Critical study for SIDS endpoint							
17.11.2003								(22) (26)	

**3.1.1 PHOTODEGRADATION**

<b>Deg. product</b>	:	
<b>Method</b>	:	other (calculated): expert judgement
<b>Year</b>	:	2004
<b>GLP</b>	:	
<b>Test substance</b>	:	other TS: Phosphorus trichloride
<b>Result</b>	:	Photodegradation due to OH radicals in the atmosphere is not calculable with AOPWIN v. 1.90 (2000) (expert judgement). Direct photolysis of gaseous phosphorus trichloride is not expected due to the lack of adsorption of light with a wavelength above 225 nm (Jan-Khan and Samuel 1936). Photodegradation in water cannot be calculated because the substance is not stable in water due to hydrolysis (Bayer Chemicals 2004).
<b>Reliability</b>	:	(2) valid with restrictions Accepted calculation method
<b>Flag</b>	:	Critical study for SIDS endpoint
22.09.2004		(24) (27)

**3.1.2 STABILITY IN WATER**

<b>Type</b>	:	abiotic
<b>t1/2 pH4</b>	:	at °C
<b>t1/2 pH7</b>	:	at °C
<b>t1/2 pH9</b>	:	at °C
<b>Deg. product</b>	:	yes
<b>Method</b>	:	other: pH monitoring
<b>Year</b>	:	2003
<b>GLP</b>	:	no
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given
<b>Deg. products</b>	:	7647-01-0 231-595-7 hydrogen chloride
<b>Method</b>	:	The reaction of phosphorus trichloride and water was studied by bringing a small amount of neat phosphorus trichloride into contact with an excess of well stirred water and following the generation of acidic reaction products using a pH electrode. Quantitative analysis (redox titration with AgNO <sub>3</sub> and pH titration) after completion of reaction confirmed that all reaction products have been captured by this method.
<b>Result</b>	:	The experimental set up could not distinguish the apparent reaction rate from, e.g., the mixing delay or the inertia of the measuring system. However, the half-life of phosphorus trichloride in water was estimated to be less than 10 seconds at 23 °C. Quantitative analysis (redox titration with AgNO <sub>3</sub> and pH titration) after completion of reaction confirmed that all reaction products were captured by this method. In the chloride titrations with AgNO <sub>3</sub> 99 % of the chloride expected to be generated by hydrolysis of the phosphorus trichloride were recovered. In the pH titrations, 94 % of the expected total acidity were recovered.
<b>Reliability</b>	:	(2) valid with restrictions Basic data given
<b>Flag</b>	:	Critical study for SIDS endpoint
22.09.2004		(24)
<b>Type</b>	:	abiotic
<b>t1/2 pH4</b>	:	at °C
<b>t1/2 pH7</b>	:	at °C

<b>t1/2 pH9</b>	:	at °C
<b>Deg. product</b>	:	
<b>Method</b>	:	other: see test conditions
<b>Year</b>	:	1925
<b>GLP</b>	:	no
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given
<b>Remark</b>	:	Study is reliable in regard to time course of acidity release. In regard to postulated intermediates up to date by the time the study was performed but outdated in regard to analytical methods of the 21st century
<b>Result</b>	:	<p>The kinetics and the intermediates of the hydrolysis of phosphorus trichloride were examined. A fast release of acidity (98 %) was observed within 1 min, but this acidity still increased within about half an hour. An "abnormally pronounced reducing property" was observed that became normal after some hours. The author postulated that P(OH)<sub>2</sub>Cl occurred as an intermediate which would account for increased acidity for 1/2-hour. The increased reducing power for hours is due to the second form of phosphorus acid, P(OH)<sub>3</sub> which does not immediately (after nascency from PCl<sub>3</sub>) change to the ordinary form HPO(OH)<sub>2</sub>. It also reacted as a reductant for iodine. However:</p> <ul style="list-style-type: none"> <li>- The speed of the iodine reaction might vary depending on the specific reaction conditions</li> <li>- The structural formula of the suggested intermediate is not consistent with the observation, that 98 % of the acidity was released within one minute (which means that most of this intermediate - if it was formed as a stoichiometric intermediate of hydrolysis - would have hydrolysed also during about 1 min but was virtually stable afterwards)</li> <li>- It is observed that when phosphorus trichloride reacts with water, the solution initially produced had much stronger reducing properties than after several hours.</li> <li>- The duration of increased reducing power is far longer than can be attributed to the intermediate oxychloride. It is suggested by the author that this may be due to the second form of phosphorus acid, P(OH)<sub>3</sub>, which does not immediately change to the ordinary form, HPO(OH)<sub>2</sub>.</li> <li>- The hydrolysis was always accompanied by a slight smell of phosphine which became more pronounced at much higher temperatures.</li> </ul>
<b>Test condition</b>	:	<p>The presence of reducing intermediates (postulated to be phosphorus (III) oxychloride [P(OH)<sub>2</sub>Cl] and P(OH)<sub>3</sub>) was examined by titration with iodine:</p> <ul style="list-style-type: none"> <li>- Up to 3.2 g of phosphorus trichloride were put in glass bulbs, which were submerged in 200 ml water</li> <li>- After breaking the bulbs and mixing the phosphorus trichloride with water (= start of the experiments), a iodine solution was added after a defined hydrolysis time (e.g. 15 s, 24 h)</li> <li>- Definite volumes of the solutions were titrated by a sodium thiosulphate solution at different times</li> </ul> <p>To examine the time course of acidity formation the solution containing phosphorus trichloride and water was titrated by sodium hydroxide</p>
<b>Reliability</b>	:	(2) valid with restrictions Basic data given
<b>Flag</b>	:	Critical study for SIDS endpoint
22.09.2004		(28)
<b>Type</b>	:	abiotic
<b>t1/2 pH4</b>	:	at °C
<b>t1/2 pH7</b>	:	at °C
<b>t1/2 pH9</b>	:	at °C
<b>Deg. product</b>	:	
<b>Method</b>	:	other: Calorimetry
<b>Year</b>	:	1989

<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Method</b>	:	The rate of hydrolysis measured in dilute aqueous-organic solvent (dioxan) using calorimetric techniques. - Precision calorimeter LKV-8700 (Sweden). The response time of the instrument was about 0.5 s, suited to determine half-lives of about 2 s - Heat released (value taken from literature): 68,5 kcal/Mol for $\text{PCl}_3 + 3 \text{H}_2\text{O} > \text{H}_3\text{PO}_3 + 3 \text{HCl}$ - After thermic equilibrium at 18 °C, an ampulla containing $\text{PCl}_3$ in dioxan was broken in dioxan/water (= start of experiment) and the heat released was measured for 1 h	
<b>Result</b>	:	The reaction is of second order. The half-life of phosphorus trichloride was found to be 1-13 seconds, depending on reaction conditions for $\text{PCl}_3$ -concentration in the range $6.1\text{-}18 \times 10^{-4}$ mol/l and water concentrations in the range $4\text{-}350 \times 10^{-3}$	
<b>Reliability</b>	:	(2) valid with restrictions Basic data given	
<b>Flag</b>	:	Critical study for SIDS endpoint	
22.09.2004			(29)
<b>Type</b>	:	abiotic	
<b>t1/2 pH4</b>	:	at °C	
<b>t1/2 pH7</b>	:	at °C	
<b>t1/2 pH9</b>	:	at °C	
<b>Deg. product</b>	:		
<b>Method</b>	:		
<b>Year</b>	:	1872	
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Result</b>	:	Reaction products of phosphorus trichloride in water were examined. In hot water hydrolysis led to hydrochloric acid, phosphorus acid and a small amount of red phosphorus (with the unproven postulation of the formation of phosphine as an intermediate).	
<b>Reliability</b>	:	(4) not assignable Unsuitable test system	
<b>Flag</b>	:	Critical study for SIDS endpoint	
22.09.2004			(30)
<b>Type</b>	:	abiotic	
<b>t1/2 pH4</b>	:	at °C	
<b>t1/2 pH7</b>	:	at °C	
<b>t1/2 pH9</b>	:	at °C	
<b>Deg. product</b>	:		
<b>Method</b>	:	other: Computer modeling	
<b>Year</b>	:	2001	
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Method</b>	:	The dangers caused by accidental releases of phosphorus trichloride (including spill behaviour) were examined using a computer model (REACTPOOL).	
<b>Remark</b>	:	Not relevant for assessment	
<b>Result</b>	:	The model suggests possible effects depending on the way phosphorus trichloride reacts with water, the amount of water available, surface roughness and wind speed. Under stoichiometric or excess water conditions the reaction products between phosphorus trichloride and water are hydrochloric acid and	

- phosphorus acid with 3 mol of hydrochloric acid forming for every mole of phosphorus trichloride.  
The hydrolysis is highly exothermic, raising both the temperature and the vapour evolution rates. Hydrochloric acid vapour will be evolved due to its high volatility. The amount of phosphorous acid evolved is negligible due to its extremely low volatility.  
When phosphorus trichloride is in excess of the stoichiometric amount of water essential for complete hydrolysis, liquid phosphoric acid, solid red phosphorus and hydrochloric acid were formed. The hydrolysis reaction may produce solid particles of red phosphorus.  
Increasing roughness and wind speed results in increasing vapour evolution rates.
- Reliability** : (2) valid with restrictions  
Basic data given
- 14.07.2005 (31)
- Type** : abiotic  
**t1/2 pH4** : at °C  
**t1/2 pH7** : at °C  
**t1/2 pH9** : at °C  
**Deg. product** :  
**Method** : other: Calorimetry  
**Year** : 1998  
**GLP** : no data  
**Test substance** : other TS: Phosphorus trichloride, purity not given  
**Deg. products** : 13598-36-2 237-066-7 phosphonic acid  
7647-01-0 231-595-7 hydrogen chloride  
7664-38-2 231-633-2 orthophosphoric acid  
7723-14-0 231-768-7 phosphorus
- Method** : Small scale experiments on the reaction characteristics of phosphorus trichloride and water, including both closed and open test cells. The impact of variables on reaction rates including the interface surface area, layer depth, and stirring, were investigated experimentally.
- Result** : -Phosphorus trichloride hydrolyzes according to:  $PCl_3 + 3 H_2O > H_3PO_3 + 3 HCl$ .  
-The reaction between water and phosphorus trichloride begins immediately.  
-When hydrochloric acid is added rather than water, the maximum temperature achieved is less (maximum approximately 75°C compared to 100°C) and the reaction appears to proceed more slowly.  
-From the stoichiometric case, three stages were apparent following the addition of water. Firstly, a rapid temperature rise and little loss of water is observed as the reaction proceeds and all hydrochloric acid enters into solution. Then the temperature falls due to further dilution with water and phosphorus trichloride vapour is evolved. Finally the temperature rises again and only hydrochloric acid is liberated.  
-No phosphine odour is detected.
- Test condition** : Experiments were carried out using ARC (Accelerating Rating Calorimeter), RSST (Reactive System Screening Tool -pseudo-adiabatic calorimeter) and VST (Vent Sizing Package).  
Four different reactor vessels were used during the experiments: a Dewar flask, a 50 ml beaker, a 100 ml and a 25 ml graduated cylinder. An electrically driven glass rod with a right angle bend was used as a stirrer in four of the cases to investigate the effect of mechanical mixing on the reaction. Only one stirring speed, estimated to be around 300 rpm, was used. The desired volume of phosphorus trichloride was measured, heated to about 50°C and then added to the reactor vessel. The reactor was then placed on an electronic mass balance and a thermocouple was inserted, so that it penetrated the phosphorus trichloride layer prior to water addition.

		The stirrer was started and a measured volume of distilled water or HCL (aqueous) at ambient temperature was added. The temperature and mass of the reactor was recorded throughout the water addition and reaction stages using a computer data logging system.	
<b>Reliability</b>	:	(2) valid with restrictions	
22.09.2004		Study meets generally accepted scientific principles	(32)
<b>Type</b>	:	abiotic	
<b>t1/2 pH4</b>	:	at °C	
<b>t1/2 pH7</b>	:	at °C	
<b>t1/2 pH9</b>	:	at °C	
<b>Deg. product</b>	:	not measured	
<b>Method</b>	:	other: see test conditions	
<b>Year</b>	:	1942	
<b>GLP</b>	:	no	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Remark</b>	:	Stock solution contains toluene	
<b>Result</b>	:	At 20 °C the constant for the reaction rate was found to be $k = 0.093$ corresponding to a half life of $t_{1/2} = 7$ s.	
		At 35 °C the constant for the reaction rate was found to be $k = 0.13$ corresponding to a half life of $t_{1/2} = 5$ s.	
<b>Test condition</b>	:	Stock solution was prepared with 137.4 g of phosphorus trichloride dissolved in 1000 cm <sup>3</sup> toluene. An aliquot of 10 cm <sup>3</sup> was added in 10 cm <sup>3</sup> bidistilled, not buffered water and introduced in a reactor provided with a thermostat. Two tests were performed with different temperatures: 20 °C and 35 °C. For each temperature 3 replicates were conducted (with slight different surface of reaction). Solution was analysed in an interval of approx. 7 min till 47 min (20 °C) and 34 min (25 °C). Analytical method: The sample was titrated with NaOH N/10. Phenolphthalein was used as an indicator.	
<b>Reliability</b>	:	(2) valid with restrictions	
22.09.2004		Basic data given	(33)
<b>Type</b>	:	abiotic	
<b>t1/2 pH4</b>	:	at °C	
<b>t1/2 pH7</b>	:	at °C	
<b>t1/2 pH9</b>	:	at °C	
<b>Deg. product</b>	:		
<b>Method</b>	:	other: see below	
<b>Year</b>	:	1987	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Method</b>	:	Vapor phase hydrolysis of phosphorus trichloride in argon was examined at about 1.000 K.	
<b>Remark</b>	:	High temperature hydrolysis not relevant for environmental assessment.	
<b>Result</b>	:	In the equilibrium gas mixture, co-condensed with an excess of argon, OPCI is detected. After addition of water in excess molecular phosphoric acid (HOPO) is found.	
<b>Reliability</b>	:	(2) valid with restrictions	
22.09.2004		Study meets generally accepted scientific principles	(34) (35)
<b>Type</b>	:	abiotic	

## 3. ENVIRONMENTAL FATE AND PATHWAYS

ID: 7719-12-2

DATE: 13.02.2006

**t1/2 pH4** : at °C  
**t1/2 pH7** : at °C  
**t1/2 pH9** : at °C  
**Degradation** : .2 % after 39 minute(s) at pH and 5 °C  
**Deg. product** : yes  
**Method** : other: non-homogeneous system according to Carrara and Zoppellari (1894)  
**Year** : 1896  
**GLP** : no  
**Test substance** : other TS: Phosphorus trichloride, purity not given  
**Deg. products** : 7647-01-0 231-595-7 hydrogen chloride  
  
**Result** : Authors report that within about 39 min at 5 °C only 0.2 % (= 1/500 molecular weight) of phosphorus chloride was completely degraded (K = 0.0071). Only 2-3 °C above the temperature the observation took place (5 °C), the reaction was too vigorous to make measurements.  
**Test condition** : Examination of the reaction of phosphorus trichloride with water in non-homogeneous system according to Carrara and Zoppellari (1894). Phosphorus trichloride hydrolyses according to  $PCl_3 + 3 H_2O > PO_3H_3 + 3HCl$ .  
- Cylindrical recipient with a thermostate  
- Constant temperature (5 °C)  
- The amount of acid released was analyzed with alkali  
- Test period: 70 min  
**Reliability** : (4) not assignable  
Documentation insufficient for assessment

22.09.2004

(36) (21)

**Type** : abiotic  
**t1/2 pH4** : at °C  
**t1/2 pH7** : at °C  
**t1/2 pH9** : at °C  
**Deg. product** :  
**Method** : other: Description of the method is not given  
**Year** : 1924  
**GLP** : no  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Remark** : The nature of the water - phosphorus trichloride reaction is described. It is suggested that the oxygen atom in a water molecule would attach itself to the lone pair of electrons on the phosphorus atom and a hydrogen and chlorine atom would then react to form hydrochloric acid. This would be repeated until all chlorine atoms were displaced from the phosphorus.

**Reliability** : (4) not assignable  
Documentation insufficient for assessment

22.09.2004

(37)

**Type** : abiotic  
**t1/2 pH4** : at °C  
**t1/2 pH7** : at °C  
**t1/2 pH9** : at °C  
**Deg. product** :  
**Method** :  
**Year** : 1947  
**GLP** :  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Remark** : It is not clearly stated, how the humidity of the gas phase was controlled  
**Result** : Hydrolysis of phosphorus trichloride in vapour phase was investigated. No

- reaction between phosphorus trichloride and water vapour in the vapour phase was detected. At 25 °C the reaction was shown to be heterogeneous by comparing rates in waxed and unwaxed vessels. Authors speculate that the rate of hydrolysis is governed by the rate of absorption of the gases on the phosphoric acid deposited on the reaction vessel surface. At higher temperatures no hydrolysis could be detected.  
Description and experimental details are not sufficient.
- Reliability** : (4) not assignable  
Documentation insufficient for assessment
- 22.09.2004 (38)
- Type** : abiotic  
**t1/2 pH4** : at °C  
**t1/2 pH7** : at °C  
**t1/2 pH9** : at °C  
**Deg. product** :  
**Method** : other: description of the method is not given  
**Year** : 1897  
**GLP** : no  
**Test substance** : other TS: Phosphorus trichloride, purity not given
- Remark** : Not relevant for environmental assessment  
**Result** : The reaction products of phosphorus trichloride and water were hydrochloric acid and phosphorous acid. When phosphorus trichloride was in excess of water the author postulates the formation of some phosphoryl monochloride as well as P2O5, P2O3 and P2O.
- Reliability** : (4) not assignable  
Documentation insufficient for assessment
- 22.09.2004 (39) (40)
- Type** : abiotic  
**t1/2 pH4** : at °C  
**t1/2 pH7** : at °C  
**t1/2 pH9** : at °C  
**Deg. product** :  
**Method** :  
**Year** : 1871  
**GLP** :  
**Test substance** : other TS: Phosphorus trichloride, purity not given
- Remark** : For excess phosphorus trichloride the formation of some red phosphorus, hydrochloric acid and phosphoric acid was observed. It is suggested that initially phosphorus acid was formed which then reacted with phosphorus trichloride. Overall, 12 mol of water were required per 5 mol of phosphorus trichloride as indicated by the suggested total reaction.
- Reliability** : (4) not assignable  
Original reference not available
- 22.09.2004 (41)
- Type** : abiotic  
**t1/2 pH4** : at °C  
**t1/2 pH7** : at °C  
**t1/2 pH9** : at °C  
**Deg. product** :  
**Method** :  
**Year** : 1811  
**GLP** :  
**Test substance** : other TS: Phosphorus trichloride, purity not given

<b>Remark</b>	:	It is noted that a rise in temperature and the gradual formation of hydrochloric acid and phosphorus acid occurred, when water reacted with phosphorus trichloride. It is suggested that three moles of water are required per mole of phosphorus trichloride for hydrolysis.	
<b>Reliability</b>	:	(4) not assignable Original reference not available	
22.09.2004			(42)
<b>Type</b>	:	abiotic	
<b>t1/2 pH4</b>	:	at °C	
<b>t1/2 pH7</b>	:	at °C	
<b>t1/2 pH9</b>	:	at °C	
<b>Deg. product</b>	:		
<b>Method</b>	:	other: modeling	
<b>Year</b>	:	2001	
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Method</b>	:	A case study modeling the indoor release of phosphorus trichloride in a major accident. It is assumed that when phosphorus trichloride reacts with the moisture in air hydrochloric acid and phosphorous acid are formed. For modeling purposes the duration of exposure, the mass of water for reaction, the pool growth, the mass flow rate of hydrochloric acid produced and the mass release rate of hydrochloric acid at different circumstances of accident were determined.	
<b>Remark</b>	:	Not relevant for environmental assessment	
<b>Result</b>	:	The hydrochloric acid release rate is the same for releases from storage, no matter the size of the leak from the storage tank. This is because it is assumed that the banded area is immediately filled and thus the surface area over which the reaction with moisture in air takes place is always constant. In reality the liquid pool would grow to its limiting size and there would also be the surface area of the liquid as it is released from the hole.	
<b>Reliability</b>	:	(2) valid with restrictions Study meets generally accepted scientific principles	
22.09.2004			(43)
<b>Type</b>	:	abiotic	
<b>t1/2 pH4</b>	:	at °C	
<b>t1/2 pH7</b>	:	at °C	
<b>t1/2 pH9</b>	:	at °C	
<b>Deg. product</b>	:		
<b>Method</b>	:		
<b>Year</b>	:	1996	
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Remark</b>	:	In general, adding small amounts of phosphorus trichloride to water (molar ratio of water to phosphorus trichloride > 3), the hydrolysis products of phosphorus trichloride are phosphonic acid and hydrochloric acid. Is the molar ratio of water to phosphorus trichloride between 2.5 and 3, a mixture of phosphorus and pyrophosphonic acid is formed. A molar ratio below 2.5 results in a product of indefinite composition, called lower oxides of phosphorus. When the molar ratio of water to phosphorus trichloride is low, traces of phosphine may be formed in hot water.	
<b>Reliability</b>	:	(2) valid with restrictions Data from handbook or collection of data	
<b>Flag</b>	:	Critical study for SIDS endpoint	
22.09.2004			(4)

**3.1.3 STABILITY IN SOIL**

**Deg. product** :  
**Method** :  
**Year** :  
**GLP** :  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Remark** : not stable, hydrolysis with moisture in soil  
 22.09.2004 (25)

**3.2.1 MONITORING DATA****3.2.2 FIELD STUDIES****3.3.1 TRANSPORT BETWEEN ENVIRONMENTAL COMPARTMENTS**

**Type** : adsorption  
**Media** :  
**Air** : % (Fugacity Model Level I)  
**Water** : % (Fugacity Model Level I)  
**Soil** : % (Fugacity Model Level I)  
**Biota** : % (Fugacity Model Level II/III)  
**Soil** : % (Fugacity Model Level II/III)  
**Method** : other: Expert judgement  
**Year** : 2003

**Result** : Models of fate and behaviour in the environment require values for  $K_{ow}$ , water solubility and vapour pressure. Since the substance is highly unstable in water, environmental distribution modelling of either the substance itself or its hydrolysis products (mineral acids, the anions of which are ubiquitous in the environment) is not relevant in this case.

**Test substance** : Phosphorus trichloride, purity not given  
**Reliability** : (2) valid with restrictions  
 Basic data given

**Flag** : Critical study for SIDS endpoint  
 14.07.2005 (24)

**3.3.2 DISTRIBUTION**

**Media** : air - biota - sediment(s) - soil - water  
**Method** : other (calculation): Expert judgement  
**Year** : 2003

**Result** : Models of fate and behaviour in the environment require values for  $K_{ow}$ , water solubility and vapour pressure. Since the substance is highly unstable in water, environmental distribution modelling of either the substance itself or its hydrolysis products (mineral acids, the anions of which are ubiquitous in the environment) is not relevant in this case.

**Test substance** : Phosphorus trichloride, purity not given  
**Reliability** : (2) valid with restrictions  
 Basic data given

**Flag** : Critical study for SIDS endpoint  
14.07.2005 (24)

### 3.4 MODE OF DEGRADATION IN ACTUAL USE

### 3.5 BIODEGRADATION

**Type** : aerobic  
**Inoculum** :  
**Deg. product** :  
**Method** : other: expert judgement  
**Year** : 2004  
**GLP** : no  
**Test substance** : other TS: Phosphorus trichloride, purity not given

**Remark** : "Endpoint Biodegradation" not applicable to inorganics.  
Since phosphorus trichloride hydrolyzes rapidly in water (Bayer Chemicals 2004), no biodegradation can be measured. The hydrolysis products chloride, phosphonate and hydrogen ions, are inorganic end products of biodegradation.

**Reliability** : (2) valid with restrictions  
Basic data given

**Flag** : Critical study for SIDS endpoint  
22.09.2004 (24)

### 3.6 BOD5, COD OR BOD5/COD RATIO

### 3.7 BIOACCUMULATION

### 3.8 ADDITIONAL REMARKS

**4.1 ACUTE/PROLONGED TOXICITY TO FISH**

<b>Type</b>	:	static	
<b>Species</b>	:	Brachydanio rerio (Fish, fresh water)	
<b>Exposure period</b>	:	96 hour(s)	
<b>Unit</b>	:	mg/l	
<b>LC0</b>	:	>= 1000	
<b>Limit test</b>	:		
<b>Analytical monitoring Method</b>	:	no	
<b>Method</b>	:	other: UBA-Verfahrensvorschlag "Letale Wirkung beim Zebraerbling Brachydanio rerio" ( LC 0, LC 50, LC 100; 48-96 Stunden) (Mai 1984)	
<b>Year</b>	:	1991	
<b>GLP</b>	:	yes	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Remark</b>	:	Although not explicitly mentioned in the original report, the test was conducted with pH-neutralised medium. LC0 >= 1000 mg/l PCI3 corresponds to a LC0 of >= 597 mg/l of (neutralised) phosphonic acid as hydrolysis product. The accepted scientific name for Brachydanio rerio is Danio rerio.	
<b>Test condition</b>	:	- Brachydanio rerio from West-Aquarium (Bad Lauterberg) - pH: 7.3-7.7 adjusted - Temperature: 22 °C - Oxygen: 7.8-8.6 mg/l - Test concentration: 1000 mg/l (nominal concentration), 1000 mg/l PCI3 correspond to 597 mg/l HP(O)(OH) <sub>2</sub> - 5 l test medium, 10 fishes/tank - No carrier - Control: synthetic tap water, hardness 14.8 °dH - No analytical monitoring because test substance hydrolyses rapidly into hydrochloric acid and phosphonic acid	
<b>Reliability</b>	:	(1) valid without restriction Test procedure in accordance with national standard methods	
<b>Flag</b>	:	Critical study for SIDS endpoint	
14.07.2005			(44)
<b>Type</b>	:	semistatic	
<b>Species</b>	:	Leuciscus sp. (Fish, fresh water)	
<b>Exposure period</b>	:	10 day(s)	
<b>Unit</b>	:	mg/l	
<b>LC100</b>	:	25	
<b>Limit test</b>	:		
<b>Analytical monitoring Method</b>	:	no data	
<b>Method</b>	:	other	
<b>Year</b>	:	1970	
<b>GLP</b>	:	no	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Remark</b>	:	Original reference in Russian, cited according to German translation The pH was presumably pH 3-4, which is not tolerated by several fish species (compare OECD-SIDS Hydrochloric Acid (2002)). The recommended for fish tests is pH 6.0 to pH 8.5 according to OECD Guidelines	
<b>Result</b>	:	5d-LC100 = 30 mg/l 10d-LC100 = 25 mg/l	
<b>Test condition</b>	:	Tests were performed under the following conditions: -Test species: Leuciscus leuciscus -Dilution water: dechlorinated tap water	

	-Test temperature: 22 - 24 °C	
	-Oxygen concentration: 5.8 - 7.8 mg/l	
	-Concentration of test substance: 10, 20, 30, 40 mg/l, pH not adjusted	
	-Test duration: 5-10 days	
<b>Reliability</b>	: (2) valid with restrictions	
	Study acceptable for assessment	
<b>Flag</b>	: Critical study for SIDS endpoint	(45)
22.07.2004		
<b>Type</b>	: semistatic	
<b>Species</b>	: other: species name not stated, it is assumed to be <i>Acipenser stellatus</i>	
<b>Exposure period</b>	: 5 day(s)	
<b>Unit</b>	: mg/l	
<b>NOEC</b>	: ca. 20	
<b>LC100</b>	: ca. 70	
<b>EC1 (length)</b>	: ca. 20	
<b>EC5 (body weight)</b>	: ca. 20	
<b>Limit test</b>	:	
<b>Analytical monitoring</b>	: no data	
<b>Method</b>	: other	
<b>Year</b>	: 1970	
<b>GLP</b>	: no data	
<b>Test substance</b>	: other TS: Phosphorus trichloride, purity not given	
<b>Remark</b>	: Original reference in Russian, cited according to German translation	
<b>Result</b>	: 70 mg/l (highest concentration tested): 100 % malformation of larvae, death after 3d (LC100)	
	60 mg/l: 15 % reduction of length, 30 % reduction of body weight, slight pigmentation	
	20 mg/l (lowest concentration tested): 1.2 % reduction of length (EC1), 5.4 % reduction of body weight (EC5). EC1 and EC5 were used as NOEC (20 mg/l).	
<b>Test condition</b>	: Test was performed under the following conditions:	
	-Test species: larvae of <i>Acipenser stellatus</i>	
	-Test vessels: enamelled basins of 1 l or aquaria of 10 l	
	-Dilution water: water from the river Wolga	
	-Test temperature: 20.2 - 24.2 °C	
	-Oxygen concentration: 5.9 - 8.14 mg/l	
	-Concentration of test substance: 20, 60, 70 mg/l, pH not adjusted	
	-Test duration: 5 days	
<b>Reliability</b>	: (2) valid with restrictions	
	Study acceptable for assessment	
<b>Flag</b>	: Critical study for SIDS endpoint	(45)
22.09.2004		
<b>Type</b>	: semistatic	
<b>Species</b>	: other: species name not stated, it is assumed to be <i>Acipenser stellatus</i> (eggs)	
<b>Exposure period</b>	: 3 day(s)	
<b>Unit</b>	: mg/l	
<b>NOEC</b>	: ca. 60	
<b>LC100</b>	: ca. 75	
<b>LC30</b>	: ca. 70	
<b>Limit test</b>	:	
<b>Analytical monitoring</b>	: no data	
<b>Method</b>	: other	
<b>Year</b>	: 1970	
<b>GLP</b>	: no data	
<b>Test substance</b>	: other TS: Phosphorus trichloride, purity not given	

<b>Remark</b>	:	Original reference in Russian, cited according to German translation	
<b>Result</b>	:	LC0 ca. 60 mg/l LC100 ca. 75 mg/l	
<b>Test condition</b>	:	Test was performed under the following conditions: -Test species: eggs of <i>Acipenser stellatus</i> -Test vessels: enamelled basins of 1 l -Dilution water: water from the river Wolga -Test temperature: 20.2 - 24.2°C -Oxygen concentration: 5.9 - 8.14 mg/l -Concentration of test substance: 20, 60, 70, 74, 80, 100, 150 mg/l, pH not adjusted -Test duration: 3 days	
<b>Reliability</b>	:	(2) valid with restrictions Study acceptable for assessment	
<b>Flag</b> 22.09.2004	:	Critical study for SIDS endpoint	(45)
<b>Type</b>	:	semistatic	
<b>Species</b>	:	other: species name not stated, it is assumed to be <i>Esox lucius</i> and <i>Perca fluviatilis</i>	
<b>Exposure period</b>	:	30 day(s)	
<b>Unit</b>	:	mg/l	
<b>LC0</b>	:	> 40	
<b>Limit test</b>	:		
<b>Analytical monitoring</b>	:	no data	
<b>Method</b>	:	other	
<b>Year</b>	:	1970	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Remark</b>	:	Original reference in Russian, cited according to German translation	
<b>Result</b>	:	No mortality and no deviations from normal appearance of fish observed at 40-50 mg/l	
<b>Reliability</b>	:	(4) not assignable Documentation insufficient for assessment	
<b>Flag</b> 22.09.2004	:	Critical study for SIDS endpoint	(45)
<b>Type</b>	:	semistatic	
<b>Species</b>	:	<i>Carassius carassius</i> (Fish, fresh water)	
<b>Exposure period</b>	:	30 day(s)	
<b>Unit</b>	:	mg/l	
<b>NOEC</b>	:	ca. 40	
<b>Limit test</b>	:		
<b>Analytical monitoring</b>	:	no data	
<b>Method</b>	:	other	
<b>Year</b>	:	1970	
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given	
<b>Remark</b>	:	A NOEC of ca. 40 mg/l was derived from EC0 Original reference in Russian, cited according to German translation Test cannot be used to derive a PNECaqua, because it is not clear, whether early life stages have been covered	
<b>Result</b>	:	10-40 mg/l: Continuous increase of body weight during whole test period, no difference in comparison to control 50-58 mg/l: Decrease in body weight up to day 15, than increase of body weight 60 mg/l: No increase of body weight up to day 30, than decrease to 0.35 mg body weight	

<b>Test condition</b>	<p>65 mg/l: 100 % death after 2 days data of body weight development of control are not given</p> <p>: Tests were performed under the following conditions:</p> <ul style="list-style-type: none"> <li>-Test vessel 10 l aquarium</li> <li>-Semistatic incubation with change of incubation solution every 24 h</li> <li>-Dilution water: dechlorinated tap water</li> <li>-Test temperature: 22 - 24 °C</li> <li>-Oxygen concentration: 5.8 - 7.8 mg/l</li> <li>-pH 3.3- 7.0</li> <li>-Concentration of test substance and corresponding values for HCl and HP(O)(OH)<sub>2</sub>:</li> </ul> <table border="0" style="margin-left: 20px;"> <thead> <tr> <th>PCl<sub>3</sub> (mg/l)</th> <th>HCl (mg/l)</th> <th>cal HP(O)(OH)<sub>2</sub></th> </tr> </thead> <tbody> <tr><td>10</td><td>2.7</td><td>6.0</td></tr> <tr><td>20</td><td>5.3</td><td>11.9</td></tr> <tr><td>30</td><td>8.0</td><td>18.0</td></tr> <tr><td>40</td><td>10.6</td><td>23.9</td></tr> <tr><td>50</td><td>13.3</td><td>29.9</td></tr> <tr><td>55</td><td>14.3</td><td>32.8</td></tr> <tr><td>58</td><td>15.4</td><td>34.6</td></tr> <tr><td>60</td><td>16.0</td><td>35.8</td></tr> <tr><td>65</td><td>17.3</td><td>38.8</td></tr> </tbody> </table> <ul style="list-style-type: none"> <li>-Fish were weighthed every five days</li> <li>-Fish were fed with earthworms and gammarids</li> <li>-Prior to incubation, fish were acclimated to test conditions for 10-15 days in dechlorinated tap water</li> </ul>	PCl <sub>3</sub> (mg/l)	HCl (mg/l)	cal HP(O)(OH) <sub>2</sub>	10	2.7	6.0	20	5.3	11.9	30	8.0	18.0	40	10.6	23.9	50	13.3	29.9	55	14.3	32.8	58	15.4	34.6	60	16.0	35.8	65	17.3	38.8
PCl <sub>3</sub> (mg/l)	HCl (mg/l)	cal HP(O)(OH) <sub>2</sub>																													
10	2.7	6.0																													
20	5.3	11.9																													
30	8.0	18.0																													
40	10.6	23.9																													
50	13.3	29.9																													
55	14.3	32.8																													
58	15.4	34.6																													
60	16.0	35.8																													
65	17.3	38.8																													
<b>Reliability</b>	<p>: (2) valid with restrictions Study acceptable for assessment</p>																														
<b>Flag</b> 22.07.2004	<p>: Critical study for SIDS endpoint</p>																														

(45)

#### 4.2 ACUTE TOXICITY TO AQUATIC INVERTEBRATES

<b>Type</b>	: static
<b>Species</b>	: Daphnia magna (Crustacea)
<b>Exposure period</b>	: 48 hour(s)
<b>Unit</b>	: mg/l
<b>EC0</b>	: 25
<b>EC50</b>	: 35.4
<b>EC100</b>	: 50
<b>Analytical monitoring</b>	: no
<b>Method</b>	: Directive 92/69/EEC, C.2
<b>Year</b>	: 2003
<b>GLP</b>	: yes
<b>Test substance</b>	: other TS: Phosphorus trichloride, purity not given
<b>Method</b>	<p>: Method is in most parts equivalent to the OECD TG 202 Daphnia sp., Acute immobilisation test and reproduction test, Part I -The 24h EC50 Acute immobilisation test.</p>
<b>Result</b>	<p>: The following values were determined:</p> <p>24h-EC0 = 25 mg/l 24h-EC100 = 50 mg/l 48h-EC0 = 25 mg/l 48h-EC100 = 50 mg/l</p> <p>Under pH-adjusted conditions no immobilisation of the daphnids has been observed at a nominal concentration up to 100 mg/l. Geometric mean (EC0/EC100) = 35.4 mg/l Measured pH-values: concentration 0h 48h</p>

	control	7.9	7.8
	12.5 mg/l	6.7	7.7
	25 mg/l	6.1	7.2
	50 mg/l	3.6	3.7*
	100 mg/l	2.9	2.9*
	50 mg/l	7.9	7.9 (with adjusted pH)
	100 mg/l	8.0	7.8 (with adjusted pH)
	The results of the respective replicates clearly demonstrate that the immobilisation observed was caused by pH-effects.		
<b>Test condition</b>	:	-50 ml glass beakers holding 10 neonates in 20 ml of test medium -dilution water: reconstituted water, total hardness, measured at test start: 14.8°dH -10 neonates per vessel, 2 replicates per concentration/control -temperature during the test: 18-22 °C -pH and oxygen values measured at the end of the test -experimental design: 6 test concentrations plus 1 control -no feeding during the exposure period -lighting: 16 h light/8 h dark -nominal test concentrations: 12.5, 25, 50 and 100 mg/l without adjustment of pH-value, additionally 50 and 100 mg/l with adjustment of the pH-value, since extreme pH decreases were observed due to hydrolysis of the test substance -criteria of effects: item-induced alteration of the normal mobility behaviour and loss of locomotory actions of the neonates, observed at 24 h and 48 h -no chemical analysis has been performed, as the test substance phosphorus trichloride hydrolyses rapidly in aqueous medium.	
<b>Reliability</b>	:	(1) valid without restriction Comparable to guideline study	
<b>Flag</b>	:	Critical study for SIDS endpoint	
14.07.2005			(46)

#### 4.3 TOXICITY TO AQUATIC PLANTS E.G. ALGAE

<b>Species</b>	:	Scenedesmus subspicatus (Algae)
<b>Endpoint</b>	:	growth rate
<b>Exposure period</b>	:	72 hour(s)
<b>Unit</b>	:	mg/l
<b>NOEC</b>	:	12.5 measured/nominal
<b>LOEC</b>	:	25
<b>EC50</b>	:	33.41
<b>Limit test</b>	:	yes
<b>Analytical monitoring</b>	:	no
<b>Method</b>	:	Directive 92/69/EEC, C.3
<b>Year</b>	:	2003
<b>GLP</b>	:	yes
<b>Test substance</b>	:	other TS: Phosphorus trichloride, purity not given
<b>Method</b>	:	Method is in most parts equivalent to the OECD TG 201 Alga, Growth inhibition test
<b>Remark</b>	:	Accepted new scientific name for Scenedesmus subspicatus: Desmodesmus subspicatus.
<b>Result</b>	:	The following results were observed: -Effect concentrations based on biomass growth (b): EC 10 = 21.30 mg/l EC 50 = 30.24 mg/l Determined NOEC and LOEC-values based on biomass growth (b): NOEC = 12.5 mg/l LOEC = 25 mg/l

Under pH-adjusted conditions no inhibition of the algae growth has been observed at a nominal concentration of 100 mg/l.

-Effect concentrations based on population density growth rate (r):

EC 10 = 24.89 mg/l

EC 50 = 33.41 mg/l

Determined NOEC and LOEC values based on population density growth rate (r):

NOEC = 12.5 mg/l

LOEC = 25 mg/l

Under pH-adjusted conditions no inhibition of the algae growth has been observed at a nominal concentration of 100 mg/l.

Measured pH-values:

concentration	0h	72h	av.growth rate
control	8.2	10.5	1.12
3.13 mg/l	7.8	10.4	1.17
6.25 mg/l	7.6	10.3	1.16
12.5 mg/l	7.3	10.0	1.17
25 mg/l	6.9	9.2	1.02
50 mg/l	3.5	3.5	0.0
100 mg/l	2.9	2.9	0.0
100 mg/l	8.0	10.7	1.31 (with adjusted pH)

The results of the respective replicates clearly demonstrate that the inhibitory effects observed were caused by pH-effects.

**Test condition**

- : -Static conditions
- Algal inoculum about 10E+04 cells/ml initial cell density
- 300 ml Erlenmeyer flasks with stoppers as test vessels
- Temperature during the test: 21-25 °C
- Lighting 60-120 µE/m²/s
- pH is measured at the beginning of the test and at 72 h
- Experimental design: 6 test concentrations plus 1 control, 3 replicates per concentration, 6 replicates per control, highest test concentration without algae
- Nominal test concentrations: 3.13, 6.25, 12.5, 25, 50 and 100 mg/l without adjustment of pH-value, 100 mg/l PCl<sub>3</sub> correspond to 59.7 mg/l HP(O)(OH)<sub>2</sub>
- As extreme pH-decreases were observed due to hydrolysis of the test substance additional replicates of the highest test concentration (100 mg/l) were investigated after pH-adjustment
- Cell densities measured at 24 h intervals using a microcell counter
- Inhibition of algal population measured as reduction in biomass growth (index b) and population density growth rate (index r), relative to control cultures under identical conditions
- The 72 h-EC50 values are calculated or read from the concentration/percentage response curve.
- No chemical analysis has been performed, as the test substance phosphorus trichloride hydrolyses rapidly in aqueous medium.

**Reliability**

- : (1) valid without restriction
- Comparable to guideline study

**Flag**

14.07.2005

- : Critical study for SIDS endpoint

(47)

**4.4 TOXICITY TO MICROORGANISMS E.G. BACTERIA**

- Type** : aquatic
- Species** : activated sludge
- Exposure period** : 3 hour(s)
- Unit** : mg/l
- EC50** : 9450
- EC05** : 3520

<b>Analytical monitoring</b>	: no	
<b>Method</b>	: ISO 8192 "Test for inhibition of oxygen consumption by activated sludge"	
<b>Year</b>	: 1991	
<b>GLP</b>	: yes	
<b>Test substance</b>	: other TS: Phosphorus trichloride, purity 99.7 %	
<b>Test condition</b>	: Inoculum: Activated sludge from laboratory waste water treatment plant, inoculum contained 6 g/l TS Test concentrations of Phosphorus trichloride: 1000, 1800, 3200, 5600, and 10000 mg/l, Reference substance: 3,5-dichlorophenol No analytical monitoring because test substance hydrolyses into hydrochloric acid and phosphonic acid	
<b>Reliability</b>	: (1) valid without restriction Test procedure in accordance with national standard methods	
<b>Flag</b>	: Critical study for SIDS endpoint	
09.02.2004		(44)

**4.5.1 CHRONIC TOXICITY TO FISH****4.5.2 CHRONIC TOXICITY TO AQUATIC INVERTEBRATES****4.6.1 TOXICITY TO SEDIMENT DWELLING ORGANISMS****4.6.2 TOXICITY TO TERRESTRIAL PLANTS****4.6.3 TOXICITY TO SOIL DWELLING ORGANISMS****4.6.4 TOX. TO OTHER NON MAMM. TERR. SPECIES****4.7 BIOLOGICAL EFFECTS MONITORING****4.8 BIOTRANSFORMATION AND KINETICS****4.9 ADDITIONAL REMARKS**

<b>Memo</b>	: Aquatic toxicity	
<b>Remark</b>	: Aquatic toxicity range 10-100 mg/l	
<b>Test substance</b>	: Phosphorus trichloride, purity not given	
<b>Reliability</b>	: (4) not assignable Original reference not available	
09.02.2004		(48)
<b>Memo</b>	: Harmful effects	
<b>Remark</b>	: harmful effects through shift of pH in water	
<b>Test substance</b>	: Phosphorus trichloride	
<b>Reliability</b>	: (4) not assignable Manufacturer data without proof	
09.02.2004		(22)

### 5.0 TOXICOKINETICS, METABOLISM AND DISTRIBUTION

<b>In Vitro/in vivo</b>	:	In vitro	
<b>Type</b>	:		
<b>Species</b>	:		
<b>Number of animals</b>			
<b>Males</b>	:		
<b>Females</b>	:		
<b>Doses</b>			
<b>Males</b>	:		
<b>Females</b>	:		
<b>Vehicle</b>	:		
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Result</b>	:	detailed investigations on the reaction between water and PCI <sub>3</sub> were performed. PCI <sub>3</sub> hydrolyses quickly and exothermally to give hydrochloric and phosphorous acid. Reaction is complete within few seconds when stirred.	
<b>Reliability</b>	:	(4) not assignable	
21.09.2004			(49)
<b>In Vitro/in vivo</b>	:	In vitro	
<b>Type</b>	:		
<b>Species</b>	:		
<b>Number of animals</b>			
<b>Males</b>	:		
<b>Females</b>	:		
<b>Doses</b>			
<b>Males</b>	:		
<b>Females</b>	:		
<b>Vehicle</b>	:		
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Remark</b>	:	Hydrolysis of PCI <sub>3</sub> is reported under 5.11 Metabolism	
21.09.2004			

#### 5.1.1 ACUTE ORAL TOXICITY

<b>Type</b>	:	LD50
<b>Value</b>	:	= 550 mg/kg bw
<b>Species</b>	:	rat
<b>Strain</b>	:	
<b>Sex</b>	:	no data
<b>Number of animals</b>	:	
<b>Vehicle</b>	:	other: vegetable oil
<b>Doses</b>	:	
<b>Method</b>	:	
<b>Year</b>	:	
<b>GLP</b>	:	no data
<b>Test substance</b>	:	other TS: phosphorus trichloride

<b>Result</b>	: LD50: 550 mg/kg (470-643) LD16: 430 mg/kg LD84: 675 mg/kg	
	Signs of intoxication: depression, anorexia, disturbance of movement co-ordination, fatigue, weakness, diarrhoea, appearance of an ichorous discharge from the eyes, respiratory frequency: 50-80 per minute. 20 to 40 minutes after application of the LD50: cyanosis, weakness, sweat, convulsions, short-windedness.	
	Necropsy: lungs of deceased animals were intensely red discolored. The livers were dark-gray-brown and full of blood when sectioned, the stomach was distended and hemorrhagic	
<b>Test condition</b>	: 6 Animals per dose group;	
<b>Reliability</b>	: (2) valid with restrictions short report, detailed description of toxicity	
<b>Flag</b> 21.09.2004	: Critical study for SIDS endpoint	(50) (51)
<b>Type</b>	: LD50	
<b>Value</b>	: = 18 mg/kg bw	
<b>Species</b>	: rat	
<b>Strain</b>	: Sprague-Dawley	
<b>Sex</b>	: male/female	
<b>Number of animals</b>	: 20	
<b>Vehicle</b>	: other: undiluted	
<b>Doses</b>	: 12.6 - 15.8 - 20.0 - 25.1 mg/kg	
<b>Method</b>	:	
<b>Year</b>	:	
<b>GLP</b>	: no	
<b>Test substance</b>	: other TS: phosphorus trichloride >99%	
<b>Result</b>	: Dose / Mortality 12,6 / 0/5 15,8 / 2/5 20,0 / 3/5 25,1 / 5/5	
	LD50: 18 (15.6 - 20.7) mg/kg	
	Symptoms: Reduced appetite and activity (3 to 10 days in survivors), increasing weakness, collapse, death	
	Necropsy: lung congestion, liver discoloration, acute gastrointestinal inflammation (stomach ulceration, also in some survivors)	
<b>Reliability</b>	: (2) valid with restrictions short, tabular report;	
<b>Flag</b> 21.09.2004	: Critical study for SIDS endpoint	(52) (53)
<b>Type</b>	: LD50	
<b>Value</b>	: = 200 mg/kg bw	
<b>Species</b>	: rat	
<b>Strain</b>	: Sprague-Dawley	
<b>Sex</b>	: male/female	

**Number of animals** : 50  
**Vehicle** : other: corn oil  
**Doses** : 100-200-300-400-500 mg/kg  
**Method** : other: no data  
**Year** : 1977  
**GLP** : no data  
**Test substance** : other TS: PCI3

**Remark** : Male and female rats were fasted for 24 hours before administration of the test substance. The test material was administered orally, by intubation, as a 10% solution in corn oil. Animals were observed at 1, 3, 6, 24, 48, 72 hours then daily up to 14 days. The oral LD50 was calculated. All surviving animals were killed, autopsied and observed for gross pathological organ changes.

**Result** : Mortality

100	3/10
200	5/10
300	8/10
400	9/10
500	10/10

LD50: 200 ± 29 mg/kg bw

LD16: 110 mg/kg

LD84: 365 mg/kg

**Signs of toxicity:**

Decreased locomotor activity, piloerection, ptosis, wet underside and death. Normal body activity returned within 9 days in all surviving animals.

**Necropsy:**

Rugation of the pyloric mucosa, enlarged and darkened spleen, irregular thickening of the pyloric mucosa, stomach fused to liver or liver and pancreas, chronic pulmonary disease.

**Reliability** : (2) valid with restrictions  
 Summary report, detailed description

**Flag** : Critical study for SIDS endpoint

16.01.2006

(54)

**Type** : other  
**Value** :  
**Species** : other: mice and rats  
**Strain** :  
**Sex** :  
**Number of animals** :  
**Vehicle** :  
**Doses** :  
**Method** :  
**Year** :  
**GLP** :  
**Test substance** : other TS: phosphorus trichloride

**Result** : by intragastric intubation to rats and mice PCI3 showed a median degree of toxicity, through the respiratory tract to mice, it was highly toxic. Threshold for acute toxicity to rat pulmonary macrophages was 686.8 mg/m<sup>3</sup>. PCI3 was a strong irritant to guinea pigs (no further data).

**Reliability** : (4) not assignable  
 Abstract only, no further data

21.09.2004

(55)

5.1.2 ACUTE INHALATION TOXICITY

**Type** : LC50  
**Value** : = 226 mg/m<sup>3</sup>  
**Species** : rat  
**Strain** :  
**Sex** :  
**Number of animals** :  
**Vehicle** :  
**Doses** :  
**Exposure time** :  
**Method** :  
**Year** :  
**GLP** :  
**Test substance** : other TS: phosphorus trichloride

**Remark** : exposure period not known  
**Result** : Symptoms:  
 agitation, pawing of nose, sedation, coordination disturbances, lateral position, fibrilar twitching, convulsions, reduced and strenuous respiration  
 Additionally: corrosion of lips, lacrimation, and corneal opacity. Loss of appetite and weight and weakness were recorded in survivors

**Reliability** : (2) valid with restrictions  
 limited documentation; very few details given

**Flag** : Critical study for SIDS endpoint

21.09.2004

(50) (51)

**Type** : LC50  
**Value** : = 592 mg/m<sup>3</sup>  
**Species** : rat  
**Strain** :  
**Sex** :  
**Number of animals** :  
**Vehicle** :  
**Doses** :  
**Exposure time** : 4 hour(s)  
**Method** :  
**Year** :  
**GLP** : no data  
**Test substance** : other TS: phosphorus trichloride

**Remark** : No further data given  
**Result** : LC50s of PCI3 was 104.3 ppm (592 mg/m<sup>3</sup>) for rats. The slope of the dose response curve was 4.3 +- 0.7.  
 Hydrolysis of PCI3 was about 40 percent.

Animals showed signs of irritation (agitated, restless, irregular breathing, eyes closed, chromodakryorhea) during exposure to PCI3.  
 All deaths occurred within 10 days.  
 Histopathology revealed necrosis in the epithelium and its supporting structures in the nostrils. Pulmonary damage was negligible. The main site of pathologic action ws the kidney with nephrosis of the tubules of the cortico-medullary region.

**Test condition** : 20 female rats per group; whole body  
 Animals were observed and deaths were recorded up to 14 days post exposure. Median lethal concentrations (LC50) were computed.  
 Hydrolysis of the test substance in the atmosphere: ca. 40%

**Reliability** : (2) valid with restrictions  
 few details reported, number of groups and dose regimen missing

**Flag** : Critical study for SIDS endpoint

21.09.2004

(56)

**Type** : LC50  
**Value** : > 118 ppm  
**Species** : rat  
**Strain** : Sprague-Dawley  
**Sex** : male/female  
**Number of animals** :  
**Vehicle** :  
**Doses** :  
**Exposure time** : 4 hour(s)  
**Method** : other: EPA TSCA  
**Year** : 1982  
**GLP** : yes  
**Test substance** : other TS: phosphorus trichloride:>99.9%

**Remark** : There is a large discrepancy between nominal and measured concentrations  
 e.g. high dose  
 nominal concentration: 452.9 ppm = 2540 µg/l  
 gravimetric measured conc.: (80 ppm) = 449 µg/l  
 chloride measured: 159 ppm = (893 µg/l)  
 inorganic phosphorus conc.: 118 ppm = (627 µg/l)  
 Values in brackets are calculated. Other figures are stated in the report.

The correct value cannot be drawn from the report. Possibly the unit (µg/l) for the gravimetric measured concentrations is wrong and should be read as "ppm" analogous to the other tables. This would give a gravimetric value near the nominal concentration. It is unclear how the nominal concentration was determined.

If one accepts the measured values inspite of the large deviations, then the LC50 should be stated with regard to some measured value, not as a nominal value

**Result** : Mortality:  
 453 ppm: nose only: 1m+1f (day 7)  
 whole body: 2m+1f (day 7+8)  
  
 LC50 > 453 ppm (nominal)(2582 mg/m³)  
 gravimetric: 449 mg/m³;  
 analytic: 159,2 ppm (based on chloride); 118,2 ppm (based on inorg. P)

Body weight: transient decrease at concentrations >= 278 ppm (1585 mg/m³)

Clinical observation:  
 dose dependent incidence and severity in all treated groups: wheezing, labored respiration, localized sores at nose, mouth and paws, swollen nose, blocked nostrils, red crusts around the eyes.

**Source** : Astaris  
**Test condition** : Chamber analysis :  
 Group: C- low1- mid - high - low2  
 Nominal Concentrations: 0-103.4-278.1- 452.9 -28.5 ppm  
 0- 580 - 1560- 2540 - 160 µg/l  
 gravimetric concentration: 0-281.8-354.3- 449.5 -57.9 µg/l

	Analysis	
	Chloride:	0-101.7-219.6- 159.2 -16.2 ppm
	inorg. phosphorus:	0- 81.8-149.6- 118.2 -14.3 ppm
	mass median aerodynamic diameter:	
	< 0.65 µm	in groups 2-4;
	= 1.75 µm	in group 5
	Animals: 5m+5f per group per exposure type	
	Nose only as well as whole body exposure were employed	
	Duration: 4h	
<b>Reliability</b>	:	(2) valid with restrictions
		Guideline study, full report available, GLP
		Analysis of atmosphere gives variable results, depending on method
<b>Flag</b>	:	Critical study for SIDS endpoint
22.09.2004		(57)
<b>Type</b>	:	LC50
<b>Value</b>	:	> 453 ppm
<b>Species</b>	:	rat
<b>Strain</b>	:	Sprague-Dawley
<b>Sex</b>	:	male/female
<b>Number of animals</b>	:	40
<b>Vehicle</b>	:	
<b>Doses</b>	:	
<b>Exposure time</b>	:	4 hour(s)
<b>Method</b>	:	
<b>Year</b>	:	
<b>GLP</b>	:	
<b>Test substance</b>	:	other TS: phosphorus trichloride
<b>Remark</b>	:	453 ppm = 2582 mg/m <sup>3</sup>
<b>Result</b>	:	LC50 (4h): >453 ppm (nominal) 3 male and 2 female animals died. Clinical signs: reduction of body weight, wheezing, labored respiration, localized sores, swollen nose, blocked nostrils, red crusts around the eyes.
<b>Test condition</b>	:	4 Groups of 5m+5f Sprague-Dawley rats were exposed for 4 h to PCI3. Afterwards they were observed for 14 days
<b>Reliability</b>	:	(2) valid with restrictions
		Summary, no further data
<b>Flag</b>	:	Critical study for SIDS endpoint
21.09.2004		(53)
<b>Type</b>	:	LC50
<b>Value</b>	:	132 mg/m <sup>3</sup>
<b>Species</b>	:	rat
<b>Strain</b>	:	
<b>Sex</b>	:	
<b>Number of animals</b>	:	
<b>Vehicle</b>	:	
<b>Doses</b>	:	
<b>Exposure time</b>	:	
<b>Method</b>	:	
<b>Year</b>	:	
<b>GLP</b>	:	
<b>Test substance</b>	:	other TS: phosphorus trichloride
<b>Reliability</b>	:	(4) not assignable
		RTECS data; Report not available

21.09.2004 (58) (59)

**Type** : LC50  
**Value** : > 20.29 mg/l  
**Species** : rat  
**Strain** :  
**Sex** :  
**Number of animals** : 10  
**Vehicle** :  
**Doses** : 20.29 mg/l  
**Exposure time** : 1 hour(s)  
**Method** :  
**Year** : 1977  
**GLP** :  
**Test substance** : other TS: PCI3

**Remark** : 10 rats were exposed to an aerosol of the test material. Observation period not given.

**Result** : Mortality  
 20.29 0/10

Signs of toxicity:

Face pawing, shovel nosing, salivation, nasal discharge, escape motions, decreased locomotor activity, redness on eye membrane and ears, lacrimation

**Reliability** : (2) valid with restrictions  
 Short report, limited description

**Flag** : Critical study for SIDS endpoint

16.01.2006 (60)

**Type** : other  
**Value** :  
**Species** : rat  
**Strain** :  
**Sex** :  
**Number of animals** :  
**Vehicle** :  
**Doses** :  
**Exposure time** :  
**Method** :  
**Year** :  
**GLP** :  
**Test substance** : other TS: phosphorus trichloride

**Result** : The threshold of acute toxicity of PCI3 was 686.8 mg/m<sup>3</sup> in rat lung macrophages

**Reliability** : (4) not assignable  
 Abstract only, no further data

21.09.2004 (55)

**Type** : other: Inhalation hazard test  
**Value** :  
**Species** : rat  
**Strain** : Wistar  
**Sex** : male/female  
**Number of animals** : 60  
**Vehicle** :  
**Doses** :  
**Exposure time** :  
**Method** :  
**Year** : 1977

<b>GLP</b>	:	no	
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Result</b>	:	Mortality: none after 0.5 and 1 minute; 100% after 2, 5, and 10 minutes exposure within 5 to 60 minutes Symptoms: agitation, lethargy, irregular gasping, wheezing, sneezing, prostration, ptosis Necropsy: discoloration of lungs, after 2 and more minutes: corneal opacity and swelling of nose.	
<b>Test condition</b>	:	Animals: 6 male and 6 female Wistar rats per group Exposure time: 0.5, 1, 2, 5, 14 minutes Observation time: up to 14 days after exposure Observation: clinical findings, necropsy	
<b>Test substance</b>	:	chemically pure substance	
<b>Reliability</b>	:	(2) valid with restrictions Detailed report available; non-standard test,	
<b>Flag</b>	:	Critical study for SIDS endpoint	
21.09.2004			(61)
<b>Type</b>	:	other	
<b>Value</b>	:		
<b>Species</b>	:	mouse	
<b>Strain</b>	:		
<b>Sex</b>	:		
<b>Number of animals</b>	:		
<b>Vehicle</b>	:		
<b>Doses</b>	:		
<b>Exposure time</b>	:		
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Result</b>	:	PCl3 was "highly toxic"	
<b>Reliability</b>	:	(4) not assignable Abstract only, no further data	
21.09.2004			(55)
<b>Type</b>	:	other: LClo	
<b>Value</b>	:	= 600 ppm	
<b>Species</b>	:	mouse	
<b>Strain</b>	:		
<b>Sex</b>	:		
<b>Number of animals</b>	:		
<b>Vehicle</b>	:		
<b>Doses</b>	:		
<b>Exposure time</b>	:	10 minute(s)	
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Remark</b>	:	600 ppm = 3420 mg/m <sup>3</sup>	
<b>Reliability</b>	:	(4) not assignable Report not available	
21.09.2004			(62)

<b>Type</b>	:	LCLo	
<b>Value</b>	:	3500 mg/m <sup>3</sup>	
<b>Species</b>	:	rabbit	
<b>Strain</b>	:		
<b>Sex</b>	:		
<b>Number of animals</b>	:		
<b>Vehicle</b>	:		
<b>Doses</b>	:		
<b>Exposure time</b>	:		
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Reliability</b>	:	(4) not assignable RTECS data; Report not available	
21.09.2004			(58) (59)
<b>Type</b>	:		
<b>Value</b>	:		
<b>Species</b>	:	rabbit	
<b>Strain</b>	:		
<b>Sex</b>	:		
<b>Number of animals</b>	:	7	
<b>Vehicle</b>	:		
<b>Doses</b>	:	4-30, 40-330, 930-3870 ppm (22.8-171, 228-1881, 2109- 22059 mg/m <sup>3</sup> )	
<b>Exposure time</b>	:		
<b>Method</b>	:		
<b>Year</b>	:	1904	
<b>GLP</b>	:	no	
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Result</b>	:	Symptoms: low dose: Sedation, reduced respiratory frequency medium dose: agitation, nasal discharge, rhinitis, sedation, reduced respiratory frequency, irregular respiration, corneal corrosion, dyspnea, tremor, lacrimation, salivation, high dose: sneezing, agitation, closed eyes, white secretion from eyes, corneal opacity (at 118 min), death, weight loss  Necropsy: medium dose: hyperemia of trachea, slight lung edema, partial infiltration of right lung high dose: severe lung edema, emphysema, severe catarrh, corrosion of tongue and cornea	
<b>Test condition</b>	:	2-3 rabbits per concentration range were exposed to PCI3 at concentrations of 4-30, 40-330, or 370-3870 ppm and observed for clinical signs for 6, 6-10, or 3-4 hours respectively	
<b>Reliability</b>	:	(2) valid with restrictions Early study, detailed description of findings, non-standard test animal	
<b>Flag</b>	:	Critical study for SIDS endpoint	
21.09.2004			(63)
<b>Type</b>	:	LCLo	
<b>Value</b>	:	500 mg/m <sup>3</sup>	
<b>Species</b>	:	cat	
<b>Strain</b>	:		
<b>Sex</b>	:		
<b>Number of animals</b>	:		
<b>Vehicle</b>	:		

<b>Doses</b>	:		
<b>Exposure time</b>	:		
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Reliability</b>	:	(4) not assignable RTECS data; Report not available	
21.09.2004			(58) (59)
<b>Type</b>	:	other: TCLo	
<b>Value</b>	:	10 - 40 mg/m <sup>3</sup>	
<b>Species</b>	:	cat	
<b>Strain</b>	:		
<b>Sex</b>	:		
<b>Number of animals</b>	:		
<b>Vehicle</b>	:		
<b>Doses</b>	:		
<b>Exposure time</b>	:		
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Remark</b>	:	40 mg/m <sup>3</sup> for 6h 10 mg/m <sup>3</sup> for 1h	
<b>Reliability</b>	:	(4) not assignable RTECS data; Report not available	
21.09.2004			(58) (59)
<b>Type</b>	:		
<b>Value</b>	:		
<b>Species</b>	:	cat	
<b>Strain</b>	:		
<b>Sex</b>	:	no data	
<b>Number of animals</b>	:	13	
<b>Vehicle</b>	:		
<b>Doses</b>	:	4-30, 40-330, 370-3870 ppm (22.8-171, 228-1881, 2109- 22059 mg/m <sup>3</sup> )	
<b>Exposure time</b>	:		
<b>Method</b>	:		
<b>Year</b>	:	1904	
<b>GLP</b>	:	no	
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Remark</b>	:	According to the author the findings were almost identical to findings produced by HCl vapors in earlier studies. PCI3 was more effective at low doses	
<b>Result</b>	:	Symptoms: low dose: salivation, irregular respiration, dyspnea, cough, lacrimation, sedation medium dose: immediate agitation, reduced irregular respiratory frequency, cough, sneezing, breathing through open mouth, foamy salivation, nasal discharge, rhinitis, conjunctivitis, dyspnea, loss of weight high dose: as above but earlier onset and more severe; death  Necropsy: low dose: discoloration of nasal cavity, injection of trachea, lung edema, ecchymosis,	

	medium dose: corneal opacity, conjunctivitis, nasal corrosion, trachea contain foamy red liquid, edema of epiglottis, severe emphysema	
	high dose: as medium dose	
<b>Test condition</b>	: 1-5 cats per concentration range were exposed to PCI <sub>3</sub> at concentrations of 4-30, 40-330, or 370-3870 ppm and observed for clinical signs for 3-6, 6-10, or 3-6.5 hours respectively	
	additionally a gross necropsy was performed	
<b>Reliability</b>	: (2) valid with restrictions	
	Early study, detailed description of findings, non-standard test animal	
<b>Flag</b>	: Critical study for SIDS endpoint	
21.09.2004		(63)
<b>Type</b>	: LC50	
<b>Value</b>	: = 285 mg/m <sup>3</sup>	
<b>Species</b>	: guinea pig	
<b>Strain</b>	:	
<b>Sex</b>	:	
<b>Number of animals</b>	:	
<b>Vehicle</b>	:	
<b>Doses</b>	:	
<b>Exposure time</b>	: 4 hour(s)	
<b>Method</b>	:	
<b>Year</b>	:	
<b>GLP</b>	:	
<b>Test substance</b>	: other TS: phosphorus trichloride	
<b>Remark</b>	: no further data given	
<b>Result</b>	: LC50s of PCI <sub>3</sub> was 50.1 ppm (285 mg/m <sup>3</sup> ) for guinea pigs. The slope of the dose response curve was 5.7 +/- 0.9. Hydrolysis of PCI <sub>3</sub> was about 40 percent. Animals showed signs of irritation (agitated, restless, irregular breathing, eyes closed, chromodakryorhea) during exposure to PCI <sub>3</sub> . All deaths occurred within 10 days. Histopathology revealed necrosis in the epithelium and its supporting structures in the nostrils. Pulmonary damage was negligible. The main site of pathologic action was the kidney with nephrosis of the tubules of the cortico-medullary region.	
<b>Test condition</b>	: 10 male guinea pigs per group; whole body Animals were observed and deaths were recorded up to 14 days post exposure. Median lethal concentrations (LC50) were computed.	
<b>Reliability</b>	: (2) valid with restrictions the differentiation between the study in rats and the guinea pig study regarding the observed symptoms is not possible	
<b>Flag</b>	: Critical study for SIDS endpoint	
21.09.2004		(56)
<b>Type</b>	: LC50	
<b>Value</b>	: 63.5 mg/m <sup>3</sup>	
<b>Species</b>	: guinea pig	
<b>Strain</b>	:	
<b>Sex</b>	:	
<b>Number of animals</b>	:	
<b>Vehicle</b>	:	
<b>Doses</b>	:	
<b>Exposure time</b>	:	
<b>Method</b>	:	
<b>Year</b>	:	
<b>GLP</b>	:	
<b>Test substance</b>	: other TS: phosphorus trichloride	

**Reliability** : (4) not assignable  
RTECS data; Report not available

21.09.2004

(58) (59)

### 5.1.3 ACUTE DERMAL TOXICITY

**Type** : LD50  
**Value** : > 1000 mg/kg bw  
**Species** : rabbit  
**Strain** : New Zealand white  
**Sex** : male/female  
**Number of animals** : 5  
**Vehicle** : other: undiluted  
**Doses** : 1000 - 1260 - 2000 mg/kg  
**Method** :  
**Year** :  
**GLP** : no  
**Test substance** : other TS: phosphorus trichloride: >99%

**Result** : Dose / Mortality  
1000 / 0/2  
1260 / 1/2  
2000 / 1/1

LDI<sub>0</sub> = 1260 mg/kg

**Symptoms:**

Reduced appetite and activity (3 to 15 days in survivors), increasing weakness, collapse, death

**Necropsy:**

dead animals: lung hyperemia, liver discoloration, kidney and spleen darkened, enlarged gall bladder, slight gastrointestinal inflammation  
survivors: kidney and spleen discolored, slight gastrointestinal inflammation

**Test condition** : semi-occlusive; 24 h exposure

**Reliability** : (2) valid with restrictions  
short report; few details given

**Flag** : Critical study for SIDS endpoint

21.09.2004

(52) (53)

**Type** : LD50  
**Value** : > 250 mg/kg bw  
**Species** : rabbit  
**Strain** : New Zealand white  
**Sex** : male  
**Number of animals** : 12  
**Vehicle** : other: none  
**Doses** : 250 mg/kg

**Method** :  
**Year** : 1977  
**GLP** :  
**Test substance** : other TS: PCI3

**Remark** : Range finder:  
Two animals were used, one at each of 250 and 500 mg/kg bw.

**Main study:**

The test material was administered undiluted to intact skin for 6 animals

and to abraded skin for the other 6 animals. Animals were observed at 1, 3, 6, 24, 48, 72 hours then daily up to 14 days.

**Result** : Mortality (range finder)  
                   250 0/1  
                   500 1/1  
 Based on the corrosive effects at these doses, 250 mg/kg was used in the main study.  
 Main Study  
 250 0/12  
 LD50: > 250 mg/kg bw  
 Signs of toxicity:  
     In the range finding study the test substance produced oedema, erythema, necrosis, eschar, and death at 500 mg/kg.  
     In the range finder and main study, the animals exposed to 250 mg/kg showed oedema, erythema, necrosis and eschar but no mortality.

**Reliability** : (2) valid with restrictions  
 Short report, detailed description

16.01.2006 (64)

#### 5.1.4 ACUTE TOXICITY, OTHER ROUTES

##### 5.2.1 SKIN IRRITATION

**Species** : rabbit  
**Concentration** :  
**Exposure** :  
**Exposure time** :  
**Number of animals** :  
**Vehicle** :  
**PDII** :  
**Result** : highly corrosive  
**Classification** :  
**Method** :  
**Year** : 1984  
**GLP** :  
**Test substance** : other TS: phosphorus trichloride

**Result** : PCI3 caused skin corrosion after 60 seconds of contact to skin even when the skin was cleaned using water, lutrol, or paraffin oil  
**Test condition** : 100 µl of PCI3 were applied to the shaved skin of rabbits (number not stated). The treated area was washed 60 seconds thereafter with water, lutrol, or paraffin oil. Changes were recorded

**Reliability** : (2) valid with restrictions  
**Flag** : Critical study for SIDS endpoint

21.09.2004 (65)

**Species** : rabbit  
**Concentration** :  
**Exposure** :  
**Exposure time** :  
**Number of animals** :  
**Vehicle** :

<b>PDII</b>	:		
<b>Result</b>	:		
<b>Classification</b>	:		
<b>Method</b>	:	other: no data	
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Remark</b>	:	thickening in skin folds, bleeding	
<b>Test condition</b>	:	4 drops were applied to the shaved skin	
<b>Reliability</b>	:	(4) not assignable Short notice; no details given	
21.09.2004			(50)
<b>Species</b>	:	rabbit	
<b>Concentration</b>	:		
<b>Exposure</b>	:		
<b>Exposure time</b>	:		
<b>Number of animals</b>	:		
<b>Vehicle</b>	:		
<b>PDII</b>	:		
<b>Result</b>	:		
<b>Classification</b>	:		
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Result</b>	:	A correlation between the inhalation irritation threshold for humans and rats on one hand, and skin irritation for rabbits on the other, was assessed for PCI3 and other chemicals. The degree of hyperemia following the dermal application to rabbits was correlated with an increase in the thickness of the skin fold. The skin irritation was concn.-dependent. The inhalation toxicity may be approx. assessed from skin irritation tests.	
<b>Test condition</b>	:	Literature review; no experimental details	
<b>Reliability</b>	:	(4) not assignable non standard evaluation and comparison scheme; no experimental data	
21.09.2004			(66)
<b>Species</b>	:	rabbit	
<b>Concentration</b>	:	undiluted	
<b>Exposure</b>	:	Semiocclusive	
<b>Exposure time</b>	:	24 hour(s)	
<b>Number of animals</b>	:		
<b>Vehicle</b>	:		
<b>PDII</b>	:		
<b>Result</b>	:	corrosive	
<b>Classification</b>	:		
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride: >99%	
<b>Result</b>	:	Loosening about edge of scab in ten to fourteen days showing injury in depth	
<b>Test condition</b>	:	New Zealand White rabbits were treated on the shaved skin	
<b>Reliability</b>	:	(2) valid with restrictions	

	Abstract only, no further data	
<b>Flag</b> 21.09.2004	: Critical study for SIDS endpoint	(52) (53)
<b>Species</b>	: rabbit	
<b>Concentration</b>	: undiluted	
<b>Exposure</b>	:	
<b>Exposure time</b>	: 24 hour(s)	
<b>Number of animals</b>	: 6	
<b>Vehicle</b>	: other: none	
<b>PDII</b>	:	
<b>Result</b>	: corrosive	
<b>Classification</b>	:	
<b>Method</b>	:	
<b>Year</b>	: 1977	
<b>GLP</b>	:	
<b>Test substance</b>	: other TS: PCI3	
<b>Remark</b>	: 0.5 ml of the test material was administered undiluted to abraded skin. Animals were observed for signs of dermal irritation on removal of dressing and at 72 hours.	
<b>Result</b>	: Application of the test material caused immediate tissue destruction. Due to the severity of tissue destruction, a primary irritation index could not be calculated.	
<b>Reliability</b>	: (2) valid with restrictions Short report, detailed description	
<b>Flag</b> 16.01.2006	: Critical study for SIDS endpoint	(67)
<b>Species</b>	: rat	
<b>Concentration</b>	:	
<b>Exposure</b>	: no data	
<b>Exposure time</b>	:	
<b>Number of animals</b>	:	
<b>Vehicle</b>	:	
<b>PDII</b>	:	
<b>Result</b>	:	
<b>Classification</b>	:	
<b>Method</b>	:	
<b>Year</b>	:	
<b>GLP</b>	: no data	
<b>Test substance</b>	: other TS: phosphorus trichloride	
<b>Remark</b>	: observation in acute inhalation study	
<b>Result</b>	: The irritant effect of PCI3 is more pronounced, which is demonstrated by turbidity of the cornea, appearance of skin ulcers around the mouth and nose of the experimental animals and by a well developed pathomorphological picture of irritation of the respiratory passages.	
<b>Reliability</b>	: (2) valid with restrictions short report, few detail, no data on study design	
<b>Flag</b> 21.09.2004	: Critical study for SIDS endpoint	(51)

### 5.2.2 EYE IRRITATION

<b>Species</b>	: rabbit
<b>Concentration</b>	:

**Dose** :  
**Exposure time** :  
**Comment** :  
**Number of animals** :  
**Vehicle** :  
**Result** :  
**Classification** :  
**Method** : other: no data  
**Year** :  
**GLP** :  
**Test substance** : other TS: phosphorus trichloride  
  
**Remark** : irreversible necrosis in the eye causing loss of vision.  
**Reliability** : (4) not assignable  
 limited documentation; no experimental details given  
**Flag** : Critical study for SIDS endpoint  
 21.09.2004 (50)  
  
**Species** : rabbit  
**Concentration** : undiluted  
**Dose** : .1 ml  
**Exposure time** :  
**Comment** :  
**Number of animals** :  
**Vehicle** :  
**Result** : corrosive  
**Classification** :  
**Method** :  
**Year** :  
**GLP** :  
**Test substance** : other TS: phosphorus trichloride:>99%  
  
**Result** : corrosive in 5 seconds  
 immediate discomfort with severe pawing, squealing, trashing about the  
 stocks, eyes tightly closed  
**Test condition** : PCI3 was applied to the conjunctival sac. Effects were scored according to  
 Draize  
**Reliability** : (2) valid with restrictions  
 Abstract only, few details given  
**Flag** : Critical study for SIDS endpoint  
 21.09.2004 (52) (53)  
  
**Species** : rabbit  
**Concentration** : undiluted  
**Dose** : .1 ml  
**Exposure time** :  
**Comment** :  
**Number of animals** : 6  
**Vehicle** : none  
**Result** : corrosive  
**Classification** :  
**Method** :  
**Year** : 1977  
**GLP** :  
**Test substance** : other TS: PCI3  
  
**Remark** : The test material was instilled undiluted into the conjunctival sac of one eye  
 of each rabbit. Ocular reactions were graded at 1, 24, 48, 72 hours, 4 and 7  
 days.

**Result** : Scoring could not be done due to irreversible damage to eye tissue on contact.  
**Reliability** : (2) valid with restrictions  
 Short report, detailed description  
**Flag** : Critical study for SIDS endpoint  
 16.01.2006 (67)

### 5.3 SENSITIZATION

**Type** : Guinea pig maximization test  
**Species** : guinea pig  
**Number of animals** :  
**Vehicle** :  
**Result** : not sensitizing  
**Classification** :  
**Method** : other  
**Year** :  
**GLP** :  
**Test substance** : other TS: 1% hydrochloric acid in 70% ethanol.  
**Remark** : Sensitization was not induced in 15 guinea pigs that were given two intradermal injections and a covered application (48-hr) of 1% HCl (in ethanol of undefined concentration) and challenged 2 weeks later by a similar 24-hr covered.  
 No. of animals with skin reaction at challenge:  
 Treated: 0/15 Control group: 0/6  
 22.07.2004 (68)

**Type** : Mouse ear swelling test  
**Species** : mouse  
**Number of animals** :  
**Vehicle** :  
**Result** : not sensitizing  
**Classification** :  
**Method** : other  
**Year** :  
**GLP** : no  
**Test substance** : other TS: 1% hydrochloric acid in 70% ethanol.  
**Remark** : Four consecutive daily uncovered applications of 1% HCl solution in 70% ethanol to the abdominal skin were followed 7 days later by a challenge with a 5% uncovered application to the ear. No evidence of sensitization was seen.  
 No. of animals: Not stated Control group: Not stated  
 22.07.2004 (68)

### 5.4 REPEATED DOSE TOXICITY

**Type** : Sub-acute  
**Species** : rat  
**Sex** : male/female  
**Strain** : Sprague-Dawley  
**Route of admin.** : inhalation  
**Exposure period** : 4 weeks  
**Frequency of treatm.** : 5 days/week  
**Post exposure period** : none  
**Doses** : 0.5, 3.0, and 10 ppm

<b>Control group</b>	:	yes	
<b>NOAEL</b>	:	= 3 ppm	
<b>Method</b>	:	other: EPA/TSCA	
<b>Year</b>	:	1982	
<b>GLP</b>	:	yes	
<b>Test substance</b>	:	other TS: phosphorus trichloride: >99.9 %	
<b>Remark</b>	:	histopath: squamous metaplasia of respiratory epithelium; focal suppurative inflammation of the anterior nasal region.	
<b>Result</b>	:	1) 0.5 ppm: no effects 2) 3 ppm: no effects 3) 10 ppm: nasal cavity and turbinates, histopathology:proteinaceous fluid with suppurative inflammation; inflammation and squamous metaplasia of respiratory epithelium	
<b>Test condition</b>	:	Animals: 15 per sex per group Exposure: 6h*5d*4W; whole body Parameters: mortality, ophthalmoscopy, clinical findings, body weights, clinical pathology (hematology; clinical chemistry; urinalysis), organ weights (absolute; relative to brain and body weight), gross pathology, histopathology	
<b>Reliability</b>	:	(1) valid without restriction Guideline study, full report available	
<b>Flag</b>	:	Critical study for SIDS endpoint	
08.07.2005			(69)
<b>Type</b>	:	Sub-chronic	
<b>Species</b>	:	rat	
<b>Sex</b>	:	male/female	
<b>Strain</b>	:	other: F344/CrlBr and Sprague-Dawley	
<b>Route of admin.</b>	:	inhalation: aerosol	
<b>Exposure period</b>	:	90 days	
<b>Frequency of treatm.</b>	:	6h/d x 5d/w	
<b>Post exposure period</b>	:		
<b>Doses</b>	:	0-10-20-50ppm (nominal)	
<b>Control group</b>	:	yes, concurrent vehicle	
<b>Method</b>	:		
<b>Year</b>	:	1983	
<b>GLP</b>	:	yes	
<b>Test substance</b>	:	other TS: HCl	
<b>Result</b>	:	NOAEC = 20 ppm (disregarding irritation) Mortality: none  Body weight (50 ppm) and food consumption (20 and 50 ppm) were reduced.  clinical pathology: no effects in hematology, clin. chemistry, urinalysis  Histopathology: Rhinitis in the nasal cavity, detected in all treatment groups, accompanied with occasional hyperkeratosis, was the only finding. No treatment related findings were present in other tissues , e.g. the reproductive organs testes, ovaries, etc. (tissues investigated: see TC).	
<b>Test condition</b>	:	Age of animals at start: 6 -7 weeks Number of animals: 31 m +21 f per group Exposure: whole body; (6h*5d/w)*90d Vehicle : air  Parameter examined: Clinical signs (daily), body weight (weekly), food consumption (weekly),	

urinalysis, hematology, clinical chemistry, necropsy, organ weights, histopathology [nasal turbinates, trachea, lung, brain, heart, kidney, liver, testis, adrenal, duodenum, eyes and optic nerve, mesenteric lymph nodes, aorta, sternum bone, ear canal, bone marrow, colon, epididymis, jejunum, mandibular lymph nodes, oviducts, ovaries, prostate, skin, pituitary glands, spinal cord, sciatic nerve, peripheral nerve, salivary gland, spleen, thyroid glands, urinary bladder, uterus, thymus, fore and glandular stomach, pancreas, parathyroid, skeletal muscle, seminal vesicle, tongue, femur bone, cecum, esophagus, ileum, lacrimal gland, mammary gland, larynx]

Statistics:

ANOVA for parametric data; significant values were studied with Tukey's or Scheffe's test of multiple comparison

Non parametric data were evaluated according to Kruskal- Wallis ANOVA discontinuous data were compared using Fisher' Exact or Chi-square Probability Test

**Reliability** : (2) valid with restrictions  
**Flag** : Critical study for SIDS endpoint  
13.02.2006 (70)

**Type** : Sub-chronic  
**Species** : mouse  
**Sex** : male/female  
**Strain** : B6C3F1  
**Route of admin.** : inhalation  
**Exposure period** : 90 days  
**Frequency of treatm.** : 6h\*5d/w\*90d  
**Post exposure period** : none  
**Doses** : 0-10-20-50 ppm  
**Control group** : yes  
**Method** :  
**Year** :  
**GLP** : yes  
**Test substance** : other TS: HCl

**Result** : Moratlity: 4/52 (low dose)  
0/52 (mid dose)  
2/52 (high dose)  
According to authors death was not related to exposure

Body weight gain: all exposed high dose animals affected  
clinical pathology: no effects in hematology, clin. chemistry, urinalysis

Organ weights: decreased liver weight (high dose)

Pathology: gross lesions in perioral tissue and toes (swelling, ulcerative dermatitis); cheilitis with accumulating hemosiderin-laden macropahges and eosinophilic globules in the ephthelium of nasal turbinates were observed in exposed mice

**Test condition** : Animals: 31 m +m 21 f per group  
Exposure: whole body; (6h\*5d/w)\*90d  
**Reliability** : (2) valid with restrictions  
**Flag** : Critical study for SIDS endpoint  
27.09.2004 (70)

5.5 GENETIC TOXICITY 'IN VITRO'

<b>Type</b>	:	Ames test	
<b>System of testing</b>	:	S. typhimurium TA1535, TA100, TA1537, D3052, TA1538, TA98, C3076, G46	
<b>Test concentration</b>	:	0.1-1000µg/mL	
<b>Cycotoxic concentr.</b>	:		
<b>Metabolic activation</b>	:	with and without	
<b>Result</b>	:	negative	
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Test condition</b>	:	positive and negative controls used in both activated and non-activated tests	
<b>Reliability</b>	:	(2) valid with restrictions Non-standard test: gradient test	
<b>Flag</b>	:	Critical study for SIDS endpoint	(71)
21.09.2004			
<b>Type</b>	:	Bacterial gene mutation assay	
<b>System of testing</b>	:	E. coli WP2, WP2uvrA-	
<b>Test concentration</b>	:	0.1 - 1000 µg/mL	
<b>Cycotoxic concentr.</b>	:		
<b>Metabolic activation</b>	:	with and without	
<b>Result</b>	:	negative	
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Test condition</b>	:	positive and negative controls used in both activated and non-activated tests	
<b>Reliability</b>	:	(2) valid with restrictions Non-standard test: gradient test	
<b>Flag</b>	:	Critical study for SIDS endpoint	(71)
21.09.2004			
<b>Type</b>	:	Ames test	
<b>System of testing</b>	:	S. typhimurium TA1535, TA100, TA1537, TA1538, TA98 S. cerevisiae D4	
<b>Test concentration</b>	:	0.001 to 5.0 µl per plate	
<b>Cycotoxic concentr.</b>	:	5.0 µl per plate	
<b>Metabolic activation</b>	:	with and without	
<b>Result</b>	:	negative	
<b>Method</b>	:		
<b>Year</b>	:	1977	
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: PCI3	
<b>Reliability</b>	:	(2) valid with restrictions short report; limited description	
<b>Flag</b>	:	Critical study for SIDS endpoint	(72)
16.01.2006			

#### 5.6 GENETIC TOXICITY 'IN VIVO'

**Type** : Cytogenetic assay

<b>Species</b>	:	human	
<b>Sex</b>	:	no data	
<b>Strain</b>	:		
<b>Route of admin.</b>	:		
<b>Exposure period</b>	:		
<b>Doses</b>	:		
<b>Result</b>	:	negative	
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Result</b>	:	The incidence of chromosomal aberrations in workers with contact to PCI3 was 1.58 % The incidence of chromosomal aberrations in workers without contact to PCI3 was 1.4 % There was no significant difference	
<b>Test condition</b>	:	Human peripheral lymphocytes were investigated  24 Worker with contact to PCI3 and 10 workers without contact to PCI3 were examined; there were smokers in both groups;  0.45 ml blood were collected, heparinized and cultured at 37 C for 72 hours. 4 hours before end of culture colchicine was added. Slides were prepared and exyamined microscopically	
<b>Reliability</b>	:	(2) valid with restrictions short report	
<b>Flag</b>	:	Critical study for SIDS endpoint	(73)
21.09.2004			
<b>Type</b>	:	Micronucleus assay	
<b>Species</b>	:	mouse	
<b>Sex</b>	:	male	
<b>Strain</b>	:	other: Kunming	
<b>Route of admin.</b>	:	i.p.	
<b>Exposure period</b>	:	5 days	
<b>Doses</b>	:	10,94-21,88-43,75 mg/kg	
<b>Result</b>	:	negative	
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: phosphorus trichloride	
<b>Result</b>	:	PCI3 did not induce chromosome aberration or micronucleus rate increases in bone marrow cells or abnormalities in sperm of mice. Chromosomal aberrations: 0,8%, 0,6% or 0,6% in treated animals (P >0,05; X2 = 1,7362) = non significant negative control: 0,2% positive control: 11,7%. P <0,01 (X2 = 33,31) =significant.	
<b>Test condition</b>	:	Animals were injected 5 times and killed on day 6. 3-4 h before death animals were treated with colchicine (ip) both femora were prepared and the bone marrow collected for the determination of chromosomal aberrations and micro-nuclei Vehicle: cod liver oil pos. control: cyclophosphamide For chromosomal aberrations 100 cells were scored and for micro-nuclei 1000 polychromatic erythrocytes were examined. Statistics: Chi square test	

**Reliability** : (2) valid with restrictions  
Detailed publication  
**Flag** : Critical study for SIDS endpoint  
27.09.2004 (73)

**5.7 CARCINOGENICITY**

**Species** : rat  
**Sex** : male  
**Strain** : Sprague-Dawley  
**Route of admin.** : inhalation  
**Exposure period** : Maximum, 128 weeks (for life)  
**Frequency of treatm.** : 6 hours/day, 5 days/week  
**Post exposure period** : No  
**Doses** : 10.0 ppm (14.9 mg/m<sup>3</sup>)  
**Result** :  
**Control group** : yes, concurrent vehicle  
**Method** : other  
**Year** :  
**GLP** : no data  
**Test substance** : other TS: Hydrogen chloride, purity 99.0% grade, Matheson Gas Products

**Remark** : Method: Three groups of 100 male rats, nine weeks old, were unexposed (colony controls), exposed by inhalation to air (air control) or exposed to 10 ppm of hydrogen chloride. Complete necropsy was performed on each animal and particular attention was give to the respiratory tract.

**Result** : Mortality did not significantly differ from air control group. No preneoplastic or neoplastic nasal lesion was observed in any group, but hyperplasia of the larynx and trachea was observed in treated animals (22/99 and 26/99, respectively). Tumor responses were similar in the treated and control groups, the total incidences of tumours at various sites being 19/99, 25/99 and 24/99 in treated, air control and colony control animals, respectively.

	Observation	HCl	Air	Colony
	No. examined	99	99	99
Larynx	Hyperplasis	22	2	2
	Squamous metaplasia	0	0	0
Trachea	Hyperplasis	26	6	2
	Squamous metaplasia	0	0	0
Nasal mucosa	Rhinitis	81	72	70
	Epithelial or squamous hyperplasia	62	51	45
	Squamous metaplasia	9	5	6
	Total no. of tumors	19	25	24

**Reliability** : (2) valid with restrictions  
main study details reported  
**Flag** : Critical study for SIDS endpoint  
21.09.2004 (74)

**Species** : rat  
**Sex** : male/female  
**Strain** : Sprague-Dawley  
**Route of admin.** : inhalation  
**Exposure period** : 588 days (19.4 months)  
**Frequency of treatm.** : 6 hours/day, 4.7 days/week or two-thirds of each week  
**Post exposure period** : No  
**Doses** : Average concentration; 10.2 ppm  
**Result** :

<b>Control group</b>	:	yes, concurrent vehicle	
<b>Method</b>	:	other	
<b>Year</b>	:		
<b>GLP</b>	:	no data	
<b>Test substance</b>	:	other TS: Hydrogen chloride, purity; 99.0% grade, Matheson Gas Product	
<b>Remark</b>	:	Method: 20 rats were treated with Hydrogen chloride gas or air shamexposed as control. Complete necropsy was performed on each animals and particular attention was given to the respiratory tract.	
<b>Result</b>	:	No carcinogenic response was observed. Mortality after 588 days (19.4 months) was 29% in treated rats and 28% in airsham-exposed rats; at necropsy no nasal cancers were observed. There was no significant weight loss in treated group.	
<b>Test condition</b>	:	Animals: 20 rats per group Treatment: whole body hydrogen chloride gas or air sham-exposed as control. Duration: 6h*5d/w for lifetime Observation: All animals were observed daily and weighed monthly Complete necropsy was performed on each animal and particular attention was given to the respiratory tract	
<b>Reliability</b>	:	(2) valid with restrictions publication, main features of study reported	
<b>Flag</b>	:	Critical study for SIDS endpoint	
13.02.2006			(75)
<b>Species</b>	:	rat	
<b>Sex</b>	:	male/female	
<b>Strain</b>	:	other: Charles River CD	
<b>Route of admin.</b>	:	oral feed	
<b>Exposure period</b>	:	27 months	
<b>Frequency of treatm.</b>	:		
<b>Post exposure period</b>	:		
<b>Doses</b>	:	0 - 2000 - 8000 - 32000 ppm	
<b>Result</b>	:	negative	
<b>Control group</b>	:	yes	
<b>Method</b>	:		
<b>Year</b>	:		
<b>GLP</b>	:		
<b>Test substance</b>	:	other TS: mono sodium phosphite	
<b>Remark</b>	:	Due to its corrosive properties and high reactivity PCI3 will be bio-available exclusively at the portal of entry. Products of hydrolysis (HCl and phosphorous acid) will form at a high rate and be neutralised immediately. Therefore, at longer term exposure at low doses only the salts (anions) will be systemically available and data on these salts can be used as a surrogate for PCI3 in this context.	
<b>Result</b>	:	There was no evidence of oncogenic response in the urinary bladder, adrenal medulla or at any other site	
		The following symptoms were detected:	
		reduction of body weight: 32000 ppm (m: -13,8%; f:-9,4%) 8000 ppm (m: -15,4%) 2000 ppm (m: - 9,5%) reduced utilization of food: 32000 + 8000 ppm, males only soft stool: 32000 ppm, males slight reduction of urinary pH: 32000 ppm, males	

increase in relative weight:  
liver: males  
kidney: males + females  
heart: males + females  
**Test condition** : Animals: 60 per sex/group  
**Test substance** : mono sodium phosphite  
**Reliability** : (2) valid with restrictions  
**Flag** : Critical study for SIDS endpoint  
28.11.2005 (76) (77)

**Species** : mouse  
**Sex** : male/female  
**Strain** : C3H  
**Route of admin.** : oral unspecified  
**Exposure period** : 11 months  
**Frequency of treatm.** : First 5 months of the experiment injections were made 5 days/week. Because of great mortality among the treated mice, the number of injections was decreased to 3 days/week on alternate days.  
**Post exposure period** : No  
**Doses** : 90-360 mg/kg b.w.  
**Result** :  
**Control group** : yes, concurrent vehicle  
**Method** : other  
**Year** :  
**GLP** : no  
**Test substance** : other TS: Hydrochloric acid

**Remark** : Method: Hydrochloric acid was orally given to mice with (40 mice) or without (58 mice) 1,2,5,6-dibenzanthracene, which was administered once a week during a later period. Probably, only the gastrointestinal tract was examined.  
**Result** : The treatment did not apparently increase the incidence of tumors. The treatment lead to stomach damage. Probably no other tissue examined.  
**Reliability** : (2) valid with restrictions  
short notice, few details

21.09.2004 (78)

#### 5.8.1 TOXICITY TO FERTILITY

#### 5.8.2 DEVELOPMENTAL TOXICITY/TERATOGENICITY

**Species** : rat  
**Sex** : female  
**Strain** : other: no data  
**Route of admin.** : gavage  
**Exposure period** : day 6-15  
**Frequency of treatm.** :  
**Duration of test** :  
**Doses** : 0 - 0 - 6.44 - 9.7 - 19.3 mg/kg bw  
**Control group** : other: positive + vehicle control groups  
**NOAEL teratogen.** : = 19.3 mg/kg bw  
**Method** :  
**Year** :  
**GLP** :  
**Test substance** : other TS: phosphorus trichloride

**Remark** : positive control substance described as: Dikushuang, the chinese name of

some pesticide  
The dose was 7 mg/kg bw

The stated number of resorptions 67 is 47.5% of 141; i.e. The number of resorptions is included in the number of fetuses. It is possible that the heading "Number of fetuses" should have been translated as "number of implantations" and is a translation mistake

**Result** : Body weight gain of dams:  
no effect in treated animals; reduction in pos. controls

Effects on fertility:	neg	low	med	high	pos
Number of pregnant rats:	16	15	15	14	15
number of fetuses:	136	126	132	124	141
Corpora lutea:	218	205	217	206	150
corpora lutea per rat:	13.6	13.7	14.5	14.7	10
number of pups/litter:	8.13	7.73	8.47	8.06	8.2

dead fetuses:	3	1	3	10	0
% dead fetuses:	2.3	0.79	2.27	8.06	0

resorptions:	3	9	2	1	67
% resorptions:	2.2	7.1	1.52	0.81	47.5*

litters with resorptions:	3	3	1	1	11
% Litter w. resorptions:	12.3	20	6.67	7.14	73.3*

\* = significant

	neg	low	med	high	pos
pups examined:	130	116	127	113	74
size (cm):	3,84	3,89	3,84	3,79	3,64
tail (cm):	1,29	1,29	1,25	1,25	1,13
weight of placenta (g):	0,43	0,49	0,46	0,47	0,51
frontal fontanella (mm):	3,14	2,95	3,06	2,97	2,93

There were no significant differences between treated and negative control animals. No malformations were detected.  
Skeletal development in treated fetuses was retarded but without a dose effect relation.  
Pos. control showed clear effects on fertility indices but no malformations.

**Test condition** : Animals were killed on day 20 of pregnancy and fetuses collected by cecarean section.  
the positive control group was exposed on days 9 - 11

**Reliability** : (2) valid with restrictions  
Detailed publication

**Flag** : Critical study for SIDS endpoint  
13.02.2006 (73)

### 5.8.3 TOXICITY TO REPRODUCTION, OTHER STUDIES

**Type** : other:structural aberrations in sperm cells  
**In vitro/in vivo** : In vivo  
**Species** : rat  
**Sex** : male  
**Strain** :  
**Route of admin.** : inhalation  
**Exposure period** : 4 h  
**Frequency of treatm.** : daily

**Duration of test** : 45 or 60 days  
**Doses** : 2 - 33,5 - 97,76 mg/m<sup>3</sup>  
**Control group** :  
**Result** : negative  
**Method** :  
**Year** : 1989  
**GLP** :  
**Test substance** : other TS: phosphorus trichloride

**Remark** : The authors regard the result as "not significant"  
**Result** : 45 days exposure:  
group: 97,25 mg/m<sup>3</sup> 2 mg/m<sup>3</sup> Control  
(97,76)  
findings(%): 1,44\* 0,56 0,4  
hooks (%): 16 (44,4) 5 (35,7) 4 (40,0)  
banana (%): 16 (44,4) 7 (50) 5 (50)  
amorphous (%): 4(11,1) 2 (14,3) 1 (10)

60 days exposure:  
group: 33,5 mg/m<sup>3</sup> 2 mg/m<sup>3</sup> Control  
findings(%): 1,1 1,3 0,86  
hooks (%): 42 (45,2) 26 (46,4) 27 (62,8)  
banana (%): 42 (45,2) 28 (50) 14 (32,6)  
amorphous (%): 8 (8) 2 (3,6) 2 (4,7)  
folding (%): 1 (1,0)

Total structural aberrations were increased significantly only at 97.25 mg/m<sup>3</sup> after 45 days. No change was seen after 60 days.

Comment: The original paper contains both figures (97,25 and 97,76mg/m<sup>3</sup>). It is not clear which of these is correct.

**Test condition** : 48 male rats were randomly allocated to 4 groups:  
neg. control;  
PCl<sub>3</sub> (2 mg/m<sup>3</sup>)  
PCl<sub>3</sub> (33,5 mg/m<sup>3</sup>)  
PCl<sub>3</sub> (97, 25 - 97,76 mg/m<sup>3</sup> \*)

Body weight: 154-180 g

Animals were treated for 45 or 60 days

Sperm cells were collected from epididymides and slides prepared 500 cells per animal, a total of 2500 cells per group, were evaluated

\*) Comment: The original paper contains both figures (97,25 and 97,76mg/m<sup>3</sup>). It is not clear which of these is correct. No information regarding a positive control is given.

**Reliability** : (2) valid with restrictions  
Detailed publication

**Flag** : Critical study for SIDS endpoint

(73)

**Type** : other:structural aberrations in sperm cells  
**In vitro/in vivo** : In vivo  
**Species** : mouse  
**Sex** : male  
**Strain** :  
**Route of admin.** : gavage  
**Exposure period** : 5 days  
**Frequency of treatm.** : daily

**Duration of test** :  
**Doses** :  
**Control group** :  
**Method** :  
**Year** :  
**GLP** :  
**Test substance** : other TS: phosphorus trichloride

**Result** : control | T R E A T E D  
group: water oil pos | 45 mg/kg 90 mg/kg 178 mg/kg  
total: 1,24 1,16 2,64 | 3,56 2,60 3,40  
hooks: 4 3 1 | 1 10 11  
(%): 12,9 10,4 1,5 | 10,1 11 21,6  
banana: 7 5 37 | 23 5 6  
(%): 22,4 17,2 56 | 25,8 5,5 11,8  
amorph: 15 15 25 | 49 60 21  
(%): 48,4 51,7 37,9 | 55,1 65,9 41,2  
folds: 5 6 3 | 8 16 13  
(%): 16,1 20,7 4,6 | 9 17,6 24,5

**Test condition** : There were no significant findings  
60 male mice were randomly allocated to 6 groups:  
neg Control-water;  
neg. control- vegetable oil;  
pos. control-cyclophosphamide (30 mg/kg);  
PCI3 (45 mg/kg)  
PCI3 (90 mg/kg)  
PCI3 (178 mg/kg)

Body weight: 20-25 g

Animals were treated for 5 days and killed after 4 weeks

**Reliability** : Sperm cells were collected from epididymides and slides prepared  
500 cells per animal, a total of 2500 cells per group, were evaluated  
(2) valid with restrictions  
Detailed publication

**Flag** : Critical study for SIDS endpoint

23.09.2004

(73)

## 5.9 SPECIFIC INVESTIGATIONS

## 5.10 EXPOSURE EXPERIENCE

**Type of experience** : Human - Medical Data

**Remark** : workers exposed to phosphorus-chloride compounds: increased incidence  
of asthmatics

**Test substance** : phosphorus trichloride

**Reliability** : (4) not assignable  
PCI3 mentioned without detailed evidence

14.09.2005

(79)

**Type of experience** : Human

<b>Remark</b>	:	threshold of the irritant effects similar in man and animals. Lim ir. 0.004 or 0.005 mg/l in man or animal.	
<b>Test substance</b>	:	phosphorus trichloride	
<b>Reliability</b>	:	(2) valid with restrictions limited documentation, non standard evaluation scheme	
<b>Flag</b> 12.05.2004	:	Critical study for SIDS endpoint	(51)
<b>Type of experience</b>	:	Human	
<b>Remark</b>	:	workers employed in the production of phosphorus trichloride: concentration was 10-20 mg/m <sup>3</sup> under normal condition and 80-150 mg/m <sup>3</sup> at times when the plant was out of order. In acute poisoning after 2-6 hours burning sensation in eyes and throat, photophobia, feeling of chestoppression, dry cough, irritation of mucous membranes. In subacute poisoning the symptoms occur after 1-8 weeks with signs of irritation and asthmatic bronchitis.	
<b>Test substance</b>	:	phosphorus trichloride	
<b>Reliability</b>	:	(2) valid with restrictions limited documentation; case report	
<b>Flag</b> 12.05.2004	:	Critical study for SIDS endpoint	(80)
<b>Type of experience</b>	:	Human	
<b>Remark</b>	:	patients accidentally exposed to PCI <sub>3</sub> and its degradation products after a railroad accident: symptoms: burning eyes (85%; mild conjunctivitis in 36%), shortness of breath (59%), throat irritation (59%) and lacrimation (59%), headache and nausea (48%), burning skin (44%), sputum production (41%), generalized or pleuric chest pain (33%) and rash/itch (33%), wheezing (26%), blurred vision (22%), vomiting (15%) and abdominal pain (15%). Lactic dehydrogenase was increased in 22% of patients and recovered within four weeks after exposure. Pulmonary function tests showed a decrease of vital capacity in patients within 1/8 mile of the event. Hypoxemia was recorded in these patients after two months.	
<b>Test substance</b>	:	phosphorus trichloride	
<b>Reliability</b>	:	(2) valid with restrictions limited documentation; case report	
<b>Flag</b> 19.02.2004	:	Critical study for SIDS endpoint	(81) (82)
<b>Type of experience</b>	:	Human - Medical Data	
<b>Remark</b>	:	accident: conjunctivitis, pain in nose and throat, difficulty in swallowing, sore throat, feeling of chest oppression.	
<b>Test substance</b>	:	no data	
<b>Reliability</b>	:	(4) not assignable Report not available	
13.05.2004			(83)
<b>Type of experience</b>	:	Human	
<b>Result</b>	:	PCI <sub>3</sub> after inhalation (1 single breath) caused painful irritaion of eyes, nose and throat, strained respiration, difficulty to swallow lasting pain in the throath and severe catarrh. Recovery was achieved wihtin 8 days.	

- Test substance** : no data  
**Reliability** : (4) not assignable  
report not available  
19.02.2004 (84)
- Type of experience** : Human - Medical Data
- Result** : PCI3 caused irritation of eyes and mucous membranes in workers. Symptoms appeared immediately or delayed for up to 1 day. One patient died after several days due to an asthmatic fit caused by the exposure to PCI3  
in concentrations less than 1 mg/l air rhinitis, conjunctivitis, pain in nose and throat: in higher concentrations dyspnea and death.
- Test substance** : phosphorus trichloride  
**Reliability** : (2) valid with restrictions  
Case report; limited documentation  
**Flag** : Critical study for SIDS endpoint  
19.02.2004 (85)
- Type of experience** : Human - Medical Data
- Result** : PCI3 is quickly hydrolysed in water. It causes immediate irritation to mucous membranes. In patients PCI3 therefore causes immediate defence and effects are mainly found in the upper respiratory tract
- Test substance** : phosphorus trichloride  
**Reliability** : (4) not assignable  
secondary literature  
19.02.2004 (86)
- Type of experience** : Human
- Result** : Symptoms reported by exposed workers were: respiratory tract and eye irritation, cough, asthma, loss of voice
- Test substance** : phosphorus trichloride  
**Reliability** : (2) valid with restrictions  
limited documentation; few details given  
**Flag** : Critical study for SIDS endpoint  
26.02.2004 (87)
- Type of experience** : Human
- Result** : PCI3 is mentioned as a cause of occupational asthma: 48/170 (28,2%) exposed to phosphorus chlorides (PCI3, PCI5, POCl3) are reported to have asthma
- Test substance** : phosphorus trichloride  
**Reliability** : (4) not assignable  
Short remark within a review  
**Flag** : Critical study for SIDS endpoint  
23.03.2004 (88)
- Type of experience** : Human - Epidemiology
- Result** : 37 exposed and 22 unexposed workers were investigated regarding their pulmonary function. Exposed employees had a significantly higher prevalence (65%) of occasional respiratory discomfort than unexposed employees (5%). No consistent association between pulmonary function parameters and exposure could be documented.

<b>Test substance</b>	:	phosphorus trichloride	
<b>Reliability</b>	:	(2) valid with restrictions	
19.05.2004			(89) (90)
<b>Type of experience</b>	:	Human - Epidemiology	
<b>Result</b>	:	Follow up of NTIS PB 81-170920 26 exposed and 11 non-exposed 13 of the xposed worker reported intermittent respiratory distress (wheezing, breathlessness, chest tightness). Although a significantly larger group of exposed workers exhibited symptoms, a decrement of pulmonary function due to exposure could not be demonstrated.	
<b>Test substance</b>	:	phosphorus trichloride	(91) (92)
14.09.2005			
<b>Result</b>	:	IgE specific for phosphorus trichloride was determined in an occupational surveillance program for 5 years No specific IgE against phosphorus trichloride was seen, neither in the surveillance program nor in any case of product contact	
<b>Test substance</b>	:	phosphorus trichloride	(93)
14.09.2005			

#### 5.11 ADDITIONAL REMARKS

<b>Type</b>	:	Metabolism	
<b>Result</b>	:	Hydrolysis of PCI <sub>3</sub> is mostly complete within 4-6 seconds in excess water. Products are hydrochloric acid (HCl) and phosphorous acid (H <sub>3</sub> PO <sub>3</sub> ) The rate of pH-change is comparable to the addition of concentrated hydrochloric acid	
<b>Test condition</b>	:	The change of pH of water was recorded during addition of PCI <sub>3</sub> at room temperature	
<b>Test substance</b>	:	phosphorus trichloride	
<b>Reliability</b>	:	(1) valid without restriction Full report available	
<b>Flag</b>	:	Critical study for SIDS endpoint	(24)
16.06.2004			
<b>Type</b>	:	Neurotoxicity	
<b>Result</b>	:	Serum BChE but not brain AChE was inhibited in vivo 1 h after exposure ED <sub>50</sub> : 14 mg/kg Mortality: >100 mg/kg Symptoms: tremors and muscle fasciculation	
<b>Test condition</b>	:	Animals: male Swiss Webster mice Dose: 0-10-30-100 mg/kg ip in corn oil	
		Tissue samples were removed for AChE determination 1 or 24 hours after treatment or at death. Blood, skeletal muscle, diaphragm, and brain were examined.	
		The in vivo experiments were supplemented by in vitro studies using different sources of AChE	
<b>Test substance</b>	:	phosphorus trichloride	
<b>Reliability</b>	:	(4) not assignable irrelevant route of exposure (i.p.); only few experimental details;	(94)
13.02.2004			

<b>Type</b>	:	Neurotoxicity	
<b>Remark</b>	:	Species not relevant for mammalian toxicity determination; insufficient documentation	
<b>Result</b>	:	ED 50 : 6-20 mg/L IC 50 : 6 mg/L (brain) Mortality was associated with > 90% inhibition	
<b>Test condition</b>	:	adult house flies were exposed to vapours of POCl <sub>3</sub> in a 120 ml glass chamber. The inability to walk or fly was recorded and at 15 minutes mortality was determined. Animals were frozen on dry ice, heads removed and AChE activity assayed.	
<b>Test substance</b>	:	phosphorus trichloride	
<b>Reliability</b>	:	(4) not assignable	
13.02.2004			(94)
<b>Type</b>	:	other: Acute toxicity of phosphous acid	
<b>Result</b>	:	LD50 : Male rats: 2844,4 (2261,7 - 3427,1) mg/kg Female rats: 1895,3 (1521,3 - 2269,3) mg/kg Mice: 2172,3 (1548,5 - 2796,1) mg/kg	
		Death occurred within few days after treatment (mean time to death 14,3 h for rats)	
<b>Test substance</b>	:	phosphorous acid; degradation product of PCI <sub>3</sub>	
<b>Reliability</b>	:	(2) valid with restrictions limited documentation	
<b>Flag</b>	:	Critical study for SIDS endpoint	
01.08.2005			(95)
<b>Type</b>	:	other: Corrosion	
<b>Remark</b>	:	corrosive changes in animals exposed to PCI <sub>3</sub> vapours.	
<b>Test condition</b>	:	1	
<b>Test substance</b>	:	phosphorus trichloride	
<b>Reliability</b>	:	(2) valid with restrictions no experimental details given	
19.02.2004			(56)
<b>Type</b>	:	other: Exposure limit	
<b>Result</b>	:	Recommended Exposure Limit (REL, NIOSH) = 0.2 ppm Permissible Exposure Limit (PEL, OSHA) = 0.5 ppm Immediately dangerous for Life or Health (IDLH) = 25 ppm	
<b>Test substance</b>	:	phosphorus trichloride	
<b>Reliability</b>	:	(4) not assignable secondary literature	
19.02.2004			(96)
<b>Type</b>	:	other: MAK value	
<b>Result</b>	:	MAK = 0.5 ppm = 3 mg/m <sup>3</sup> (MAK= maximum concentration in air at the workplace (8h/d; 5d/w))	
<b>Test substance</b>	:	phosphorus trichloride	
<b>Reliability</b>	:	(4) not assignable secondary literature	
19.02.2004			(97)
<b>Type</b>	:	other: Maximum allowable concentration in the workplace air	

**Result** : The maximum allowable concentration are mentioned as:  
2 ppp (12 mg/m<sup>3</sup>) for 1 minute  
1 ppm for "short exposure"  
0.5 ppm for "satisfactory conditions"

**Test substance** : phosphorus trichloride

**Reliability** : (4) not assignable  
secondary literature

19.02.2004 (98)

**Type** : other: Repeated dose toxicity of phosphorous acid

**Remark** : limited documentation

**Result** : In the subacute study all doses created toxic effects. The largest changes occurred in the levels of choline esterase in blood and liver as well as the activity of monoamine oxidase in liver.  
Statistically significant reductions were seen in liver (day 20: high dose; day 30: mid+high dose), kidney (day 10: high dose; day 20 mid+high dose; day 30: all doses) and blood (day 20+30: all doses). Reductions reached a level of up to 48% in the high dose group after 30 days.  
Monoamine oxidase showed an increase on days 10 and 20 (up to 75%) and a decrease (up to 25%) on day 30 in mid and high dose groups.  
In the mid and high doses a reduction of alkaline phosphatase in blood and of sulfhydryl groups in liver and kidneys were recorded. Motoractivity and body weight were also reduced. Increases were observed in the activity of ALAT in blood.  
Microscopic examination of the liver revealed dilation of sinuses, infiltration of connective tissue and necrotic foci. In kidneys destruction of tubular and glomerular cell occurred, accompanied by lymphocytic infiltration.  
The LOEL was 2,45 mg/kg based on the effect on choline esterase in kidney tissue.

The chronic treatment of animals with 1/250 LD50 (11.4 mg/kg) caused increases in motroactivity and choline esterase activity.  
At the histopathologic examination after 90 and 180 days of treatment livers showed dilation and lymphocytic infiltration, focal destruction of liver trabecula.

A dose of 1/2000 LD50 (1.4 mg/kg) was without any effect.

In both studies no functional effects on spermatogenesis were detected. Only a dose of 1/10 LD50 caused necrosis of testicular epithelium.

**Test condition** : Animals: male rats

Dose ranges:  
subacute test: 0.1 - 0.02 - 0.004\*LD50 ( 284-11.4 mg/kg)  
chronic test: 0.004 - 0.0005\*LD50 (11.4-1.4 mg/kg)

Animals were examined after 10, 20 and 30 days in the subacute test and after 60, 90, 120, 150, and 180 days in the chronic test

Parameters:  
Body weight; haematology; choline esterase activity in brain, liver, kidneys and blood; monoamineoxidase in liver and brain; clinical chemistry in blood and tissue;  
open field test; behavior;  
histopathology of liver, kidney and testes.

**Test substance** : phosphoric acid

**Reliability** : (2) valid with restrictions

23.03.2004 (95)

<b>Type</b>	: other: Review	
<b>Test substance</b>	: phosphorus trichloride	
<b>Reliability</b>	: (4) not assignable secondary literature	
19.05.2004		(99)
<b>Type</b>	: other: Review	
<b>Test substance</b>	: phosphorus trichloride	
<b>Reliability</b>	: (4) not assignable secondary literature	
01.08.2005		(100)
<b>Type</b>	: other: Review	
<b>Test substance</b>	: phosphorus trichloride	
<b>Reliability</b>	: (4) not assignable secondary literature	
14.09.2005		(101)
<b>Result</b>	: In subacute and chronic studies no functional effects on spermatogenesis were detected. Only a dose of 1/10 LD50 caused necrosis of testicular epithelium.	
<b>Test condition</b>	: Embryotoxicity test: The number as well as the weight of pups was reduced. No effects on fertility index, pre- and post-implantation loss. No malformations. Dose ranges: (male rats) subacute test: 0.1 - 0.02 - 0.004 LD50 (284-11.4 mg/kg) chronic test: 0.004 - 0.0005 LD50 (11.4-1.4 mg/kg )  Animals were examined after 10, 20 and 30 days in the subacute test and after 60, 90, 120, 150, and 180 days in the chronic test  Embryotoxicity Test: (female rats) Dose: 284 mg/kg Parameters: Fertility index; maternal, pre- and post implantation mortality; fetal survival;	
<b>Test substance</b>	: phosphorus trichloride	
<b>Reliability</b>	: (2) valid with restrictions limited documentation	
14.09.2005		(102)
<b>Result</b>	: negative	
<b>Test condition</b>	: Guinea pigs were treated by subcutaneous implantation at doses of 28 and 57 mg/kg	
<b>Test substance</b>	: phosphorus trichloride	
14.09.2005		(102)
<b>Result</b>	: Comprehensive review on properties, uses, exposure and effects of PCI3	
<b>Test substance</b>	: phosphorus trichloride	
14.09.2005		(103)
<b>Result</b>	: TLV-TWA = 0,2 ppm (1,1 mg/m <sup>3</sup> )	

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**Test substance** : TLV-STEL = 0,5 ppm (2,8 mg/m<sup>3</sup>)  
14.09.2005 : phosphorus trichloride (12)

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