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## An Efficient Approach to Sulfate Metabolites of Polychlorinated Biphenyls

Xueshu Li<sup>a</sup>, Sean Parkin<sup>b</sup>, Michael W. Duffel<sup>c</sup>, Larry W. Robertson<sup>a</sup>, and Hans-Joachim Lehmler<sup>a,\*</sup>

<sup>a</sup> Department of Occupational and Environmental Health, College of Public Health, University of Iowa, Iowa City, IA, 52242, USA

<sup>b</sup> Department of Chemistry, College of Arts and Sciences, University of Kentucky, Lexington, KY, 40506, USA

<sup>c</sup> Division of Medicinal and Natural Products Chemistry, College of Pharmacy, University of Iowa, Iowa City, IA, 52242, USA

### Abstract

Polychlorinated biphenyls (PCBs), a major class of persistent organic pollutants, are metabolized to hydroxylated PCBs. Several hydroxylated PCBs are substrates of cytosolic phase II enzymes, such as phenol and hydroxysteroid (alcohol) sulfotransferases; however, the corresponding sulfation products have not been isolated and characterized. Here we describe a straightforward synthesis of a series of ten PCB sulfate monoesters from the corresponding hydroxylated PCBs. The hydroxylated PCBs were synthesized by coupling chlorinated benzene boronic acids with appropriate brominated (chloro-)anisoles, followed by demethylation with boron tribromide. The hydroxylated PCBs were sulfated with 2,2,2-trichloroethyl chlorosulfate using DMAP as base. Deprotection with zinc powder/ammonium formate yielded the ammonium salts of the desired PCB sulfate monoesters in good yields when the sulfated phenyl ring contained no or one chlorine substituent. However, no PCB sulfate monoesters were isolated when two chlorines were present *ortho* to the sulfated hydroxyl group. To aid with future quantitative structure activity relationship studies, the structures of two 2,2,2-trichloroethyl-protected PCB sulfates were verified by X-ray diffraction.

### Keywords

Biaryls; Polychlorinated biphenyls (PCBs); Sulfates; 2,2,2-Trichloroethyl (TCE) group; Metabolites

### 1. Introduction

Polychlorinated biphenyls (PCBs) were manufactured commercially in large quantities and used in numerous technical applications, for example as lubricants, cooling fluids, flame retardants, adhesives, and plasticizers (Hansen 1999; Robertson and Hansen 2001). PCBs are still in use as dielectric fluids in capacitors and transformers. Their wide-spread industrial use

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Corresponding Author: Hans-Joachim Lehmler, Department of Occupational and Environmental Health, College of Public Health, University of Iowa 100 Oakdale Campus #221 IREH Iowa City, IA 52242-5000, Phone +1 (319) 335-4211, Fax: +1 (319) 335-4290, e-mail: hans-joachim-lehmler@uiowa.edu.

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and physicochemical properties, such as lipophilicity, semi-volatility and stability towards biological, chemical and thermal degradation, have resulted in widespread environmental contamination. PCBs have also been associated with a broad range of adverse human health effects, such as (neuro-)developmental toxicity (Kodavanti 2004) and carcinogenicity (Silberhorn et al. 1990). The production of PCBs was banned in the United States in the late 1970s because of these environmental and public health concerns.

Especially lower chlorinated PCBs undergo oxidative metabolism to hydroxylated PCBs catalyzed by cytochrome P-450 enzymes (Letcher et al. 2000). Some hydroxylated PCB metabolites persist in blood, liver and other tissues of humans, where they can reach levels that are comparable to PCB blood levels (Bergman et al. 1994; Hovander et al. 2006; Park et al. 2007). They potently inhibit the activity of phenol sulfotransferases (SULT) and, thus, may interfere with the sulfation of endogenous and exogenous compounds (Kester et al. 2000; Schuur et al. 1998a; Schuur et al. 1998b; Schuur et al. 1998c; van den Hurk et al. 2002; Wang et al. 2005; Wang et al. 2006). Similarly, hydroxylated PCBs are inhibitors of hydroxysteroid (alcohol) sulfotransferases, such as human SULT2A1 (Liu et al. 2006). There is also evidence that some hydroxylated PCBs are substrates for SULTs. Sacco and James demonstrated that several hydroxylated PCBs are sulfated by polar bear liver cytosol (Sacco and James 2005). Liu et al. recently reported that two hydroxylated PCBs, 4-hydroxy-2',3,5-trichlorobiphenyl and 4'-hydroxy-2,3',4,5'-tetrachloro-biphenyl, are substrates for SULT2A1 (Liu et al. 2006).

Very little is currently known about the biological properties and metabolic disposition of PCB sulfates. While many phenyl sulfates are water soluble and readily excreted, calculated octanol/water partition coefficients indicate that PCB sulfates may retain significant lipophilic properties (James 2001). The first chemical synthesis of a series of lower chlorinated PCB sulfate monoesters described herein provides a source of sufficient quantities of these metabolites for detailed study of their chemical and biochemical properties, and how these properties may relate to the metabolic disposition and detoxication of PCBs and hydroxylated PCBs.

## 2. Materials and methods

### 2.1. Chemicals and Instruments

All of the chlorinated benzene boronic acids, the brominated phenols and tetrakis (triphenylphosphine)palladium(0) were obtained from Fisher Scientific (Fairlawn, New Jersey, USA). 4-Hydroxy biphenyl (**7a**) was purchased from Sigma-Aldrich chemical company (St. Louis, MO, USA). All hydroxylated PCBs **7** were synthesized as described previously (Lehmler and Robertson 2001; McLean et al. 1996). Chlorosulfuric acid 2,2,2-trichloroethyl ester was synthesized according to the method by Hedayatullah and co-workers (Hedayatu et al. 1971). The  $^1\text{H}$  and  $^{13}\text{C}$  NMR spectra were recorded on a multinuclear Bruker DRX 400 Digital NMR Bruker spectrometer at ambient temperature. All  $^1\text{H}$  and  $^{13}\text{C}$  chemical shifts are reported in parts per million (ppm) relative to internal tetramethylsilane ( $\text{Me}_4\text{Si}$ ). Melting points were determined using a MelTemp apparatus and are uncorrected. The gas chromatography-mass spectra (GC-MS) were recorded using a Thermo Voyager EI instrument. High-resolution mass spectra (HR-MS) were measured using an Autospec ESI-MS instrument at the University of Iowa Mass Spectrometry Facility. Infrared spectra (IR) were recorded on a NEXUS 670 FT-TR instrument. UV/Vis spectra were measured using a Perkin Elmer Lambda 650 UV/Vis spectrometer at 23 °C (UV/Vis spectral data of the corresponding hydroxylated PCBs **7** are shown in parentheses for comparison). The characterization of selected compounds is provided below. The characterization of all other compounds is provided in the supplementary material.

## 2.2. General procedure for the synthesis of sulfuric acid 2,2,2-trichloroethyl (TCE) esters of hydroxylated phenols and PCBs

A solution of 2,2,2-trichloroethyl chlorosulfate (3.2 mmol) in anhydrous DCM (5 mL) was added slowly at 0°C to a solution of phenols **1a,b** or hydroxylated PCB **7a-l** (Lehmler and Robertson 2001; McLean et al. 1996) (3 mmol) and 1.5 equivalents of 4-N,N'-dimethylaminopyridine (DMAP, 4.5 mmol) in anhydrous DCM (15 mL) (Liu et al. 2004). The reaction mixture was stirred for 30 minutes at 0°C, allowed to warm to ambient temperature and stirred for an additional 10 hours. The solvent was removed under reduced pressure and the residue was dissolved in ethyl acetate (20 mL). The ethyl acetate solution was washed with distilled water (20 mL), 1M HCl solution (2 × 20 mL) and distilled water (20 mL). The organic phase was dried over anhydrous Na<sub>2</sub>SO<sub>4</sub>. The solvent was removed under reduced pressure, and the residue was purified by column chromatography on silica gel using a mixture of *n*-hexanes and chloroform (8 : 1 to 5 : 1, v/v) as eluent. The TCE esters **2a-b** and **8a-l** were obtained in good yields ranging from 75% to 94%.

**2.2.1. Sulfuric acid 4'-chloro-biphenyl-4-yl 2,2,2-trichloroethyl ester (8b)**—White solid; mp: 108–109 °C; <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>): δ/ppm 4.86 (s, 2H, CH<sub>2</sub>), 7.42 (AA'XX' system, 2H, *J* ~ 8.6 Hz), 7.43 (AA'XX' system, 2H, *J* ~ 8.6 Hz), 7.47 (AA'XX' system, 2H, *J* ~ 8.6 Hz), 7.59 (AA'XX' system, 2H, *J* ~ 8.6 Hz). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>): δ/ppm 80.4 (CH<sub>2</sub>), 92.3 (CCl<sub>3</sub>), 121.5 (2×CH), 128.4 (2×CH), 128.6 (2×CH), 129.1 (2×CH), 134.1, 137.9, 139.9, 149.6 (C<sub>Ar</sub>-OSO<sub>3</sub>). IR (film): 3027, 2975, 1480, 1409, 1389, 1215, 1153, 1095, 989, 821 cm<sup>-1</sup>. EI-MS *m/z* (relative intensity, %): 414 (35, C<sub>14</sub>H<sub>10</sub>Cl<sub>4</sub>O<sub>4</sub>S<sup>++</sup>), 284 (25), 217 (10), 203 (100), 175 (46), 149 (15), 139 (35).

## 2.3. General procedure for the synthesis of ammonium salts of chlorinated biphenyl sulfates

Ammonium formate (0.77 g, 12 mmol) was added to a solution of the (chlorinated) biphenyl TCE sulfate **2** or **8** (2 mmol) in methanol (5 mL) (Liu et al. 2004). Zinc dust (0.26 g, 4 mmol) was added after the ammonium formate had dissolved completely, and the reaction mixture was stirred until the TCE ester **2** or **8** was consumed completely as determined by TLC (usually within 30 minutes). The solution was filtered through Celite and concentrated under reduced pressure at temperatures below 35 °C. The product was purified by column chromatography on silica gel using a mixture of chloroform, methanol and ammonium hydroxide (8 : 1 : 0.2, v/v) as eluent. The solvent was removed under reduced pressure at temperature below 35 °C to yield the final products as a white solid with yields ranging from 83% to 97%. The R<sub>f</sub> values of all PCB sulfates were approximately R<sub>f</sub> = 0.3 (CHCl<sub>3</sub> : CH<sub>3</sub>OH : NH<sub>4</sub>OH = 10 : 2 : 0.5, v/v).

**2.3.1. Sulfuric acid mono-(4'-chloro-biphenyl-4-yl) ester, ammonium salt (9b)**—White solid; mp: 250 °C (dec.); <sup>1</sup>H NMR (400 MHz, CD<sub>3</sub>OD): δ/ppm 7.38 (AA'XX' system, 2H, *J* ~ 9.0 Hz), 7.40 (AA'XX' system, 2H, *J* ~ 8.9 Hz), 7.56 (2 overlapping AA'XX' systems, 4H, *J* ~ 8.8 Hz). <sup>13</sup>C NMR (100 MHz, CD<sub>3</sub>OD): δ/ppm 122.9 (2×CH), 128.7 (2×CH), 129.4 (2×CH), 129.9 (2×CH), 134.3, 137.7, 140.6, 153.9 (C<sub>Ar</sub>-OSO<sub>3</sub>). IR (KBr): 3235, 3079, 1246, 1061 cm<sup>-1</sup>. UV/Vis: λ<sub>9b,max</sub>(MeOH) = 258 nm, ε<sub>9b</sub> = 2.42 × 10<sup>4</sup> L·mol<sup>-1</sup>·cm<sup>-1</sup> (λ<sub>7b,max</sub>(MeOH) = 267 nm, ε<sub>7a</sub> = 2.29 × 10<sup>4</sup> L·mol<sup>-1</sup>·cm<sup>-1</sup>). HRMS (ESI, negative): [M-NH<sub>4</sub>]<sup>-</sup> found *m/z* 282.9844, calculated for C<sub>12</sub>H<sub>8</sub>(35)ClO<sub>4</sub>S 282.9832.

## 2.4. Single crystal structure determination of **8e** and **8k**

Crystals of the TCE-protected PCB sulfates **8e** and **8k** suitable for crystal structure analysis were obtained by slow crystallization from methanol. X-ray diffraction data were collected at 90.0(2) K on a Nonius KappaCCD diffractometer. Raw data were integrated, scaled, merged and corrected for Lorentz-polarization effects using the Denzo-SMN package (Otwinowski

and Minor 1997). The structures were solved by direct methods (Sheldrick 2008) and missing atoms were located in difference Fourier maps (Sheldrick 2008). Refinement was carried out against  $F^2$  by weighted full-matrix least-squares. Hydrogen atoms were found in difference maps but subsequently placed at calculated positions and refined using appropriate riding models. Non-hydrogen atoms were refined with anisotropic displacement parameters. Atomic scattering factors were those of SHELXL (Sheldrick 2008), as taken from the International Tables for Crystallography vol. C (Wilson 1992). The crystal data and the related parameters are summarized in Table 2. Additional crystallographic data have been deposited with the Cambridge Crystallographic Data Center as Supplementary Publications CCDC 687167 (**8e**) and CCDC 719235 (**8k**). Copies of the data can be obtained free of charge on application to the CCDC, 12 Union Road, Cambridge CB2 1EZ, U.K. (fax, (+44)1223-336- 033; e-mail, deposit@ccdc.cam.ac.uk).

### 3. Results and discussion

#### 3.1. Synthesis

A variety of sulfation reagents and conditions have been reported in the literature. For examples, sulfur trioxide complex with pyridine or tertiary amines are commonly used for the sulfation of different alcohols, including saccharide derivatives (Nishino and Nagumo 1992; Petitou and van Boeckel 2004; Pires et al. 2001) and organic compounds with phenolic moieties, such as phenols (Hanson et al. 2006; Hearse et al. 1969; Ragan 1978), flavanoids (Gunnarsson and Desai 2002), flavonoids (Gunnarsson and Desai 2003) and steroids (Santos et al. 2003). However, these sulfation methods have drawbacks, such as tedious purification procedures and low yields. In recent years, novel sulfation methods have been developed to overcome these problems. In particular substituted alkyl chlorosulfates, including isobutyl, neopentyl and 2,2,2-trichloroethyl chlorosulfate (TCE-Cl), have been used to obtain sulfate diesters which form the desired sulfate monoesters upon deprotection in good-to-excellent yields (Liu et al. 2004; Simpson and Widlanski 2006).

In order to obtain PCB sulfates to investigate their biological role in the metabolism and toxicity of PCBs, we employed the 2,2,2-trichloroethyl (TCE)-protection method to synthesize a series of sulfate metabolites of several lower chlorinated PCBs. As shown in Scheme 1, we initially synthesized the brominated TCE sulfate diester **2b** and investigated its coupling with chlorinated benzene boronic acids **4** to obtain the desired PCB TCE sulfate diesters. However, the Suzuki coupling of **2b** with chlorinated benzene boronic acids **4** failed because the TCE group is unstable under the reaction conditions employed.

In an alternate approach, we first prepared the hydroxylated biphenyl derivatives **7** and introduced the sulfate group in the final steps of the synthesis. As shown in Scheme 2, a series of hydroxylated PCB derivatives (**7a-l**) were synthesized using the Suzuki-coupling of chlorinated benzene boronic acids **4** and appropriate brominated (chloro)-anisoles **5**, followed by demethylation with boron tribromide (Lehmler and Robertson 2001; McLean et al. 1996). Subsequently, TCE-protected PCB sulfates diesters **8a-j** were synthesized from the hydroxylated PCBs **7a-l** by sulfation with 2,2,2-trichloroethyl chlorosulfate as sulfation reagent and DMAP as base. The sulfation reactions proceeded in good-to-excellent yields ranging from 75 to 94% (Table 1).

Zinc powder-ammonium formate is an efficient and mild deblocking system and can be employed with halogenated aromatic compounds without dehalogenation (Liu et al. 2004). Therefore, the TCE group was removed in the last step of the synthesis by reductive elimination using this system, with yields ranging from 83% to 97% for PCB TCE esters **8a-j** (Table 1). Although the deprotection of the PCB TCE sulfate diesters **9k** and **9l** yield PCB sulfate monoesters according to TLC analysis, we were unable to isolate the desired product after

column chromatography on silica gel with chloroform-methanol-ammonium hydroxide as eluent. Instead, the product rapidly degraded in solution to the hydroxylated starting materials **7k** and **7l**. Similarly, the chlorinated phenol TCE diester **2b** did not yield the desired sulfate monoester, which suggests that the sulfate monoesters of phenolic compounds with two chlorine atoms in *ortho* position to the sulfated phenol are unstable under the conditions required for the isolation of solid products.

One possible explanation for this observation is the increasing degree of chlorination in the hydroxylated phenyl ring, which increases the acidity ( $pK_a$  value: di-*ortho* < mono-*ortho* < non-*ortho* chloro (Tampal et al. 2002)). As a result, the hydroxylated PCBs **7k** and **7l** are excellent leaving groups, thus resulting in the instability of the corresponding PCB sulfate monoesters. Similarly, the stability of phenolic sulfate monoesters has been shown to correlate with the  $pK_a$  of the phenol and the length of the C<sub>Ar</sub>-O and O-S bond length of the sulfate monoester (Brandao et al. 2005). This interpretation is also supported by the C-O and S-O bond length observed in the molecular structures of TEC sulfate diesters **8e** and **8k** (see below).

The structures of the sulfate esters were confirmed by NMR, IR, and UV/Vis. In the IR spectra, we observed typical absorption bands at 1200–1220 cm<sup>-1</sup> and 1420–1460 cm<sup>-1</sup> (S=O of TCE-protected sulfate diesters **8**), 1000–1010 cm<sup>-1</sup> (O-S-O of TCE-protected sulfate diesters **8**), 1230–1250 cm<sup>-1</sup> and 1440–1490 cm<sup>-1</sup> (S=O of sulfate monoester ammonium salts **9**), and 1060–1070 cm<sup>-1</sup> (O-S-O of sulfate monoester ammonium salts **9**) (Ragan 1978). In the UV/Vis spectra, the  $\lambda_{max}$  values of the hydroxylated PCBs **7** were always greater than the  $\lambda_{max}$  values of the corresponding sulfates **9**, with the difference ranging from 2 to 10 nm. These differences in  $\lambda_{max}$  may be useful for the detection of PCB sulfate monoesters **9** with UV/Vis detectors.

### 3.2. Solid state molecular structure of PCB TCE sulfate diesters

The availability of structural information of PCB sulfate monoesters would be valuable for quantitative structure activity relationship (QSAR) studies of their interaction with SULTs. Unfortunately, we were unable to obtain crystals of the PCB sulfate monoesters suitable for X-ray crystal structure determination due to their instability under the conditions employed for crystallization. Instead, we obtained single crystals of two PCB TCE sulfate diesters. The molecular structure and the labeling scheme of selected PCB sulfate diesters (**8e** and **8k**) is shown in Figure 1. Relevant X-ray crystallographic data and selected bond lengths, angles and dihedral angles are reported in Tables 2 and 3, respectively. To the best of our knowledge, no crystal structures of similar mixed alkyl aryl esters of sulfuric acid have been reported previously.

The length of the C4-O1 and S1-O1 bonds of diesters **8e** and **8k** differed significantly (Table 3). While the C4-O1 bond of **8k** was significantly shorter compared to **8e** (1.405 Å versus 1.426 Å), the S1-O1 bond of **8k** was longer compared to **8e** (1.600 Å versus 1.585 Å). These differences in the bond lengths of **8e** and **8k** are due to the two electronegative chlorine substituents *ortho* to the sulfate group of **8k**. This results in a more positive partial charge on the C-4 carbon atom and, ultimately, a shorter C4-O1 bond length. At the same time, the reduced electron density on the O1 atom contributes to a longer and weaker S1-O1 bond. The increasing weakness of the S1-O1 bond is due to an increasing number of electronegative substituents (i.e., chlorines) in the phenyl ring system, and explains, at least in part, why we were unable to isolate the PCB sulfate monoesters corresponding to the di-*ortho* substituted TCE sulfate diesters **8k** and **8l**. Similarly, the C-O bond length of aromatic sulfuric acid monoesters (and ultimately their stability) correlated with the C-O bond length and, ultimately, the  $pK_a$  value of the corresponding phenol (Brandao et al. 2005).



The O1 atom of the two PCB TCE sulfate diesters was in the plane of the phenyl ring, with deviations from the ring plane of 0.092 Å and 0.128 Å for **8e** and **8k**, respectively. The bond lengths of the S1-O1 and S1-O2 ester bonds were similar for both compounds and ranged from 1.564 Å to 1.600 Å. In contrast, the S-O bond length in aromatic sulfate monoesters ranged from 1.611 to 1.653 Å (Brandao et al. 2005), which is slightly longer than the bond length observed for **8e** and **8k**. The S1-O3 and S1-O4 bond lengths were shorter in the diesters (1.409 Å to 1.416 Å) compared to monoesters (1.427 to 1.445 Å) (Brandao et al. 2005) due to the double bond character of these bonds.

The dihedral angle between the phenyl rings of a PCB congener determine its three dimensional structure and, thus, its affinity to cellular targets, such as nuclear transcription factors (Lehmler et al. 2002; Vyas et al. 2006a). The solid state dihedral angles between the two phenyl rings of PCB TCE sulfate diesters were smaller compared to the structurally-related PCB metabolites. For example, the solid state dihedral angle between the two phenyl rings of **8e** (52.13°) was smaller compared to the corresponding methoxylated PCB (59.92°) (Vyas et al. 2006b). These deviations from the energetically most favorable conformations of **8e** and **8k** are likely due to crystal packing effects, which allow the molecule to adopt an energetically unfavorable conformation (i.e., dihedral angle) to maximize intermolecular interactions and, thus, the lattice energy in the crystal.

## 4. Conclusions

Hydroxylated PCBs are emerging as an important, but frequently overlooked, aspect of PCB toxicity. Little is known about the disposition of this group of PCB metabolites and their toxicity, both in rodent animal models and humans. In recent years, some hydroxylated PCBs **7** have been documented to inhibit cytosolic SULTs, whereas other hydroxylated PCBs **7** appear to be substrates for SULTs. Here, we report the first chemical synthesis of a series of PCB sulfate monoesters **8** in a four step synthesis from chlorinated benzene boronic acids **4** and brominated (chloro-)anisoles **5**. Suzuki coupling of boronic acids **4** with brominated anisoles **5**, followed by demethylation with BBr<sub>3</sub> yielded the desired hydroxylated PCBs **7**. Subsequently, sulfation with 2,2,2-trichloroethyl chlorosulfate and deprotection gave the desired PCB sulfate monoesters **9a-j** in good-to-excellent yields. The ammonium sulfate monoesters **9** are unstable over extended periods of time and degrade to the corresponding hydroxylated PCBs. This is in particular true for PCB sulfate monoesters with two chlorine substituents *ortho* to the sulfate group. Most likely this is due to the increased acidity of the phenolic ring system. In summary, this series of PCB sulfate monoesters is available to study their physicochemical properties, their disposition *in vivo* and their interaction with SULTs and other enzymes.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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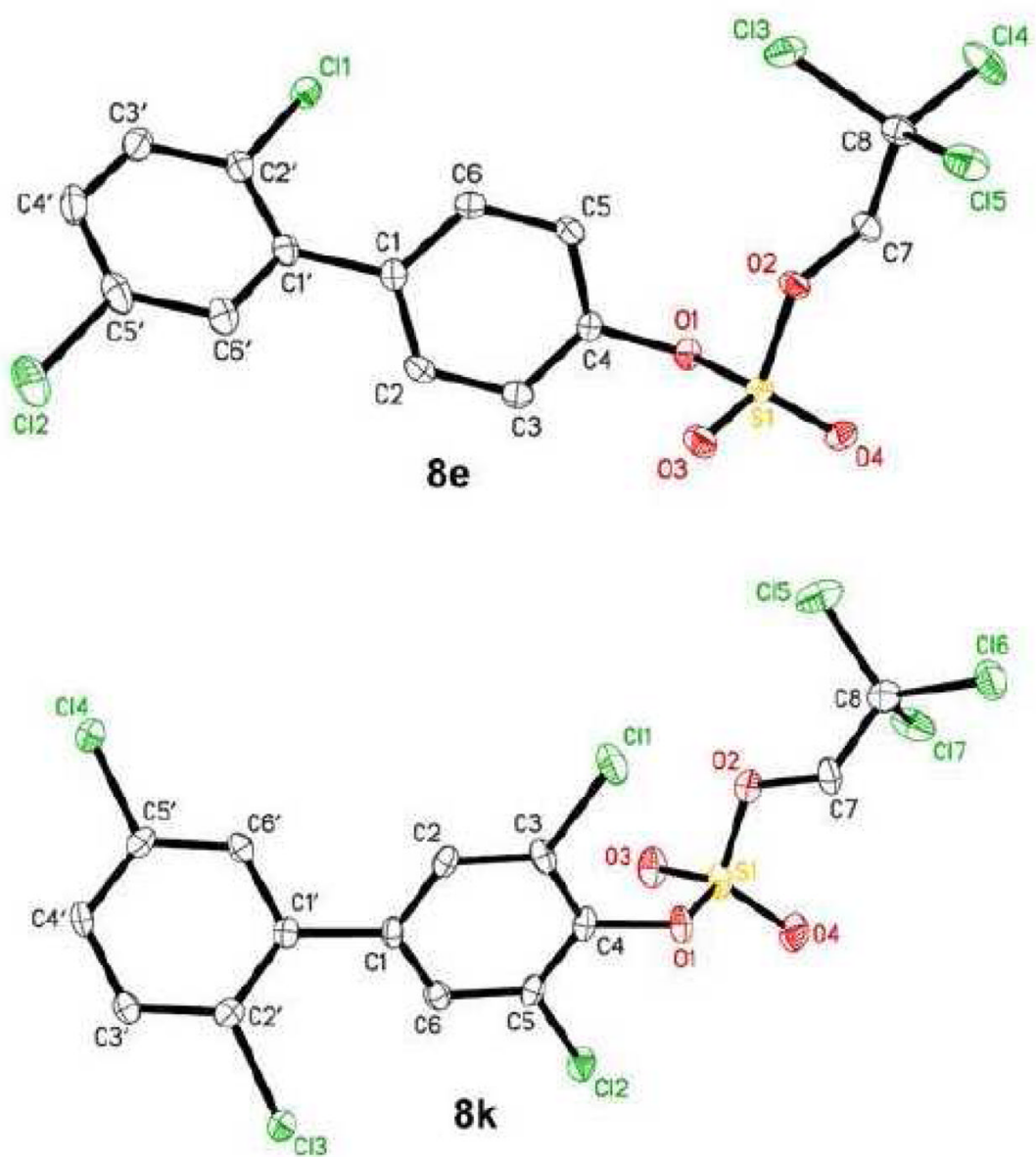
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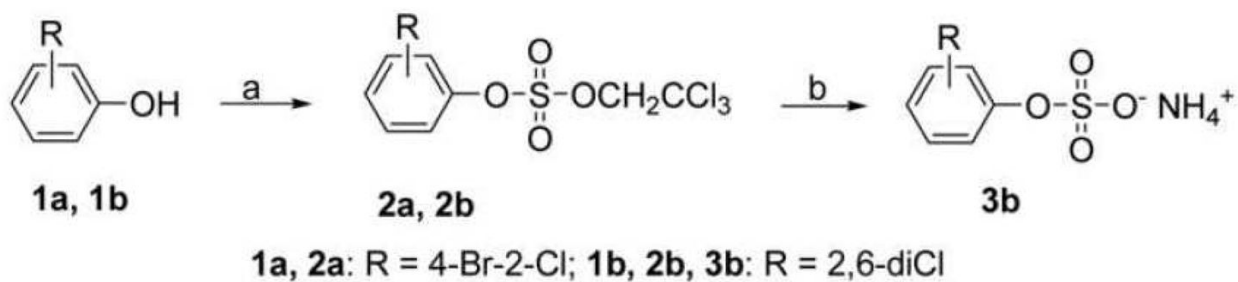
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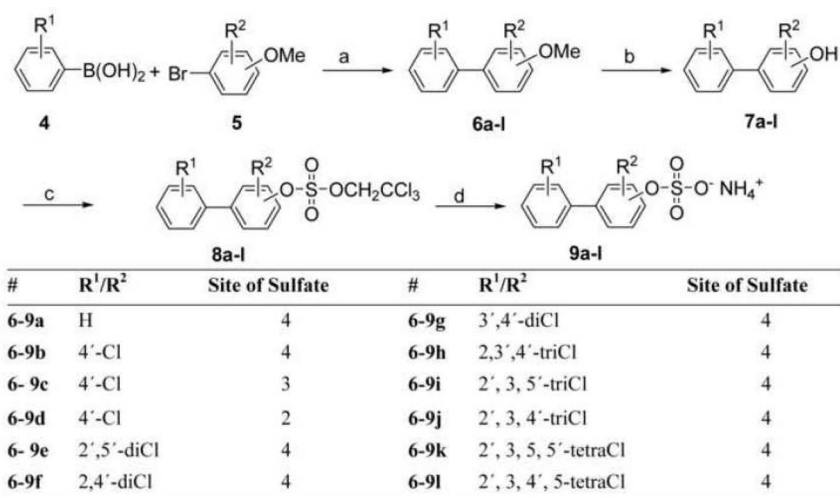




**Figure 1.** Molecular structure of sulfuric acid 2',5'-dichloro-biphenyl-4-yl 2,2,2-trichloroethyl ester (**8e**) and 2',3,5',5-tetrachloro-biphenyl-4-yl 2,2,2-trichloroethyl ester (**8k**) showing the atom labeling scheme. Displacement ellipsoids of **8e** and **8k** are drawn at the 50% probability level.

**Scheme 1.**

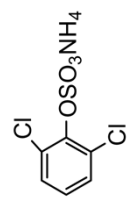
Sulfation of substituted phenols ((a) 2,2,2-trichloroethyl chlorosulfate, DMAP, dry  $\text{CH}_2\text{Cl}_2$ , 10 h; (b) Zn powder,  $\text{HCO}_2\text{NH}_4$ , MeOH). Sulfate monoester **3b** was identified *in situ* using TLC, but could not be isolated.

**Scheme 2.**

Synthesis of hydroxylated PCB sulfate ammonium salts ((a) 2 mol% Pd(PPh<sub>3</sub>)<sub>4</sub>, K<sub>2</sub>CO<sub>3</sub>, Toluene, 80 °C, 24h; (b) 1M BBr<sub>3</sub> in CH<sub>2</sub>Cl<sub>2</sub>, 10h; (c) 2,2,2-trichloroethyl chlorosulfate, DMAP, dry CH<sub>2</sub>Cl<sub>2</sub>, 10 h; (d) Zn powder, HCO<sub>2</sub>NH<sub>4</sub>, MeOH).

Yield/%	#	Sulfate monoester (3 and 9)	Yield/%
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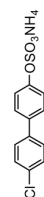
83



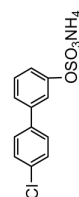
\_a



97



90

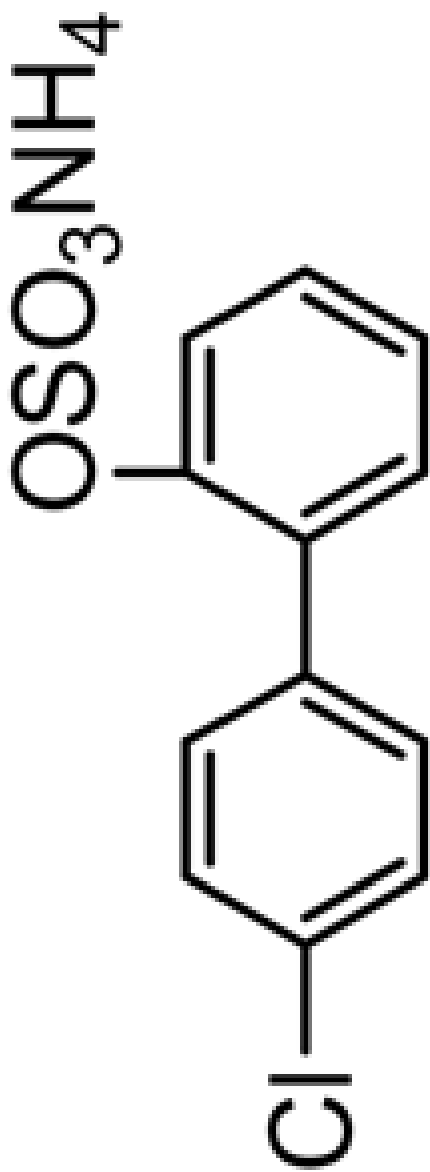


93

Yield/%	#	Sulfate monoester (3 and 9)	Yield/%
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75

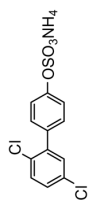
9d



83

81

9e



92

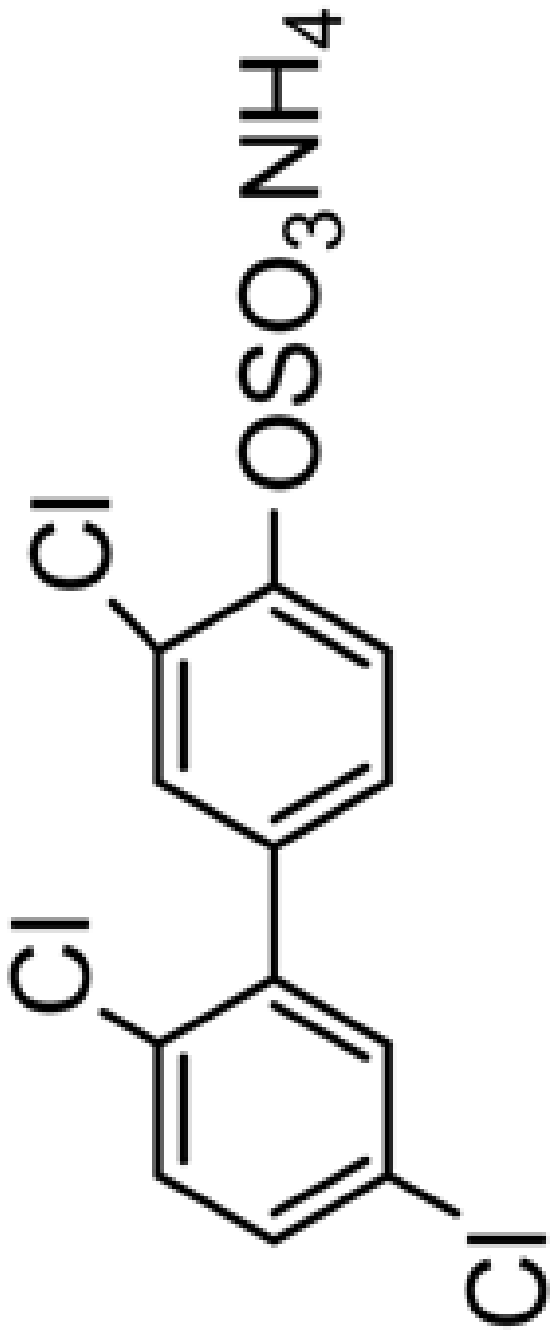
CE

3TCE



Yield/%	#	Sulfate monoester (3 and 9)	Yield/%
77	9f	 <chem>Clc1ccc(cc1)-c2ccc(cc2)S(=O)(=O)N</chem>	97
79	9g	 <chem>Clc1ccc(cc1)-c2cc(Cl)ccc2S(=O)(=O)N</chem>	94
83	9h	 <chem>Clc1ccccc1-c2cc(Cl)ccc2S(=O)(=O)N</chem>	96

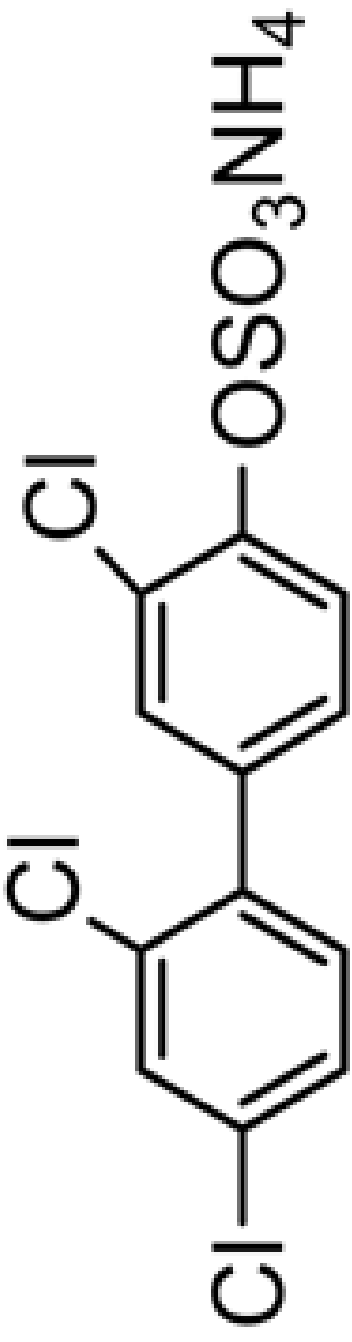
Yield/%	#	Sulfate monoester (3 and 9)	Yield/%
85	91		97
85	9j		85



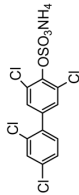
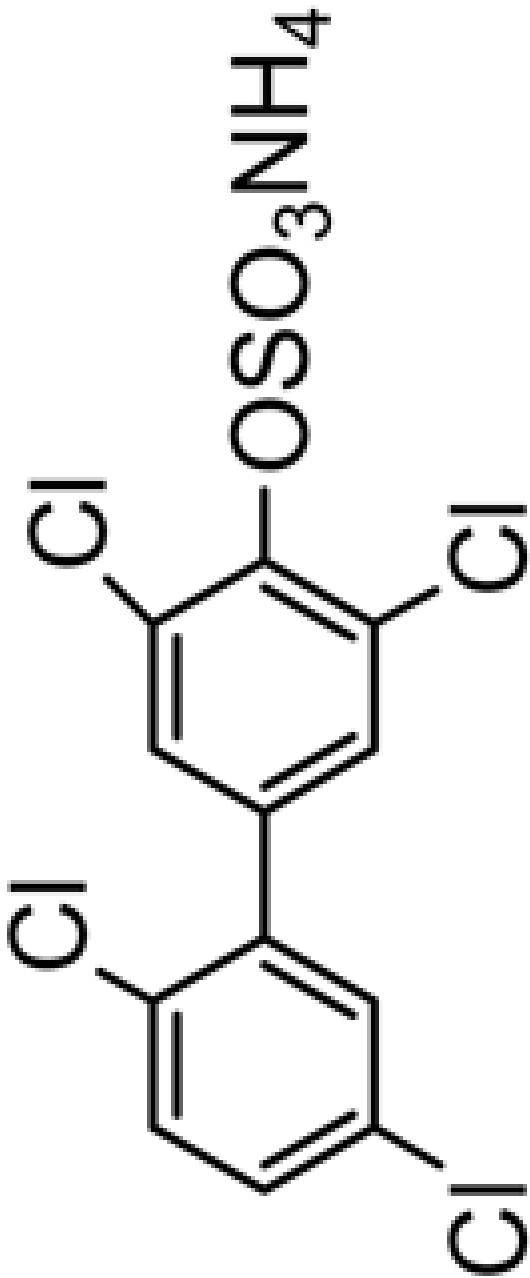
85

9j

85

O<sub>3</sub>TCE

Yield/%	#	Sulfate monoester (3 and 9)	Yield/%
83	9k		— <sup>a</sup>
94	9l		— <sup>a</sup>

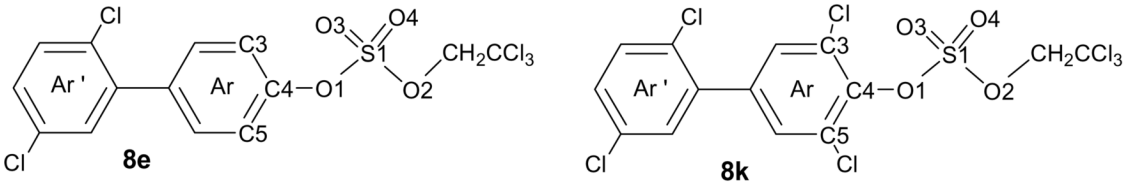


3TCE

O<sub>3</sub>TCE

**Table 2**X-ray crystallographic data for PCB TCE sulfate diesters **8e** and **8k**.

Property	<b>8e</b>	<b>8k</b>
Formula	C <sub>14</sub> H <sub>9</sub> Cl <sub>5</sub> O <sub>4</sub> S	C <sub>14</sub> H <sub>7</sub> Cl <sub>7</sub> O <sub>4</sub> S
<i>M</i>	450.52	519.42
<i>T</i> /K	90.0(2)	90.0(2)
wavelength	0.71073 Å	0.71073 Å
Space group	Monoclinic, P21/c	Monoclinic, P21/n
<i>a</i> (Å)	9.0188(2)	8.8951(2)
<i>b</i> (Å)	10.5961(2)	17.8153(4)
<i>c</i> (Å)	18.2405(3)	12.3093(3)
$\alpha$ (°)	90	90
$\beta$ (°)	90.9292(8)	104.5587(11)
$\gamma$ (°)	90	90
<i>V</i> (Å <sup>3</sup> )	1742.91(6)	1888.01(8)
<i>Z</i>	4	4
Calculated density (mg·m <sup>-3</sup> )	1.717	1.827
Absorption coefficient (mm <sup>-1</sup> )	0.968	1.181
F(000)	904	1032
Crystal size (mm)	0.37×0.33×0.26	0.25×0.24×0.20
$\theta$ range (°)	2.22 to 27.48 -11 ≤ <i>h</i> ≤ 11	2.06 to 27.48 -11 ≤ <i>h</i> ≤ 11
Limiting Indices	-13 ≤ <i>k</i> ≤ 12 -23 ≤ <i>l</i> ≤ 23	-23 ≤ <i>k</i> ≤ 17 -15 ≤ <i>l</i> ≤ 15
Reflections collected/unique	23838/3984	22255/4327
<i>R</i> <sub>(int)</sub>	0.0353	0.0337
Completeness to $\theta = 25.00$	99.9%	99.9%
Max. and minx. transmission	0.787 and 0.654	0.798 and 0.757
Data/restraints/parameters	3984/0/217	4327/0/235
Goodness-of-fit on <i>F</i> <sup>2</sup>	1.070	1.098
Final <i>R</i> indices <i>I</i> >2σ( <i>I</i> )	<i>R</i> 1 = 0.0306; <i>wR</i> 2 = 0.0694	<i>R</i> 1 = 0.0481; <i>wR</i> 2 = 0.1153
<i>R</i> indices (all data)	<i>R</i> 1 = 0.0403; <i>wR</i> 2 = 0.0741	<i>R</i> 1 = 0.0853; <i>wR</i> 2 = 0.1330
Largest diff. peak and hole (eÅ <sup>-3</sup> )	0.322 and -0.484	0.888 and -0.668

**Table 3**Selected bond length, bond angles and dihedral angles of PCB TCE sulfate monoesters **8e** and **8k**.


Property		<b>8e</b>	<b>8k</b>
Bond length (Å)	C4-O1	1.426(2)	1.405(4)
	S1-O1	1.5853(13)	1.600(2)
	S1-O2	1.5684(12)	1.564(3)
	S1-O4	1.4162(13)	1.415(3)
Bond angles (°)	O1-S1-O2	102.82(7)	101.65(13)
	O1-S1-O3	110.14(7)	110.29(14)
	O1-S1-O4	104.55(7)	104.36(14)
	O2-S1-O3	105.25(7)	105.42(15)
	O2-S1-O4	110.17(7)	110.25(15)
Dihedral angles (°)	O3-S1-O4	122.32(8)	122.93(16)
	Ar-Ar'	52.13(6)	50.03(10)
Deviation of O1 from Ar plane (Å)		0.092(5)	0.128(5)

Note: Ar and Ar' represent the aromatic rings of the biphenyl moiety.