### PCB IN RECYCLED PAPER PRODUCTS

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# ABSTRACT

PCB mixtures occur in amounts of 5 to 6000 ppb in paper products made of recycled material. Contents in samples made from Central-European raw material were higher than those from Finnish material. However, origin of some samples with high contents remained unknown.

#### INTRODUCTION

This study was initiated because we observed an intensive smell of PCB in letter envelopes made by a Finnish company from recycled "environmentally friendly" paper for our University. Determination of PCB gave contents of mixture analogous to Aroclor 1242 in a large envelope as high as 4 ppm (µg/g). Therefore, various products made from recycled paper were analyzed and the producers asked for origin of their raw material and possible other sources of PCB in production.

# SAMPLES AND ANALYSES

The samples studied are listed together with the determination results in Table 1. The samples were cut in small pieces, weighed, 2,4,6-trichlorobiphenyl added as an internal standard, and extracted in Soxhlet with a solvent mixture diethyl ether - acetone - petroleum ether 60-80 - hexane (1:5.5:9:2.5 v/v/v/v) for six hours. The extract was evaporated in Rotavapor and finally with nitrogen gas stream. One ml of hexane was added and the solution shaken three times with 2 ml of conc. sulfuric acid. Then, the solution was transferred to a column of 1 g of neutral alumina (activated in  $800^{\circ}C$  and deactivated with 5 % of water) in a Pasteur pipette. The column was eluted with 10 ml of hexane and the eluate evaporated to a small volume and 1  $\mu$ l of it injected to gas chromatograph (MICROMAT HRGC). Dual column system including quarz capillaries  $25 \text{ m} \log and 0.32 \text{ mm}$  ID and 0.25 µm phase thickness was used. The stationary phase in one column was SE-54 and in another OV-1701. EC detectors were used. Carrier gas applied was He 1 ml/min. The temperature program used was  $150^\circ - +5^\circ/\text{min} - 260^\circ\text{C}$ , and stay there 10 minutes. The quantitation was done by comparing intensities of 12 main peaks to the corresponding peaks of weighed standard of Aroclor 1242 (Fig. 1). Limit of determination was 1 mg/g.

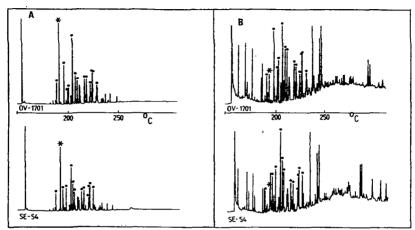


Fig. 1. Gas chromatograms of PCB fractions. The internal standard peak is marked by an asterisk and the peaks taken into quantitation by a dark spot. A. Reference mixture of Aroclor 1242. B. Tissue paper; 25.3 g sample containing 327 ppb of PCB.

| Sample          | Origin of raw material       | PCB ng/g (ppb) |
|-----------------|------------------------------|----------------|
| Letter envelope | Federal Republic of Germany  | 4093           |
|                 |                              | 1489           |
|                 | 1f                           | 1129           |
| 11              | "                            | 80             |
|                 |                              | 388            |
| ••              |                              | 433            |
| Small envelope  | Finland                      | 30             |
| Toalet paper    | Federal Republic of Germany  | 359            |
| Tource "Febor   |                              | 466            |
|                 |                              | 5              |
| **              | Austria                      | 127            |
|                 | Finland                      | 25             |
| •1              | "                            | 23             |
| **              | 11                           | 41             |
|                 | 11                           | 74             |
| Ticcue namer    | Finland                      | 327            |
| Tissue paper    | "                            | 22             |
| Cardboard box   | outer surface Unknown origin |                |
| Cardboard box   | central layer "              | 1010           |
| *1              | inner surface "              | 4787           |

TABLE 1. Samples and their contents of PCB as Aroclor 1242.

Organohalogen Compounds 4

# DISCUSSION

Paper products have been shown to contain 10-100 pg/g amounts of chloroaromatic ethers, such as PCDDs and PCDFs.<sup>1-3</sup> Origin of these compounds was first assumed to be chlorobleaching of pulp. However, other sources play a significant role, as was seen in the elevated concentrations of some PCDDs and PCDFs in recycled paper.<sup>1</sup> Also other chloroaromatics from pulp bleaching, like R-PCDFs and UPACs appear in papers.<sup>3,4</sup> Present analyses show that PCBs can occur in recycled paper products at levels which are several orders of magnitude higher than those of PCDDs and PCDFs. Because some PCB components are similarly toxic as 2,3,7,8-chlorosubstituted PCDDs and PCDFs, our observation calls for further control analyses of recycled papers. In addition, other EC-active persistent (H<sub>2</sub>SO<sub>4</sub>-resistent) compounds appear in these samples (Fig. 1) requiring more consideration.

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