Organochlorine pesticides (DDTs and HCHs) in soils from the outskirts of Beijing, China

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Abstract

Concentrations of HCH (hexachlorocyclohexane) and DDT (Dichlorodiphenyltrichloroethane) were determined in shallow subsurface (5–30 cm depth) and deep soil layers (150–180 cm depth) from the outskirts of Beijing, China. Concentrations of total HCHs (including α, β, γ, δ-isomers) and total DDTs (including p,p’-DDT, p,p’-DDE, p,p’-DDD, o,p’-DDT) in shallow subsurface soils ranged from 1.36 to 56.61 ng/g dw (median 5.25 ng/g), and from 0.77 to 2178 ng/g (median 38.66 ng/g), respectively, and those in the deeper layers were approximately an order of magnitude less. The spatial distribution of HCHs and DDTs reflected the known historical usage of these pesticides. No correlation between the concentrations of pesticides and soil organic matter content or clay content can be found. The factors affecting residue levels and compositions of DDT and HCH were discussed. The contour maps of β/γ ratios and DDT/DDE ratios for both the shallow subsurface and deep layer soils were drawn.

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Keywords: DDT; HCH; Soil residues; Spatial distribution; Beijing outskirts

1. Introduction

Organochlorine pesticides (OCPs) are of worldwide concerns because of their persistence (Dimond and Owen, 1996), bioaccumulation (Nakata et al., 2002), and toxicity to human and other animal and plant lives (Jones and de Voogt, 1999). 1,1,1-Trichloro-2,2-bis-(p-chlorophenyl) ethane (DDT) and 1,2,3,4,5,6-hexachlorocyclohexane (HCH) have been among the most widely used OCPs throughout the world. DDT was listed on the Stockholm Convention as one of 12 persistent organic pollutants (POPs). The origin and fate of OCPs in soils with different land use have been extensively studied in many countries. Although the usage of OCPs was phased out for decades, the elevated concentrations were still observed in many agricultural soils (Szeto and Price, 1991; Harris et al., 2000) and the relationship between sites of greatest application and current residue levels was found strong (Shivaramaiah et al., 2002). The release of OCPs from soils continues to be a source to the environment (Munn and Gruber, 1997; Meijer et al., 2003).

Technical HCH and DDT consist of about 50% of pesticide products in the 1970s (Zhang, 1996) and were heavily used for agricultural purposes in the Beijing area. From 1972 to 1974, 208.5 tonnes of HCH and 52.2 tonnes of DDT were sold annually in the eastern suburbs of Beijing, 399.3 tonnes of HCH and 98.7
tonnes of DDT in the western suburbs, and 2.25 tonnes of HCH and 39.59 tonnes of DDT in the remaining areas. An earlier survey has reported HCHs and DDTs residues to range from 22.0 to 254.0 ng/g dw (mean 107.4 ng/g), respectively, in wheat fields and those in orchard soils were generally in the range of 48 to 1375 ng/g and 1000 to 2000 ng/g, respectively. HCH and DDT were used in limited amounts in orchard soils in the late 1970s and applied at 22.5 kg ha\(^{-1}\) in wheat fields before harvest every year (Cooperation group, 1980). The usage of technical HCH and DDT was banned in 1983 in China.

In the past two decades, there is little information on whether these agrochemicals accumulate in soils in Beijing, and, if they do, to what extent. Recently, systematic investigations have been carried out in soils from the outskirts of Beijing to assess the soil quality in the Beijing area. This paper presents the current status of OCPs residues. The dataset generated will serve as a baseline for further studies.

2. Materials and methods

2.1. Sample collection

This was an intensive soil-sampling program. The sampling area was 1100 km\(^2\): 5 km east, 14 km south, 7 km west, and 6 km north from Beijing’s fourth-ring road. The shallow subsurface soil samples (5–30 cm depth) were collected every 1 km, and 16 cores were pooled to obtain one representative sample for every 16 km\(^2\) area. The deep layer soil samples (150–180 cm depth) were collected every 16 km\(^2\) area. The map of soil survey and sampling sites in the Beijing outskirts is shown in Fig. 1. Together, 47 shallow subsurface soil and 46 deep layer soil samples were analyzed. The soil samples are brown and mostly sandy loam and silt loam. Each fully mixed sample was air dried under a hood at room temperature and sieved to pass 35 mesh sieve. All samples were placed into amber glass bottles and refrigerated at –4 °C until analysis.

2.2. Extraction and cleanup

All solvents obtained from the Beijing Chemical Factory were of analytical grade and were redistilled in all-glass system before use. The reference standard mixtures of eight OCPs at a concentration of 100 mg/l were purchased from National Research Center for Certified Reference Materials, China. Florisil (Aldrich, USA) was activated for 16 h at 130 °C and kept in a desiccator before use. 2,4,5,6-Tetrachloro-m-xylene was used as the surrogate standard (Supelco, Bellefonte, PA, USA).
Ni$^{13}$ electron capture detector and a HP-5 column (25 m × 0.25 mm i.d., and 0.25 μm film thickness) with ultrapure nitrogen as carrier gas and make-up gas. GC conditions were: injector and detector temperature 275 and 300 °C, respectively, initial oven temperature 80 °C held for 1 min, increased at the rate of 10 °C/min to 150 °C and then ramped at 4 °C/min to 280 °C and held for 10 min. A 1 μl of the sample was injected in the splitless mode and the purge valve was opened at 0.75 min. Compounds were identified by retention time using external standards and quantified using peak area integration.

Selected samples were re-analyzed by a Hewlett Packard 6890 GC coupled to a 5973 MSD operated in the selected ion monitoring mode to further confirm the identification of compounds. A DB-5 MS (25 m × 0.25 mm i.d., 0.25 μm film thickness) capillary column was used. The GC oven temperature program was as described above. The transfer line and quadrupole temperatures were 250 and 150 °C, respectively. Helium was used as carrier gas. Data acquisition and processing were controlled by a HP Chem-Station data system.

For quality assurance and quality control, the procedural blanks, matrixes spiked by the standard solution were analyzed. None of the target compounds were detected in the procedural blanks. The recoveries were 250 and 150 % as described above. The transfer line and quadrupole temperatures were 250 and 150 °C, respectively. Helium was used as carrier gas. Data acquisition and processing were controlled by a HP Chem-Station data system.

Data for the GC oven temperature program were as directed by the surrogate from the soils ranged from 0.14 to 11.8 ng/g. The detection limits of the method were in the range of 4.3–2400 and 0.36–110 ng/g in Poland soils, respectively (Falandy these values were low because of long-term weathering. The concentrations of DDTs ranged from nondetectable to 11,800 ng/g with a geometrical mean of 9.63 ng/g in US soils (Aigner et al., 1998). In comparison with residual levels in archived background soils without any direct applications of pesticides in UK (0.1–10 ng/g; Meijer et al., 2001), in pristine areas such as Tibet plateau (DDTs: n.d.-2.83 ng/g; HCHs: 0.18–5.38 ng/g; Fu et al., 2001) and European high altitude mountains (DDTs: 1.7–13 ng/g; HCHs: 0.08–0.49 ng/g; Grimailt et al., 2004), and in some sites with relatively low application in South Korea (<3 ng/g; Kim and Smith, 2001), these values were high. When compared with data reported several years ago or recent data from places where degradation and dissipation of OCPs were retarded, these values were low because of long-term weathering. The concentrations of DDTs in orchard soils from New Zealand (median: 1170 ng/g; Gaw et al., 2003) were about 1–2 orders of magnitude higher than the values in our report.

The spatial distribution of HCHs (Fig. 2) and DDTs (Fig. 3) for both the shallow subsurface soils and deep layer soils was rather similar, with higher levels found at the east, west and northwest sites for both pesticides, reflecting the known historical usage in the region. Moreover, there was a HCH factory once located at the south part of the city; therefore, the relative high residue levels of HCHs in the shallow subsurface soils were detected in the south. Correlation of log-transformed concentrations of HCHs with those of DDTs exhibited high coefficients (shallow subsurface soils: $r^2 = 0.544$, $P < 0.01$), indicating the similar application history of technical HCH and DDT. One location with the greatest DDTs concentration (16–22) was not included due to a suspected recent source, which is discussed below.

4. Discussion

4.1. Factors affecting residue levels

A number of factors appear to control the environmental fate and behavior of organochlorine pesticides. Residue levels of organochlorine pesticides are affected by application history, agricultural practices, soil phys-
Table 1
The range, mean and median of organochlorine pesticides (ng/g dw) and selected soil properties

<table>
<thead>
<tr>
<th></th>
<th>Shallow subsurface soil (n = 47)</th>
<th>Deep layer soil (n = 46)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Minimum</td>
<td>Maximum</td>
</tr>
<tr>
<td>OC (%)</td>
<td>0.66</td>
<td>3.25</td>
</tr>
<tr>
<td>Clay (%)</td>
<td>6.80</td>
<td>22.33</td>
</tr>
<tr>
<td>α-HCH</td>
<td>0.32</td>
<td>8.19</td>
</tr>
<tr>
<td>β-HCH</td>
<td>0.37</td>
<td>45.11</td>
</tr>
<tr>
<td>γ-HCH</td>
<td>n.d.</td>
<td>12.86</td>
</tr>
<tr>
<td>δ-HCH</td>
<td>n.d.</td>
<td>5.85</td>
</tr>
<tr>
<td>HCHsb</td>
<td>1.36</td>
<td>56.61</td>
</tr>
<tr>
<td>β[\gamma]</td>
<td>0.09</td>
<td>29.47</td>
</tr>
<tr>
<td>p,p′-DDE</td>
<td>0.22</td>
<td>832.37</td>
</tr>
<tr>
<td>p,p′-DDD</td>
<td>0.15</td>
<td>50.38</td>
</tr>
<tr>
<td>o,p′-DDT</td>
<td>n.d.</td>
<td>533.23</td>
</tr>
<tr>
<td>p,o′-DDT</td>
<td>n.d.</td>
<td>1924.76</td>
</tr>
<tr>
<td>DDTsc</td>
<td>0.77</td>
<td>2178.55</td>
</tr>
<tr>
<td>DDT/DDE</td>
<td>0.006</td>
<td>63.9</td>
</tr>
</tbody>
</table>

* a Nondetectable.
* b Sum of HCH isomers.
* c Sum of DDT isomers and metabolites.

Fig. 2. The spatial distribution of HCHs in soils from the outskirts of Beijing.
ico-chemical properties, as well as physico-chemical properties of agrochemicals and meteorological factors such as temperature and rainfall etc. (Szeto and Price, 1991; Boul et al., 1994; Spencer et al., 1996).

OCPs are hydrophobic and considered to be easily adsorbed by soil organic matter. A study carried out in mountain soils from the subtropical Atlantic (Teide, Tenerife Island) found that DDTs and HCHs exhibited a high dependence on soil total organic carbon (Ribes and Grimalt, 2002). However, in our present study, no correlation between the concentrations and soil organic carbon content was found, indicating that residue levels were a reflection of OCPs application history and dissipation rates rather than air–soil equilibrium, which is similar to agricultural soils from Alabama (Harner et al., 1999) and from Georgia and South Carolina (Kannan et al., 2003). Residues of OCPs dissipated in loamy sand much faster than in the muck and higher levels of OCPs were often detected in the muck (DDTs: 3997 ng/g, Szeto and Price, 1991; 11,800 ng/g, Aigner et al., 1998). However, in our study, Sample Nos. 16–22, in which the highest DDTs was found (2178 ng/g), had both the soil organic carbon content (1.01%) and the clay content (7.03%) lower than the median values.

OCP residues have been studied in different soil horizons (Boul et al., 1994; Miglioranza et al., 1999; Weiss, 2000; Feng et al., 2003), and much attention was paid to the shallow subsurface soils. There are very limited data available for the deep soil layers, which is an indicator of potentiality of OCPs present in the groundwater contamination. In our study, the concentrations of HCHs and DDTs in the deep layer soils (150–180 cm depth) were about an order of magnitude lower than those in the shallow subsurface soils, because of mechanical mixing and irrigation in the agricultural soils as well as long-term leaching.

Fields with crops that require higher OCPs use usually contain higher residue levels even after long-term weathering. Our shallow subsurface soil samples were pooled from 16 soil cores with different land uses, and
it was very difficult to assess the effect of different land use on soil residues of OCPs. As for the deep layer soils, the highest concentrations of \( p,p'-DDE \) (range: 0.12–2.26 ng/g; median: 1.26 ng/g) were found in orchard soils among wheat fields (range: 0.06–0.69 ng/g; median: 0.17 ng/g), vegetable gardens (range: 0.09–2.24 ng/g; median: 0.17 ng/g), wood soils (range: 0.07–1.87 ng/g; median: 0.19 ng/g), and maize fields (median: 0.05–0.21 ng/g; median: 0.16 ng/g). No significant difference was found for HCH isomers. The results are consistent with the application history of OCPs (Cooperation group, 1980).

Beijing is located in the temperate region. It is cold and dry in winter with the lowest temperature of \(-15.4\) °C, while in summer it is hot and humid with the highest temperature of 38 °C. Mean yearly precipitation is about 630 mm. The persistence of OCPs in soil is greater in temperate climates compared to the tropics (Iwata et al., 1994).

4.2. Compositions of OC residues

Technical HCH mixtures containing \( \alpha-(67\%) \), \( \beta-(10\%) \), \( \gamma-(15\%) \), \( \delta-(8\%) \) HCH had been used in China. \( \beta\)-HCH has all chlorines in equatorial positions and the lowest vapor pressure. It is the most persistent isomer and tends to accumulate in soil (Li, 1999). The mean percentages of HCH isomers in the shallow subsurface soils were as follows: \( \beta > \gamma > \alpha > \delta \), which confirmed that the technical HCH had not been in use in the Beijing area for quiet a long time, while those in the deep layer soils were: \( \gamma > \alpha \approx \beta > \delta \) (Fig. 4A), indicating that \( \gamma\)-HCH and \( \alpha\)-HCH had the higher leaching ability than \( \beta\)-HCH and \( \delta\)-HCH (Cooperation group, 1980). The ratios of \( \beta\)-HCH to \( \gamma\)-isomer have been used in this study to indicate historical use and trace pollution sources. Forty five out of 47 shallow subsurface soils had \( \beta/\gamma \) ratios ranging from 0.09 to 29.47 with a median of 2.30. As for the deep layer soils, the \( \beta/\gamma \) ratios were in the range of n.d. to 7.78 with a median of 0.52, lower than those of the shallow subsurface soils because of \( \gamma\)-HCH being readily leached into the soil (Simonich and Hites, 1995; Miglioranza et al., 1999). The contour maps of the ratios of \( \beta/\gamma \) isomers provide a clear overview of the proportions of two HCH isomers in the studied area (Fig. 5). The greatest values for both shallow subsurface soils and deep layer soils were located at the northwest sites where high total HCHs were also observed.

Technical DDT is typically composed of 77.1% \( p,p'\)-DDT, 14.9% \( o,p'\)-DDT, 4% \( p,p\)-DDE, and some other trace impurities. The mean percentages of individual compounds in both shallow subsurface soils and deep layer soils were as follows: \( p,p'-DDE > p,p'-DDT > p,p'-DDD > o,p'-DDT \) (Fig. 4B). The mean percentage of \( p,p'-DDE \) in the deep layer soils was lower than that in the shallow subsurface soils, but that of \( p,p'-DDD \) was higher, because DDT is likely to be dechlorinated to DDD in the anaerobic condition in the deep layer soils. It has been suggested that \( o,p'-DDT \) and \( p,p'-DDT \) have declined at similar rates in Alabama, USA soils (Harner et al., 1999). However, in our present study, the median values of \( o,p'/p,p'-isomers \) are 1:2.5 and 1:1.3 for the shallow subsurface soils and deep layer soils, respectively, which are different from its "starting" value of 1:5.2.

The ratio of parent compound to metabolites was used to infer sources and qualitatively judge the age of contaminant residues in soil. DDT/DDE ratios were quite variable, ranging from 0.006 to 63.9 with a median of 0.38 for the shallow subsurface soils, and from 0.05 to 231.9 with a median of 0.86 for the deep layer soils. Most soils contained more DDE than DDT, indicating DDT residues are from "old" sources. The contour maps (Fig. 6) showed that highest DDT/DDE ratios for the shallow subsurface soils and deep layer soils were located at the southwest and southeast sites, respectively, where high DDTs levels were found. Extremely high ratios were found in the deep layer soils, probably because of the long-term leaching of \( p,p'-DDT \) and retarded degradation of DDT to DDE in the deep layer soils (Miglioranza et al., 1999; Harris et al., 2000). Great proportions of \( p,p'-DDT \) were also observed in agricultural soils of central Germany (Manz et al., 2001) and of southern Ontario (Meijer et al., 2003), and in some western USA soils (Hitch and Day, 1992). It was found that high copper (Van Zwieten et al., 2003) or arsenic (Gaw et al., 2003) content could inhibit the degradation of...
DDT to DDE in soil. However, the total metal contents (data provided by Professor H.X. Cheng) including copper and arsenic in the shallow subsurface soils were generally at the same levels, no correlation between DDT/DDE ratios and metal contents can be found. Another study carried out in Beijing reported that accumulative levels of DDT in the mother’s milk declined irregularly, and the authors suggested that this might result from the possible use of dicofol which contains 5–10% DDT as impurity (Yu et al., 2001). Dicofol is classed as organo-
chlorine pesticides, and DDT is one of the intermediate products during its manufacture. Now the use of dicofol was restricted in China because relatively high levels of DDT contamination were found in soils where dicofol was applied.

5. Conclusions

The usage of the technical HCH and DDT has been illegal for decades in China, and OCPs residues in most sites were relatively low. The residual levels in the deep layer soils were an order of magnitude lower than those in the shallow subsurface soils. The spatial distributions of HCHs and DDTs in the outskirts of Beijing were quite similar, which reflected the known historical usage. No correlation of residue levels with soil organic carbon content and clay content was found, indicating that the soil–air system was out of equilibrium. In some typical sites, the relative high residue levels of DDTs were observed, which may result from a combination of heavy historical use and retarded degradation of DDT in soils. Moreover, it was quite possible that dicofol may be a “new” source, which remains to be further studied.

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