Sediment contamination of an urban canal—a case study approach for an integrated assessment of organic sediment contamination

M. Ricking¹,² & K. Frische³ & J. Schwarzbauer³

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Abstract

Purpose The pesticide DDT and its metabolites represent a contamination risk for the aquatic environment, especially the polar metabolite DDA. The study provides a quantitative assessment of long-term pollution risks from sedimentary DDT residues with a special focus on DDA. It presents an overview of the contamination range of different DDX compounds in the sediments of a canal in Berlin (Germany), resulting from a former industrial point source that has implications for drinking water resources in the nearby area. The comprehensive analysis scheme provides information on free accessible and potentially metabolized precursors also in the non-extractable residues. This allows a quantitative assessment of the DDA pollution potential derived from the sedimentary DDT residues.

Materials and methods The area was investigated for fine-grained sediment by means of a geo-electric mapping. Twelve sediment cores were taken in four areas (three in each section). A wider range of precursor metabolites has been included due to their transformation potential to the polar metabolite DDA. The sediments were analysed quantitatively for extractable and easily releasable fractions by application of a variety of degradation techniques as well as a dispersion extraction procedure on the sediment samples. These extracts were fractionated and subsequently analysed by GC-MS.

Results and discussion Concentrations were obtained for extractable and bound metabolites. Different scenarios for the calculation of the amount of contaminated sediment are displayed as a tool for contamination assessment. The formation potential of DDA as the water-soluble metabolite is presented. Several precursor metabolites, e.g. DDD and DDMS, extractable from the sediment organic matter, revealed a high potential for a long-term formation of DDA, especially in the easily releasable fraction (via hydrolysis) with a mean concentration of up to 11,000 μg g⁻¹ dry sediment. The resulting DDA contamination potential represents a significant pollution risk for the groundwater from a downstream waterworks area and by remobilisation into the whole ecosystem and adjacent rivers.

Conclusions The application of the presented methods provides a tool for a quantitative assessment of the long-term release potential of DDA under different scenarios by a comprehensive analysis of contaminated sediments (and soils). This approach can be transferred to pollutants that are also characterized by a complex metabolism accompanied by bound residue formation.

Keywords DDX-related compounds · Sequential degradation · Non-extractable residues (NER) · DDA formation potential

1 Introduction

In sediments, organic pollutants can become incorporated by various ways into the organic matter and metabolized microbially. With respect to the aquatic environment, organic contaminants can be associated with the suspended particulate matter and released again by desorption processes (Northcott and Jones, 2000; Semple et al. 2003). Within the sedimentary organic matter (SOM), which consists amongst others of humic substances, an extractable and non-extractable fraction (NER) of pollutants can be differentiated and analysed separately by sequential degradation reactions (Eschenbach et al. 1994). Operationally defined, the extractable fraction...
represents the direct bio-available/bio-accessible contamination potential for the aquatic environment due to the weak bonding to the SOM by van der Waals forces or ionic attraction. The non-extractable fraction can be divided into reversible bound and irreversible bound residues, representing temporary or final depositions, respectively (see Section 2.4).

The chemical interactions of organic substances with sediments and soils (natural polymers or geo-sorbents) may result in strong binding and slow subsequent release rates that significantly affect remediation rates and endpoints (Scheunert and Schröder 1998). Hence, this is an important effect in particular for the risk assessment of pollutants. The underlying physical and chemical processes or phenomena potentially responsible for this apparent sequestration of persistent, bioaccumulative and toxic substances (PBTs) by geo-sorbents are only partially understood (Nowak et al. 2014; Kästner et al. 2016; Schäffer et al. 2018; Taylor et al. 2019). Recent observations suggest that PBT interactions with geo-sorbents comprise different organic and inorganic surfaces and matrices, and distinctions are clearly visible along these lines particularly with regard to the roles of inorganic micropores (e.g. in soot or clay minerals), natural organic matter components, combustion residue particulate carbon and spilled organic liquids. Definitely, SOM is not uniformly spread over all solid phases and not universally treatable as a well-defined organophilic phase with different $K_{om}$ values, as more than one order of magnitude difference in calculated distributions between $K_{om}$ and solubility or log $K_{ow}$ are reported (Takada and Eganhouse 1998; Ehlers and Loibner, 2006). Partition-like interaction with a “microscopic non-polar organic environment”, humic-membrane-like structure, polar hydrophobic exterior and non-polar hydrophobic interior is to be expected.

Detailed information related to the formation of non-extractable residues has been published by Luthy et al. (1997), Northcott and Jones, (2000), Schwarzbauer et al. (2003), Ehlers and Loibner (2006) and Kästner et al. (2016). The formation of NER in geo-polymers is realized through stronger ionic and covalent bonding, entrapment (inclusion in cavities) and biological processes. The reversibility of sorption-desorption is based on geochemical parameters like functional groups, quality of the SOM, pH, Eh and water chemistry. Important for the microbial degradation are the strength of the bonding and the degree of incorporation into the geo-polymers. Toxic effects are reduced due to limited bio-availability of bound pollutants resulting in a higher persistence and modified degradation rates relative to laboratory model systems (Garrison et al. 2014). Sequential chemical treatments are applied to selectively release compounds depending on the character of incorporation (Semple et al. 2003; Schwarzbauer et al. 2003; Kalathoor et al. 2015).

To address the complete contamination of riverine systems, the non-extractable fraction, characterized by incorporation due to covalent bonding, can reveal a long-term contamination risk due to their remobilisation potential of organic compounds to the environment. Especially, polar organic compounds exhibiting considerable solubility represent a contamination risk for the aquatic ecosystem.

According to Zhu et al. (2019), the degradation of DDT in the NER was differentiated into three categories (including the initial DDT) in order to clarify not only the further cleavage of the aromatic ring structure but also the degree of degradation by dehalogenation, functional group formation and loss of a C-atom. Halogenated organic compounds are prone to a high environmental persistence. Pesticides such as DDT, exhibiting the main contamination revealed in this study, are well-known pollutants. They are characterized by high enrichment factors in sediment (Eganhouse and Pontolillo, 2008; Eganhouse et al. 2018, Taylor et al. 2019), resulting in a strong tendency for geo-accumulation, act as sinks and long-term sources for aromatic organic halogens (AOHs).

DDT has been applied worldwide between the 1940s and 1986 as an insecticide in agriculture and is still allowed for treatment of vectors such as malaria (WHO 2011). It is now classified as one of the “Dirty Dozen” according to the Stockholm Convention. DDT is also an intermediate in the formulation of Dicofol as an acaricide, miticide and antifoulant treatment. Due to the high persistence and chronic toxicity, its use as a pesticide has been prohibited worldwide since the early 1970s worldwide. Even very recent publications from Asia (Babut et al. 2019) and Europe (Qu et al. 2019) report a high percentage of DDX (sum of all analysed DDT metabolites) in the line of organochlorine pesticides (OCPs). A recent review states the effort of DDX-removal from environmental samples is still a technical topic in the EU (Vagi and Petsas 2020).

The water solubilities of DDT, DDD and DDE at 25 °C are 5.5 μg L$^{-1}$, 20.0 μg L$^{-1}$ and 14 μg L$^{-1}$, respectively. According to Lee et al. (1994), the log $K_{ow}$ values are 6.91, 6.22 and 6.96, respectively.

Environmental contamination by persistent organic compounds is referred to both point sources and diffusive sources. Point sources may be related to industrial production effluents, municipal emissions or agricultural activities. Diffusive sources may be surface run-off, deposition and atmospheric transport including long-range atmospheric transport (LRAT) which is the major pathway for the transfer of persistent pollutants to remote sites (Kallenborn 2006; Becker et al. 2008).

The global distribution of DDT and its metabolites has been studied intensively (e.g. Marei et al. 1978; de la Cal et al. 2008; Bettinetti et al. 2016; Mitra et al. 2019), but DDA has been neglected widely. Only a few studies considered this metabolite (Heberer and Dünnbier 1999; Schwarzbauer et al. 2003; Heim et al. 2005; Wan et al. 2005; Chen et al. 2012; Garrison et al. 2014), despite an early identification in 1945 (Strohlman and Smith 1945; White and Sweeny 1945).
In the present study, subaquatic sediments of the Teltow Canal in Berlin (Germany) were intensively sampled and analysed close to a point source (see Fig. 2).

The map shows the Berlin area within the Elbe River Basin and a detailed view of the sampling area on the Teltow Canal with the point source at the eastern end of the canal. The factory (BC) discharged production wastewater from pesticides into the canal, where the Johannisthal waterworks some 6 km downstream produced drinking water.

The canal is used as a drainage for rainwater, industrial wastewater, wastewater treatment plants and as a shipping canal for industrial supply, with a length of approximately 38 km and a mean depth of 2 m. It is characterized by a very slow flow velocity and discharge (WSA 2018, average discharge at the lock Kleinmachnow: 8.97 m$^3$ s$^{-1}$), occasional flow direction reversal, a high sedimentation rate in favoured accumulation areas and mostly anoxic conditions at the sediment-water interface. High accumulation rates and undisturbed deposition are present in these subaquatic sediments. The former industrial plant is located at the eastern border of the canal, close to the Dahme River. Between the 1940s and 1986, about 60,000 t of DDT were synthesized and formulated on site (Heim et al. 2005). Emission of untreated neutralised effluents leads to a strong contamination of canal sediment by DDT and other halogenated compounds for decades. Therefore, from 1995 to 1999, after removal of an underwater barrier, an excavation of approx. 180,000 t of contaminated sediment (~135,000 m$^3$) was realized, which led to long-term remobilisation processes (Heim et al. 2005; Frische et al. 2010). The production data are published in Heim et al. (2005).

A few kilometres downstream of the study area, a waterworks is located using riverbank filtration, posing a risk of water contamination by the DDT-metabolite DDA and all precursor metabolites (see Fig. 1) through remobilisation processes via professional shipping activity.

The aim of this study was to obtain not only an overview on the actual contamination range referring to the formation potential of water-soluble DDA (water solubility at pH 5–10 is 20–59 g L$^{-1}$; Frische et al. 2010) from the remaining DDT-contaminated sediment of the canal but also the quantitative estimation of the sedimentary contamination release potential. The release behaviour of DDA under dynamical conditions was experimentally investigated to get a more precise insight into the environmental fate. Additionally, further information about the degradability/persistence of DDA is deduced (Fig. 2).

In the literature, the kinetics in biota (animals, humans) are described as fast; the environmental kinetics are mostly unknown.

DDT serves as a representative for compounds for other environmentally relevant compounds with similar toxicity, bio-accumulation potential, physicochemical features, degradability and formation potential of NER like methoxychlor and low molecular PAHs (Eschenbach et al. 2001).

2 Materials and methods

2.1 Field mapping and sampling

The determination of the sediment thickness was a prerequisite for targeted sediment sampling in the Teltow Canal and the information about the sediment structure and the distribution of organoleptic muds. With the help of geophysical measuring and interpretation methods, the spreading of muds in a 1.25-km long section of the canal could be inquired. For this purpose, a combination of hydro acoustic measurements (echo sounder) and special methods of water geo-electric was applied.

2.1.1 Preliminary remarks on echolocation

The study concept is based on a combination of hydro acoustic measurements (dual-frequency sediment echo sounder) and geo-electrical water bottom mapping. Already during the measurements, it was found that the sediments in the study area were relatively “sound-reflecting” sediments with an unexpectedly hard sediment surface. The echolocation was carried out mainly to determine the depth of the water and for the evaluation of the sandy basis of the canal for the interpretation of the measurement data.

2.1.2 Methodological basics

The special process of water geo-electrics was realized through measuring cables that are towed directly above the sediment surface. From these results a high spatial resolution for the uppermost sediment layer regardless of the water depth, whose thickness can be determined with an accuracy of approx. 10%.

As a physical principle, geo-electrics uses the different behaviour of substances with regard to their resistivity, in order to be able to withstand an approximate horizontal storage of the layers being able to determine material properties and spatial limitations. The specific electrical resistance of a layer is essentially determined by the following components:

- Solid sand, gravel grains: = infinitely high
- Pure clay: mostly under 10 $\Omega$ m$^{-1}$
- Average waters 10 ... 40 $\Omega$ m$^{-1}$
- Sediments with high organic content: sludge $\leq$ water

The resistance of a loose sediment depends on the water saturation of the water, and the particle size distribution and
charged surfaces like clay minerals present in mud. This relationship forms the basis for a layer structure of the substrate. The field mapping of the sediment area was performed by geo-electric measurements in cooperation with Inc. FUGRO. Three profiles were taken along a stretch of 1250 m. Distinctive areas in the sediment body are rich in humic matter with elevated TOC contents (see Fig. 4; see Table 1) with a geo-electric resistance of less than 35 Ω m$^{-1}$ in the sediment.
body. The completely contaminated sediment represents ~22% of the area. It was divided into sub-areas A–D in order to facilitate the calculation of the total sediment depot. Three selected sediment cores were taken from each sub-area—according to the data by the echo-sounding profiles and to cover a wide range of organic sediment in each sub-area.

In total, twelve subaquatic sediment cores with a maximum length of 65 cm were taken from the canal by means of a gravity coring system according to Eijelkamp (8 cm diameter, modified). Each sediment core was divided into a maximum of four parts; the upper sandy layer with TOC contents \(\leq 0.01\%\) was abolished. Aliquots of the thoroughly homogenised sediment samples were freeze-dried including the pore water and stored at 4 °C.

For the later calculation of the risk potential, two different geometries of the sediment body were used, one with a trapezoid shape (bottom layers are smaller than the surface layers) and one with a column shape (no difference with depth). The dry mass of each layer was calculated via the dry mass of the sediment material and bulk density of the sediment core multiplied with the integrated area from the mapping. The total calculated sediment masses are given in the supporting information (see Table SI-2).

The TOC content was analysed by means of the evolving CO\(_2\) by heating under oxygen at 1000 °C in a C-Mat (LECO, Mönchengladbach, Germany).

### 2.2 Release experiments

For the short-time release experiment in duplicate, 20 g wet contaminated sediment from the canal, taken in 2006 (stored under anoxic conditions at 4 °C in the dark) was eluted with 480 mL water under oxic (through addition of ascorbic acid) and anoxic conditions in brown glass bottles in the darkness.

### Table 1: Lithology of the sediment cores with TOC content and bulk density

| Area | Core | Layer | Depth (cm) | TOC (%) | Bulk density (g cm\(^{-3}\)) |
|------|------|-------|------------|---------|-----------------------------|
| D    | 1    | Top   | 20–35      | 9.29    | 0.70                        |
|      | 1    | Middle| 35–45      | 6.91    | 0.72                        |
|      | 1    | Base  | 45–60      | 4.52    | 0.76                        |
|      | 2    | Top   | 09–26      | 1.30    | 0.92                        |
|      | 2    | Middle| 26–43      | 0.86    | 0.83                        |
|      | 3    | Middle| 26–44      | 3.82    | 0.82                        |
|      | 3    | Base  | 44–62      | 1.82    | 0.92                        |
| C    | 4    | Top   | 12–27      | 9.46    | 0.67                        |
|      | 4    | Middle| 27–44      | 10.31   | 0.65                        |
|      | 4    | Base  | 44–59      | 11.16   | 0.66                        |
|      | 5    | Top   | 11–26      | 9.11    | 0.60                        |
|      | 5    | Middle| 26–43      | 6.12    | 0.71                        |
|      | 5    | Base  | 43–58      | 3.33    | 0.82                        |
|      | 6    | Top   | 06–21      | 7.53    | 0.60                        |
|      | 6    | Middle| 21–49.5    | 5.13    | 0.65                        |
|      | 6    | Base  | 49.5–64.5  | 2.73    | 0.76                        |
| B    | 7    | Middle| 18–29      | 4.60    | 0.94                        |
|      | 8    | Middle| 18–31.5    | 9.62    | 0.67                        |
|      | 8    | Base  | 31.5–44.5  | 0.07    | 1.03                        |
|      | 9    | Middle| 16–28      | 0.87    | 0.90                        |
|      | 9    | Base  | 28–40      | 0.01    | 1.12                        |
| A    | 10   | –     | –          | –       | –                           |
|      | 11   | Top   | 9–19       | 8.81    | 0.56                        |
|      | 11   | Middle| 19–29      | 0.02    | 0.81                        |
|      | 12   | Middle| 16–32      | 9.34    | 0.69                        |
|      | 12   | Base  | 32–47.5    | 5.96    | 0.73                        |
|      | 13   | Top   | 10–22      | 7.09    | 0.70                        |
|      | 13   | Middle| 22–34      | 1.14    | 1.03                        |

Fig. 2 Map of the Elbe River with an insert of the sampling area in Berlin, located at the eastern end of the Teltow Canal (detail view of the sampling area and the production site is marked in red)
The water was stirred gently to simulate a current flow for resuspension. Water sampling was performed three times, after 1, 7 and 42 days with a subsequent replacement of the water. All the water was extracted via a liquid-liquid-extraction in triplicate according to Frische et al. (2010) and separately analysed.

2.3 Chemicals and reagents

All chemicals, reagents and solvents were purchased from Sigma–Aldrich (Taufkirchen, Germany), Merck (Darmstadt, Germany) and Roth (Karlsruhe, Germany). Solvents were distilled and purity was checked by gas chromatography. All glassware was pre-cleaned with acetone and n-hexane prior to usage.

2.4 Degradation procedures

The samples were chemically degraded by a sequential procedure to reveal the extractable and different non-extractable fractions (see Fig. 3). Recoveries were taken from Schwarzbauer et al. (2003) and Heim et al. (2005), determined by spiking experiments on annealed sediments. Details of the method are published in Schwarzbauer et al. (2003) and summarized in Fig. 3. The procedures applied to obtain the different fractions are described briefly in the following:

**Extraction (extractable fraction)** As a first step prior to chemical degradation, all sediment samples were extracted sequentially with 40 mL of methanol and mixtures of hexane/acetone (1:1 v/v) by shaking for 24 h in the dark and ultrasonication at 2 × 450 W (Fa. Badelin, Berlin, Germany) in a water bath for 30 min. After centrifugation, the combined crude extract was reduced in volume and followed separating by liquid chromatography as described in Schwarzbauer et al. (2003).

**Chemical degradation** Prior to applying the sequential chemical degradation procedure on sediment samples, a possible formation of analytical artefacts was checked. For this purpose, various reference substances (2,4′- and 4,4′-DDD, 2,2-bis(4-chlorophenyl)acetonitrile - 4,4′-DDCN, 2,2-bis(4-chlorophenyl)-1,1-dichloroethene - 4,4′-DDE, 2,2-bis(4-chlorophenyl)-1-chloroethene - 4,4′-DDMU and 2,2-bis(4-methoxyphenyl)-1,1,1-trichloroethane - 4,4′-MDT) were subjected individually to the chemical degradation procedure but formation of artefacts have not been observed.

**Hydrolysis (easily releasable fraction)** Pre-extracted sediment samples were mixed with 120 mg of KOH dissolved in a solution of 0.2 mL of water and 8 mL of methanol. The mixture was sealed and heated at 105 °C for 24 h. After this alkaline hydrolysis procedure, the mixture was filtered, acidified to pH 3–5 with 10% hydrochloric acid solution and, subsequently, extracted three times with dichloromethane. Thereafter, the extract was dried, reduced in volume to 0.5 mL and separated into two fractions by column chromatography using dichloromethane (fraction 1) and methanol (fraction 2) as eluent. Prior to GC/MS analysis, 50 μL of an internal standard was added.
standard solution containing 90 ng μL⁻¹ d₃₄-hexadecane was added to each fraction.

**BBR₃ treatment (hardly releasable fraction)** Five milliliters of a 1.0 M boron tribromide solution in dichloromethane was added to approximately 150 mg of pre-extracted or pre-extracted/saponified dry samples. The extraction procedure followed the procedure for the extractable fraction. After 24 h of stirring at room temperature, a second ultrasonication step was added. Two milliliters of diethyl ether was added, and the supernatant liquid was decanted and filtered using a Whatman glass fibre filter (0.7 μm pore diameter). The solid residue was washed twice with diethyl ether, and the combined organic layers were washed twice with 5 mL of pre-extracted water and dried with anhydrous granulated sodium sulphate. After volume reduction, the crude extracts were separated into three fractions by column chromatography (Baker, 2 g silica gel 40 m) using the following eluent mixtures: fraction: 1-n-pentane/dichloromethane (95/5 v/v), fraction 2: dichloromethane and fraction 3: methanol. Prior to analysis, the extracts were fortified with the same internal standard solution as for the hydrolysis fraction.

**RuO₄ oxidation (hardly releasable fraction)** A mixture of 8 mL of tetrachloromethane, 8 mL of acetone and 1 mL of pre-extracted water was added to aliquots of pre-extracted or pre-extracted/saponified dry samples and additionally fortified with 5000 mg of sodium periodate and extracted or pre-extracted water was added to aliquots of pre-extracted water and dried with anhydrous granulated sodium sulphate. After volume reduction, the crude extracts were washed twice with 5 mL of pre-extracted water as well as the extracts were saturated with water of the same temperature and subsequently washed with diethyl ether. All organic solutions were combined, dried by means of anhydrous sodium sulphate and reduced to 0.5 mL. The fractionation was realized on the silica gel microcolumns into 2 fractions with fraction 1: dichloromethane and fraction 2: diethyl ether/methanol (40/60 v/v). Prior to GC-MS analysis, the extracts were fortified with the same internal standard solution as for the BBr₃-treatment.

### 2.5 Analysis by GC-MS

Gas chromatographic mass spectrometric measurements were performed on a Carlo Erba Mega Series GC 5160 with a 30 m × 0.25 mm × 0.25 μm capillary column linked to a Thermoquest Trace Quadrupole-MS. Carrier gas was helium (purity 99.999%) set to a velocity of 36 cm s⁻¹. The oven temperature was held at 60 °C for 3 min, then programmed at 5 °C min⁻¹ to 180 °C and then programmed at 3 °C min⁻¹ to 310 °C and held for 15 min. A 1 μL hot splitless injection at 270 °C was applied with a splitless time of 60 s. The mass spectrometer operated in EI⁺ full scan mode (70 eV) scanning from 35 to 740 m/z with a scan time of 0.5 s. Source temperature was set to 250 °C and interface temperature to 200 °C.

Identification of compounds was based on comparison of gas chromatographic and mass spectral properties of detected substances with those of authentic reference material. The degradation of DDX compounds was checked routinely to be less than 15% in the injector (US-EPA recommendation). The limit of detection (LOD) was approx. 0.3 ng g⁻¹ with a limit of quantification (LOQ) of 1 ng g⁻¹ d.w. The ion traces recorded are given in the supporting information (see Table SI-1).

### 3 Results and discussion

#### 3.1 Sediment mapping and raster sampling

In principle, it was obvious in each of the three profiles that in the area of the uppermost meter, the specific electrical resistances are more or less significantly reduced compared with the deeper subsurface. Whereas at depths of more than 1 m below the sediment surface specific electrical resistances of more than 65 Ω m⁻¹, mostly more than 95 Ω m⁻¹ were measured, corresponding with the sandy base of the canal. The resistances in the uppermost meter of the sediment layer were mainly at values < 65 Ω m⁻¹, often even below 40 Ω m⁻¹. The specific electrical resistance of the water was 21.3 Ω m⁻¹ at the time of measurement.

Based on the assumption that the sediment in a first approximation is saturated with water of the same temperature and electrical conductivity, the resistances observed in the sediment can be interpreted as follows:

- Areas with an electrical resistivities of more than 65 Ω m⁻¹ are likely to be predominantly affected by sands in natural storage and normal compaction.
- Areas with electrical resistivities below 65 Ω m⁻¹ are categorized as sandy mud with an elevated organic matter and clay mineral content.

The combination of the above two facts points to the conclusion that the near-surface sediment layers were weakly consolidated sediments exhibiting a predominantly sandy matrix with an increased porosity, washing of fine grain into the pore water and an increased content of organic substances in the pore space and the matrix. They were identified with a largest thickness of approximately 1 m at the western end of the profiles and an average thickness of 0.4 m.
Based upon these data, sediments with an electrical resistance below 35 $\Omega\text{m}^{-1}$ were selected for the sampling of the cores, which are predominantly located in the eastern part of the investigation area, mostly on the northern shore area (see Fig. 4). TOC contents < 0.01% were not included in the selection. The lithology is presented in Table 1.

The length of the sediment cores (see Table 1) and extrapolation of the dry weight acted as a basis for the calculation of the total sediment mass and the amount of incorporated DDA and relevant DDT metabolites. Calculation of the total sediment body was performed with the two different sediment body geometries, resulting in a total dry sediment mass of 1035 t with the trapezoid approach and 4820 t with the column approach. Details of each core and layer are given in the supporting information (see Table SI-2).

### 3.2 Quantification of particle-associated DDT metabolites

The sequential chemical analysis of the freeze-dried sediment samples (including pore water) revealed a high variety of DDT metabolites that were determined in both non-bound and bound fractions. Noteworthy, the majority of DDT derivatives represents precursor metabolites of DDA (see Fig. 1). The mean content of DDX (as sum of all DDT metabolites) was analysed for the sedimentation area of the canal, subdivided into A–D sections away from the source in area B (see Fig. 4). No differentiation into o,p´- and p,p´-DDX was applied. All basic data are given in the supporting information (see Table SI-3).

The DDT metabolites are divided into freely available (EF), easily (ER) and hardly releasable fractions (HR, according to Fig. 3). The individual concentrations in the different samples varied by a factor of up to more than 20,000 in the EF fraction and up to more than 1,000,000 in the ER fraction within one layer and up to more than 200,000 and 30,000,000 within one area, pointing to a high heterogeneity. This must be considered for all quantifications.

Comparing the four areas, area C was comprised by the highest DDX contents and the highest variation of compounds content (see Figs. 5 and 6, and Table SI-3). In the following, the occurrence in the extractable and non-extractable DDT metabolites is discussed separately.

#### 3.2.1 Extractable fraction

In almost every sample, DDX were detected, except for DDT, DDE, DDEt, DDA, DDM and DBP in some samples (see Table SI-4). Especially, the parent compound DDT ranged from not detected, through low concentrations about a few hundred nanograms per gram up to 15,000 ng g$^{-1}$ in the middle layer close to the source, in accordance with earlier investigations (Schwarzbauer et al. 2003). Downstream from the source, the concentrations mostly decrease. The freely available contaminants were dominated by DDD (183,500 ng g$^{-1}$) and DDMS (52,000 ng g$^{-1}$; see Table SI-3). DDA was only detected in low concentrations close to the source and not detected in the areas C and D. This observation follows strictly the anaerobic metabolic pathway of DDT as described in Fig. 1.

A comparison of the sum DDX in the top, middle and base layer of the areas A–D is given in Fig. 5. The highest concentrations were measured in the base layer of area A, close to the source region. Figure 5 comprises the min and max values and the median on a logarithmic scale. Unfortunately, no top sample of area B was gained due to sandy material covering the contaminated layers.

#### 3.2.2 Easy releasable fraction

The metabolite composition of the free available fractions differs in comparison to the easily releasable fraction obtained by alkaline hydrolysis. Additionally, the composition of the metabolites is highly different in comparison to the free available fraction.

Main metabolites received from hydrolysis reaction are DDA, DDNU, DBP and DDM. In contrast to the free the source-inflow into the Teltow Canal and the red dots are 500 m indicators. Most of the contaminated sediment was located on the northern riverbank due to the low flow velocity of the canal.
available fraction, these metabolites of the non-extractable residues are dominated by DDA with a mean maximum of 17,000 μg g⁻¹ (see Table SI-4). The sample from area C (middle layer) was characterized by maximum concentrations of DDA (see Table SI-4).

A comparison of the sum DDX in the ER in the top, middle and base layer of the areas A–D is given in Fig. 6. The highest concentrations are measured in area C, close to the source region. The regular flow direction is along the areas A to D. Analogous to Fig. 5, the min and max values and the median values are presented on a logarithmic scale.

The existence of DDT and metabolites in the investigation area can be explained by the highly reducing environmental conditions (Taylor et al. 2019), for which e.g. a half-life time of up to 65 years has been calculated (Clow et al. 2017). In addition, the decreasing degradation rate due to lesser bioavailability via formation of NER needs to be considered. The contribution of the partially aerobic metabolites to the sum of DDX is in the range of 2–3% for DDE and up to 28% in maximum for DDMU in the ER fraction.

As the compounds were introduced as a chemical cocktail, partially with solvent contents in the lower % degree, the mixture may be preserved. Unfortunately, no information about the emitted number of compounds is available.

For comparison with earlier studies in this investigation area, the date of sampling and the different depths of the sediment cores have to be considered (Schwarzbauer et al. 2003; Heim et al. 2005). In the study presented here, 12 sediment cores represent a wider range of data within the regarded sampling location.

In comparison with reported concentration ranges in other regions, the canal sediments are highly contaminated, partially above those reported for Superfund Sites in the USA (e.g. Garrison et al. 2014; Taylor et al. 2019).

The available Sediment Quality Guidelines (SQG) for freshwater sediments are highly exceeded. Available criteria for freshwater sediments for the preloaded Elbe River System indicate target values of <5 ng g⁻¹ d.w. for p,p'-DDT/DDD/DDE, respectively. The intervention value as the worst case is a factor of 10 higher (Ahlf et al. 2002).
The Canadian SQG states probable effects above 4.77 ng g\(^{-1}\) for p,p\(^{\prime}\)-DDT and 8.51 ng g\(^{-1}\) and 6.75 ng g\(^{-1}\) d.w. for p,p\(^{\prime}\)-DDD and DDE, respectively (Canadian Council of Ministers of the Environment 2002). MacDonald et al. (2000) published a consensus-based SQG TEC for total DDTs (sum of p,p\(^{\prime}\)-DDT/DDD/DDE) of 5.28 ng g\(^{-1}\) d.w. There are no specific sediment criteria defined and no EQS is available within the WFD. Comparing the top, middle and base sediment layers for each area, the middle layer in area C reveals the highest contents followed by the base layers, illustrating the complex system of the sediment body (see Figs. 5 and 6). The reason for these varying concentrations in ER may be the different degradation procedure applied and the different sediment type and sampling period, keeping in mind that the formerly cited samples were taken in 1997/1999, potentially consisting of sediment of an undisturbed system in contrast to the sediment sampled in 2007.

The hardly releasable fraction (HR) is obtained by degradation of the pre-hydrolysed sediment with BBr\(_3\) and oxidation by K\(_2\)MnO\(_4\) or RuO\(_4\) (steps 3 and 4; see Fig. 3). Cleavage of ether bonds and covalent carbon bond is applied by these operations. The hardly releasable metabolites represent a long-term contamination potential for DDA due to the strong incorporation into the particulate matter or organic matter in the sediment. These NER are not addressed in this paper. Results obtained from BBr\(_3\) treatment are lower than in Schwarzbauer et al. (2003). Comparable data for DDA were reported in Schwarzbauer et al. (2003) with a factor 10 higher.

The presented and applied method was compared with newer approaches like pyrolysis-GC-MS and TMAH-chemolysis GC-MS (Kronimus et al. 2006; Zhu et al. 2019). Considering potential artefacts, the combination serves as a tool for the assessment of a contamination.

In natural ecosystems, the prevailing sorption of these compounds with a low water solubility occurs on/in fine-grained SOM and clay minerals usually coated with humic matter, leading to the formation of NER. Therefore, the accumulation in sediments and organisms leads routinely to the application of a normalisation to the content of total organic carbons (% TOC), independent of the source, quality and degree of degradation. Often, the log \(K_{oc}\) is calculated. There was no correlation observed for DDx content and TOC as well as for the contents in the extractable fraction (EF) with the easily releasable fraction (ER). Neither a correlation with the sum of DDx was visible (data not shown).

The metabolism of DDT is proceeded, a long-standing pollution is attested. The semi-final metabolites DDA, DBP, DDCN and DDM are available in noticeable amounts, especially DDA.

### 3.3 Calculation of the DDA potential

As already presented, the DDA contamination potential was calculated using data from precursor metabolites that can be converted to DDA over time and the DDA concentrations themselves. These values have been considered in both the free available and the easily releasable DDA fractions. Hardly releasable proportions have been neglected due to their low potential of a short-time remobilisation.

Further on, a risk for long-term contamination is the existence of the high concentrations of precursor metabolites which can form DDA by metabolism. The very rough estimation is based upon the mean concentrations of each layer. For the base layer in area A, only one concentration was analysed, the data are extrapolated to the other two layers. The estimated DDA release potential shows a wide range of values, given in the brackets as minimum and maximum values.

#### 3.3.1 Calculation of the maximum free available amounts of DDA and precursor metabolites

Data in Table SI-3 were applied to calculate the directly available amounts for the formation of DDA via the precursor metabolites DDT to DDEt. DDA itself was added to the total formation and the final metabolites DDCN to DBP are included in the sum DDx. These were calculated separately obtained by sediment extraction (step 1). They are delivering the percentage of precursors and the direct available percentage of DDA releasable.

The span of the precursors ranges from 88 to 100% in the EF fraction (see Table SI-4) with a slightly higher formation potential in the middle and deeper layers, considering no data are available for the top layer of area B.

The amounts of calculated maximum DDA contamination are extrapolated by multiplying the maximum concentrations obtained for each sediment area of each layer. The amounts of the labile-bond fraction are directly available (in the case of DDA) or can be released after microbial degradation of precursor metabolites (DDT to DDEt) to DDA.

Summarizing all concentrations up to DDA for all layers and areas, applying the trapezoid model a concentration of 48 kg (32–82 kg) results for this fraction. The application of the column model resulted in 278 kg (214–354 kg) DDA absolute (see Tables SI-3 and SI-5).

#### 3.3.2 Calculation of the maximum easily releasable amounts of DDA and precursor metabolites

Applying the scheme proposed by Zhu et al. (2019), the highest variation occurred in the middle layer of area C with a factor of more than 10,000 (see Table SI-4). The scheme differentiates the processes into dehalogenation, functional groups formation and loss of one carbon atom for the potential
precursors down to DDNU. The dehalogenation formation of DDA and metabolites from DDA are included. The release potential is estimated via the progressive release potential and the subsequent formation of metabolites.

The fractions of easily releasable amounts were obtained by basic hydrolysis (step 2). The data of both steps were added for representing the total easily releasable amounts. Especially, in the middle layer of area C, the sum of all precursors plus DDA results in 16,700 μg g⁻¹ of the total 16,800 μg g⁻¹, equal to 99.5% (see Table SI-5).

The highest DDA release potential is calculated for the middle layer of area C. Except for the top layers of the areas, A and D, the data show a release potential between 65 and almost 100% (see Table SI-4).

### 3.3.3 Calculation of the total content and the relative contribution to the DDA contamination range

Applying the two geometric models for the sediment layers (described in Section 3.1), an extrapolation of the data to the total sediment of 1035 t (trapezoid model) and 4820 t (column model) was realized, being aware of the rough calculation approach. The trapezoid model resulted in 48 kg DDA formation via precursors for the extractable fraction (EF) and 13.7 kg via the easy releasable fraction (ER). The direct DDA release potential is 0.4 kg for the EF and 1346 kg for the ER (see Table 2), considering the mean contents. The column model resulted in 278 kg DDA formation via precursors for the extractable fraction (EF) and 47.0 kg via the easy releasable fraction (ER). The direct DDA release potential is 0.74 kg for the EF and 9800 kg for the ER fraction, respectively, considering the mean contents.

The relative contribution of precursors is relatively high in areas A and D at the top layer and decreases to almost 1% in the middle and base layer in the extractable fraction. The calculation excludes the potential DDA degradation products DDCN, DDM and DBP. In the easy releasable fraction, the contribution of DDA is in the middle and base layers around 99%, except for the top layers in areas A and D. The highest total contribution is given by the middle layer C with 75%, depending on the model applied.

| Table 2 | Overview of the sum of DDA precursors and DDA direct release content for the areas A–D referring to the applied trapezoid and column model (kg layer⁻¹) |
|----------------|---------------------------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|
| **Trapezoid model** | | | | | | | | | | | | | | | |
| **Area** | **EF** | **ER** | **Area** | **EF** | **ER** | **Area** | **EF** | **ER** | **Area** | **EF** | **ER** | **Area** | **EF** | **ER** |
| A | Top | Top | B | Top | Top | C | Top | Top | D | Top | Top |
| Sum DDA precursor | 3.8 | 1.8 | No sample | No sample | 1.9 | 1.3 | 5.1 | 6.1 |
| DDA direct | 0.0 | 0.4 | No sample | No sample | 0.0 | 122.9 | 0.0 | 2.2 |
| Middle | Middle | Middle | Middle | Middle | Middle | Middle | Middle | Middle |
| Sum DDA precursor | 0.8 | 0.1 | 0.8 | 0.1 | 3.0 | 2.3 | 3.7 | 1.6 |
| DDA direct | 0.0 | 0.9 | 0.0 | 1.6 | 0.0 | 984.7 | 0.4 | 88.6 |
| Base | Base | Base | Base | Base | Base | Base | Base | Base |
| Sum DDA precursor | 22.4 | 0.0 | 0.1 | 0.0 | 3.7 | 0.4 | 2.8 | 0.1 |
| DDA direct | 0.0 | 5.6 | 0.0 | 5.9 | 0.0 | 130.4 | 0.0 | 2.7 |
| Sum DDA precursor | 48.1 | 13.7 |
| DDA direct | 0.4 | 1345.9 |
| **Column model** | | | | | | | | | | | | | | | |
| **Area** | **EF** | **ER** | **Area** | **EF** | **ER** | **Area** | **EF** | **ER** | **Area** | **EF** | **ER** | **Area** | **EF** | **ER** |
| A | Top | Top | B | Top | Top | C | Top | Top | D | Top | Top |
| Sum DDA precursor | 9.2 | 2.4 | No sample | No sample | 14.5 | 10.1 | 7.9 | 9.3 |
| DDA direct | 0.0 | 0.5 | No sample | No sample | 0.0 | 940.7 | 0.0 | 3.3 |
| Middle | Middle | Middle | Middle | Middle | Middle | Middle | Middle | Middle |
| Sum DDA precursor | 0.9 | 0.1 | 5.7 | 0.9 | 23.1 | 17.7 | 5.7 | 2.4 |
| DDA direct | 0.0 | 1.0 | 0.0 | 11.7 | 0.0 | 7534.6 | 0.6 | 136.4 |
| Base | Base | Base | Base | Base | Base | Base | Base | Base |
| Sum DDA precursor | 151.3 | 0.0 | 0.9 | 0.0 | 32.9 | 3.4 | 24.9 | 0.6 |
| DDA direct | 0.0 | 37.7 | 0.0 | 52.2 | 0.0 | 1166.9 | 0.1 | 24.7 |
| Sum DDA precursor | 276.9 | 47.0 |
| DDA direct | 0.7 | 9909.7 |
DBP to the sum of DDX is given in Table SI-5 and illustrated in Fig. 7 as difference S-DDX.

The directly available DDA contamination is very low. For the EF fractions, the precursors dominated the contribution. For the ER fraction, the direct available DDA concentration exceeds the formation potential via precursors, except an increased precursor contribution for the top layers in areas A and D.

The main amount of the DDA range is expected by remobilisation of easy-bound amounts through degradation of the SOM or by microbial transformation of free available amounts of precursor metabolites. The easily available DDT metabolites in the ER fraction represent a formation potential of DDA after remobilisation and transformation.

Important is the consideration that the easy releasable fraction can form DDA by degradation processes, providing a huge risk for the aquatic environment. The degradation kinetics are not yet studied in detail; a slow formation of DDA and a subsequent slow further metabolization of DDA is evident. There is almost no contribution of the EF fraction, except for middle layer of area A with a contribution of 22.4 kg layer$^{-1}$ and
150 kg layer\(^{-1}\), for the trapezoid and column model, respectively.

Summarizing the calculations for the different models for the investigation area, the trapezoid model resulted in 48 kg DDA formation potential in the EF fraction (32–82 kg) and 1360 kg in the ER fraction (11–4010 kg). By applying the column model, a 278 kg DDA formation potential in the EF fraction (214–354 kg) and 9800 kg in the ER fraction (250–29,000 kg) is ascertained. In both cases, the middle layer of area C contributed around 75% to the total release potential.

### 3.4 Preliminary remobilisation experiments

Based upon earlier investigations on the sediment contamination, preliminary lab-experiments with real sediment samples were performed to simulate the short-time resuspension within the water body. Especially, the influence of the professional shipping results in dynamic conditions with a potential of resuspension of the upper layers and accordingly building up of a concentration gradient out of the sediment pore water. Heim et al. (2005) published data on the upper most layers of a sediment core clearly illustrating a diffusion gradient out of the sediment into the overlaying water column. Dredging is another opportunity to remobilise sediment layers, only applied exclusively for navigability. The results of the experiments reveal information about the transport of DDA between different environmental compartments. The transport of sedimentary DDA via pore water into the surface water and the transport from water through the soil into the groundwater area are the decisive factors for a possible contamination of the infiltration area of the drinking water plant.

#### 3.4.1 Release experiment

Initial laboratory release experiments, simulating the remobilisation/degradation of extractable and non-extractable (NER) residues and the land disposal after dredging in aquaria resulted in a release due to aerobic redox conditions and no differences upon land disposal. Especially, the water phase was rapidly depleted of DDA in the remobilisation scenario with a subsequent adsorption and incorporation into the SPM of the ecosystem.

Under both redox conditions, the maximum of the DDA concentration was released after the first day (93 and 98%), with no marked differences between oxic and anoxic conditions, indicating that under dynamic conditions, DDA is released by hydrolysis processes (cleavage of the ester bond) and uncoiling of the SOM, independently of the redox potentials (see Fig. 8). An aliquot of the experimental sediment was analysed for DDA, resulting in 24 ng g\(^{-1}\) in the EF fraction and 40,000 ng g\(^{-1}\) in the ER fraction (TOC 8.65%). The pore water concentrations were analysed in the range of 160 μg L\(^{-1}\). The released DDA concentrations are given in Table 3.

### 4 Conclusions

The approach offers a tool for the analysis of recent and older contaminations of sediments, including the (bio) available fraction and different fractions from easily to hardly releasable under different environmental conditions.

The different composition of the bound DDT-related compounds shows a high amount of DDA in the hydrolysable (“easily releasable” ER) fraction, in accordance with earlier results (Schwarzbauer et al. 2003). The degradative sediment treatment proves the binding of DDA to the sediment by ester bonds and potential release by hydrolysis reactions.

Summarizing the calculations for the different models within the 1.25 km long investigation area, the trapezoid model resulted in 48 kg DDA formation potential in the EF fraction and 1360 kg in the ER fraction. By applying the column model, a 278 kg DDA formation potential in the EF fraction and 9800 kg in the ER fraction is determined. In both cases, the middle layer of area C contributed around 75% to the total release potential.

The applied degradation scheme allows the determination of the directly releasable contents as well as the determination of the

### Table 3 Experimental DDA release from sediment

| Sampling (days) | Oxic treatment DDA (ng g\(^{-1}\)) | Anoxic treatment DDA (ng g\(^{-1}\)) |
|----------------|----------------------------------|----------------------------------|
| 1              | 8.938                            | 11.442                           |
| 7              | 385                              | 31                               |
| 42             | 254                              | 125                              |

![Fig. 8 Remobilisation of DDA from sediment under oxic and anoxic conditions](image-url)
contents that lead to the contaminants by metabolism. Especially, those compounds which tend easily to metabolism (such as p,p’-DDT and p,p’-DMDT) are subjected to degradation and entrapment in the SOM. A further linking via covalent bonds is possible and evaluable with the approach.

The results of the release experiment offer a tool for the first estimation of the environmental behaviour of DDA. The high release behaviour of contaminated sediments for DDA shows that the contamination of water is to be expected, so that DDA must be regarded as a water pollutant.

In the case of dynamic conditions, e.g. while dredging work or by commercial shipping, DDA would be released immediately and in high amounts. Very recent sediment sampling in area B (November 2018) revealed a comparable sandy layer relative to the sampling in 2007. There is no monitoring program performed within the affected canal area water body and sediment.

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