

Uptake and translocation of organophosphates and other emerging contaminants in food and forage crops

Trine Eggen · Eldbjørg S. Heimstad ·
Arne O. Stuanes · Hans Ragnar Norli

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Abstract Emerging contaminants in wastewater and sewage sludge spread on agricultural soil can be transferred to the human food web directly by uptake into food crops or indirectly following uptake into forage crops. This study determined uptake and translocation of the organophosphates tris(1-chloro-2-propyl) phosphate (TCPP) ($\log K_{ow}$ 2.59), triethyl-chloro-phosphate (TCEP) ($\log K_{ow}$ 1.44), tributyl phosphate (TBP) ($\log K_{ow}$ 4.0), the insect repellent *N,N*-diethyl toluamide (DEET) ($\log K_{ow}$ 2.18) and the plasticiser *N*-butyl benzenesulfonamide (NBBS) ($\log K_{ow}$ 2.31) in barley, wheat, oilseed rape, meadow fescue and four cultivars of carrot. All species were grown in pots of agricultural soil, freshly amended contaminants in the range of

0.6–1.0 mg/kg dry weight, in the greenhouse. The bioconcentration factors for root (RCF), leaf (LCF) and seed (SCF) were calculated as plant concentration in root, leaf or seed over measured initial soil concentration, both in dry weight. The chlorinated flame retardants (TCEP and TCPP) displayed the highest bioconcentration factors for leaf and seed but did not show the same pattern for all crop species tested. For TCEP, which has been phased out due to toxicity but is still found in sewage sludge and wastewater, LCF was 3.9 in meadow fescue and 42.3 in carrot. For TCPP, which has replaced TCEP in many products and also occurs in higher residual levels in sewage sludge and wastewater, LCF was high for meadow fescue and carrot (25.9 and 17.5, respectively). For the four cultivars of carrot tested, the RCF range for TCPP and TCEP was 10–20 and 1.7–4.6, respectively. TCPP was detected in all three types of seeds tested (SCF, 0.015–0.110). Despite that DEET and NBBS have $\log K_{ow}$ in same range as TCPP and TCEP, generally lower bioconcentration factors were measured. Based on the high translocation of TCPP and TCEP to leaves, especially TCPP, into meadow fescue (a forage crop for livestock animals), ongoing risk assessments should be conducted to investigate the potential effects of these compounds in the food web.

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T. Eggen (✉)
Bioforsk, Norwegian Institute for Agricultural
and Environmental Research, Postveien 213,
4353 Klepp St., Norway
e-mail: Trine.Eggen@bioforsk.no

E. S. Heimstad
Norwegian Institute of Air Research (NILU),
Hjalmar Johansens gate 14,
9296 Tromsø, Norway

A. O. Stuanes
Norwegian University of Life Sciences, Box 5003,
1432 Ås, Norway

H. R. Norli
Bioforsk, Norwegian Institute for Agricultural
and Environmental Research, Høgskoleringen 7,
1432 Ås, Norway

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Introduction

Food safety is an important global issue receiving high priority worldwide. Transfer of contaminants from soil, water and air to the food chain is one aspect of food safety, and identification of sources, transfer pathways and

environmental residue levels of emerging contaminants are attracting great attention. For instance, pharmaceuticals, musk compounds and organophosphates up to micrograms per liter or milligrams per kilogram dry weight (dw) have been found in water or sewage sludge (Calderón-Preciado et al. 2011; Lee et al. 2010; Muñoz et al. 2009; Reemtsma et al. 2006). Residue levels in waste water and sewage sludge for selected emerging contaminants studied in the present work is summarized in Table 1 (Glassmeyer et al. 2005; Green et al. 2008; Huppert et al. 1998; Leonards et al. 2011; Marklund et al. 2005; Nakada et al. 2006; Terzic et al. 2008).

Contaminants can be transported to soil via several routes. Manure and sewage sludge are used as fertilisers and soil conditioner on agricultural soils while effluent from wastewater treatment plants (WWTPs) is used for irrigation. In addition, manure and sewage are used to manufacture commercial mature compost and soil-based growing mediums, which are commonly used in domestic gardens. Sewage sludge and wastewater are known to contain a large mixture of different legacy and emerging contaminants including high-volume human pharmaceuticals and personal care additives, e.g. review by Harrison et al. (2006) and detected in different screening projects (Calderón-Preciado et al. 2011; Díaz-Cruz et al. 2009; Duarte-Davidson and Jones 1996; Kolpin et al. 2002; Muñoz et al. 2009), while manure can contain residues of veterinary pharmaceuticals from medication of livestock animals (Campagnolol et al. 2002; Furtula et al. 2010; Kolpin et al. 2002; Zhao et al. 2010). Examples of such compounds which also are found to be taken up by plants are galaxolide, tonalide, triclosane, enrofloxacin, carbamazepine, metformin and trimethoprim (Boxall et al. 2006; Eggen and Lillo 2012; Macherius et al. 2012; Migliore et al. 2003).

Emerging contaminants are new substances found or expected to be found in the environment and which may have potential toxic effects but yet not regulated due to lack of persistent, toxicity and bioaccumulation data. Many of these compounds are additives used widely in everyday industrial and household products, such as flame retardants for textiles and other products, surface-active substances used as detergents or water and oil repellent products, fragrances used in hygienic, cosmetic and cleaning products and plasticisers used in products such as toys and food containers (Eriksson et al. 2003; Goldman 1998; Marklund et al. 2003; Slack et al. 2005). Emerging contaminants cover a wide range of properties, and unlike many legacy organic hydrophobic contaminants (e.g. persistent organic pollutants), many of these new compounds tend to be more polar and water-soluble but are still persistent in the environment.

Chemical substances recognised as an environmental or human threat are phased out and replaced with less hazardous substances. For instance, triethyl-chloro-phosphate

(TECP) has been phased out in Europe (Andresen et al. 2004) due to its toxicity (European Commission 2009; WHO World Health Organization 1998). However, tris(1-chloro-2-propyl) phosphate (TCPP), which has replaced TCEP in many products (Quednow and Püttmann 2009), is also considered to be potentially carcinogenic and is undergoing a health and environmental risk assessment (European Commission 2008). In addition, the detergent tributyl phosphate (TBP), the insect repellent *N,N*-diethyl toluamide (DEET), which is widely used in consumer products such as anti-mosquito agents or certain types of sportswear, and the plasticiser *N*-butyl benzenesulfonamide (NBBS) are all emerging contaminants that are being evaluated for their potential environmental and human health risks (Aronson et al. 2011; OECD April 2001; Strong et al. 1991).

Transport of water and solutes, including contaminants, from soil via plant roots to aboveground compartments is driven by the water potential gradient created by plant transpiration (McFarlane 1995). It has been shown that many of the legacy and less hydrophilic organic pollutants, e.g. polychlorinated biphenyls, dichlorodiphenyltrichloroethane and its metabolites, polyaromatic hydrocarbons and dioxins, can be taken up from the soil via roots (Inui et al. 2008b; White 2010; Whitfield-Åslund et al. 2008; Zohair et al. 2006). However, except for some plant species-dependent difference for instance for *Vivica cracca* (Ficko et al. 2010) and certain varieties of *Cucurbita pepo ssp* (pumpkin and zucchini) (White 2010; Whitfield-Åslund et al. 2008), uptake of these compounds via roots is general low. Due to their higher polarity, the emerging compounds might have a greater capability to be taken up by plant roots and further translocated within plants. However, knowledge of if and how they transfer into the terrestrial food web is still scarce. Many polar emerging contaminants have a high potential to pass through treatment processes commonly used for landfill leachates or in WWTPs and can thus be detected in effluents and the environment (Nakada et al. 2010). Thus, more knowledge of environment–food web transfer of such compounds is important.

In recent decades, a number of plant uptake models ranging in scope from simple steady-state equations with one input parameter to compartment models containing several dynamic uptake, intra-plant processes and input parameters have been established to predict uptake of compounds (Briggs et al. 1982; Chiou et al. 2001; Dettenmaier et al. 2009; Rein et al. 2011; Ryan et al. 1988; Trapp 2000). However, in order to verify or adjust existing uptake models for emerging contaminants, experimental or controlled field data are needed.

The main objective of the present work was to compare uptake and translocation of selected polar and semipolar emerging organic contaminants with different

Table 1 Reported residual levels in wastewater $\mu\text{g/L}$ and sewage sludge (mg/kg dw) of the selected emerging contaminants investigated in the present study: average or median values (depended on what is given), min and max values in brackets

TBP	TCPP		TCEP		NBBS		DEET
	Sludge mg/kg	Wastewater $\mu\text{g/L}$	Sludge mg/kg	Wastewater $\mu\text{g/L}$	Sludge mg/kg	Wastewater ^p $\mu\text{g/L}$	Wastewater ^p $\mu\text{g/L}$
13 (6.6–52) ^{a,d}	0.094 (0.026–0.350) ^e	2.5(1.1–18.0) ^{a, d}	2.58 (0.56–7.20) ^e	0.42 (0.09–1.0) ^{a, d}	1.28 (0.030–0.276) ^e	1.35 (0.3–2.2) ^{a,e}	(0.67–0.89) ^{a,i}
0.41(0–0.316) ^{b, c}	0.28 (0.0096–0.85) ^d	0.60 (0.01–1.16) ^{b, c}	0.87 (0.06–1.90) ^d	0.62 (0.004–0.27) ^{b, c}	0.035 (0.001–0.110) ^d	0.82 (0.24–1.7) ^{b, e}	(0.90–1.02) ^{b, i}
2.7 (0.36–6.1) ^{b, d}	1.7–2.22 ^{b,f}	1.7–2.22 ^{b,f}		0.47 (0.35–0.89) ^{b,d}			0.84 (max 6.9) ^{b,g}
	2.0 (1.5–24.0) ^{b, d}	2.0 (1.5–24.0) ^{b, d}		0.19 (max 0.5) ^{b, g}			0.18 (max 2.1) ^{b,h}
	0.46 (max 2.5) ^{b, g}	0.46 (max 2.5) ^{b, g}					

^a Inlet^b Outlet^c KLIF 2011^d Marklund 2005^e Huppert 1998^f Green 2008^g Terzic 2008^h Glassmeyer 2005ⁱ Nakada 2006^j No data found for sludge

structures and properties in an experimental growth study using different high-volume and agriculturally important crop plants. This is important knowledge related to human health risk assessments where transfer of contaminants from soil to edible plant compartments is included. Verification of the experimental data in dynamic plant uptake models was performed in a separate study (Trapp and Eggen, in press *Environmental Science and Pollution Research*). The crop plants included in the study were cereals (barley, *Hordeum vulgare*; wheat, *Triticum aestivum*), a grass forage (meadow fescue, *Festuca pratense*), oily rape seed (*Brassica rapa*) and root vegetable (carrot, *Daucus carota*). The emerging organic contaminants analysed were: the flame retardants TCPP and TECP, the detergent TBP, the plasticiser NBBS, and the insect repellent DEET. A summary of the main structure and chemical and physical properties of these five substances at relevant pH is presented in Table 2. In order to compare uptake and translocation of contaminants in different plant species and organs, it is necessary to apply soil concentration which is analytical

measurable. Thus, clean soil with artificial added contaminants was used as experimental approach.

Materials and methods

Plant uptake experiment

The study was a greenhouse pot experiment conducted at Bioforsk Vest Særheim between December 2008 and April 2009. A detailed description of the experimental procedure is given elsewhere (Eggen et al. 2011), and it is only briefly summarised below.

Soil characterisation

Loamy sand soil from an agricultural field in West Norway was sieved (<4 mm) and mixed with a controlled-release fertiliser (3 g/kg soil, Multicote 4, (N/K/P) 15:7:15 (2+) TE, Haifa Chemicals Ltd.) using a cement mixer for approximately 10 min. The soil had 0.7 g kg⁻¹ total organic carbon,

Table 2 Selected physico-chemical properties of the test compounds used in the present study

Test compound and application	Structure (at pH 5.5)	CAS-no.	MW	log K _{ow}	Half-life in soil (d)	Henry's Law constant (atm·m ³ ·mole ⁻¹)	Water solubility (mg/L)	Polarisability ^d (Å ³)	Electron Affinity ^d (eV)	Electro-negativity ^d (eV)
Tris(2-Chloroethyl) Phosphate (TCEP). Flame retardant		115-96-8	285.5	1.44 exp ^a	120 est ^b	3.29E-06 est 25°C ^a	7000 exp 25°C ^a	21.72	1.992	6.494
Tris(1-chloro-2-propyl) phosphate (TCPP). Flame retardant		13674-84-5	327.6	2.59 exp ^a	120 est ^b	5.96E-08 est 25°C ^a	1200 exp 25°C ^a	27.293	1.916	6.394
Tributyl phosphate (TBP). Detergent		126-73-8	266.3	4.00 exp ^a	17 est ^b	1.41E-06 est 25°C ^a	280 exp 25°C ^a	26.836	1.242	5.805
N,N-diethyl toluamide (DEET). Insect repellent		134-62-3	191	2.18 exp ^{a,c}	75 est ^b	2.08E-08 est 25°C ^a	912 est 25°C ^a	23.843	0.157	4.63
N-butyl benzenesulfonamide (NBBS). Plasticiser		3622-84-2	213.3	2.31 est ^{b,c}	30 est ^b	2.17E-06 est 25°C ^b	398 est 25°C ^b	21.199	0.763	5.451

^a Exp=experimental data and est=estimated data from ChemIDPlus Advanced <http://chem.sis.nlm.nih.gov/chemidplus/chemidheavy.jsp>

^b Estimated values calculated by EpiSuite 4.xr SRC Interactive PhysProp Database Demo <http://www.syrres.com/what-we-do/databaseforms.aspx?id=386>

^c Compounds are dissociable but at the relevant pH range they exist as neutral compounds

^d Calculated properties with the software Cache, Fujitsu Limited

pH 6.0 ($v/v=1:2.2$) and cation exchange capacity of $46.6 \text{ mmol}_c \text{ kg}^{-1}$.

Test compounds

The selected test substances were TCPP, TECP, TBP, DEET and NBBS. All organophosphates were supplied by Chiron AS, Trondheim, Norway, and DEET and NBBS by Sigma-Aldrich, Norway, with purity better than 98 % for all compounds.

Spiking procedure

Stock solutions of the test compounds (40 mg/mL) were dissolved in acetone (approximately 5–10 mL). One millilitre stock solution was then diluted in 50 mL distilled water (all test compounds added together), after which it mixed thoroughly by hand with 4.0 kg dw soil and added to each 4-L pot. Each pot was prepared separately. The nominal estimated concentration of test compound in each pot was 1 mg/kg. Soil samples were taken directly after spiking and stored at 4 °C, approximately 3 weeks, until analysis to determine the actual initial soil concentration.

Selected plants and growth conditions

The selected plant species are all important crop or forage plants: barley (*H. vulgare* cv. Edel) (root, leaf, grain); wheat (*T. aestivum* cv. Bjarne) (grain); and meadow fescue (*F. pratense* cv. Fure) (root, leaf); oilseed rape (*B. rapa* cv. Valo) (seed); and four carrot cultivars (*D. carota* ssp. *sativus* cvs): Napoli (root and leaf), Amagar (root), Nutri-Red (root) and Rothild (root). Barley, meadow fescue and carrot cv. Napoli were chosen as model-plants, and all plant compartments were analysed. Only the edible compartments were analysed in the other plant species.

The solvent (acetone) residues in soil were allowed to evaporate for 3 days before seeds were sown. The number of plants per pot, selected based optimal biomass of plants in 4 L pots, was 5, 7, 10, 10 and 20, for carrot, barley, rape, wheat and fescue, respectively. After germination, growth conditions were set to 20/14 °C (day/night) and 16 h day length. The pots (individual trays) were irrigated when necessary to keep them moist, at least once a day, with water fertilised to electrical conductivity 1.5 mS/cm and pH 7.4. Control pots without test compounds were grown for all plant species. All treatments were conducted in triplicate.

Harvesting

Both control and exposed plant materials were harvested when mature or ripe (after 2–3 months). Leaf was cut while

root still was in the pots. Roots were carefully washed in tap water. All plant materials were dried (1 day at 50 °C, 2 days at 40 °C) (controls and exposed material in separate ovens to prevent cross-contamination) immediately after harvesting and stored in paper bags at room temperature until analysis, approximately 3 weeks. Biomass of the plant compartments root and leaf in each pot was weighed before and after drying. A small test to compare concentration levels in dried and not-dried seeds ($n=3$) was performed to check for significant evaporation during the drying processes (data not shown).

Analytical methods

Sample preparation followed the QuEChERS (quick, easy, cheap, effective, rugged and safe) method (Lehotay et al. 2005). In brief, plant and soil samples were spiked with 2-brom-biphenyl as an internal standard and extracted with double-distilled water and acetonitrile. All samples were initially cleaned up with primary–secondary amine. Further clean-up of seeds (DSC-18 sorbent) and of carrot, meadow fescue and barley (Envi-Carb) was performed. The extracts were analysed using an Agilent 6890 N gas chromatograph (GC) connected to an Agilent 5973 mass spectrometer with an inert ion source operated in selected ion monitoring mode. The GC was equipped with a Gerstel Programmable Temperature Vaporising Injector (Mühlheim Ruhr, Germany). Separation was performed using a fused silica J&W Scientific HP-5MSI (0.25 mm i.d. × 30 m) with 0.25 μm film thickness. For details of sample preparation, analysis and information about retention time, quantification ions and recovery, please see Table S1 and S2 in the Electronic Supplementary Material. Except for DEET in meadow fescue leaf and TBP in carrot root and meadow fescue leaf, the limit of quantification (LOQ) was set to 0.01 μg/g. In meadow fescue leaf, the LOQ had to be increased to 0.05 μg/g. Unfortunately, no LOQ was set at 0.01 μg/g for DEET in meadow fescue leaf and TBP in carrot root due to interferences. However, in real samples, the concentrations were relative high ($\text{DEET} \geq 0.08 \text{ μg/g}$ and $\text{TBP} \geq 0.92 \text{ μg/g}$) and the interferences became insignificant. All results for plant and soil concentrations were calculated based on dry weight.

Statistical analysis

Differences in concentrations of the compounds between species were tested using the software PROC GLM in SAS 9.0 (SAS Institute, Cary, NC, USA) with Ryan–Einot–Gabriel–Welsch Q multiple-comparison test. For all the tests, the significance level was set at $p < 0.05$.

Results and discussion

The nominal initial soil concentration was 1 mg/kg, but the actual measured initial soil concentration ($n=4$, average \pm standard deviation) for TBP, TCEP, TCPP, DEET and NBBS was 0.62 ± 0.05 , 0.85 ± 0.11 , 0.72 ± 0.12 , 1.00 ± 0.12 and 1.03 ± 0.12 mg/kg, respectively. The present study sought to compare root uptake and translocation of different potential contaminants to leaves and seeds, so it was important to select an initial soil concentration that was realistic for detection of the contaminants in plant material, although potentially higher than a realistic exposure situation. The initial soil values selected are comparable to those used in other studies, and spiking of soil is also a commonly used technique (Boxall et al. 2006; Gao et al. 2005; Winker et al. 2010; Wu et al. 2010). Data on residual levels of emerging contaminants in sewage sludge, particularly regarding DEET and NBBS, are scarce, and the concentrations in sewage sludge can vary widely (Harrison et al. 2006; Clarke and Smith 2011). Therefore, realistic concentrations in agricultural soil are actually not known. However, while many previous studies have been short-term, e.g. hydroponic cultures with incubation commonly up to a few weeks (Briggs et al. 1982; Murano et al. 2010), the present study involved plant growth over 17 weeks, providing a more realistic picture that accounted for concentration dilution during growth, soil degradation and possible *in planta* metabolism (Schröder et al. 2007).

Plant uptake and translocation

The bioconcentration factors for root (RCF), leaf (LCF) and seed (SCF) were calculated as concentration in plant compartments (milligrams per kilogram dry weight) over actual measured initial soil concentration (milligrams per kilogram dry weight). Despite high variations between species and cultivars, the concentrations in roots were generally lower than those in leaves for all test compounds except TBP (log $K_{ow}=4$) in carrots (Fig. 1a–c). For roots (Fig. 1a), a general higher uptake in carrots than in barley and meadow fescue was observed, with TCPP showing particularly high uptake in carrot (RCF=10–20). RCF for TBP and DEET in carrot was in the range 1.7–4.6 and 0.4–2.3, respectively, while it was even lower (range, 0.2–0.7) in NBBS and TCEP.

Comparison of RCF between the four cultivars of carrot revealed a significant ($p<0.05$) difference for TBP, with higher levels in cv. Napoli and cv. Nutri Red (RCF, 4.4–4.6) than those of cv. Amagar and cv. Rothild (1.6–2.5) and for DEET where cv. Nutri Red was higher than Amagar (Fig. 1a). However, there was a general trend for the highest average RCF for all compounds tested to be found in cv. Napoli and the lowest in cv. Amagar (Fig. 1a). In contrast, an opposite trend is reported for metformin (a cationic

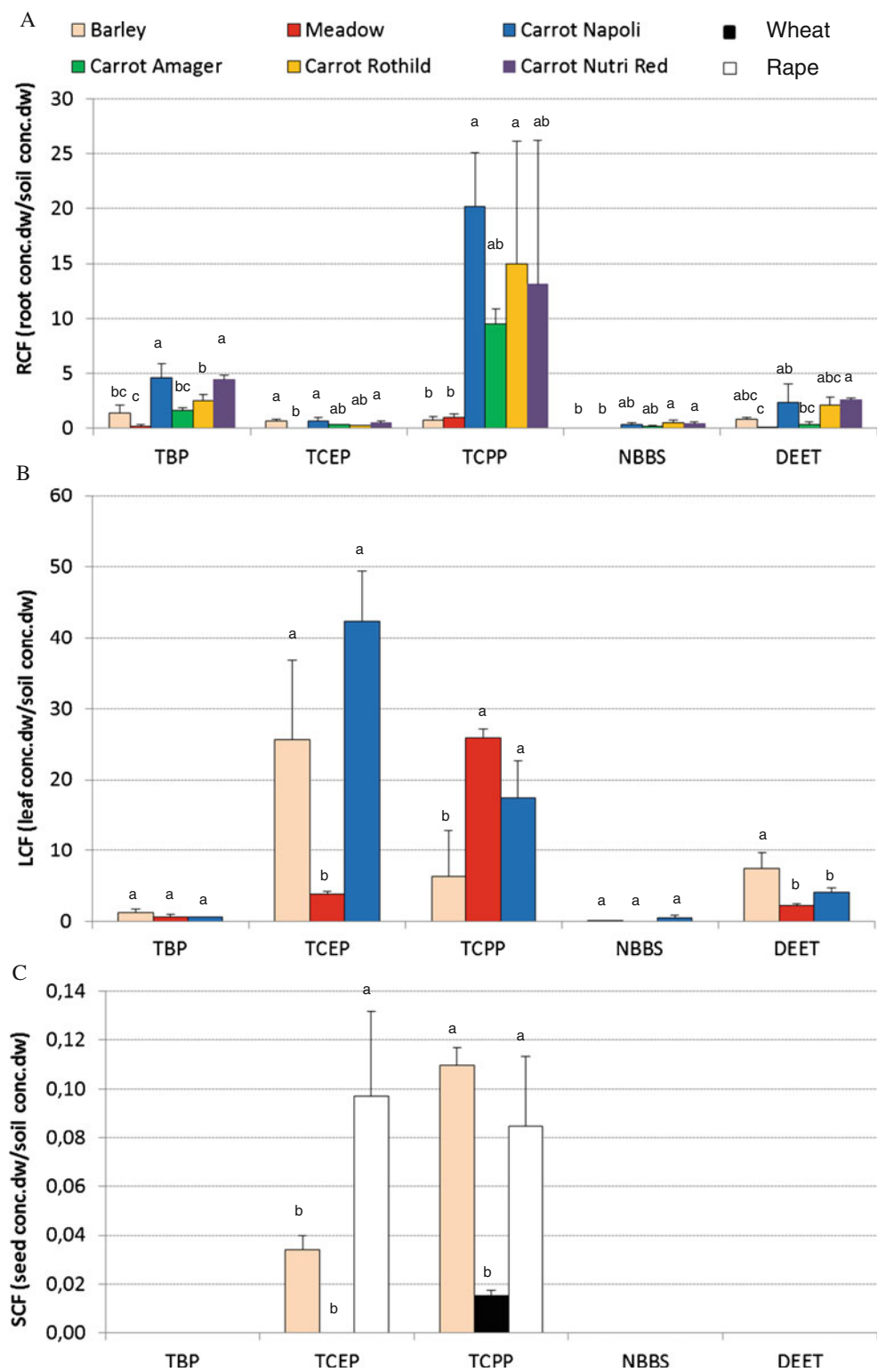
pharmaceutical) which showed lower RCF in carrot cv. Napoli than in cv. Amagar, RCF 2 and 10, respectively (Eggen et al. 2011). The highest measured RCF in barley and meadow fescue was 1.4 for TBP in barley and 0.9 for TCPP in meadow fescue, respectively (Fig. 1a). The concentration was below the LOQ for NBBS in both plant species and for TCEP in meadow fescue.

The concentrations in leaves were generally higher than those in roots and also showed a different pattern for different compounds (Fig. 1b). TCEP demonstrated high translocation to leaves, with LCF ranging from 3.9 in meadow fescue to 26 and 42 in barley and carrot, respectively, while the RCF was <1 for all three plant species. Ratio leaf/root TCPP, also with $RCF<1$ for barley and meadow fescue, showed significantly higher uptake to leaves of meadow fescue than barley (LCF 25.6 and 6.4, respectively) (Fig. 1b). In leaves of carrot cv. Napoli, the concentration of TCPP was comparable to that in roots (RCF and LCF in range 10–20). The LCF for DEET ranged from 2.3 to 7.4 (RCF 0.1–2.6), and barley leaves showed significant higher uptake than meadow fescue and carrot (Fig. 1b). Uptake of TBP to leaves was low, $LCF<1.2$, for all three plant species with no significant differences. NBBS which was not measurable in roots of barley or meadow fescue was detected in leaves of both barley and carrot cv. Napoli (LCF 0.08–0.5). The high difference between leaf (average 3.3 mg/kg) and root (<0.05 mg/kg) for TCEP in meadow fescue (ratio $>1,000$ if concentration in root is estimated to half of $LOQ=0.025$ mg/kg) is the highest root–leaf translocation in this study. High root–leaf ratio was also observed for TCEP in barley and carrot (range of 45–75), TCPP in meadow, barley and carrot and DEET in meadow fescue (range, 20–30).

The translocation of test compounds to seeds was low, with only TCPP being detected in wheat, barley and rape and TCEP in barley and rape (Fig. 1c). The concentration of TCPP was significantly higher in barley and rape seeds than in wheat, with SCF 0.110, 0.085 and 0.015, respectively. The TCEP concentration in rape seeds was significantly higher than that in barley, with SCF 0.097 and 0.034, respectively (Fig. 1c).

The control pots were standing close to the exposed pots, but none of the test compounds were detected above the LOQ in the control plants except for TCPP in control rape seeds, in which had concentrations of 0.010–0.014 mg/kg (compared with 0.060–0.120 mg/kg in exposed pots). TCPP and TCEP can both occur in indoor and outdoor air samples (Marklund et al. 2003; Reemtsma et al. 2008), but analysis of leaves and seeds from control plants showed that the greenhouse air atmosphere was not a significant source in the present study. In addition, the Henry's law constant values are low for the test compounds (Table 2), and evaporation from soil to leaves is not expected. Thus, the results

Fig. 1 Bioconcentration factors in roots (RCF) (a), leaves (LCF) (b) and seeds (SCF) (c) presented as milligram test compound per kilogram dry weight plant material ($n=3$) per milligram test compound per kilogram dry weight soil ($n=4$). Average and standard deviation ($n=3$ pots) is present. The $\log K_{ow}$ for TBP=4.0, TCEP=1.44; TCEP=2.59; NBBS=2.31 and DEET=2.18



for control leaves and seeds support the assumption that root uptake and translocation was the main transport pathway to aboveground plant compartments.

A high plant species variation was observed in our study, e.g. general higher uptake in carrot roots than meadow fescue and barley, lower uptake of TCEP and DEET in

leaves of meadow fescue than barley, and the opposite pattern, higher in meadow fescue than barley leaves was measured for TCPP. A high variation in uptake between species, or even between cultivars, is not unlikely and has previously been reported in several studies (Gonzalez et al. 2005; Inui et al. 2008a; Lunney et al. 2004; White 2002;

Zhang et al. 2009). Suggested explanations for such variations are differences in quantity and quality of root exudates, plant composition, root structure, biomass, endophytes populations and multi-species interaction (Kelsey and White 2005; Mattina et al. 2006; White et al. 2003a, b) (Khan and Doty 2011; Li et al. 2012). While the focus in food safety is to avoid high uptake and translocation to edible plant compartments, the opposite is the case for phytoremediation. The high translocation of chlorinated organophosphates to leaves of carrot, meadow and barley indicate that species from these plant families might be particularly suitable for phytoremediation for such compounds (Fig. 1, Table 3). High translocation to leaves is also reported for sulfolane (estimated by EPISuit, log K_{ow} -0.24, water solubility 292.8 g/L) to cattail, *Typha latifolia*, with leaf/root ratio > 150 (Doucette et al. 2005) and for carbamazepine (estimated by EPISuite, log K_{ow} 2.25, water solubility 17.7 mg/L) to ryegrass, *Lolium perenne* (Winker et al. 2010). Since there is a clear plant species variance for uptake and translocation for different contaminants, it is necessary to investigate which plant species is most optimal in each case.

The experimental bioconcentration factors found in this study indicate that there is not a clear relationship between log K_{ow} and plant uptake and translocation. Generally, LCF was higher (20–42) for TCEP (log K_{ow} 1.44) and TCPP (log K_{ow} 2.59) than for DEET and NBBS (log K_{ow} 2.18 and 2.31, respectively) (Fig. 1b). Biodegradation influences the environmental fate of contaminant compounds and TCEP and TCPP, both chlorinated compounds, showed the highest estimated half-lives (Table 2). Thus, since soil biodegradation rates also influence a compounds' potential for transfer to plants, plant uptake models should include degradation kinetic parameters. In theoretical structure–activity studies, the polarisability (\AA^3) of contaminants has been shown to have a good correlation with various physico-chemical properties, including bioconcentration factors (Hong et al. 2009; Papa et al. 2007; Staikovaa et al. 2004) and chemico-biological interactions (Hansch et al. 2003; Karelson and Lobanov 1996; Verma et al. 2005). The passage of xenobiotics through endodermic pores in plant roots is reported to be dependent on chemical polarity and molecular

configuration (van Leeuwen and Vermeire 2007). TCEP and TCPP also have higher water solubility, electronegativity and electronaffinity (Heimstad et al. 2001) than the other compounds studied here (Table 2). However, the species-dependent differences seen in the present study and in several other studies (Collins and Willey 2009; Eggen and Lillo 2012; Zhang et al. 2009) indicate complex biological effects that are not yet understood. For instance, several studies show that both hydrophobic organic compounds and hydrophilic dissociable organic compounds can be present in higher concentrations in roots than in leaves (Herklotz et al. 2010; Migliore et al. 1996), and a hydrophilic cationic pharmaceutical can be accumulated in oily rape seeds (Eggen and Lillo 2012). It is important to reveal regulation of contaminants or emerging contaminants with potential to high uptake and translocation to edible plant compartments like carrot, seeds or forage grasses. Today, it is no regulation or guidelines for emerging contaminants content in food items.

Growth effects

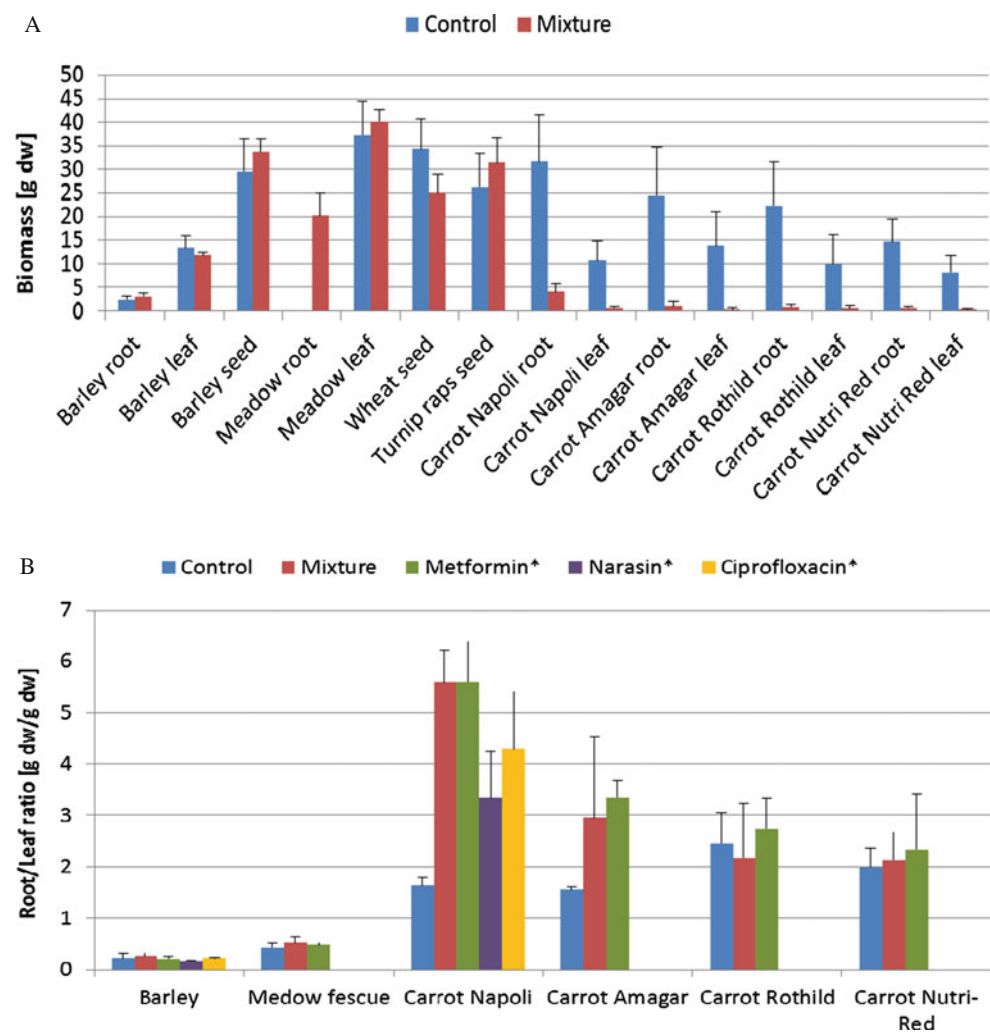
Plant growth and mortality were visually and quantitatively measured in terms of decline of plant biomass per pot (Fig. 2). Comparison of biomass of root and leaf (given as grams dry weight per pot) grown in control pots and in pot exposed to a mixture of the investigated emerging contaminants is shown in Fig. 2a. Under optimal plant growth conditions, the root–shoot ratio is quite specific for each plant species, but a number of external factors, e.g. nutrient and water supply, can alter this ratio (Marschner 1995). In the present experiment, no difference in the root/leaf ratio was observed for exposed meadow fescue and barley compared with the control (Fig. 2b). However, for the carrot cvs. Napoli and Amagar, the root–leaf ratio was higher in exposed plants (5.6 and 3.0, respectively) than in control plants (1.6). No clear differences were seen for the other two carrot cvs. Rothild and Nutri-Red. The same pattern of higher root–leaf ratio in cvs. Napoli and Amagar exposed to metformin, ciprofloxacin and narasin has been reported in a recently published plant uptake study (Eggen et al. 2011). In addition, root vegetables, e.g. carrot and radish, have been found to be more

Table 3 Summary of measured bioconcentration factor trends in root (RCF), leaf (LCF) and seed (SCF) for the different test compounds independent of statistical significance is shown

RCF _{Barley}	TBP ^a >DEET ^a >TCPP ^a >TCEP ^a	LCF _{Barley}	TCEP ^a >DEET ^b >TCPP ^b ~TBP ^b >NBBS ^b	SCF _{Barley}	TCPP ^a >TCEP ^b
RCF _{Meadow}	TCPP ^a >TBP ^b >DEET ^b	LCF _{Meadow}	TCPP ^a >TCEP ^b >DEET ^c >~TBP ^d ,NBBS ^d	SCF _{Wheat}	TCPP
RCF _{Carrot-Napoli and Amagar}	TCPP ^a >TBP ^b >DEET ^b >TCEP ^b ~NBBS ^b	LCF _{Carrot}	TCEP ^a >TCPP ^b >DEET ^c	SCF _{Rape}	TCEP ^a >TCPP ^a
RCF _{Carrot-Rothild}	TCPP ^a >TBP ^b >DEET ^b >TCEP ^b ~NBBS ^b				
RCF _{Carrot-Nutri Red}	TCPP ^a >TBP ^b >DEET ^c >TCEP ^d ~NBBS ^d				

Results of multiple comparison is shown by superscripted letters
Significant differences ($p < 0.05$) are marked

Fig. 2 Biomass of control and exposed plants given in dry weight (a) and root–leaf ratio based on dry weight (b). Biomass data from plants exposed to emerging contaminants in a previous experiment (Eggen et al. 2011) are marked with an *asterisk*. ‘Mixture’ is results from the present study where test compounds were added in a mixture cocktail of TBP, TCEP, TCP, DEET and NBBS. Published data of metformin, narsin and ciprofloxacin were tested in separate pots



sensitive to phytotoxins (EuropeanFoodSafetyAuthorities 2008; Migliore et al. 2003). The results of the present study confirm that there can be variation between different species and even cultivars of the same species after exposure to these emerging contaminants.

Environmental and food safety relevance

There is an increasing number of emerging contaminants in the environment, including pharmaceuticals and different kinds of additives used in everyday products. These compounds can reach the terrestrial or aquatic food web through transfer from consumer products into wastewater from homes, hospitals and industries or by leaching into groundwater when disposed of in municipal landfills (Table 1). WWTPs discharge to lakes and seas and thus influence the aquatic food web, while sewage sludge is applied to soil or soil mixtures used for cultivation of crops. Organophosphates have been found in marine and freshwater biota (mussels, crab, fish) (Evenset et al. 2009; Leonards et al. 2011; Sundkvist et al. 2010), human milk (Sundkvist et al. 2010) and drinking water

(Galassi et al. 1989 TCP, which in the present study shows high uptake in carrot root and forage grass, has been detected in fish muscles in capelin and in milk and plasma in harbour seal (Sagerup et al. 2011; Sundkvist et al. 2010).

A TCEP risk assessment from 2009 claimed that “since there is no indication that TCEP may show a bioaccumulation potential, a risk characterization for exposure via the food chain is not necessary” (European Commission 2009). Similarly, a risk assessment for TCP states that owing to “... lack of any significant bioaccumulation potential of TCP, it is reasonable to conclude that there are no risks” (European Commission 2008). Based on the experimental bioconcentration factors for TCEP and TCP, the high variation between species found in the present study and the relatively long half-life in soil and a persistency potential, there is reason to investigate the transfer and possible bioaccumulation of these compounds in food webs more deeply.

More generic knowledge about the relationship between the chemical properties of various compounds, uptake mechanisms into crops and plant composition is necessary

in order to perform health risk assessments where soil–plant transfer is part of the exposure route. Such knowledge is also important for identification and prediction of compounds with potentially high transfer to human and livestock food webs. Regulatory authorities should pay special attention to these compounds, and measures to reduce or remove sources should be introduced in an early phase. This is also valuable knowledge for food authorities devising restrictions or recommendations for cultivation of certain crops in areas with enhanced levels of organic compounds.

Conclusions

In this 17-week pot experiment, the organophosphates TCEP and TCPP generally exhibited higher uptake and translocation in crop plants than TBP, the insect repellent DEET and the plasticiser NBBS, despite DEET and NBBS having comparable $\log K_{ow}$ values as TCEP. Although TCEP and TCPP had similar properties, there were clearly species-specific uptake patterns in meadow fescue, barley and carrot. The surprisingly high translocation of TCPP into leaves of meadow fescue, a livestock forage species, is of particular concern and highlights the necessity for further studies investigating the effects of these compounds in the food web, to improve regulatory guidelines.

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