



Endocrine disruptors in breeding ponds and reproductive health of toads in agricultural, urban and natural landscapes

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HIGHLIGHTS

- Anthropogenic contaminants can cause lifelong reproductive impairment in wildlife.
- We found endocrine disruptor chemicals in rarely studied amphibian breeding habitats.
- Toads from agricultural, urban and natural ponds reproduced equally well in the lab.
- However, offspring fitness was reduced in agricultural and urban populations.
- Thus, transgenerational effects of pollution may contribute to amphibian declines.

GRAPHICAL ABSTRACT



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ABSTRACT

Many chemical pollutants have endocrine disrupting effects which can cause lifelong reproductive abnormalities in animals. Amphibians are the most threatened group of vertebrates, but there is little information on the nature and quantity of pollutants occurring in typical amphibian breeding habitats and on the reproductive capacities of amphibian populations inhabiting polluted areas. In this study we investigated the occurrence and concentrations of endocrine disrupting chemicals in the water and sediment of under-studied amphibian breeding habitats in natural, agricultural and urbanized landscapes. Also, we captured reproductively active common toads (*Bufo bufo*) from these habitats and let them spawn in a 'common garden' to assess among-population differences in reproductive capacity. Across 12 ponds, we detected 41 out of the 133 contaminants we screened for, with unusually high concentrations of glyphosate and carbamazepine. Levels of polycyclic aromatic hydrocarbons, nonylphenol and bisphenol-A increased with urban land use, whereas levels of organochlorine and triazine pesticides and sex hormones increased with agricultural land use. Toads from all habitats had high fecundity, fertilization rate and offspring viability, but the F1 generation originating from agricultural and urban ponds had reduced development rates and lower body mass both as larvae and as juveniles. Females with small clutch mass produced thicker jelly coat around their eggs if they originated from agricultural and urban ponds

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compared with natural ponds. These results suggest that the observed pollution levels did not compromise reproductive potential in toads, but individual fitness and population viability may be reduced in anthropogenically influenced habitats, perhaps due to transgenerational effects and/or costs of tolerance to chemical contaminants.

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1. Introduction

Chemical contaminants enter the hydrosphere from multiple sources. By drift, runoff and leaching, surface and ground waters receive pesticides and fertilizers from agricultural areas and various other pollutants from domestic and industrial wastewater discharges (Holt, 2000). An enormous body of literature shows that these contaminants are ubiquitously present in freshwater lakes and streams, usually in minute concentrations (Hoffman et al., 2003; Murray et al., 2010). Many hundreds of these chemicals are known to be harmful to animals and humans, even at very low concentrations, by interfering with the endocrine system and causing abnormalities in somatic and sexual development and reproductive physiology (Guillette and Edwards, 2008; Hoffman et al., 2003; Orton and Tyler, 2015). The WHO-IPCS defines exogenous substances that alter function of the endocrine system and consequently cause adverse health effects in an intact organism or its progeny or (sub)population as endocrine disruptors (Damstra et al., 2002). Such endocrine-disrupting chemicals (EDCs) can have life-long negative effects that permanently compromise reproductive potential. For example, perinatal exposure to the pesticides vinclozolin and methoxychlor increased the incidence of male infertility in adult rodents (Anway et al., 2005), whereas decreased fecundity and fertilization success was observed in adult fish after early-life exposure to ethynyl-estradiol, a synthetic estrogen found in contraceptive pills that contaminates sewage effluents (Maack and Segner, 2004).

Species that develop in water early in their life are especially vulnerable to the life-long effects of EDCs. One group at particular risk is amphibians, as most of their species have aquatic larvae, and their early ontogeny and sexual differentiation is sensitive to perturbation by water-borne substances with hormone-like activity (Orton and Tyler, 2015). Amphibians are also a group of serious conservation concern because their populations are disappearing around the globe at an alarming rate (Campbell Grant et al., 2016; Stuart et al., 2004). While the suspected reasons for these declines are linked to anthropogenic impacts such as habitat degradation, climate change and diseases, the role of chemical contaminants in amphibian diversity loss is not well understood (Campbell Grant et al., 2016; Orton and Tyler, 2015). Clarifying the latter is hindered by at least two major gaps in our knowledge.

On one hand, little information is available on the types and quantities of EDCs in the water bodies used by amphibians for spawning and larval development. One reason for this is that many amphibian species avoid flowing waters and large lakes with scarce shoreline vegetation, preferring relatively small, shallow pools instead (Wells, 2007). The input, dilution, dispersion and transformation of pollutants may be different in these water bodies than in large lakes and rivers, yet the former have been much less frequently surveyed for EDCs than the latter (Lorenz et al., 2017). Furthermore, amphibians persisting in anthropogenically influenced landscapes often breed in privately owned ponds, like angling ponds or residential garden ponds (Hassall, 2014). These kinds of water bodies are also rarely included in EDC surveys, despite the fact that they may be particularly exposed to certain contaminants, e.g. leaking from untreated wastewater (Smits et al., 2014). Even less is known about EDCs in the sediment of these ponds, even though many chemicals persist much longer in sediment than in water, and thereby sediment analyses can reveal past pollution events. Although the chemicals absorbed to sediment are considered to have little bioavailability, they can get re-suspended by disturbances to the pond bottom (Knott et al., 2009) and re-enter the food chain via bottom-grazing

and filter-feeding animals such as the larvae of many amphibian species (Wells, 2007).

On the other hand, it is poorly known if the amphibians living in habitats polluted by EDCs actually have reduced reproductive capacities. Typically, the effects of EDCs on reproductive health have been inferred in two ways (Hoffman et al., 2003; Orton and Tyler, 2015). First, laboratory ecotoxicology experiments usually expose developing larvae to EDCs and assess their reproductive anatomy, histology or physiology before or shortly after metamorphosis. Second, field studies usually look at correlations between land use or pollution levels and sex ratios or indirect indices of reproductive abilities such as sex hormone levels and secondary sexual characteristics in adults. The most frequently used endpoint in both kinds of studies is the incidence of intersex, a condition where an individual's gonads contain both female and male tissue. However, intersex can be a natural phase of ontogeny in some species (Orton and Tyler, 2015), and next to nothing is known about the reproductive success of intersex individuals (Jobling et al., 2002). Also, measurement of the above indices often requires invasive, even deadly techniques (e.g. sacrificing the animals for histological examination), which hinders such studies in dwindling populations where information on reproductive potential would be most needed.

In this study, we combined two objectives to contribute to filling these knowledge gaps. Firstly, we investigated the presence and concentration of EDCs in under-studied types of amphibian breeding habitats, i.e. ponds that are relatively small and shallow and/or privately owned. We sampled the water and sediment of these ponds in spring when amphibian larvae are in their early, sensitive stages of development, and we tested whether land use (i.e. agricultural, urban, or natural landscape) predicts the quality and quantity of EDC pollution. Secondly, we directly studied the reproductive capacity of common toads (*Bufo bufo*) that breed in these ponds. This species is widespread in Europe and occupies a broad range of habitat types, including landscapes influenced by agriculture and urbanization (Agasyan et al., 2009). It is protected in many countries, and although its populations are presumed large and stable, localized declines have been observed recently (Agasyan et al., 2009). Common toads are highly philopatric, with most individuals breeding in the pond in which they developed as larvae (Reading et al., 1991). Thus we hypothesized that, if breeding ponds in anthropogenically influenced landscapes are contaminated by higher levels and/or more potent kinds of EDCs than more natural ponds, then as a consequence of larval exposure to these contaminants we can detect reduced reproductive capacities in the adults breeding in anthropogenically influenced habitats. To evaluate reproductive capacity while ruling out the effects of actual exposure during breeding, we allowed adult toads to spawn in a non-polluted environment in which they could realize their full reproductive potential. We tested whether fecundity, fertilization rate, and the offspring's viability, development and growth were decreased in toads originating from habitats that are characterized by land use associated with higher pollution by EDCs.

2. Methods

2.1. Study sites and land use

We studied 12 ponds in Hungary (Table 1, Supplementary KMZ file), chosen based on three criteria: 1) common toads and other amphibians were known to breed there (from either our field experiences or personal communications by herpetologist colleagues), 2) we were able

Table 1
Characteristics of the 12 ponds in the study. See Supplementary KMZ file for a map of locations.

| Pond | Habitat type | Pond size (ha) | Proportion of landscape cover | | | | | | | Number of EDCs | Number of toad clutches in the lab |
|---------------------|--------------|----------------|-------------------------------|----------|------------|-------------|---------------|-------|------------|----------------|------------------------------------|
| | | | Arable fields | Pastures | Vegetation | Residential | Public, built | Roads | Rail-roads | | |
| Bajdázó (B) | Natural | 0.39 | 0 | 0.022 | 0.970 | 0 | 0 | 0.024 | 0 | 19 | 7 |
| Szárazfarkas (S) | Natural | 0.05 | 0 | 0 | 0.988 | 0 | 0 | 0.012 | 0 | 25 | 15 |
| János-tó (J) | Natural | 0.46 | 0 | 0 | 0.987 | 0 | 0 | 0.012 | 0 | 28 | 14 |
| Merzse (M) | Natural | 5.11 | 0.341 | 0.068 | 0.584 | 0 | 0 | 0.011 | 0 | 26 | 0 |
| Határrét (H) | Agricultural | 5.31 | 0.484 | 0.137 | 0.284 | 0.070 | 0 | 0.026 | 0 | 31 | 11 |
| Perőcsény (P) | Agricultural | 5.43 | 0.346 | 0.141 | 0.498 | 0 | 0 | 0.014 | 0 | 25 | 4 |
| Anyácsapuszta (A) | Agricultural | 3.38 | 0.802 | 0.051 | 0.145 | 0 | 0 | 0.007 | 0 | 31 | 2 |
| Gyermely (GY) | Agricultural | 17.10 | 0.735 | 0.014 | 0.213 | 0.001 | 0.027 | 0.011 | 0 | 29 | 0 |
| Pesthidegkút (PH) | Urban | 0.45 | 0.013 | 0 | 0.156 | 0.724 | 0.031 | 0.077 | 0 | 28 | 8 |
| Pilisszentiván (PS) | Urban | 5.15 | 0 | 0 | 0.282 | 0.455 | 0.173 | 0.076 | 0.015 | 31 | 10 |
| Pilisvörösvár (PV) | Urban | 9.47 | 0.004 | 0.024 | 0.270 | 0.531 | 0.083 | 0.077 | 0.014 | 33 | 7 |
| Göd (G) | Urban | 0.70 | 0 | 0 | 0.248 | 0.431 | 0.033 | 0.053 | 0.011 | 32 | 12 |

to subjectively categorize the surroundings of the pond as natural, urban or agricultural habitat, such that we had 4 ponds per habitat type; 3) logistics. The distance between ponds varied between 3.8 and 68.7 km, except for two urban ponds (Pilisvörösvár and Pilisszentiván) that were ca. 1 km apart, separated by a fragmented landscape of roads, railroads and buildings (Supplementary KMZ file).

To verify our subjective categorization of the three habitat types, we quantified land use around each pond using QGIS 2.18 (QGIS Development Team, 2017) as follows. We chose a buffer-zone width of 500 m following Cothran et al. (2013), as the migration distance of common toads rarely exceeds 500 m (Reading et al., 1991; Sinsch, 1988). We created the 500 m wide buffer zone around the shoreline of each pond based on the maps of Google Satellite, Bing Aerial, and OpenStreetMap Thunderforest Outdoors, using the Projected Coordinate System for Hungary (HD72/EOV). We obtained land-use polygons in the buffer zones from the Budapest shape file of the Urban Atlas 2012 (Copernicus Land Monitoring Service, European Environment Agency), using the Geoprocessing Tool of QGIS. Then we manually updated the land-use categorization and/or the shape of some polygons based on maps of the Hungarian Land Parcel Identification System (MEPAR), the satellite imagery of Google Satellite and Bing Aerial, the Time Laps feature of Google Earth, and our field experience. We assigned each polygon into one of seven land-use categories: “natural” vegetation (e.g. woodlands, non-agricultural meadows), arable fields, pastures, residential areas, public built areas (e.g. commercial and industrial areas), roads with vehicular traffic, and railroads. We measured the area of each polygon and each buffer zone using the Geometry Tools of QGIS.

For each pond, we calculated seven landscape variables as the total area of each land-use category divided by the area of the entire buffer zone (Table 1). Using these landscape variables we performed a principal component analysis (PCA), which yielded two axes with >1 eigenvalue, explaining 80.8% of variation in total; urban landscape areas loaded positively on the first axis whereas agricultural landscape areas

loaded positively on the second axis (Table 2). These two axes separated the 12 ponds clearly into three groups that matched our initial, subjective categorization of natural, urban and agricultural habitats, except for one pond (Merzse swamp, a nature-conservation area surrounded by agricultural fields) that received higher scores on the second axis than the other three natural ponds (Fig. 1).

2.2. Toad reproductive capacity

We captured adult toads at the start of their spawning season, between 16 and 28 March 2017 (91% captured over 3 days between 20 and 23 March). We aimed to capture at least 10 pairs from each pond; however, because the spawning season of common toads is extremely short (a few days), we could not capture gravid females at two ponds (Merzse and Gyermely). Captured animals were transported to our laboratory in Budapest, where temperature was 20 ± 1.55 °C and artificial light-dark cycles mimicked the natural photoperiod. We measured the body mass (± 0.1 g) of each toad, and housed each pair in $52 \times 37 \times 33$ cm plastic boxes filled with 15 L reconstituted soft water (RSW; 48 mg NaHCO_3 , 30 mg $\text{CaSO}_4 \times 2 \text{H}_2\text{O}$, 61 mg $\text{MgSO}_4 \times 7 \text{H}_2\text{O}$, 2 mg KCl added to 1 L reverse-osmosis filtered, UV-sterilized tap water) and

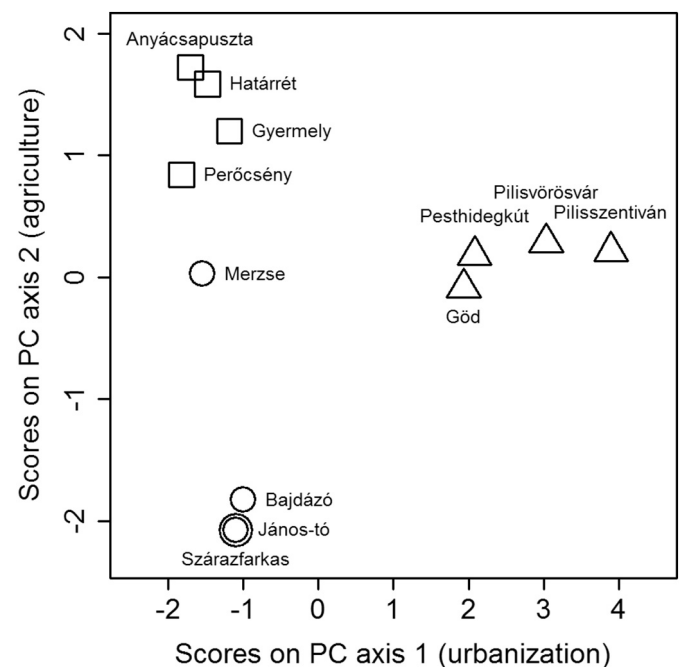


Fig. 1. Grouping of 12 ponds along the first two axes of the principal component analysis; symbol shape indicates the original, subjective habitat categorization (circles: natural, squares: agricultural, triangles: urban).

Table 2
The first two axes (PC1 and PC2) of the principal component analysis, and their Pearson correlations with the seven landscape variables.

| | PC1 | PC2 |
|--|-------|-------|
| Eigenvalue | 2.00 | 1.28 |
| % variance explained | 57.34 | 23.47 |
| Correlation of PC scores and landscape variables | | |
| Natural vegetation | -0.41 | -0.88 |
| Arable fields | -0.58 | 0.74 |
| Pastures | -0.52 | 0.54 |
| Public built areas | 0.86 | 0.13 |
| Residential areas | 0.91 | 0.11 |
| Roads | 0.96 | 0.07 |
| Railroads | 0.87 | 0.07 |

containing 4 vertical wooden sticks as spawning substrates. Each box housed one male and one female haphazardly chosen from the individuals captured at the same pond. Ninety out of 101 pairs spawned within one week (mostly within 3 days), after which all animals were released at the pond where they had been captured. Because capture success varied across ponds, and not all captured pairs spawned in the laboratory, we finally obtained 36 natural, 17 agricultural and 37 urban clutches (Table 1).

To quantify reproductive capacity, we took several measurements from each pair. On the day after spawning, we measured the parents' body mass again, and we calculated an estimate of fecundity (i.e. clutch mass before water uptake) by taking the difference between the female's pre-spawning and post-spawning body mass (note that post-hibernation toads do not feed before spawning). It is likely that the females laid all their eggs, because they appeared lean after spawning and the males completely lost interest in them. Only a single female had an appearance after spawning that implied that she might have retained some of her eggs, but her clutch mass was not an outlier (she had the 20th lowest residual clutch mass) and our conclusions on fecundity are not altered by excluding this clutch (results not shown). We also measured the mass of the entire clutch including not only the mass of the eggs but also the water absorbed by their jelly coats (i.e. clutch mass after water uptake). The difference between clutch mass before and after water uptake provides a proxy for jelly thickness (which is very difficult to measure directly in toad egg strings), assuming that the more water the jelly absorbs the thicker it gets; this trait is important for fitness because thicker jelly coats may provide greater protection from exogenous chemical stress (Edginton et al., 2007; Licht, 1985; Shu et al., 2015).

To estimate fertilization rate and offspring survival, from each clutch we placed ca. 30 eggs, taken from 3 haphazardly chosen parts of the egg string, into a 21 × 16 × 12 cm plastic box filled with 0.5 L RSW. We did not have identical numbers of eggs in all containers because ensuring that would have required longer manipulation of the egg string, risking the eggs falling out of the jelly and jeopardizing their further development. Five days after spawning, we measured fertilization rate as the proportion of eggs that started to develop, and removed the non-fertilized eggs (i.e. completely spherical eggs that had started to mold). Two weeks after spawning, when the embryos became free-swimming tadpoles (developmental stage 25, according to Gosner, 1960), we counted the proportion of embryos that survived to this stage, and started to feed the tadpoles with chopped and slightly boiled spinach. On day 17, we estimated the young tadpoles' average body mass by measuring the total mass (± 0.01 g) of four randomly chosen tadpoles from each family and dividing it by four; then we selected one healthy-looking individual from each family and moved it into a 2-L plastic box filled with 1 L RSW (in total, 88 tadpoles from 36 natural, 16 agricultural, and 36 urban families). The remaining eggs and tadpoles were released at the pond where their parents had been captured.

To measure the tadpoles' rate of development and growth, we raised them to metamorphosis; twice a week we changed their rearing water and fed them ad libitum with chopped spinach. When a tadpole started metamorphosis (i.e. appearance of forelimbs, developmental stage 42), we decreased the water level to 0.1 L and slightly tilted the container to allow the animal to leave the water. When it completed metamorphosis (i.e. disappearance of the tail, developmental stage 46), we measured its body mass (± 0.1 mg) and moved it into a clean rearing box.

To measure juvenile survival and growth, we raised the toadlets for ca. 5 months after metamorphosis. Each rearing box contained wet paper towels as substrate and a piece of egg carton as shelter, which were changed every two weeks. Toadlets were fed ad libitum with springtails and small crickets, amended with a 3:1 mixture of CaCO₃ and Promotor 43 powder (Laboratorios Calier S.A., Barcelona, Spain) containing vitamins and amino acids. Between 6th October and 10th November 2017, we measured the toadlets' body mass (± 0.01 g) and, as part of another experiment, we euthanized them using a water

bath containing 5.4 g/L MS-222 buffered with the same amount of Na₂HPO₄. The timing of this procedure was balanced among the animals from the three habitat types such that natural, agricultural and urban individuals were systematically rotated during the one-month period. This time of the year corresponds to the time right before the beginning of first hibernation and is relevant to toadlets' fitness because their pre-hibernation body mass predicts survival during hibernation and post-hibernation body mass (Üveges et al., 2016).

All experimental procedures were carried out according to the permits issued by the Government Agency of Pest County (Department of Environmental Protection and Nature Conservation) and the Budapest Metropolitan Municipality (Department of City Administration, FPH061/2472-4/2017). The experiments were further approved by the MTA ATK NÖVI Ethical Commission.

2.3. Water and sediment samples

During 25–27 April 2017, ca. one month after the peak spawning period of toads, we visited each pond once and collected samples for chemical analyses. The date of sampling corresponded to the early larval development of free-living toads, when the gonads are not yet differentiated (Falconi et al., 2004) and thus vulnerable to perturbations by EDCs: the tadpoles we found during sample collection were in developmental stages 26–33 (Gosner, 1960). From each pond, we collected 10 L water and 1 L sediment into 1-L amber glass bottles, and 3 L water into 1-L amber PET flasks (for measuring glyphosate, to avoid its adsorption to glass) in the areas where toad tadpoles were present and/or where the adults had been captured. Within these areas, samples were collected at several locations from the water close to the surface and from the upper layers of the bottom sediment (using a tube sampler). All samples were transported to the laboratory within 24 h, where they were stored at -18 °C until analyses.

We selected relevant groups of EDCs, and specific compounds within each group, by consulting databases and reviews (Hoffman et al., 2003; Molander et al., 2009; Orton and Tyler, 2015; USEPA, 2017). We also included several pharmaceuticals that frequently occur in surface waters but their endocrine disrupting effects are poorly known. In total, we analyzed our samples for 133 potential EDCs (including some of their metabolites): 60 pesticides, 22 phenolics, 19 polycyclic aromatic hydrocarbons (PAHs), 7 polychlorinated biphenyls (PCBs), 4 phthalates, 16 pharmaceuticals, and 5 natural sex hormones (see Supplementary Material for a complete list).

Chemical analyses were conducted at the Bálint Analitika accredited laboratory. All compounds were detected and measured by gas chromatography with mass spectrometry (GC-MS), except that pharmaceuticals, hormones, glyphosate and its metabolite aminomethylphosphonic acid (AMPA) were analyzed by reversed-phase high-performance liquid chromatography coupled to tandem mass spectrometry (RP-HPLC-MS/MS). All GC-MS analyses were carried out according to standards (PAHs and PCBs: MSZ 1484-6:2003 and MSZ 1484-11:2003, organochlorine pesticides: EPA 8081B:2007, other pesticides, phthalates, and bisphenol-A: EPA 8270D-2007, alkyl-phenols: MSZ EN ISO 18857-1:2007, chlorophenols: MSZ-EN 12673:2000 and MSZ 21470-97:2009). All compounds were detected in selected ion monitoring (SIM) mode using Agilent 5890 and 6890 GC-MS instruments. Analysis of pharmaceutical residues was performed in accordance with the EPA 1694:2007 standard. For natural hormones, glyphosate and AMPA, in-house methods were used. In case of hormones, water samples were extracted with dichloromethane (DCM); then the organic phase was evaporated, and the residues were dissolved in methanol. Sediment samples were extracted with acetone prior to extraction with DCM to dehydrate the samples. In case of glyphosate and AMPA, a derivatization with 9-fluorenylmethyl chloroformate was applied prior to analysis. The conditions of the derivatization reaction were optimized as: pH = 9 (borate), 10% acetonitrile, agitation for 60 min at room temperature; the reaction was then stopped by adding 500 μ L of 10% formic acid to each sample.

After filtration, the samples were analyzed using RP-HPLC-MS/MS; all compounds were detected by multiple reaction monitoring (MRM) using a Sciex 6500 QTrap mass spectrometer. Detection limits are given in the Supplementary Material.

2.4. Statistical analyses

All analyses were run in the R 3.3.1 environment (R Core Team, 2016), using the following packages: cluster (Maechler et al., 2017),

Table 3

EDC concentrations in a) water and b) sediment ('nd': not detected; abbreviations of pond names as in Table 1).

| a) Chemicals in water (ng/L; [*] µg/L) | | Natural ponds | | | | Agricultural ponds | | | | Urban ponds | | | | |
|---|--|-------------------------|------|------|------|--------------------|------|------|------|-------------|------|------|------|-----|
| Group | Compound | B | S | J | M | H | P | A | GY | PH | PS | PV | G | |
| Polycyclic aromatic hydrocarbons (PAH) | Naphthalene | 15 | 80 | 90 | 23 | 67 | 19 | 17 | 22 | 23 | 20 | 78 | 25 | |
| | 2-methyl-naphthalene | 15 | 62 | 94 | 13 | 47 | 15 | 15 | 18 | 13 | 16 | 62 | 15 | |
| | 1-methyl-naphthalene | 6 | 25 | 39 | 7 | 21 | 6 | 6 | 8 | 6 | 8 | 26 | 8 | |
| | Acenaphthylene | nd | 2 | 5 | 1 | 3 | nd | 1 | 1 | 1 | 2 | 9 | 1 | |
| | Acenaphthene | 1 | 4 | 43 | 1 | 4 | 1 | 1 | 1 | 1 | 2 | 6 | 2 | |
| | Fluorene | 40 | 149 | 19 | 3 | 13 | 3 | 3 | 3 | 44 | 7 | 18 | 5 | |
| | Phenanthrene | 25 | 176 | 34 | 9 | 24 | 6 | 6 | 6 | 42 | 11 | 37 | 11 | |
| | Anthracene | 2 | 20 | 3 | nd | 2 | nd | nd | nd | 8 | 1 | 2 | 1 | |
| | Fluoranthene | 2 | 15 | 18 | 3 | 15 | 2 | 3 | 3 | 5 | 4 | 22 | 4 | |
| | Pyrene | 1 | 7 | 11 | 2 | 8 | 1 | 1 | 2 | 3 | 2 | 11 | 2 | |
| | Benz(a)anthracene | 1 | 4 | 7 | 1 | 5 | 1 | 1 | 1 | 2 | 1 | 6 | nd | |
| | Chrysene | 1 | 3 | 5 | 1 | 3 | 1 | nd | 1 | 1 | 1 | 3 | nd | |
| | Benzo(b)fluoranthene+ Benzo(k)fluoranthene | 1 | 8 | 17 | 2 | 8 | 1 | 1 | 2 | 4 | 2 | 14 | 1 | |
| | Benzo(e)pyrene | 1 | 6 | 7 | nd | 3 | 1 | 1 | nd | 2 | 1 | 6 | 1 | |
| | Benzo(a)pyrene | nd | 2 | 3 | nd | 2 | nd | nd | nd | 1 | nd | 4 | nd | |
| | Phthalates | Diethyl-phthalate | nd | nd | 10 | 10 | 10 | nd | 10 | 10 | 30 | 20 | 30 | 20 |
| | | Dibutyl-phthalate | 30 | nd | 250 | nd | 40 | nd | 120 | 140 | 180 | 110 | 90 | 80 |
| | Pesticides | Di-ethylhexyl phthalate | 730 | 180 | nd | 300 | nd | 520 | 270 | nd | 310 | 230 | 240 | 570 |
| | | o,p-DDE + p,p-DDE | nd | nd | nd | nd | nd | nd | 3 | 1 | nd | nd | nd | nd |
| | | Terbutylazine | nd | nd | nd | nd | nd | nd | nd | 330 | nd | nd | nd | nd |
| Terbutryn | | nd | nd | nd | nd | nd | nd | nd | 30 | nd | nd | nd | nd | |
| Hormones | Glyphosate* | 5.18 | 15 | 2.36 | 10.9 | 8.85 | 7.06 | 10.4 | 14.8 | 6.52 | 8.12 | 10.9 | 9.5 | |
| | Aminomethylphosphonic acid* | 2.62 | 25.9 | 3.39 | 6.21 | 17.6 | 6.67 | 7.33 | 9.94 | 14 | 19.1 | 15.2 | 4.98 | |
| Pharmaceuticals | Testosterone | nd | nd | nd | nd | nd | 10 | nd | 9 | nd | nd | nd | nd | |
| | Caffeine | nd | nd | 90 | 40 | 60 | nd | 40 | nd | nd | 50 | 70 | 90 | |
| | Sulfamethoxazole | nd | nd | 1 | nd | nd | nd | 1 | 1 | nd | nd | 1 | nd | |
| | Carbamazepine* | nd | 0.06 | 90.5 | nd | 276 | 11.2 | nd | nd | 1.3 | 50.2 | 65.3 | 16.1 | |
| | Carbamazepine epoxide | nd | nd | nd | nd | <5 | nd | nd | nd | nd | <5 | <5 | nd | |
| Ketoprofen | nd | nd | nd | nd | nd | nd | nd | nd | nd | nd | nd | <5 | | |
| b) Chemicals in sediment (µg/kg) | | Natural ponds | | | | Agricultural ponds | | | | Urban ponds | | | | |
| Group | Compound | B | S | J | M | H | P | A | GY | PH | PS | PV | G | |
| Polycyclic aromatic hydrocarbons (PAH) | Naphthalene | 6 | 8 | 20 | 7 | 8 | 5 | 7 | 8 | 7 | 20 | 10 | 20 | |
| | 2-methyl-naphthalene | 5 | 6 | 16 | 6 | 7 | 4 | 5 | 7 | 6 | 12 | 9 | 18 | |
| | 1-methyl-naphthalene | 2 | 3 | 9 | 3 | 4 | 2 | 3 | 3 | 3 | 7 | 5 | 10 | |
| | Acenaphthylene | nd | nd | 1 | 1 | nd | nd | nd | nd | 1 | 2 | 5 | 2 | |
| | Acenaphthene | nd | nd | 1 | nd | 1 | nd | nd | nd | nd | 1 | 7 | 5 | |
| | Fluorene | 1 | 1 | 2 | 1 | 3 | 1 | 1 | 1 | 1 | 3 | 13 | 12 | |
| | Phenanthrene | 1 | 3 | 10 | 7 | 4 | 1 | 3 | 3 | 6 | 27 | 115 | 35 | |
| | Anthracene | nd | nd | 1 | 1 | 1 | nd | nd | nd | 1 | 4 | 23 | 10 | |
| | Fluoranthene | 1 | 6 | 30 | 19 | 11 | 2 | 6 | 5 | 18 | 69 | 226 | 146 | |
| | Pyrene | nd | 4 | 16 | 13 | 7 | 1 | 4 | 3 | 14 | 42 | 164 | 99 | |
| | Benz(a)anthracene | nd | 2 | 9 | 8 | 4 | 1 | 2 | 2 | 9 | 26 | 112 | 57 | |
| | Chrysene | nd | 3 | 13 | 9 | 3 | 1 | 3 | 3 | 4 | 23 | 67 | 56 | |
| | Benzo(b)fluoranthene+ Benzo(k)fluoranthene | 1 | 14 | 139 | 25 | 17 | 3 | 9 | 7 | 24 | 98 | 261 | 153 | |
| | Benzo(e)pyrene | nd | 6 | 42 | 10 | 7 | 1 | 4 | 3 | 9 | 37 | 96 | 56 | |
| | Benzo(a)pyrene | nd | 3 | 10 | 8 | 5 | 1 | 2 | 2 | 10 | 27 | 107 | 50 | |
| | Indeno(1,2,3-cd)pyrene | nd | 4 | 34 | 5 | 6 | nd | 2 | 2 | 8 | 27 | 82 | 46 | |
| | Dibenzo(a,h)anthracene | nd | nd | 4 | 1 | 1 | nd | nd | nd | 1 | 5 | 15 | 5 | |
| | Benzo(g,h,i)perylene | nd | 5 | 24 | 9 | 7 | 1 | 3 | 2 | 8 | 28 | 70 | 48 | |
| | Phthalates | Dibutyl-phthalate | 10 | nd | nd | nd | 10 | 20 | 20 | nd | nd | 10 | 10 | 10 |
| | | Di-ethylhexyl phthalate | 30 | 30 | 60 | 50 | 60 | 20 | 30 | 10 | 30 | 100 | 40 | 140 |
| Phenolics | Nonylphenol | 127 | 135 | 124 | 113 | 194 | 143 | 94 | 70 | 172 | 323 | 179 | 306 | |
| | Bisphenol A | 3 | 7 | nd | 4 | 43 | 4 | 5 | 4 | 5 | 16 | 5 | 8 | |
| Pesticides | o,p-DDE + p,p-DDE | nd | 1 | 3 | 1 | 25 | 79 | 139 | 8 | 6 | 5 | 11 | 8 | |
| | o,p-DDD + p,p-DDD | nd | nd | nd | nd | 5 | 4 | 11 | nd | 1 | 2 | 40 | 13 | |
| | Dicofol | nd | nd | nd | nd | nd | nd | 1 | nd | nd | nd | 4 | 3 | |
| | Aldrin | nd | nd | nd | nd | nd | nd | 7 | nd | nd | nd | nd | nd | |
| Hormones | Dieldrin | nd | nd | nd | nd | nd | nd | 7 | nd | nd | nd | nd | nd | |
| | Estrone (E1) | nd | nd | nd | nd | 1.03 | nd | 0.94 | <0.5 | nd | 1.53 | 0.78 | 0.55 | |
| Pharmaceuticals | 17 α -ethynylestradiol (EE2) | nd | 4.78 | nd | nd | nd | nd | nd | nd | nd | nd | nd | nd | |
| | Caffeine | nd | nd | nd | nd | nd | 2.76 | nd | nd | nd | nd | nd | nd | |
| | Carbamazepine | nd | nd | nd | nd | 332 | nd | nd | nd | nd | nd | 514 | 106 | |

vegan (Oksanen et al., 2018), psych (Revelle, 2017), smatr (Warton et al., 2012), geepack (Halekoh et al., 2006), and lsmeans (Lenth, 2016). We set the level of statistical significance at 0.05, and we report means with standard errors.

2.4.1. Does land use predict EDC pollution?

We compared the three habitat types in a linear model to test if the number of detected EDC compounds (either in the water or in the sediment or both) differs between natural ponds and anthropogenically influenced ponds. To analyze the concentration of chemicals, we used the concentration of each EDC detected in water and in sediment as separate variables, because both the unit of measurement and the biological implications differ between water and sediment concentrations. Thereby we had 60 variables describing the concentrations of 41 EDCs either in the water or in the sediment (Table 3). Because many of these variables had zero-inflated distribution, we used non-parametric multivariate methods to analyze whether the concentrations of various EDCs correlated with urban or agricultural land use, as follows.

First we calculated the pairwise dissimilarity of all ponds based on either the 7 landscape variables or the 60 EDC variables. We used the Gower distance as metric of dissimilarity (function 'daisy' in package 'cluster'), because it does not ignore negative matches (i.e. when a certain landscape type or EDC is absent at both sites) and allows the assignment of differential variable weights. The number of compounds detected in our samples varied greatly across different chemical groups (Table 3), and we did not want our analysis to be biased by EDC groups with many compounds; therefore during the calculation of Gower distances we assigned to each compound a weight inversely proportional to the number of detected compounds in the respective group of EDCs (i.e. the total number of compounds, 60, divided by the number of compounds in the respective group). We did not apply differential weights for the landscape variables because even a small landscape area (e.g. a sewage plant) can be a considerable source of pollution.

To assess whether the two distance matrices (i.e. dissimilarities of ponds by landscape and by EDCs) were significantly correlated, we used the Mantel-test with 999 permutations (function 'mantel' in package 'vegan'). Then, to interpret this correlation (i.e. which EDCs are associated with which landscapes), we reduced the dimensionality of the data as follows. For the landscape variables we used the two PCA axes as described above, quantifying urban and agricultural land use respectively. For the 60 EDC variables, we summed the concentrations of compounds belonging to the same group, separating the following 6 groups: PAHs, phenolics, phthalates, pesticides, pharmaceuticals, natural hormones. For each group, concentrations measured in water and in sediment were summed separately, yielding 11 variables. Then we tested the pairwise relationships between the scores along each PCA axis and the concentrations in each EDC group using Spearman rank-correlations, and calculated the 95% confidence interval of each correlation coefficient by bootstrapping with 999 permutations (function 'cor.ci' in package 'psych').

2.4.2. Does land use predict toads' reproductive capacity?

As measures of each pair's reproductive capacity, we analyzed the following 8 variables. Because fecundity is known to increase with female size (Banks and Beebee, 1986; Reading, 1986), we statistically controlled for this effect by calculating residuals from a standardized major axis regression (function 'sma' in package 'smatr') with female post-spawning body mass as the explanatory variable and clutch mass before water uptake (i.e. the female's body mass loss from before to after spawning) as the dependent variable. This approach is favorable over simple ratios or ordinary least-squares regression residuals when the goal is to obtain accurate predicted values rather than merely testing if the slope differs from zero (Peig and Green, 2009, 2010). We used the residuals from this regression as measure of fecundity corrected for female size.

As a proxy for jelly thickness, we subtracted clutch mass before water uptake (i.e. female weight loss during egg laying) from clutch mass measured after water uptake. This variable, referred to henceforth as jelly mass, increases with the total mass of eggs, but our data indicated a non-linear association and we had no a priori knowledge on the function by which this relationship can be adequately described. Therefore, to take variation in clutch mass into account, we categorized clutch mass into three groups (i.e. small, medium and large clutches, before water uptake) with equal sample size, and analyzed the effects of habitat within these three groups (see below).

We measured fertilization rate as the proportion of eggs that started embryonic development, and embryo viability as the proportion of embryos that survived to the free-swimming tadpole stage. We did not analyze the survival of tadpoles and juveniles because only one out of 88 tadpoles died before metamorphosis and 5 out of 87 toadlets died before the termination of the study. Out of these 6 cases of mortality, 4 individuals were of urban origin while only one originated from each of the other two habitat types. We analyzed offspring body mass measured at three times: on day 17 (early tadpole stage), at completion of metamorphosis, and at termination (ca. 5 months after metamorphosis). We also analyzed the time to metamorphosis (number of days from spawning until the completion of metamorphosis).

Each of the above variables was used as a dependent variable in Generalised Estimation Equations models (function 'geeglm' in package 'geepack'). In these models, we allowed for the non-independence among pairs captured from the same pond using the 'exchangeable correlation' (or 'compound symmetry') association structure (Zuur et al., 2009). We used habitat type as a fixed factor, and we parameterized the design matrix such that we estimated the differences between the natural habitat type (used as intercept) and each of the two anthropogenically influenced habitat types (i.e. agricultural and urban). We did not use the PCA scores of the landscape variables as explanatory variables because their distribution was strongly bimodal and hence unsuitable for testing linear relationships (Table 1, Fig. 1). Note, however, that our habitat categories are in good agreement with the PCA scores (Fig. 1), and we had no clutches from site Merzse which turned out to be halfway between natural and agricultural ponds along the second PCA axis. In the model of juvenile body mass, we included the individual's age (i.e. number of days from completion of metamorphosis to termination; mean-centered) as a covariate. In the models of fertilization rate, embryo viability, offspring mass, and time to metamorphosis, we also included the body mass of both parents (measured after spawning; mean-centered) as covariates. In the model of jelly thickness, we included clutch mass (categorized as small, medium and large; see above) as fixed factor and its interaction with habitat type, then we performed pairwise post-hoc comparisons by calculating linear contrasts between natural and either agricultural or urban habitat within each clutch-mass category and correcting the significance level for multiple testing by the Tukey method (function 'lsmeans' in package 'lsmeans').

3. Results

3.1. Does land use predict EDC pollution?

Out of the 133 EDCs screened for, we detected 41 compounds in at least one of the 12 ponds: 19 PAHs, 3 phthalates, 2 phenolics, 9 pesticides, 6 pharmaceuticals, and 2 natural hormones (Table 3). Out of these, 21 compounds were present both in water and sediment samples; 9 compounds were found only in water while another 11 compounds were found only in sediment (Table 3). No polychlorinated phenols or PCBs were detectable in any of the ponds. The number of EDCs was slightly higher in agricultural ponds (29.0 ± 1.4 ; $p = 0.066$) and significantly higher in urban ponds (31.0 ± 1.1 ; $p = 0.014$) than in natural ponds (24.5 ± 1.9 ; Table 3).

Pond dissimilarity based on EDC concentrations in water and sediment correlated significantly with pond dissimilarity based on landscape

variables (Mantel-test: $r = 0.41$, $p = 0.008$). Pairwise correlations showed that the intensity of urbanization (i.e. pond scores along the first PCA axis) correlated positively with the sediment concentrations of PAHs and phenolics (nonylphenol and bisphenol-A), whereas the intensity of agriculture (i.e. pond scores along the second PCA axis) correlated positively with the sediment concentrations of organochlorine pesticides and estrone (Table 4). Furthermore, testosterone and triazine pesticides were only detected in the water of agricultural ponds (Table 3).

3.2. Does land use predict toads' reproductive capacity?

Fecundity corrected for female size did not differ between the three habitat types (Table 5, Fig. 2a). Jelly mass was significantly larger in agricultural and urban clutches when clutch mass was small, whereas medium and large clutches had relatively large jelly mass irrespective of habitat type (Table 6, Fig. 2b). Pairs from agricultural and urban ponds had similarly high fertilization rate (Table 5, Fig. 2c) and embryo viability as pairs from natural ponds (Table 5, Fig. 2d).

Tadpoles from natural ponds were larger at the start of larval life than tadpoles with urban and agricultural origin (Table 5, Fig. 2e). Despite taking less time to develop into metamorphosed toadlets (Table 5, Fig. 2f), tadpoles from natural ponds did not have smaller mass at metamorphosis compared to tadpoles from anthropogenically influenced ponds (Table 5, Fig. 2g). Furthermore, toadlets from natural ponds reached larger body mass by October compared to the animals with agricultural or urban origin (Table 5, Fig. 2h).

4. Discussion

Our chemical analyses of pond water and sediment revealed a complex picture on the EDC levels in amphibian breeding habitats. On one hand, 92 out of 133 pollutants were not present in detectable quantities in any of the studied water bodies (Supplementary Material), including many compounds known as dangerous EDCs, such as PCBs and various pesticides (Hoffman et al., 2003). On the other hand, several EDCs were present in many or all ponds, and even the small forest ponds that represent natural breeding habitats of many amphibians were contaminated. The concentrations we found generally fall within the range of values reported from other surface waters and sediments (see Table S1 for a non-comprehensive review), with a few notable exceptions. The levels of glyphosate, an herbicide used in very large amounts worldwide and also in Hungary, were higher in our ponds than those typically measured in large rivers and lakes; they were similar to the concentrations observed in urban runoff (Table S1). This finding, coupled with the ubiquity of glyphosate in our ponds irrespective of land use, suggests that a significant part of the glyphosate load in surface waters originates not only from agricultural applications but from other anthropogenic sources like weed control of railways, roadsides, and backyards (Hanke et al., 2010). Carbamazepine, an anti-epileptic and

Table 4

Spearman rank-correlations (with 95% confidence intervals; CI) between EDC concentrations and PCA scores of landscape composition. Correlation coefficients for which the CI excludes zero are highlighted in bold.

| EDC group | PC1 (urbanization) | PC2 (agriculture) |
|-----------------------------|--------------------------|--------------------------|
| PAHs in water | 0.52 (−0.18, 0.89) | −0.55 (−0.94, 0.13) |
| PAHs in sediment | 0.59 (0.03, 0.90) | −0.19 (−0.7, 0.43) |
| Phenolics in sediment | 0.60 (0.09, 0.97) | −0.06 (−0.61, 0.66) |
| Phthalates in water | 0.22 (−0.43, 0.69) | −0.17 (−0.84, 0.66) |
| Phthalates in sediment | −0.21 (−0.72, 0.64) | 0.53 (−0.04, 0.90) |
| Pesticides in water | 0.32 (−0.25, 0.80) | 0.24 (−0.46, 0.93) |
| Pesticides in sediment | −0.29 (−0.78, 0.45) | 0.80 (0.44, 0.96) |
| Pharmaceuticals in water | 0.24 (−0.33, 0.72) | 0.08 (−0.57, 0.70) |
| Pharmaceuticals in sediment | 0.11 (−0.50, 0.69) | 0.09 (−0.47, 0.68) |
| Hormones in water | −0.47 (−0.83, 0.00) | 0.38 (0.04, 0.78) |
| Hormones in sediment | 0.22 (−0.49, 0.78) | 0.62 (0.10, 0.85) |

Table 5

Reproductive capacity of toad pairs in relation to habitat type (Generalised Estimation Equations models).

| Dependent variable | Model parameters | Estimate ± SE | p |
|--------------------------------|-------------------------|-----------------|--------|
| Fertilization rate (logit) | Natural habitat, mean | 2.012 ± 0.245 | <0.001 |
| | Agricultural – natural | 0.265 ± 0.520 | 0.610 |
| | Urban – natural | −0.06 ± 0.291 | 0.840 |
| | Mother's body mass | 0.001 ± 0.009 | 0.950 |
| Embryo viability (logit) | Father's body mass | 0.006 ± 0.069 | 0.930 |
| | Natural habitat, mean | 4.097 ± 0.235 | <0.001 |
| | Agricultural – natural | −0.053 ± 0.300 | 0.860 |
| | Urban – natural | −0.132 ± 0.254 | 0.600 |
| Tadpole mass (mg) | Mother's body mass | −0.007 ± 0.006 | 0.250 |
| | Father's body mass | 0.010 ± 0.020 | 0.610 |
| | Natural habitat, mean | 40.855 ± 1.183 | <0.001 |
| | Agricultural – natural | −5.320 ± 2.705 | 0.049 |
| Time to metamorphosis (days) | Urban – natural | −4.355 ± 2.137 | 0.041 |
| | Mother's body mass | 0.007 ± 0.037 | 0.853 |
| | Father's body mass | 0.028 ± 0.136 | 0.838 |
| | Natural habitat, mean | 57.17 ± 0.209 | <0.001 |
| Mass at metamorphosis (mg) | Agricultural – natural | 2.585 ± 0.289 | <0.001 |
| | Urban – natural | 0.876 ± 0.284 | 0.002 |
| | Mother's body mass | 0.008 ± 0.010 | 0.461 |
| | Father's body mass | −0.011 ± 0.035 | 0.747 |
| Juvenile mass (g) | Natural habitat, mean | 164.294 ± 4.351 | <0.001 |
| | Agricultural – natural | −7.637 ± 6.448 | 0.240 |
| | Urban – natural | −4.030 ± 6.364 | 0.530 |
| | Mother's body mass | 0.063 ± 0.160 | 0.700 |
| Age at mass measurement (days) | Father's body mass | −0.734 ± 0.496 | 0.140 |
| | Natural habitat, mean | 4.561 ± 0.049 | <0.001 |
| | Agricultural – natural | −0.269 ± 0.065 | <0.001 |
| | Urban – natural | −0.436 ± 0.070 | <0.001 |
| | Age at mass measurement | 0.036 ± 0.007 | <0.001 |
| | Mother's body mass | 0.001 ± 0.002 | 0.778 |
| Father's body mass | 0.031 ± 0.011 | 0.006 | |

Sample sizes: 90 for fecundity and fertilization rate, 88 for embryo viability, 87 for tadpole mass, 86 for time to and mass at metamorphosis, and 82 for juvenile mass. The juveniles' age at mass measurement (days), and the mother's and father's body mass (g) were mean-centered before analysis.

anti-depressant drug which is very stable in the environment, was found at extraordinarily high concentrations in some of our ponds, in 5 samples exceeding the highest level reported for European rivers (Table S1). This indicates that persistent contaminants can accumulate heavily in small standing waters exposed to frequent input, e.g. due to wastewater leaching and runoff, or direct deposition associated with human activities such as tourism and angling.

Whether the detected EDC concentrations are dangerous to amphibians is difficult to assess due to a general paucity of data (reviewed in Table S1). Although almost all compounds detected in our ponds have been demonstrated to have reproduction-related EDC effects (Hoffman et al., 2003), there is large variation among chemical groups in the amount of experimental data, especially on aquatic vertebrates. For some intensively studied compounds, like bisphenol-A and ethynyl-estradiol, there is evidence that even very small, environmentally relevant concentrations can interfere with the sexual development of amphibians (Tamschick et al., 2016a, 2016b, 2016c). For some other compounds, like phthalates and glyphosate, only a few experiments were published so far on their effects on amphibian sexual development, which applied much higher concentrations than those found in our ponds and in other water bodies (Table S1). Disquietingly, we found no relevant publications at all on amphibians for most of the compounds we detected, including PAHs, the non-hormonal pharmaceuticals, and the pesticides excluding glyphosate and DDT-derivatives. Given the widespread occurrence of these chemicals in aquatic habitats (Lorenz et al., 2017; Murray et al., 2010), we urgently need more data on their endocrine-disrupting effects at environmentally realistic concentrations.

Most of the EDCs we found in sediments are environmentally persistent, and several groups of these chemicals showed a concentration gradient increasing from natural towards anthropogenically influenced ponds. Specifically, urban land use was associated with higher levels

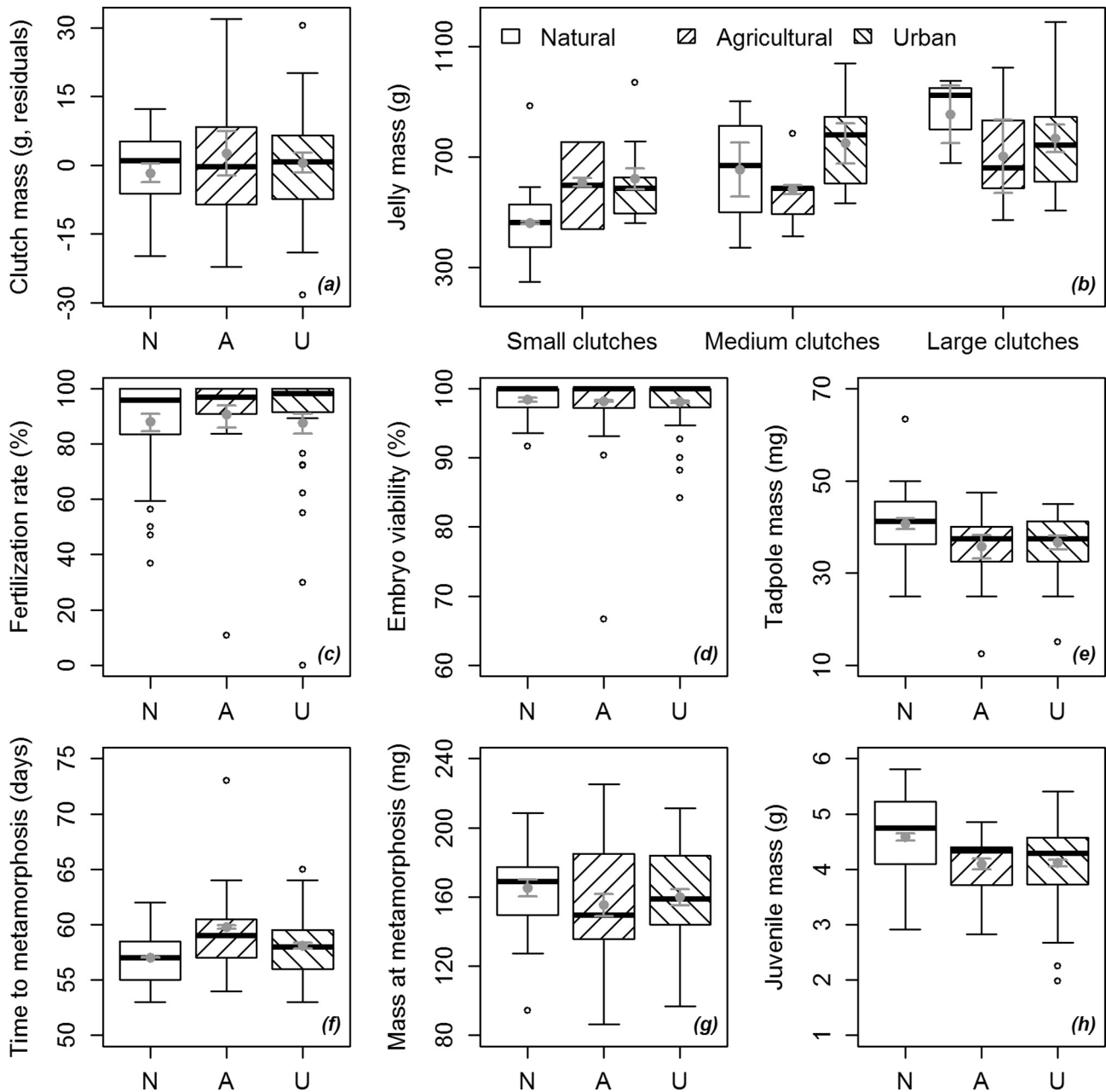


Fig. 2. Reproductive capacity of toad pairs in relation to habitat type (N: natural, A: agricultural, U: urban). Boxplots show the distribution of observed data (thick middle line: median, box: interquartile range; whiskers extend to the most extreme data points within $1.5 \times$ interquartile range from the box); error bars show the mean \pm SE fitted by Generalised Estimation Equations models (see Tables 5–6).

of PAHs (industrial and domestic combustion byproducts) and two phenolics widely used as industrial additives (nonylphenol and bisphenol-A), whereas agricultural land use was associated with higher levels of pesticides (mostly banned organochlorines that were used in large amounts 50 years ago) and natural sex hormones. The latter may be

Table 6

Jelly mass (g) in relation to habitat type (Generalised Estimation Equations model, $n = 90$ clutches).

| Clutch mass | Contrast | Estimate \pm SE | <i>p</i> |
|-------------|------------------------|-------------------|----------|
| Small | Natural – agricultural | -147.0 ± 7.8 | <0.001 |
| | Natural – urban | -158.5 ± 19.0 | <0.001 |
| Medium | Natural – agricultural | 72.7 ± 49.0 | 0.590 |
| | Natural – urban | -94.6 ± 60.6 | 0.531 |
| Large | Natural – agricultural | 152.2 ± 84.7 | 0.363 |
| | Natural – urban | 87.5 ± 58.1 | 0.572 |

due to animal excreta used as fertilizers or originating from livestock grazing (Kolodziej and Sedlak, 2007; Lange et al., 2002) or aquaculture in fish ponds (Barel-Cohen et al., 2006). These pollution gradients demonstrate that aquatic wildlife in anthropogenically influenced habitats have been exposed to higher pollution loads in the recent past than their counterparts in more natural habitats. Since many amphibian species can live as long as 10–15 years in nature (Smirina, 1994) and common toads live up to 6–12 years (Hemelaar, 1988), adverse effects of pollution may be detectable in the reproductive health of amphibian populations even a decade after exposure.

Our results on the toads' reproductive capacities are also complex. On the bright side, pairs from all ponds had high fecundity, fertilization rate and offspring viability under ideal environmental conditions, suggesting that the populations living in urban and agricultural habitats are not suffering from long-term reproductive impairments despite the more frequent occurrence and higher concentrations of EDCs in

these habitats. This means that individual reproductive success and population viability may also be high if acute exposure to pollutants during breeding does not compromise the adults' reproductive output or the offspring's survival and development. Embryos and young larvae can be particularly sensitive to chemical insults (Hoffman et al., 2003; Mikó et al., 2017), so it may be adaptive for females to provide their spawn with extra protection in environments where contamination load is higher. This might explain our finding that female toads from agricultural and urban ponds produced large jelly mass even when clutch size was relatively small, as if the minimum thickness of jelly needed was larger than for females from natural ponds. The jelly coat around the eggs can restrict the uptake of waterborne pollutants and thereby reduce embryo mortality (Edginton et al., 2007; Licht, 1985), so we hypothesize that females living in polluted habitats may produce thick jelly coats as a pre-emptive measure to buffer the effects of expectable contamination events. Interestingly, another environmental stressor, acidification has been found to exert strong local selection on embryonic acid tolerance in frog populations, which is mediated by the jelly's enhanced ability to retain water due to its increased content of negatively charged glycans (Shu et al., 2015, 2016). It would be worth investigating whether chemical pollution favors similar alterations in the macromolecular composition of egg jelly.

On the down side, however, our results indicated reduced performance in the offspring originating from agricultural and urban ponds compared to those originating from natural ponds. Despite being raised in a contaminant-free environment that allowed for maximal investment into development and growth, individuals originating from anthropogenically influenced habitats took longer to complete metamorphosis and had smaller body mass both as larvae and as juveniles. These traits are critical determinants of fitness and population size in amphibians (Wells, 2007): for example, early metamorphosis increases survivorship to maturity by allowing to reach reproductive size earlier (Smith, 1987), and larger juveniles are more likely to survive the first hibernation (Úveges et al., 2016) and maintain their size advantage as adults (Berven, 1990) which in turn affects female fecundity and male mating success (Banks and Beebe, 1986; Davies and Halliday, 1979; Höglund, 1989; Reading, 1986). Thus, our results suggest that the offspring of toads in anthropogenically influenced habitats have reduced chances of becoming successfully reproducing adults. Similarly, recent studies on British populations of the common toad reported that embryos collected from agricultural habitats grew slower in captivity than embryos from reference sites (Orton and Routledge, 2011), and adult males were smaller at higher intensity of anthropogenic land use (Orton et al., 2014).

The lower mass and slower development of toads from anthropogenically influenced habitats may be due to several, mutually non-exclusive phenomena. One possibility is that reduced offspring performance is a lasting consequence of chemical pollution. It is now well established that EDCs can have transgenerational effects, such that early-life exposure of parents leads to adverse health outcomes in offspring and later generations (Anway et al., 2005; Bhandari et al., 2015). Such effects can occur even when the exposed parents show no phenotypic abnormalities (Bhandari et al., 2015). Alternatively, reduced offspring size may be a cost of an adaptive resistance to contaminants. For example, several urban fish populations have evolved tolerance to toxic pollutants (Meyer and Di Giulio, 2003; Whitehead et al., 2012), but their offspring had reduced growth rates in clean water and were more susceptible to other stressors compared with the offspring of conspecifics from a non-contaminated site (Meyer and Di Giulio, 2003). Similarly, in frogs, evolved pesticide tolerance along an agricultural land-use gradient was found to be associated with susceptibility to parasites (Hua et al., 2017). Similar trade-offs may have been responsible for poor performance in the offspring of our toads from anthropogenically influenced habitats. For example, if the latter produce more or different jelly material to provide tolerance against EDCs or other pollutants (see above), they might have less resources to allocate into the eggs (Podolsky, 2004), and their offspring might not be able to

catch up from this initial handicap in egg size or quality (Loman, 2002). Furthermore, because oocytes mature while females are on their post-spawning feeding grounds (Wells, 2007), egg nutrient content may be affected by terrestrial EDC exposure of females; research on such effects on vitellogenesis is virtually absent and highly needed.

Besides pollution, several other environmental factors may differ between anthropogenically influenced and natural habitats, and many of these environmental differences may select for local adaptation in life-history traits. For example, common-garden experiments have shown that tadpole development is faster in populations breeding in temporary pools where desiccation favors earlier metamorphosis than in permanent pools (Lind et al., 2008), and growth efficiency is higher in populations living in harsh environments such as northern latitudes and high altitudes (Lindgren and Laurila, 2005). If anthropogenically influenced ponds are less likely to dry out or if larval competition is low (e.g. due to small population size in fragmented landscapes), this may relax the selection on growth and developmental rates. Also, compared to natural forest habitats, ponds in urbanized and agricultural landscapes may have less closed-canopy vegetation and shade, which would result in higher temperatures, more food for tadpoles, and altered predator fauna, all of which might result in relaxed selection on growth and development speed (Van Buskirk and Arioli, 2005). Disentangling these possible effects of human-induced habitat changes on life-history evolution, and thereby population viability, is an important challenge for ecology and conservation biology.

Taken together, we have documented that small water bodies and private ponds representing typical amphibian breeding habitats contain various EDCs, sometimes in remarkably high concentrations, and show distinct pollution gradients with increasing influence by agriculture and urbanization. This contamination is relevant not only for amphibians but also for other aquatic animals, terrestrial wildlife, and even human health. Although we found no sign of pollution-related reproductive failure in adult toads, our results suggest reduced vigor in offspring originating from anthropogenically influenced habitats. Our study thus highlights that investigating transgenerational EDC effects and resistance to contaminants may be crucially important for furthering our understanding of the consequences of chemical pollution from evolutionary, ecological and conservationist points of view.

Author contributions

VB and AH designed the study. BÜ and VB collected water and sediment samples. OC performed HPLC analyses. BÜ and VV collected adult toads in the field. VB, BÜ, NU, VV and EN tended to the animals in the laboratory. VB, NU and VV digitized the data. NU conducted geoinformatics measurements. VB performed the statistical analyses and wrote the first draft. All authors contributed to writing the manuscript and gave approval of the final version.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found in the online version, at doi:<https://doi.org/10.1016/j.scitotenv.2018.03.363>. These data include Google map of the most important areas described in this article.

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