Use of Biochemical Markers for the Assessment of Organic Pollutant Contamination of the Vltava River, Czech Republic

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Abstract

In this study, selected biochemical markers - cytochrome P450 (CYP 450), ethoxyresorufin-O-deethylase (EROD), glutathione S-transferase (GST) and glutathione (GSH) - were measured in chub (Leuciscus cephalus L.) liver samples obtained from three locations on the Vltava River (Podolí, Podbaba and Vraňany) and from a control location not known as their inducers. These include polychlorinated biphenyls, hexachlorobenzene and octachlorostyrene measured in chub muscle and polyaromatic hydrocarbons measured in bear Vodňany. The levels of selected biomarkers should correlate with the concentration of cheottom sediments obtained from the same locations. The highest EROD activity (median 101.37 pmol·min⁻¹·mg⁻¹ protein), GST activity (median 42.82 nmol·min⁻¹·mg⁻¹ protein), and GSH concentration (median 8.01 nmol·mg⁻¹ protein) were found in fish liver from the Podbaba location. There were no significant differences in CYP 450 level or EROD activity among the different locations. In Podbaba, GST activity (P < 0.01) and GSH concentrations (P < 0.001) were significantly higher than in the control location. The results of these analyses were correlated and a significant correlation was found between biochemical markers and their inducers. The results show that the use of biochemical markers in water quality assessment is a convenient method that can supplement classical chemical monitoring.

Cytochrome P450, EROD, GST, GSH, Leuciscus cephalus, river pollution

The Vltava river (376.7 km) is the longest river in the Czech Republic. The river passes through the capital, Prague, which is the largest metropolitan area in the country with approximately 1.2 million inhabitants. This conurbation is a major source of pollution for the adjacent aquatic environment. The Vltava river, as the major water collector of Prague, presents an ideal scenario to study the discharge of anthropogenic pollutants into the aquatic environment.

There have been many reports of chemical monitoring and aquatic environment pollution assessment carried out on the Vltava River (e.g., Stachel et al. 2003; Žlábek et al. 2005; Hajšlová et al. 2007). To evaluate the relevance of the results obtained by chemical monitoring, it is helpful to assess the effects of anthropogenic pollution on aquatic organisms by measuring specific indicators (biochemical markers). These biomarkers are measurable biochemical indicators, and their level or activity changes are in response to the exposure to substances with similar mechanisms of toxic action. The advantage of analyzing biomarkers lies in their ability to provide comprehensive information on the effects of pollution and also to reflect the synergistic and antagonistic effects of individual components contributing to pollution.
In this study, elements of both phase I (cytochrome P450 and ethoxyresorufin-O-deethylase) and phase II (glutathione S-transferase and glutathione) of xenobiotic metabolism were used for the assessment of aquatic environment pollution in the Vltava river. These biomarkers were also used in previous studies conducted along the Vltava river (Široká et al. 2005; Randák et al. 2006; Havelková et al. 2007).

Cytochrome P450 (CYP 450) is an important biochemical marker of aquatic environment contamination. The most useful isoform of CYP 450 is the 1A subfamily used in water pollution monitoring. The strongest inducers of this isoform are polychlorinated biphenyls (PCB), polycyclic aromatic hydrocarbons (PAH), nitrated polycyclic aromatic hydrocarbons (NPAH) and dioxins (e.g. 2,3,7,8-TCDD) that are commonly found in water (White et al. 1997; Nilsen et al. 1998; Schlenk and Di Giulio 2002). Induction of the CYP1A family happens via the aryl hydrocarbon receptor (AhR). Generally, the toxicity of a pollutant is related to the degree of its affinity to AhR. Following interaction with previously mentioned xenobiotic substances, AhR is carried to the nucleus, where it enhances gene expression in CYP1A and results in an increased synthesis of cytochrome proteins.

The presence of CYP1A is closely associated with the activity of enzyme ethoxyresorufin-O-deethylase (EROD). Enzymatic activity of EROD is highly inducible and an extremely sensitive indicator for the presence of AhR agonists in the environment. Its activity is usually one of the first detectable and quantifiable responses of organisms to the presence of AhR agonists. The strongest and most important inducers of EROD activity seem to be polychlorinated dibenzo-p-dioxins (PCDDs) and dibenzofurans (PCDFs), polychlorinated biphenyls (especially non-ortho-substituted PCBs), polycyclic aromatic hydrocarbons (PAHs) or pesticides (Whyte et al. 2000).

Glutathione and enzyme glutathione S-transferase (both of them involved in phase II of xenobiotic metabolism) were used as well for aquatic environment quality determination. Intracellular tripeptide glutathione (L-γ-glutamyl-cysteinyl-glycine: GSH) represents the most important molecule involved in the defense of cells against oxidative stress (Otto and Moon 1996; Mannervik and Danielson 1988). Catalyzed by an enzyme glutathione S-transferase (GST) conjugative reaction with GSH, this is the principal detoxication pathway for electrophilic xenobiotics. Glutathione S-transferase facilitates the nucleophilic attack of glutathione against an electrophilic part, thus transferring the substrate to the immediate vicinity of glutathione (Eaton and Bammler 1999). The result of this process usually forms a less reactive and more readily soluble product.

Glutathione S-transferase (GST) and GSH can also be used as biomarkers in fish, but their usefulness is less sustainable than in the case of CYP 1A (Van der Oost et al. 2003). The spectrum of potential GST inducers is variable. Increased overall GST activity has been demonstrated in the liver of various fish species after exposure to PCBs (Perez-Lopez et al. 2000; Perez-Lopez et al. 2002), PAHs (Noble et al. 1998; Henson et al. 2001), or pesticides (Frasco and Guilhermino 2002).

The aim of this study was to assess the environmental impact of Prague conurbation, a major source of aquatic pollution, on the Vltava river, using the biochemical markers in chub liver (Leuciscus cephalus L.). The results of these biochemical analyses were compared with the chemical analyses of chub muscle and bottom sediments for specific inducers of the measured biomarkers. Contamination levels of the selected localities were assessed on the basis of the results and the most polluted localities were revealed.

Materials and Methods

This study was conducted in accordance with national and institutional guidelines for the protection of human subjects and animal welfare (the law “Protection of Animals against Cruelty Act” No. 246/92 Collection of Law).
Collection of fish samples
As the indicator-fish species, the chub (*Leuciscus cephalus* L.) was chosen because it represents an omnivorous species and was available at all sampling sites. From 10 to 12 males of chub were caught at each location from April to May 2005 by electrofishing and killed by severing the spinal cord after stunning. Biometric details of the fish are summarized in Table 1. The chubs were caught at three locations along the Vltava river and at the control site (a pond near Vodňany). The locations along the Vltava river were Podolí (upstream of Prague, 57 r. km), Podbaba (downstream of Prague, and downstream of the wastewater treatment plant, 40.3 r. km), and Vraňany (downstream of Prague, and a chemical factory in Kralupy nad Vltavou, 10 r. km). Fish from the ponds near Vodňany (South Bohemia, Czech Republic) were used as control group. Long-term monitoring of the quality of market-ready fish harvested in the Vodňany region has demonstrated only minimum concentrations of exogenous substances in fish from local ponds (Svobodová et al. 2003; Svobodová et al. 2004), thus making this location a suitable control site. The sampling sites are shown in Fig. 1.

<table>
<thead>
<tr>
<th>Location (Date of capture)</th>
<th>n</th>
<th>Mean weight ± S.D. (g)</th>
<th>Age (min-max) (years)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vodňany pond (control locality) (4. 5. 2005)</td>
<td>12</td>
<td>279 ± 144</td>
<td>3–6</td>
</tr>
<tr>
<td>Podolí (upstream of Prague) (25. 4. 2005)</td>
<td>10</td>
<td>331 ± 96</td>
<td>3–5</td>
</tr>
<tr>
<td>Podbaba (downstream of Prague) (29. 4. 2005)</td>
<td>10</td>
<td>290 ± 146</td>
<td>3–8</td>
</tr>
<tr>
<td>Vraňany (downstream of Prague) (29. 4. 2005)</td>
<td>12</td>
<td>718 ± 296</td>
<td>4–8</td>
</tr>
</tbody>
</table>

Fig. 1. Sample-collecting locations along the Vltava river and the control locality

Collection of tissue samples
Immediately after the fish were killed, individual liver samples (approximately 2 g) were collected from each chub for the determination of its biomarkers. Samples were immediately frozen in liquid nitrogen and stored at -80 °C until the enzyme assessments could be performed. Individual samples of fish muscle were collected for chemical analysis of the specific inductors of the mentioned biomarkers. Muscle samples were placed in thermo-boxes filled with dry ice, and stored at -18 °C until they could be analyzed. The PAH levels in bottom sediment at all of the investigated locations were determined in sediment drawn near the river bank at a depth of 0.05–0.15 m by metal core samplers. After the particles larger than 0.5 mm were removed, the sediment samples were homogenized, refrigerated, and then transported (4 °C) to the laboratory, where they were stored at -18 °C (Široká et al. 2005). The collection of bottom sediment samples was carried out according to the international norm ISO 5667-12, which provides guidance on the sampling of sediments from rivers, streams, lakes and similar standing waters and estuarines.

Determination of CYP 450 and EROD activity in liver samples
Before assessing the enzymes, microsomal protein concentrations were determined for each sample according to Lowry (1951). Depending on the protein concentrations, the required amounts of microsomal suspension were taken for the quantification of CYP 450 and the determination of EROD activity.
Before the analysis, a thawed sample of liver tissue (from 0.5 to 2.0 g) was placed into a homogenization buffer (pH 7.4). The homogenized samples were centrifuged at 10 000 g for 20 min at a temperature of 4 °C. The supernatant was then carefully pipetted into ultracentrifugation tubes and re-centrifuged at 100 000 g for 60 min at 4 °C (the ultracentrifuge Beckman OPTIMA - LE - 80 K, USA). This second supernatant was removed and the microsomal fraction was washed with a homogenization buffer twice and then re-suspended in the buffer. This suspension was put into Eppendorf tubes and stored at -80 °C until the enzymatic analysis could be performed. Visible light spectrophotometry (390–490 nm) was used to determine the total quantity of CYP 450. Measurements were made after cytochrome reduction by sodium dithionite, after the complex with carbon monoxide was formed. The suspension (300 μl) was added into 6 ml of phosphate buffer (pH 7.8). A small amount of dithionite was added and the solution was stirred and poured into two 3.5 ml glass cells. The first of these was used as a standard measurement, and the second was bubbled with carbon monoxide for 20 s (Široká et al. 2005). Each sample was analyzed in triplicate and the mean value was used for statistical analysis. Measurements were made using the spectrophotometer UVIKON XS, Secomam, France.

The EROD catalytic activity was determined by fluorescence spectrophotometry (Perkin-Elmer Fluorescence Spectrophotometer 203, USA). The spectrophotometer monochromators were set at 535 nm and 585 nm for excitation and emission wavelength, respectively. In the presence of NADPH (nicotinamide adenine dinucleotide phosphate), EROD enzymatic activity converts the substrate ethoxyresorufin to resorufin, which is a fluorescent product. A phosphate buffer (pH 7.8), NADPH and microsomal suspension adequate for 0.2 mg/ml 1 protein were put into a cell. Ethoxyresorufin was added, and the increase in fluorescence was recorded for 5 min. EROD activity was subsequently calculated based on a comparison of fluorescence to the standard (resorufin) of known concentration. Each sample was analyzed in triplicate and the mean value was used for statistical analysis.

**Determination of GSH and GST in liver samples**

Before the analysis, frozen liver samples were extracted with a phosphate buffer. Four ml of the phosphate buffer (pH 7.2) were added to the samples, which were homogenized and then centrifuged for 10 min at 2 400 g and 4 °C. The resulting supernatant was used for the determination of GST, GSH and protein concentration.

The catalytic activity of GST was measured spectrophotometrically by a modified method of Habig et al. (1974) using a biochemical analyzer Cobas Emira (Roche Diagnostics, Switzerland) at 340 nm. The supernatant with a phosphate buffer (pH 7.2), 0.02 M CDNB (1,2-dichloro-4-nitrobenzene in methanol), and 0.1 M reduced glutathione was pipetted into the cuvette of the biochemical analyzer. Specific activity was expressed as the nmol of the formed product, per minute per milligram of protein. Tripelptide glutathione was determined by Ellman’s method (1959) using the biochemical analyzer Cobas Emira. The amount of 50 μl of 25% trichloroacetic acid was added into the 500 μl of supernatant to precipitate the protein. This was vortexed for 5 min and, after 15 min of incubation at room temperature, it was centrifuged for 10 min at 2 400 g at 4 °C. The supernatant and reaction agents (buffer – 0.8 M Tris/HCl, 0.02 M EDTA pH 7.2 and 0.01 M 2,2-dinitro-5,5-dithiobenzoic acid in ethanol) were pipetted into sample cells. The absorption of coloured product was determined at 414 nm and concentrations (nmol·mg⁻¹ protein) were calculated according to a standard calibration. Protein concentrations were determined by a Bicinchoninic Acid Protein Essay Kit (Sigma-Aldrich) using bovine serum albumin as the standard (Smith et al. 1985). Each sample was analyzed in triplicate (for both GST and GSH) and the mean value was used for statistical analysis.

**Determination of persistent organic pollutants in muscle samples**

Polychlorinated biphenyl (PCB) congeners – IUPAC numbers 28, 52, 101, 118, 138, 153, 180, hexachlorobenzene (HCB), and octachlorostyrene (OCS) were determined in individual chub muscle samples using two-dimensional capillary gas chromatography. The isolation of target analytes from fish muscle was carried out by Soxhlet extraction into a hexanediichloromethane (1:1, v/v) solvent mixture. Purification of the extracts was performed by Gel Permeation Chromatography on a Bio- Beads S-X3 column and mobile phase ethylacetate:cyclohexane (1:1, v/v) (Hajšlová et al. 1995). An HP 5890 Ser. II, gas chromatograph (Agilent Technologies, USA) was used for persistent organic pollutants assessment.

**Determination of PAH in sediments**

Before the assessments, sediment obtained from one location was mixed and a composite sample was prepared. These four sediment samples (200 g of sediment) were dried at 105 °C for 16 h to a constant weight. Five grams of each sample were then mixed with anhydrous sodium sulphate and extracted with 50 ml of a dichloromethane-methanol mixture. The combined extracts were evaporated on a rotary vacuum evaporator until dry, and the residue was dissolved in 4 ml of chloroform. This crude extract was cleaned on the bio-Beads S-X3 column. Elution solvent was evaporated using a rotary vacuum evaporator until dry, and the residue was dissolved in 1 ml of acetonitrile. Determination of PAHs in the purified sample was accomplished by reverse-phase HPLC with the column temperature at 35 °C and the flow-rate of 1.2 ml·min⁻¹. Detection of analyte was carried out by programmable fluorescence detection (Hosnedl et al. 2003). The HPLC system used for the determination of target analytes was composed of the Hewlett-Packard 1050 pumping system, HP 1050 autosampler, and HP 1046 A fluorescence detector.

The PAHs assessed were fluorene, naphthalene, acenaphthylene, phenanthrene, anthracene, fluoranthene, pyrene, benzo(a)anthracene, chrysene, benzo(b)fluoranthene, benzo(k)fluoranthene, benzo(a)pyrene, indeno (1,2,3-c,d) pyrene, benzo (g,h,i) perylene, and dibenzo (a,h) anthracene. These 15 PAHs are European Union priority congeners (EPA 610).
Statistical methods

Because normality of the distribution of the data obtained was not demonstrated, nonparametric techniques were used. The Kruskal-Wallis test was used to compare contaminant concentrations as well as the biochemical markers of contamination found at the individual locations. When the Kruskal-Wallis test showed significant differences between the locations \( (P < 0.05) \), multiple comparisons of all location pairs were subsequently performed. The Spearman’s rank correlation coefficient was used to monitor the relationships between the selected biomarkers and organic pollutants within the various locations.

Results

Biochemical monitoring

The results of the phase I enzymes (CYP 450, EROD) in fish liver from four locations showed no significant differences between the investigated locations. Despite the fact that the highest level of CYP 450 (median 0.18 nmol·mg\(^{-1}\) protein) was found in the Vraňany samples (downstream of Prague conurbation), and the lowest level (median 0.15 nmol·mg\(^{-1}\) protein) was found in Podoli (upstream of the city), the differences in the levels assessed were very low. Increased EROD activity was found in chub liver from Podbaba (median 101.37 pmol·min\(^{-1}\)·mg\(^{-1}\) protein) compared to the activity found in samples from Podolí (median 60.75 pmol·min\(^{-1}\)·mg\(^{-1}\) protein). Just as in the recorded CYP 450 levels, EROD activity levels found in fish liver from the four locations were not significantly different among the sampling sites.

In fact, the results from phase II (enzyme GST and peptide GSH) seem to be more variable. The highest level of GST activity was found in chub liver from Podbaba (median 42.82 nmol·min\(^{-1}\)·mg\(^{-1}\) protein), while the lowest activity was found at the control location, Vodňany (median 12.60 nmol·min\(^{-1}\)·mg\(^{-1}\) protein), as expected. The activity of GST in samples from Podbaba was significantly higher \( (P < 0.01) \) than GST levels recorded at Vodňany, Podolí or Vraňany. As for the GSH concentration in chub liver, the results mimic those of GST activity. The highest GSH concentration was found in fish from Podbaba (median 8.01 nmol·mg\(^{-1}\) protein), and the lowest concentration was found at the control locality Vodňany (median 0.97 nmol·min\(^{-1}\)·mg\(^{-1}\) protein). Glutathione concentrations in livers from Podbaba and Vraňany were significantly higher \( (P < 0.001 \text{ and } P < 0.01, \text{ respectively}) \) than those from Vodňany. Glutathione concentrations in liver from Podbaba were significantly higher \( (P < 0.001) \) than GSH concentrations at Podolí.

The main characteristics of all the measured biomarkers in chub liver are summarized in Fig. 2.

Chemical monitoring

The results of chemical monitoring are summarized in Table 2 and Fig. 2. In all of the studied locations, concentrations of the selected chemical pollutants measured in chub muscle (PCB, HCB and OCS) were significantly higher \( (P < 0.01) \) than those in samples from the control location (Vodňany).

The location most contaminated by PCB was Vraňany, where the median of PCB concentration found in chub muscle samples was 123.10 μg·kg\(^{-1}\) muscle, wet weight (w. w.). The lowest PCB concentration was found at the control location Vodňany (median 8.15 μg·kg\(^{-1}\) muscle, w. w.). The PCB concentrations in the muscle samples of chub from Podolí, Podbaba, and Vraňany were significantly higher \( (P < 0.001) \) than those from the control location Vodňany.

The results of HCB determination in chub muscle obtained from Podbaba and Vraňany have the same value (median 1.45 μg·kg\(^{-1}\) muscle, w. w.), as these two locations present the most contaminated sites. The lowest amount of contamination was found at the control site at Vodňany (median 0.56 μg·kg\(^{-1}\) muscle, w. w.). The HCB concentrations in chub muscle from Podbaba and Vraňany were significantly higher \( (P < 0.01) \) than those from Podolí, and from the control site.
Fig. 2. Biochemical characteristics and POPs concentrations in muscle samples of chub from three locations along the Vltava river and from the control locality Vodňany
Referring to OCS, the situation is very similar to that in PCB; the Vraňany location was confirmed to be the most affected by octachlorostyrene. The levels of OCS in chub muscle samples from this location were 0.20 μg·kg⁻¹ muscle w. w. (median value), and the lowest OCS level was found at the control site V odňany (median 0.03 μg·kg⁻¹ muscle), again. The OCS concentrations in chub muscle from Podolí, Podbaba (P < 0.05) and Vraňany (P < 0.001) were significantly higher than those from the control location.

The concentrations of individual PAH congeners and the sum of all congeners’ concentration in bottom sediments from all four locations are shown in the Table 2. The highest values of Σ 15 PAH were found at Podbaba (12 195 μg·kg⁻¹ dry matter of bottom sediment) and Podolí (10 846 μg·kg⁻¹ dry matter of bottom sediment). PAH concentrations in bottom sediment from Podolí, Podbaba, and Vraňany were significantly higher (P < 0.001 and P < 0.05, respectively) than those from the control location. The most abundant congeners found at the assessed locations were fluoranthene and pyrene.

Correlations between biochemical and chemical monitoring
Spearman rank correlations between biochemical markers and chemical monitoring in the indicator fish are summarized in Table 3. Significant correlations (P < 0.05) were found between the levels of phase I enzymes – CYP 450 and EROD. Significant correlations (P < 0.05) were also found between GST and GSH in chub liver, and persistent organic pollutants in chub muscle samples (PCB, HCB, OCS). Correlations between biochemical markers and PAH in sediment could not be determined, as PAH were assessed only in composite samples, and according to statistics, such data (composite sample versus individual sample) can not be correlated.

Discussion
Enzymes of phase I of xenobiotic detoxification
The biochemical assessment of phase I enzymes (CYP 450 and EROD activity) revealed no significant differences between the four locations. Although certain substances (e.g. PCB, PAH, dioxins or pesticides) have been shown to induce isoform CYP1A and EROD

Table 2. Content of 15 PAHs in bottom sediments

<table>
<thead>
<tr>
<th>PAH compounds*</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Vodňany</td>
</tr>
<tr>
<td>Fluorene</td>
<td>7.3</td>
</tr>
<tr>
<td>Naphthalene</td>
<td>4.4</td>
</tr>
<tr>
<td>Acenaphthylene</td>
<td>5.0</td>
</tr>
<tr>
<td>Anthracene</td>
<td>221.9</td>
</tr>
<tr>
<td>Fluoranthene</td>
<td>146.6</td>
</tr>
<tr>
<td>Pyrene</td>
<td>114.7</td>
</tr>
<tr>
<td>Benzo(a)anthracene</td>
<td>64.6</td>
</tr>
<tr>
<td>Chrysene</td>
<td>59.3</td>
</tr>
<tr>
<td>Benzo(b)fluoranthene</td>
<td>74.3</td>
</tr>
<tr>
<td>Benzo(k)fluoranthene</td>
<td>37.5</td>
</tr>
<tr>
<td>Benzo(a)pyrene,</td>
<td>68.6</td>
</tr>
<tr>
<td>Indeno(1,2,3-c,d)pyrene</td>
<td>71.4</td>
</tr>
<tr>
<td>Benzo(g,h,i)perylene</td>
<td>59.6</td>
</tr>
<tr>
<td>Dibenzo(a,h)anthracene</td>
<td>7.5</td>
</tr>
<tr>
<td>Σ PAH</td>
<td>791</td>
</tr>
</tbody>
</table>

* PAHs are reported as μg·kg⁻¹ dry matter of bottom sediment
activity in fish in both the laboratory and natural conditions (extensive reviews written by Van der Oost et al. 2003 and Whyte et al. 2000), no significant correlation was found between the phase I biochemical markers and the results of chemical analyses in our study. One possible explanation for our non-correlative results could be the differences in the predicate values of various isoforms of CYP 450. Only the total content of CYP 450 was assessed in this study, not specifically the 1A isoform. The response of total CYP 450 content is less sensitive than that in the case of the selected isoenzymes (Bucheli and Font 1995). It was demonstrated that single xenobiotic compounds can act as inducers of specific isoenzymes, but can inhibit others. This may result in a considerable alteration of isoenzyme levels, whereas the amount of total CYP 450 is not always affected.

However, the results of this present study suggest that increased PCB and PAH concentrations do not always result in an increase of CYP 450 levels. Results similar to those found in our study were reported in chub by Široká et al. (2005). In a study of contamination in the Labe and Vltava rivers (Czech Republic), they failed to find any significant increases in the activity of CYP 450 or EROD even at demonstrably increased PCB and PAH concentrations. The contents of both measured biochemical markers and organic pollutants were comparable to our results. On the other hand, Randák et al. (2006) and Havelková et al. (2007) demonstrated a positive correlation between phase I liver enzymes and organic pollutants (PCB) in the muscle of fish from the Vltava river (localities situated upstream and downstream of Prague) in their field studies of chub. The localities situated downstream of Prague showed the highest levels of CYP 450 content and EROD activity. These studies also confirmed the negative impact of the city of Prague on the aquatic environment.

The explanation for CYP 450 levels not responding to contaminants in the aquatic environment might possibly be due to the high concentrations of PCB and PAH substances (Brammel et al. 2004) or the presence of specific CYP 450 inhibitors (e.g. Cu, Zn, Pb, Cd, Cr or Ni) in the environment (Forlin et al. 1986; Fent and Bucheli 1994; Risso-de Faverney et al. 2000; Bozcaarmutlu and Arinc 2004). This inhibitory activity appears to result from competition with iron heme porphyrin binding. Moreover, inhibitors may be selective for a particular CYP 450 isoenzyme. Occasionally, different compounds are able to act as selective inhibitors towards CYP 450 within the same subfamily (Lewis 2001). However, low or increased concentrations of selected organic pollutants (PAH, PCB, TCDD/F) can act as CYP1A inducers, whereas high concentrations of these pollutants can act as inhibitors of the CYP1A subfamily (Stegeman et al. 1997; Schlezinger and Stegeman 2001). Another factor that can influence the levels of measured biomarkers is the duration of exposure of water organisms to an efficient concentration of pollutants. The rapid rise of EROD activity after 1–3 days of exposure is typical for compounds that are easily metabolized (PAH) (Sleiderink and Boon 1996; Wolkers et al. 1996). On the other hand, in the case of halogenated inducers (e.g. PCDD/F or PCB), the induction of

<table>
<thead>
<tr>
<th></th>
<th>CYP</th>
<th>EROD</th>
<th>GST</th>
<th>GSH</th>
<th>PCB (muscle)</th>
<th>HCB (muscle)</th>
<th>OCS (muscle)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CYP</td>
<td></td>
<td>0.344</td>
<td>0.079</td>
<td>0.089</td>
<td>0.086</td>
<td>0.028</td>
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</tr>
<tr>
<td>EROD</td>
<td>0.191</td>
<td></td>
<td>0.253</td>
<td>0.168</td>
<td>0.129</td>
<td>0.209</td>
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<tr>
<td>GST</td>
<td></td>
<td></td>
<td>0.718</td>
<td>0.261</td>
<td>0.478</td>
<td>0.433</td>
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<tr>
<td>GSH</td>
<td></td>
<td></td>
<td></td>
<td>0.390</td>
<td>0.411</td>
<td>0.481</td>
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<tr>
<td>PCB (muscle)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.766</td>
<td>0.907</td>
<td></td>
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<tr>
<td>HCB (muscle)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<td>0.813</td>
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<tr>
<td>OCS (muscle)</td>
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</tr>
</tbody>
</table>
EROD activity or CYP 450 content lasts much longer (from 3 to 16 weeks) (Whyte et al. 2000). Finally, the effects of stress or defense mechanisms which allow the fish to survive in a highly contaminated environment should also be considered.

Enzymes of phase II of xenobiotic detoxification

The results of GST activity and GSH concentration indicate that the localities downstream of Prague are exposed to these quantities of pollutants inducing these markers, including well-known inducers such as PCB and PAH. Chub liver from Podbaba (downstream of Prague) showed approximately 3 times higher amount of GST activity and 10 times higher amount of GSH concentration than fish from the control site. Increased GST activity caused by the presence of PAH and PCB in the aquatic environment has been corroborated by many laboratory and field studies at different rivers involving various fish species (Henson et al. 2001; Perez-Lopez et al. 2002; Pandey et al. 2003; Ahmad et al. 2006). Similarly, increased concentrations of liver GSH compared with the control sites have been observed in English sole (*Pleuronectes vetulus*) collected from a PCB- and PAH-polluted section in Washington (Nishimoto et al. 1995) and in brown bullhead (*Ameiurus nebulosus*) from a PAH-polluted section in New York (Eufemia et al. 1997).

In contrast to a certain number of studies demonstrating increased GST activity in fish from various locations contaminated with organic pollutants (PAH, PCB, PCDD/PCDF), most studies did not show any significant differences between the fish from the control and polluted sites.

Regarding comparison of the sensitivity of phase I and phase II biomarkers, our results showed that phase II biomarkers were more sensitive in this case, which is extremely rare. These biomarkers tested on various fish species have usually proved to be less dependable as an indicator of environmental contamination (Van der Oost et al. 2003). A possible explanation for our differing results may be due to variations in the sensitivity of fish species to the effects of organic contaminants or to the presence of the phase I enzyme inhibitors and GST and GSH inducers at the monitored localities. Sen and Semiz (2007) found that ionic detergents strongly inhibit EROD activity, whereas much less inhibition was observed with GST catalyzed activities. Therefore, this inhibiting potential of detergents suggests that their contribution to the CYP1A induction has to be taken into account for a better interpretation of their environmental impact. The most abundant sources of ionic detergents are from households and sewage water from Prague.

Chemical monitoring

Increasing environmental pollution makes the bottom sediments a “sink” for hydrophobic chemicals, which can then be released back into the water. In this study, we focused on organic pollutants that are typically bio-accumulated in the environment and which can influence biomarkers.

The highest PAH concentrations (a total of 15 PAHs) in bottom sediments were found at Podbaba, which were consistent with the values of GST and GSH found in fish livers from this location. This site is situated downstream of Prague (and downstream of the wastewater treatment plant). The most abundant PAH congeners in bottom sediment samples were fluoranthene, phenanthrene and pyrene. Fluoranthene and pyrene are also the most abundant PAH congeners in the bottom sediment all over the world, e.g. in the Baltic Sea (Pikkarainen 2004), in Hong Kong (Ke et al. 2005) or in the Barents Sea, Russia (Savinov et al. 2003). Fluoranthene is found in many combustible products, along with other PAHs. Even if the highest concentration of fluoranthene in bottom sediment could be determined, this PAH congener seems to inhibit EROD activity, which was demonstrated in various studies on fish *in vivo* (Willett et al. 2001) or *in vitro* (Bols et al. 1999; Fent and Batscher 2000). Because of fluoranthene inhibition effect, CYP1A based bio-assessments may have underestimate the exposure of PAH mixtures.
The highest concentrations of PCB, HCB and OCS were detected at the localities situated downstream of Prague - Vraňany and Podbaba. All of these contaminants are of primary concern due to their persistence and bio-accumulation in the environment, and their toxicity to aquatic organisms. The use of PCBs was very wide (coolants and insulating fluids for transformers and capacitors; additives in PVC coatings for electrical wiring and electronic components; pesticide extenders, cutting oils, flame retardants, hydraulic fluids, adhesives, paints, and in carbonless copy paper). Hexachlorobenzene was used as a fungicide in agriculture, however, the main source of this chemical is presently the industry. Hexachlorobenzene is released into aquatic environment as a byproduct of chlorinated compounds (Bailey 2001). Octachlorostyrene is not commercially manufactured but has been reported to be an inadvertent byproduct of processes that combine carbon and chlorine at high temperatures. The probable source of these compounds could be both the Prague agglomeration and heavy industry near Vraňany (a chemical plant at Kralupy nad Vltavou).

Nevertheless, the highest levels from the majority of assessed biomarkers and chemical pollutants were found in Vraňany and Podbaba. Under normal conditions, levels of pollutants present in wastewaters successively decrease according to the increased distance from the source of contamination. However, this was not the trend in our particular case; therefore, we must presume that pollution in the aquatic ecosystem downstream of Prague is disseminated over a large distance from the Prague agglomeration. Additionally, the negative impact on the aquatic environment could possibly stem from the chemical plant situated near Vraňany, and its influence on the environment deserves more investigation.

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