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# PCBs in fish and their cestode parasites in Lake Victoria

John Oluoch-Otiego · Elijah Oyoo-Okoth · Kipkorir Koross Godfrey Kiptoo · Emily J. Chemoiwa · Charles C. Ngugi · Gelas Simiyu · Elijah S. Omutange · Veronica Ngure · Mary A. Opiyo

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**Abstract** Polychlorinated biphenyls (PCBs) are classified as persistent organic pollutants (POPs) regulated by the Stockholm Convention (2001). Although their production and use was stopped almost three decades ago, PCBs are environmental persistent, toxic, and bioaccumulate in biota. We assessed the levels of 7 PCB congeners (IUPAC

nos. 28, 52, 101, 118, 138, 153, and 180) in sediment and fish (*Oreochromis niloticus*, *Lates niloticus*, and *Rastrineobola argentea*) and evaluated the potential of cestode fish endoparasite (*Monobothrioides* sp., *Proteocephalaus* sp., and *Ligula intestinalis*) as biomonitors of PCBs in Lake Victoria, Kenya. The median concentration of  $\Sigma$ 7PCBs in sediments and fish were 2.2–96.3  $\mu\text{g}/\text{kg}$  dw and 300–3,000  $\mu\text{g}/\text{kg}$  lw, respectively. At all the sampling sites, CB138, CB153, and CB180 were the dominant PCB congeners in sediment and fish samples. Compared to the muscle of the piscine host, *Proteocephalaus* sp. (infecting *L. niloticus*) biomagnified PCBs  $\times$ 6–14 while *Monobothrioides* sp. (infecting *O. niloticus*) biomagnified PCBs  $\times$ 4–8. Meanwhile, *L. intestinalis* (infecting *R. argentea*) biomagnified PCBs  $\times$ 8–16 compared to the muscle of unparasitized fish. We demonstrate the occurrence of moderate to high levels of PCB in sediments and fish in Lake Victoria. We also provide evidence that fish parasites bioaccumulate higher levels of PCBs than their piscine hosts and therefore provide a promising biomonitor of PCBs. We urge further a long-term study to validate the use of the above cestode fish parasites as biomonitoring tools for PCBs.

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**Keywords** PCBs · Bioaccumulation · Parasite · Lipid content · Biomagnification · Lake Victoria

## Introduction

Polychlorinated biphenyls (PCBs) as commercial formulations were employed in the past as dielectric fluids

in power transformers and capacitors, as insulators, coolants, plasticizers in plastic and rubber products, and as hydraulic fluid (Meijer et al. 2003; Johnson et al. 2006). They may be introduced into the environment through accidental leaks and fires in electrical equipment, past disposal in dumps, accidents in transportation of equipment with PCBs, and leakage from hazardous waste sites (Yang et al. 2009). Since they are regarded as persistent organic pollutants (PoP) by Stockholm Convention, they could seriously threaten the environment, animal, and human health (Brown et al. 1994; Berg et al. 2013; Pizarro-Aranguiz et al. 2015; UNEP 2009), depending on their concentrations, the type of PCB congener, and extent of exposure (Zhang et al. 2014). There are concerns about their persistent and toxic nature that led to their eventual ban on production throughout the world in the late 1970s (Breivik et al. 2007). Despite the ban, their presence is still detectable due to their high biostability and hydrophobicity and because they are resistant to both chemical and biological degradation (Ribas-Fito et al. 2001). Also due to the uncontrolled use of the compounds in the industry and agriculture, PCBs are now ubiquitous contaminants in the environmental media worldwide (Meijer et al. 2003; Xing et al. 2005; Yang et al. 2008).

Once in the aquatic ecosystems, small amounts of PCBs may be re-dissolved at the water–sediment interface or incorporated into sediments (Karvonen et al. 2013; Huang et al. 2015; Colombo et al. 2005) and suspended particulate matter (Eisenreich et al. 1989). They can be taken up by sediment-dwelling organisms (McLeod et al. 2008) or build up in aquatic organisms including fish accumulating to higher levels than those in water and sediments (Sures et al. 1999). Bioaccumulation of PCBs in water biota increases over time depending on PCB concentration in the environment and the type of species (Fu and Wu 2006). Fish that are close to the top of the aquatic food web have a relatively long life span and concentrate high amounts of PCBs (Brázová et al. 2012b). Hence, the concentration of PCBs accumulated in fish may be used as a good tool to assess the degree of PCB pollution of the environment (Fang et al. 2009; Hu et al. 2009; Brázová et al. 2012b).

Parasite infection is potential stressor to organisms and may disrupt a number of physiological processes in their host (Frank et al. 2013). The interactions between parasites and their fish hosts have attracted increasing

interests from ecological viewpoint (see reviews in (Oyoo-Okoth et al. 2010b; Lima et al. 2012; Karvonen et al. 2013; Seppälä et al. 2009; Bellay et al. 2015), while studies on the interaction between parasites, their hosts, and pollutants have generated information suitable for developing sentinels for contaminant biomonitoring (Marcogliese and Pietrock 2011; Sures 2007; Oyoo-Okoth et al. 2010a, b). Fish parasites have also previously been used in monitoring inorganic pollution (e.g. see (Sures 2001; Huang et al. 2015; Sures et al. 1999), but research on the accumulation of organic compounds, including PCBs (e.g., Brázová et al. 2012a), is scant in parasitic organisms (Sures 2004, 2008; Marcogliese and Pietrock 2011; Le et al. 2014; Morrill et al. 2014). To this purpose, the aim of this study was to assess the levels of PCBs in sediments and commercial fish species and also evaluate the potential of fish parasites as bioindicator of PCBs in Lake Victoria, Kenya. The host–parasite assemblages from different sites in the lake were analyzed to characterize spatial changes in the ability of the parasites and their hosts to accumulate PCBs.

## Materials and methods

### Study area

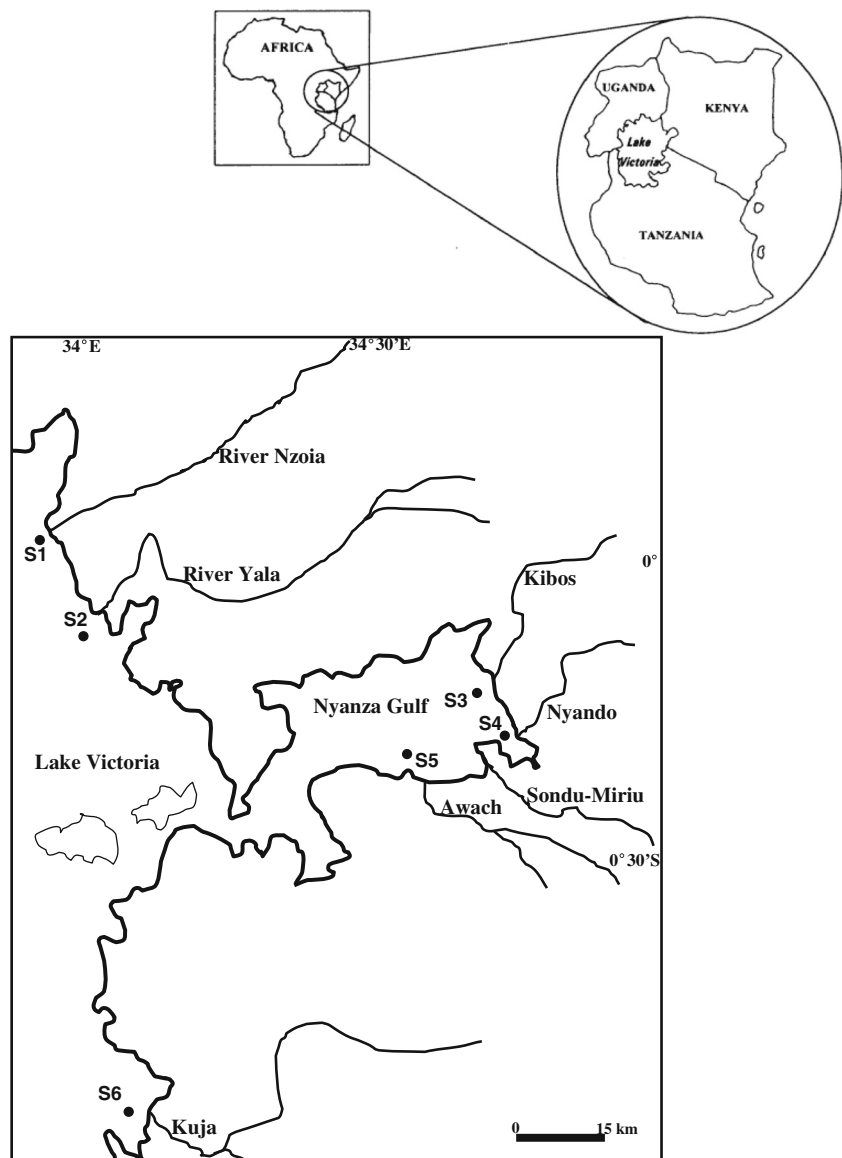
Lake Victoria is the largest tropical lake in the world, having a surface area of about 68,800 km<sup>2</sup> and a maximum depth of about 70 m. The lake is situated at an altitude of 1,134 m above sea level and is bordered by Tanzania, Kenya, and Uganda (Fig. 1). The mean annual ambient temperatures in areas adjacent the lake range between 20 and 34 °C. Bimodal rainfall pattern is experienced in the area; long and heavy rains (1,200 to 1,600 mm) occur from March to May and the short rains (350 to 550 mm) from November to December. There are three commercially exploited fish species: Nile tilapia (*Oreochromis niloticus*), Nile perch (*Lates niloticus*), and silver sardine (*Rastrineobola argentea*). Six sites that receive water from the inflowing rivers from the Kenya side of the lake were selected for this study (Fig. 1). Human activities that include fishing, agriculture, livestock grazing, sugarcane farming, and industrial activities in towns at these sites are described elsewhere (Oyoo-Okoth et al. 2012).

### Sampling strategies

Fieldwork was carried out between 3rd February to 15th November 2014 covering the dry and rainy season at each of the six selected sampling sites (Fig. 1). Most of the sampling work was conducted in the morning between 7 am to 11 am. On each sampling date: sediment, fish, and fish parasite samples were collected. Sediments were collected at the surface (up to 20 cm) using Ekman Grab Sampler. The sediments were transferred to marked polythene bags, placed in an icebox ( $\sim 4^{\circ}\text{C}$ ) and transported to the laboratory within 30 min for

analyses. In the field, fish were sampled using pelagic gill nets (mesh size 0.5") and rinsed (ultrapure water) and carefully dissected dorso-ventrally using stainless steel instruments for parasitological examinations of endoparasites. Each dissection instrument was used for specific fish. The internal organs were examined for endoparasites under a dissecting and light microscope. The fish muscles were trimmed carefully to expose any embedded parasite cysts. The observed parasites were isolated, counted and each transferred to a labeled vial, then fixed in 70 % alcohol and sent to the Department of Parasitology, University of Eldoret for identification.

**Fig. 1** Map of the Nyanza Gulf in Lake Victoria, Kenya showing the location of the sampling sites



The cestodes were identified morphologically using standard identification keys and pictorial guides, e.g., Yamaguti (1959), Colombo et al. (2005), Sures (2007), and Xing et al. (2005). Data of the fish and parasite samples collected from the six sampling sites of Lake Victoria is presented in Table 1. Other parasites occurred in fish but their prevalence were low (< 2 %) while the cestode parasites that infected one species was not found in the other fish species (i.e., parasite–host specificity). The cestodes for PCB analysis were freeze dried before analysis. Fish which did not have any parasite after field observations were placed into separately marked polythene bags, stored in ice boxes, and transported to the laboratory for further chemical analyses.

### Chemical analysis

All samples were freeze dried prior to analysis. In the sediment, the PCBs were measured in the freeze-dried

samples; while in fish and fish parasites, the PCBs were measured in the lipid-extracted fractions of the samples. Extraction of lipids followed methods described in (Campbell et al. 2004) with slight modification. Briefly, 2 g of the freeze-dried sample (sediment, fish, or fish parasites) was ground with 10 g of anhydrous sodium sulfate using a mortar and pestle. The samples were then extracted on an accelerated solvent extractor device (ASE 300, Dionex, USA) using 1:1 v/v mixture of *n*-hexane/dichloromethane. For the sediments, activated copper granules were added to the extract to remove elemental sulfur before purification with column chromatography. The extract was concentrated to 2 mL using a rotavapor. The concentrated extract was divided into two sub-samples for fish (0.3 and 1.7 mL). The 0.3 mL was used for gravimetric determination of lipid content while the remaining part was kept for clean up. The extract was cleaned in a silica column to make the extract suitable for injection into the gas chromatograph

**Table 1** Data of the fish and parasite samples collected from the six sampling sites of Lake Victoria (mean values  $\pm$  standard error)

Fish/fish parasites		Sampling sites					
		Site S1	Site S2	Site S3	Site S4	Site S5	Site 6
<i>Oreochromis niloticus</i>	Number of fish sampled	189	201	184	192	188	205
	Mean length (cm)	40.3 $\pm$ 9.2	43.8 $\pm$ 11.3	37.2 $\pm$ 7.1	38.3 $\pm$ 8.3	41.2 $\pm$ 10.2	43.6 $\pm$ 9.9
	Mean dry weight (g)	306.1 $\pm$ 45.3	420.2 $\pm$ 67.8	314.3 $\pm$ 60.2	355.6 $\pm$ 63.2	398.3 $\pm$ 55.2	415.4 $\pm$ 70.2
	Prevalence of <i>Monobothrioides</i> spp. (%)	23.3	22.4	20.2	24.1	21.8	20.3
	Number of fish analyzed	12	12	12	11	12	12
<i>Lates niloticus</i>	Number of fish sampled	133	136	125	144	119	116
	Mean length (cm)	50.2 $\pm$ 11.3	55.3 $\pm$ 19.2	48.3 $\pm$ 10.1	49.3 $\pm$ 12.3	52.3 $\pm$ 13.2	56.4 $\pm$ 14.3
	Mean dry weight (g)	401.2 $\pm$ 70.3	459.4 $\pm$ 81.2	435.6 $\pm$ 65.6	430.4 $\pm$ 61.4	475.6 $\pm$ 69.3	487.2 $\pm$ 90.2
	Prevalence of <i>Proteocephalus</i> spp. (%)	37.6	40.4	40.0	37.2	37.8	42.8
	Number of fish analyzed	11	12	12	12	12	12
<i>Rastrineobola argentea</i>	Number of fish sampled	234	142	201	197	210	219
	Mean length (mm)	37.9 $\pm$ 9.4	41.3 $\pm$ 11.2	36.4 $\pm$ 9.1	36.4 $\pm$ 10.2	37.9 $\pm$ 8.9	40.4 $\pm$ 10.1
	Mean dry weight (g)	0.54 $\pm$ 0.21	0.58 $\pm$ 0.22	0.51 $\pm$ 0.31	0.49 $\pm$ 0.23	0.54 $\pm$ 0.19	0.56 $\pm$ 0.28
	Prevalence of <i>Ligula intestinalis</i> (%)	28.6	34.5	24.9	23.4	21.4	23.7
	Number of fish analyzed	14	12	14	13	12	12
<i>Monobothriodes</i> spp.	Number analyzed (n)	15	14	15	14	15	15
	Mean abundance in fish	4.45 $\pm$ 1.16	5.01 $\pm$ 2.42	4.31 $\pm$ 1.49	2.45 $\pm$ 1.19	3.83 $\pm$ 1.56	3.23 $\pm$ 1.99
<i>Proteocephalus</i> spp.	Number analyzed (n)	11	12	12	12	12	12
	Mean abundance in fish	3.65 $\pm$ 1.19	4.21 $\pm$ 1.82	4.31 $\pm$ 1.09	3.45 $\pm$ 1.31	2.44 $\pm$ 1.41	3.23 $\pm$ 1.92
<i>Ligula intestinalis</i>	Number analyzed (n)	15	15	15	14	15	15
	Mean abundance in fish	1.45 $\pm$ 0.16	2.41 $\pm$ 0.82	2.31 $\pm$ 0.09	2.45 $\pm$ 1.21	2.43 $\pm$ 0.92	1.23 $\pm$ 1.02



(GC). The column contained alternating layers of neutral and acidic silica gel. The column was pre-eluted with hexane, the sample was applied, and then the PCBs were eluted with hexane. Finally, the sample was concentrated to a final volume of 100  $\mu\text{L}$ .

The 7 “indicator” congeners (IUPAC nos. 28, 52, 101, 118, 138, 153, and 180) have been chosen on the basis of their persistence in food web and their tendency to biomagnification. The analysis of 7 indicator PCBs (IUPAC numbers 28, 52, 101, 118, 138, 153, and 180) was done using high-resolution gas chromatography coupled with high-resolution mass spectrometry (HRGC/HRMS; AutoSpec Ultima, Waters, USA). Chromatographic separation was achieved by injecting 1  $\mu\text{L}$  of sample on a fused silica capillary column (DB5MS, 60 m  $\times$  0.25 mm i.d.  $\times$  0.25  $\mu\text{m}$  film thicknesses). The oven temperature of the GC was maintained at 120  $^{\circ}\text{C}$  for 1 min, then increased to 150  $^{\circ}\text{C}$  at a rate of 30  $^{\circ}\text{C min}^{-1}$ , and finally, to 300  $^{\circ}\text{C}$  at 2.5  $^{\circ}\text{C min}^{-1}$ . Helium was used as a carrier gas at a flow rate of 1.0 mL/min.

#### Quality assurance and quality control

To ensure the quality of data, surrogate PCB standards, blanks, replicates, and a certified reference material (NMIJ CRM 7404-a, Japan) were included in the analysis. The PCBs reference standards and other organic solvents and chemicals were of analytical grade and purchased from commercial suppliers. Recoveries for majority of the surrogate standards met the requirements of US EPA methods 1668A (were in the range of 80–117 %). The recoveries resulting from triplicate determinations ( $N = 3$ ) of the certified reference material were between 81 and 112 % for the PCB congeners. The limit of detection (LOD) which was calculated as three times the signal-to-noise ratio, varied from 0.04 to 0.32 ng/kg for sediments and 0.02 to 0.60 ng/kg for fish, and 0.03 to 0.56 ng/kg parasites.

#### Biomagnification (BMF) of PCBs in fish parasites

The biomagnification factor (BMF) of the PCBs in the parasite tissue was calculated according to the formula: Adopted from (Sures et al. 1999).  $\text{BMF} = \text{Concentration of the PCBs in the parasite tissue} / \text{Concentration of the PCBs in the fish muscle}$ .

#### Data analysis

All statistical analyses were performed with STATISTICA 10.0 (StatSoft, Inc., Tulsa, OK, USA). Data on PCBs concentration are presented as mean  $\pm$  SD per site. One-way ANOVA was used to compare concentrations among sites. For each tested data set, the assumption of normality prior to ANOVA was verified using the Shapiro–Wilk test. Significantly different means were analyzed by post hoc Tukey’s HSD test. Relationship between PCBs concentration in sediment and fish were analyzed using Pearson’s correlation. Significance was declared at  $P < 0.05$  for all analysis.

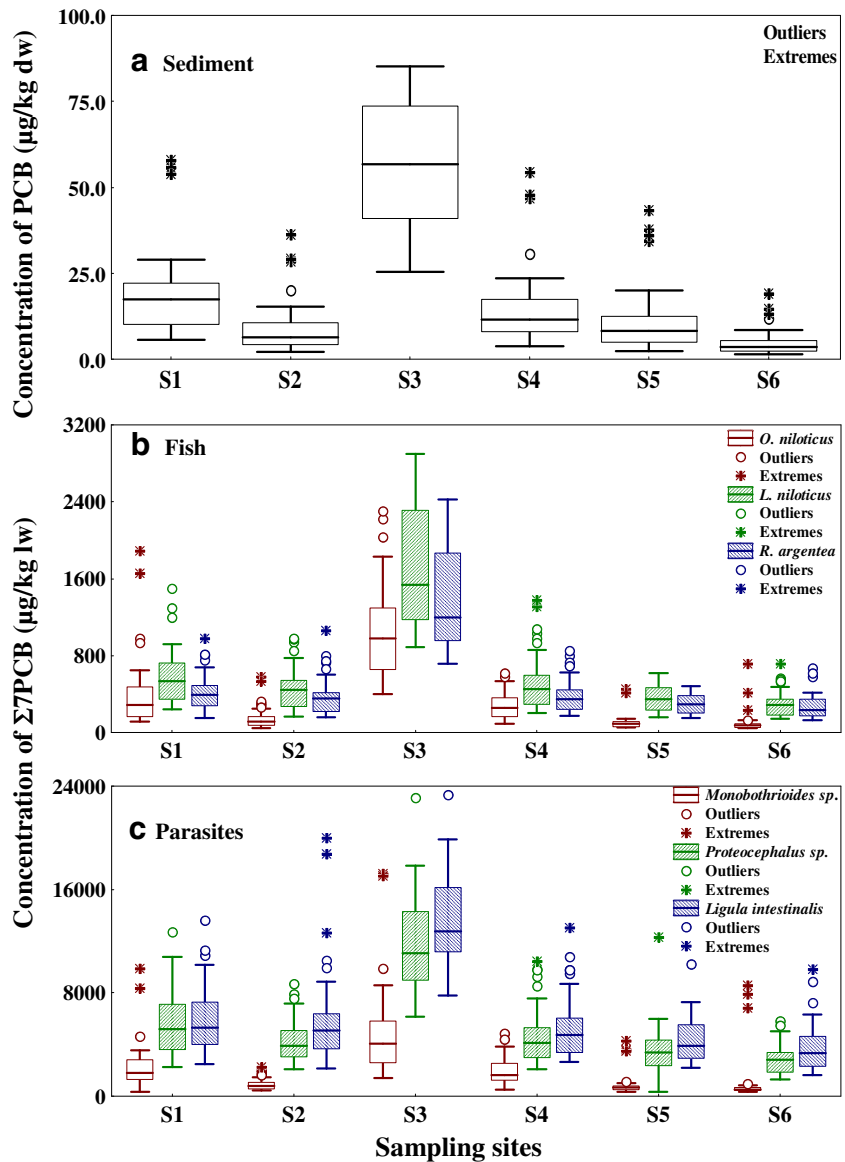
#### Results

Lipid content of freeze-dried wet weight of the fish were: (mean  $\pm$  SEM [range]): *O. niloticus* 3.2  $\pm$  0.4 [2.9–3.7]%, *L. niloticus* 5.9  $\pm$  1.1 [5.2–6.4]%, and *R. argentea* 4.9  $\pm$  0.6 [4.8–5.6]% while the lipid content of the parasites were: *Monobothrioides* sp. 4.4  $\pm$  0.6 [4.1–4.7]%, *Protocephalus* sp. 5.8  $\pm$  0.9 [5.8–6.4]%, and *Ligula intestinalis* 6.1  $\pm$  0.8 [5.5–6.7]%. The prevalence of *Monobothrioides* sp. ranged from 20.2 to 24.1 % in *O. niloticus*, *Protocephalus* sp. ranged from 37.2 to 42.8 %, while for *L. intestinalis* the prevalence was 21.4–34.5 %.

Concentrations of  $\Sigma 7\text{PCBs}$  in sediments, fish, and parasites samples at the six sites is presented in Fig. 2. The mean concentration of  $\Sigma 7\text{PCBs}$  in sediment (a) ranged from 2.2 to 96.4  $\mu\text{g/kg dw}$  (Fig. 2a). There was a significant spatial variation of PCBs in sediments ( $P < 0.001$ ). S3 had the highest concentration of PCBs (mean = 73.5  $\mu\text{g/kg dw}$ ; 31.2–96.4  $\mu\text{g/kg dw}$  range). The PCB levels at S1 (mean = 24.7  $\mu\text{g/kg dw}$ ; 17.3–28.7  $\mu\text{g/kg dw}$  range) followed that of S3. Site S6 (mean = 4.6  $\mu\text{g/kg dw}$ ; 2.2–8.1  $\mu\text{g/kg dw}$  range) had the lowest sedimentary PCB levels.

Concentration of PCBs in fish (b) ranged from 300 to 3,000  $\mu\text{g/kg lw}$  (Fig. 2b). There were notable significant spatial differences in PCB levels ( $P < 0.001$ ); S3 had 2–3 $\times$  higher concentration of PCBs (mean = 1450  $\mu\text{g/kg lw}$ ; 600–3,000  $\mu\text{g/kg lw}$  range) compared to other sites (300–550  $\mu\text{g/kg lw}$  range). The pattern of PCB concentration at all the sampling sites followed the pattern: *L. niloticus* > *R. argentea* > *O. niloticus*. PCB levels in parasites (c) ranged from 1,200 to 18,000  $\mu\text{g/kg lw}$ .

**Fig. 2** Box plots of mean  $\pm$  SD of PCBs concentration in: **a** surface sediments ( $\mu\text{g}/\text{kg dw}$ ), **b** fish ( $\mu\text{g}/\text{kg lw}$ ), and **c** fish parasites ( $\mu\text{g}/\text{kg lw}$ ) at six sampling sites of Nyanza Gulf in Lake Victoria. Number of samples per station is presented in Table 1

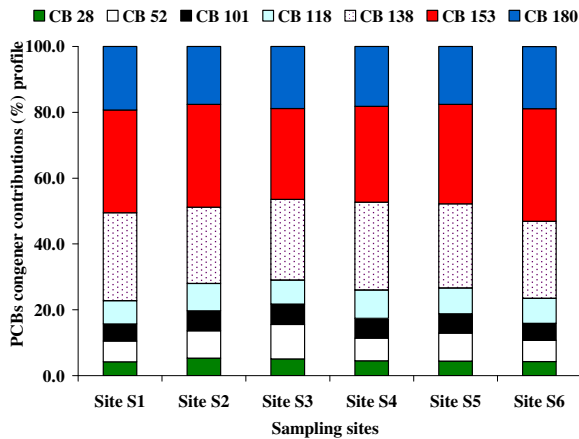


Again there were discernable spatial differences ( $P < 0.001$ ), parasite at S3 had the highest concentration of PCBs (mean = 13,500–14,500  $\mu\text{g}/\text{kg lw}$ ; 6,000–18,000  $\mu\text{g}/\text{kg lw}$  range). The correlation between PCBs in sediments and in fish was significant for all species of fish ( $r > 0.82$ ,  $P < 0.005$ ). Accumulation of PCBs in *L. intestinalis* (parasite infecting *R. argentea*; mean = 12,000  $\mu\text{g}/\text{kg lw}$ ; 6000–16,000  $\mu\text{g}/\text{kg lw}$ ) was in similar ranges ( $P > 0.05$ ) to accumulation of PCBs in *Protocephalus sp.* (parasite infecting *L. niloticus*; mean = 12,000  $\mu\text{g}/\text{kg lw}$ ; 4,500–13,500  $\mu\text{g}/\text{kg lw}$ ) while *Monobothrioides sp.* (infecting *O. niloticus*) had the

lowest PCB accumulation (mean = 5,400  $\mu\text{g}/\text{kg lw}$ ; 3,000–7,000  $\mu\text{g}/\text{kg lw}$ ).

We established similar pattern of PCB congeners in sediments, fish, and in parasites. The general concentration patterns (calculated as % of the  $\Sigma 7\text{PCBs}$ ) is shown in Fig. 3 for the sediment and Fig. 4 for the fish and fish parasites. At all the sampling sites, CB138 (25–32 %), CB153 (27–35 %), and CB180 (17–25 %) were the dominant PCB congeners in sediment, fish, and parasites. Other congeners (CB28, CB52, CB101, and CB118) occurred at concentrations that were  $< 10$  % each of the congeners at all the sampling sites.





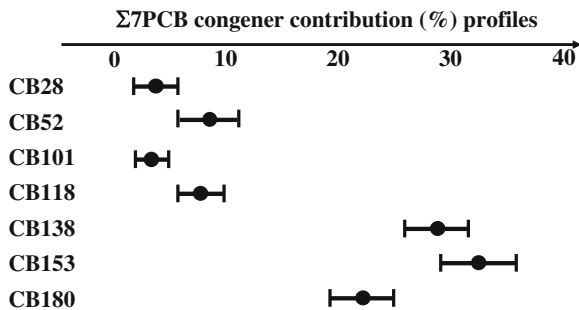
**Fig. 3** Pattern of PCB congener contributions (%) profile in surface sediments at the six sampling sites of the Nyanza Gulf in Lake Victoria, Kenya

The BMF of PCBs in parasite compared to the con-specific fish species is presented in Fig. 5. The fish parasites biomagnified PCBs  $\times 4\text{--}16$  without any noticeable spatial differences. Accumulation of PCBs in *L. intestinalis* (parasite infecting *R. argentea*; BCF  $\sim \times 8\text{--}16$ ) was significantly ( $P < 0.05$ ) higher compared to bioaccumulation in *Proteocephalus* sp. (BCF  $\sim \times 6\text{--}15$ ) and *Monobothrioides* sp. (BCF  $\sim \times 4\text{--}8$ ).

**Discussion**

PCBs in sediments

The concentration of  $\Sigma 7$ PCBs in sediments were higher than those measured in River Pangani, Tanzania (1.3–7.0  $\mu\text{g}/\text{kg dw}$ ; Hellar–Kihampa et al. 2013), Murchison Bay, Lake Victoria, Uganda (0.41–4.652  $\mu\text{g}/\text{kg dw}$ ;



**Fig. 4** General pattern of PCB congener contribution (%) profiles in fish and fish parasites at the six sampling sites of the Nyanza Gulf in Lake Victoria, Kenya. Horizontal bars represent ranges (in %) of congeners relative to  $\Sigma 7$ PCBs

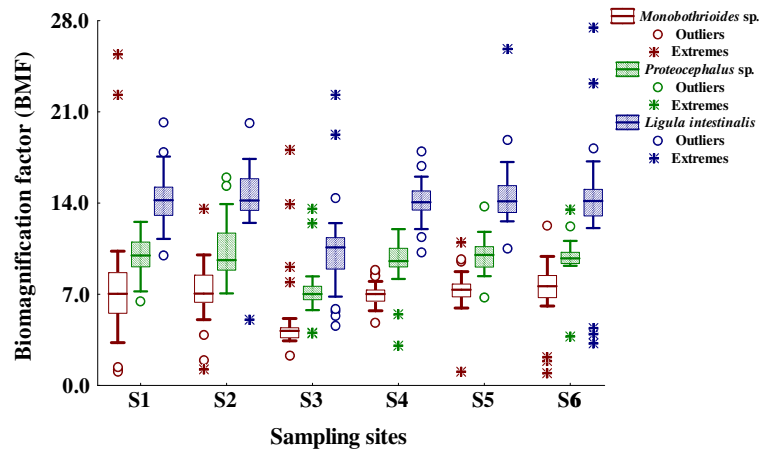
Ssebugere et al. 2014b), and Pearl River Delta, South China (0.43–1.77  $\mu\text{g}/\text{kg dw}$ ; Wang et al. 2011). These values were also consistently very high compared to River Congo, DRC ( $< 0.005\text{--}0.65 \mu\text{g}/\text{kg dw}$ ; Verhaert et al. 2013), in the Napoleon Gulf of Lake Victoria, Uganda (0.04–1.02  $\mu\text{g}/\text{kg dw}$ ; Ssebugere et al. 2014a), and those from Guanabara Bay (0.018–0.184  $\mu\text{g}/\text{kg dw}$ ; de Souza et al. 2008). PCBs in contaminated site such as S3 were comparable to values reported in Dagu Drainage River in China (9.687–22.148  $\mu\text{g}/\text{kg dw}$ ; Liu et al. 2007) and Lake Qarun in Egypt (1.48–137.200  $\mu\text{g}/\text{kg dw}$ ; Barakat et al. 2013) but were lower than the upper limits of PCBs in Cortiou, Marseille in France (11.2–1,412.2  $\mu\text{g}/\text{kg dw}$ ; Wafo et al. 2006).

The rivers that discharge into the *L. victoria* are located near urban centers with numerous small-scale motor vehicle repair workshops releasing contaminants such as oils and paints which may release contaminants into the rivers draining Lake Victoria. Currently, there is very little enforcement of the regulations on disposal of the wastes into the nearby water bodies (personal observation). Possible PCB contamination sources at the sampling sites also include leakages of contaminated fluids from the electric transformers, motor vehicle lubricants, and heat transfer fluids associated with transportation activities (Ssebugere et al. 2014a). Sites S1, S3, S4, and S5 are also located in catchment areas that practice biomass burning of sugarcane and agro-industrial discharge from sugar industries which is done almost every harvesting season and may be potential source of PCBs (Eckhardt et al. 2007). Site S3 had the highest concentration of PCBs since it is located downstream of urbanized Kisumu City with several chemical manufacturing factories, agro-industries, industrial waste treatment plants, and municipal solid waste incinerators. Its close vicinity to industrial facilities where ships dock are other potential sources of PCBs. The current results indicate higher concentration of PCBs in sediments influenced by human activities in the catchment areas.

PCBs accumulation in fish

In commercial fish species, the PCB ranges (300–3,000  $\mu\text{g}/\text{kg lw}$ ) were higher than the  $\Sigma 7$ PCBs concentration in fish from freshwater bodies in Africa such as Lake Tanganyika (24.3–77.7  $\mu\text{g}/\text{kg lw}$ ; Manirakiza et al. 2002), Eastern Slovakia (mean = 108  $\mu\text{g}/\text{kg lw}$ ; Brázová et al. 2012a), and Van Region, Turkey

**Fig. 5** Box plots of mean  $\pm$  SD of biomagnification factor (BMF) of PCBs in fish parasites relative to fish at the six sampling sites of the Nyanza Gulf in Lake Victoria, Kenya



(<LOD – 277  $\mu\text{g}/\text{kg}$  lw; Aksoy et al. 2011). The piscine levels of PCBs were systematically much higher than fish species sampled from upper River Thames (<0.77–3.32  $\mu\text{g}/\text{kg}$  lw; Yamaguchi et al. 2003), Yongxin Island, South China Sea (6.3–89  $\mu\text{g}/\text{kg}$  lw; Sun et al. 2014), and Napoleon Gulf, Lake Victoria, Uganda (0.042–0.778  $\mu\text{g}/\text{kg}$  lw; Ssebugere et al. 2014a). The PCBs were comparable to fish species from Negro Basin, Argentinean Patagonia (24.3–77.7  $\mu\text{g}/\text{kg}$  lw; Ondarza et al. 2014), European eel (*Anguilla anguilla*) from Loire Estuary (80–4,500  $\mu\text{g}/\text{kg}$  lw; Blanchet–Letrouvé et al. 2014), and wild brown trouts (*Salmo trutta*) sampled from Marche rivers, Central Italy (80–3,100  $\mu\text{g}/\text{kg}$  lw) (Frank et al. 2013). The spatial patterns of PCBs were consistent with PCBs analyzed in the sediments; thus, it is possible that exposure to PCB-contaminated environment via direct contact or ingestion of sediment-dwelling organisms may constitute a major exposure route for fish. When analyzing these differences, one must take into account the trophic niche of the three species (Ojwang et al. 2004; Campbell et al. 2004), since they show different feeding habits, and thus, different positions in the trophic web.

In the present study, PCBs were higher in *L. niloticus*, a carnivorous species when compared to omnivorous *O. niloticus* and *R. argentea*. The present observations suggest that these compounds can biomagnify throughout the food web to higher levels in the top predators. The higher PCB accumulated by *L. niloticus* than other fish species therefore appear related to the diet in its trophic position. *L. niloticus* consume all available fish species including its own siblings, insects, crustacea, and molluscs whereas *O. niloticus* and *R. argentea* rely on zooplankton and macroinvertebrates. The high levels

of PCB 138, 153, and 180 most likely demonstrate that more of these congeners were released into the environment. Also, the high degree of chlorination resulting in lower chemical degradation rates may make them be retained in aquatic systems to a greater degree, resulting in bioaccumulation in sediments and biota (Lavandier et al. 2013). These results indicate considerable quantities of PCBs in the environment and therefore, there is a need to search for suitable tools to monitor the PCB concentrations.

#### PCBs accumulation by fish parasites

Previous study in Lake Victoria have identified that the populations of *R. argentea* in Lake Victoria, exhibit a high degree of infestation with the tapeworm, *L. intestinalis*, a cestode of the family Diphyllbothriidae (Oyoo-Okoth et al. 2010a, c). Information on the cestode parasitic infections of *O. niloticus* and *L. niloticus* are absent. In the current study, the prevalence of *Monobothrioides* sp. occurred in high abundance in *O. niloticus* samples (20.2 to 24.1 %), *Protocephalus* sp. in *L. niloticus* (37.2–42.8 %), and *L. intestinalis* in *R. argentea* samples (21.4–34.5 %; Table 1). Numerous studies have highlighted parasites as excellent bioaccumulators of inorganic pollutants such as metals compared to their fish hosts. This has often been achieved by calculating BMF, reflecting the ratio between the concentrations in the parasite and in the host. Yet the few studies that have attempted to measure organic compounds such as PCBs bioaccumulation in parasites compared their fish hosts. The studies available have established that these contaminants are lower in parasites than that in host tissues when

measurement are done as dry weight or wet weight basis, e.g., Brázová et al. (2012a) and Le et al. (2014).

In this study, we used lipid-normalized concentrations of PCBs in the host and compared with that of the cestodes to establish the potential role of parasites as bioaccumulators of PCBs. The cestode infecting the three different fish species contained higher concentration of PCBs than their host by a BMF ranging from  $\times 4$  to 16 that was consistent among the sampling sites. The present result where PCBs are higher in parasites than in the hosts are consistent with studies of PCB accumulation in perch (*Perca fluviatilis*) compared to their parasites (*Acanthocephalus lucii*; Brázová et al. 2012a) but are inconsistent with studies of the accumulation of PCBs in salmon (*Salmo salar*) infected with the tapeworm *Eubothrium crassum* (Persson et al. 2007). A characteristic feature of all cestodes is the absence of an alimentary canal (Karvonen et al. 2013). The lack of an alimentary tract means that substances enter the cestodes across the tegument. This structure is well adapted for transport functions, since it is covered with numerous microvilli resembling those lining the lumen of the mammalian intestine (Bellay et al. 2015). It has also been hypothesized that because cestodes cannot synthesize their own cholesterol and fatty acids, they efficiently obtain them from their host's intestinal lumen (Yang et al. 2009). Absorption of PCBs with the content of the host intestinal lumen is thus probable.

In studies of metal accumulation, a biokinetic model that explores the variability in metal accumulation in situ has been applied to explain the variable bioaccumulation in organisms (McLeod et al. 2008). According to the model, uptake and depuration parameters can be estimated experimentally to provide testable predictions. We previously showed that *R. argentea* had lower uptake rate constant for cadmium which was apparently sequestered well, while *L. intestinalis* had higher uptake rates of metals and low depuration rate resulting to accumulation of higher amounts of Cd in parasites than the fish host (Johnson et al. 2006). Currently, we lack information on the uptake and depuration kinetics of PCBs in fish and their respective parasites, and thus, we have very little knowledge on the mechanistic explanation of high bioaccumulation of PCBs in different fish species and parasites to warrant any conclusion. The current results indicate higher concentration of PCBs in parasites compared to the muscle of the fish host and offer a possibility of studying further the prospect of developing parasite-fish model for biomonitoring PCBs.

## Conclusion

In conclusion, this study provide data indicating that the PCBs levels in Lake Victoria are moderate to high and in some sites, it is comparable with ranges of the PCB values found in other places of the world. The high PCB in the sediments due to increase from diverse human activities are taken up by the fish and their parasites. Comparatively, fish endoparasites biomagnified PCBs than levels in the fish hosts. For this, cestode parasites are potential biomonitors of the degree of accumulation of PCBs in the aquatic environment and show corresponding increase with levels of PCBs in sediments and fish.

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