The major and trace element chemistry of fish and lake water within major South African catchments

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ABSTRACT

Chemical elements in lake water are incorporated into fish tissues through bioconcentration and biomagnification. Lake water and fish tissue samples from 23 lakes, located within 4 major South African catchments, were analysed to investigate the link between element concentrations in lake water and otolith, fin spine, muscle, liver and gill tissues. The comparison is complicated by the seasonal variation in water chemistry as well as the large natural variation between individual fish within a lake. Comparisons between fish from different lakes can also only be done within the same species, which may not occur within all the lakes within the project area. This may be further complicated by erratic anthropogenic contamination. It is therefore more successful to use inter-element ratios for comparison than absolute element concentrations. Using the Sr/Ca elemental ratio, a species-specific correlation was identified between lake water, otolith, fin spine and gill tissue samples. The best discrimination between fish from different lakes was achieved using a Na/Ca versus Mg/Ca elemental ratio diagram of gill tissues. The best discrimination between fish from different lakes was achieved using a Ba/Mg versus Sr/Mg elemental ratio diagram for fin spine tissue.

Keywords: trace elements, water chemistry, fish tissue, elemental ratios, lake water, bioaccumulation

INTRODUCTION

According to Gibbs (1970), atmospheric precipitation, rock dominance and an evaporation-crystallization process are the three major mechanisms controlling surface water chemistry. The waters of rock-dominated systems are in partial equilibrium with the materials in their catchments and their position within this group is dependent on the relief, chemical composition and climate of the catchment. Gorham (1961) stated that Ca, Mg, Na and K account for most of the ionic composition of the world's freshwater systems. Freshwater bodies should therefore have characteristic chemistries related to the underlying geology. Fish living in these water bodies should thus also have characteristic chemistries related to these water bodies.

Dallas and Day (2004) define bioaccumulation as the ability of an organism to accumulate and concentrate substances directly from the surrounding water (bioconcentration) or indirectly via the food chain (biomagnification). Bioaccumulation of metals by fish is influenced by a number of factors. The physico-chemical water quality conditions, e.g. pH, influence bioaccumulation indirectly by changing the solubility of metal compounds, or directly by damage to epithelia, making them more permeable. Hardness and salinity also reduce metal uptake through competition for binding sites (Jezierska and Witeska, 2006). The hydrological period is known to influence metal bioaccumulation mostly due to a concentration of salts in the water during the dry season. Robinson and Avenant-Oldewage (1997) noted significant differences in metal content of fish tissues between wet and dry seasons. Kotze et al. (1999) noted significant differences in metal content of tissues

* To whom all correspondence should be addressed. e-mail: wikusj@geoscience.org.za Received: 17 December 2014; accepted in revised form 20 November 2015 during different surveys at Lake Loskop and Mamba Weir in the Olifants River.

Coetzee et al. (2002) reported that Zn, Pb, Cr and Ni showed significant positive correlations with fish size. Nussey et al. (2000) found that the smaller the fish the higher the bodyload of metals due to various bioaccumulation processes. Coetzee et al. (2002) found few significant differences in the metal content of fish tissues between males and females. Different species may accumulate different amounts of metals due to different living and feeding habits (Jezierska and Witeska, 2006). Kotze et al. (1999) noted species differences in metal content of tissues from fish in the Olifants River. Robinson and Avenant-Oldewage (1997), Kotze et al. (1999) and Coetzee et al. (2002) noted significant differences in metal content of fish tissue samples from distinct localities in the Olifants River.

Traditionally, bioaccumulation monitoring is used as an indicator of pollutant exposure by comparing the concentrations of toxicologically important metals (e.g. Cd, Pb) in an indicator species and tissue type between sites or surveys (Wepener et al., 2011). However, for the purposes of this paper, the relationship between the ratios of selected metals in fish tissue types and the water, rather than individual metal concentrations, were used to determine if the fish tissue reflected the major geological features of the catchment. Seasonal variations in the metal concentrations of lake water due to dilution by relatively clean rainwater are largely negated by the use of elemental ratios rather than absolute concentrations. Different fish tissues incorporate and retain chemical elements from the environment for varying time periods. Spine tissue bioaccumulates metals from the environment over a longer period than blood and muscle tissue. The seasonal variation in water chemistry is therefore more likely to be represented in the blood and muscle tissue than in the spine tissue. A multi-year sampling campaign may therefore produce better correlation than a single sampling exercise.

Otoliths are acellular and metabolically inert structures that permanently retain any chemical elements that are accreted onto their growing surfaces (Campana, 1999). Secor et al. (1995) and Bath et al. (2000) noted that some otolith element ratios (Sr/Ca and Ba/Ca) are proportional to their ratios in the surrounding water. Whitledge (2008) could distinguish fish from 4 American rivers based on their otolith Sr/Ca and Ba/Ca ratios. Wells et al. (2003) investigated westslope cutthroat trout (Oncorhyncus clarki lewisi) from the Coeur d'Alene River in Idaho and found that otolith and scale chemistries were linearly related to the Sr/Ca and Ba/Ca ratios in the water. Walther and Thorrold (2006) calculated that water contributed 83% of Sr and 98% of Ba in otoliths formed in spiked seawater, implying that the chemical signatures recorded in otoliths of marine fishes should reflect the ambient water composition of these elements at the time of deposition.

Otoliths are made of calcium carbonate. Sr^{2+} , Ba^{2+} and Mg^{2+} have very similar ionic radii to Ca^{2+} and can thus substitute for Ca^{2+} . In addition, elements like Sr, Ba, Mn, Fe, Pb, Li, Mg, Cu and Ni are metabolically inert and are not resorbed from otoliths (Sako et al., 2005). Limburg (1995) noted that the use of Sr/Ca ratios in freshwater systems is limited relative to seawater systems, due to lower element concentrations (up to 10 times). Recent studies by Krause and Secor (2004) have however indicated that the range of Sr in freshwater systems may approach that of seawater. Campana (1999) observed that the concentrations of the most common elements (Ca, Na, K, Mg and Cl)

differ substantially between fresh and salt water, yet do not appear to be reflected in the otoliths. Trace elements like Sr, Zn, Pb, Mn, Ba and Fe in fresh and seawater are however consistent with an environmental effect.

Gillanders (2005) indicated that it is plausible that the effect of ambient Sr outweighs that of salinity. Secor et al. (1995) and Kawakami et al. (1998) positively linked the Sr/Ca ratio in sagittal otoliths to salinity. Martin et al. (2004) found significantly elevated Sr/Ca ratios in otoliths of marine larval spot (*Leiostomus xanthurus*) at a salinity of 25‰ vs. 15‰.

Campana (1999) noted that existing literature does not support an overall relationship between otolith Sr/Ca ratio and temperature for either fresh or seawater fish. Townsend et al. (1992) suggested that temperature-dependent Sr/Ca fractionation only happens at low water temperatures (< 10°C). Martin et al. (2004) found a significant linear relationship between temperature and Sr/Ca ratios in otoliths of marine larval spot (*Leiostomus xanthurus*). Elsdon and Gillanders (2002, 2004) investigated juvenile black bream (*Acanthopagrus butcheri*) and found that water temperature significantly influenced the Sr/Ca and Ba/Ca ratios in otoliths while salinity alone did not influence these ratios. Bath et al. (2000) found that temperature significantly influenced Sr incorporation but not Ba incorporation into otoliths of marine fish.

The primary aim of this paper is therefore to investigate whether a simplified analytical technique will be able to relate fish tissue chemistry to water chemistry as well as to



Figure 1

Map of South Africa indicating secondary catchments of the project area. The catchments represented are the Upper Vaal River catchment, the Mgeni River catchment, the Crocodile River (West) catchment and the Upper Olifants River catchment. Codes for the lakes are explained in Table 1.

distinguish between fish from different lakes. The secondary aim is to determine whether any deviation from the expected fish tissue chemistry could be linked to either pollution events or the sampling of fish introduced from another catchment. The study was undertaken to develop a scientific method to minimize illegal entries at major South African freshwater fishing tournaments (Jordaan, 2015).

MATERIALS AND METHODS

Description of the project area

The project area consisted of selected lakes within the Vaal, Mgeni, Crocodile (West) and Olifants River catchments (Fig. 1). The catchments were selected due to different sizes, different sources of pollution and different underlying geological composition.

Collection and preparation of samples

Samples included: water taken from the surface of lakes, and fish samples taken mainly by bank and boat angling as well as gill netting (Table 1). Water samples were collected in 2 ℓ high-density polyethylene (HDPE) containers, not acidified, cooled and sent to the laboratory for analyses within 24 h.

Four major species of fish were targeted, i.e., common carp (Cyprinus carpio), sharptooth catfish, (Clarias gariepinus), largemouth bass, (Micropterus salmoides), and Mozambique tilapia (Oreochromis mossambicus). Some minor species were also included in the analysis. Fish samples were collected in plastic containers, packed in ice and brought to the laboratory, where they were frozen to -5°C. Fish muscle samples were removed from the fillets of each fish. Fish spine samples were collected from the dorsal, ventral or pectoral fins of each fish. Otoliths and gills were extracted by dissecting the fish skulls from the ventral side. Liver samples were collected by ventral dissection. Samples were oven-dried at approx. 80°C for 14 days. All soft tissues were removed from spine samples, whereafter they were pulverized in a swing mill. Gill filaments were manually removed from dried gill samples before crushing.

Chemical analyses

Trace element concentration analyses of water samples consisted of first filtering samples through 0.45 μ m cellulose nitrate filters. Water samples were then diluted 5 times to add the internal standards (In and Ir) and to reduce total dissolved solids. The samples were made up in 2 ml/100 ml HNO₃ to keep analyte elements in solution. Analytical grade HNO₃ and ultra-pure water were used in all preparations. Samples were analysed on a Perkin Elmer SCIEX ELAN DRCII ICP-MS with AS 93 plus auto-sampler (Jordaan and Maritz, 2010).

Trace element concentration analyses of fish samples consisted of crushing the dried material with an agate mortar and pestle, or a swing mill for spine samples. Preparation of tissue samples for analyses followed a simplified version of the method described by Wepener and Vermeulen (2005). A 0.3 g portion of sample was digested in HNO₃ and HClO₄ for 2 h at 85°C in an aluminium heating block. Samples were diluted to add the internal standards (In and Ir), and analysed on a Perkin Elmer SCIEX ELAN DRC II ICP-MS with AS 93 plus autosampler (Jordaan and Maritz, 2010).

Quality assurance

To evaluate the water method, liquid samples from the SABS Water-Check (group 1) inter-laboratory proficiency test (South African Bureau of Standards, 2010a, 2010b, 2010c) were regularly analysed. Elements analysed included Al, Ba, Be, B, Cd, Cr, Co, Cu, Fe, Pb, Mn, Hg, Mo, Ni, Si, Sr, V, Zn, As and Se. Average *Z*-scores obtained were as follows: April 2010, 0.73; June 2010, 0.73 and October 2010, 0.69. All Z-scores between -2 and 2 are considered satisfactory.

The certified reference materials DOLT-4 (Dogfish Liver) and DORM-3 (Fish Protein) (National Research Council Canada, 2007, 2008) were analysed in quadruplet to evaluate the performance of the analytical method compared to analyses by Aygun and Abanoz (2011), De La Calle et al. (2012) and Sannac et al. (2012) of the same reference material. Dilution factors were chosen to include both major and trace elements in the same analyses, which implies that the method was not sensitive enough to analyse some elements at ultra-trace levels. Only those elements that were mostly above the detection limit (more than 93% of analyses in total dataset) were considered for scientific interpretation (Appendix 1: Table A1). The Rb concentration in otolith samples and Ba and U in gill samples were also included even though between 10 and 20% of the data were below the detection limit. The Merck VI certified calibration standard, with the addition of certified Merck single element standards, was used to calibrate both water and fish analysis methods.

Statistical analysis

Summary statistics (mean, standard deviation) of metal concentrations were determined using Microsoft Excel. Scatter plots of elemental ratios were manually developed to visually establish the best parameters to maximize separation of dissimilarity and minimize the separation of similarity.

RESULTS AND DISCUSSION

Sample identification

In Table 1 the number of species sampled and tissues collected from each lake are represented. The codes provided for the different lakes and fish species are used in the figures and tables throughout the paper. The Olifants River catchment was sampled during both the wet and dry season while most of the lakes in the other catchments were sampled during only one of the seasons for water analyses. For the purpose of this project August–November is considered the dry season and December–July is considered the wet season.

Otolith tissue

From the literature it is evident that no single model has been presented that generally explains the link between metal concentrations in otoliths and water chemistry for all species across the freshwater and marine salinity and temperature range. This is however possible for specific examples. The dissolved metal concentrations in water samples and the concentrations of metals in the otolith tissue of carp from the project area are presented in Table 2. Discrimination between lakes is only possible if there are measurable differences in water chemistry between these lakes, usually as a result of different catchment geology of anthropogenic input. Figure 2 is a diagram of

TABLE 1

Codes for the different lakes and fish species are indicated. Numbers in matrix indicate the number of target species analysed per lake and tissue type. 'No sample' is indicated by ns. Otolith tissue is indicated by o. Spine tissue is indicated by s. Muscle tissue is indicated by m. Liver tissue is indicated by l. Gill tissue is indicated by g. The sum of otolith, spine, muscle, liver and gill tissue samples is indicated by o, s, m, l, g combined. Lake water samples were mainly collected between 2007 and 2009. Fish samples were mainly collected between 2007 and 2011. Both sexes were collected during wet and dry

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						arget	specie	S					I		specie	s 	1	1
Catchment	Lake	Code			Cyprinus carpio (CC-common carp)			Clarias gariepinus (CG-sharptooth catfish)	Micropterus salmoides (MS-largemouth bass)	Oreochromis mossambicus (OM-Mozambique tilapia)	Chetia flaviventris (CF-canary kurper)	Labeo capensis (LA-Orange River mudfish)	Labeo cylindricus (LC - redeye mudfish)	Labeo ruddi (LU-silver mudfish)	Labeo rosae (LR-rednose mudfish)	Labeo umbratus (MO-moggel)	Labeobarbus aeneus (LN-smallmouth yellowfish)	Labeobarbus polylepis (LP-smallscale yellowfish)
Tiss	sue		0	s	m	1	g		S				0, 8	s, m, l, g	combi	ned		
	Bon Accord	BA	4	4	4	3	4	3	3	4	ns	ns	ns	ns	ns	ns	ns	ns
	Roodeplaat	RP	3	3	3	3	3	5	3	5	4	ns	ns	ns	ns	ns	ns	ns
Vest	Klipvoor	KL	5	5	5		5	4	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns
e (V	Marais	MA	6	6	5	6	6	6	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns
odil	Rietvlei	RV	3	3	3	3	3	5	ns	ns	8	ns	ns	ns	ns	ns	ns	18
Lo I	Hartbeespoort	HB	2	3	3	3	3	4	ns	5	11	ns	ns	ns	ns	ns	ns	ns
	Roodekopjes	RO	5	6	5	5	5	4	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns
	Vaalkop	VK	5	5	5	5	5	5	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns
	Bronkhorstspruit	BS	3	3	4	4	5	ns	4	ns	ns	ns	ns	ns	ns	ns	ns	ns
	Witbank	WI	ns	1	1	1	5	1	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns
	Doornpoort	DP	1	1	1	1	5	1	1	ns	ns	ns	ns	ns	ns	ns	ns	ns
ants	Middelburg	MB	1	ns	ns	ns	ns	ns	2	ns	ns	ns	ns	ns	ns	12	10	ns
olif	Loskop	LK	14	48	16	16	16	4	ns	15	ns	ns	ns	ns	16	ns	ns	ns
	Rust de Winter	RW	2	4	4	4	4	2	4	5	ns	ns	ns	ns	ns	ns	ns	ns
	Rhenosterkop	RK	ns	ns	ns	ns	ns	ns	5	5	ns	ns	6	9	ns	1	ns	ns
	Arabie	AR	2	2	3	2	3	1	ns	6	ns	ns	ns	ns	ns	ns	ns	ns
_	Sterkfontein	ST	ns	ns	ns	ns	ns	ns	5	ns	ns	ns	ns	ns	ns	ns	15	ns
Vaa	Vaal	VA	15	28	16	15	17	12	ns	ns	ns	28	ns	ns	ns	ns	ns	ns
	Bloemhof	BL	9	31	22	22	22	8	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns
	Midmar	MI	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns
eni	Albert Falls	AF	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns
Mg	Nagle	NA	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns	ns
	Inanda	IN	1	1	1	1	1	ns	ns	6	ns	ns	ns	ns	ns	ns	ns	ns

the Sr/Ca elemental ratio of lake water and fish otoliths from the project area, as used by Elsdon and Gillanders (2003) and Bath et al. (2000). There are large variations in Sr/Ca ratios of the lake water (all freshwater lakes) and overlaps between lakes from the four catchments. When considering the Olifants River catchment, the Highveld lakes have lower Sr/Ca ratios that combine to form an average for Lake Loskop. Downstream the Olifants River mixes with water from the Elands River with a much higher Sr/Ca ratio to produce an intermediate ratio at Lake Arabie. The Vaal River catchment shows an increase in Sr/Ca ratio along the length of the project area. In the Crocodile River catchment trends are opposite for the Lake Rietvlei and Lake Roodeplaat systems.

The largest variation in Sr/Ca ratios of the otoliths is related to the specific fish species, as indicated by the linear regressions of the data from the project area (Fig. 2). Even though data were collected over several years during both the wet and dry seasons, which implies that there may be significant variation in the data from a single lake, the data are still relatively constant and separated mainly according to species. According to the literature, water (i.e. environmental exposure) has the major influence on otolith chemistry and food the least (Walther and Thorrold, 2006). Fish behaviour may thus not play a significant role in otolith chemistry. Salinity should not play a major role in this study since all lakes are freshwater lakes even though some are moderately polluted. A clear trend from lakes in colder areas to lakes in warmer areas was also not observed. The major extrinsic contributing factor thus appears to be the chemistry of the lake water while an intrinsic factor may be due to physiological mechanisms of otolith formation that possibly differs between fish species (Campana, 1999). Fish must however be in equilibrium with lake water and must not be able to migrate between different lakes or catchments.

	lakes in the project area (codes for lakes as in Table 1; hs = ho sample)										
tch	e	Wate	r			Otolith	tissue				
Cat	Lal	Ca (μg/ℓ)	Sr (µg/ℓ)	Ca (g/100g)	Mg (mg/kg)	Fe (mg/kg)	Ni (mg/kg)	Sr (mg/kg)	Co (µg/kg)		
	BA	35 833±1 311	155±3	39.4±4.1	854±94	2 038±362	17±2	93±34	684±75		
	RP	30 460±3 872	100±5	39.9±3.1	903±52	2 209±374	23±7	156±42	855±219		
đ	KL	34 001±2 814	136±3	35.9±1.4	833±49	1 635±64	15±1	69±7	607±28		
odilo	MA	30 981	112	31.8±4.3	879±86	2 637±270	17±2	98±27	675±54		
roc	RV	2 9131±564	102±2	39.0±2.9	928±155	1 829±558	18±2	102±24	626±93		
0	HB	35 992±1 449	139±6	38.8±1.9	849±70	1 814±216	17±1	94±1	659±52		
	RO	29 229±5 229	163±11	41.5±2.5	944±53	1 718±87	16±1	135±17	631±44		
	VK	39 348±4 053	284±27	38.0±0.9	988±33	2 036±415	17±1	161±25	673±59		
	BS	19 472±1 900	80±9	41.3±3.2	927±206	2 121±553	18±3	105±25	680±71		
	WI	46 394±12 046	375±90	39.7	839	2 745	21	212	746		
	DP	38 962±3 297	310±29	40.9	711	1756	18	182	719		
ants	MB	7 0832±17 933	524±137	ns	ns	ns	ns	ns	ns		
olifa	LK	30 788±6 933	180±35	42.6±2.3	842±253	1 615±219	14±2	201±290	586±81		
	RW	6 772±974	39±4	35.3±3.1	977±80	2 720±111	17±1	120±14	724±41		
	RK	14 574±1 036	97±4	ns	ns	ns	ns	ns	ns		
	AR	30 702±6 580	180±34	41.6±2.4	1 032±108	2 052±501	18±2	158±13	678±58		
	ST	8 509±71	63±1	ns	ns	ns	ns	ns	ns		
Vaal	VA	14 033±3 283	89±25	42.3±5.8	753±106	3 218±1902	31±19	161±24	1 109±592		
	BL	30 259±6 847	123±48	43.2±2.1	731±53	1 407±140	13±2	89±7	528±55		
	MI	4 089±674	32±5	ns	Ns	ns	ns	ns	ns		
eni	AF	4 639±418	33±3	ns	Ns	ns	ns	ns	ns		
Mg	NA	5 935±810	39±5	ns	Ns	ns	ns	ns	ns		
	IN	12 393±1 152	64±6	37.4	766	2 140	17	221	649		





Figure 2

Comparison between average Sr/Ca (mmol/mol) elemental ratio of fish otoliths and lake water from the project area (codes for lakes and fish species as in Table 1). The error bars indicate one standard deviation below and above average. Only standard deviations for OM and MS are indicated as it shows the typical ranges for all the data. Solid lines indicate linear regression lines of the average ratios of the target species per lake.

Together with the modified alkali-lime index [Na₂O+K₂O-CaO] and the aluminium saturation index [Al/(Ca-1.67P+Na+K)], the Fe/Mg ratio is often used as a means of rock classification in igneous geology (Frost et al., 2001). This primary variation in the Fe/Mg ratio may induce variation in weathered/transported materials and the water that contributes to these processes. Lake sediments and the otoliths of fish within these lakes may thus show variation in their Fe/Mg ratio, in part due to the chemistry and weathering processes in the upper catchment. Figure 3 is a diagram of Fe versus Mg of fish otoliths from the project area normalized to Ca, used as a discrimination diagram. Solid symbols indicate carp otoliths and are divided according to the lakes from where they were collected. Open symbols indicate the rest of the otoliths and are only divided according to species. The Mg/Ca parameter distinguishes between 3 groups of species (carp; labeo; tilapia, bass and catfish). The Fe/Ca parameter distinguishes mostly between different lakes even though much overlap exists. A similar distinction between lakes is seen in all species investigated. Fe and Mg are thus not incorporated into otoliths in exactly the same manner for all species while the chemistry of the environment is roughly reflected. A clear distinction can also not be made between otoliths from different catchments.

Cobalt and Ni can sometimes be used as indicators of genetic processes (Herd et al., 2009) and ore-forming processes (Bralia et al., 1979) in geology. According to the Goldschmidt classification (Battey, 1981) both Co and Ni are siderophile elements and are thus normalized to Fe. Figure 4 represents the Co/Fe elemental ratio versus the Ni/Fe elemental ratio of fish otoliths from the project area. A positive correlation is observed between these two components. The diagram does not distinguish between different fish species. Only carp (large solid symbols), bass, tilapia and catfish otoliths are indicated on the diagram, but all samples obtained from the project area follow the same trend. Distinction between otoliths from different lakes can broadly be made although large overlaps do occur. Co and Ni are thus incorporated into otoliths in a fashion indicating environmental chemistry rather than differences between species.

The bass populations from Lake Middelburg and Lake Witbank contain one individual each that has extremely elevated Ni/Fe otolith ratios (Fig. 4). These lakes are directly downstream from a severely polluted coal-mining area. The area also contains coal-fired power stations and steel mills utilizing iron ore from the Bushveld Igneous Complex, providing an ample Ni source.

Plotting the different elements detected in the fish otoliths from the project area in a similar fashion as in Figs 3 and 4 allow distinction between 2 groups of elements: Sr, Ba, Na, K, P and Mg, which best distinguish between different fish species and to a lesser degree between different lakes, and Fe, Co, Ni, Mn, Ga, V and Ca which only broadly distinguish between lakes. It was not possible to use La, Pr and Sm to distinguish between lakes.

Spine tissue

Biomineralization of otoliths, discussed above, differs from vertebrate bone in that otolith epithelium is not in direct contact with the region of calcification (Campana, 1999). However, in an experiment where juvenile snapper (*Pagrus auratus*) were exposed to water with elevated Sr concentrations (10 x ambient) for 5 days it was shown by Pollard et al. (1999) that Sr is absorbed from the water and deposited in the dorsal spines where it was persistent for at least 36 days and showed no sign of decay during the experiment.

The concentrations of metals in the spine tissue of the four target fish species from the different lakes are presented in Table 3. Using a plot of the Sr/Ca elemental ratio of lake water versus fish otoliths, as used by Bath et al. (2000) and Elsdon and



Fe/Ca (mmol/mol) elemental ratio versus Mg/Ca (mmol/mol) elemental ratio of fish otoliths from the project area (codes for lakes and fish species as in Table 1)

Gillanders (2003) for plotting spine samples from the project area, provides similar results as obtained for otoliths (Fig. 5). The regression of the carp spine data however exhibits a steeper slope while the data spread is greater. Carp spine samples from Lake Inanda, Lake Vaalkop and Lake Roodeplaat are more displaced from the regression line than the other carp spine samples (Lake Vaalkop and Lake Roodeplaat carp samples are overlain by catfish samples). The distinction between different species is not as clear as when using the otolith Sr/Ca ratio.



Figure 4

Co/Fe (mmol/mol) elemental ratio versus Ni/Fe (mmol/mol) elemental ratio of carp, bass and catfish otoliths from the project area (codes for lakes and fish species as in Table 1).



Comparison between average Sr/Ca (mmol/mol) elemental ratio of fish spine and lake water from the project area (codes for lakes and fish species as in Table 1). The error bars indicate one standard deviation below and above average. Only standard deviations for OM and MS are indicated as it shows the typical ranges for all the data. Solid lines indicate linear regression lines of the average ratios of the target species per lake.

	Average (± star	ndard dev	iation) metal conto (codes fo	TABLE 3 ent of carp, catfish, or lakes and fish sp	bass and tilapia spi ecies as in Table 1)	ne tissue from proj	ect area lakes
					Spine tissue		
Spp.	Catchment	Lake	Ca (g/100g)	Na (mg/kg)	Mg (mg/kg)	Sr (mg/kg)	Ba (g/kg)
		BA	21.4±0.9	6 177±804	3 367±200	360±29	40 364±6 663
		RP	19.7±1.4	6 146±671	3 290±298	282±50	47 084±19 723
		KL	20.0±0.5	5 200±193	2 748±158	348±12	38 019±3 988
		MA	18.9±0.7	6 434±348	3 041±183	324±36	17 739±4 826
	Crocodile	RV	19.9±0.5	6 478±207	2 977±64	311±44	17 155±4 276
		HB	20.2±1.5	5 977±729	3 488±396	288±83	15 418±2 496
		RO	20.5±0.2	6 924±260	3 332±241	461±30	30 384±5 031
		VK	20.7±1.6	6 191±731	3 412±398	674±72	70 383±15 180
CC		BS	20.7±1.9	6 505±1 137	3 472±519	380±37	74 751±14 731
		WI	18.0	5 722	2 876	574	27 721
		DP	20.7	6 133	3 740	570	51 547
	Olifants	LK	20.1±3.5	6 437±1 093	3 369±680	493±103	63 819±44 801
		RW	16.8±0.9	6 702±274	3 887±291	372±27	12 2042±17 431
		AR	16.1±1.2	6 185±638	4 398±281	398±18	59 661±3 453
		VA	26.5±8.8	9 866±3 794	4 369±1292	657±247	79 476±27 888
	Vaal	BL	20.1±3.4	5 948±993	3 510±581	333±67	41 572±10 665
	Mgeni	IN	23.3	6 647	3 737	666	85 860
		BA	24.2±0.6	6 467±620	3 493±140	257±37	22 949±9 767
		RP	23.7±1.1	6 733±474	3 391±189	239±26	13 278±4 917
		KL	23.5±0.6	5 083±492	3 135±65	279±38	19 961±5 887
		MA	21.0±0.8	5 729±341	3 143±106	190±36	18 826
	Crocodile	RV	19.6±2.5	5 095±350	3 613±192	148±10	11 972±5 744
		HB	20.2±3.6	6 198±307	4 080±385	185±32	10 856±1 268
		RO	22.6±0.5	6 122±774	3 486±187	300±50	11 992±2 865
CG		VK	20.8±0.8	5 737±629	3 150±131	424±131	16 521±2 685
		WI	19.4	4 357	3 082	524	18 825
		DP	25.8	4 807	2 749	497	<6 281
	Olifants	LK	20.8±1.1	6 189±671	3 138±139	365±68	19±3
		RW	19.2±0.4	6 835±94	3 897±51	405±18	10 5732±6 819
		AR	23.0	5 370	4 015	257	21 069
	371	VA	22.1±3.3	6 745±953	3 447±378	458±92	20 649±9 461
	Vaai	BL	24.5±2.0	6 800±935	3 603±374	273±50	22 684±15 147
	Crocodile	RP	22.6±1.0	5 536±769	3 925±132	210±38	10 680±2 554
		BS	23.6±1.0	5 650±502	4 495±245	279±32	19 300±2 456
		DP	22.2	4 334	3 930	619	<6 281
MS	Olifants	MB	17.0±5.5	3 825±1 785	2 784±1 243	448±164	14 017±0
		RK	21.1±2.4	5 331±489	4 161±476	453±97	30 846±2 848
		RW	22.5±2.2	5 556±597	4 097±594	395±65	87 277±25 354
	Vaal	ST	24.0±2.9	6 476±798	4 078±529	530±82	24 668±5 347
		BA	22.4±2.2	5 301±447	2 851±165	475±27	18 986±1 987
	C == = = 1:1.	RP	25.6±0.3	7 655±276	3 119±121	367±13	15 286±2 129
	Crocoalle	HB	21.4±3.6	5 679±404	3 289±241	337±89	12 725±2 681
OM		RK	23.3±0.2	4 929±242	2 451±140	829±41	95 406±14 503
	Olifunt	AR	23.1±1.8	6 218±635	3 143±554	673±115	59 454±5 723
	Olifants	LK	21.0±4.6	6 598±1 322	2 527±580	542±112	61 156±42 752
	Mgeni	IN	28.4±1.4	7 902±453	3 647±383	700±34	73 998±11 967

Using Fe as a discriminator between spine samples from the project area, as was done for the otolith samples (Fig. 3), did not produce the expected results. A diagram of Na versus Mg normalized to Ca did however distinguish better between fish species as well as individual lakes (Fig. 6).

Following the identification of elements from otoliths that are more suitable to discriminating between species (Sr, Ba, Na, K, P and Mg), a diagram of Ba versus Sr normalized to Mg produced some separation between fish spine samples from different lakes and to a lesser extent from different species (Fig. 7). Barium versus Sr normalized to Ca gives similar results, but with less separation of the tilapia spine samples. Figures 8, 9 and 10 show catfish, bass and tilapia spine samples, respectively, plotted per lake for comparison with Fig. 7 where carp samples are plotted per lake. In all three species the samples from a specific lake cluster together and are separated from samples from other lakes, indicating that the Ba/Mg versus Sr/Mg plot can successfully distinguish between fish from different lakes for at least 4 species.



Figure 6

Na/Ca (mmol/mol) elemental ratio versus Mg/Ca (mmol/mol) elemental ratio of fish spine tissue from the project area (codes for lakes and fish species as in Table 1).



Figure 7

Ba/Mg (mmol/mol) elemental ratio versus Sr/Mg (mmol/mol) elemental ratio of fish spine tissue from the project area. Carp samples from individual lakes are indicated by solid symbols. Open symbols indicate catfish, bass and tilapia sample groups (codes for lakes and fish species as in Table 1).



Figure 8

Ba/Mg (mmol/mol) elemental ratio versus Sr/Mg (mmol/mol) elemental ratio of fish spine tissue from the project area. Individual catfish samples are indicated by solid symbols (codes for lakes and fish species as in Table 1).



Figure 9

Ba/Mg (mmol/mol) elemental ratio versus Sr/Mg (mmol/mol) elemental ratio of fish spine tissue from the project area. Individual bass samples indicated by solid symbols (codes for lakes and fish species as in Table 1).

Muscle tissue

Metal concentrations in the muscle tissue of carp from the different lakes are presented in Table 4. Rb is the only element that shows some differentiation between fish muscle from the different lakes. When plotted against P and

normalized to Mg (Fig. 11), a weak separation between lakes can be seen in the Rb/Mg ratio and a broad distinction between species in the P/Mg ratio. The Rb/Mg ratio distinguishes better between carp muscle tissue samples from lakes in the Crocodile catchment than in the Olifants catchment.



Figure 10

Ba/Mg (mmol/mol) elemental ratio versus Sr/Mg (mmol/mol) elemental ratio of fish spine tissue from the project area. Individual tilapia samples indicated by solid symbols (codes for lakes and fish species as in Table 1).



Figure 11

Rb/Mg (mmol/mol) elemental ratio versus P/Mg (mmol/mol) elemental ratio of fish muscle from the project area (codes for lakes and fish species as in Table 1).

Liver tissue

Liver tissue contained higher concentrations of trace elements than other tissues, which made additional comparison of samples at trace element level possible. The concentrations of metals in the liver tissue of carp from the different lakes are presented in Table 5. Major elements did not distinguish between different species or between liver samples from different lakes. Figure 12 is a diagram of the Co/Mo elemental ratio versus the Zn/Rb elemental ratio of liver samples from the project area. These ratios were empirically chosen purely to give the best distinction between species as well as between samples from the same species collected from different lakes. Figure 12 shows some distinction between tilapia, bass, carp, labeo and catfish. Among the carp samples, it also made some distinction between different lakes. In the Olifants River

TABLE 4
Average (± standard deviation) metal content of carp
muscle tissue from the different lakes in the project area
(codes for lakes as in Table 1; ns = no sample)

Catch		Muscle tissue							
Catch.	Lake	Mg (g/kg)	P (mg/kg)	Rb (mg/kg)					
	BA	1 169±149	8 629±1 034	8±2					
	RP	1 033±234	10 357±1 137	12±3					
	KL	773±28	7 465±796	9±1					
Cura es dilla	MA	989±123	10 575±705	15±1					
Crocodile	RV	965±215	9 170±1 584	12±2					
	HB	1 106±342	9 438±2 393	12±4					
	RO	787±117	7 142±1 299	7±1					
	KL	1 415±136	7 142±1 299 7 10 751±1 388 9	9±2					
	BS	793±369	8 862±2 668	8±4					
	WI	1239	10 655	12					
0l:fants	DP	797	nd	7					
Ulliants	LK	1 013±183	8 973±928	10±2					
	RW	945±92	9 339±2 061	14±3					
	AR	1 143±86	9 240±506	11±2					
Vaal	VA	1 015±274	8 804±1 027	5±3					
vaai	BL	965±147	7 991±823	5±2					
Mgeni	IN	1 315	9 252	12					

	(codes	for lakes a	s in Table 1;	ns = no sam	ple)				
		Liver tissue							
Catch.	Lake	Zn (mg/kg)	Rb (µg/kg)	Mo (μg/kg)	Co (µg/kg)				
	BA	2 008±967	3 940±924	1 638±199	387±71				
	RP	691±356	6 722±896	715±430	112±69				
	MA	805±326	7 247±896	1 759±942	348±199				
Crocodile	RV	904±136	6 345±865	1 741±525	365±84				
	HB 880±345		8 093±2311	822±505	224±70				
	RO	612±221	4 817±898	683±200	584±538				
	VK	1 175±404	4 439±1 386	1 302±363	560±227				
	BS	1 539±1401	4 969±1 555	459±147	226±202				
	WI	3 077	5 294	Liver tissue Rb Mo (µg/kg) 940±924 1638±199 722±896 715±430 247±896 1759±942 345±865 1741±525 993±2311 822±505 817±898 683±200 439±1 386 1 302±363 969±1 555 459±147 5 294 418 4 940 489 48±3 200 944±352 751±3 544 1 216±212 925±538 753±164 449±2 194 1 037±451 108±1 221 765±309 4 020 552	450				
Ol:fanta	DP	755	4 940	489	224				
Ulitants	LK	587±299	6 448±3 200	944±352	422±259				
	RW	225±90	8 751±3 544	1 216±212	234±67				
	AR	522±56	6 925±538	753±164	226±105				
Vaal	VA	885±418	3 449±2 194	1 037±451	632±368				
Vdal	BL	710±448	3 408±1 221	765±309	399±216				
Mgeni	IN	1 152	4 020	552	691				

TABLE 5 Average (± standard deviation) metal content of carp liver tissue from the different lakes in the project area



Figure 12

Co/Mo (mol/mol) elemental ratio versus Zn/Rb (mol/mol) elemental ratio of fish liver from the project area (codes for lakes and fish species as in Table 1).

catchment 3 groups can be identified: (i) Lake Rust de Winter, (ii) Lake Witbank and (iii) Lakes Bronkhorstspruit, Loskop and Arabie. If a connection between water quality and fish tissue elemental concentrations is assumed than the separation may be explained as Lake Rust de Winter is in the Elands River, a tributary of the Olifants River and Lake Witbank is subjected to pollution from adjacent coal-mining activity. In the Crocodile River 4 groups can be identified: (i) Lake Roodekopjes, (ii) Lake Bon Accord, (iii) Lake Vaalkop and (iv) Lakes Roodeplaat, Hartbeespoort, Marais and Rietvlei. Except for Lake

Roodeplaat the division is based on different tributaries within the catchment. A clear distinction is not found for the two lakes in the Vaal River catchment. If the liver is considered a transient location of elements (higher throughput rate) then it may capture shorter-term chemical variations in the environment better than bone or muscle tissue.

Gill tissue

Gill metal loadings have good potential for correlation with metals dissolved in lake waters as gills are in direct contact with the lake water. This is also the basic assumption behind the biotic ligand model used to predict toxicity due to dissolved metals (Niogi and Wood, 2004). The concentrations of metals in the gills of carp from the different lakes are presented in Table 6. Using a plot of the Sr/Ca elemental ratio of lake water versus fish otoliths as used by Bath et al. (2000) and Elsdon and Gillanders (2003) for plotting gill samples from the project area, again exhibits similar results to otoliths (Fig. 13). A clear correlation is observed between lake water and fish gills with good separation between the different species. Labeos correspond with either carp or tilapia while bass and catfish greatly overlap.

Figure 14 represents the Mg/Ca elemental ratios versus the Na/Ca elemental ratios of fish gill samples from the project area. It completely separates carp, bass, tilapia and catfish, as well as most of the labeo species. Both ratios span a much wider range than for the spine samples (Fig. 6) and thus produce a better separation between species. Within species the Mg/Ca and Na/Ca ratios can however not separate between samples from individual lakes.

Figure 15 is a diagram of the Co/Ni elemental ratio versus the Sr/Ca elemental ratio of fish gill samples from the project area. In this case a clear separation is observed between different species as well as between fish from specific lakes as demonstrated by the carp samples. The Co/Ni ratio is the main factor separating the bass, tilapia and catfish samples while the Sr/Ca ratio is the prominent ratio in defining the carp as a group as well as separating carp samples from individual lakes. This diagram produces the best discrimination between Lakes Vaal and Lake Bloemhof. Distinction between carp samples from the Olifants River catchment is not clear while carp samples from the Crocodile River catchment show much less overlap.

CONCLUSIONS

When comparing lake water chemistry to fish tissue chemistry there are several important factors to consider. Firstly there must be a measurable difference in water chemistry between the different lakes within the project area, which is predominantly controlled by the underlying geology and the anthropogenic activity within the catchment. Secondly the fish must be in equilibrium with the lake water and must not be able to migrate between different lakes or catchments.

The elemental ratios that can be used for comparison depend on the ability to accurately analyse these elements in water or fish samples as well as on the specific fish tissue type. The Sr/Ca has traditionally been used to compare fish with lake water. In the project area, the Sr/Ca ratio also correlated very well between individual fish species and water from specific lakes. A summary of the effectiveness of elemental ratio

	Average	e (± standard dev	iation) metal conte (codes for la	TABLE 6 nt of carp liver tiss kes as in Table 1; n	ue from the differ s = no sample)	ent lakes in the pro	oject area				
		Gill tissue									
Catch.	Lake	Ca (g/100g)	Na (mg/kg)	Mg (mg/kg)	Sr (µg/kg)	Ni (µg/kg)	Co (µg/kg)				
	BA	9.7±1.1	4 653±777	7 405±607	309±38	3654±507	698±459				
	RP	8.0±1.0	4 646±559	5 745±1 306	238±85	4 258±846	265±43				
	KL	7.9±1.4	1 840±173	5 817±801	304±50	3 846±820	403±92				
C	MA	8.4±1.0	3 842±568	6 432±963	303±26	7 933±3 582	589±269				
Crocodile	RV	10.2±0.8	5 240±1 924	7 741±1 453	310±34	3 508±473	601±143				
	HB	9.1±1.2	5 267±1 811	6 746±1 141	3 310±34 3 508±473 1 304±48 3 586±283 3 338±23 2 073±172 0 587±97 3 340±855	303±74					
	RO	8.0±0.7	3 611±393	5 877±414	338±23	2 073±172	616±379				
	VK	9.3±1.4	4 049±740	7 330±850	587±97	3 340±855	697±229				
	BS	7.4±4.0	3 770±1 554	6 176±3 017	303±26 7 933±3 582 589±26 310±34 3 508±473 601±14 304±48 3 586±283 303±7 338±23 2 073±172 616±37 587±97 3 340±855 697±22 257±156 3 927±515 552±34 636 3 118 741 612 5 758 515	552±341					
	WI	8.7	4 645	7 365	636	Ni Co (μg/kg) 3654±507 698±459 4 258±846 265±43 3 846±820 403±92 7 933±3 582 589±269 3 508±473 601±143 3 586±283 303±74 2 073±172 616±379 3 340±855 697±229 3 927±515 552±341 3 118 741 5 758 515 3 872±1 965 1 043±749 4 535±647 469±166 3 522±1 040 435±179 7 529±5 015 1 111±404 8 609±19 223 1 122±799 3 423 476					
0116	DP	11.2	5 077	8 490	612	5 758	515				
Olitants	LK	9.0±2.7	5 536±1 590	6 910±1 698	424±118	3 872±1 965	1 043±749				
	RW	7.1±0.7	4 407±481	5 815±677	358±66	4 535±647	469±166				
	AR	8.6±1.5	3 970±504	6 136±1 420	389±83	3 522±1 040	435±179				
Vl	VA	10.0±1.8	6 826±2 873	7 879±1 445	458±84	7 529±5 015	1 111±404				
vaai	BL	8.8±1.0	4 422±1 217	6 975±743	291±55	8 609±19 223	1 122±799				
Mgeni	IN	8.3	4 480	5 166	368	3 423	476				



Figure 13

Comparison between average Sr/Ca (mmol/mol) elemental ratio of fish gills and lake water from the project area (codes for lakes and fish species as in Table 1). The error bars indicate one standard deviation below and above average. Only standard deviations for OM and MS are indicated as it shows the typical ranges for all the data. Solid lines indicate linear regression lines of the average ratios of the target species per lake.



Na/Ca (mmol/mol) elemental ratio versus Mg/Ca (mmol/mol) elemental ratio of fish gills from the project area (codes for fish species as in Table 1)



Figure 15 Diagram of the Sr/Ca (mmol/mol) elemental ratio versus Co/Ni (mmol/mol) elemental ratio of gills from the project area (codes for lakes and fish species as in Table 1)

diagrams in discriminating between fish species and lakes is given in Table 7. Using other elemental ratios was less successful, mainly due to the low abundance of these elements in lake water.

Considering only fish tissue samples, the best separation between species was achieved using a Na/Ca versus Mg/Ca elemental ratio diagram for gill tissue, followed by a Co/Mo versus Zn/Rb elemental ratio diagram for liver tissue. The best discrimination between fish from different lakes was achieved using a using a Ba/Mg versus Sr/Mg elemental ratio diagram for spine tissue from carp, catfish, bass and tilapia.

ACKNOWLEDGEMENTS

The authors would like to acknowledge Bella Honeybourne, the organizing committee of the 'Three Species Bonanza', EcoCare Trust, Waterlab and the various nature conservation organizations for their support during fieldwork as well as the Council for Geoscience and the University of Johannesburg for financial and technical support.

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TABLE 7												
Summary of effectiveness of elemental ratio diagrams for discriminating between species and lakes												
(codes for lakes and fish species as in Table 1). 1 = Effective, 2 = Partial, 3 = Ineffective												
Ratios	Tissue	Species	Lakes	Comment	Figure							
Sr/Ca vs. Sr/Ca	Otolith, water	1	2		2							
Fe/Ca vs. Mg/Ca	Otolith, water	3	3		3							
Co/Fe vs. Ni/Fe	Otolith	3	2		4							
Sr/Ca vs. Sr/Ca	Spine, water	2	2		5							
Na/Ca vs. Mg/Ca	Spine	2	2		6							
Ba/Mg vs. Sr/Mg	Spine (CC, CG, MS, OM)	2	1	Best for lakes	7, 8, 9,10							
Rb/Mg vs. P/Mg	Muscle	2	2		11							
Co/Mo vs. Zn/Rb	Liver	1	2	Best for species	12							
Sr/Ca vs. Sr/Ca	Gill, water	2	2		13							
Na/Ca vs. Mg/Ca	Gill	1	3	Best for species	14							
Sr/Ca vs. Co/Ni	Gill	1	2		15							

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Certifie	ed reference ma	aterial compared	to metal concen mass. Data no	TABLE A1 trations obta ot available ir	nined during ana Indicated by na.	lysis. Concentra	ations in μg/k	g dry tissue		
		DOLT-4		DORM-3						
Element	Certified	Analysed	De La Calle et al. (2012)	Certified	Analysed	De La Calle et al. (2012)	Aygun and Abanoz (2011)	Sannac et al. (2012)		
Ва	na	301	na	na	3 788	na	na	na		
Са	680 000	664 040	516 000	na	2 318 522	1 909 000	na	na		
Со	250	246	na	na	267	na	na	na		
Fe	1 833 000	1 705 838	1 908 000	347 000	326 196	351 000	338 700	368 000		
Mg	1 500 000	1 405 539	na	na	983 594	na	na	na		
Мо	1 000	991	na	na	258	na	na	na		
Na	6 800 000	6 975 287	na	na	15 069 208	na	na	na		
Ni	970	312	na	1 280	1 317	na	na	1 420		
Р	na	11 249 293	11 126 000	na	7 899 404	5 716 000	na	na		
Rb	na	3 204	na	na	5 940	na	na	na		
Sr	5 500	5 116	na	na	8 275	na	na	na		
Zn	116 000	123 015	125 000	51 300	52 390	53 700	50 100	47 500		