

Seasonal variations in metal content of two *Unio pictorum mancus* (Mollusca, Unionidae) populations from two lakes of different trophic state

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ABSTRACT

To assess the impact of lake trophy on trace element accumulation by *Unio pictorum mancus*, a population from the meso-oligotrophic Lake Maggiore was compared with a population from the eutrophic Lake Candia. The element content in soft tissue and shell biomass, the seasonal variations of element concentrations in soft tissues and shell, and the relationship between element concentrations in the water and those in the tissues were estimated in pursuit of this objective. Thirteen mussel samplings were performed in Lake Maggiore between May 2003 and September 2004 and in Lake Candia between June 2003 and August 2004. Filtered water samples were collected seasonally. Water, shell and soft tissue samples were analysed by ICP-OES and ICP-MS for the measurement of following elements: Zn, Cu, Fe, Mn, Ni, As, Pb, Co, Cr, Mo, V, Cd, Be and Ca. The element concentrations in the mussel tissues and shell from Lake Maggiore were higher than those from Lake Candia. Due to the higher population density and higher element concentrations in the mussels of Lake Maggiore, the element content in the biomass per m² in this lake far exceeded that in Lake Candia. The element concentrations in the mussel tissues, but not in their shells, of both lakes showed a seasonal pattern of variation; this was to be expected as a result of the slower turnover time of the elements in the shell than in the tissues. For all the elements, except Mo, Cr and Be, the "concentration factor" (C.F.) value was higher in Lake Maggiore than in Lake Candia mussels, ranging from 10 to 10⁴ and 10 to 10³ respectively. The higher concentrations in the mussel tissues from Lake Maggiore were justified not merely by the higher element concentrations in the lake water, but probably also because other causes, such as element concentrations in food and element abundance in available forms, combine to affect the concentrations of the elements in the tissues.

Key words: freshwater bivalve, seasonal variations, metals, shell, soft tissues, concentration factor

1. INTRODUCTION

The great capacity of bivalves to concentrate various toxic and non-toxic trace elements in their bodies with no evident danger to themselves is often exploited to identify pollutants in an environment, including those in which the pollutants are so diluted that they cannot be analyzed by the methods commonly used (e.g., Merlini *et al.* 1965; Ferrington *et al.* 1983; Doherty *et al.* 1993; Oertel 1998; Ravera *et al.* 2003a). As an example, radiomanganese (Mn-54) from the fall-out of nuclear tests in the Pacific area was detected for the first time in Europe by gamma spectrometry in *Unio pictorum* from Lake Maggiore (Ravera & Vido 1961; Gaglione & Ravera 1964). Besides having great accumulation capacity and low discrimination power (Bryan 1979), bivalves are regarded as useful pollutant indicators because of their wide geographical distribution, the presence of adults in all seasons and their sedentary way of life. In addition, they can easily be sampled and several species can live in both clean and polluted water bodies.

The shell and soft tissues composing the mussel body typically differ in their chemical composition, and the turnover time of the same element is much shorter in

the tissues than in the shell. Consequently, the chemical composition of the tissues is a reflection of the recent situation of the environment, while the shell composition reflects the integrated situation over a time period corresponding to the age of the mussels. Since several mussel species have a life span of many years (e.g., up to 200 years for *Margaritifera margaritifera*), the variations over time of some environmental characteristics (e.g., nutrient enrichment, pH lowering, temperature variations, metal concentration) have been recorded through shell analyses (Mutvei & Westermark 2001; Price & Pearce 1997; Lazareth *et al.* 2003).

Mussels can play an important role in the environments where they are very abundant (Dame *et al.* 2002), particularly because of their influence on biogeochemical cycles. For example, the shells are a reserve of calcium carbonate and associated elements (e.g., *strontium*) which, when the mussel dies, are dissolved in soft water environments in a relatively short time, while in hard water they are preserved for a long time and may give rise to thanatocoenosis. During the mussel's life, there is a continuous flux of elements from the environment (water and food) to the tissues and *vice versa*. When a mussel dies, its tissues become available as food for other organisms or decompose in a relatively

short time. Thus in the former case the trace elements are transferred from the mussel tissues to other organisms, in the latter the elements are released to the water. In evaluating the importance of mussels in biogeochemical cycles, the first step is to quantify the element content in the shell and tissue biomasses. There is a huge amount of information on the concentration of a number of elements in marine and freshwater mussels, but the available information on the element contents in their biomass is unfortunately poor.

The present study, dealing with 13 trace elements (Mn, Fe, Zn, Cu, Pb, Ni, Cr, Cd, Co, V, Be, Mo, As) plus calcium measured in mussels (shell and soft tissues) and filtered water from Lake Maggiore and Lake Candia, had three major aims.

The first was to compare element contents in the shell and soft tissue biomass of two *Unio pictorum mancus* populations: one from the oligo-mesotrophic Lake Maggiore, the other from the eutrophic Lake Candia.

The second aim was to compare seasonal variations of element concentrations in the mussel shell and tissues of both populations and to estimate the annual mean concentration of each element and its variability. In addition, the similarity level of the seasonal variations of the elements in the mussel tissues was tested.

Thirdly, since mussels are commonly used in bio-monitoring surveys as accumulator organisms, we wanted to evaluate the relationship between element concentrations in mussel soft tissues and those in the water of the two lakes.

2. MATERIAL AND METHODS

2.1. Study sites

The study was carried out on individuals of *Unio pictorum mancus* (Bivalve, Unionidae) from the deep meso-oligotrophic Lake Maggiore and the other from the shallow eutrophic Lake Candia, respectively (Tab. 1). One sampling station (at a place called Sabbie d'Oro) was located on the south-eastern side of Lake Maggiore (latitude N 45°50'19", longitude E 8°37'17") near the village of Brebbia (Province of Varese). The station is a gently sloping bay with silty-muddy sediments. The other station, on the south-eastern side of Lake Candia (latitude N 45°19'32", longitude E 7°54'38", Province of Turin), was located in the littoral zone, also gently sloping, with sediments made up of mud, silt and sand. Further details are reported in another paper (Ravera *et al.* 2007, this issue).

2.2. Sampling and sample preparation

Six mussels from each sample collected monthly from the two lakes for a population biology study (Ravera *et al.* 2007) were used to analyse 13 trace elements and Ca concentration in the soft tissues and shell. The mussels were chosen at random within a shell

length range from 64 to 71 mm. This size was preferred because specimens of this size were abundant in both lakes. The mussel shells were carefully cleaned by scrubbing the surface with a nylon nailbrush to eliminate sediment particles and periphyton coating the periostracum, and to remove attached zebra mussels (*Dreissena polymorpha*).

Tab. 1. Morphometric parameters of Lake Maggiore (Ambrosetti *et al.* 1992) and Lake Candia (Giussani & Galanti 1992).

	Lake Maggiore	Lake Candia
Mean lake level (m a.s.l.)	194	226
Watershed area (km ²)	6599	9.9
Lake area (km ²)	212.5	1.5
Mean depth (m)	177	3.8
Maximum depth (m)	370	7.7

The cleaned mussels were preserved in plastic bags placed in an icebox and brought to the laboratory where they were kept at -20 °C until analysis. After separation of the soft tissues from their shell, tissues and shells of the six mussels were separately pooled to obtain two sub-samples each. The soft tissues were freeze-dried and then kept at 40 °C for 24 hours before pulverisation in a Planetary Micro Mill Fritsch (Fritsch GmbH, Idar-Oberstein, Germany) Pulverisette 7 model with jars and balls in agate. The shells were broken into several pieces to facilitate pulverisation. The powder obtained was passed through a sieve with 0.2 mm opening size. The samples of shells and soft tissues were mineralised by a Microwave Digestion System CEM (Matthews, NC, USA) MDS-2000 model, with an operating frequency of 2450 MHz and a maximum power of 620 W. Prior to each digestion, the teflon PFA vessels were washed with 2 mL of hot concentrated nitric acid (heated for 10 min at 620 W) and then thoroughly rinsed with deionized water. About 500 mg of sample, 8 mL of HNO₃ (65% m/v), and 1 mL of H₂O₂ (30% m/v) were transferred into microwave vessels and left in contact for two hours. Each time, six samples were placed in the microwave carousel, together with a blank prepared with suprapure reagents, and were submitted to the following heating program: 5 min at 40% power; 5 min at 60% power; 15 min 80% power; last cooling. All these solutions were transferred and filtered (Whatman n° 42) into a 50 mL flask and brought to volume. A certified mussel tissue, reference sample CRM 278 (*Mytilus edulis*), prepared by the Community Bureau of Reference (BCR) was used as reference material to evaluate the accuracy of the analytical procedure (Tab. 2).

Water samples were collected seasonally from each lake: 10 double samples from Lake Maggiore and 7 from Lake Candia. Soon after collection the samples were filtered on a 0.45 µm (pore size) Millipore filter and preserved in plastic bottles by the addition of a few drops of 65% (w/v) nitric acid.

Tab. 2. Determination of metals in BCR (Community Bureau of Reference) mussel tissue CRM 278; values found are means of 4 replicates \pm SD. *: Means of 3 replicates \pm SD; values in brackets are reported by BCR and are indicative only.

Element	Found ($\mu\text{g g}^{-1}$)	Certified ($\mu\text{g g}^{-1}$)
Al	37 \pm 8	(70)
As*	5.88 \pm 0.08	5.9 \pm 0.2
Be*	0.06	
Ca	934 \pm 48	(1000)
Cd*	0.335 \pm 0.007	0.34 \pm 0.02
Co*	0.35 \pm 0.01	(0.34)
Cr*	0.83 \pm 0.1	0.80 \pm 0.08
Cu	9.07 \pm 0.42	9.6 \pm 0.16
Fe	121 \pm 4	133 \pm 4
Mn	6.8 \pm 1.4	7.3 \pm 0.2
Mo*	0.36	
Ni*	0.97 \pm 0.02	(1.0)
Pb*	1.89 \pm 0.01	1.91 \pm 0.04
Tl*	<0.001	
V*	0.55 \pm 0.03	
Zn	75 \pm 3	76 \pm 2

2.3. Reagents and calibration

The reagents used, nitric acid (65% w/v) and hydrogen peroxide (30% w/v), were suprapure (Suprapur, Merck, Darmstadt, FRG). High-purity water (electrical resistivity $>18 \text{ M}\Omega \text{ cm}$) was produced with a Milli-Q system (Millipore, MA, USA). Calibration for ICP-OES was obtained with external standards. Standard solutions were prepared by diluting a 1000 mg L^{-1} multielement solution (ICP Multielement Standard IV, Merck, Darmstadt, FRG) with the same acid amount used for sample dissolution. Glassware was cleaned by soaking with the contact overnight in a 10% (w/v) nitric acid solution and then rinsed with deionized water.

Calibration for ICP MS was also obtained with external standards. Standard solutions were prepared diluting by weight a 1000 mg L^{-1} multielement solution (Romil Ltd, Cambridge, UK). Ultra pure nitric acid, also from Romil (Romil UpA), was twice redistilled before use. The containment materials were in FEP (Fluorinated Ethylene Propylene) from NALGENE (Thermo Fisher Scientific Inc., Waltham, MA 02454, USA) and used after an internal, well-established 2 week cleaning procedure. All the chemical operations were performed in a Clean Room (<100 Class).

2.4. Chemical analyses and instrumentation

The analytical determination of Ca, Cu, Fe, Mn and Zn was carried out using the ICPOES Jobin Yvon (Jobin Yvon Emission Horiba Group, Long Jumeau, Cedex, France) JY 24 model. Table 3 gives some basic information about the operating conditions of the ICP-OES spectrometer. The other elements (Be, V, Cr, Co, Ni, As, Mo, Cd, Pb) present in tissue samples at trace level concentration were measured by ICP-MS using an Agilent 7500ce (Agilent Technologies, Palo Alto, CA 94306, USA) equipped with the Collision /Reaction

Cell. The instrumental operating conditions are summarized in table 4. During ICP-MS measurements, isobaric interferences can arise from both the argon used to sustain the plasma and the reagents involved in the sample preparation. In particular, there are well-known interferences in tissue samples that can overlap the masses of Cr, V and As at trace levels. As reported above, the sample preparation method must obviate this problem by avoiding the introduction of chloride-based interferences. In addition, the combined use of the collision cell and the measurement of two separate isotopes for elements like Cr and Cu results in real agreement with the expected values certified in the Reference Standard Material used (Tab. 2).

Tab. 3. Instrument operating parameters of ICP-OES JY24.

Parameter	
Incident power (W)	950
Reflected power (W)	<5
Ar gas flow rate (L min^{-1})	
Outer gas	13
Auxiliary gas	<1
Aerosol gas	0.9
Nebulizer	
Model	Cross-flow – Scott chamber
Sample flow rate (mL min^{-1})	1.5
Pump of sample injection	Gilson miniplus 2

Tab. 4. ICP-MS Agilent 7500ce Operating Conditions.

Parameter	
Plasma RF Power (W)	1500
Carrier Gas Flow (L min^{-1})	1.1
Sample depth (mm)	9.5 from load coil
Spray Chamber Temperature ($^{\circ}\text{C}$)	2 \div 3
Sample flow rate ($\mu\text{L min}^{-1}$)	250
Nebulizer	Agilent microflow PFA
Interface	Nickel simple and skimmer cones

2.5. Calculations

To evaluate the similarity between the seasonal trends of concentrations of the different metals in the tissues of mussels from each lake, considered separately, the significance level for the 91 combinations of the 14 metals was calculated using the Pearson correlation matrix. The significance levels were $p < 0.01$ (highly significant) and $p < 0.05$ (significant). The same method was used to evaluate the seasonal variations of each single metal in the tissues of molluscs from the two lakes, considered together for comparative purposes.

The concentration factor (C.F.) was calculated as the ratio between the concentration of the element in the mussel tissues (related to wet weight) and its concentration in the filtered water. For each element the mean annual concentration in the mussel tissues was divided by the mean concentration in the water. The wet weight/dry weight ratio (9.66 for Lake Maggiore and

Tab. 5. Mean concentration standard deviation and coefficient of variation (% CV) of elements in tissues and shell of mussels from Lake Maggiore and Lake Candia. The mean values are expressed in $\mu\text{g g}^{-1}$ d.w. except for calcium (mg g^{-1} d.w.) and beryllium (ng g^{-1} d.w.).

	Lake Maggiore						Lake Candia					
	tissue			shell			tissue			shell		
	mean	SD	% CV	mean	SD	% CV	mean	SD	% CV	mean	SD	% CV
Ca	60.9	11.0	18	365	18	4.9	16.1	2.5	16	365	21	5.7
Mn	6470	1200	19	189	19	10	3690	620	17	737	86	12
Fe	2390	570	24	108	50	46	1840	290	16	245	92	37
Zn	840	194	23	3.61	0.88	24	120	12	10	1.82	0.44	24
Cu	28.0	7.06	25	5.51	0.99	18	12.7	6.35	50	3.21	1.25	39
Pb	12.0	6.41	54	0.49	0.16	34	2.64	1.48	56	0.26	0.05	18
As	9.05	1.48	16	0.52	0.15	28	2.68	0.34	13	0.43	0.17	40
Ni	6.95	3.34	48	6.48	2.04	31	4.22	1.70	40	5.50	1.15	21
Cr	3.97	1.00	25	0.64	0.33	52	1.48	0.41	28	0.46	0.19	41
Cd	3.56	1.05	30	0.05	0.02	33	0.75	0.13	18	0.04	0.02	52
Co	0.89	0.09	10	0.81	0.04	5.4	0.59	0.09	15	0.61	0.05	8.7
V	0.65	0.28	42	0.12	0.04	33	0.21	0.08	37	0.08	0.04	55
Mo	0.53	0.08	16	0.07	0.04	57	0.27	0.05	18	0.09	0.07	78
Be	21.5	11.4	53	4.50	3.28	73	7.92	5.75	73	1.55	0.86	55

8.65 for Lake Candia mussels, Ravera *et al.* 2007) was applied to calculate the element concentrations in the tissue wet weight from the concentrations measured in the dried material.

The flesh condition index (F.C., tissue dry weight/shell dry weight ratio, Lobel and Wright 1982) and the condition index (C.I., ratio between the tissue dry weight and the product of shell length, width and height, Lobel *et al.* 1991) were calculated to estimate the level of dilution of the elements produced by the mussel soft tissues. The value of both indices generally increases as the metal concentration in the tissues decreases. We devised another index, the Influence index (*I.i.*), as an attempt to estimate the potential influence of the metal contents in the mussel population biomass on its habitat. This index is calculated as the ratio between the metal content in the mussel (soft tissue and shell) biomass per sediment surface unit and the metal concentration in the lake water. The value of the index is expressed as the volume of water (m^3) with the same metal concentration as in the lake water.

$$I.i. = AB/C \times 10^3$$

where: *A* = metal concentration in the biomass ($\mu\text{g g}^{-1}$); *B* = dry weight of the biomass (g m^{-2}); *C* = metal concentration in the lake water ($\mu\text{g L}^{-1}$).

3. RESULTS

3.1. Element concentrations in shell, soft tissues and water

The mean concentration, standard deviation and coefficient of variation (%CV) of elements in the soft tissues and shell of mussels from Lake Maggiore and Lake Candia are reported in table 5. In both lakes, most element concentrations showed wide variations over time in both mussel shell and tissues, as can be seen from the relatively high coefficients of variation. Altogether, most of the elements examined revealed higher

concentrations in both shell and tissues of the Lake Maggiore mussels.

The mean concentrations of elements in filtered water are given in table 6. Element concentrations were generally higher in Lake Maggiore than in Lake Candia, except for Mn, Fe and Ni, which were higher in Lake Candia, and for Co and Cd concentrations, which were similar in the two lakes.

Tab. 6. Element concentrations (mean \pm SD) in filtered (0.45 μm) lake water.

Element	Lake Maggiore	Lake Candia
mg Ca L ⁻¹	26.4 \pm 5.30	15.1 \pm 2.8
$\mu\text{g Zn L}^{-1}$	77.2 \pm 52.6	40.1 \pm 14.2
$\mu\text{g Fe L}^{-1}$	37.2 \pm 48.7	93.3 \pm 93.2
$\mu\text{g Mn L}^{-1}$	20.0 \pm 39.4	55.1 \pm 52.8
$\mu\text{g Cu L}^{-1}$	2.10 \pm 0.32	2.00 \pm 0.00
$\mu\text{g Ni L}^{-1}$	1.60 \pm 0.55	1.85 \pm 0.44
$\mu\text{g As L}^{-1}$	1.36 \pm 0.33	0.51 \pm 0.11
$\mu\text{g Mo L}^{-1}$	0.71 \pm 0.16	0.19 \pm 0.09
$\mu\text{g Pb L}^{-1}$	0.48 \pm 0.22	0.29 \pm 0.09
$\mu\text{g Cr L}^{-1}$	0.33 \pm 0.60	0.12 \pm 0.03
$\mu\text{g V L}^{-1}$	0.28 \pm 0.12	0.12 \pm 0.07
$\mu\text{g Co L}^{-1}$	0.09 \pm 0.05	0.09 \pm 0.01
$\mu\text{g Cd L}^{-1}$	0.02 \pm 0.01	0.02 \pm 0.01
ng Be L ⁻¹	0.98 \pm 0.83	0.19 \pm 0.23

The decreasing sequences of the mean element concentrations in the soft tissues, shell and filtered water from Lake Maggiore and Lake Candia were quite similar, the greatest differences being in the elements with the lowest concentrations, such as Co and Pb in shells and tissues from both lakes (Tab. 7).

The F.C. and the C.I. were both lower in Lake Maggiore mussels than in the Lake Candia mussels (Tab. 8). The C.F. for each element from both lakes and their ratios are listed in table 9.

Tab. 7. The decreasing sequences of the mean element concentrations in the soft tissues, shell and filtered water from Lake Maggiore and Lake Candia.

L. Maggiore	tissues	Ca>Mn>Fe>Zn>Cu>Pb>As>Ni>Cr>Cd>Co>V>Mo>Be
	shell	Ca>Mn>Fe>Ni>Cu>Zn>Co>Cr>As>Pb>V>Mo>Cd>Be
	water	Ca>Zn>Fe>Mn>Cu>Ni>As>Mo>Pb>Cr>V>Co>Cd>Be
L. Candia	tissues	Ca>Mn>Fe>Zn>Cu>Ni>As>Pb>Cr>Cd>Co>Mo>V>Be
	shell	Ca>Mn>Fe>Ni>Cu>Zn>Co>Cr>As>Pb>Mo>V>Cd>Be
	water	Ca>Fe>Mn>Zn>Cu>Ni>As>Pb>Mo>V>Cr>Co>Cd>Be

Tab. 8. Mean value of the tissues and shell dry weight and mean volume of mussel calculated on length, width and height of shell. These variables refer to mussels ranging from 64 mm to 71 mm length, the size interval of the specimens analysed for trace elements. F.C. = flesh condition index, C.I. = concentration index.

	Lake Maggiore	Lake Candia
tissues (g)	1.32	1.74
shell (g)	13.87	11.20
volume (ml)	46.77	46.79
F.C.	0.095	0.155
C.I.	0.028	0.037

Tab. 9. Concentration factors calculated for the mussel tissues from Lake Maggiore and Lake Candia and the ratios between the concentration factors.

Element	Concentration factor		CF _{LM} /CF _{LC}
	Lake Maggiore	Lake Candia	
Mn	33473	7732	4.33
Fe	6661	2282	2.92
Cd	15796	5523	2.86
Pb	2564	1049	2.44
Be	2275	4821	0.47
Ca	239	123	1.94
Cu	1382	732	1.89
Ni	450	264	1.70
Co	1023	803	1.27
V	241	202	1.19
As	689	601	1.15
Cr	1229	1407	0.87
Mo	77	168	0.46
Zn	1126	346	3.26

3.2. Element contents in the biomass of mussel tissues and shell

The results from the biomass of mussel populations from the two lakes (Ravera *et al.* 2007) were used to calculate the mean element contents in mussel soft tissues and shell per m⁻² (schematized in Fig 1).

The "Influence index" (*I_i*) was applied to estimate the potential influence on the environment of the metal contents in the mussels from the two lakes. The values are reported in table 10. In both lakes, 8 elements (Ca, Cu, Ni, Cr, Co, V, Mo, Be) have index values higher for the shell than for the tissues; conversely, 4 elements (Fe, Zn, Pb, Cd) have higher values for the tissues than for the shell. This is due to the metal concentrations being so high in the tissues that they exceed those of the large

biomass of the shell and its metal contents. The index values calculated for the soft tissues and shell of the Lake Maggiore mussels are higher than those of Lake Candia mussels. In Lake Maggiore the index values of Mn and As for the tissues are higher than those for the shell, while in Lake Candia the opposite is the case.

3.3. Seasonal variation patterns of element concentrations in the soft tissues and shell

The monthly variations of element concentrations in the shell and soft tissues are schematized in Figs 2a, 2b and 2c. All the element concentrations in the tissues of Lake Maggiore mussels were higher than those in Lake Candia mussels and generally showed a wider range of variation.

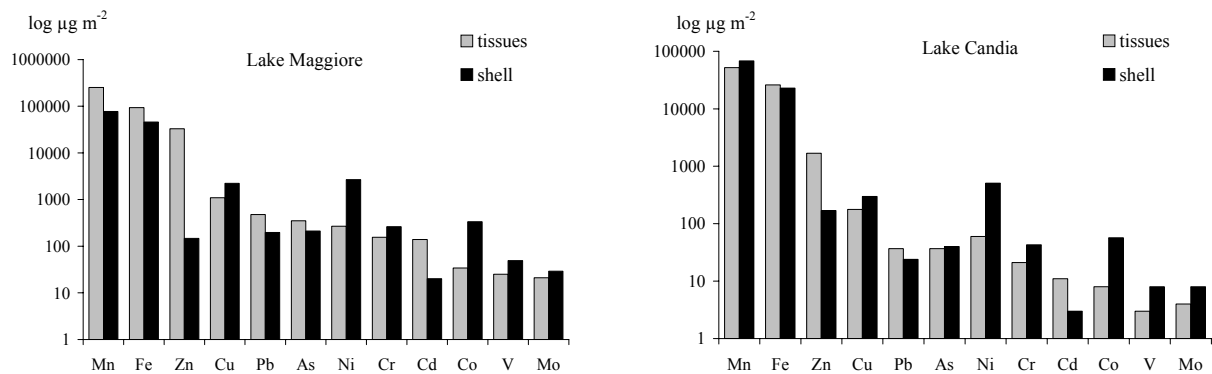


Fig. 1. Mean element contents in the shell and soft tissue biomass (dry weight) of mussels from Lake Maggiore and Lake Candia.

Tab. 10. Values of the *I.i.* (influence index) expressed in m^3 of water corresponding to the ratio between the metal content in the tissues and shell biomass and the mean concentration of the same metal in lake water.

Element	Lake Maggiore		Lake Candia	
	tissues	shell	tissues	shell
Ca	0.09	5.64	0.01	2.22
Mn	12.51	3.86	0.94	1.23
Fe	2.48	1.18	0.28	0.24
Zn	0.42	0.02	0.04	0.004
Cu	0.52	1.07	0.09	0.15
Pb	0.97	0.42	0.13	0.08
As	0.26	0.16	0.07	0.08
Ni	0.17	1.65	0.03	0.27
Cr	0.46	0.79	0.18	0.35
Cd	6.90	1.00	0.50	0.20
Co	0.38	3.67	0.09	0.62
V	0.09	1.75	0.02	0.06
Mo	0.03	0.04	0.02	0.04
Be	0.85	1.87	0.58	0.75

The similarity levels of the seasonal patterns of the different elements in the mussel tissues from each lake are reported in table 11. A highly significant correlation between the tissue concentrations of 14 pairs of elements, and a significant correlation between that of 17 pairs, were observed in Lake Maggiore; in Lake Candia 11 element pairs were correlated at a highly significant p -level, 8 pairs at a significant p -level. The element pairs with a high degree of correlation showed a very similar pattern of seasonal variation, characterized by generally low concentrations in summer 2003 followed by high concentrations in autumn-winter, a decrease in early spring and an increase in summer 2004. A less evident, but similar, pattern of variation characterized the element pairs correlated with a lower degree of significance. All the significantly correlated pairs ($p < 0.01$ and $p < 0.04$) show a positive relationship, the sole exception being the highly significant negative correlation between Mo and Ni concentrations in the tissues of Lake Candia mussels, which showed a mirrored pattern of seasonal variation.

The between-lakes similarity level of each element in the mussel tissues was also tested, but only two ele-

ments (Ni and Pb) showed a close similarity in the two lakes.

4. DISCUSSION AND CONCLUSIONS

The aim of this study was to compare some important biological and ecotoxicological characteristics of two *U. pictorum mancus* populations living in two lakes of different trophic state. As this study was carried out simultaneously on the same monthly samplings of populations of *Unio pictorum mancus* from two lakes (the mesooligotrophic Lake Maggiore and the eutrophic Lake Candia), it was possible to compare the biological and chemical characteristics of both populations.

Generally, element concentrations were higher in the tissues than in the shell of both populations. The ratio between element concentrations in the tissues and those in the shell of Lake Maggiore mussels was always higher than those in Lake Candia. This is the result of high metal concentrations in the tissues calculated for Lake Maggiore mussels rather than low concentrations in the shell.

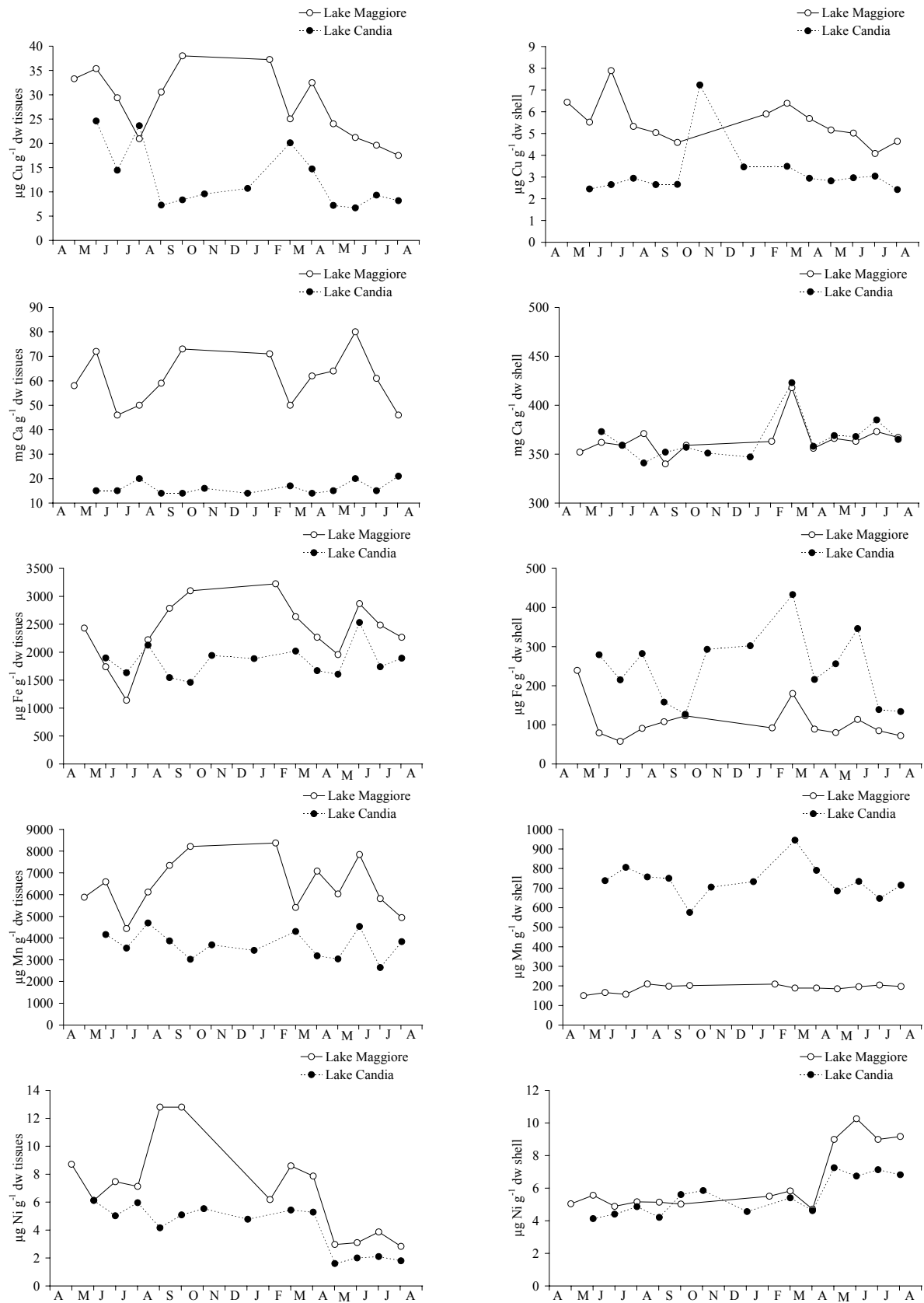


Fig. 2a. Seasonal variations of element concentrations in shell and soft tissues of *Unio pictorum mancus* from Lake Maggiore and Lake Candia.

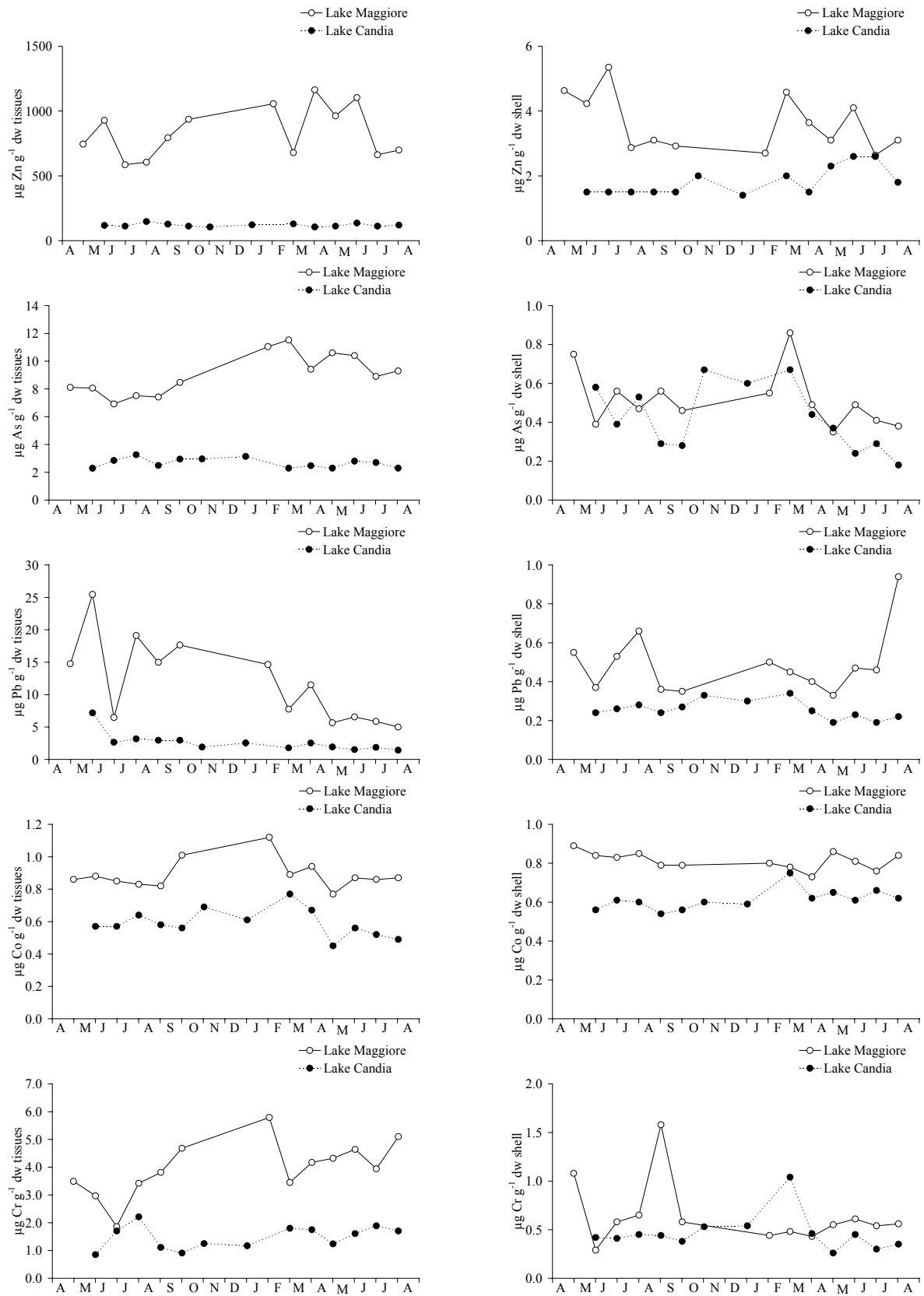


Fig. 2b. Seasonal variations of element concentrations in shell and soft tissues of *Unio pictorum mancus* from Lake Maggiore and Lake Candia.

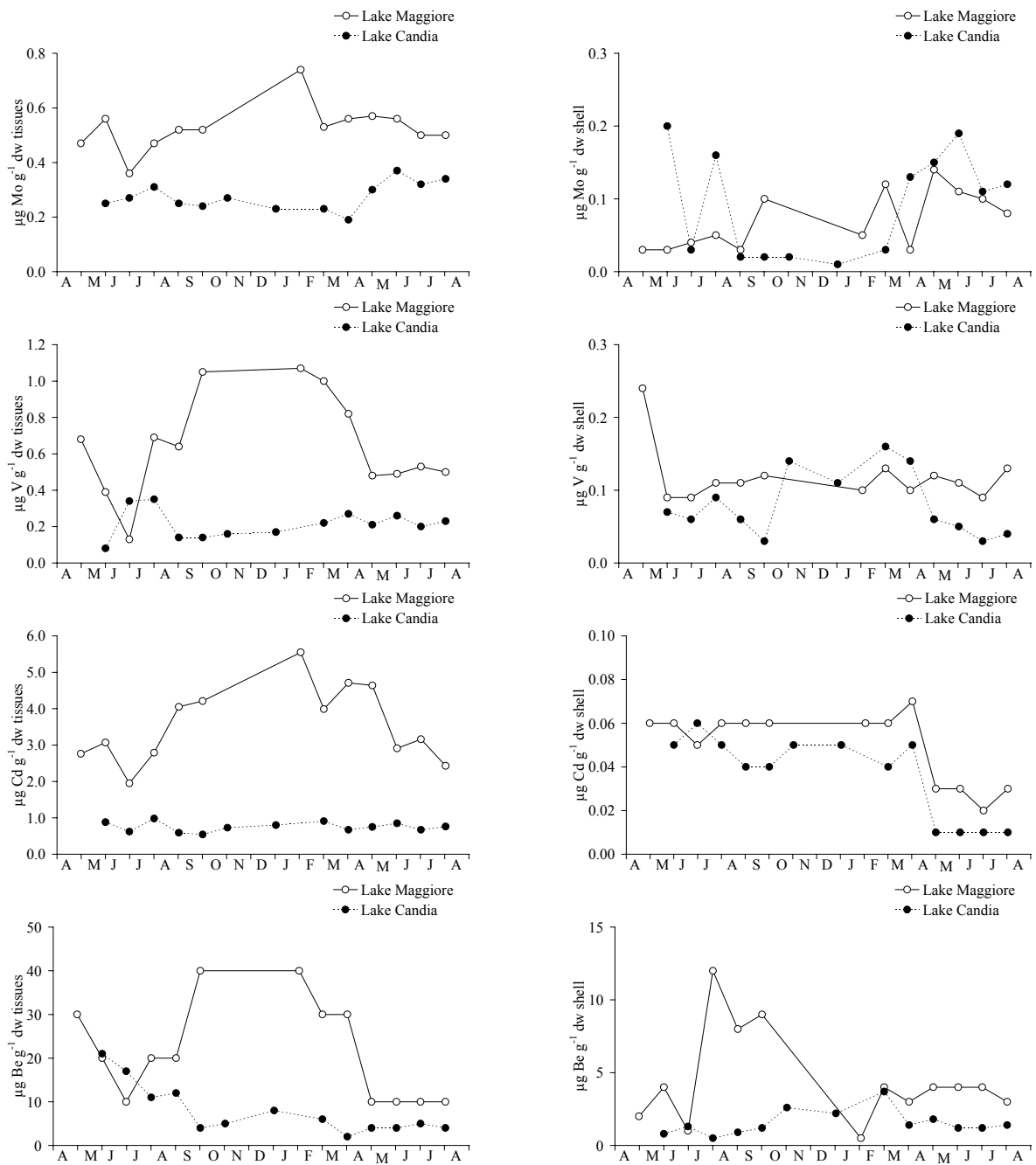


Fig. 2c. Seasonal variations of element concentrations in shell and soft tissues of *Unio pictorum manicus* from Lake Maggiore and Lake Candia.

Tab. 11. Correlation matrix calculated for 91 combinations of elements in mussel tissues from Lake Maggiore (above diagonal) and lake Candia (below diagonal). ** = highly significant ($p < 0.01$), * = significant ($p < 0.04$).

	Zn	Cu	Ca	Fe	Mn	Ni	As	Pb	Co	Cr	Mo	V	Cd	Be
Zn		n.s.	0.801**	n.s.	0.760**	n.s.	n.s.	n.s.	n.s.	n.s.	0.714**	n.s.	0.643*	n.s.
Cu	n.s.		n.s.	n.s.	n.s.	0.635*	n.s.	0.630*	0.602*	n.s.	n.s.	n.s.	n.s.	0.759**
Ca	0.621*	n.s.		n.s.	0.833**	n.s.	n.s.	n.s.	n.s.	n.s.	0.617*	n.s.	n.s.	n.s.
Fe	0.628*	n.s.	0.738**		0.746**	n.s.	n.s.	n.s.	0.570*	0.761**	0.619*	n.s.	n.s.	n.s.
Mn	0.801**	n.s.	0.661*	0.739**		n.s.	n.s.	n.s.	0.607*	0.602*	0.703**	n.s.	0.652*	n.s.
Ni	n.s.	0.704**	n.s.	n.s.	n.s.		n.s.	n.s.	n.s.	n.s.	n.s.	n.s.	n.s.	0.633*
As	n.s.	n.s.	n.s.	n.s.	n.s.	n.s.		n.s.	n.s.	0.606*	0.684**	n.s.	n.s.	n.s.
Pb	n.s.	0.634*	n.s.	n.s.	n.s.	n.s.	n.s.		n.s.	n.s.	n.s.	n.s.	n.s.	n.s.
Co	n.s.	n.s.	n.s.	n.s.	n.s.	0.726**	n.s.	n.s.		n.s.	0.626*	0.683*	n.s.	0.770**
Cr	n.s.	n.s.	n.s.	n.s.	n.s.	n.s.	n.s.	n.s.	n.s.		0.761**	n.s.	0.586*	n.s.
Mo	n.s.	n.s.	0.736**	n.s.	n.s.	-0.699**	n.s.	n.s.	n.s.	n.s.		0.576*	0.808**	n.s.
V	n.s.	n.s.	n.s.	n.s.	n.s.	n.s.	n.s.	n.s.	n.s.	0.844**	n.s.		0.712**	0.870**
Cd	0.654*	0.621*	0.600*	0.771**	0.729**	n.s.	n.s.	n.s.	n.s.	n.s.	n.s.	n.s.		0.609*
Be	n.s.	0.583*	n.s.	n.s.	n.s.	n.s.	n.s.	0.782**	n.s.	n.s.	n.s.	n.s.	n.s.	

The capacity of taxa such as mussels for accumulating various pollutants (for example, trace elements) in their body tissues has been exploited to develop the so-called concentration factor (C.F.). The C.F.s calculated for mussel tissues from Lake Maggiore were higher than those of the corresponding elements from Lake Candia, with the exception of the C.F.s for Be, Mo and Cr, which were higher in Lake Candia. Consequently, except for these three elements, the ratios between the C.F.s of Lake Maggiore and those of the same elements of Lake Candia were higher than 1 for all the elements. While a ratio equal to 1 would suggest that tissue accumulation depends solely on water concentration, a departure from unity indicates that other factors, such as the abundance of available element forms in water and/or food, can affect metal intake and accumulation.

A drawback of the C.F. index is that the concentration in the tissues, which results from the accumulation of elements over a relatively long time, must be compared with the concentration in the water, which reflects the environmental situation at the moment of sampling. To minimize this difficulty we compared the mean element concentrations in the tissues with the mean values in the water. Although the water element concentrations were similar in both lakes, the C.F. of the Lake Maggiore population was distinctly higher for most of the elements. Similar observations have been made by other authors (Johnson *et al.* 1993; Ravera *et al.* 2003), and Metcalfe-Smith *et al.* (1992) observed that mussels from uncontaminated environments were frequently more highly contaminated than those from clean waters.

One of the main reasons for the lack of correlation between element concentration in the water and in the tissues of the organism is the tendency to consider the total element concentration in the water instead of the element forms available to the mussel (e.g., Ravera 2004; Maruo & Orians 2006). Generally, the ionic form of an element is the most readily available, though there are exceptions (e.g., Hg, Pb, Sn). The particulate form may also be important for filter feeding animals like mussels. The relative proportion of the various element

forms varies in relation to the element as well as to environmental conditions. For example, Muntau (1981) found in water samples from an area of Lake Maggiore close to our sampling station that the percentages of elements in particulate form were: Cu 15%; Zn 18% and Cd 39%, and in ionic or weakly chelated form: Cu 62%, Zn 25% and Cd 56%. In spite of continually improving methods for metal species identification and quantification, field studies on aquatic invertebrates in relation to element forms are very scarce (e.g., Lee *et al.* 2006). In spite of drawbacks, however, the use of accumulator organisms in biomonitoring yields reliable results in identifying new pollutants (e.g., artificial radioisotopes) or in estimating temporal and spatial differences in trace element concentrations.

The values of the flesh condition index (F.C.) and those of the condition index (C.I.) were both lower in Lake Maggiore than in Lake Candia mussels, in which the concentration of all the metals in the tissues was lower than in Lake Maggiore (Tab. 8). This is in agreement with the general rule of these indices, which is that their values decrease as the metal concentrations in the tissues increase.

Information on trace element concentrations in mussel tissues is very abundant, less so for the shell and very poor for trace element contents in the shell and tissue biomass. The higher element contents in the tissue and shell biomass of mussels from Lake Maggiore than in the mussels from Lake Candia are due mainly to greater population density, and to a lesser extent to higher element concentrations in the tissues. Mean individual weight, which is greater in Lake Candia mussels, is of negligible importance for mussel biomass. Indeed, the population density of mussels in Lake Maggiore (46.86 individuals m^{-2}) exceeded that in Lake Candia (6.93 individuals m^{-2}) by a factor of about 7. In Lake Maggiore, therefore, mussels can be expected to exert a greater influence on the biogeochemical cycles of elements than in Lake Candia.

Monitoring the seasonal variations of element concentrations in mussel tissues, besides providing infor-

mation on element metabolism over the year, is useful for calculating the actual mean concentration of the elements. Certainly, the wide seasonal variations in element concentrations show the inadequacy of calculating element concentration from samples collected in a single season. The differences between the patterns of variation over time of the same element in shell and tissues may be explained by the turnover time of the elements, which is markedly shorter in the soft tissues than in the shell. Consequently, element concentrations in the tissues may vary over the seasons in relation to the physiological needs of the mussel and the abundance of the elements in available forms in water and food. Conversely, the greatest part of the elements present in the shell is adsorbed from the water to the periostracum, while a relatively smaller amount (except for Ca) is transferred from the mantle to the shell during the whole life-span of the mussel. It follows that, although no seasonal pattern of element concentrations in the shell can be expected, a knowledge of these concentrations over time is essential for estimating the element content in the shell biomass related to the surface unit of the sediments. We found evidence of higher concentrations of several elements (e.g., Fe, Mn, Co, V, Be) in the soft tissues during the cold season. Other authors observed lower metal concentrations in the soft tissues of marine and freshwater bivalves in summer than in winter (e.g., Savari *et al.* 1991). Klaric *et al.* (2004) observed in winter and spring an As concentration increase in the soft tissues, due at least partly to the low nutrition status of mussels in these seasons.

A closer similarity of the seasonal pattern of the metal concentrations in the mussel tissues was observed in Lake Maggiore than in Lake Candia.

The positive correlation between some element pairs could be the result of their metabolic analogy and/or of seasonal variations of the available forms in the environment. Only one negative correlation was observed between the tissue concentrations of Ni and Mo in Lake Candia mussels. This may be due to a competitive inhibition for metabolic sites between these two elements with a similar ionic radius ($\text{Ni}^{+2} = 69\text{\AA}$ and $\text{Mo}^{+4} = 70\text{\AA}$). Markich & Jeffree (1994) demonstrated this mechanism clearly in laboratory experiments. This fact highlights the strong influence of environmental conditions on the seasonal variations of element concentrations in mussels.

In contrast with the similarity in the seasonal pattern of several metal pairs in the same lake, only two metals (Ni and Pb) showed a similar seasonal pattern in the mussel tissues of both lakes, a fact which demonstrates that the behaviour of a given metal in the mussel can vary widely in different environments. Further research into this important problem is clearly required.

The metal content in a population represents the amount of metals immobilized in its biomass, which is equivalent to that transferred to the environment after

the death of the population organisms. This influence on the environment, and particularly on biogeochemical cycles, was estimated by the Influence index (*I.i.*), which shows values higher for both the soft tissues and shell of Lake Maggiore mussels than for those of Lake Candia mussels.

Populations of *U. pictorum mancus* adapted to environments of different trophic state are characterized by marked biological and chemical differences. The population of the meso-oligotrophic Lake Maggiore showed a higher density and a larger total biomass per surface unit despite smaller individual size and a slower growth rate (Ravera *et al.* 2007), as well as higher trace element concentrations in the shell and tissues, compared to the population of the eutrophic Lake Candia. We can therefore conclude that, at least in our case, high trophic can reduce the impact of mussels on their habitat and particularly on biogeochemical cycles. Although the influence of eutrophication on mussels is evident (e.g. Patzner & Müller 2001), it is rather difficult to identify the most important variables affecting the mussel and its capacity to accumulate metals. In the literature there is much more information on the influence of biological factors on metal accumulation in marine mussels (e.g., *Mytilus*) than in freshwater species.

Previous research (Ravera *et al.* 2003b) showed that trace element concentrations (Cu, Zn, Fe and Mn) in *U. pictorum mancus* tissues from eutrophic lakes were lower than those from specimens inhabiting oligo- and meso-oligotrophic lakes, even though the concentrations in the eutrophic lake water were higher than those in the less productive lakes. The present study revealed higher element concentrations in both water and mussel tissues of the meso-oligotrophic Lake Maggiore than in the eutrophic Lake Candia. Two conclusions can be drawn from this comparison: 1) higher trace element concentrations in the water are not necessarily related to the high trophic state of the water body, and 2) high element concentrations in mussel tissues cannot be explained only by high element concentrations in the water; other causes must be involved, such as a high level of food contamination and/or a high percentage of available forms of the element in water.

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