Environmental variables, pesticide pollution and meiofaunal community structure in two contrasting temporarily open/closed False Bay estuaries

S Bollmohr[1,2]*, PJ van den Brink[3,4], PW Wade[5], JA Day[6] and R Schulz[2]

[1]*Zoology Department, University of Johannesburg, 2006 Auckland Park, Johannesburg, South Africa*
[2]*Institute for Environmental Sciences, University Koblenz-Landau, D-76829 Landau/Pfalz, Germany*
[3]*Alterra, Centre for Water and Climate, Wageningen University and Research Centre, PO Box 47,
6700AA Wageningen, The Netherlands*
[4]*Wageningen University, Department of Aquatic Ecology and Water Quality Management, Wageningen
University and Research Centre, PO Box 47, 6700AA Wageningen, The Netherlands*
[5]*Environmental Geoscience Unit, Council for Geoscience, Private Bag X112, Pretoria, 0001, South Africa*
[6]*Freshwater Research Unit, University of Cape Town, Private Bag, Rondebosch 7701, South Africa*

Abstract

Environmental variables (including natural and anthropogenic stressors) and meiobenthic communities were sampled in a 'natural' (Rooiels) and a 'disturbed' (Lourens) estuary in the Western Cape, South Africa, bimonthly for 20 months. A primary aim of the study was to assess if the meiobenthic community structure is driven by different variables when comparing 'natural' versus 'disturbed' system. Due to the much smaller catchment of the Rooiels Estuary, many environmental variables were significantly different ($p < 0.001$) from the variables in the Lourens Estuary, e.g. salinity, temperature, pH, total suspended solids, nitrate and depth. No pesticide concentrations were expected in the Rooiels Estuary due to the absence of agricultural development in the catchment. However, chlorpyrifos (8.9 μg/kg), prothiofos (22.0 μg/kg) and cypermethrin concentrations (0.42 μg/kg) were detected frequently, with the highest concentrations recorded during the summer months. Principal response curve analysis showed that temporal variability between sampling dates explained 42% of the variance in environmental variables and pesticide concentrations and spatial variability between the 2 estuaries explained 58%. Variables contributing most to the differences were higher concentrations of endosulfan, p,p-DDE and nitrate concentrations in the Lourens Estuary and larger grain size and higher salinity at the bottom in the Rooiels Estuary. In general the meiofaunal community in the Rooiels Estuary showed a significantly higher number of taxa ($p < 0.001$), a significantly higher Shannon Wiener Diversity Index ($p<0.001$) and a generally lower meiofaunal abundance with less variability than in the Lourens Estuary. The differences were mostly explained by a higher abundance of *Cypretta* and *Darcythompsonia* in the Rooiels Estuary and a higher abundance of *Thermocyclops* and *Canthocamptus* in the Lourens Estuary. The variables explaining a significant part (14%) of the variance in meiofaunal abundance in the Rooiels Estuary were salinity and temperature, with the Redundancy Analysis indicating that the abundance of most of the taxa increased with higher salinity and temperature, e.g. *Upogebia*, *Nereis*, *Uroma* and nematodes were clearly positively correlated to salinity and temperature. The variables explaining a significant part of the variance in the dataset (43%) within the Lourens Estuary were also salinity and temperature but included chlorpyrifos, nitrate and flow (including river and tidal flow).

Keywords: particle-associated insecticides; meiofauna; estuaries; South Africa

Introduction

Temporarily open/closed estuaries (TOCEs) are the dominant estuary type in South Africa, comprising 73% of all estuaries (Whitfield, 1995). Similar systems, sometimes referred to as 'blind', 'intermittently open', or 'seasonally open' estuaries, are also found in Australia, on the west coast of the USA, in South America and in India (Ranasinghe and Pattiaratchi, 1999). TOCEs are often closed off from the sea, and the lack of freshwater input and tidal exchange may result in changes in various physicochemical variables (e.g. salinity, temperature, flow) and anthropogenic stressors (e.g. pesticides). This spatial and temporal variability within TOCEs can have direct and indirect impacts on the biota (e.g. zooplankton, meiofauna, ichthyofauna, phytoplankton) (Kibirige and Perissinotto, 2003).

Limited long-term monitoring data exists for TOCEs, making it difficult to recognise the degree of variability in these estuaries and the factors influencing their communities. A few studies have sampled different variables on a monthly basis (Nozais et al., 2005, Kibirige and Perissinotto, 2003) and others on a seasonal basis over a period of up to a year (Perissinotto et al., 2000) or several years (James et al., 2008). These studies indicate that there is considerable temporal variability within these systems. The present study is one of the first to examine natural variables, anthropogenic stressors and community responses in such estuaries over a period of 2 years. One of the factors limiting our understanding of South African estuaries is the lack of long-term monitoring data. The National Water Act (Act No. 36 of 1998) in South Africa recognizes basic human water requirements as well as a need to sustain the country's freshwater and estuarine ecosystems in a healthy condition for present as well as future generations (Adams et al., 2002). A National River Monitoring Programme has been in place in South Africa for many years (DWAF, 2006) but no equivalent has yet been designed for estuaries.

Many estuarine and coastal management initiatives worldwide (e.g. North America, Europe and Australia) are required to derive and use environmental quality indices; examples include implementation of the European Water Framework Directive and the UC Clean Water Act (Quintino et al., 2006). Those indices are then used to test for departure from a reference or control condition. Thus to be able to identify the impacts of anthropogenic stressors such as pesticides, it is important to understand the natural spatial and temporal variability in physical and chemical conditions and their impacts on community structure. It is therefore crucial to understand the driving variables by examining natural estuarine systems whose catchments are not associated with any anthropogenic activity. To achieve this, the dynamics of abiotic and biotic variables of a natural system need to be compared with those of a disturbed system using a multivariate approach such as the principal response curve (PRC). The PRC approach has been shown to be valuable in statistically analysing multiple endpoints (Van den Brink and Ter Braak, 1999) and has already been used for biomonitoring by Den Besten and Van den Brink (2005) and Van den Brink et al. (2009).

For the purposes of the current study the Lourens Estuary represents a disturbed system and the Rooiels Estuary an undisturbed or reference system. Previous research (Bollmohr and Schulz, 2009; Schulz et al., 2001) has shown that pesticides contribute significantly to the pollution and associated effects in the Lourens River. Particles entering the estuary are associated with high concentrations of chlorpyrifos (19.6 µg/kg), endosulfan (18.6 µg/kg) and prothiofos (34.0 µg/kg) (Bollmohr et al., 2007), with highest concentrations occurring during the spring and summer season. The present study focused on changes in the epibenthic, hyperbenthic and demersal meiobenthos and zooplanktonic organisms living in the flocculent layer between the sediment and water column.

The importance of these organisms in anthropogenic stressor studies is discussed in Bollmohr et al. (2009c). International meiofaunal studies have tended to focus on community structure in relation to variables such as sediment composition (Sherman and Coull, 1980), salinity (Miliou, 1993) and food sources (e.g. Decho and Castenholz, 1986) that drive its composition. To ascertain the effect of pesticides on meiobenthic abundance, various studies have been performed using sediment toxicity tests (e.g. Chandler and Green, 2001; Bollmohr et al., 2009b) or microcosm experiments (Chandler et al., 1997) but only a few have used biomonitoring in the field (Warwick et al., 1990, Bollmohr et al., 2009c). Very few studies have focused on seasonal fluctuations in meiofauna or zooplankton in South African estuaries in relation to the distribution of physical factors (Nozais et al., 2005, Pillay and Perissinotto, 2009), the state of the mouth (Kibirige and Perissinotto, 2003) and food-web interactions (Perissinotto et al., 2000), and virtually no studies exist on pollutants (Bollmohr et al., 2007; Bollmohr et al., 2009c).

The aims of this study were twofold: To identify the driving environmental variables (including natural and anthropogenic stressors) in a reference and a disturbed system, and to identify the extent to which these variables reflect differences in meiofaunal assemblage structure within the flocculent layer.

Material and methods

Study area

The study took place in the False Bay region of the Western Cape Province, South Africa (Fig. 1), an area characterised by cool, wet winters and warm dry summers. Measurements within the 2 TOCE study sites occurred in the lower reaches of the estuaries and always during low tide.
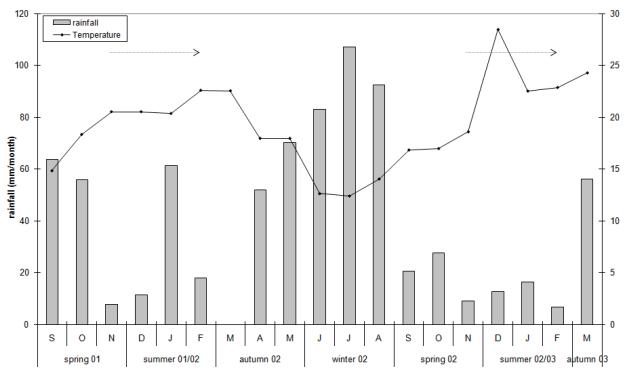
Figure 2
Monthly rainfall and temperature data (obtained from Verelegen weather station at 80 m altitude) during the study period (September 2001 to March 2003). The dotted lines indicated the spraying season of pesticides.

The Lourens Estuary

The Lourens Estuary mouth is located at 34°06′S and 18°49′E; the river (20 km long) rises in the Hottentots Holland Mountains, flows through intensive agricultural areas, and the town of Somerset West, after which it enters False Bay at the Strand (Cliff and Grindley, 1982). The estuary is about 0.71 km² in area with a tidal range of 1.48 m. Mean annual runoff (MAR) is approximately 122 x 10⁶ m³ (Whitfield and Bate, 2007). Land use in the Lourens River sub-catchment of 140 km² comprises forestry, agriculture, residential areas and light industries. A large section of the upper catchment is privately-owned agricultural land with vineyards and apple, pear and plum orchards on which pesticide application takes place annually between August and February (Schulz, 2001).

The Rooiels Estuary

The Rooiels Estuary mouth is situated at 34°18′S and 18°49′E, with a catchment that lies within the southern extension of the Hottentots Holland Mountain Reserve and flows through the Kogelberg Nature Reserve. The total distance from the river mouth to the end of the longest tributary is only 9 km (Whitfield and Bate, 2007). The estuary is about 0.122 km² in area with a tidal range of 1.48 m. The MAR is 10 x 10⁶ m³ and the entire catchment (21 km²) lies within a nature reserve, with a small holiday resort located near the mouth (Heydorn and Grindley, 1982).

Pesticide application and seasonal weather data

The 20-month investigation from August 2001 to March 2003 included dry summer months (November to March) with infrequent rainfall (Fig. 2) and wet winter months from May to August. Organophosphates, such as chlorpyrifos (686 kg/ha) and prothiofos (87 kg/ha), organochlorines such as endosulfan (158 kg/ha) and pyrethroids such as cypermethrin (8 kg/ha) and fenvalerate (5 kg/ha), are frequently applied to pears, plums and apples between August and February before fruit is harvested (Dabrowski et al., 2002); frequency of application is highest during the dry summer season from November to January (Schulz, 2001) (dotted lines in Fig. 2).

Physico-chemical measurements

Various physicochemical parameters were measured every

2 weeks during the study period. Measurement always took place in the morning during low tide. The flow rate, which included the river flow and the ebb tidal flow, was measured every metre across each estuary using a flow meter (Höntzsch Co, Waiblingen, Germany) and the discharge towards the sea was calculated.

Dissolved oxygen, pH, temperature and conductivity of the estuarine waters were measured with electronic meters from Wissenschaftliche Technische Werkstätten GmbH, Weilheim, Germany. Salinity was established by converting conductivity values into salinity using an automatic converter (Fofonoff and Millard, 1983). Total suspended solids (TSS) were measured with a turbidity meter (Dr Lange, Düsseldorf, Germany) and nutrients were measured using photometric test kits from Macherey & Nagel, Düren, Germany. Total organic carbon (TOC) of the sediment was quantified by removal of all organic matter, followed by gravimetric determination of sample weight-loss (ASTM, 2000). Grain-size analysis was performed by dry-fraction sieving of sediment samples previously treated with hydrogen peroxide. Sediment was divided into 7 fractions: silt and clay (< 0.063 mm), very fine sand (0.063 to 0.125 mm), fine sand (0.125 to 0.25 mm), medium sand (0.25 to 0.5 mm), coarse sand (0.5 to 1 mm), very coarse sand (1to 2 mm) and gravel (>2 mm). Each fraction was weighed and expressed as a percentage of the total mass. Four sediment samples (top 15 cm) per site (within a 1 m² transect) were taken for analysis.

Micro-algal biomass was determined in the water column and in the sediment by measuring the concentration of chlorophyll *a*. Four replicates of 2 ℓ water samples were filtered through Whatman (GF/C) glass-fibre filters. The chlorophyll *a* was extracted by grinding the filters with 10 mℓ of 99% acetone. After centrifuging, the light absorbance at 665 nm of the supernatant was determined before and after adding 2 drops of 0.1 N HCl. Chlorophyll *a* concentrations were calculated using the equation of Hilmer (1990). Nine replicates of benthic micro-algal samples were taken within a 1 m² transect with a 15 mℓ syringe. Sediment chlorophyll *a* was extracted from the top 1.5 cm of sediment in the same way as the water column chlorophyll *a*.

Pesticide analyses

Insecticides associated with suspended particles were accumulated continuously by a suspended-particle sampler, from which samples were collected every 2 to 3 weeks. The suspended-particle sampler consisted of a plastic container (500 mℓ) with a screw-on lid pierced in the centre by a hole 20 mm in diameter and an open glass jar suspended directly under the hole in the lid. The samplers were attached approximately 50 mm above the river bed to a metal stake fixed into the sediment.

Suspended particles were extracted twice with methanol and concentrated using C18 columns. Insecticides were eluted with hexane and dichlormethane and analysed at the Forensic Chemistry Laboratory, Department of National Health, Cape Town. Measurements were made with gas chromatographs (Hewlett-Packard 5890, Avondale, PA, USA) fitted with standard Hewlett-Packard electron-capture, nitrogen phosphorus and flame-photometric detectors, with a quantification limit of 0.1µg/kg and overall mean recoveries between 79 and 106% (Schulz et al., 2001). Further details of extraction and analysis of pesticides is described in Schulz et al. (2001).

Table 1
Mean (± standard error) physicochemical parameters measured in the Rooiels and Lourens estuaries from September 2001 to March 2003 (n=44). Asterisks indicate significant differences between the 2 estuaries (*$p < 0.05$; **$p < 0.01$; ***$p < 0.001$))

	Unit	Lourens River estuary	Rooiels River estuary
Outflow	m³/sec	1.00 ± 1.15	0.46 ± 0.19
Mean depth ***	m	0.51 ± 0.18	1.07 ± 0.14
Salinity (surface)***	g/kg	1.69 ± 2.07	5.19 ± 5.74
Salinity (bottom)***	g/kg	6.29 ± 8.18	19.5 ± 11.6
Temperature (surface)***	°C	19.2 ± 4.4	18.0 ± 4.2
Temperature (bottom)***	°C	19.4 ± 4.4	16.8 ± 3.3
Nitrate***	mg/ℓ	3.5 ± 3.4	0.00 ± 0.00
Nitrite*	mg/ℓ	0.077 ± 0.124	0.003 ± 0.006
Ammonia	mg/ℓ	0.007 ± 0.013	0.01 ± 0.02
Ortho-phosphate*	mg/ℓ	0.39 ± 0.32	0.17 ± 0.25
Oxygen (surface)	mg/ℓ	9.33 ± 1.88	8.75 ± 1.28
Oxygen (bottom)*	mg/ℓ	8.80 ± 2.00	7.75 ± 1.01
pH***		7.6 ± 0.5	6.7 ± 1.00
Total suspended solids***	mg/ℓ	63.9 ± 39.6	1.03 ± 1.21
Total organic carbon	%	1.29 ± 0.62	1.29 ± 0.54
Grain size**	μm	164 ± 79.9	239 ± 30.1
Chlorophyll *a* (water)**	mg/ℓ	3.76 ± 3.98	0.39 ± 0.46
Chlorophyll *a* (sediment)**	mg/ℓ	8.98 ± 8.45	2.29 ± 1.75

Meiobenthic sampling and community analysis

The sampling approach attempted to reduce the sampling error caused by small-scale variability in the sediments (Livingston, 1987) and to sample the organisms living in the flocculent layer between the sediment and overlying water column. An area of 10 m x 300 mm was actively sampled just above the bottom using a zooplankton net of 63 μm mesh size. Four replicates were collected and within each replicate the zooplanktonic and meiobenthic organisms were identified to the lowest possible taxonomic level. Estimates of abundance were based on 8 sub-samples in a matrix box with 64 chambers, of which 8 chambers were randomly selected for identification and counting of all organisms. To assess temporal and spatial patterns of benthic faunal assemblages, the community metrics density (number of individuals/m³), number of taxa, and H′ diversity (Shannon and Wiener, 1948) were calculated.

Statistics

Multivariate techniques are often used to link field concentrations of chemicals and other environmental parameters with invertebrate responses (Leps and Smilauer, 2003). In this study ordination methods were used for dimension reduction, i.e. to reduce the data set to a 2-dimensional summary, which can be presented as an ordination diagram. In this way a graphical summary of the data set is obtained. Redundancy analysis (RDA) is the direct form of principal component analysis (PCA) that focuses the analysis on that particular part of the variance that is explained by external explanatory variables. The significance of the environmental variables was evaluated using Monte Carlo permutation tests (Van den Brink et al.,

2003). Redundancy analysis was performed separately for the 2 estuaries.

The differences in community composition between the Lourens and Rooiels Estuaries were visualised by principal response curves (PRC) (Van den Brink and Ter Braak, 1999) using the CANOCO software package Version 4.5. The PRC is based on the RDA ordination technique, a constrained form of PCA. The analysis results in a diagram showing the first principal component of the differences in community structure on the y-axis and deviations in time between the 2 sites on the x-axis, with the Rooiels Estuary as reference. The species weights are shown in a separate diagram, and indicate the affinity each has with the major determinant indicated. The species with high positive weights show responses similar to those indicated by the PRC; those with negative weights, show responses opposite to the response indicated by PRC. Species with near-zero weight show a response very dissimilar to the response indicated by PRC, or no response at all.

The data sets of the physicochemical variables and pesticides were tested for significant differences between the 2 estuaries using the student t-test and were also analysed using PRC. In these analyses the variables were centred and standardised before analysis to account for differences in measurement scale (Kersting and Van den Brink, 1997). Guidance on the use of PRC to analyse monitoring data is described in Van den Brink et al. (2009).

Results

Temporal and spatial comparison of environmental variables and particle- associated pesticides

Due to the different type and size of catchment, the magnitudes of many environmental variables differed significantly between the 2 estuaries (Table 1). The most significant differences ($p<0.001$) were for depth, salinity, temperature, nitrate, pH and TSS. In general, the Lourens Estuary is a shallower system when compared to the Rooiels (0.51 ± 0.18 m compared to 1.07 ± 0.14 m), with lower salinities in the bottom (6.29 ± 8.18 compared to 19.5 ±11.6) and surface waters (1.69 ± 2.07 compared to 5.19 ± 5.74). Both bottom (19.4 ± 4.43°C) and surface temperatures (19.2 ± 4.44°C) were higher in the Lourens than in the Rooiels (18.0 ± 4.20°C and 16.8 ± 3.30°C, respectively). The water contained significantly more total suspended solids in the Lourens (63.9 ± 39.6 mg/ℓ) than in the Rooiels (1.03 ± 1.21 mg/ℓ), and the pH was significantly higher in the Lourens than in the Rooiels Estuary.

As a result of anthropogenic inputs, nitrate, nitrite and orthophosphate concentrations were significantly higher in the Lourens than the Rooiels Estuary ($p<0.01$), as was phytoplankton biomass measured as chlorophyll *a* concentrations in the water column (3.76 ± 3.98 mg/ℓ compared to 0.39 ± 0.46 mg/ℓ) and sediment (8.98 ± 8.45 mg/m² compared to 2.29 ± 1.75 mg/m²).

Cypermethrin (CYP) , fenvalerate (FEN), endosulfan total (END), p,p-DDE, chlorpyrifos (CPFS) and prothiofos (PTF)
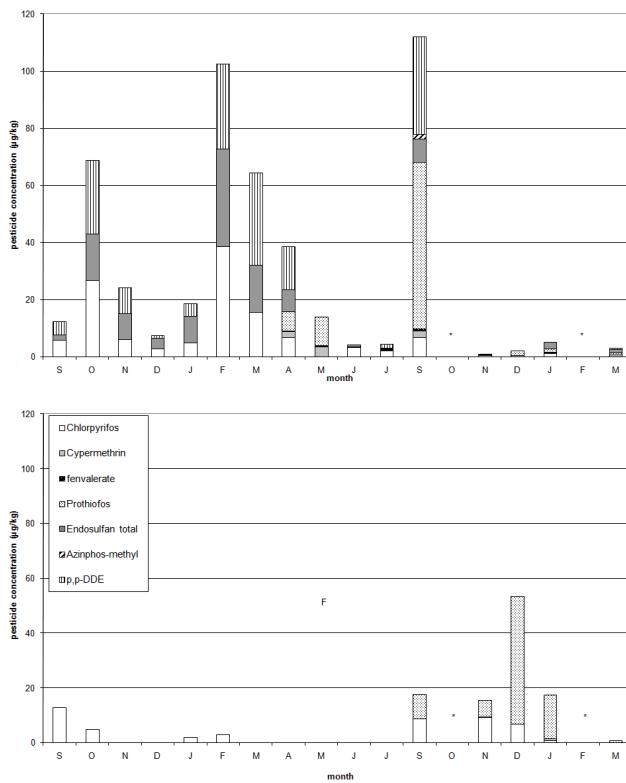
Figure 4
Diagram showing the first component of the PRC of the differences in measured environmental variables and pesticides between Lourens and Rooiels River estuaries. Forty-two percent of the total variation could be attributed to differences between sampling dates, and the other 58% to differences between the estuaries; 43% of the latter are displayed in the diagram. The parameter weights shown in the right part of the diagram represent the affinity of each parameter with the response shown in the diagram. For clarity, only parameters with a weight > 0.5 or < -0.5 are shown.

Figure 3
Monthly composition of particle-bound pesticide concentrations detected at Lourens and Rooiels estuaries from September 2001 to March 2003. Asterisks indicate that no samples were collected.

were found in varying concentrations during almost all months within the Lourens Estuary (Table 2, Fig. 3), with the highest concentrations occurring during spring (September-November) and autumn (March-May). Despite the absence of agricultural activities in the catchment of the Rooiels, chlorpyrifos, PTF and CYP were frequently detected. The highest concentrations occurred during the summer months (November-January) and PTF concentrations were even higher than those in the Lourens Estuary (Table 2, Fig. 3). However, concentrations of FEN, total - END, p,p-DDE and azinphos-methyl (AZPM) were found to be significantly higher in the Lourens Estuary.
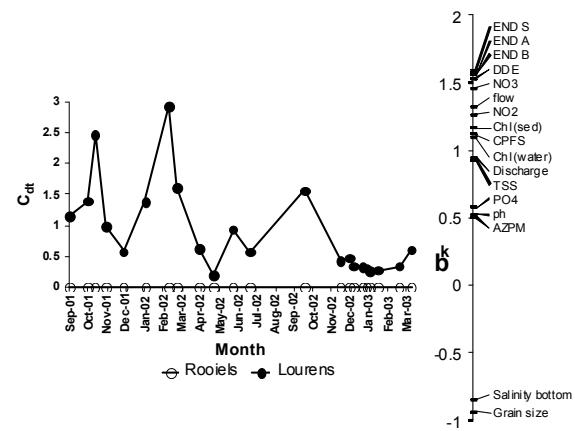
In the Lourens River PTF (34 µg/kg) occurred in the highest concentrations, followed by p,p-DDE (31.0 µg/kg), CPFS (19.6 µg/kg), total - END (18.6 µg/kg), CYP (2.84 µg/kg) and fenvalerate (0.64 µg/kg) (Table 2). Cypermethrin (92%) and CPFS (92%) were most frequently detected, followed by total - END (86%) and PTF (85%). In the Rooiels Estuary concentrations of PTF (22.0 µg/kg) were highest, followed by CPFS (8.9 µg/kg) and CYP (0.42 µg/kg). Chlorpyrifos showed the highest frequency of detection (50%), followed by PTF(39%) and CYP (23%).

The PRC diagram (Fig. 4) indicates differences in environmental variables and particle-bound pesticide concentrations between the 2 estuaries and their importance in contributing to the overall differences between them. There is high temporal variability, with 42% of the variation being attributable to differences between sampling dates. Spatial differences that were not linked to temporal variability explained 58% of the variation. Variables that contributed most to the differences between the estuaries were higher concentrations of END, p,p-DDE and

Table 2
Mean (± standard error) concentrations (µg/kg) of particle-bound pesticides in the Lourens and Rooiels estuaries and 90th percentile from September to March 2003 (n=27). Asterisks indicate significant differences between the 2 estuaries (*$p < 0.05$; **$p < 0.01$)

	Lourens Estuary			Rooiels Estuary		
	Mean	% of detection[1]	90th percentile	Mean	% of detection[1]	90th percentile
Cypermethrin	0.92 ± 1.09	92	2.84	0.31 ± 0.84	23	0.42
Fenvalerate*	0.19 ± 0.22	77	0.64	0.00 ± 0.00	0	0.00
DDE**	8.42 ± 13.6	5	31.0	0.00 ± 0.00	0	0.00
Endosulfan total**	6.03 ± 9.40	86	18.6	0.00 ± 0.00	0	0.00
Azinphos-methyl	0.07 ± 0.31	50	0.00	0.00 ± 0.00	0	0.00
Chlorpyrifos	7.04 ± 10.5	92	19.6	3.21 ± 5.46	50	8.90
Prothiofos	6.74 ± 15.8	85	34.0	12.90 ± 28.03	39	22.0

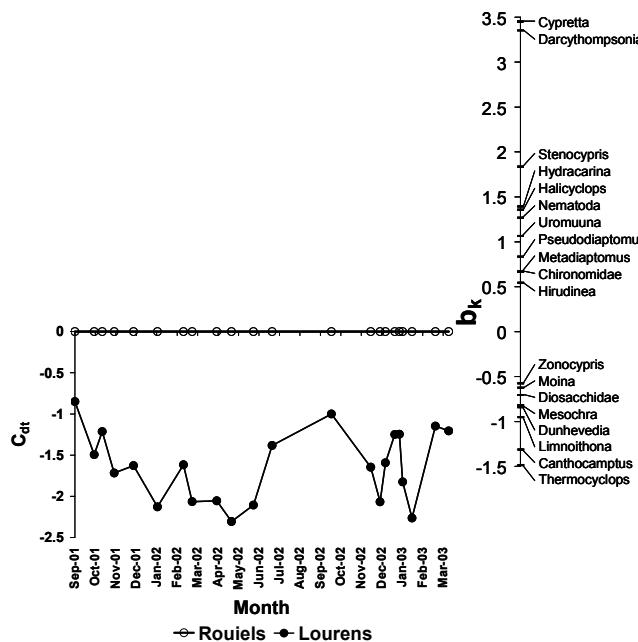[1] *Percentage of samples in which the pesticide was detected*

Figure 5
Diagram showing the first component of the PRC of the differences in species composition between the Rooiels and Lourens estuaries. Fifty-two per cent of the total variation in species composition could be attributed to differences between sampling dates, and the other 48% to differences in species composition between the estuaries; 44% of the latter are displayed in the diagram. The taxa weights shown in the right part of the diagram represent the affinity of each taxon with the response shown in the diagram. For clarity, only species with a weight > 0.5 or < −0.5 are shown.
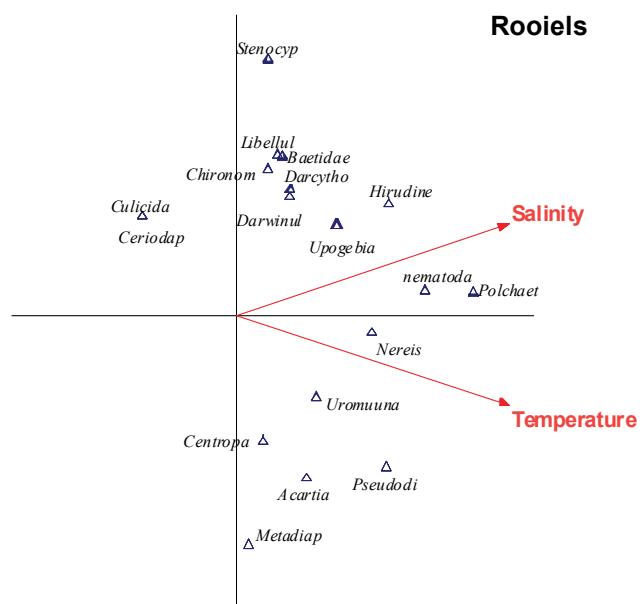

Figure 6
RDA triplot resulting from an analysis using the environmental variables and pesticide concentrations that explain a significant part of the seasonal variation in the species composition in the Rooiels Estuary as explanatory variables and meiofauna taxa composition as taxa data. The 2 significant variables explain 14% of the seasonal variation in species composition, of which 71% is displayed on the first axis and another 29% on the second axis. For clarity only the 18 most important taxa out of 45 are shown.

nitrates in the Lourens and larger grain size and higher salinity at the bottom of the Rooiels. Flow, nitrite, CPFS and chlorophyll were also important variables contributing to differences between the 2 estuaries.

Temporal and spatial comparison of meiofauna communities

In general the meiofaunal community in the Rooiels Estuary showed a significantly higher number of taxa ($p < 0.001$), a significantly higher Shannon Wiener Diversity Index ($p < 0.001$) and a generally lower abundance, with less variability over time, than the Lourens Estuary (Table 3). The PRC analysis (Fig. 5) indicated that the 2 community structures were very different from one another; 52% of the difference was explained by temporal differences and 48% by differences between the 2 estuaries. Differences in the fauna were explained mostly by more *Cypretta* and *Darcythompsonia* in the Rooiels, and more *Thermocyclops* and *Canthocamptus* in

the Lourens. Overall the difference is attributable to the greater number of species in the Rooiels Estuary, which correlates with the higher diversity index (Table 3).

The meiofaunal community and its driving variables

Within the RDA biplot the most important taxa and significant environmental variables are shown in Figs. 6 and 7. The variables explaining a significantly part (14%) of the seasonal variation in invertebrate abundance in the Rooiels Estuary were salinity and temperature (Fig. 6). The RDA indicated that the abundance of most of the taxa were increasing with higher salinity and temperatures and taxa such as *Upogebia*, *Nereis*, *Uroma* and the nematodes were positively correlated with salinity and temperature.

The meiofaunal community in the Lourens Estuary is significantly influenced by a number of variables that are probably the result of anthropogenic impacts (Fig. 7). The variables explaining a significant part of the variance in the dataset (43%) were salinity and temperature, CPFS, nitrate and flow. *Mesochra* decreased with higher CPFS and nitrate concentrations, and increased with higher salinity and temperature.

Table 3
Mean (± standard error) number of taxa, total abundance and Shannon Wiener Diversity Index in the Lourens and Rooiels estuaries and 90th percentile from September to March 2003 (n=27). Asterisks indicate where there are significant differences (p < 0.001) between the 2 estuaries.

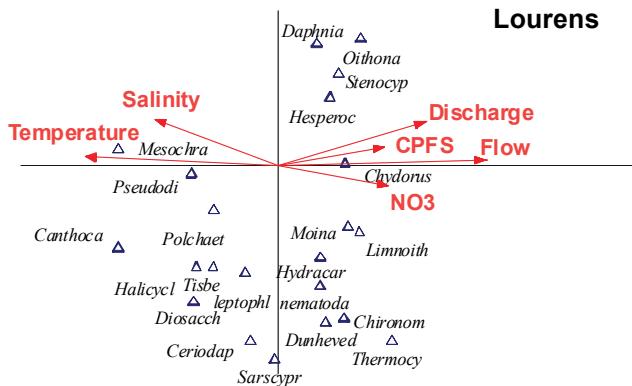|	Lourens Estuary		Rooiels Estuary	
	Mean	90th percentile	Mean	90th percentile
Number of taxa***/m³	6.8 ± 2.7	10	10.3 ± 2.7	14
Total abundance/m³	6 600 ± 12 391	16 423	2 442 ± 1 318	3 951
Shannon Diversity Index***	1.1 ± 0.4	1.6	1.6 ± 0.3	1.9

Figure 7
RDA triplot resulting from an analysis using the environmental variables and pesticide concentrations that explain a significant part of the seasonal variation in the species composition in the Lourens Estuary as explanatory variables and meiofauna taxa composition as taxa data. The 6 significant variables explain 43% of the seasonal variation in species composition, of which 69% are displayed on the first axis and another 12% on the second axis. For clarity only the 20 most important taxa out of 45 are shown.

Furthermore, the taxa *Pseudodiaptomus*, *Canthocamptus* and polychaetes were negatively correlated with CPFS and nitrate concentrations. In general, oligohaline taxa (e.g. *Moina* and *Limnoithona*) decreased in abundance at higher salinities.

Discussion

Temporal and spatial comparison of environmental variables and particle-associated pesticides

Due to a lack of agricultural and urbanisation influences in its catchment the Rooiels Estuary is more pristine than the Lourens River, with lower particle-bound pesticide concentrations and also lower nutrient concentrations resulting in lower chlorophyll *a* values. The lower MAR in the Rooiels Estuary results in a more saline environment with evident stratification in salinity, temperature and oxygen and with higher salinity and lower oxygen values in bottom waters, a typical feature for natural, relatively deep TOCEs during the dry season (Kibirige and Perissinotto, 2003)

Despite the fact that no agricultural activities are documented for its catchment, pesticides such as CPFS, PTF and CYP were sometimes recorded in the Rooiels, particularly during the 2003 summer when concentrations were actually higher than those in the Lourens. The input of pesticides to the Rooiels Estuary could originate from one or more of the following sources:

Studies have shown that volatilisation can cause off-site movement of pesticides via atmospheric transport and subsequent contamination of surface waters (Harman-Fetcho et al., 2000; LeNoir et al., 1999). The Western Cape climate is characterised by strong south-easterly winds during summer with wind directions from agricultural activities southeast towards the Rooiels Estuary. Most pesticide application using mist-blow techniques occurs during the summer season (Schulz, 2001), which also coincides with the period of high concentrations found in the estuary. However, high mountains (Hottentots Holland Mountain) are situated between the 2 catchments, which could serve as a partial barrier for atmospheric pesticide transport. Additionally it is puzzling why endosulfan

was not found in the Rooiels Estuary, since it is also heavily applied during the summer season. Due to the south-easterly wind direction, the current within False Bay is directed from the Lourens and other estuaries with 'agricultural' catchments towards the Rooiels Estuary (Taljaard et al., 2000). Particle-bound pesticides with a high K_{oc} and low solubility (as prothiofos, CPF and cypermethrin) could be transported by the currents in False Bay and enter the Rooiels Estuary on the incoming tide. However, pesticides tend to bind to particles which in turn are likely to settle either within the estuary (Bollmohr et al., 2009c) or within the bay. Additionally it is hard to explain why p,p-DDE, which is present in high concentrations in the Lourens River, and has the highest K_{oc} and the longest DT_{50}, has not been detected in the Rooiels Estuary.

Residential areas are situated on the southern side of the estuary and the houses are mainly occupied during the holidays (Heydorn and Grindley, 1982). The primary holiday period occurs during summer (December and January), which coincides with the time when high pesticide concentrations were found. Many household products contain CPFS and CYP.

Bollmohr et al. (2007) ranked the pesticide concentrations detected in the Lourens River in terms of their risk towards the environment, by comparing these with various effect concentrations (Chandler and Green, 2001; Chandler et al., 1994; Leonard et al., 2001). Chlorpyrifos and total-END posed the highest risk towards both freshwater and marine communities in the Lourens Estuary. Within the Rooiels Estuary there is a possibility that CPF may pose a risk towards harpacticoid species during certain summer months (Bollmohr et al., 2009b; Chandler and Green, 2001). Toxicity data and threshold values for PTF are inadequate; therefore it is not possible to assess a probable risk for this pesticide towards the aquatic environment in either estuary.

Temporal and spatial comparison of meiofaunal communities

The prolonged period of mouth closure that most TOCEs experience generally leads to low levels of taxonomic diversity (Perissinotto et al., 2000). Hartwell and Claflin (2005) classified a Shannon Wiener Index of <1.5 for macrobenthos in permanently open estuaries as 'low', and above 2.4 as 'high', while a Shannon Wiener index for zooplankton in the impacted, temporarily-open Mpenjati Estuary ranged between only 1.0 and 1.20 (Kibirige and Perissinotto, 2003). Although the Lourens Estuary supports more freshwater species than the Rooiels (Bollmohr et al., 2009c) due to a higher MAR, overall meiofaunal diversity is lower than in the Rooiels, which is characterised by a greater number of taxa, most of which are of marine or estuarine origin. The higher meiofaunal abundance in the Lourens Estuary can be explained by the higher organic input arising from a greater MAR when compared to the Rooiels. The lower variability in meiofaunal abundance in the Rooiels Estuary is probably a result of the more stable estuarine environment when compared to the Lourens.

Species contributing most to spatial differences were the ostracod *Cypretta* and harpacticoid copepod *Darcythompsonia* for the Rooiels Estuary (with the highest positive weighting), and *Thermocyclops* and *Canthocamptus* for the Lourens Estuary (with the highest negative weighting). Ostracods are mainly influenced by temperature, salinity, depth and grain size (Curry, 1999), and sewage pollution as a result of low oxygen (Mezquita et al., 1999). Based on toxicity data they are not significantly more sensitive to pollution than other crustaceans

(Sanchez-Bayo, 2006) but have a slower recovery rate after pollution events (Dernie et al., 2003). The combination of low oxygen, salinity, low grain size (mud), higher temperature and more frequent pesticide concentrations might, however, have resulted in the loss of ostracods from the Lourens Estuary. No information is available on the harpacticoid genus *Darcythompsonia* and further work on the sensitivity of this genus is required. *Thermocyclops* and *Canthocamptus* are known as freshwater and/or estuarine species and might have been carried into the Lourens Estuary from upstream, whereas the salinity in the Rooiels Estuary might have been too high for these taxa to survive. Insufficient toxicity data are available for these genera to reach conclusions regarding the causality of the correlations of their abundances with the pesticide levels in either estuary.

The meiofaunal community and its driving variables

The RDA analysis identified salinity and temperature as the only significant variables driving meiofaunal community structure in the Rooiels Estuary. These variables have been identified as the most important for natural estuarine systems by various other authors (e.g. Austen, 1989; Willmer et al., 2000; Nozais et al., 2005). Grain size and water depth are often regarded as primary variables in determining aspects of community structure in meiobenthic organisms (Soetaert et al., 1994; Curry, 1999; Sherman and Coull, 1980), but these variables showed little variation within the Rooiels Estuary.

The community structure of the Lourens Estuary is, however, significantly influenced by a number of variables, including salinity, temperature, flow and anthropogenic stressors such as nitrate and CPFS. The above result suggests interactions between the effects of both natural and anthropogenic drivers on the meiofaunal community. The abundance of the genus *Mesochra*, as an example, increased with decreasing CPFS concentrations and increasing salinity. A microcosm toxicity test on intact estuarine sediment with the marine copepod *Amphiascus tenuiremis*, by Chandler et al. (1997), showed acute effects after 96 h at CPFS levels ranging from 21 to 33 μg/kg. A related study by Chandler and Green (2001) found that chronic full-life-cycle exposures to concentrations of 11 to 22 μg/kg sediment-associated CPF resulted in consistently significant reductions in production of copepodites and nauplii by *A. tenuiremi*s. These concentrations were exceeded by the 90th percentile found in the Lourens Estuary (19.6 μg/kg). A laboratory study by Bollmohr et al. (2009b) indicated that exposure to CPFS of 5.9 to 6.4 μg/kg, which was exceeded on 37% of the sampling occasions during the current field programme, was detrimental to the survival of *Mesochra parva*. The above laboratory study also revealed that *M. parva* survival decreased by 60 to 90% due to CPFS exposure and increased by 10 to 50% as salinity increased from 3 to 15.

The frequency of occurrence and mean concentration of CPFS in the Rooiels Estuary (FO = 50% of samples, mean concentration = 3.21 ± 5.46 μg/kg) was very different to the frequency of occurrence and mean concentration of CPFS in the Lourens Estuary (FO = 92% of samples, mean concentration = 7.04 ± 10.5 μg/kg). These results clearly indicate an elevated and more prolonged pollution problem in the Lourens when compared to the Rooiels system. However, more work is required to understand the response of various meiofaunal taxa to this pollutant under different salinity conditions.

References

ADAMS JB, BATE GC, HARRISON TD, HUIZINGA P, TALJAARD S, VAN NIEKERK L, PLUMSTEAD EE, WHITFIELD AK and WOOLDRIDGE TH (2002) A method to assess the freshwater inflow requirements of estuaries and application to the Mtata estuary, South Africa. *Estuar. Coast. Shelf Sci.* **25** 1382-1393.

ASTM (2000) *Standard Test Methods for Moisture, Ash and Organic Matter of Peat and Other Organic Soils.* Method D 2974-00. American Society for Testing and Materials. West Conshohocken, PA.

AUSTEN MC (1989) Factors affecting estuarine meiobenthic assemblage structure: a multifactoria; microcosm experiment. *J. Exp. Mar. Biol. Ecol.* **130** 167-187.

BOLLMOHR S, VAN DEN BRINK PJ, WADE PW, DAY JA and SCHULZ R (2009) Spatial and temporal variability in particle-bound pesticide exposure and their effects on benthic communities structure in a temporarily open estuary. *Estuar. Coast. Shelf Sci.* **82** 50-60.

BOLLMOHR S and SCHULZ R (2008) Seasonal changes of macroinvertebrate community in a Western Cape River receiving nonpoint-source insecticide pollution. *Environ. Toxicol. Chem.* **28** 809-817.

BOLLMOHR S, DAY JA and SCHULZ R (2007) Temporal variability in particle-bound pesticide exposure in a temporarily open estuary, South Africa. *Chemosphere* **68** 479-488.

BOLLMOHR S, HAHN T and SCHULZ R (2008) Interactive effect of salinity decrease, salinity adaptation and chlorpyrifos exposure on an estuarine harpacticoid copepod, *Mesochra parva*, in South Africa. *Ecotoxicol. Environ. Saf.* **72** 756-764.

CHANDLER GT, COULL BC, SCHIZAS NV and DONELAN TL (1997) A culture-based assessment of the effects of chlorpyrifos on multiple meiobenthic copepods using microcosms of intact estuarine sediment. *Environ. Toxicol. Chem.* **16** 2339-2346.

CHANDLER GT and GREEN AS (2001) Developmental stage specific life-cycle bioassay for assessment of sediment-associated toxicant effects on benthic copepod production. *Environ. Toxicol. Chem.* **20** 171-178.

CLIFF S and GRINDLEY JR (1982) Report No.17 Lourens (CSW7). In: Heydorn AEF and Grindley JR (eds.) *Estuaries of the Cape (Part 2) Synopsis of Available Information on Individual Systems.* CSIR Report No. 416. CSIR, Stellenbosch, South Africa. 39 pp.

CURRY BB (1999) An environmental tolerance index for ostracodes as indicators of physical and chemical factors in aquatic habitats. *Palaeogeogr. Palaecol.* **148** 51-63.

DABROWSKI JM, PEALL SKC, REINECKE AJ, LIESS M and SCHULZ R (2002) Runoff -related pesticide input into the Lourens River, South Africa: basic data for the exposure assessment and risk mitigation at the catchment scale. *Water Air Soil Pollut.* **135** 265-283.

DECHO AW and CASTENHOLZ RW (1986) Spatial patterns and feeding of meiobenthic harpactiocid copepods in relation to resident microbial flora. *Hydrobiol.* **131** 87-96.

DERNIE KM, KAISER MJ, RICHARDSON EA and WARWICK RM (2003) Recovery of soft sediment communities and habitats following physical disturbance. *J. Exp. Mar. Biol. Ecol.* **285-286** 415-434.

DEN BESTEN PJ and VAN DEN BRINK PJ (2005) Bioassay response and effects on benthos after pilot remediations in the delta of the rivers Rhine and Meuse. *Environ. Pollut.* **136** 197-208.

FOFONOFF TL and MILLARD JR RC (1983) Algorithms for computation of fundamental properties of seawater. UNESCO Technical papers in marine science No. 44.

HARMAN-FETCHO JA, McCONELL LL, RICE CP and BAKER JE (2000) Wet deposition and air-water gas exchange of currently used pesticides to a subestuary of the Chesapeake Bay. *Environ. Sci. Technol.* **34** 1462-1468.

HARTWELL SI and CLAFIN LW (2005) Cluster analysis of contaminated sediment data: Nodal analysis. *Environ. Toxicol. Chem.* **24** 1816-1834.

HEYDORN AEF and GRINDLEY JR (1982) Estuaries of the Cape, Part II. Synopses of Available Information on Individual Systems. Report No. 7: Gamtoos (CMS48). *CSIR Research Report No.* **406**.

HILMER T (1990) Factors Influencing the Estimation of Primary Production in Small Estuaries. Ph.D. Thesis, University of Port Elizabeth, South Africa.

JAMES NC, WHITFIELD AK and COWLEY PD (2008) Long-term stability of the fish assemblages in a warm-temperate South African estuary. *Estuar. Coast. Shelf Sci.* **76** 723-738.

KERSTING K and VAN DEN BRINK PJ (1997) Effects of the insecticide Dursban4E (active ingredient chlorpyrifos) in outdoor experimental ditches: responses of ecosystem metabolism. *Environ. Toxicol. Chem.* **16** 251-259.

KIBIRIGE I and PERISSINOTTO R (2003) The zooplankton community of the Mpenjati Estuary, a South African temporarily open/closed system. *Estuar. Coast. Shelf Sci.* **58** 727-741.

LENOIR JS, MCCONNELL LL, FELLERS GM, CAHILL TM and SEIBER JN (1999) Summertime transport of current-use pesticides from California's central valley to the Sierra Nevada Mountain Range, USA. *Environ. Toxicol. Chem.* **18** 2715-2722.

LEPS J and SMILAUER P (2003) *Multivariate Analysis of Ecological Data using Canoco.* Cambridge University Press, Cambridge, UK. 269 p.

LEONARD AW, HYNE RV, LIM RP, LEIGH KA, LE J and BECKETT R (2001) Fate and toxicity of endosulfan in Naomi River Water and Bottom Sediment. *J. Environ. Qual.* **30** 750-759.

MEZQUITA F, HERNANDEZ JR and RUEDA J (1999) Ecology and distribution of ostracods in a polluted Mediterranean river. *Palaeogeogr. Palaeol.* **148** 87-103.

MILIOU H (1993) Temperature, salinity and light induced variations on larval survival and sex ration of *Tisbe holothuriae* Humes (Copepoda: Harpactiocida). *J. Exp. Mar. Biol. Ecol.* **173** 95-109.

NOZAIS C, PERISSINOTTO R and TITA G (2005) Seasonal dynamics of meiofauna in a South African temporarily open/closed estuary (Mdloti Estuary, Indian Ocean). *Estuar. Coast Shelf Sci.* **62** 325-338.

PERISSINOTTO R, WALKER DR, WEBB P, WOOLDRIDGE TH and BAILEY R (2000) Relationships between zoo- and phytoplankton in a warm temperate, semi-permanently closed estuary, South Africa. *Estuar. Coast Shelf Sci.* **51** 1-11.

PILLAY D and PERISSINOTTO R (2009) Community structure of epibenthic meiofauna in the St. Lucia Estuarine Lake (South Africa) during a drought phase. *Estuar. Coast Shelf Sci.* **81** 94-104.

QUINTINO V, ELLIOT M and RODRIGUES AM (2006) The derivation, performance and role of univariate and multivariate indicators of benthic change: Case studies at differing spatial scales. *J. Exp. Mar. Biol. Ecol.* **330** 368-382.

RANASINGHE R and PATTIARATCHI C (1999) Circulation and mixing characteristics of a seasonally open tidal inlet: a field study. *Mar. Freshwater Res.* **50** 281-290.

SÁNCHEZ-BAYO F (2006) Comparative acute toxicity of organic pollutants and reference values for crustaceans. I. Branchiopoda, Copepoda and Ostracoda. *Environ. Pollut.* **139** 385-420.

SCHULZ R (2001) Rainfall-induced sediment and pesticide input from orchards into the Lourens River, Western Cape, South Africa: importance of a single event. *Water Res.* **35** 1869-1876.

SCHULZ R, PEALL, SKC, DABROWSKI JM and REINECKE AJ (2001) Current-use insecticides, phosphates and suspended solids in the Lourens River, Western Cape, during the first rainfall event of the wet season. *Water SA* **27** 65-70.

SHANNON CE and WIENER N (1948) A mathematical theory of communication. *A.T. & T. Tech. J.* **27** 379-423.

SHERMAN KM and COULL BC (1980) The response of meiofauna to sediment disturbance. *J. Exp. Mar. Biol. Ecol.* **45** 59-71.

SOETAERT K, VINEX M, WITTOECK J, TULKENS M, VAN GANSBEKE D (1994) Spatial patterns of the Westerschelde meiobenthos. *Estuar. Coast. Shelf Sci.* **39** 367-388.

VAN DEN BRINK PJ and TER BRAAK CJF (1999) Principal response curves: Analysis of time-dependent multivariate responses of a biological community to stress. *Environ. Toxicol. Chem.* **18** 138-148.

VAN DEN BRINK PJ, DEN BESTEN PJ, BIJ DE VAATE A and TER BRAAK CJF (2009) The use of the principal response curves techniques for the analysis of multivariate time series from biomonitoring studies. *Environ. Monit. Assess.* **152** 271-281.

VAN DEN BRINK PJ, VAN DEN BRINK NW and TER BRAAK CJF (2003) Multivariate analysis of ecotoxicological data using ordination: Demonstrations of utility on the basis of various examples. *Australas. J. Ecotox.* **9** 141-156.

WARWICK RM, PLATT HM, CLARKE KR, AGARD J and GOBIN J (1990) Analysis of macrobenthic and meiobenthic community structure in relation to pollution and disturbance in Hamilton Harbour, Bermuda. *J. Exp. Mar. Biol. Ecol.* **138** 119-142.

WHITFIELD AK (1995) Available Scientific Information on Individual South African Estuarine Systems. WRC Report No. 577/1/95. Water Research Commission, Pretoria, South Africa. 204 pp.

WHITFIELD AK and BATE G (2007) The Freshwater Requirements of Intermittently Open Cape Estuaries. WRC Report No. 1581/1/2007, Pretoria, South Africa.

WILLMER P, STONE G and JOHNSTON I (2000) *Environmental Physiology of Animals.* Blackwell Science, Oxford, UK. 644 pp.

Appendix 1 Taxon list (average ± standard deviation)			
Taxa	Abbreviation #	Lourens River estuary (n=31)	Rooiels River estuary (n=29)
Halicyclops	*Halicycl*	51 ± 122	128 ± 177
Thermocyclops	*Thermocy*	218 ± 509	1 ± 7
Macrocyclops		0 ± 0	3 ± 11
Paracyclops		3 ± 13	0 ± 0
Hesperocyclops	*Hesperoc*	9 ± 33	0 ± 0
Limnoithona	*Limnoith*	53 ± 155	2 ± 11
Oithona		3 ± 12	0 ± 0
Smirnoviella		1 ± 6	0 ± 0
Acartia		5 ± 28	2 ± 13
Pseudiaptomus	*Pseudodi*	80 ± 204	228 ± 498
Centropagidae	Centropa	0 ± 0	43 ± 231
Metadiaptomus	*Metadiap*	5 ± 28	33 ± 87
Canthocamptus	*Canthoca*	2768 ± 4326	175 ± 245
Darcythompsonia	*Darcytho*	0 ± 0	575 ± 558
Mesochra	*Mesochra*	9142 ± 22978	235 ± 496
Diosacchidae	Diosacch	111 ± 284	0 ± 0
Tisbe	*Tisbe*	52 ± 134	0 ± 0

Stenocypris	*Stenocyp*	1 ± 3	104 ± 181
Sarscypridopsis	*Sarscypr*	11 ± 24	12 ± 64
Darwinula	*Darwinul*	0 ± 0	5 ± 13
Zonocypris		5 ± 9	0 ± 0
Cypretta	*Cypretta*	0 ± 0	558 ± 642
Daphnia	*Daphnia*	5 ± 19	0 ± 0
Moina	*Moina*	166 ± 771	0 ± 0
Chydorus	*Chydorus*	50 ± 283	0 ± 0
Ceriodaphnia	*Ceriodap*	52 ± 224	2 ± 12
Dunhevedia	*Dunheved*	676 ± 3662	2 ± 11
Hydracarina		4 ± 14	35 ± 37
Chironomidae	Chironom	24 ± 53	31 ± 36
Nereis		2 ± 6	20 ± 35
Polchaetlarvae	Polychaet	10 ± 24	120 ± 249
Uromuuna	*Uromuuna*	0 ± 0	27 ± 53
Turbellaria		0 ± 0	4 ± 12
Rotifera		14 ± 82	0 ± 0
Culicidae		0 ± 2	0 ± 2
Oligochaet		3 ± 9	5 ± 14
Perlidae		1 ± 3	0 ± 0
Baetidae	Baetidae	6 ± 15	2 ± 8
Leptophlebiidae	Leptophl	1 ± 6	2 ± 11
Porifera		0 ± 0	5 ± 18
Hirudinea	Hirudine	0 ± 0	15 ± 33
Libellulidae	Libellul	0 ± 2	1 ± 5
Upogebia	*Upogebia*	0 ± 0	9 ± 20
Nematoda	Nemtatoda	22 ± 43	143 ± 170
Amphipoda		0 ± 0	4 ± 9

#*appearing in Figs. 6 and 7*