

THE MULTIVARIATE STATISTICAL ANALYSIS OF ABIOTIC PARAMETERS OF THE LAKE TANGANYIKA SUB CATCHMENT

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Abstract

Results from multivariate statistical analysis of abiotic parameters conducted during the dry season at 20 accessible sites on 8 rivers, 2 lakes and a dam covering the Lake Tanganyika sub catchment are discussed. Standard methods were used to determine the levels of abiotic parameters from water samples. Physical parameters including DO, EC, Eh, turbidity, temperature, pH and secchi transparency were measured in situ while chlorophyll a was determined in the laboratory. Nutrients such as NO_3^- , SiO_2 , PO_4^{3-} and Fe^{2+} were determined along with HCO_3^- . Factor analysis resulted in four factors including increased primary productivity, redox conditions, dissolution, and reduction processes. Processes including dissolution, diffusion, adsorption, absorption, nitrification, denitrification, mixing and reduction along with the anthropogenic activities, increased photosynthetic activity of algae and the geomorphology of the ecosystems contribute to the variation of the abiotic parameters. It is recommended that quantification of river flows, sediment load and nutrient budget at various sampling points be determined seasonally for proper evaluation of the hydrologic and limnological functioning of the ecosystem.

Keywords: Multivariate statistics, Malagarasi Wetland, Abiotic parameters

INTRODUCTION

The limnology of tropical rivers and lakes along with dams has recently received a paramount and growing attention due to pressures from land, atmospheric, and riverine inputs. The Lake Tanganyika sub-catchment includes the Malagarasi-Muyovozi Wetland Ecosystem in the Western Tanzania, a polymictic fresh water ecosystem designated as first Tanzanian Ramsar site and the world's 1024th Ramsar site since 2000 (Nkotagu and Athuman, 2008).

It is estimated that during the last 10 years the depth of the wetland has systematically been reduced to a maximum of about 2 m. The wetland is located in western Tanzania about 200 km NE of Lake Tanganyika as shown in figure 1. The wetland is characterized with biodiversity including varieties of flora and fauna such as fish of which 50 species are known with some being endemic, crocodiles, hippopotamus and other microorganisms.

These tropical ecosystems support the life of many people and other organisms within the Lake Tanganyika sub catchment and hence form a unique ecosystem. However, very little

information on the assessment of limnological data in this sub-catchment is known.

The present study therefore trucked down the multivariate statistical analysis of the abiotic parameters from various surface water bodies including rivers, lakes and a dam in the Lake Tanganyika sub-catchment as detected at various accessible points.

MATERIALS AND METHODS

Study area

The study was conducted for six weeks during the dry season commencing September to October 2004. The study covered the rivers Igombe, Lugufu, Luiche, Makere, Malagarasi, Moyowosi, Ruchugi, and Ugalla along with the lakes Sagara and Nyamagoma as well as the Igombe Dam within the eastern side of the Lake Tanganyika sub-catchment (Fig. 1).

Geology of the study area

The river is located predominantly at the mesoproterozoic sandstones of about 1,200 million years old with lithology covering shales, quartzites, dolomitic limestones, gneisses and basalts along with quaternary sediments at some places (Pina *et al.*, 2004). These rocks are exposed differently at various

places along the riverbed thus influencing the geomorphologic of the river behaviour accordingly. Shallow portions of the river are observed to be underlain by hard quartzite, quartzitic sandstones and or gneissic rocks that resist both chemical and physical weathering. However, the deepest parts of the river are observed to be dominated by dolomitic limestones and basalts as bedrocks that are easily chemically weathered. The river is located within the western arm of the Eastern African Rift System therefore its geomorphologic behaviour is essentially both geologically and tectonically controlled.

Sampling

Sampling sites were initially decided basing on water depths as measured at various positions on the rivers, lakes and the dam using a SCUBA device. Water samples were collected at selected depth intervals up to maximum depths in half and one-litre plastic bottles filled to the brim using a 2-l water sampler. However, at some rivers, sampling was conducted in a horizontal transect across the river at three positions including the centre, right and left banks.

Aliquots of 400 ml and 800 ml of each water sample were filtered using pore size glass fibre 47 mm size filter papers. The filtrate aliquots were transported on ice to the laboratory for Silica (SiO₂), Nitrate (NO₃⁻), Phosphate (PO₄³⁻) and Iron (Fe²⁺) determinations. The remaining unfiltered water samples were similarly transported to the laboratory for alkalinity determination. Filters were placed into a test tube on to which 10 ml of 90% ethanol were added. The test tubes were then wrapped with aluminium foil, marked and stored overnight in a cooler at 4°C ready for chlorophyll *a* readings the following day.

In situ physical parameters including electrical conductivity (EC), dissolved oxygen (DO), pH and temperature were measured at each sampling site using a Multi Probe meter 340i model. In addition, turbidity and water transparency were also measured using a HACH turbidimeter 2100P model and a 20 cm diameter Secchi disk respectively.

Laboratory work

The filtered water samples were stored at 4°C before analysis. Fluorescence readings for chlorophyll *a* were taken using a Fluorometer before and after acidifying the sample residues with 0.1 N HCl. The absorbance readings were then converted into concentrations as chlorophyll *a* using the formula according to McIntyre (2004 unpublished data):

$$\text{Chl } a \text{ (}\mu\text{g l}^{-1}\text{)} = 0.003 \times (F_{\text{before acid}} - F_{\text{after acid}}) \times \frac{\text{Ethanol Extraction Volume (ml)}}{\text{Volume of Filtered Water (L)}}$$

Unfiltered water samples were tested for alkalinity using a titrimetric method with 0.1 N HCl and results expressed as HCO₃⁻ (mg l⁻¹) as explained by APHA (1998). Nutrients including SiO₂, NO₃⁻, PO₄³⁻ and Fe²⁺ were determined from the filtered water samples using a HACH Spectrophotometer DR/2010 model according to HACH (2002).

Data analysis and interpretation

The data were analysed by the SPSS 11.0 package and other statistical analyses following Davis (1986).

RESULTS AND DISCUSSION

Descriptive Statistics

The results for the descriptive statistics of the abiotic parameters (Tab. 1) indicate high mean salinity from the 64 samples. This is attributed to increased dissolution of minerals and washing in nutrients from anthropogenic sources (Nkotagu and Athuman, (2008) Accepted. However, the mean pH of 7.72 is a good indicator that the water is well buffered as shown by the high mean alkalinity (59.726 mg l⁻¹).

The mean levels of nutrients show a significant variation consequent to the photosynthetic activities of algae and the anthropogenic activities in the catchment area along with several processes such as absorption, adsorption, nitrification, denitrification in the water column under redox conditions.

Correlation Analysis

The results from the correlation matrix between the abiotic parameters (Tab. 2) using the two tailed Pearson correlation show positive and negative correlation between any two parameters consequent to ecological and geomorphologic differences. Temperature correlates positively with pH, weakly with transparency, salinity, HCO_3^- , and SiO_2 . This indicates that warm surface waters result in increased photosynthetic activity as indicated by high abundance of chlorophyll *a* and dissolved oxygen and thus leading to increased consumption of CO_2 .

The positive correlation between pH and the HCO_3^- supports the observed relationship. High positive correlation between chlorophyll *a* and turbidity suggest primary productivity to be the main cause of the later in many parts of the study area. This suggestion is supported by the high negative correlation between chlorophyll *a* and transparency. The positive correlation among the nutrients SiO_2 , PO_4^{3-} and NO_3^- indicates that these nutrients have a common source (mainly anthropogenic). The redox potential correlates negatively with dissolved oxygen, pH, temperature, salinity and alkalinity consequent to redox conditions.

Factor Analysis

Factor analysis (Thurstone, 1931) is used to present the structure of studied data by means of their grouping and classification as well as for space dimension reduction of the analyzed parameters.

The Principal Component Analysis (PCA) and Varimax rotation with Kaiser Normalization (Kaiser 1958) were used to obtain results for the factor analysis. Four rotated factors were extracted (Tab. 3) as the controlling measures to the abiotic parameters within the sub-catchment.

Factor 1

Factor 1 is highly positively loaded with NO_3^- , DO, chlorophyll *a* and turbidity and negatively loaded with water transparency. This factor may generally be referred to as a primary productivity factor showing that water

transparency was low due to high turbidity caused by the increased photosynthetic algae as indicated by a positive relationship between chlorophyll *a* and turbidity. The concentration of NO_3^- increased with DO and chlorophyll *a* indicating that primary productivity increases with increased nutrient concentration. .

Factor 2

Factor 2 is highly loaded with DO, pH, temperature and salinity and showed that the DO relates directly to the pH, temperature and the salinity of water. However, these parameters in this factor are observed to relate negatively with the redox potential of water showing that the redox conditions affect the oxygen levels leading to high CO_2 in the water as a result lowering the pH of water.

Factor 3

The factor is highly positively loaded with PO_4^{3-} , SiO_2 , NO_3^- , temperature and salinity along with the alkalinity of water. This can be thought as a generalized dissolution process factor indicating that the dissolution of HCO_3^- , PO_4^{3-} , SiO_2 and NO_3^- as facilitated by high temperature through water mass overturns lead to high salinity.

Factor 4

This factor is positively loaded with Fe^{2+} and water depth. The factor may be referred to as a reduction process factor. This indicates that the reduction of ferric iron to ferrous iron increases with increasing depth of water.

Cluster Analysis

Cluster analysis (Tryon, 1939) combines different algorithms for classification. In this study three major clusters of abiotic parameters were resulted (Fig. 2). The clustering indicates that parameters are clustered according to biological process favouring turbidity, NO_3^- as a favoured nutrient in the process and dissolved oxygen.

The second cluster favours salinity to be primary due to HCO_3^- , pH, SiO_2 , PO_4^{3-} and water temperature implying that dissolution of anthropogenic salts under favourable

temperatures and pH may be responsible for the salinity of the water in the area.

Finally, the third cluster involves transparent, depth, Eh and Fe^{2+} indicating that the depth of water as expected influences the concentration of the above parameters. Light penetration decreases with depth due to increased chlorophyll *a* and suspended sediments. However, Eh and Fe^{2+} increase with depth due to reduction process.

CONCLUSIONS AND RECOMMENDATIONS

The data in the present study conclude that the variation of the abiotic parameters at the Lake Tanganyika sub-catchment is controlled by factors including increased primary productivity, redox conditions, dissolution, nitrification, denitrification, mixing and reduction along with anthropogenic activities.

The suspended sediments close to the Malagarasi delta and non-point pollutants such as fertilizers at the Kasulu-Kibondo Bridge and animal excreta near Igombe dam could also be contributing heavily to the fluctuation of nutrient within the sub-catchment.

It is recommended that future work focuses on the statistical approaches in the quantification of flow, sediment load and nutrient budget at various points along the rivers, lakes and the dam in both dry and rain seasons in order to fully understand the hydrological and limnological functioning of the Lake Tanganyika sub-catchment.

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NOMENCLATURE

SPSS 11.0 = Statistical Package for Social Scientists version 11.0

F = Fluorescence of the sample, which is equivalent to the absorbance on a Spectrophotometer and

Chl *a* = Chlorophyll *a* in $\mu\text{g l}^{-1}$.

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List of tables

Table 1.0: Descriptive statistics of abiotic parameters

Parameter	N	Minimum	Maximum	Mean	Std. Deviation
EC ($\mu\text{S cm}^{-1}$)	64	32.00	544.00	294.793	127.445
HCO_3^- (mg l^{-1})	64	18.30	140.30	59.726	26.770
Temp ($^{\circ}\text{C}$)	64	20.23	27.28	25.254	1.491
Turb (NTU)	64	2.13	68.52	16.800	18.039
Chl <i>a</i> ($\mu\text{g l}^{-1}$)	64	0.01	144.18	16.365	37.194
SiO_2 (mg l^{-1})	64	2.70	35.30	14.271	5.532
pH	64	6.39	8.52	7.719	0.518
DO (mg l^{-1})	64	0.08	9.16	4.652	2.387
NO_3^- (mg l^{-1})	64	0.44	2.64	1.196	0.549
Trans (m)	64	0.13	1.93	1.041	0.616
Depth (m)	64	0.00	4.00	0.970	1.034
PO_4^{3-} (mg l^{-1})	64	0.01	0.16	0.045	0.034
Fe^{2+} (mg l^{-1})	64	-0.01	0.02	0.003	0.008
Eh (mV)	64	-101.25	30.00	-51.850	32.488

Table 2.0: Correlation matrix showing levels of relationships between parameters

	Trans	Depth	Turb	pH	DO	Temp	EC	Eh	HCO_3^-	SiO_2	NO_3^-	PO_4	Fe^{2+}	Chl <i>a</i>
Trans	1													
Depth	.368	1												
Turb	-.627	-.087	1											
pH	.169	-.019	.055	1										
DO	-.211	.027	.261	.427	1									
Temp	.392	.063	-.084	.511	.287	1								
EC	.417	.193	-.193	.575	.087	.642	1							
Eh	-.180	.002	-.055	-.983	-.423	-.467	-.546	1						
HCO_3^-	-.084	.096	.181	.451	.256	.496	.708	-.410	1					
SiO_2	-.166	-.178	.247	.367	.365	.433	.465	-.327	.634	1				
NO_3^-	-.534	-.272	.454	.081	.264	.050	.056	-.058	.391	.404	1			
PO_4	-.303	-.201	.228	.133	.088	.086	.331	-.080	.512	.439	.405	1		
Fe^{2+}	-.218	.063	.142	-.241	.063	-.239	-.149	.213	-.057	-.088	-.100	-.049	1	
Chl <i>a</i>	-.431	-.065	.736	.092	.426	.207	-.024	-.065	.323	.431	.576	.301	-.072	1

Table 3.0: Rotated component matrix^a showing factor loadings with abiotic parameters

Parameter	Component			
	1	2	3	4
Turb (NTU)	0.853			
Trans (m)	-0.823	0.269		0.208
Chl <i>a</i> (µg l ⁻¹)	0.786	0.146	0.245	
NO ₃ ⁻ (mg l ⁻¹)	0.652		0.381	-0.287
Eh (mV)		-0.914	-0.159	0.117
pH		0.908	0.217	-0.140
DO (mg l ⁻¹)	0.506	0.605		0.181
Temp (°C)	-0.167	0.588	0.489	
HCO ₃ ⁻ (mg l ⁻¹)	0.172	0.288	0.842	0.111
EC (µS cm ⁻¹)	-0.326	0.445	0.757	0.111
PO ₄ ³⁻ (mg l ⁻¹)	0.281	-0.148	0.712	-0.238
SiO ₂ (mg l ⁻¹)	0.328	0.288	0.668	-0.112
Depth (m)	-0.234			0.816
Fe ²⁺ (mg l ⁻¹)	0.221	-0.271	-0.104	0.547

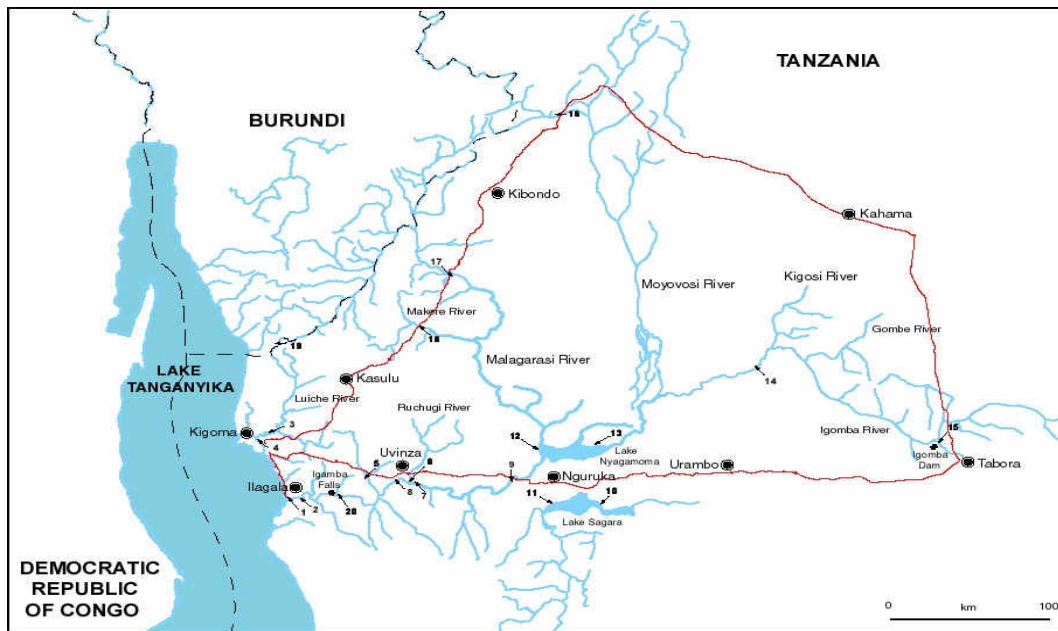


Fig. 1.0: The Lake Tanganyika sub catchment showing the sampling sites 1 to 20.

Key: 1=Malagarasi River at the Delta, 2=Malagarasi River at the Ilagala Ferry, 3=Luiche River at the Kigoma-Kasulu Bridge, 4=Luiche River at the Delta in Ujiji, 5=Lugufu River at Jackobsen's Farm, 6=Ruchugi River at the Kigoma-Uvinza Bridge, 7=Malagarasi River at the Nyanza Salt Mine, 8=Malagarasi River at the Uvinza-Mpanda Bridge, 9=Malagarasi River at the Uvinza-Nguruka Bridge, 10=Lake Sagara at Amerika, 11=Ugalla River at Katumba, 12=Moyovosi River mouth at the Lake Nyamagoma, 13=Igombe River at Mtega in Kaliua, 14=Igombe River at Uyowa in Urambo, 15=Igombe Dam in Tabora, 16=Moyovosi River at the Kagera-Kigoma Bridge, 17=Malagarasi River at the Kasulu-Kibondo Bridge, 18=Makele River at the Bridge, 19=Malagarasi River at the Burundi-Tanzania Border, 20=Malagarasi River at the Igamba Water Fall

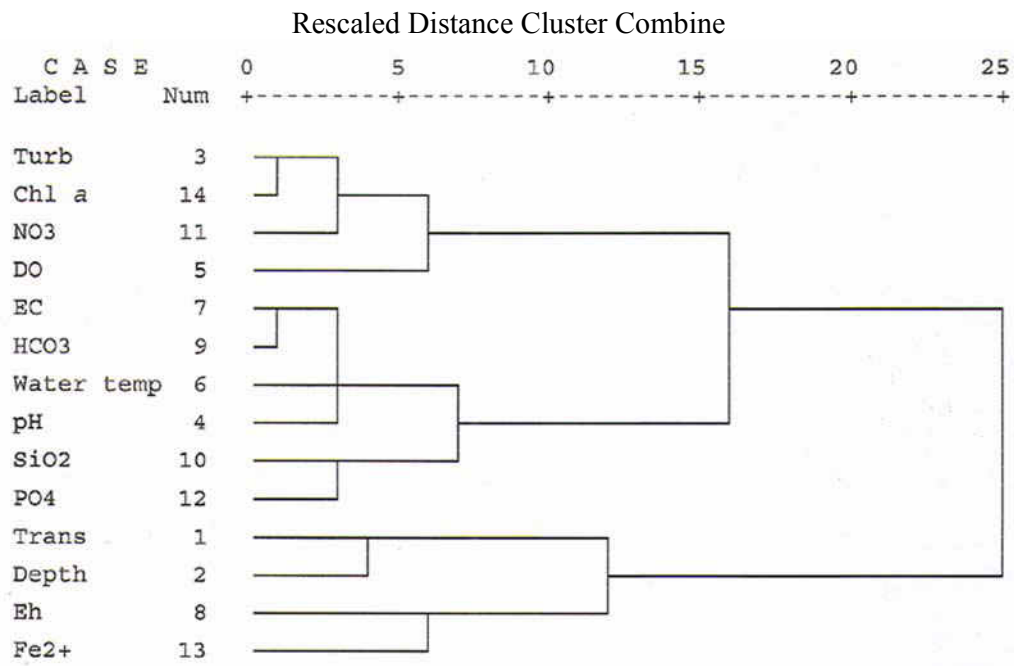


Fig. 2.0: The dendrogram results for the sampling sites