Water Balance and Variations of Nutrients and Major Solutes along a River Transect Through the Okavango Delta, Botswana

Mangaliso John Gondwe*, Wellington R.L. Masamba§ and Michael Murray-Hudson♣

Abstract
Wetland goods and services provide a means of survival for many riparian rural communities in developing countries, particularly in the Sub-Saharan Africa. To sustain the provision of goods and services for human use, spatial and temporal variations in the quantity and quality of water in a given wetland should be adequately understood and managed. In this study we monitored the quantity and quality of surface water through the Okavango Delta for 2 years, from June 2008 to June 2010. Specifically we monitored precipitation over the Delta and surface inflow and outflow (via the Okavango and Thamalakane Rivers respectively) for water quantity, nutrients, cation and anion levels in the surface water along a river transect from Mohembo through the Delta to Lake Ngami. The study revealed that the water balance during the study period was similar to earlier reports. Most of the water inflow was via the Okavango River while surface outflow via the Boro River accounted for about 2% of total inflow. Calcium (Ca) and bicarbonate (HCO$_3^-$) were the most dominant cation and anion respectively across the river transect sampled except in Lake Ngami where sodium (Na) was the dominant cation. Total nitrogen (0.222±0.018 to 0.698±0.035 mg L$^{-1}$) and total phosphorus (0.042±0.005 to 0.131±0.024 mg L$^{-1}$) levels across the Delta were generally low. The N:P ratios suggest spatial and temporal P and N growth limitations in the Delta. N limitation seems to be ameliorated by biological nitrogen fixation. Since nutrient poor aquatic systems tend to be very sensitive to pollution, excessive P input into the Delta should be avoided to prevent eutrophication.

Keywords: Okavango Delta, tropical wetland, major ions, nutrient levels, nutrient limitation

Introduction
Wetlands have historically been regarded as waste lands. Consequently, many wetlands around the world have been extremely vulnerable to degradation or reclamation for agricultural and urban developments. The world has lost over 50% of its original wetlands over the past century (Barbier 1993). For instance, the United States of America and South Africa have respectively lost about 54% and 50% of their original wetlands during the last century. Wetlands perform many important functions which include water storage, water purification, modification of local climate, groundwater recharge, chemical attenuation and transformation and biological diversity (plants and animals) (Junk 2002). Due to high human population density and rampant poverty in many African countries, wetland goods and services provide a means of survival for many riparian rural communities (Schuyt 2005).

Hydrology is an integral component of wetland ecosystems. Water quantity and quality determine the capacity of wetlands to perform the various functions and services. While hydrological variations, such as in flood-pulsed systems, ensure wetland functioning, anthropogenically induced changes in these systems tend to be destructive of the wetland’s functioning. For instance, high volume discharge of wastewater can increase wetland productivity which may affect other wetland functions such as main-
tenance of biological diversity. It is therefore important to monitor water quantity and quality in natural wetland ecosystems vulnerable to human perturbations such as wastewater discharge.

In this study we monitored the quantity and quality of surface water through the Okavango Delta for two years from June 2008 to June 2010. Specifically we monitored precipitation over the Delta and surface inflow and outflow via the Okavango and Thamalakane Rivers respectively, for water quantity and nutrient, cation and anion levels in the surface water along a river transect from Mohembo through the Delta to Lake Ngami. Other physicochemical variables measured during the study have been reported in Gondwe and Masamba (2016). The objective of this study was to re-evaluate the water balance and water quality in the Okavango Delta in order to provide a baseline for a long term monitoring programme.

Materials and Methods

Study area

The freshwater Okavango Delta (19° 17’ S, 22° 54’ E) in northern Botswana is one of the world’s largest inland deltas and is situated at the distal end of the Cubango-Okavango River Basin (CORB). The total catchment area of the CORB is 165,000 km², of which 82% is located in Angola drained by the Cubango and Cuito rivers which join near the Angola-Namibia border to form the Okavango River (Wilk et al. 2006). Much of the water that sustains the Delta comes through the Okavango River as runoff from the central Angolan highlands which experience about 1200 mm of rainfall per annum. Local precipitation over the Delta which falls between November and May is lower at approximately 490 mm/a (McCarthy et al. 2000). However, because of the large Delta area, the total volume of rainwater that annually falls over the Delta area is approximately 30-40% of the flood water brought in via the Okavango River (Snowy Mountains Engineering Corporation 1987, 1990). At the lower end of the Panhandle, the flood water spreads out into an alluvial fan (Delta) of swamps, channels and islands covering between 6,000 and 13,000 km² depending on prevailing floods and local precipitation (Gieske 1997). Currently the main distribution channel after the bifurcation of the Okavango River is the Jao-Boro channel which more or less bisects the Delta into western and eastern areas (Figure 1). The Jao-Boro channel passes through the urban village of Maun as the Thamalakane River and goes on to Lake Ngami. It has been estimated that the surface outflow in the Thamalakane River measured at Maun is only 2% of the total annual inflow via the Okavango River and precipitation over the Delta. This is because approximately 98% of the Delta’s total inflow water is lost to the atmosphere through evapotranspiration estimated at 1500 mm y⁻¹ (Wilson and Dincer 1976; Snowy Mountains Engineering Corporation 1987; Ramberg and Wolski 2008).

Study sites

Water quality parameters were monitored along the Okavango-Jao-Boro-Thamalakane River system at ten (10) sites which were: Mohembo, Drosky, Sepopa, Guma, Nxaraga, Boro, Maun, Tsanekona, Toteng and Lake Ngam (Figure 1). The justification for the selection of these sites was given in Gondwe and Masamba (2016), and was based on the belief that management and conservation of the Okavango Delta requires a clear understanding of the quantity and quality of the flood water as it flows the length of the Delta (from Mohembo to Lake Ngami). While water discharge into and out of the Delta is measured at Mohembo and Maun respectively, the quality of the water needs to be regularly monitored at several points across the Delta. Water quality as recorded at Mohembo portrays the baseline quality and reflects upstream activities and processes affecting the water before it enters the Delta. The Drosky and Sepopa study sites in the Panhandle area were chosen to capture changes in the water quality due
to human activities (e.g., livestock grazing) before the water spreads over the Okavango Delta proper which is represented by the Guma study site. Nxaraga study site in the seasonal swamp and Boro site at the southern edge of the Delta were chosen in order to determine the effect of the preceding wetland on the water quality. Wetlands are generally known to purify flow-through water. Further sites selected downstream included Maun, Tsanekona, Toteng and Lake Ngami were intended to capture any impacts of human activities in the river system.

Figure 1: Map of the Okavango Delta showing locations of water sampling sites (from Gondwe and Masamba, 2016).

**Sampling**

Collection of water samples for this study was done collaboratively by field staff from the Okavango Research Institute (ORI), with Department of Water Affairs (DWA), Drosky’s Cabins, Sepopa Water Works and Guma Lagoon Camp. All field workers attended a training workshop on data/sample collection and handling prior to the start of the sampling campaign in order to generate quality data. Field staff from the DWA, Drosky’s Cabins, Sepopa Water Works and Guma Lagoon Camp collected samples from Mohembo, Drosky, Sepopa and Guma study sites respectively, while ORI field staff sampled Nxaraga, Boro, Maun, Tsanekona, Toteng and Lake Ngami.

Sampling was conducted almost fortnightly for two years between June 2008 and June 2010.
On site measurements of surface water temperature (°C), electrical conductivity (EC, μS cm⁻¹), dissolved oxygen (DO, mg L⁻¹), pH and turbidity (NTU) were conducted at all the ten study sites. These measurements were reported in Gondwe and Masamba (2016). For this study water samples were collected from below the water surface at each site into acid-washed polyethylene bottles. The water samples were transported within 24 hours of collection to the environmental laboratory at ORI for further sample processing and storage until analysis. Water samples for cation and anion analyses were filtered through 0.45μm GF/F filters within 24 hours of collection into clean polyethylene bottles which were immediately stored at 4°C until analysis. Filtered water samples for major cation analysis were acidified before storage to pH 2 with nitric acid for preservation.

**Analysis of water samples**

All chemical analyses of water samples were done at the environmental laboratory at the Okavango Research Institute. Water samples were analyzed using standard methods for major cations which included calcium (Ca), magnesium (Mg), sodium (Na), potassium (K) and iron (Fe); anions including carbonates (CO₃²⁻), bicarbonates (HCO₃⁻), chlorides (Cl⁻), and sulphates (SO₄²⁻); dissolved silica and plant nutrients as total nitrogen (TN) and total phosphorus (TP). Total dissolved metals were determined using a SpectraAA 220 atomic absorption spectrophotometer (Varian Australia Pty Ltd, Victoria, Australia) equipped with a Varian GTA-110 furnace, either using acetylene/air for Na, K, Fe, Mn or acetylene/nitrous oxide for Ca and Mg. Chlorides (Cl⁻) and sulphates (SO₄²⁻) were determined using a Dionex ICS-3000 DC ion chromatography (Dionex Corp., USA). Total alkalinity as carbonate and bicarbonate were determined by direct titration of water samples with standardised HCl to pH 4.5 using a Mettler Toledo DL50 Autotitrator equipped with a Rondolino autosampler (Mettler-Toledo Inc., USA). Dissolved silica was measured using the reduced molybdosilicate spectrophotometric method. TN and TP were analysed using the continuous-flow AutoAnalyzer 3 (Bran+Luebbe, Germany). Total dissolved ions (TDI, mg L⁻¹), which is a measure of water salinity, was estimated from EC by multiplying by a conversion factor of 0.64 (i.e., TDI = 0.64*EC). Since dissolved silicic acid (H₄SiO₄⁻) remains largely undissociated (i.e., unionized) in natural waters except at pH>9 and therefore does not significantly contribute to EC, the silica concentration (mg L⁻¹) was added to the TDI concentration to estimate total dissolved solids (TDS) concentration in surface waters along the study transect across the Delta.

**Table 1**: Characteristics of the meteorological stations which provided precipitation data used in this study.

<table>
<thead>
<tr>
<th>Station</th>
<th>Latitude</th>
<th>Longitude</th>
<th>Elevation (m)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Shakawe Meteorology Station</td>
<td>18° 22'</td>
<td>21° 50'</td>
<td>1030</td>
</tr>
<tr>
<td>Seronga Police Station</td>
<td>18° 49'</td>
<td>22° 24'</td>
<td>985</td>
</tr>
<tr>
<td>Gumare Agriculture Station</td>
<td>19° 22'</td>
<td>22° 10'</td>
<td>962</td>
</tr>
<tr>
<td>Maun International Airport</td>
<td>19° 59'</td>
<td>23° 25'</td>
<td>945</td>
</tr>
</tbody>
</table>

**Precipitation and river discharge data**

Precipitation data used in this paper were collected by rain gauges at Shakawe meteorological station, Seronga police station, Gumare agriculture station and Maun airport (Table 1). The areal mean precipitation over the Delta was determined using the arithmetic-mean method (equation 1) by dividing the sum of rain depths (mm) recorded at Shakawe, Seronga, Gumare and Maun by the number of stations (4 stations) (Chow et al. 2003)
Results and Discussion

Precipitation over the Delta area during the study period has been presented in Figure 2a. Precipitation over the Delta generally occurred between October and April/May while June-September was generally dry. Analysis of variance of means of annual precipitation totals for the 4 meteorological stations (Table 1) between 2006 and 2012 showed no significant differences (p=0.515). Consequently, annual totals were averaged to estimate average precipitation over the Delta (see above). The areal mean precipitation over the Delta generally shows high interannual variation, ranging from 400 to 720 mm y\(^{-1}\) for the period 2007-2010. The area of maximum inundation for the period 2007-2010 varied from 8,120-11,600 km\(^2\) (http://www.okavangodata.ub.bw/ori/monitoring/flood_maps/), and a mean of 9,644 km\(^2\). Assuming the mean area of maximum inundation of 9,644 km\(^2\) represents the Delta swamp size, we estimated the annual precipitation over the Delta to have contributed approximately 5.8 \times 10^9 m\(^3\) over the 2007/08 rainy season, 4.1 \times 10^9 m\(^3\) over 2008/09 rainy season and 5.0 \times 10^9 m\(^3\) of water during the 2009/10 rainy season. These annual precipitation rates are consistent with estimates reported in previous studies (e.g., Gumbricht et al. 2004).

River inflow and outflow at Mohembo and Maun respectively for the study period, 2008 to 2010, are presented in Figure 2b. There were high intra- and inter-annual variations for both inflow and

\[
P = \frac{P_1 + P_2 + P_3 + \ldots + P_n}{n} = \frac{1}{n} \sum_{i=1}^{n} P_i
\]

where \(P\) is average precipitation over the catchment area for a given period, and \(P_1, P_2, P_3, \ldots, P_n\) are the precipitations in a given time period at stations 1,2, 3,\ldots, \(n\) respectively within the catchment (\(n = 4\) in our study). Discharge (m\(^3\) s\(^{-1}\)) data for the study period for Mohembo and Maun were obtained from the DWA, Botswana. For both data sets (precipitation and discharge), an annual cumulative input was calculated from October to September of the following year.

Data analysis

Graphical plots presented in this paper were prepared using Sigmaplot 12.0 software (Systat Software Inc, USA). Linear relationships were tested for significance at alpha level of 0.05 in SigmaPlot 12.0 software.

Figure 2: (a) Monthly precipitation over the Okavango Delta and (b) water inflow (solid line) at Mohembo (Okavango River) and outflow (dashed line) at Maun (Thamalakane River) from 2007 to 2010.
outflow measurements. The mean inflow at Mohembo and outflow at Maun between October 2007 and September 2010 were 330 and 6 m$^3$s$^{-1}$, respectively. Although the annual discharge at Mohembo varied considerably, the mean was slightly higher than previously reported by McCarthy et al. (1998) at 292 m$^3$s$^{-1}$ and Bauer (2004) at 300 m$^3$s$^{-1}$. Due to the Delta’s low topographic gradient of about 1:3500, peak inflows observed at Mohembo in April reached the southern distal area of Maun in September (Figure 2b), which is again consistent with earlier studies (Wolski et al. 2006). For instance, McCarthy and Ellery (1998) and Gumbricht et al. (2004) have previously reported that the flood water takes about 3-5 months to traverse the 250 km distance from Mohembo to the distal areas of the Delta. During the study period, total inflow at Mohembo increased by over 60% from 7.9 x 10$^9$ m$^3$ in the 2007/08 flood season to 12.8 x 10$^9$ m$^3$ in the 2009/10 flood season. Precipitation contributed 42%, 28% and 28% to the total water inflow into the Delta in 2007/08, 2008/09 and 2009/10 flood seasons, respectively. An inflow of ~9 x 10$^9$ m$^3$ per annum has been reported by Gumbricht et al. (2004). Total outflow from the Delta through the Thamalakane River at Maun concomitantly increased from 0.04 x 10$^9$ m$^3$ in 2007/08 to 0.36 x 10$^9$ m$^3$ in 2009/10. During the whole study period annual outflow recorded at Maun on the Thamalakane River was ≤3% of the total annual inflow at Mohembo.

**Major ions**

**Anions**

Mean concentrations for HCO$_3^-$, Cl$^-$ and SO$_4^{2-}$ anions have been presented in Figure 3a-c. At the prevailing neutral to slightly acidic pH levels in the river system (Gondwe and Masamba 2016), HCO$_3^-$ was the most dominant anion in the river system accounting for 97.2±0.6% of the total anionic concentration in the water, while CO$_3^{2-}$ was consistently below detection except on a few occasions at Lake Ngami when pH was higher than 8.4 (Figure 3d). Mackay et al. (2011) have reported HCO$_3^-$ dominance of over 90% throughout the Delta. HCO$_3^-$ concentration and its anionic dominance generally increased from Mohembo to Lake Ngami (Figure 3a), while Cl$^-$ concentrations at anionic dominance of 1.9±0.3% (Figure 3c) remained stable at 0.80±0.06 mg L$^{-1}$ between Mohembo and Tsanekona but more than doubled thereafter to 1.88±0.46 mg L$^{-1}$ at Lake Ngami. Sulphate (SO$_4^{2-}$) concentrations, which had a mean anionic dominance of 0.8±0.3%, declined from around 1.08±0.34 mg L$^{-1}$ at Mohembo to 0.13±0.03 mg L$^{-1}$ between Guma and Tsanekona but slightly increased thereafter to Lake Ngami (Figure 3b). While Cl$^-$ is considered a conservative element in the Delta and therefore tended to remain stable across the sampled river system, the decline in SO$_4^{2-}$ concentration (Fig 5b) between Mohembo and Nxaraga could be partly due to the dissimilatory sulphate reduction by obligate anaerobes such as *Desulfovibrio*, which use SO$_4^{2-}$ as a terminal electron acceptor during catabolism of organic compounds such as ethanol and carbohydrates in reducing ecotones in the permanently flooded swamps. Sulphate reducing bacteria can be found in almost all anaerobic organically rich environments including marine, estuarine and freshwater sediments (Brown 1982). In aquatic systems where SO$_4^{2-}$ concentrations are high, dissimilatory SO$_4^{2-}$ reduction may account for a significant proportion of the total carbon mineralized there (Jørgensen 1977). In the current study SO$_4^{2-}$ concentration increased between Nxaraga and Lake Ngami probably because water turbulence facilitated aeration of the river water which led to the re-oxidation of reduced sulphides (S$^{2-}$) to SO$_4^{2-}$. Sulphate concentrations along the river transect were generally higher during the wet than dry season. Generally the hot but rainy season in the flood-pulsed Okavango Delta is characterised by lower flood water levels, compared to the dry season when flooding occurs, which allows aeration of the water column and subsequent oxidation of the reduced sulphides to SO$_4^{2-}$.  

31
Figure 3: Concentrations of (a) bicarbonate ($\text{HCO}_3^-$), (b) sulphate ($\text{SO}_4^{2-}$), (c) chloride ($\text{Cl}^-$) anions and (d) their respective dominance along the study transect across the Delta.

Cations
Concentrations of Na, K, Mg, Ca and Fe have been presented in Figure 4a-e. Similar to earlier studies (e.g., MacKay et al. 2012), Ca was the most abundant cation at all the study sites (i.e., Ca>Na>Mg>K>Fe) except in Lake Ngami where Na was the most dominant (i.e., Na>Ca>Mg>K>Fe) (Figure 4f). Concentrations of all five cations in the inflowing water at Mohembo were generally low at < 4 mg L$^{-1}$ as also reported by McCarthy and Ellery (1994) and MacKay et al. (2011). Although the concentrations of the cations were low, they increased from Mohembo to Lake Ngami (Figure 4) partly due to the high evapotranspirative water loss which exceeds annual precipitation over the Delta area by a factor of three (Wilson and Dincer 1976; Dincer et al. 1978). A much higher increase of approximately 3-4 fold in the cation concentrations was observed between the upper Panhandle (Mohembo and Drosky) and the distal (Boro and Maun) areas of the Delta. The evapotranspiration varies spatially in the Delta in that it is generally higher in seasonal and occasional floodplains than in permanent swamps due to higher water temperatures and shallower water levels (Mackay et al. 2011; Akoko et al. 2013; Gondwe and Masamba 2016).
Figure 4: (a) Sodium (Na), (b) Potassium (K), (c) Calcium (Ca), (d) Iron (Fe), (e) Magnesium (Mg) as concentrations and (f) their dominance along the study transect across the Delta. The legend for (f) is Ca=●, Na=○, K=▼ and Mg=△.

Total dissolved ions (TDI)
The TDI concentrations, as estimated from Electrical Conductivity (EC), were lower, but slightly increased with distance down the river from Mohembo to Guma, downstream of Guma TDI steeply increased to Lake Ngami during both dry and wet seasons (Figure 5a). Since TDI concentration represents the sum concentration of dissolved major cations and anions (i.e., $\Sigma K^+ + Na^+ + Mg^{2+} + Ca^{2+} + CO_3^{2-} + HCO_3^- + SO_4^{2-} + Cl^-$) in a water sample, we tested the relationship between the two variables in the surface waters along the study transect. TDI was positively correlated to the sum of the major ions (i.e., $\Sigma$[cations + anions]) in the surface water (Figure 5c; Table 2). Separate strong positive correlations were also observed between TDI and the sum of cations ($\Sigma[K^+ + Na^+ + Mg^{2+} + Ca^{2+}]$) (Figure 5d; Table 2) and anions ($\Sigma[CO_3^{2-} + HCO_3^- + SO_4^{2-} + Cl^-]$) (Figure 5d; Table 2) which show that both cations and anions evaporate as water flows down the study transect from Mohembo to Lake Ngami (see Atekwana et al. 2016 and references therein). As discussed above, $CO_3^{2-}$ concentrations were only sporadically detected at Toteng and Lake Ngami where much higher pH values were observed. Data in Figure 5d and Table 2 show that cations contribute much less to TDI than anions. The gradients of the
plots also indicate that both cations and anions contribute more to TDI during the rainy season than dry season. This can be attributed to evapoconcentration and higher wet- than dry-atmospheric deposition of cations and/or anions in the Delta.

Silica, which is also a dissolved solute, varied from a mean concentration of 7.37±0.18 mg L\(^{-1}\) in the upper Delta (Mohembo to Guma) to 24.06±0.49 mg L\(^{-1}\) between Boro and Lake Ngami (Figure 5b). Dissolved silica exists as unionised SiO\(_2\) or Si(OH)\(_2\) forms at pH<9 and as a result it was not included in EC measurements because of poor, or lack of, correlation with EC at all sites sampled except at Sepopa (\(r^2=0.7456\)) which is probably also explained by evapoconcentration.

**Figure 5:** TDI (mg L\(^{-1}\)) along the river transect (a) and relationship between TDI and the sum of major ions (b), the sum of cations (c) and the sum of anions (d). See Table 2 for linear equations and correlation coefficients.

Using the daily discharge rates as measured at Mohembo and Maun between 2008 and 2010, TDS (i.e., TDS=TDI (mg L\(^{-1}\)) + SiO\(_2\) (mg L\(^{-1}\))) input into the Delta via the Okavango River was estimated at 256,197-415,104 t y\(^{-1}\). Previous studies have reported TDS input at 400,000 t y\(^{-1}\) (McCarthy and Metcalfe 1990; Gumbricht et al. 2004; Zimmermann et al. 2006). Loss of solutes through surface outflow in the Thamalakane River measured at Maun varied between 3,994-45,926 t y\(^{-1}\), which is approximately 2-11% of the total solutes brought in by the Okavango River. McCarthy and Metcalfe (1990), Gumbricht et al. (2004) and Zimmermann et al. (2006) estimated the average solute loss via surface outflow at 7% of total input measured at Mohembo. In view of the large amounts of solutes deposited in the Delta each year, it is surprising that the Delta has remained a freshwater ecosystem instead of a saline water body or salt pan like the downstream Makgadikgadi Pans. The Delta’s freshwater status is attributable to a simple but important characteristic feature of the Okavango Delta (McCarthy et al 1986, 1991), which is the permanent sequestration of most solutes under the numerous
tree islands in the Delta facilitated by evapotranspiration of the woodland trees on the islands fringes. The evapotranspiration of the trees lowers the groundwater table under the islands which creates groundwater gradients driving large lateral flows of groundwater towards the centre of the islands from the surrounding floodplains. The water loss to the centre of islands is probably higher in the seasonal floodplains than in the permanent swamps primarily because of the absence of a significant peat layer in the seasonal floodplains that could potentially reduce infiltration. For instance an infiltration study in the Delta in Phelo’s seasonal floodplain near Nxaraga observed rapid recession of the groundwater table in the floodplain after the cessation of the floods which was primarily attributed to transpirative loss of the water by floodplain vegetation and woodland trees (Ramberg et al. 2006). Due to the lateral flow of the groundwater, several studies have shown high concentrations of solutes in the order of 25,000 mg L\(^{-1}\) (Gieske 1996) under these islands in the seasonal swamps. As fresh swamp water flows under the islands, the dense saline water flows deeper into the centre of the islands stabilizing the whole system. Generally older and larger forested islands are common in the seasonal and occasional swamps of the Delta while smaller and younger grassland islands dominate the permanent swamps (Gumbricht et al. 2004). As a result most of the chemical sedimentation beneath islands due to transpiration by deep rooted trees primarily occurs in the seasonal and occasional swamps of the Delta.

This study observed a continued increase of approximately 38±12% in Na, K, Mg and Ca concentrations with distance along the river from Boro to Lake Ngami which is probably due to the continued evaporative loss of surface river water and to a lesser extent anthropogenic pollution of the water, such as Cl\(^-\), SO\(_4^{2-}\) and TSS brought in with effluent water.

<table>
<thead>
<tr>
<th>Y-axis</th>
<th>Season</th>
<th>Equation</th>
<th>r(^2)</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>(\Sigma) (cations+anions)</td>
<td>Dry</td>
<td>(y = 1.8595*\text{TDI} + 10.0739)</td>
<td>0.9734</td>
<td>139</td>
</tr>
<tr>
<td></td>
<td>Wet</td>
<td>(y = 1.9704*\text{TDI} + 4.1926)</td>
<td>0.9474</td>
<td>167</td>
</tr>
<tr>
<td>(\Sigma) (cations)</td>
<td>Dry</td>
<td>(y = 0.3791*\text{TDI} - 1.0358)</td>
<td>0.9731</td>
<td>140</td>
</tr>
<tr>
<td></td>
<td>Wet</td>
<td>(y = 0.3978*\text{TDI} - 2.1568)</td>
<td>0.9463</td>
<td>169</td>
</tr>
<tr>
<td>(\Sigma) (anions)</td>
<td>Dry</td>
<td>(y = 1.4631*\text{TDI} + 12.8741)</td>
<td>0.9588</td>
<td>147</td>
</tr>
<tr>
<td></td>
<td>Wet</td>
<td>(y = 1.5770*\text{TDI} + 7.3227)</td>
<td>0.9051</td>
<td>169</td>
</tr>
</tbody>
</table>

**Plant nutrients**

**Nutrient concentrations in the water column**

Nitrogen (N) and phosphorus (P) are two of the most important macronutrients for primary production in both terrestrial and aquatic ecosystems (Conley et al. 2009). Most limnologists measure N and P as total nitrogen (TN) and total phosphorus (TP) respectively because these represent N and P values already in biomass and/or dissolved and available for assimilation into active biomass (Dodds 2003). Both TN and TP have been shown to be better predictors, than dissolved inorganic N and P, of nutrient status in a wide variety of aquatic systems (Dodds 2003). In fact TP is strongly correlated with phytoplankton biomass in a variety of aquatic systems (Dillon and Rigler 1974) such that current management strategies of inland water bodies of reducing P loading to control eutrophication are based on this relationship (Smith and Shapiro 1981). This is because concentrations of inorganic N tend to be typically lower in most aquatic systems including wetlands due to their high loss to denitrification processes and high assimilation rates (Cronberg et al. 1995).
TP and TN in the flood water at Mohembo were low at 0.042±0.005 and 0.222±0.018 mg L⁻¹, respectively (Figure 6a & c). These nutrient concentrations are consistent with concentrations reported by Cronberg et al. (1995), Garstang et al. (1998), Krah et al. (2006), Ramberg et al. (2010) and MacKay et al. (2011). Ramberg et al. (2010) reported an average of 0.023 mg L⁻¹ TP and 0.36 mg L⁻¹ TN in inflow water at Mohembo. As flood water flows downstream, TP declined slightly to 0.035±0.004 mg L⁻¹ at Tsanekona but increased thereafter by more than 3-fold to 0.131±0.024 mg L⁻¹ at Lake Ngami. The low TP concentrations across the Delta are again consistent with values reported by Krah et al. (2006) and MacKay et al. (2011). The observed decline in TP across the Delta may be partly explained by high uptake of the phosphorus by dense stands of Cyperus papyrus and Phragmites australis which flank the primary channels in the Panhandle and permanent swamp regions (Garstang et al. 1998; MacKay et al. 2011; Mosimane 2015). These nutrients are necessary to sustain the high productivity rates of the papyrus stands in this environment where nutrients are generally low throughout the year. The absorbed nutrients are then efficiently utilized and conserved through reallocation within the plant. The nutrient abstraction by these macrophytes, combined with other factors including high sedimentation of suspended material facilitated by high water retention time and immobilization of both P and N in peat as indicated by lower C:P and C:N ratios of peat material than in living vegetation (Garstang et al. 1998), results in nutrient deficiency in the rest of the Delta (McCarthy and Ellery 1998). In fact, a general reduction in biological productivity with distance from channels into the backwaters and from the Panhandle to distal areas near Maun has been reported from satellite imagery (McCarthy et al. 1993). Garstang et al. (1998) estimated a C. papyrus biomass of about 3.5 x 10⁴ kg ha⁻¹ on channel fringes which decreases significantly with distance to 3.5 x 10³ kg ha⁻¹ in the backwaters and distal areas of the Delta.

The slight decline in the TP concentration from 0.042±0.005 to 0.035±0.004 mg L⁻¹ across the Delta has previously been reported as well (see MacKay et al. 2011), and may be attributed to an observation (McCarthy et al. 1989) that over 70% of tissue P (compared to <50% N) is internally recycled by vegetation in the Delta, particularly by the C. papyrus in the permanent swamp, through translocation from senescing aerial shoots to younger tissues or to perennial belowground rhizome system. According to Johnston (1991) nutrient translocation, also known as internal recycling, helps perennial wetland vegetation to conserve already assimilated nutrients, which consequently reduces their net nutrient uptake from the soils and/or water.

Unlike TP, TN increased by approximately 3-fold from 0.222±0.018 mg L⁻¹ at Mohembo to 0.698±0.035 mg L⁻¹ at Maun, despite potential loss of N to denitrification in the anaerobic environments. The observed increase in TN could partly be attributed to an influx of fixed N from N₂-fixing autotrophic cyanobacteria and diazotrophs such as sulphate reducing bacteria common in anaerobic wetland environments (Šantrůčková et al. 2010). Although a wide occurrence of cyanobacteria mats has been reported in the Delta (Garstang et al. 1998; Marazzi 2014), diazotrophs have not been adequately studied but their nitrogenase activity is likely to be higher in the permanent swamps dominated by highly productive C. papyrus. This is because diazotrophs are generally associated with plant rhizosphere and sediments characterised by high availability of labile C sources (Šantrůčková et al. 2010). The potential of N₂-fixation to increase N concentrations in aquatic ecosystems varies from 5-70% of fixed N depending on physico-chemical conditions such as temperature, pH and water content in the system (e.g., Belnap 2001). However, the increase in N concentration can be significant as has been observed in Lake Victoria, East Africa (Hecky 1993; Lehman and Branstrator 1994) and in Lake 226 at the Experimental Lakes Area (ELA), northwestern Ontario, Canada, which was artificially made N-deficient by addition of NO₃⁻ and PO₄³⁻ at N:P weight ratio of 5 (Schindler 1977). In Lake 226, N-deficiency created a bloom of N₂-fixing cyanobacteria which was later succeeded by a high abundance of...
non $N_2$-fixing algal species (e.g., *Limnothrix redekei*) which gained some competitive advantage over $N_2$-fixing species when some of the fixed N was recycled back into the environment (Schindler *et al.* 2008).

**Figure 6:** (a) Total nitrogen (TN), (b) total phosphorus (TP), (c) TN and TP relationship and (d) TN:TP ratio along the study transect across the Delta.

**Limiting nutrient**

In terrestrial and aquatic systems the relative availability of N and P does not only control plant biomass, but also species composition of the plant communities (Koerselman and Meuleman 1996). This is probably because different plant species have differential requirements for N and P. The N:P ratio is commonly used to determine the nutrient, between N and P, that limits primary production in a given ecosystem. The N:P ratio is generally determined using TN and TP because dissolved nutrients, especially P, diminish rapidly when available due to uptake by algae and other aquatic plants (Håkanson *et al.* 2007; Schindler *et al.* 2008). Generally, ecosystems with TN:TP weight ratios lower than 7:1 are said to be N limited and their algal communities tend to be dominated by $N_2$-fixing species. In contrast, ecosystems with TN:TP mass ratios higher than 7:1 are P limited and tend to be dominated by non-$N_2$-fixing species. For instance, experimental eutrophication of Lake 227 at the Experimental Lakes Area, Canada with N and P fertilizers (N:P mass ratio of 12:1) led to a dominance by chrolophytes (Schindler *et al.* 1973; 2008). In contrast, $N_2$-fixing heterocystic *Anabaena spiroides* Klebs dominated in Lake 226 which was supplied with N-deficient fertilizer at N:P ratio of 5:1. When the N:P loading ratio of Lake 227 was lowered from 12:1 (1969-1974) to 5:1 in 1975, $N_2$-fixing heterocystic algae (*Aphanizomenon gracile*) became dominant in the water column (Schindler *et al.* 1977; Findlay *et al.* 1994). Thus the N:P ratio acts as an important determinant of species composition in natural phytoplankton com-
munities (Smith 1983). The boundary between N limitation and P limitation is however not very clear as indicated by conflicting conclusions from different studies around the world. For instance, Abell et al. (2010) reviewed data for 121 New Zealand lakes for nutrient limitation and classified majority of them as P limited (TN:TP >15:1) and few N limited systems (TN:TP <7.1). Guildford and Hecky (2000) associated TN:TP mass ratios >22.6 (>50 by mol) with P limitation and TN:TP ratios <9.1 indicated N limitation. On the other hand Smith (1983) compiled data from 17 lakes worldwide and concluded that N\textsubscript{2}-fixing cyanobacteria were abundant in systems with TN:TP mass ratios <29, but rare at TN:TP ratios >29.

The TN:TP mass ratios in the Okavango Delta varied from 6.73±0.87 at Mohembo to 23.39±3.69 at Boro (Figure 6d). Similar increasing TN:TP ratios from the Panhandle region to the distal regions along the Boro River have been reported by Marazzi (2014). Although both TP and TN further increased by about 3- and 10-fold respectively from Boro to Lake Ngami, the water remained at TN:TP ratio of around 21.28±2.19 (Maun to Lake Ngami) which may be associated with evaporative concentration in the river system (Figure 6d). There are several conclusions which could be drawn from these TN:TP mass ratios in terms of nutrient limitation and phytoplankton species composition. The N:P thresholds derived by Abell et al. (2010) suggest that algal growth in the Panhandle region of the Delta potentially experienced N limitation during both dry and wet seasons (TN:TP=5.97±0.81), and should favour proliferation of N\textsubscript{2}-fixing cyanobacteria which are generally better N competitors than other algal species in N limited environments (Smith 1983; Levine and Schindler 1999). The rest of the study transect probably experienced co-limitation by N and P during the dry season (TN:TP=12.17±0.73) and P limited during the wet season (TN:TP =23.35±3.00). Mackay et al. (2011) have previously suggested potential P limitation along the Boro River where they also reported mass TN:TP ratios of around 20.4. The algal assemblage in this part of the Delta would be co-dominated by diatoms, green algae and cyanobacteria especially during the dry season when N and P were co-limiting algal growth. Despite the varying TN:TP ratios, Marazzi (2014) observed widespread occurrence of N\textsubscript{2}-fixing cyanobacteria across the Delta which suggests a widespread N deficiency across the whole studied river system. The abundance of N\textsubscript{2}-fixing cyanobacteria were found to be higher in the permanent and seasonal swamps where TN:TP ratios suggested N and P co-limitation or P-limitation than in the Panhandle region where TN:TP ratios favoured cyanobacteria dominance (Marazzi 2014). It is possible that N\textsubscript{2}-fixation by cyanobacteria and other microbes might have shifted nutrient limitation from N- to P-limitation by raising N availability relative to P in the permanent and seasonal swamps as has been observed in other ecosystems around the world (Schindler 1977; Schindler et al. 2008). N\textsubscript{2}-fixation is likely to have been enhanced by higher water temperatures in the rest of the Delta than in the Panhandle region (see Marazzi 2014).

According to Koerselman and Meuleman (1996) and Bedford et al. (1999), the relationship between nutrient concentrations and N:P ratios can also be examined by plotting N concentrations against P concentrations (Figure 6c). Data in Figure 6c show that P-limitation (N:P ratio ≥22.6) seems to prevail during the wet season while N limitation (N:P ratio ≤9.1) prevails during the dry season when available N is probably higher than P. P limitation during the wet season could be attributed to input of fixed N from N\textsubscript{2}-fixation which is expected to be higher during the wet season when water temperatures in the Delta are much higher than during the dry season (see Gondwe and Masamba 2016). Similar results have been observed in tropical wetlands in Belize (Rejmánková et al. 2004) and Yucatan Peninsula, Mexico (Vargas and Novelo 2007) where N\textsubscript{2}-fixation rates were higher during warmer wet seasons than during the winter months. Warmer temperatures may also result in increased rates of other biological processes such as methanogenesis in the Okavango Delta during the warmer wet period as
compared to the cooler dry season (Gondwe and Masamba 2014; Masamba et al. 2015). This study’s results from the Okavango Delta contradict an earlier study by Mackay et al. (2011) which found TN in the Delta to be much higher during the flooding period which occurs during the dry season. Mackay et al. (2011) results suggest P-limitation, not N-limitation, during the dry season and probably the opposite during the wet season.

The low TN:TP ratios in the flood water at Mohembo (Figure 6d), which are slightly below the Redfield ratio of 7:1 by mass, are rather unexpected for the Okavango River system which is dominated by runoff from relatively undisturbed catchments in the Central Angolan Highlands. Runoff water from undisturbed catchments tend to have high TN:TP ratios because P concentrations in the soils generally tend to be very low (see Downing and McCauley 1992). It is not clear if the low TN:TP ratios can be associated with high P leaching and runoff from relatively small irrigated agricultural schemes operating along the Okavango River in the Zambezi province (formerly known as the Caprivi strip) Namibia. Further research is therefore needed to directly assess the impact of these irrigation schemes and other upstream development activities on the quality of the water of the Okavango River at Mohembo and as it flows into the Panhandle swamp.

It is also important to assess nutrient availability to wetland plants for conservation purposes. Besides the laborious traditional field fertilization experiments (Chapin et al. 1986), assessment of plant species and communities for nutrient availability is commonly done by analysing plant tissues for N:P ratios (Koerselman and Meuleman 1996). Several studies in European wetlands (see Koerselman and Meuleman 1996; and references therein) have confirmed that N:P ratios of plant tissues correlate with N:P supply ratios. Generally, those studies highlight the same finding that an N:P ratio of <14 in plant tissue indicates N limitation while an N:P ratio >16 indicates P limitation. These critical N:P ratios have been used to classify emergent macrophyte species in the Everglades as either N or P limited (Daoust and Childers 1999). Although the current study did not assess the nutrient status of vegetation in the Delta, N and P concentrations in vegetation reported by two previous studies (Bonyongo 1999; Mosimane 2015) suggest that vegetation in the Delta could be limited by N. Nitrogen and P concentrations in plant tissues from the seasonal floodplain reported by Bonyongo (1999) indicated a mean N:P ratio of 12.29±0.80 while Mosimane (2015) reported a mean N:P ratio of 11.87±0.59 (by mass). Since the research objectives of the above two studies were not primarily focused on nutrients levels in plant tissues, only a few plant samples were collected in an ad hoc manner. We therefore suggest that future studies should rigorously reassess the nutrient status of the vegetation in the Delta for purposes of informing management strategies with regard to nutrient pollution in the Delta.

Conclusion
Wetlands act as natural water purifiers and have been deliberately constructed and used to improve the quality of various effluents in many parts of the world. In the Okavango Delta the role of evapotranspiration in increasing concentrations, particularly of cations, anions and silica, was evident as also shown by earlier studies. High evapotranspiration in the Delta is consistent with the disparity between huge inflow and little outflow in the Delta. Analysis of nutrient data showed spatial and temporal N and P limitation in the Delta. N limitation is probably ameliorated by biological nitrogen fixation indicated by widespread abundance of nitrogen fixing cyanobacterial species in the Delta.

Acknowledgements
The authors would like to thank the staff from the Department of Water Affairs, Drosky’s Cabins, North West District Council and Guma Lagoon Camp for collecting water samples as well as taking field
measurements. We would also like to thank the field and laboratory staff of the Okavango Research Institute for collecting samples from the lower Delta and analysing them, respectively. This activity was funded by the BIOKAVANGO Project.

**References**


Dincer, T, Hutton, LG. and Kupee, BBJ 1979. ‘Study using stable isotopes, of flow distribution, surface/groundwater relations and evapotranspiration in the Okarango Swamp, Botswana’, Iso


Marazzi, L 2014. ‘Biodiversity and biomass of algae in the Okavango Delta (Botswana), a subtropical flood-pulsed wetland’, PhD Thesis, Department of Geography, University College London, pp. 426


