## Original Article

# Structure and seasonal dynamics of phytoplankton and zooplankton in Lake Azili, small Lake of the pond of River Ouémé, Benin 

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#### Abstract

The early reaction of the plankton communities to environmental changes makes it a useful tool for monitoring the pollution of aquatic environments. Lake Azili is a small water body, the majority of which is strongly influenced by river Ouémé during the floods time. Its ecosystem is one of the most important for the country, due to its rich biodiversity, especially that of halieutics. By following the evolution of its health, due to the strong anthropological pressure, this study aims to estimate the structure and the dynamics of its planktonic biodiversity and to assess its current state. It was carried out for seven months between May to November, 2012, according to the hydrological seasons of Benin. Plankton samplings were monthly, taken in a vertical sense from the lake, from all depths, using plankton net. The diversity indices were calculated for the compartment of zooplankton, that of phytoplankton being a preliminary evaluation. A total of 51 species of phytoplankton and 36 zooplankton species were inventoried. Instability is observed in the seasonal structure of both communities, especially for the period of transition between the floods and the floods recession. Due to the specific composition and the diversity, the ecosystem of Lake Azili is perturbed.


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## Introduction

The aquatic biodiversity has been used for several decades as an ecological health and environment pollution indicator. In this regard, several studies estimated the distribution of the aquatic flora and fauna to understand the environmental changes. Nowadays, in all of Africa's waters, the monitoring of quality is much more based on the assessment of macroscopic organism (mainly halieutic fauna), although the microscopic organisms are more sensitive to the environmental variations. In continental waters, sensibility of microorganisms is more notable due to strong human influence (Groga, 2012). The distribution of these organisms is dictated by the auto-ecological processes that result from the global changes and the anthropological factors (Dolédec et al., 1999). Several algorithms and other
statistical designs currently exist and are developed based on the plankton structures at specific abundance levels to assess the quality of the aquatics environments. But these tools are developed in temperate conditions, and it is necessary to adapt them on the tropical environments. This adaptation requires a good knowledge of the structure of microorganisms (phytoplankton and zooplankton) as well as their ecology. Thus, several African countries began the exploration of the planktonic populations in their waters. But in Benin, little study regarding plankton was carried out, while the country has several water bodies, among which Lake Azili forms a particular ecosystem. Some populations depend on this lake, in particular the village "Agonvè", surrounded with water and forming a small island within the country. Over the past few years, a

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Figure 1. Localization of the sampling sites in Lake Azili.
continuous process of degradation of the ecosystem has been recorded (Aklinon, 2005). Therefore, it is important to involve different disciplines to evaluate its current state and to identify ways of rehabilitation. Hence, this study aims to estimate the phytoplankton and zooplankton communities of the Azili Lake and to establish a base for future works of quality assessment.

## Materials and Methods

Study sites and sampling stations: This study was carried on Lake Azili, which is a small flood lake of Ouémé River in the municipality of Zangnanado, situated in the center of Benin. The Lake Azili extends in period of flood recession from $7^{\circ} 15^{\prime}$ to $7^{\circ} 20^{\prime} \mathrm{N}$ and from $2^{\circ} 20^{\prime}$ to $2^{\circ} 30^{\prime} \mathrm{E}$. Its area is 200 ha of water and reach 300 ha of permanent deep swamp. It measures an average 1.7 km from North to South and 2 km from East to West. Its volume is regularly influenced by the Olougbé River and the channel Houan. The latter is a tributary and outflow for Azili Lake, at the same time. Along 5 km , the channel Houan links the lake with the Ouémé River which, during the overflow between July and October pours
a part of its waters in the Lake through the channel. The Oulugbé River is 15 km long and $2-3 \mathrm{~m}$ wide, and it is a permanent tributary of the lake. The high water of Lake Azili is essentially caused by those of Ouémé River.
Three sampling stations (Fig. 1), with a different level of human impact and representing the various housing environments, were chosen so as to represent the lake area. Sampling have been made from upstream to downstream, as follows: Dohoundji $\quad\left(07^{\circ} 15^{\prime} \mathrm{N}, \quad 002^{\circ} 27^{\prime} \mathrm{E}\right)$, Whéliyamey $\left(07^{\circ} 15^{\prime} \mathrm{N}, \quad 02^{\circ} 27^{\prime} \mathrm{E}\right)$ and Gbekpletin $\left(07^{\circ} 15^{\prime} \mathrm{N}\right.$, $002^{\circ} 27^{\prime} \mathrm{E}$ ).
Sampling methods: The sampling was carried out monthly, according to the hydrological seasons of Benin. It covered the period from May to November, 2012. The period of recession was represented by three months (May, June and July) and the flood season by the period from August to November. To ensure the good representation of the various species, the composite sampling protocol was retained. A composite sampling unit was a mixture of three vertical lines (Lake full vertical sampling) at the same station. The sampling effort was thus nine hauls a month. The samples are then immediately fixated in 5\% formalin solution. Both organisms groups were sampled by the same conical net with a mesh size of $30 \mu \mathrm{~m}$. Thus, the study allowed to estimate exactly the zooplankton diversity, while regarding the phytoplankton, the diversity is not precisely estimated due to the fact that several species are smaller than $30 \mu \mathrm{~m}$.
The hydrological and physico-chemical conditions during the sampling are presented in Table 1. A Hanna multi-parameter and a Voltcraf oximeter have been used to survey the physico-chemical parameters of water, while the hydrology was evaluated using a Secchi disc (diameter 30 cm ) and a graded and weighted rope.
Laboratory analysis. In the laboratory, all the samples were concentrated in the same volume of 100 ml before observation on photonic microscopy. The fifth aliquots of every sample were analyzed by taking, after homogenization, 1 ml of water sample

Table 1. Physico-chemical parameters recorded during the study.

|  | Low water time |  |  |  | Flood time |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | May | June | July | August | September | October | November |
| Température $\left({ }^{\circ} \mathrm{C}\right)$ | $30.7 \pm 0.1$ | $28.4 \pm 0.1$ | $26.2 \pm 0.1$ | $27,5 \pm 0.2$ | $27.7 \pm 0.1$ | $28.2 \pm 0.3$ | $29.37 \pm 0.3$ |
| Dissolved oxygen $\left(\mathrm{mg} . \mathrm{L}^{-1}\right)$ | $6.1 \pm 0.1$ | $5.6 \pm 0.2$ | $3.8 \pm 0.1$ | $3.4 \pm 0.1$ | $3.6 \pm 0.1$ | $4.0 \pm 0.3$ | $4.2 \pm 0.3$ |
| pH | $6.8 \pm 0.1$ | $6.8 \pm 0.03$ | $6.9 \pm 0.2$ | $6.6 \pm 0.1$ | $6.3 \pm 0.2$ | $6.8 \pm 0.02$ | $6.6 \pm 0.04$ |
| TDS (ppm) | $19.0 \pm 0.1$ | $17.5 \pm 0.0$ | $26.2 \pm 2.0$ | $25.5 \pm 1.3$ | $28.7 \pm 0.3$ | $28.5 \pm 2.2$ | $26.3 \pm 0.3$ |
| Conductivity $\left(\mu \mathrm{S.cm} \mathrm{~cm}^{-1}\right)$ | $43.7 \pm 4.2$ | $34.5 \pm 0.9$ | $51.3 \pm 3.5$ | $51.8 \pm 2.3$ | $55.0 \pm 0.9$ | $56.3 \pm 4.2$ | $51.5 \pm 0.5$ |
| Secchi depth $(\mathrm{m})$ | $0.7 \pm 0.01$ | $0.5 \pm 0.05$ | $0.6 \pm 0.04$ | $0.37 \pm 0.0$ | $0.40 \pm 0.02$ | $0.34 \pm 0.02$ | $0.19 \pm 0.0$ |

( 1 ml pipette fixed). The species identification, based on the morphological characters, was carried out for phytoplankton compartment based on NogueiraCorreia and Ferreira (2000), Tsukii (2005), Kinross (2007) and Oyadomari (2011). The identification of zooplankton species implied according to Beauchamp (1965), Smith and Fernando, (1978), Pourriot and Francez (1986) and Lynne (2004). After identification, the individuals of every species were counted per compartment, on counting cell. For the algal group, three categories of species were considered in the enumeration, i) the species found in abundance in all the fields, which were counted in 20 compartments, ii) the frequent species, counted in 40 fields and iii) the rare species which were counted on all the cell. As zooplanktons, the individuals were counted in all the compartments. During the counting for both communities, only the individuals having a whole structure are considered.
Data analysis: After enumeration of the individuals of species identified in 20 ml ( $1 / 5$ of sample), the abundances related to the filtered water volume were obtained with the following formulas: For phytoplankton species, the abundance by milliliter of sample was first calculated by: $\mathrm{N}=(\mathrm{n} * 196 / \mathrm{X})$, where N is the total number by ml observed; n the number of individuals counted by ml, 196 total number of compartment on the counting cell and X number of observed compartment. Then, the density of every species in the Lake, as well for the phytoplankton as for zooplankton was calculated by:

$$
\mathrm{D}=\left[100^{*} \sum_{\mathrm{i}=1}^{20} \mathrm{Ni}\right] /[20 * 3 * \mathrm{He} * \mathrm{Sbf}]
$$

Where D is the density (ind. $\mathrm{m}^{-3}$ ), Ni the total number
of individuals by ml for the species i , He the depth of the lake, Sbf the basic area of the plankton net, 3 number of hauls by sample and 20 number of aliquot of 1 ml observed.
After calculation of the abundances, the data were transformed before any treatment with $\log (\mathrm{X}+1)$ to stabilize the variances of biological over-dispersal (Frontier, 1973). These data were projected in the factorial design of the principal components analysis (PCA) to study the seasonal dispersal. The univariate diversity was also studied only for zooplankton by calculating the indices below. It is not calculated for phytoplankton considering that all the species of the lake are not certainly represented. The diversity of Shannon-Wiener, to discriminate the various populations. It is obtained by:

$$
\mathrm{H}^{\prime}=-\sum\left[\left(\frac{\mathrm{ni}}{\mathrm{~N}}\right) * \log 2\left(\frac{\mathrm{ni}}{\mathrm{~N}}\right)\right]
$$

Where $\mathrm{H}^{\prime}$ is the index of diversity expressed in bit/individual, ni the number of the species, N the total number of individual constituting all the species, $\log 2$ the logarithm on base 2 . The diversity index of Simpson. It is obtained by:

$$
\mathrm{SP}=1-\sum\left[\frac{\mathrm{ni} \mathrm{i}^{*}(\mathrm{ni}-1)}{\mathrm{N}^{*}(\mathrm{~N}-1)}\right]
$$

Where SP is the diversity index, ni the specific effective and N the total number of individual in the sample merged all species. The diversity index of Margalef, expressed by:

$$
\mathrm{Marg}=\frac{\mathrm{Ns}-1}{\ln (\mathrm{n})}
$$

Where Ns is the total species number and n the total abundance. The Evenness index to estimate the regularity between the species. It is obtained by:

$$
\text { Evenness }=\frac{\mathrm{H}^{\top}}{\text { Log2Ns }}
$$

Where $\mathrm{H}^{\prime}$ is the Shannon index, $\log 2$ the logarithm on base 2 and Ns the specific richness.
The ANOVA one ways test is finally used to study the temporal variations in the zooplankton and phytoplankton abundance.

## Results

Populating composition and occurrences: During the study, a total of 51 phytoplankton species, share out in five classes (Table 2) are inventoried including, Diatomophyceae represented by 24 taxa, Cyanophyceae by eight taxa, Chlorophyceae by 12 taxa, Euglenophyceae by six taxa and Pyrrophyceae by one species. As indicated in the methodology, this diversity represents probably not all the phytoplankton in lake due to the used net. Nevertheless, it represents a preliminary result of this diversity study. As the presence/absence, of the diatoms species during the shallow depths season, all taxa were identified except Aulacodiscus sp. The latter was identified only during the floods, contrary to Navicula sp., which was absent in the samples only during recession period. About the Cyanobacteria taxa, all were identified as well in flood as during the recession. In the Chlorophytes group, it was noted the absence of Cladophora sp. during the flood recession while it appeared in the floods time. An opposite distribution pattern was observed for Cosmarium sp. As the Euglenophytes, all the species were represented in both seasons with the exception of Trachelomas sp. which had the same distribution as Cosmarium sp. The only one Pyrrophyceae species was identified during both seasons. According to the occurrence percentages of various taxa regarding all the algal population, it is obtained a net dominance of the diatoms Melosira genera in the lake.
As regard to zooplankton, 36 taxa made up 30 species of rotifers, three copepods and three cladocerans (Table 2) were recorded. The presence/absence showed that three rotifers taxa, including Brachionus patulus, Lepadella sp. and

Proales daphnicola were appeared in the lake during the floods time. In addition, two other rotifers i.e. Lecane leontina and Lecane sp. were disappeared with the floods. As copepods, just Acartia sp. was found only during the high depths. The cladoceran Bosmina sp., contrary to Moina dubia, had the same distribution similar to the previous copepod species. The occurrence percentage of each rotifers species regarding whole group and both seasons showed the dominance of species as Anuraeopsis navicula, Anuraeopsis sp., Asplanchna brighwelmlii, Brachionus falcatus, Cephalodella giba, Keratella tropica, Polyarthra vulgaris and Testudinella sp.. As the copepods, Mesocyclops sp. was dominant, while Daphnia sp. dominated the cladoceran population.
Plankton population's abundance: The total phytoplankton harvested with the net, during both hydrologic seasons, knew globally a distribution in which the floods time was unfavorable (Fig. 2A). The best densities were obtained between May and July with the peak in June, which was followed by a progressive decrease until October showing a slight rise in November. This temporal variation was significantly different between June and October ( $P<0.05$ ). Regarding the zooplankton (Fig. 2B), the distribution of the abundance was similar with that of phytoplankton, except in September. The two lowest densities of total zooplankton observed in October and November was significantly different and from that of all other months ( $P<0.05$ ). In addition, the total zooplankton in May (relatively low) was significantly different from that of the other months with the exception of September.
Seasonal distribution of phytoplankton: The abundances data of various phytoplankton taxa, according to the months, both hydrological seasons, reduced and centered by a factorial analysis of the main components are represented on Figure 3. With eigenvalue of 5.61, the first factor explains $80.15 \%$ of the results. It is essentially formed by the lowwater period and the month of August, representing the floods inception. All the variables are negatively selected and well-represented on it. This period (May-August) is correlated to the strong representat-

Table 2. Phytoplankton and zooplankton species occurrence according to the hydrological seasons.

| Phytoplankton | Code | Water level |  | \% | Zooplankton | Code | Water level |  | \% |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | Low | Flood |  |  |  | Low | Flood |  |
| DIATOMOPHYCEAE |  |  |  |  | ROTIFERS |  |  |  |  |
| Aulacodiscus sp. | D1 | - | + | 0.03 | Anuraeopsis navicula | R1 | + | + | 4.21 |
| Caloneis sp. | D2 | + | + | 0.36 | Anuraeopsis sp. | R2 | + | + | 11.51 |
| Cocconeis sp. | D3 | + | + | 2.27 | Asplanchna brightwelmlii | R3 | + | + | 14.25 |
| diatoma trenuis | D4 | + | + | 0.13 | Asplanchna girodi | R4 | + | + | 2.93 |
| Eunotia bilunaris | D5 | + | + | 2.03 | Asplanchna sp. | R5 | + | + | 2.57 |
| Eunotia sp. | D6 | + | + | 0.05 | Brachionus caudatus | R6 | + | + | 0.71 |
| Gomphonema amoenum | D7 | + | + | 0.42 | Brachionus falcatus | R7 | + | + | 4.73 |
| Gomphonema sp. | D8 | + | + | 1.63 | Brachionus patulus | R8 | - | + | 0.15 |
| Gomphonema vibrio | D9 | + | + | 1.13 | Brahionus quadridentatus | R9 | + | + | 0.49 |
| Melosira ambigua | D10 | + | + | 29.86 | Brachionus sp. | R10 | + | + | 2.31 |
| Melosira granulata | D11 | + | + | 2.90 | Cephalodella giba | R11 | + | + | 20.27 |
| Melosira sp. | D12 | + | + | 27.39 | Colurella uncinata | R12 | + | + | 0.33 |
| Melosira varians | D13 | + | + | 5.16 | Filinia longiseta | R13 | + | + | 1.25 |
| Navicula sp. | D14 | + | - | 0.002 | Filina opoliensis | R14 | + | + | 1.92 |
| Nitzchia paradoxa | D15 | + | + | 0.14 | Habrotrocha sp. | R15 | + | + | 1.84 |
| Nitzschia reversa | D16 | + | + | 0.27 | Hexarthra intermédia | R16 | + | + | 0.50 |
| Nitzschia sigma | D17 | + | + | 5.26 | Keratella tropica | R17 | + | + | 4.75 |
| Pinnularia cardinalis | D18 | + | + | 0.08 | Lecane leontina | R18 | + | - | 0.21 |
| Suriella capronii | D19 | + | + | 0.02 | Lecane sp. | R19 | + | - | 0.01 |
| Suriella linearis | D20 | + | + | 1.57 | Lepadella patella | R20 | + | + | 0.78 |
| Synedra acus | D21 | + | + | 2.68 | Lepadella sp. | R21 | - | + | 0.01 |
| Synedra splendens | D22 | + | + | 1.40 | Lindia sp. | R22 | + | + | 2.17 |
| Thalassiosira rotula | D23 | + | + | 0.08 | Ploesoma sp. | R23 | + | + | 2.59 |
| Thalassiosira sp. | D24 | + | + | 0.74 | Proales daphnicola | R24 | - | + | 0.46 |
| CYANOPHYCEAE |  |  |  |  | Proales decipiens | R25 | + | + | 1.60 |
| Microcystis flos-aquae | CY1 | + | + | 0.62 | Proales sp. | R26 | + | + | 1.86 |
| Microcystis sp. | CY2 | + | + | 0.29 | Polyarthra vulgaris | R27 | + | + | 5.58 |
| Microcystis wesenbergii | CY3 | + | + | 0.27 | Pompholyx sulcata | R28 | + | + | 1.42 |
| Oscillatoria sp. | CY4 | + | + | 0.02 | Rotaria sp. | R29 | + | + | 1.13 |
| Raphidiopsis méditerranea | CY5 | + | + | 0.80 | Testudinella sp. | R30 | + | + | 7.46 |
| Spirulina sp. | CY6 | + | + | 0.02 | COPEPODS |  |  |  |  |
| Stigonema sp. | CY7 | + | + | 0.75 | Acartia sp | CO 1 | - | + | 1.12 |
| Synechocystis aquatilis | CY8 | + | + | 0.89 | Afrocyclops sp. | CO 2 | + | + | 4.95 |
| CHLOROPHYCEAE |  |  |  |  | Mesocyclops sp. | CO3 | + | + | 9.95 |
| Binuclearia eriensis | CH1 | + | + | 5.11 | Nauplui | N | + | + | 83.97 |
| Cladophora sp. | CH2 | - | + | 0.003 | CLADOCERA |  |  |  |  |
| Closterium aciculare | CH3 | + | + | 0.61 | Bosmina sp. | CL1 | - | + | 12.85 |

[^1]Table 2. Continued.

| Phytoplankton | Code | Water level |  | \% | Zooplankton | Code | Water level |  | \% |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | Low | Flood |  |  |  | Low | Flood |  |
| Closterium parvulum | CH4 | + | + | 0.62 | Daphnia sp. | CL2 | + | + | 87.00 |
| Cosmarium sp. | CH5 | + | - | 0.01 | Moina dubia | CL3 | + | - | 0.15 |
| Gonatozygon sp . | CH6 | + | + | $0.06$ |  |  |  |  |  |
| Micrasterias sp. | CH7 | + | + | 0.06 |  |  |  |  |  |
| Scenedesmus quadricauda | CH8 | + | + | 0.13 |  |  |  |  |  |
| Scenedesmus sp. | CH9 | + | + | 0.03 |  |  |  |  |  |
| Staurastrum sp. | CH10 | + | + | 0.02 |  |  |  |  |  |
| Stigeoclonium aestivale | CH11 | + | + | 0.37 |  |  |  |  |  |
| Tetraedron sp. | CH12 | + | + | 0.07 |  |  |  |  |  |
| EUGLENOPHYCEAE |  |  |  |  |  |  |  |  |  |
| Euglena sp. | E1 | + | + | 0.48 |  |  |  |  |  |
| Phacus caudatus | E2 | + | + | 0.15 |  |  |  |  |  |
| Phacus longicauda | E3 | + | + | 0.68 |  |  |  |  |  |
| Strombomona sp. | E4 | + | + | 0.33 |  |  |  |  |  |
| Trachelomas bituricensis | E5 | + | + | 0.03 |  |  |  |  |  |
| Trachelomas sp. | E6 | + | - | 0.01 |  |  |  |  |  |
| PIRROPHYCEAE |  |  |  |  |  |  |  |  |  |
| Peridinium bipes | P1 | + | + | 0.07 |  |  |  |  |  |

+ Presence, - Absence


Figure 2. Temporal evolution of total phytoplankton (A) and total zooplankton (B) of Lake Azili (The mustache boxes with no common letters are significantly different $(P<0.05)$ ).
-ion of diatoms (Melosira granulata, Melosira varians, Melosira ambigua, Melosira sp., Nitzschia sigma, Gomphonema sp., Synedra acus,Cocconeis sp. and Suriella linearis) and of Chlorophytes (Binuclearia eriensis), in opposition to diatoms (Nitzschia paradoxa, Navicula sp., Pinnularia
cardinalis, Suriella capronii and Aulacodiscus sp.), cyanobacteria (Oscillatoria sp.), chlorophytes (Cladophora sp., Cosmarium sp., Scenedesmus sp., Staurastrum sp.), and Euglenophytes (Trachelomas sp.).
The high-water time (October and November) and


Figure 3. Projection of phytoplankton species and the sampling time in the factorial design of Principal Component Analysis.


Figure 4. Projection of zooplankton species and the sampling time in the factorial design of Principal Component Analysis.

August compose the second axis, which explains $11.02 \%$ of the distribution. It opposes the floods period, which is positively selected with regard to the average depths. The species such as the diatoms Aulacodiscus sp., the cyanobacteria Microcystis flos-aquae, the chlorophyte Scenedesmus sp. and the euglenophyte $T$. bituricensis are well-represented during the floods, contrary to the species as Eunotia sp., Gomphonema amoenum, Thalassiosira rotula (diatoms), Microcystis wesenbergii (cyanobacteria),

Closterium aciculare, Stigeoclonium aestivale (Chlorophytes) and Peridinium bipes (Pirrophyte). The phytoplankton distribution in the factorial design of PCA, following the quality representation of various species, show that those of diatoms are represented well in both hydrological seasons, the flood recession period; however being more favorable. On the other hand, the period during which the lake receives the first influxes of River Ouémé (August-September) knew the best

Table 3. Monthly variation of the diversity index and the dominance degree of zooplankton species.

|  | RS | SP | H' | Evenness | Marg | Dominants species | $\boldsymbol{\%}$ D |
| :--- | :---: | :---: | :---: | :---: | :---: | :--- | :---: |
| May | 25 | 0.95 | 3.09 | 0.88 | 3.95 | Brachionus falcatus, Cephalodella giba | 12.8 |
| June | 28 | 0.96 | 3.28 | 0.95 | 4.28 | Asplanchna girodi, Cephalodella giba | 10.3 |
| July | 27 | 0.96 | 3.28 | 0.98 | 4.12 | Polyarthra vulgaris, Testudinella sp. | 8.9 |
| August | 29 | 0.96 | 3.29 | 0.92 | 4.46 | Anuraeopsis navicula, Asplanchna <br> brightwelmlii | 10.3 |
| September | 27 | 0.95 | 3.20 | 0.91 | 4.30 | Filina opoliensis, Asplanchna <br> brightwelmlii | 12.4 |
| October | 19 | 0.93 | 2.74 | 0.82 | 3.46 | Habrotrocha sp., Testudinella sp. | 21.2 |
| November | 18 | 0.93 | 2.74 | 0.82 | 3.02 | Keratella tropica, Testudinella sp. | 18.9 |

RS: specific Richness, SP: Simpson diversity, H': Shannon diversity, Marg: Margalef diversity
representations of other groups (Cyanobactera, Chlorophytes, Euglenophytes and the only Pirrophyte species). The high-water months there were not so selective.

## Seasonal distribution of the zooplankton:

 Zooplankton distribution submitted to analysis in principal components is presented on Figure 4. First axis which explains $53.10 \%$ of the results is characterized by the shallow depths season, all the variables being negatively correlated; it is composed by June, July and August. September is also well represented on it. Therefore, it explains the microfauna distribution during the recession and a part of that of the transition period between both seasons. It includes the species such as rotifers Testudinella sp., Anuraeopsis sp., Keratella tropica, Polyarthra vulgaris and nauplii of copepods, which oppose their strong representation of shallow depths to that of: i) Brachionus patulus, Brahionus quadridentatus, Lecane leontina, Lecane sp. and Lepadella sp. (rotifers), ii) Acartia sp. (copepod) and iii) Bosmina sp. and Moina dubia (cladoceran).The second axis, which expresses $21.71 \%$ of the results, is formed by the floods period (September, October and November). It selects positively the last two months and negatively September, the latter being very well-represented in the factorial design. It includes essentially the rotifer species, among which Filinia longiseta, Filina opoliensis, Pompholyx sulcata, Brachionus falcatus, Asplanchna brightwelmlii, Proales decipiens and Lindia sp. are strongly correlated to September. They oppose their spread to Testudinella sp., Ploesoma sp., Brahionus
quadridentatus and Habrotrocha sp., which are selected by the floods period. According to the quality representation of various zooplankton species, selected and not selected by one or the other one, both axes of the PCA design, it was observed that rotifers were distributed well on both seasons. Nevertheless, August (at the floods beginning) was favorable to the abundance of several species. The same period was the one with the best representations of two other zooplankton groups (copepods and cladoceran), while the high depths were unfavorable.
Zooplankton specific diversity and dominance: The diversity indices of zooplankton populations are presented in Table 3. Several indices were calculated to assess the diversity. In none of the studied months, the specific richness was equal to the total value observed on the lake. Thus, the ecological preference of the different species varied over the sampling period. Regarding the diversity indices, no high variations have been observed between both hydrological seasons, nor between constituent months. In that, the specific richness varied between 18 and 29 species, while the Simpson diversity is near of unity for all the study period. The Shannon Wiener index is varied between 2.74 bit.ind $^{-1}$ and 3.29 bit.ind $^{-1}$, the highest value coinciding with the biggest specific richness of August, the biggest Simpson diversity and also that of Margalef (29, 0.96 and 4.46, respectively). The samples uniformity measured by Evenness, remained very high during both seasons. All the index consideration, shows a good diversity of zooplanktonic population in the
lake. The best diversifications were recorded during the transition period (August-September) while lower diversities were recorded during the high depths (October-November). This diversity has resulted in low levels of dominance by the next most represented species in the different months of the study (Table 3). Thus, the percentage of dominance varied between $8.9 \%$ and $21.2 \%$. The lowest dominance does not correspond exactly to the most diversified, but altogether the tendency remains the same.

## Discussion

Although the study of the phytoplankton compartment is a preliminary evaluation of the diversity of the microflora of Lake Azili, a significant number of species was identified. Totaling 51 species, the found richness is widely below that of other watercourses of the sub-region (111 and 192 species, respectively on the Lake Guiers in Senegal (Ngansoumana, 2006) and in the coastal river of Ivory Coast (Niamien-Ebrottié et al., 2013)). The microflora sampled in the present study showed a structure dominated by diatoms, similar to Onozeyi (2013) on Ogun River in southwest of Nigeria. Azili Lake is a particular ecosystem with its important vegetation cover and the strong anthropological pressure; it has a muddy bottom due to the organic decomposition. This, therefore, justifies the important presence of diatoms species, which, in continental environment, knows good diversifications due to the organic mineralization and re-suspension caused by human activities (Groga, 2012). In the diatoms group the species selective of polluted environment were identified. The strong representation of the genera Nitzschia, Gomphonema, Navicula, Pinnularia, Caloneis, Eunotia, etc., shows that the ecosystem is perturbed and rich in organic elements. The cyanobacteria presence, generally characteristic of high nitrogenous and phosphated mineral concentration in thermal conditions from 25 to $30^{\circ} \mathrm{C}$ (Erkaya et al., 2011), is not characteristic of the lake, even if the thermal conditions are filled all year (tropical
environment). Some identified species were more representative during the transition period between recession and floods. This dynamics gives two hypotheses. Given that in this period, the lake receives the influxes of its tributary (the river Ouémé), whether it had a direct contribution in specimens by the river or the mineral conditions were improved by its jet. The first hypothesis would give more a gradient upstream-downstream in the species distribution, not being the case in the results of this study. The second hypothesis thus remains the most convincing even if a more spatial study remains to be carried out. It would thus be a mineralized material re-suspension and a nutriment contribution while the physico-chemical conditions of the river are poor (Hounsou, 2011). Nevertheless, the presence and temporal distribution selective species for high eutrophication conditions (Erkaya et al., 2011), shows the ecosystem disturbance even if a planktonic bloom is not observed at present. In addition, the chlorophytes have presented an ecological preference for the big floods period selecting particularly the transition period. Thus, the arrival of new species and the disappearance of others previously inventoried species can be observed, which was the case in four phytoplankton phyla, the most representative of inventoried population. An instability in the algal community structure in the lake due to exchanges with its tributary is subsequently noted.
Regarding the zooplankton populations, the biodiversity of the lake is globally cosmopolitan. Filinia longiseta, Filina opoliensis, Testudinella sp., Brachionus caudatus, Keratella tropica, etc. are generally well-represented in diverse environments (Bozkurt and Akin, 2012). Its wealth is important, being slightly superior than that of Cross River of Nigeria (Offem et al., 2009). It presents a characteristic structure of tropical fresh water by the wide dominance of rotifers. The seasonal structure of this community shows a tendency towards the transition period for several species. The diversity is also unstable due to the exchanges with the Ouémé River, a classic structure being, nevertheless,
observed with the peak abundance shortly after that of phytoplankton. In the population of inventoried rotifers, many species are characteristic of polluted environment. Selecting the strongly eutrophic ecosystems (Bozkurt and Akin, 2012), the distribution and temporal dominance of the species from the genera Filina and Lepadella, as of some phytoplankton species, show that the lake is in a degradation process. The health of an aquatic ecosystem can be estimated by several methods one of them being the calculation of the diversity index. It is recognized that species diversity increases with succession, while the characteristics of different species vary the spatial and temporal distribution according to the classification level of the food chain. A diversity index to explain the community characteristics will thus have to consider the functional structure of this one both at a spatial and temporal level. The work which we realized has a temporal consideration reduced and is valid only for the considered period. Therefore, for all the calculated indices, the population presents an average diversity. The Shannon index, dependent on the size of samples and on the type of habitat (Grall and Coïc, 2005), interpreted with Evenness, classifies the lake in the category of the moderately polluted ecosystems during the floods recession and the transition period and strongly polluted during the floods, on the scale of the sandy/muddy ecosystems according to Simboura and Zenetos (2002), showing again the impact that the Ouémé River has on the Azili Lake. The plankton biodiversity of the Lake is a classic population of tropical environment with an unstable structure due to the rejection of the tributary Ouémé River. Considering the ecosystem established by the lake, pressures which are exercised and its planktonic biodiversity, a tendency to eutrophication is observed.
The distribution of species along the two hydrological seasons, submitted with the classification of the lake on the Simboura and Zenetos (2002) pollution scale, highlighted in both group of phytoplankton and zooplankton, some species potentially bio-indicators of pollution.

Therefore, Aulacodiscus sp., Microcystis flos-aquae, Scenedesmus sp. and Trachelomas bituricensis are phytoplankton species candidates for bio-indication in the lake Azili subject to a further study in this direction. Regarding the zooplankton group, those are species such as: Filinia longiseta, Filina opoliensis, Pompholyx sulcata, Brachionus falcatus, Asplanchna brightwelmlii, Proales decipiens, Lindia sp., Testudinella sp., Ploesoma sp., Brahionus quadridentatus and Habrotrocha sp. A complementary study assessing the actual state of the lake pollution and the direct effects of pollutants on these species so will confirm or refute species as bioindicator of pollution in the lake Azili environment.

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