FRESHWATER DIATOMS DIVERSITY OF NATIONAL PARKS IN NIGERIA I: OKOMU NATIONAL PARK, SOUTH-SOUTH, NIGERIA.

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ABSTRACT

This study represents the first taxonomic and ecological description of the diatom flora of water and sediments in two freshwater lakes within Okomu National Park Edo State, Nigeria, using light and scanning electron microscopy (SEM). Twenty-nine species of diatoms belonging to fifteen genera, *Achnanthes, Achnanthidium, Chamaepinnularia, Cymbella, Discotella, Eolimna, Eunotia, Frustulia, Gomphonema, Luticola, Nitzschia, Pinnularia, Placoneis Sellaphora* and *Stauroneis* were identified. The taxa were dominated by *Eunotia* spp. Water pH of 5.9-6.3 and surface temperature of 28 °C probably implies that the diatoms especially *Eunotia* spp are exploring an ecological niche that is probably favourable for its growth. Okomu National Park appears to be a hotspot for *Eunotia* species diversity, and further study of the oligotrophic waters of this site is recommended.

Keywords: Diatom, Diversity. Freshwater, National Park, Okomu, Season, Tropical.

INTRODUCTION

Diatoms are single-celled algae and the most common phytoplankton with their cell walls made of silica (Wehr and Sheath, 2003). According to Silva et al. (1994), Ruhland et al. (2003), and Perera and Yatigammana (2015), diatoms are diverse eukaryotic unicellular microorganisms characterized by siliceous cell walls. Their ecological diversity is reflected by their occurrence in almost all aquatic habitats, where they play an important role as primary producers and in geochemical cycling of various naturally occurring elements, in particular carbon and silicon (Perera and Yatigammana, 2015). Diatoms grow abundantly in rivers, colonising almost all suitable habitats. They have been used extensively in studies related to the monitoring of water quality (Round, 1991). Diatoms can help build healthy stream environments by stabilizing the substrata and providing food and habitat for other biota within the ecosystem. Without diatoms and benthic algae, the stream ecosystem could become dysfunctional and collapse (Blinn and David, 2003) because diatoms can also tolerate variable environmental conditions, which make them extremely valuable in determining environmental disturbances by comparing pristine ecosystems versus impacted (Blinn and David, 2003). Diatoms are responsible for as much as 20% of carbon fixation globally by means of photosynthesis. This process of cycling materials through the areolae is what characterizes diatoms as excellent components for nutrient recycling and uptake (Ellie, 2014). The density and the diversity of phytoplankton are biological indicators for evaluating water quality and the degree of eutrophication (Shekhar et al., 2008). Some national parks diatoms that have been documented elsewhere in the world include, Spaulding et al. (2008) on stalk forming diatoms in Grand Teton national park Wyoming, Hobbs et al. (2009) on epipelic diatoms from Yellowstone national park Wyoming, Bartozek et al. (2013) gave report of eighty four diatoms from Iguaçu National Park in Brazil, Zhang et al. (2016) documented Oricymba xianjuensis sp. nov. in Xianju national park, China, Veselá and Johansen (2014) reported three new Eunotia spp from Acadia National park Maine while Ortiz-Lerin and Cambra (2007) discussed the distribution and taxonomy of 397 epilithic diatoms notes in rivers and streams of Northern Spain. In Nigeria, algological studies on lotic system are gaining interest but necessarily that of national parks and include that of Kadiri and Azomani (2000) who studied the effect of brewery effluent on the growth of two chlorophytes in Ikpoba River, Kadiri and Omozusi (2002) reported the phytoplankton of River Okhuahe in Benin, Kadiri (2007) reported the phytoplankton of River Ethiope. Most recent phycological information in similar freshwater bodies are available in Akoma and Opute (2010), Adesalu and Nwankwo (2010, 2012), Alika and Akoma (2012), Adesalu et al.

(2010, 2016a, 2016b), Ekhator et al. (2014, 2015), Adesalu (2016), Adesalu and Kunrunmi (2016), Akinyemi and Adesalu (2016). Of all mentioned work above, Okomu National Park water bodies and several other national parks except River Oli in Kainji Lake National park (Adesalu, 2010) water bodies remain without phycological information hence such studies remain important because majority of the riverine inhabitant most of the time, depend on their surrounding water (River) apart from rain water for their water needs. In this work, scanning electron micrographs of the most common species were also presented. The author hopes this will facilitate future taxonomic work.

MATERIALS AND METHODS

Description of study site

The Okomu National Park, (Figure 1) formerly known as the Okomu Wildlife Sanctuary, is a forest block within the 1,082 km² Okomu Forest Reserve in the Ovia South-West Local Government Area of Edo State in Nigeria (Birdlife International, 2016). The park is about 40 km West of Benin City (Williams 2008). The park holds a small fragment of the rich forest that once covered the region, and is the last habitat for many endangered species. It continues to shrink as villages encroach on it, and is now less than one third of its original size. The park holds a remnant of the Nigerian lowland forests that once formed a continuous 50-100 km wide belt from the Niger River west to the Dahomey Gap in Benin. The south and south-east of the forest is separated from the coast by mangrove and swamp forests, while to the north it merged into the Guinean Forest-Savanna Mosaic ecoregion. The park is drained by the Osse River which defines its eastern boundary. The Okomu River forms the western boundary. Among the common trees are Kapok, Celtis zenkeri, Triplochiton scleroxylon, Antiaris africana, Pycnanthus angolensis and Alstonia congoensis (Birdlife International, 2016). The park is probably the best example of mature secondary forest in south-west Nigeria. The park is accessible to tourists and has well marked trails.

Collection of sample

The study was based on a single sampling strategy during one week field trips to Okomu National Park, Edo state. Samples were collected on 25 April, 2015 from two locations each from two lakes called Lake 36 (N06° 23' 056; E005° 21' 165; N06° 23' 063; E005° 21' 156) and Lake 61 (N06° 67' 932; E005° 37' 794; N06° 19' 994; E005° 20' 718). Few of the physical parameters analyzed include surface water temperature which was measured using mercury in glass thermometer, the pH was determined with a Phillips pH meter (Model PW950) while surface water conductivities and salinity were determined using HANNA instrument (H18733) and hand refractometer respectively. Biological samples were stored in 5 L, concentrated and fixed with 4 % unbuffered formalin and analyzed with the aid of Olympus XSZ-N107 photomicroscope and JOEL JSM 6060LV for scanning micrographs.

Scanning Electron Microscopy preparation of the specimen and Light Microscopy

Biological samples preserved in 4% unbuffered formalin were allowed to settle for 24 hours in the laboratory, after which the supernatant was decanted until a concentrate of 100 ml was achieved. To facilitate diatom identification sub samples of the original collections were acidcleaned using the method suggested by Barber and Harworth (1981). A portion of the cleaned sample was added to distilled water. This slurry was filtered onto a 0.45 µm Millipore type HA filter. The filters were allowed to dry. A portion of the filter was cut and mounted to an aluminum stub using adhesive carbon tape. The aluminum stub was sputter coated with 20 nm of gold or goldpalladium. All scanning electron microscope observations were performed with a JOEL ISM 6060LV using a 10 kV accelerating voltage at St. Cloud State University, U.S.A. Description of some taxa frustule structures, valve sizes (diameter, length, width), number of striae/10 µm) and other taxonomic data were extracted from previous literatures of Ort'ız-Ler'ın and Cambra (2007) and Diatoms of United States (Online) accessed on the stated dates. This was necessary for proper understanding of the taxa. For photomicrographs, taxanomic keys employed in the identification included Hustedt (1930); Patrick and Reimer (1966, 1975); Prescott (1982); Krammer and Lange-Bertatot (1991); Whitford and Schumacher (1973), Metzeltin and Witkowski (1996), Lange-Bertalot and Metzeltin (1996).



Figure 1: Parts of Edo state map showing Okomu National Park and its environs.

RESULTS

The diatom composition of Okomu National Park is shown in table 1, a total of 29 diatom taxa belonging to 15 genera with more pennate forms than centric diatoms were observed. Out of the 15 genera, *Discotella* species was the only centric diatom observed. The pennate form which was dominated by *Eunotia* species include *Eunotia faba* Ehr.; *E. glacialis* Meister; *E.incisa* W. Smith ex.W.Gregory; *E. intermidia* (Krasske) Nörpel and Lange-Bertalot; *E. minor* (Kützing) Grunow; *E. monodon* Ehr.; *Eunotia pectinalis* (Kützing) Rabenh and *E. subarcuatoides* Alles, Nörpel and Lange-Bertalot (Figures 2 and 3) followed by *Pinnularia*

species which recorded four; Pinnularia braunii (Grunow) Cleve; P. brevicostata Cleve; P. gibba Ehr. and P. microstauron (Ehr.) Cleve (Plate 3). Other species include Achnanthidium minutissimum (Kützing) Czarnecki; Gomphonema parvulum (Kützing) Kützing; Frustulia rhomboides (Ehr.); Placoneis gastrum (Ehrenberg) Mereschkowsky; Sellaphora sp.; Eolimna minima (Grunow) Lange-Bertalot and W.Schiller; Chamaepinnularia mediocris (Krasske) Lange-Bertalot in Lange-Bertalot and Metzeltin; Luticola sp. and Achnanthes sp. (Figures 4 and 5). Nitzschia sigma (Kützing) W. Smith and Stauroneis anceps Ehr.

Table 1: Taxonomic classification of diatom flora at the Okomu National Park, Nigeria.

	Division: Bacillariophyta
	Class: Bacillariophyceae
	Order: Stephanodiscales
	Family: Stephanodiscaeae
1	Discotella sp.
	Order: Mastogloiales
	Family: Achnanthaceae
2	Achnanthes sp.I
3	Achnanthes sp.II
	Order: Cymbellales
	Family: Cymbellaceae
4	Cymbella sp
	Family: Gomphonemataceae
5	Gomphonema parvulum
6	Placoneis sp.
	Order: Cocconeidales
	Family: Achnanthidiaceae
7	Achnanthidium minutissimum (Kützing) Czarnecki
	Order: Eunotiales
	Family: Eunotiaceae
8	Eunotia faba Ehr.
9	E. glacialis Meister
10	E.incisa W.Smith ex.W.Gregory
11	E. intermidia (Krasske) Nörpel and Lange-Bertalot
12	E. minor (Kützing) Grunow
13	E. monodon Her
14	Eunotia pectinalis (Kützing) Rabenh
15	E. subarcuatoides Alles, Nörpel and Lange-Bertalot
16	Eunotia sp. I
17	Eunotia sp.II
	Order: Naviculales
	Chamaepinnularia mediocris (Krasske) Lange-Bertalot in Lange-
18	Bertalot and Metzeltin
19	Eolimna minima (Grunow) Lange-Bertalot and W.Schiller
	Family: Amphipleuraceae
20	Frustulia rhomboides (Ehr.) De Toni
	Family: Diadesmidiaceae

Description of some of the taxa

Eunotia pectinalis (Dillwyn, O. F. Müller),

Kützing) Rabenhorst 1864

Krammer and Lange-Bertalot 1991, 193, Fig. 141: 1-7, Fig. 143: 1. Metzeltin and Witkowski 1996, 98, Fig. 33:1; Lange-Bertalot and Metzeltin 1996, 146, Fig. 14: 5-7.

(Figure 2 A and B)

As described by Krammer and Lange-Bertalot (1991) and cited by Ortiz-Lerin and Cambra (2007), Frustules rectangular in girdle view. Valve elongates, slightly curved or bent. Ventral margin straight or slightly concave, sometimes slightly swollen at the centre of the valve. Valve narrowed toward broadly attenuated, truncate rounded ends. Striae parallel, slightly radiate at the ends. Striae: 7-12/10µm at the centre of the valve, to 14 in 10µm at the ends of the valve.

Eunotia monodon Ehrenberg 1843

Krammer and Lange-Bertalot 1991, 210, Fig. 158:1-6.

(Figure 2 C and D)

Ventral margin usually strong with short shells, forms with a hump in the middle. Dorsal edge most strongly convex, margin often parallel to ventral in the middle part, gradually declining and in front of the ends sometimes weakly drawn in. Terminal nodes and simply arched distal raphe close to the pole.

Eunotia subarcuatoides E. Alles, M. Nörpel and Lange-Bertalot 1991

Krammer and Lange-Bertalot 1991, 214, Fig. 138: 1-9, Fig. 145:6.

(Figure 2E)

Narrow linear in girdle view. Valves slender, bowshaped, as a rule strongly curved, with more or less parallel margins. The dorsal line somewhat strongly sloping toward the ends, valves therefore slightly narrowed towards the end; ends rounded. End nodules small, raphe slightly developed. Small forms, rarely over 30 µm long, strongly narrowed near the ends. Valves therefore strongly lanceolate, often only weakly bent. Striae: 18-23/10 µm. (Ortiz-Lerin and Cambra 2007).

Eunotia intermedia (Krasske ex Hustedt) Nörpel and Lange-Bertalot 1991 Krammer and Lange-Bertalot 1991, 215, Fig.143: 10-15.

(Figure 2 F and H)

Ventral margin straight, wall thickened in areas halfway between the centre of the valve and the ends. Dorsal margin convex. Ends not distinctly formed, but confluent with the rest of the valve. Terminal nodules distinct, near the ends of the valve. Striae parallel, 14-16/10 µm at the center of the valve, more numerous toward the ends (Ortiz-Lerin and Cambra 2007).

Eunotia glacilais Meister 1912

Krammer and Lange-Bertalot 1991, 207, Fig. 151:1-10A.

(Fig 2. I)

Frustule in girdle view, broad, rectangular with few bends, proximal raphe of a long and relatively wide hyaline area, ventral margin concave, dorsal margin parallel or slightly convex, gradually declining, the ends are slightly narrowed to slightly distended, bluntly rounded and usually bent posteriorly. Length (15) 30 approximately 200µm.

Eunotia minor (Kützing) Grunow in Van

Heurck 1881

Krammer and Lange-Bertalot 1991, 196,

Fig.142: 7-15, Fig. 144: 5

(Figure 2J)

Cells in girdle view thick, rectangular; laterally asymmetrical (lunate) in valve view along the apical axis. Valves bent with sub-rostrate large round apices. Ventral margin slightly concave but slightly inflated at the centre; dorsal margin convex. Striae uniseriate, on the ventral mantle more numerous and alternating, $9-15/10\,\mu m$.

Eunotia faba Ehrenberg1838

Krammer and Lange-Bertalot 1991, 225, Fig. 164:1-10.

(Figure 3A)

Bahls (2014) described *Eunotia faba* valves as asymmetric to the apical axis and variably asymmetric to the transverse axis. Dorsal margins are moderately arched. Ventral margins are weakly concave. The apices are broadly rounded, with a deep notch on the ventral side. Helictoglossae are removed some distance from the apices. Terminal raphe fissures are very short at the junction of the valve face and mantle. Striae are radiate and very finely punctate. Forked, or short, costae occur along the dorsal margin in some specimens.

Areolae are difficult to resolve in LM and number 30-35 in $10 \mu m$ (Bahls, 2014).

Eunotia incisa Gregory 1854

Krammer and Lange-Bertalot 1991, 221, Fig.161: 8-19, Fig. 162: 1-2, Fig. 163: 1-7 (Figure 3B)

According to Krammer and Lange-Bertalot 1991 as cited by Ortiz-Lerin and Cambra (2007), this species is characterized by the apices of the valve undifferentiated from the main body of the valve and by the very fine striae. Frustules in girdle view are rectilinear while frustules in valve view are with straight ventral margin and convex dorsal margin. Apices of the valve usually acute, sometimes rounded. Striae have more distance at the centre of the valve than at the ends, $13-17/10\,\mu m$.

Stauroneis anceps Ehrenberg 1843

Krammer and Lange-Bertalot 1991, 240, Fig. 87:3-9, Fig. 88:2-4 (Figure 3 E and F)

Stauroneis anceps valves are lanceolate to linear-lanceolate. The apices are protracted and narrowly rostrate. The axial area is moderately wide and linear, expanding slightly near the central area. The central area is a narrow rectangular stauros, hardly expanded toward valve margins and sometimes with shortened striae. The raphe is lateral. Proximal raphe ends are weakly expanded, weakly deflected, and widely spaced (not extend into the central area). Striae are radiate throughout. Areolae are variably spaced and number 20-28/10 µm. (Bahls, 2011).

Nitzschia palea (Kützing) W. Smith 1853 (Figure 3])

According to Kociolek (2011), *Nitzschia palea* valves are lanceolate with sides parallel and tapering rapidly at the poles, terminating with subcapitate apices. Fibulae are distinct, with a distinct central nodule and number 11-13 in 10 µm. Striae barely visible in LM. **Striae**, 36-38 /10 µm (Kociolek, 2011).

Pinnularia gibba Ehrenberg 1841

Krammer and Lange-Bertalot 1991, 423, Fig. 189:1-3,8,9.

(Figure 4A)

Tariq et al. (2006) described *Pinnularia gibba* valve as linear, gradually tapering to the subcapitate ends; raphe filiform with curved terminal fissures; axial

area linear, widened around the central nodule; striae divergent in the middle and convergent at ends; length of frustules 75-77 μm and breadth 10-13 μm ; striae 10-11 within 10 μm . (Tariq-Ali *et al.*, 2006)

Pinnulari microstauron (Ehrenberg) Cleve 1891

Krammer and Lange-Bertalot 1991, 425, Fig. 191:1-6; Fig.192: 1-16 (Figure 4B)

As described by (Bahls 2014), valves are linear with parallel or slightly convex sides. The apices are rostrate (much narrower than the valves). The axial area is narrow and widens abruptly just short of the central area. The central area is a rhombus, which widens along the apical axis to form a variably narrow to broad transverse fascia. The slightly curved raphe is lateral, proximal raphe ends are deflected to one side and terminate with small pores. Distal raphe fissures are shaped like commas or question marks. The distal raphe fissures are surrounded by striae at the poles. Striae are strongly radiate near the valve center, becoming strongly convergent near the apices. Occasionally an isolated short striae will occur on one side of the central fascia (Bahls, 2014).

Pinnularia brevicostata Cleve 1891

Krammer and Lange-Bertalot 1991, 410, Fig. 182: 4-7, 9

(Figure 4C)

Johnson (2006) described *Pinnularia brevicostata* valves as linear with bluntly rounded apices. Axial area broad, one-fourth to one-half of the valve breadth. Central area a narrow bilateral fascia. Striae nearly parallel throughout most of the valve, slightly radiate at the central area, parallel to slightly convergent at the apices, 9 in 10 μm. Longitudinal bands indistinct, occasionally visible near the valve margin. Raphe slightly undulates. Apical raphe fissures large, "comma" shaped.

Pinnularia braunii (Grunow) Cleve 1895

Krammer and Lange-Bertalot 1991, 416, Fig. 187:1-5

(Figure 4D)

Valves linear to linear-lanceolate with slightly convex or convex margins. Valve apices usually capitate, occasionally subcapitate. Axial area narrows at the apices, becoming progressively broader toward the central area. Central area large, elliptical a unilateral fascia or a bilateral fascia. Striae radiate at the central area, convergent at the apices, 10-11 in $10 \, \mu m$. Longitudinal bands absent (Jackson 1980).

Eolimna minima (Grunow) Lange-Bertalot and W.Schiller 1997

(Figure 4G)

As described by Spaulding and Edlund (2008), the cells of *Eolimna* are small, generally less than 20 µm in length. A narrow central sternum is present, and the raphe is straight. Valves lack apical pore fields. SEM features show areolae are positioned medially within the frustule wall and are covered internally by hymens.

Chamaepinnularia mediocris (Krasske) Lange-Bertalot in Lange-Bertalot and Metzeltin (Figure 4H)

Described by Otu and Spaulding (2011), Chamaepinnularia mediocris have linear to broadly oval valves, with rounded apices and a gibbous middle. The axial area is narrow near the apices, becoming wider near the central area. The central area is a transverse fascia: in some specimens it extends to the valve margins, in other specimens short irregular striae are present at the margins. The raphe is filiform and slightly arched. Terminal nodules are distinct. The striae are parallel and composed of a single, elongate areola on the valve face. A smaller areola is positioned on the valve mantle. Externally, the proximal raphe ends are straight. Internally, the proximal raphe ends are slightly deflected unilaterally.

Gomphonema parvulum, (Kützing) Kützing 1849

Krammer and Lange-Bertalot 1991, 358, Fig. 154:1-25

(Figure 5A, B and C)

Valves only slightly asymmetrical to trans apical axis (heteropolar), symmetrical to apical axis. Cells box-shaped in girdle view with pseudosepta visible. Apices rounded, sub-rostrate or rostrate (occasionally sub-capitate). Raphe often slightly sinuous. A single stigma is present on one side of the central area. Striae coarse and often visibly punctate - one short stria opposite the central stigma. Striae often almost parallel. Very variable species.

Frustulia rhomboides (Ehrenberg) De Toni Krammer and Lange-Bertalot 1991, 240, Fig. 95:1-3

(Figure 5D)

Valves rhomboidal to lanceolate, apices bluntly rounded to rostrate. Raphe central and contained in a median rib extending most of the length of the valve. At the apices, the raphe rib has the appearance of a pencil tip. Striae fine and arranged so as to produce a pattern of apical and trans apical rows (Kelly *et al.*, 2005).

Placoneis gastrum (Ehrenberg) Mereschkowsky

(Figure 5E)

Woodell (2015) described valves of Placoneis gastrum as elliptical, with broadly rounded to rostrate apices. Valves are usually slightly asymmetric to the apical axis. The axial area is narrow and the central area is irregular or transverse due to striae of variable length and arrangement. The raphe is simple, filiform, and slightly curved toward the secondary side of the valve (Woodell, 2015). The proximal raphe ends are slightly expanded. The distal raphe ends are abruptly hooked at the valve mantle. Striae are strongly radiate, curved and irregularly shortened near the central area. Striae are distinctly uniseriate, widely spaced from one another and number between 6-10 in 10 µm. Areolae number 24-27 in 10 μm (Woodell, 2015).

Achnanthidium minutissimum (Kützing) Czarnecki

(Figure 5F)

Potapova (2009) described Cells of Achnanthidium minutissimum as solitary or in very short chains, often attached to substrate by a stalk. Frustules are monoraphid with a concave raphe valve and convex rapheless valve. Valves are linearlanceolate with slightly drawn-out or slightly capitate ends, 1.5-3.3 µm wide, 5.6-20.8 µm long. Central external raphe ends are simple, terminal raphe fissures are short, almost straight, or absent. Internally, the central raphe ends are turned in opposite directions (Potapova 2009). Striae are radiate throughout both valves, 25-35 in 10 µm. Striae consist of one row of areolae. The striae are often interrupted in the central part of raphe valve to form a symmetrical or asymmetrical fascia. One row of elongated areolae is present on the valve

mantle. External areolae openings of areolae vary in shape from circular to transapically elongated slits. Internal openings of the areolae are elliptical, occluded by hymens perforated by small pores. Girdle bands are plain, open (Potapova, 2009).

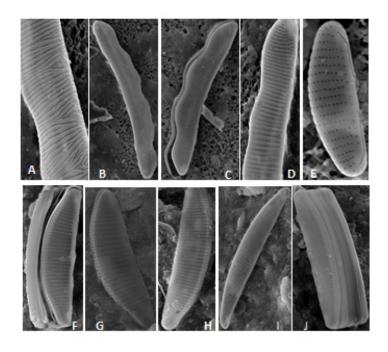


Figure 2: Valves view of some of the Eurotia species observed at the Okomu National Park Eurotia pectinalis (Kützing) Rabenhorst (A and B); E. monodonEhr (C and D); E. subarcuatoides Alles, Nörpel & Lange-Bertalot; (E); Eurotia intermedia (Krasske) Nörpel & Lange-Bertalot (F-H); E.glacialis Meister (I) and E. minor (Kützing) Grunow (J).

Scale bar is 10µm. A and D Scale bar is 5µm.

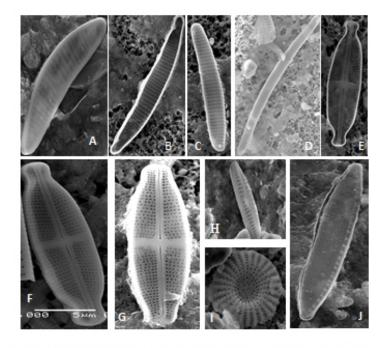


Figure 3: Eurotia faba Ehr. (A); E.incisa W.Smith ex.W.Gregory (B); Eurotia sp.I (C); Eurotia sp. II (D); Stawoneis anceps Ehr. (E and F); Stawoneis sp. (G); Cymbella sp. (H); Discotella sp. (I) and Nitzschiapaka (Kützing)W.Smith (I). Scale bar is 5µm

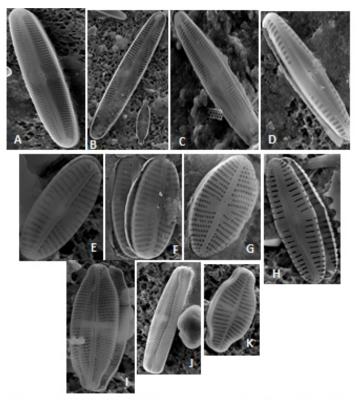


Figure 4: Pinnularia gibba Ehr (A); P. brevicostata Cleve (B); P. microstauron (Ehr.)
Cleve (C); P. braunii (Grunow) Cleve (D); Seliaphora sp. (E and F); Eolimna minima
(Grunow) Lange-Bertalot & W.Schiller (G); Chamaepinnulariamediocris (Krasske)
Lange-Bertalot in Lange-Bertalot & Metzeltin (H); Luticola sp. (I); Achmanthes sp. I (J)
and Achmanthes sp. II (K). Scale bar is 10µm (A-D); 1µm (E-H); 2µm (I-J)

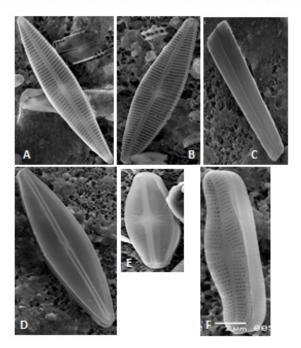


Figure 5: Gomphonema parvulum (Kützing) Kützing (A and B) valve view, G. parvulum (C) girdle view; Frustulia rhomboides (Ehr.) De Toni (D); Placoneis gastrum (Ehrenberg) Mereschkowsky (E); and Achnanthidium minutis simum (Kützing.) Czarnecki (F).

Scale bar is 10µm. (A-D); 2µm. (E. and F)

DISCUSSION

This report supported Pierre (1996) and Ortiz-Lerin and Cambra (2007) who noted that Eunotia is essentially a freshwater diatom genus frequently associated with acidic waters while Ortiz-Lerin and Cambra (2007) gave taxonomic notes on Eunotia. The frequency of Eunotia species at the Okomu National park is also in conformity with the report of Alles et al. (1991) on the tolerance of Eunotia species specifically E. subarcutoides to high and strong variations of pH values. It is of interest to note that different species of Eunotia followed by Pinnularia dominated this seasonal water body as these organisms are considered important ecological indicators (Alles et al. 1991, Kwandrans 2007). The dominance of Eunotia over the other taxa also correlates with the result of Liu et al. (2011) who reported twenty eight taxa of Eunotia in Da'erbin Lake and Surrounding Bogs in China. According to Mayama and Kobayasi (1991); Alles et al. (1991); Levkov et al. (2000, 2005); Kwandrans (2007) and Pavlov and Levkov (2013), the restriction to the freshwater oligotrophic and oligosaprobic habitats for Eunotia is a unique ecological feature of this genus, enabling it to play an important role as an indicator of water quality monitoring. It can be said that the water body within the Okomu National park is supporting ecological growth; however since this is first attempt in documenting the microalgae taxa of this Park it is recommended that more phycological work be done in future.

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