

Research Article

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Comparative Analysis of Organic and Inorganic Fertilizers Effects on Zooplankton Population Dynamics and Physicochemical Parameters of Culture-Media

*Austeen I. Obhahie^{1,2}, Johnson O. Oyewumi¹ and Andrew O. Agbon^{1,3}

¹Department of Animal and Environmental Biology, Federal University Oye-Ekiti, Nigeria ²Department of Animal and Environmental Biology, Delta State University, Abraka, Nigeria ³Department of Biology, National Open University of Nigeria, Headquarters, Jabi, Abuja, Nigeria **Corresponding Author's email*: <u>obhahie@delsu.edu.ng</u>; Phone: +2348060132602

ABSTRACT

Efficacy of inorganic (NPK 15:15:15 and Urea) and organic (poultry-droppings) fertilizers in culturing freshwater mixed-species-zooplankton were appraised for 6 weeks, using two tanks designated Tank-A and Tank-B, treated with inorganic and organic fertilizers respectively. Mixed-species-zooplankton were obtained from the wild, nursed in indoor aquaria and used to inoculate outdoor concrete tanks for zooplankton mass-production. Physicochemical parameters cumulative-mean obtained in Tank-A and Tank-B respectively were temperature [⁰C] (25.19; 24.99), pH (7.38; 7.40), electrical-conductivity (EC) [^{μs}/_{cm}] (443.66; 387.16) and transparency [cm] (25.34; 24.66), other values (in ppm) were, total-dissolved-solids (TDS), (191.1; 209.68), total-alkalinity [TA] (108.1; 150.98), dissolved-oxygen [DO] (7.80; 5.66), BOD (2.23; 2.42), free-CO₂ (23.0; 31.0), Nitrate (4.62; 3.57), Phosphate (2.47; 2.15) and Potassium (11.49; 10.28). Three zooplankton taxa; Rotifera, Cladocera and Copepoda comprising 15 species were identified. Population density average-mean in Tank-A and Tank-B respectively were combined mixed-species-zooplankton (440 inds.ml⁻¹; 389 inds.ml⁻¹), Rotifera (208; 188 inds.ml⁻¹), Cladocera (129; 110 inds.ml⁻¹), and Copepoda (103; 91 inds.ml⁻¹). Population percentage distribution in Tank-A and Tank-B were Rotifera (47.37; 48.26%), Cladocera (29.28; 28.32%) and Copepoda (23.35; 23.42%) respectively. Temperature, pH, BOD and Phosphate had no significant difference (P < 0.05), while TDS, EC, TA, DO, CO₂, Transparency, Nitrate and Potassium showed significant differences (P < 0.05). Zooplankton population density also showed significant differences (P < 0.05) between the values of Tank-A against Tank-B. However, the overall performance of both fertilizers and zooplankton response were within the optimum ranges for simulated aquatic environments' requirements for zooplankton mass-production.

Keywords: Formulated-feed; Inorganic-organic-fertilizers; Mixed-species-zooplankton; Physicochemical-parameters; Population-dynamics

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INTRODUCTION

Zooplankton are organism that are drifted by lotic waters and floats on lentic water body. Zooplankton sizes range from tenth of millimeter to four millimeter, (Eya, 2003; Shulz, 2012). They serve important role in food-chain and food-web through serving as a link between lower trophic and higher trophic level organisms, (Shulz, 2012; wikiwand, 2025). Freshwater zooplankton has the potential of replacing *Artemia* and artificial feeds in fresh-water

fish breeding, (Oladele and Omitogun, 2016; Ekelemu and Nwabueze, 2010). Zooplankton can serve as biological monitoring agents as regards toxicology and aquatic ecosystems' reaction to climate change, owing to their abundance and species diversity, due to the fact that they have short lifecycles that generally spans within weeks, (wikiwand, 2025; Shulz, 2012; Mackas and Beaugrand, 2010). They are able to provide adequate protein needed for growth and proper development of several species of fish larvae, (Ovie and Ovie, 2002; Amali and Solomon, 2001; Lubzens *et al.* 2001).

Zooplankton are categorized based on their lifecycle and size. On the basis of life-cycle, they are classified into holozooplankton (i.e. zooplankton that live their entire lifecycle as plankton) and merozooplankton (i.e. living part of their life as plankton before metamor-phosing into either nekton, sessile or benthic living), (Shiffert, 2020; wikiwand, 2025), while based on size, they are classified as picozooplankton (2um). nanozooplankton (2 - 20µm), microzooplankton (20 - 200µm) and mesozooplankton (0.2 - 20 mm). Among the larger size of zooplankton are the mesozooplankton, which are dominated by copepods, while the preceding size are the microzooplankton which are mostly rotifers and cladocerans, (wikiwand, 2025; Shulz, 2012; Eya, 2003; Schwoerbel, 1970).

Zooplankton feed on fresh-sprout bacterioplankton, phytoplankton, organic-detritus or prey on smaller animals depending on species, (Shulz, 2012; Eya, 2003). They are present and wildly distributed over the water-body column, being mostly abundant where food is available. Zooplankton biomass and species composition in ponds and shallow water fluctuates depending on interrelating driving force that may include polymixis, condition of weather, change in water level, food/feeding management and nutrients load, (Shulz, 2012; Borics et al. 2000). Studies have shown that quality and quantity of zooplankton abundance varies from one location to another under the same environment and within the same ecological conditions, (Bhuiyan et al. 2008). Season, physicochemical parameters, such as soil, watermovement and biological factors affects zooplankton distribution and abundance, (Davis et al. 2009).

Owing to the driving force and factors affecting zooplankton availability and abundance, there is need for managed simulated environment, suitable for mass production of zooplankton to meet the need of freshwater fish breeder, (Eya, 2003; Shulz, 2012). The commonly cultured zooplankton group for commercial finfish and shellfish rearing are rotifera, cladocera and copepoda, (Shulz, 2012; Eya, 2003).

Freshwater fish species spawns freely in the wild, but with low survival rate, as a result of climatic and biological setbacks, (Okogwu et al. 2006). The achievement made by induced fish breeding under managed hatchery is lost through high mortality rate at early days of fry-life (Okogwu et al. 2006). Fish larvae poor growth and high mortality rates have been traced to several factors, among which inappropriate, inadequate, lack of suitable food and poor water quality predominates, (Eya, 2003; Okogwu et al. 2006; Obhahie, 2022). Thus, comprehension of managed production of freshwater zooplankton will be of great assistance to freshwater fisheries industry in Africa and particularly in Nigeria, (Okunsebor, 2014; Ekelemu and Nwabueze, 2010).

Neonate fish larvae are fragile, minute, immature, have tiny mouth size, undeveloped chemoreceptors, eyes and digestive system that limit their proper food choice and intake at the point of exogenous first-feeding stage, (Lavens and Sorgeloose, 1996). The nutritional value of formulated feed is inadequate to supply the required nutrients for fish larva, when compared to natural fish food that from studies has shown the potential to meet the necessary nutritional requirement, (Okunsebor, 2014). These qualities inform Aquaculturists' selection criteria for suitable fish larva diet, (Figure 1[adapted from: Lavens and Sorgeloose, 1996]).

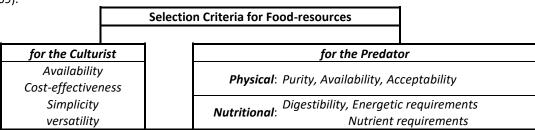


Figure 1: Criteria for fish larva food selection

Common culturable zooplankton for freshwater fish larvae are, Rotifera (e.g. *Brachionus* sp.), Cladocera (e.g. *Moina* spp.) and copepod (e.g. *Cyclopod* spp.). Freshwater zooplankton maintains a life-span of about four weeks in the freshwater habitat, (Delbare and Dhert, 1996; Eya, 2003; Shulz, 2012), and can be easily mass produced, harvested and use for freshwater larval culture (Ajah, 2010; Shulz, 2012)

Fertilization of zooplankton culture tank is a common practice for establishing adequate quantities and qualities of zooplankton production for fish. The applied fertilizer could be inorganic or organic based. Inorganic fertilizers commonly

applied are those that have components of nitrate (N) phosphorus (P) and potassium (K) or Urea (Boyd and Massaut, 1999; Obhahie, 2022; Eya, 2003). While organic fertilizers commonly applied could be sourced from wide range of animals' faeces or sourced from plants (Knud-Hansen, 1998). Suitable organic fertilizers should have low carbon-nitrogen ratio and fine particle size to allow rapid decomposition, (Geiger and Turner, 1990).

The assertion behind culture fertilization is to provide nutrients for phytoplankton increase production, which will in-turn favour increase of zooplankton population that feed on phytoplankton. Effect of fertilizers application on the quality and quantity of zooplankton culture is determined by the relative composition of zooplankton species in a particular culture pond. Rotifera, Cladocera and Copepod, species are commonly used zooplankton for fish larvae rearing. Their age at maturity, rate of egg production and quantity of eggs produced is affected by their food quality and quantity, (Eya, 2003). Thus, inorganic and organic fertilizers are applied to increase phytoplankton production that in-turn provides essential nutrient to enhance productivity of zooplankton which eventually serve as food for fish larvae, (Young and Flickinger, 1988; Boyd and Massaut, 1999; Obhahie, 2022).

MATERIALS AND METHODS

Study Area Description

The experiment was carried out in concrete tanks situated at the Hatchery in the Department of Animal and Environmental Biology, Federal University Oye-Ekiti, (FUOYE), main campus, phase II, Oye-Ekiti, Nigeria. Water sampling for physicochemical parameters and zooplankton analysis was carried-out in 6 weeks, spanning through October to December. The culture concrete tanks were designed to accommodate 1,200 litres (L) of water, but were filled to a volume of 1,000 L. The culture tanks were filled with water from borehole. The Hatchery has perimeter fence for safety and prevention from intruders.

Concrete Tanks Preparation

Two tanks designated Tank-A and Tank-B were used for the experiment. The tanks were checked for firmness and leakages by letting water into them and observed for 24 hours and drained. Noticed leakages were mended and thereafter water was let-in and left for some days for curing to be carriedout.

Concrete Tanks Curing

Tank curing was achieved by letting in water, in addition with fertilizer for rapidity. The tanks were left uninterrupted for some days, except for interval stirring to enhance nutrient circulation, until algae and insects' larvae were noticed in the tanks, indicating the tanks' ability to support life.

Zooplankton Culture Tanks Fertilization, Maintenance and Management

Fertilization of tanks for algae production was carried out by adding 20 g each of inorganic fertilizers, (NPK-15:15:15 and Urea) to Tank-A and addition of 200 g of organic fertilizer, (dried-pure poultry-droppings) to Tank-B. The inorganic fertilizers were applied via direct dissolution into Tank-A, while the organic fertilizer was dissolved in water, sieved and the solute poured into culture Tank-B. The tanks were left undisturbed, except for regulated stirring for nutrient circulation. When phytoplankton bloom was adjudged well established and rich enough to sustain zooplankton mass production, the culture tanks media were inoculated with nursed mixed-species-zooplankton samples. Culture media were renewed weekly by addition of prepared fertilizers. Zooplankton culture media were sometimes slightly drained and refilled with water in-order to cut-back excess fertilizer and achieve optimum phytoplankton bloom, in accordance to Eya, (2003); Arimoro, (2006); Obhahie, 2022.

Wild-zooplankton Collection, Nursing and Culture Tanks Inoculation

Wild mixed-species-zooplankton samples were collected from lentic ponds at Itaji, Oye, Ekiti State, with the aid of 50µm mesh-size plankton net and poured into a plastic-bucket until 10 L samples was collected and transported to Hatchery at FUOYE, where it was filtered with 1mm mesh-size net and introduced (at 5 L each) into two 30 L plasticaquaria, tagged Tank-A and Tank-B, containing algae culture media obtained from concrete Tank-A and Tank-B respectively. The wild-zooplankton samples were nursed, for acclimatization and mass multiplication in the plastic-aquaria for about a week before being gradually used to inoculate the concrete culture tanks containing 1,000 L of water, in accordance to the methods described by Eya (2003) and Obhahie, (2022), while some were reserved in the plastic-aquaria culture for continuous close range culture and multiplication. Zooplankton samples collection was done in accordance to Goswami, (2004).

Wild and Cultured Zooplankton Sample Qualitative and Quantitative Analysis

A portion of the filtered wild-zooplankton sample was transferred into a 100 ml bottle and preserved with 5% formaldehyde, while the larger portion was used to inoculate the plastic-aquaria zooplankton nursing culture media.

Zooplankton samples were harvested from the concrete tanks culture media 3 times a week in 10 L plastic bucket and filtered. A portion of it was

transferred into a 100 ml bottle and preserved with 5% formaldehyde, while the remaining larger portion was used to feed fish larvae. The preserved zooplankton samples were examined with microscope to ascertain the qualitative and quantitative nature of its composition. Zooplankton samples collection was carried-out in accordance to Goswami, (2004).

Zooplankton samples identification was carried out by examining about 25 ml of the preserved sample. A drop (1 ml) of the sample was made on countingchamber, placed on an electric-light-microscope and examined using two magnifications of x40 (to obtain a general view of the species composition) and x100 (to ascertain species identification). Zooplankton identification and enumeration was done in accordance to Jeje and Fernando, (1986), Shiel, (1995) and Goswami, (2004).

Physicochemical Parameters

Water samples were obtained from culture tanks 3 times per week and analyzed for physicochemical parameters, which includes Temperature, pH, Total Dissolved Solids (TDS), Electrical Conductivity (EC), Transparency, Total Alkalinity (TA), Dissolved Oxygen (DO), Biochemical Oxygen Demand (BOD), Free Carbon Dioxide (CO₂), Nitrate (N), Phosphate (P) and Potassium (K). Temperature, pH, TDS and EC were determined with the aid of Digital-Multiple-Parameters-Tester (DMPT). Transparency was determined with Secchi-disc. TA, BOD, Free CO2 and DO were analyzed by titrimetric method, in accordance to APHA, et al. (2017). Nitrate (N) and Phosphate (P) were determined with Spectrophoto-meter, while Potassium (K) was determined by atomic Absorption Spectrophotometer (AAS).

Data Analysis

The physicochemical parameters and zooplankton species population densities data were analyzed

using the descriptive statistics to obtain mean (M) and standard deviation (SD), while analysis of variance (ANOVA): t-Test: Paired Two Sample for Means at 95% confidence level were used to determine the significant difference among means. The effect of types of fertilizers on zooplankton species population density and physicochemical parameters values are as presented in columnchart figures.

RESULTS

Physicochemical Parameters

Table 1 records zooplankton culture media physicochemical parameters obtained in Tank-A. Column 1 – 6 contains the weekly mean (\overline{x}) and standard deviation (SD/±) of the 12 parameters, while column 7 bears Week 1 – 6 cumulative mean and SD.

Table 2 recorded the zooplankton culture media physicochemical parameters obtained in Tank-B. Column 1 – 6 contains the weekly mean (\overline{x}) and standard deviation (SD/±) of the 12 parameters, while Week 1 – 6 cumulative mean and SD are recorded in column 7.

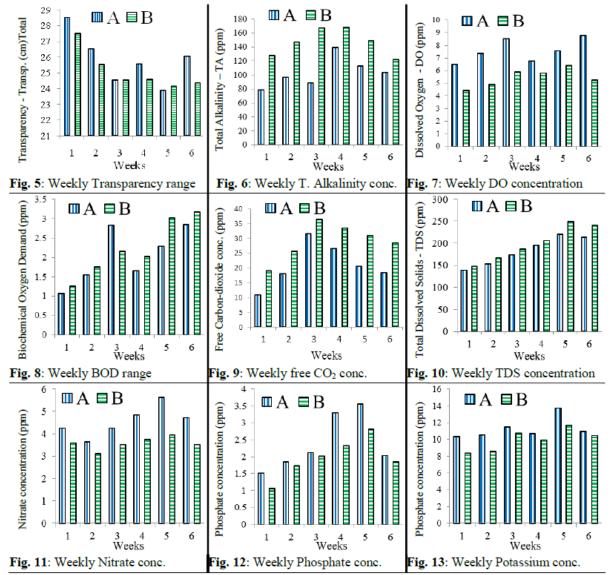
The mean values of Tank-A and Tank-B are plotted in column-chart (figures 2 – 13). The physicochemical parameters mean values fluctuated slightly. Tank-A recorded slightly higher mean values in temperature, EC, transparency, DO, nitrate, phosphate and potassium, while Tank-B had higher mean values in pH, TDS, TA, BOD and CO₂.

Temperature, pH, BOD and Phosphate exhibited no significant difference (P < 0.05) between their values in Tank-A against Tank-B, while the values of TDS, EC, TA, DO, CO_2 , Transparency, Nitrate and Potassium had significant differences (P < 0.05) between their value in Tank-A against Tank-B.

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Parameters	WEEK 1	WEEK 2	WEEK 3	WEEK 4	WEEK 5	WEEK 6	WEEK 1 - 6
Temperature (^o c)	26.54±0.53	25.53±0.42	25.57±0.47	24.65±0.94	24.67±1.25	25.52±0.41	25.19±0.48
рН	7.02±0.11	7.06±0.63	7.27±0.43	7.37±0.41	7.57±0.44	7.65±0.35	7.38±0.24
EC (µs/cm)	394.2±21.16	412.3±22.34	462.5±24.52	455.4±13.79	422.8±24.97	465.3±23.25	443.66±24.4
Transparency (cm)	28.52±0.52	26.54±0.52	24.56±0.54	25.58±1.48	23.92±0.72	26.08±1.07	25.34±1.08
TDS (ppm)	138.2±23.97	152.8±34.75	173.7±26.53	195.2±27.31	220.3±24.18	213.5±32.36	191.1±28.02
TA (ppm)	78.50±23.56	96.50±19.54	88.5±22.51	139.4±22.53	112.7±16.62	103.4±15.65	108.1±19.63
DO (ppm)	6.51±0.12	7.34±0.12	8.52±0.41	6.76±0.58	7.58±0.46	8.78±0.62	7.80±0.84
BOD (ppm)	1.06±0.29	1.55±0.23	2.83±0.38	1.65±0.64	2.28±0.36	2.85±0.32	2.23±0.62
Free CO2 (ppm)	11.00±1.41	18.00±1.42	31.50±2.12	26.50±0.71	20.50±0.71	18.50±0.71	23.00±5.83
Nitrate (ppm)	4.25±0.35	3.65±0.23	4.25±0.45	4.85±0.33	5.64±0.32	4.72±0.22	4.62±0.74
Phosphate(ppm)	1.51±0.25	1.85±0.02	2.12±0.04	3.31±0.05	3.55±0.02	2.03±0.43	2.47±0.66
Potassium (ppm)	10.36±2.03	10.57±2.01	11.51±2.12	10.67±1.08	13.77±1.03	10.92±1.03	11.49±1.33

Parameters	WEEK 1	WEEK 2	WEEK 3	WEEK 4	WEEK 5	WEEK 6	WEEK 1 - 6
Temperature (°c)	25.35±0.52	24.95±0.42	25.15±0.33	24.33±0.27	25.35±0.25	25.17±0.23	24.99±0.40
рН	7.17±0.13	7.14±0.23	7.23±0.15	7.44±0.33	7.47±0.46	7.72±0.63	7.40±0.23
EC (µs/cm)	374.1±1 4.25	386.6±23.43	396.7±25.61	383.9±21.88	364.4±14.16	404.2±22.34	387.16±15.09
Transparency (cm)	27.53±0.53	25.55±0.56	24.57±0.52	24.59±0.47	24.15±0.45	24.38±1.86	24.66±0.53
TDS (ppm)	148.1±32.86	167.2±24.64	186.1±35.42	206.1±33.29	248.6±26.27	240.4±27.45	209.68±37.76
TA (ppm)	127.5±24.55	147.5±13.53	167.5±23.52	168.3±23.54	149.3±23.37	142.3±22.54	154.98±12.07
DO (ppm)	4.45±0.22	4.92±0.21	5.88±0.43	5.82±0.52	6.42±0.24	5.26±0.62	5.66±0.58
BOD (ppm)	1.25±0.22	1.75±0.26	2.15±0.53	2.02±0.53	3.02±0.49	3.17±0.88	2.42±0.63
Free CO2 (ppm)	19.00±1.41	25.50±0.71	36.50±2.12	33.50±0.71	31.00±1.41	28.50±2.50	31.00±4.27
Nitrate (ppm)	3.60±0.25	3.10±0.35	3.51±0.15	3.75±0.43	3.95±0.64	3.52±0.12	3.57±0.32
Phosphate(ppm)	1.05±0.33	1.73±0.01	2.02±0.05	2.32±0.02	2.82±0.03	1.85±0.13	2.15±0.44
Potassium (ppm)	8.35±0.15	8.59±1.02	10.72±1.09	9.94±2.05	11.72±1.02	10.45±1.06	10.28±1.15

Abbreviations: EC [Electrical Conductivity]; Total Dissolved Solids (TDS); TA [Total Alkalinity]; BOD [Biochemical Oxygen Demand]; CO₂ [Free Carbon Dioxide]; DO [Dissolved Oxygen]



Zooplankton Population Density

Three zooplankton taxa, Rotifera, Cladocera and Copepoda comprising 15 species were identified. Table 3, column 1 -12, row 1 -3 shows the weekly zooplankton taxa popula-tion mean obtained from samples analysis and presented in column-chart, figures 14, 15 and 16.

Table 3, row 4 (Total) shows week 1 - 6 total collective mean (\bar{x}) of mixed-species-zooplankton population densities that are presented in column-chart in Figure 17. The population density mean-

values-total of mixed-species-zooplankton obtained from Tank-A and Tank-B, in each Week 1 - 6 were 432 and 388 inds.ml⁻¹; 427 and 399 inds.ml⁻¹; 451 and 401 inds.ml⁻¹; 438 and 391 inds.ml⁻¹; 447 and 365 inds.ml⁻¹; 437 and 389 inds.ml⁻¹ respectively.

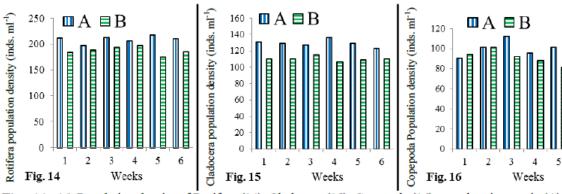
Table 3, columns 13 and 14, shows week 1 - 6 cumulative mean (\bar{x}) of mixed-species-zooplankton population densities, which are plotted in Figure 18. The population density mean values of zooplankton varied-taxa reared in Tank-A and Tank-B were, Rotifera (208.4 and 188.7 inds.ml⁻¹), Cladocera (128.8 and 110.2 inds.ml⁻¹) and Copepoda (102.7

and 91.14 inds.ml⁻¹) respectively, and its represented with column-chart by Figure 18 Columns 15 and 16 of Table 3, recorded collective average week 1 – 6 zooplankton taxa percentage (%) mean population distribution that are plotted in Figure 19. The obtained population percentage distribution in Tank-A and Tank-B were Rotifera (47.37; 48.26%), Cladocera (29.28; 28.32%) and Copepoda (23.35; 23.42%) respectively.

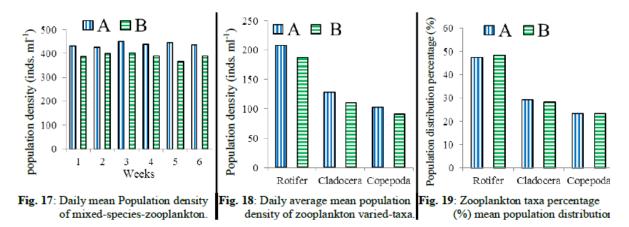
The t-test: paired two sample for means statistical analysis of the zooplankton population density showed that there was significant differences (P < 0.05) between the value obtained from Tank-A when compared to those obtained from Tank-B.

Table 3: Zooplankton Taxa Population Densit	y Mean and Percentage in Tank-A and Tank-B (i	inds.ml ⁻¹)

Zooplankton	We	ek l	We	ek 2	We	ek 3	_ We	ek 4	We	ek 5				- 6 😰	Wk 1	- 6 (%)
Taxa	A	В	A	В	A	В	A	B	A	B	A	B	A	B	A	B
Rotifer	211	184	197	188	212	194	206	197	217	175	210	185	208.4	187.8	47.37	48.26
Cladocera	131	110	129	110	127	115	136	107	129	109	123	110	128.8	110.2	29.28	28.32
Copepoda	90.3	94.3	101	101	112	92.2	95.7	87.7	101	80.8	104	93.6	102.7	91.14	23.35	23.42
Total	432	388	427	399	451	401	438	391	447	365	437	389	440	389	100.0	100.0



Figs. 14 - 16: Population density of Rotifera (14), Cladocera (15), Copepoda (16) reared on inorganic (A) and organic (B) fertilizers



DISCUSSION

Inorganic and organic fertilizers are commonly used in zooplankton culture to enhance algal blooms that in-turn serve as source nutrients to zooplankton, (Boyd and Massaut, 1999; Shulz, 2012; Ekelemu and Nwabueze, 2010). Inorganic fertilizer favours higher population density of zooplankton (Mosha *et al.* 2016) owing to its immediate release of nutrients, (Boyd and Massaut, 1999). Organic fertilizers may in contrast promote excessive phytoplankton blooms that could potentially affect water quality, survival, abundance and composition

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of zooplankton, (Boyd and Massaut, 1999). Commendable improvements were reported in inorganic and organic fertilizers treated zooplankton production tanks when compared to unfertilized culture (Mosha *et al.* 2016; Boyd, 1982). Organic fertilizer application have been reported to have creditable results on zooplankton culture, being that they are proficient in supplying the required nutrients necessary for zooplankton mass production (Orji and Chibugwu, 2010).

The zooplankton culture media physicochemical parameters obtained in Tank-A fluctuated slightly against that of Tank-B. The mean values obtained were within suitable range for tropical freshwater zooplankton production in simulated aquatic environments, when evaluated with previous works carried-out by Orji and Chibugwu, (2010), Akodogbo *et al.* (2014), Mosha *et al.* (2016), Yakubu *et al.* 2018) and several other Scientists.

The temperature mean ranges were $24.65 - 26.54^{\circ}$ C in Tank-A and 24.33 - 25.35 in Tank-B. These mean values are within the required limits for zooplankton optimum multiplication, (Arimoro and Ofojekwu, 2004), being that at a temperature below 20° C the rate of zooplankton multiplication decreases in culture ponds, (Kim, 1972). The general temperature of the experiment's vicinity is within $24 - 27^{\circ}$ C range, which is appropriate for outdoor zooplankton culture, (Ludwig, 1993; Mosha *et al.* 2016).

The pH values were alkaline, ranging from 7.02 – 7.65 in Tank-A and 7.14 – 7.72 in Tank-B, which are consistent with previous studies, (Islam *et al.* 2000; Oladele and Omitogun 2016). Research reports earmarked the optimum zooplankton culture average pH requirement to be within 6.5 – 8.5 pH range, (Eya, 2003; El-Naggar *et al.* 2008), which aligns with the result of this investigation.

Naturally transparency tends to be inverse to TDS, same tendency was observed in this investigation. The Tank-A and Tank-B highest transparency mean (28.52 and 27.53cm) coincided with the lowest TDS mean (138.2 and 148.1 ppm) in week 1, equally, transparency lowest mean (23.92 and 24.15 cm) coincided with TDS highest mean (220.3 and 248.6 ppm) at week 5, in Tank-A and Tank-B respectively. This gradual increase in TDS (turbidity) and inverse corresponding decrease of transparency is as a result of phytoplankton and zooplankton population density proliferation within the period, (Reid and Wood, 1976; Adeniji *et al.* 1997; Azionu *et al.* 2001).

The EC started with its lowest mean values (394.2 and 374.1 μ s/cm) at week 1, in both tanks. The highest values (465.3 and 404.2 μ s/cm) were achieved in week 6 in Tank-A and Tank-B respectively. There were gradual mean values rise

from week 1 to week 3, followed by a gradual decline and a sharp increase in weeks 6. This model of EC behaviour may have been influenced by the pH and alkalinity content of the culture media, being that EC, pH and alkalinity exhibits interwoven relationship, (Akindele, 2013; Ovie *et al.* 2001; Azionu *et al.* 2001).

The total alkalinity (TA) lowest mean (78.5 and 127.5 ppm) were recorded in week 1, in both tanks and the maximum mean (139.4 and 168.3 ppm) were obtained at week 4 in both tanks, followed by decreasing fluctuations in weeks 5 and 6. These decreasing trends may have been influenced by the regimes of fertilizer renewal of the culture media, (Arimoro, 2006). The TA mean values obtained in this investigation were within the natural, 40 - 41.5 ppm recorded by Ovie *et al.* (2001) at Dadin Kowa and Kiri reservoirs and the fertilized, 60 - 340 ppm obtained by El-Naggar *et al.* (2008) in an experiment conducted in earthen ponds.

Dissolved oxygen (DO) lowest mean (4.45 ppm) was recorded in Tank-B at week 1. This could be attributed to the high unutilized organic nutrients from manure at the start of the experiment. As algal bloom proliferates, the DO also improved, reaching its Tank-B maximum (6.42 ppm) at week 5. Tank-A DO lowest mean (6.51 ppm) was recorded in week 1, and it gradually improved to reach its maximum (8.78 ppm) at week 6. This improvement in DO could be attributed to the gradual algal bloom, (Boyd and Massaut, 1999; Eya, 2003).

Tank-A and Tank-B respective BOD lowest mean (1.06 and 1.25 ppm) were recorded in at week 1. This could be attributed to the relatively low and reduced metabolism at the start of work, but as algal bloom increased, metabolism also increased, thereby increasing the BOD gradually to its maximum (2.85 and 3.17 ppm) at week 6, in Tank-A and Tank-B respectively. The gradual increase of BOD, from week1 – 6 is most probably connected to the gradual accumulation of detritus resulting from increased metabolism, (Ovie *et al.* 2001; Azionu *et al.* 2001; Boyd and Massaut, 1999; Eya, 2003; Obhahie *et al.* 2007).

The free-carbon-dioxide (CO₂) lowest mean values (11.00 and 19.00 ppm) were obtained at week 1, in Tank-A and Tank-B respectively. The maximum mean values (31.50 and 36.50 ppm) were recorded at week 3 in Tank-A and Tank-B respectively, followed by a gradual decline and steady increase in weeks 4, 5 and 6.

These patterns of CO_2 behaviour could be associated with organic components gradual buildup and decline in the culture tank, in like manner as the occurrence of organic effluents build-up in water-bodies, (Boyd and Massaut, 1999; Obhahie *et al.* 2007). CO_2 is phytoplankton source of nutrient and its level in ponds exhibits diurnal fluctuation, being highest in the morning and experiencing gradual depreciation as the day progresses owing of to the process photosynthesis of phytoplankton, (Hargreaves and Brunson, 1996). The experiment started with a moderate mean value of Nitrate (N) of 4.25 and 3.60 ppm, Phosphate (P) of 1.51 and 1.05 ppm and Potassium (K) of 10.36 and 8.35 ppm, at week 1, and a gradual mean values increase culminating at week 5 with maximum mean value of N of 5.64 and 3.95 ppm, P of 3.55 and 2.82 ppm and K of 13.77 and 11.72 ppm, in Tank-A and Tank-B respectively. Week 6 experienced a slight mean values decrease.

NPK are mainly sourced from fertilizers. The regular supply of these nutrients was maintained by periodical renewal, carried-out either weekly or as determined by their available concentration in the culture media. There were situations when the culture media have to be slightly drained and refilled with water in-order to cut back the amount of fertilizer content in it, so as to achieve acceptable nutrient limits and optimum phytoplankton bloom for zooplankton consumption, (Boyd and Massaut, 1999; Eya, 2003; Arimoro, 2006).

The fertilizers reduction was based on deleterious algae bloom and density of zooplankton in the culture tanks and care was taken not to overfertilize the media, being that algae not consumed beyond a certain period degrades, increasing the level of ammonia, BOD and CO₂, while inhibiting the dissolved oxygen and pH level in the culture media, (Arimoro, 2006; Eya, 2003; Hargreaves and Brunson, 1996; Boyd, 1982). This research report revealed that inorganic and organic fertilizers culture media physicochemical parameters values were within the optimum ranges for simulated aquaculture ecosystems notwithstanding the occurrence of significant difference observed.

Three zooplankton taxa, comprising Rotifera, Cladocera and Copepoda, made-up of 15 species were identified in the culture media samples analysed from both Tank-A and Tank-B. The species composition distributions were 6 species of Rotifera, 5 species of Cladocera and 4 species of Copepoda. Quantitatively and qualitative dominance of the 3 zooplankton taxa were, Rotifera > Cladocera < Copepoda in both tanks.

The cumulative population density mean for week 1 - 6 in Tank-A and Tank-B respectively were 440 inds.ml⁻¹ and 389 inds.ml⁻¹ (Table 3, column 13 and 14). At taxa level, the mean population density of Rotifera was 208.4 inds.ml⁻¹, Cladocera was 128.8 inds.ml⁻¹, and Copepoda was 102.7 inds.ml⁻¹ in Tank-A. Recorded in Tank-B were Rotifera, 187.8 inds.ml⁻¹, Cladocera, 110.2 inds.ml⁻¹ and Copepoda 91.14 inds.ml⁻¹. The population density distribution

percentage by taxa in Tank-A and Tank-B respectively were Rotifera, 47.37 and 48.26 %, Cladocera, 29.28 and 28.32 %, while Copepoda is – 23.35 and 23.42%.

Result of this research indicated that inorganic fertilizer have more concentrations of nutrients and favoured higher phytoplankton abundance, thus more zooplankton productivity, when compared to organic fertilizer, (Kumar et al. 2014; Boyd, 1982). These findings corroborated with Mosha et al. (2016) that reported a significant higher abundance of natural-fish-food in inorganic fertilizer treated tanks when compared to organic fertilizer treated tanks. Young and Flickinger, (1988) also reported enhanced zooplankton proliferation with application of inorganic fertilizer in zooplankton production for largemouth bass fingerlings production. Zooplankton high abundance was likely due to adequate availability of nutrients in culture media, (Boyd and Massaut, 1999; Guangjun 2013.

As in this experiment, earlier works reported Rotifera to be a common dominant zooplankton group in fertilized ponds, (Kumar, *et al.* 2014). Similarly, cladocerans and copepod nauplii exhibited significant higher population densities in ponds fertilized with inorganic fertilizer than ponds fertilized with organic fertilizer, (Mischke and Zimba, 2004).

Rotifera, Cladocera and Copepoda were relatively more abundant in this investigation than those reported by Rajalakshmi et al. (2012) and Oladele and Omitogun, (2016), but less abundant than the 562 inds.ml⁻¹ reported by Arimoro and Ofojekwu, (2004), between 400 and 1,347 inds.ml⁻¹ reported by Lubzens et al. (1995) and the 1,000 inds.ml⁻¹ reported Kim, (1972), although their report was basically on Rotifera culture. Zooplankton size has been shown to determine their population density, Kim, (1972) reported 100 inds.ml⁻¹ for larger Brachionus calyciflorus and 1,000 inds.ml⁻¹ for tiny Filinia longiseta in similar ponds and culture condition. However the over-all performance of both fertilizers and zooplankton response in this investigation were within the optimum ranges for simulated aquatic environments required for zooplankton mass production.

CONCLUSION

This research experiment report indicates that inorganic fertilizer (Tank-A) resulted to relatively higher zooplankton population abundance when compared to the organic fertilizer (Tank-B). There is significant difference (P < 0.05) between the values of Tank-A against Tank-B. The average population density values of culture Tank-A (440 inds.ml⁻¹) and Tank-B (389 inds.ml⁻¹) are within the optimum range of managed zooplankton culture tank. Notwith-standing cost of inorganic fertilizers, smallholder farmers can procure and use them, being that relatively little quantity is required. Fish breeders are advised to utilize either or both inorganic and organic fertilizers. The zooplankton culture techniques applied in this study has the potential for Nigeria, in that it encourages the utilization of animal manure.

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