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## EDITORIAL COMMENT

All Glory to the Alpha and Omega, the Mighty God of all creation. The Fisheries Society of Nigeria (FISON) 27<sup>th</sup> Annual Conference and Biennial General Meeting tagged "Bayelsa 2012" was a success. All due to selfless contributions of many committed members, participants from far and near, the doggedness of the National and State Executives along with the untiring efforts of the local organising committee under the Chairmanship of Dr.M.E.Allison. It is worthy of note to appreciate the general public and special people who are passionate about the Fisheries profession. We appreciate the Secretary to the Bayelsa State Government Prof. Edmund Allison -Oguru who is a Fisheries biased Agric -Economists. The Vice-Chancellors of Niger Delta University and Federal University, Otueke, The Rector, Federal Polytechnique, Ekowe,The Deans of the Faculty of Agricultural Technology and the Faculty of Law in the Niger Delta University , all these people used their personal and official capacity to support the success of the FISON Conference in Bayelsa State. The Key note address titled "Development of Aquaculture Value Chain: A Veritable Vehicle for Self Sufficiency in Fish production and A Strategic Foreign Exchange Earner" was brilliantly delivered by Dr .O.A. Ayinla. This address dwelt on the Transformation Agenda of the Federal Government emphasizing value chain which is basically getting Agricultural products to consumers in best packaged quality and ensuring it positively impact farmers profit and livelihood.

The papers were presented in 4 sections namely 1. Aquaculture, 2. Capture Fisheries, Gear Technology and Fish Processing Technology, 3. Fish Biology and Pathology and 4. Fisheries Management, Economics, ICT and Marketing. These sections were chaired by experienced eminent Scholars in the field of Fisheries. The various papers presented brought out the enthusiasm to share information by the authors and display of professionalism which has remained the passion towards best practises . Researchers, Farmers and Academia had opportunity to bring out information and current findings for the Development of the fisheries profession. I came to realise the labours of the founding fathers of the society in holding the forth for up coming Fisheries professionals. I pray they reap the fruits of their labours and the Society to continue to grow strong to fulfil its great potentials.

The compilation of the Proceedings was quite an experience with the advancement in information technology-many papers were beautifully written and were good publishing materials, I implore authors to go the extra mile in adhering strictly to the instructions for the paper submissions in future conferences because this will go a long way in aiding early production of the proceedings.

God Bless Fisheries Society of Nigeria  
God Bless our Dear Country NIGERIA

Dr. (Mrs) A.O. Adeyemo.  
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10.	ACUTE TOXICITY OF CHICKEN MANURE TO FINGERLINGS OF <i>Oreochromis niloticus</i> . - Okwor, C. O., P. C. Ofojekwu and M.N.O.Ajima	231
11.	THE COMPARATIVE HISTOLOGICAL STUDIES OF THE TESTIS IN THE WILD AND CULTURED AFRICAN MALE CATFISH SPECIES ( <i>Clarias gariepinus</i> ) Agbebi, O. T., O. M Ademolu and A. O Adebayo	235
12.	EFFECT OF ZINC ON THE DISTRIBUTION OF MAJOR CATIONS AND ANIONS IN THE MUSCLE OF THE FRESH WATER CATFISH <i>CLARIAS GARIEPINUS</i> (BURCHELL, 1822) COLLECTED FROM EBONYI RIVER, SOUTH EAST NIGERIA. OTI, E.E., UDEH, G.N., EKWU, A.O. OKEYE, J.	240
13.	PROXIMATE COMPOSITION AND SOME MINERAL ELEMENTS OF HORSE RADISH ( <i>Moringa oleifera</i> ) SEED MEAL. Mamman, T., Shinkafi, B.A and Mohammad, A.	243
14.	INFLUENCE OF PHYSICO-CHEMICAL PARAMETERS ON THE DISTRIBUTION OF FRESHWATER SNAILS IN RIVER UKE, NASARAWA STATE, NIGERIA Obande, R. A., K.P. Dauda and Adah P. M.	266
15.	ACUTE TOXICITY OF 2, 3-DICHLOROVINYL DIMETHYL PHOSPHATE (SNIPER 1000EC) AND RESPONSE OF <i>Clarias gariepinus</i> BURCHELL UNDER LABORATORY CONDITIONS. -Idi-Ogede, A. M., Ezeri, G. N. O., Omoniyi, I. T. and Akinloye, O.A.	250
16.	EFFECT OF PROCESSING METHODS ON THE PROXIMATE COMPOSITION OF <i>Gmelina arborea</i> SEED. - Ajayi. C. T; R.N. Oladosu-Ajayi; and T.J. Adegoke	254
17.	ACUTE TOXICITY OF INDUSTRIAL EFFLUENTS FROM AGBARA ENVIRONS OF OLOGE LAGOON ON EARLY LIFE STAGES OF AFRICAN CATFISH <i>Clarias gariepinus</i> . - Adeboyejo, O.A., Fagbenro, O.A. and Adeparusi, E.O.	257
18.	HEAVY METALS CONCENTRATIONS IN <i>Chrysichthys nigrodigitatus</i> IN ARAKANGA RESERVOIR, ABEOKUTA, OGUN STATE, NIGERIA.- Adeosun, F.I., Alegbeleye W.O., Abdul, W.O., Agbon, A.O., Akinyemi, A.A., and Odebiyi, O.C.	265
19.	THE EFFECTS OF HEAVY METALS CONCENTRATION IN SOME COMMERCIAL FISH IN OGUN RIVER, OPEJI, OGUN STATE, NIGERIA. - Adeosun, F.I., Alegbeleye W.O., Akinyemi A.A., Abdul, W.O., Agbon, A.O., and Odebiyi, O.C.	268
20.	TRACE METALS IN WATER, FISH AND SEDIMENTS FROM ELECHI CREEK, PORT HARCOURT, and RIVERS STATE, NIGERIA. Vincent-Akpu, I.F. and T. Mmom	271
21.	TOXICITY OF OIL – BASED DRILLING MUD (OBM) ON THE SURVIVAL OF THE FINGERLINGS OF NIGER DELTA MUDSKIPPER, PERIOPHTHALMUS PAPILIO (BOCH AND SCHNEIDER, 1801). Nwakanma, C. and Hart, A.I.	275
22.	MICROBIOLOGICAL QUALITY OF TILAPIA RAISED IN PONDS FERTILIZED WITH RAW AND STERILIZED COWDUNG MANURES -Omojowo F.S. and Omojasola P.F.	278
23.	FOOD AND HABITS OF THE BATRACHOID( <i>Batrachoides liberiensis</i> ) OFF THE QUA IBOE RIVER ESTUARY, NIGERIA- Mfon T. Udo and Iniobong I. Brownson	281
24.	MICROBIOLOGICAL STATUS OF WATER AND FISH SAMPLES FROM JABI LAKE- ABUJA. - Atiribom, R. Y., Mbagwu, I.G., Kolndadacha, O.D. and Muazu,	284
25.	THE ANAESTHETIC EFFECTS OF CLOVE SEED IN SELECTED SPECIES OF GREY MULLETS <i>Lizafalcipinnis</i> and <i>Liza grandisquamis</i> .-Akinrotimi, O.A	287

26.	TROPHIC ECOLOGY OF FISHES: CONSOLIDATING THE BASIC (TRADITIONAL) METHODS.-Saba, A. O. and Fakoya, K. A.	292
27.	HEAVY METAL CONTENT OF SOLE, <i>Solea solea</i> and CROAKER, <i>Pseudotolithus typus</i> FROM LAGOS AND DELTA STATES, NIGERIA - George, F. O. A., R. Ogamune, D. O. Odulate and T. A. Arowolo	295
28.	LENGTH-WEIGHT RELATIONSHIP, CONDITION FACTOR AND REPRODUCTIVE BIOLOGY OF <i>Pseudotolithus senegalensis</i> . IN SELECTED COASTAL WATERS OF SIERRA LEONE. Olapade O.J and Sheku. T	300
29.	ENVIRONMENTAL IMPACT OF CRUDE OIL SPILLAGE AT AGOUBIRI COMMUNITY IN SOUTHERN IJAW LOCAL GOVERNMENT AREA OF BAYELSA STATE Anderson, Emmanuel and Adeyemo, Abiodun Oluseye	304
30.	DETECTION OF LISTERIA MONOCYTOGENES IN FROZEN FISH IN LAGOS, NIGERIA Amusan. E.E.	307
31.	INDICATIVE FISH CATCH OF BRASS RIVER AREA - Otobotekere, A. J. T	311
32.	OCCURRENCE AND DISTRIBUTION OF MACROBENTHIC INVERTEBRATES IN THE LOWER TAYLOR CREEK, BAYELSA STATE -Otobotekere A J T, and Kenigua, V S.	315
33.	MIGRATION IN FISHES: A REVIEW- Obande, R. A., Dambo A., and Adah P. M.	319
34.	HISTOPATHOLOGICAL AND HAEMATOLOGICAL EFFECTS OF ACUTE TOXICITY OF CYPERMETHRIN ON <i>Clarias gariepinus</i> JUVENILES. Asuwaju, F.P ,R.O.Ojutiku, R.J. Kolo O.O Agbelege	322
35.	HAEMATOLOGICAL CHANGES OF <i>Clarias gariepinus</i> JUVENILES FED DIFFERENT DIETARY LIPID.Oshoke J.O. , Olukunle O.A, Ajayi, A.I., Dasuki A and Saulawa L.A.	327 ✓
36.	ESTIMATED UN-IONIZED AMMONIA AT SMALL INCREMENTAL PH VALUE AND TEMPERATURES: PRACTICAL OPTION FOR FISH FARMERS – Ebonwu, B. I.	331
37.	NUTRIENT AND pH STABILITY IN LIQUID MANURE PRODUCTION AND USAGE FOR POND FERTILIZATION Ebonwu, B. I	335
38.	THE EFFECTS OF CRUDE OIL ON THE POPULATION STRUCTURE OF PLANKTON - Sikoki, F.D, Egemba, M.T. and Komi, G.W.	339
39.	GROWTH ENHANCEMENT POTENTIAL OF <i>Mucuna pruriens utilis</i> ON THE NILE TILAPIA <i>Oreochromis niloticus</i> (L.)- Komi, G.W., Sikoki, F.D., Aleleye-Wokoma, I.P. and Ekibebe, D.O.	343
40.	PREVALENCE OF <i>Camallanus cotti</i> IN <i>Poecilia reticulata</i> OBTAINED FROM SOME WASTEWATER DRAINS IN LAGOS STATE. - Akinwale, M.M.A. and Adesola.A.Hassan.	346
41.	FISH SPECIES COMPOSITION AND DIVERSITY IN THE WARRI RIVER, NIGER DELTA NIGERIA. Ogaga Augustine Aghoghovwia	349
42.	PREVALENCE OF Eustrongylides Ignotus IN <i>Poecilia Reticulata</i> OBTAINED FROM SOME WASTEWATER DRAINS OF LAGOS STATE. AKINWALE, M.-M.A and A. A. HASSAN <sup>2</sup>	353

## HAEMATOLOGICAL CHANGES OF *Clarias gariepinus* JUVENILES FED DIFFERENT DIETARY LIPID.

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### ABSTRACT

Twelve weeks feeding trial was conducted to determine the effects of different dietary lipid sources on the hematological changes in *Clarias gariepinus* juveniles. 6 iso-nitrogenous diets were formulated at 45% CP and fed to triplicate groups of 15 juveniles. The feed contained (Palm Seed Oil (PSO), Ugwu Seed (USO), Soya Bean Oil (SBO), Almond Seed Oil (ASO), Mixture Of All the vegetable oil + the fish oil (MOA) and Cod Liver Oil (CLO) which is the control. The oils were added at 5% inclusion level respectively. Fish of mean weight  $22.83 \pm 0.30$ g were fed these experimental diets in triplicate groups. The hematological analyses of fish showed that red blood cell, white blood cell, Erythrocyte sedimentation rate (ESR), Mean cell volume (MCV), Mean cell Haemoglobin (MCH) and packed cell volume were not significantly different ( $p < 0.05$ ), but haemoglobin concentration and Mean cell haemoglobin concentration (MCHC) were significantly different ( $p > 0.05$ ). The present study showed that PSO, USO, ASO, SBO and MOA can effectively replace cod liver oil without compromising the health of African catfish, *Clarias gariepinus*.

**Key words:** Dietary lipid, Juveniles, Haematology, *Clarias gariepinus*

### INTRODUCTION

Global catches from the feed-grade fisheries that provide fish oil (FO) and fish meal for aqua feed formulations have reached their sustainable limits (Pike & Barlow, 2002) and it is likely that within a decade or so there may be insufficient FO to meet the quantities required for current aquaculture growth (Tacon, 2004). Consequently, there has been considerable interest in introducing sustainable alternatives to fish meal and FO that reduce reliance on marine raw materials (Tacon, 2005).

Vegetable oils are viable alternatives as they are readily available and more cost-effective compared to FO. Many studies have reported that vegetable oils can partially or fully replace FO in fish diets without compromising growth performance as long as the essential fatty acid requirements of the fish are met (Turchini, et al., 2009). However, the adverse effect of feeding fish with vegetable oils particularly on haematological parameters is very scanty. Blood is a good indicator in determining the health of an organism (Joshi et al., 2002c), it also as a pathological indicator of the whole body, and hence haematological parameters are important in diagnosing the functional status of an animal exposed to suspected toxicant (Omitoyin, 2006).

This study therefore investigates the changes in haematological parameters of juvenile catfish fed with different dietary lipid.

### MATERIALS AND METHODS

The experimental work was carried out in the research laboratory of the Department of Wildlife and Fisheries Management teaching laboratory, University of Ibadan, Nigeria. A total of eighteen plastic circular experimental tanks of 45 litres (30 cm depth, 36 cm width and 52 cm length) covered with mosquito mesh nylon screen to prevent fish from jumping out and possible predation were used. Each of the six treatments was replicated in triplicates. Juveniles of the African catfish, *Clarias gariepinus* were obtained from a local fish hatchery and transported in oxygen bags to the laboratory. The fish were then acclimatized to laboratory conditions and fed with a commercial fish feed (35% CP) for 14 day. After acclimation, groups of fifteen *Clarias gariepinus* juveniles (mean weight  $22.82 \pm 0.30$ g) were randomly stocked into the eighteen circular plastic tanks containing 30 litres of water each for the growth trials. Experimental tanks were well aerated using air stones and aerator pump (Lawson, 1995) throughout the period of the experiment to maintain relatively uniform physiochemical parameters.

Each of the diets was fed to the fish in triplicate at 5% body weight twice daily (between 8.30am and 9.00am, and 5.30pm and 6.00pm) for 84 days. The weight of each group of fish was taken fortnightly using electronic top loading balance and the feed adjusted accordingly.

The water quality parameters of dissolved oxygen, temperature and pH were monitored on alternate days. Early in the morning (7.00 – 8.00 am) on the days when the water quality parameters were taken. Digital dissolved metre (manufactured by American Marine Inc.) was used to take the Dissolved oxygen, while the water temperature and

pH values of the experimental tanks were measured using Digital/electronic temperature probe and a pH meter respectively (Table 3). Fish meal, mineral/vitamin premix, soya bean meal, yellow maize, salt and binder used in this experiment/study were obtained from a feed miller in Ibadan, Nigeria. The Ugwu (*Telfaira occidentalis*) seeds were bought from Ojoo market and the oil from the seed was extracted using continuous soxhlet extraction technique with hexane. The almond seeds (*Terminalia catappa*) were picked from trees within the premises of University of Ibadan, Nigeria. The seeds were shelled by cracking to remove the kernels inside. The kernels were ground to powder in a hammer mill and the oil from the seed was extracted using the continuous soxhlet extraction technique with hexane. The palm nuts were also bought from Ojoo market and the oil from the seeds extracted locally. The soya bean oil was obtained from the market while cod liver oil, which is the control was obtained from the University of Ibadan pharmacy.

All dry ingredients were milled together with the hammer milling machine to obtain fine particulates. The crude protein content of the diets were kept essentially at the 45% level since this was determined as the protein requirement of juveniles catfish hybrid in a previous experiment (Eyo and Falayi, 1999). Each diet was first mixed dry and later with just enough warm water to obtain homogenous hard-paste (dough) and pelletized out into flat tray through 2mm die disc holes in different lengths.

Using the ingredients, six practical diets containing 45% crude protein, each having different lipid sources was formulated respectively. Diet CLO (control diet) contains cod liver oil, Diets PSO, contains palm oil, Diet ASO contains almond seed oil, Diet MOA contains a mixture of all the vegetable oil + fish oil, Diet USO contains ugwu seed oil and Diet SBO contains soya bean oil. The pellets were sun dried at ambient temperature of 30°C for three days and stored in air tight plastic at 26°C (Table 1).

#### Haematological evaluation

One and a half milliliters (1½ ml) of blood were collected at the beginning of the feeding trial (week 0) and at the end of trial (week 12) from the caudal peduncle of both the test and control fish. The fish from which blood for haematology was collected, were anaesthetized with 150mg/l solution of tricaine methaine sulphate (MS-222, Sigma Chemical co. St. Louis, MO, USA) (Wegner et. al., 1997). Blood samples were taken with 2 ml heparinized syringes and 21swg needles from the caudal vein of the fish from each treatment and put separately in 2ml heparinized tubes and taken to the laboratory for determination of Haematocrit (Hct), Haemoglobin (Hb), Erythrocyte Sedimentation Rate (ESR), White Blood Cell (WBC) and Red Blood Cell (RBC) using the method of Svobodova et al. (1991). The haematological indices of Mean Cell Haemoglobin Concentration (MCHC), Mean Cell Haemoglobin (MCH) and Mean Cell Volume (MCV) were calculated (Dacie and Lewis, 2001, Joshi et al 2002a). Data collected from the experiment were subjected to one way analysis of variance (ANOVA) to test the significance of variations between the means and Least Significance difference (LSD) was used to determine the level of significance ( $p < 0.05$ ) among treatments.

#### RESULTS AND DISCUSSION

Result of analysis of the hematological parameters of *Clarias gariepinus* in this study showed significant difference between the treatment values. PCV value obtained showed that CLO had the highest value. The haemoglobin (Hb) values are much higher than those obtained by Subhadra et al., (2006) for the largemouth bass with diets containing canola oil, chicken oil and menhaden fish oil, which ranged between 3.7-3.9 g dl<sup>-1</sup>. Hb value of 7.20g dl<sup>-1</sup> for the initial (pre-treatment) *Clarias gariepinus* and the mean values of 7.67 ± 1.23 and 7.07 ± 0.81 obtained for fish raised on diet (CLO, PSO) were similar to the mean Hb values of 7.44% obtained by Etim et. al., (1999) for *Chrysichthys nigrodigitatus* showing that the oxygen carrying abilities of the blood of these two catfishes are similar. The difference may be due to the fact that we used different species and the ability to utilize n-6 fatty acids presents in the vegetable oils differs from species to species. The value obtained in the study was within the recommended range value of 4.1 – 10.3 by Blaxhall and Daisley, 1973 for healthy fish. The con of Hb, WBC and RBC in fish fed diet MOA (mixture of all the oil), show clearly that the n-3:n-6 PUFA balance seems critical in the diet of the African catfish.

There were no significant difference in ESR, RBC, WBC ( $p > 0.05$ ) among the treatment. The value of the RBC ranged from  $2.30 \pm 0.68 \times 10^{12/L}$  to  $1.45 \pm 0.22 \times 10^{12/L}$  are similar to those obtained by Osuigwe et al., (2004) for *Clarias gariepinus*. The mean cell haemoglobin concentration (MCHC) differed between treatments and did not follow a clearly defined trend. Fish fed on diet USO showed significantly lower MCHC than fish raised on all the other treatments. Lie et. al., (1989), reported that an increase in MCHC and MCH values reflect a preserving mechanism in rainbow trout activated at reduced water temperatures. There was no temperature variation in this study, hence no increase relative to the initial MCHC and MCH values were observed.



## CONCLUSION AND RECOMMENDATION

In conclusion the present study revealed no inhibition to the formation of skeletal tissues, cell formation, blood formation and flow; hence the utilization of PSO, USO, ASO, SBO and MOA in the diet of *Clarias gariepinus* should be encouraged.

Table 1: Proximate Composition of diets based with different lipid source for the African catfish, *Clarias gariepinus* (% dry matter).

Ingredients	Diet PSO	Diet USO	Diet SBO	Diet ASO	Diet CLO	Diet MOA
Crude Protein (%)	46.87	45.68	46.65	45.92	45.82	46.66
Crude Fat (%)	6.79	5.64	6.12	5.98	6.87	6.48
Crude Fibre (%)	3.55	3.66	3.86	3.78	3.58	3.92
Ash (%)	16.94	16.89	17.23	15.89	17.49	17.86
Moisture (%)	6.82	7.62	8.14	8.21	6.73	7.56
NFE (%)	19.03	21.54	17.00	20.22	20.51	12.52

Table 2: Haematological parameters of African catfish *clarias gariepinus* juveniles fed with the experimental diets.

PARAMETERS	INITIAL	PSO	USO	SBO	ASO	CLO	MOA
PCV (%)	21.00	22.00±2.65 <sup>a</sup>	20.33±3.51 <sup>a</sup>	18.67±1.15 <sup>a</sup>	21.67±4.04 <sup>a</sup>	24.00±3.61 <sup>a</sup>	22.0±2.650 <sup>a</sup>
Hb (gm%)	7.20	7.07±0.81 <sup>b</sup>	3.00±3.51 <sup>c</sup>	5.57±0.38 <sup>a</sup>	6.90±1.31 <sup>d</sup>	7.67±1.24 <sup>e</sup>	7.40±1.06 <sup>e</sup>
RBC ( $\times 10^{12}/L$ )	1.84	1.79±0.46 <sup>d</sup>	1.59±0.12 <sup>d</sup>	1.45±0.22 <sup>d</sup>	1.95±0.56 <sup>d</sup>	2.03±0.68 <sup>d</sup>	1.85±0.52 <sup>d</sup>
WBC ( $\times 10^9/L$ )	16,500	17116.67 <sup>a</sup> ±1733.73	19150.00 <sup>a</sup> ±975.96	15600 <sup>a</sup> ±4850.78	19966.67 <sup>a</sup> ±7823.26	19933.33 <sup>a</sup> ±1527.53	19550.00 <sup>a</sup> ±7590.62
ESR (mm/hr)	0.20	2.00±1.00 <sup>a</sup>	1.33±0.58 <sup>a</sup>	1.33±0.58 <sup>a</sup>	1.67±0.58 <sup>a</sup>	1.67±1.15 <sup>a</sup>	1.67±0.58 <sup>a</sup>
MCV (Fl)	114.13	125.94±16.61 <sup>a</sup>	129.15±29.01 <sup>a</sup>	131.30±24.28 <sup>a</sup>	115.15±29.61 <sup>a</sup>	124.86±30.14 <sup>a</sup>	122.54±19.29 <sup>a</sup>
MCH (Pg)	39.13	40.53±6.03 <sup>a</sup>	36.83±9.53 <sup>a</sup>	39.19±7.59 <sup>a</sup>	36.67±9.52 <sup>a</sup>	39.79±9.23 <sup>a</sup>	41.49±9.72 <sup>a</sup>
MCHC (%)	34.29	32.14±0.79 <sup>a</sup>	28.41±1.51 <sup>abcd</sup>	29.81±0.32 <sup>c</sup>	31.83±0.41 <sup>e</sup>	31.91±0.37 <sup>d</sup>	33.65±3.16 <sup>b</sup>
Platelet	294000.00	137333.33 <sup>a</sup>	216333.33 <sup>a</sup>	131333.33 <sup>a</sup>	159666.67 <sup>a</sup>	246333.33 <sup>a</sup>	127333.33 <sup>a</sup>
	±28095.08	±118643.72	±29871.95	±70500.59	±137012.16	±36501.14	

a,b,c.... Means in the same row having different superscripts are significantly different ( $P < 0.05$ ), while means in the same row having same superscript are not significantly different ( $P > 0.05$ ). Values given in mean ± standard deviation of three replicates.

Table 2: Mean bi-weekly water parameters of the experimental tanks

Treatments	Parameters	Week 2	Week 4	Week 6	Week 8	Week 10	Week 12
PSO	Temp (°C)	26.02±0.05	25.03±0.45	26.03±0.00	25.70±0.00	25.20±0.16	26.00±0.01
	DO (mg/l)	6.60±0.00	6.48±0.01	6.70±0.05	6.58±0.01	6.60±0.03	6.60±0.00
	PH	7.20±0.05	6.88±0.01	7.20±0.02	6.93±0.07	6.90±0.06	7.10±0.03
USO	Temp (°C)	25.80±0.01	25.10±0.08	26.01±0.01	25.10±0.04	25.27±0.12	26.00±0.02
	DO (mg/l)	6.50±0.02	6.56±0.05	6.55±0.02	6.70±0.50	6.61±0.01	6.52±0.01
	PH	6.95±0.00	6.84±0.03	7.10±0.21	6.95±0.05	6.90±0.01	7.10±0.02
SBO	Temp (°C)	25.70±0.01	24.03±0.05	25.90±0.01	25.20±0.01	24.53±0.75	25.70±0.03
	DO (mg/l)	6.46±0.02	6.55±0.02	6.47±0.01	6.70±0.01	6.39±0.04	6.47±0.02
	PH	6.90±0.08	6.84±0.01	7.02±0.01	6.90±0.03	6.78±0.01	6.95±0.05
ASO	Temp (°C)	26.00±0.00	24.27±0.38	26.00±0.02	25.80±0.03	25.17±0.00	26.05±0.01
	DO (mg/l)	6.51±0.02	6.48±0.03	6.51±0.03	6.48±0.02	6.50±0.08	6.55±0.05
	PH	6.95±0.01	6.80±0.02	7.10±0.01	6.95±0.08	6.90±0.02	7.10±0.02
MOA	Temp (°C)	25.60±0.01	24.20±0.16	25.01±0.00	25.27±0.02	24.20±0.05	26.01±0.50
	DO (mg/l)	6.45±0.02	6.12±0.04	6.44±0.01	6.42±0.01	6.04±0.02	6.52±0.04
	PH	6.84±0.01	7.10±0.21	6.90±0.06	6.84±0.01	6.90±0.08	6.88±0.01
CLO	Temp (°C)	25.70±0.01	24.90±0.08	25.80±0.02	24.90±0.00	25.10±0.08	25.80±0.01
	DO (mg/l)	6.48±0.02	6.82±0.01	6.49±0.02	6.82±0.04	6.59±0.02	6.49±0.01
	PH	7.00±0.08	6.87±0.03	7.00±0.03	6.95±0.05	6.89±0.01	7.01±0.01

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