# Effect of fish meal replacement with Blood meal on the growth response and utilization of Hybrid (*Clarias gariepinus* ♀ X *Heterobranchus bidosarlis* ♂) fingerlings

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# 8 Abstract

This study examined the replacement of fishmeal with Bloodmeal (BM) in Practical diets of 9 clariid catfish hybrid "Heteroclarias" (Clarias gariepinus  $\bigcirc$  x Heterobranchus longifilis  $\bigcirc$ ) 10 fingerlings. Five diets containing varying levels were formulated. Diet 1, (10% BM); Diet 2, 11 (15 % BM); Diet 3, (20% BM); Diet 4 (25 % BM) and Diet 5 (30 % BM) as a replacement 12 for fishmeal were fed to three replicate of *Heteroclarias* with an initial weight of 5.9±0.01g. 13 Diet 1, (10%BM) had the best growth rate and feed utilization (p<0.05) as it had the highest 14 value of weight gain (1.60), feed intake (2.53), relative weight gain of 15.66 and Specific 15 Growth Rate of 1.58. There was no significant difference (P<0.05) in the feed conversion 16 ratio across all five treatments. Therefore, bloodmeal can replace fishmeal totally but will be 17 best at 10% replacement level in diets of Heteroclarias without compromising the growth 18 and carcass composition. 19

20 Keywords: catfish diet, *Heteroclarias*, Blood meal and fingerlings

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## 22 Introduction

23 The increased competition between the expanding aquaculture and livestock sectors for a 24 limited supply of fishmeal and fish oil continues to drive the price upwards and the price would reach a level where the use of fishmeal and fish oil may no longer be financially viable 25 26 (FAO of the United Nations, 2006). The development of a more sustainable aquaculture feed production will depend on identifying and establishing alternative feedstuffs to fishmeal 27 (Olukayode and Emmanuel, 2012). Research has shown that most imported feedstuff 28 (fishmeal) can be replaced by locally available feedstuff (Agbebi et al., 2009; Otubusin et al., 29 2009; Gabriel et al., 2007; Okanlawon and Oladipupo, 2010; Faruque et al., 2010). This, 30 therefore, entails the production of fish feed from locally available materials using local 31 32 technology in order to reduce the cost and improve availability of feed to farms.

33 Fish feed is the single most expensive input in intensive fish culture especially for catfish as they require high protein diets (Olevera-Nora, 1996). Catfishes of the family Clariidae 34 35 comprise the most commonly cultured fishes in Nigeria (Adekoya et al., 2006). Of all the ingredients needed for the formulation of the fish diet, fishmeal is considered to be the most 36 37 expensive due to its scarcity, well balanced amino acid and fatty acid, excellent quality, high protein content and also its palatability to fish (Eyo, 1985). A fish farmer would profit more if 38 a less expensive alternate protein source is used to replace the more expensive fish meal 39 without compromising the quality, acceptability and palatability of the feed (Aliu and Esume, 40 2016). Although blood meal was found to be a good substitute of fish meal (Adikwu, 1991) 41 in terms of crude protein requirement and amino acid profile there is need to conduct a 42 feeding trial to determine the performance of fish to this waste. This current study therefore is 43 designed to determine the growth performance and optimum inclusion level of blood meal in 44 the diet of *Heteroclarias* fingerlings. 45

#### 46 Materials and Methods

This study was conducted in the Wet Laboratory of Department of Fisheries and Aquaculture
Management, Faculty of Agriculture, University of Benin, Benin-city Edo state for Seventy

(70) days. One hundred *Heteroclarias* fingerlings (initial mean body weight of 5.9±0.01g) 49 50 were obtained from outdoor fish tanks of the Department of Fisheries, University of Benin, Benin city and were stocked randomly at five (5) fingerlings per aquarium in 40 litres of 51 domestic water of university of Benin in the laboratory. Temperature of water ranged from 52 27-29°C and PH of 6.9-7.6. The fingerlings were fed crumbled 2.0 mm size pellet of 53 54 experimental diets twice daily to satiation between 09.00hrs and 16.00hrs. Feeding was monitored for each unit to ensure that fishes were not underfed or overfed. Experimental 55 56 units were cleaned daily while Weekly weight gain and feed consumption were monitored weekly for 70 days. 57

58 The experimental design consists of five (5) dietary treatments with three (3) replicates each. Diet 1, (10% BM); Diet 2, (15 % BM); Diet 3, (20% BM); Diet 4 (25 % BM) and Diet 5 (30 59 60 % BM). The blood meal was collected from the Slaughter Unit of the University of Benin 61 Farm Project in Benin-City, Edo State. The blood was boiled for about 30minutes to get rid of micro-organisms that may cause certain effects like dropsy (bloated belly), popeye to the 62 63 fish and also to coagulate the blood. It was then dried in the Altona Smoking kiln for 12hours 64 at a temperature of 105°C. The dried blood was then ground finely before it was used in 65 compounding the feed.

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INGREDIENTS	DIET 1	DIET 2	DIET 3	DIET 4	DIET 5
Blood meal (80%CP)	10	15	20	25	30
Fish crumb (50% CP)	30	25	20	15	10
Brewers waste meal (23.8% CP)	8.4	8.4	8.4	8.4	8.4
SBC (44.0% CP)	35	35	35	35	35
Yellow maize (9.6% CP)	5	5	5	5	5
Palm oil	7	7	7	7	7
Bone meal	4	4	4	4	4
Vitamin premix	0.6	0.6	0.6	0.6	0.6

67 Table 1: Gross Composition of the Experimental Diets (%) on as fed basis

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69 Chemical Analysis The various diets, Bloodmeal and the experimental fish (initial and final

70 carcass) were analyzed for their proximate composition which include their moisture content,

71 nitrogen, ether extract, crude fibre and nitrogen-free extract (NFE) according to the

procedures of Association of Official Analytical Chemists (A.O.A.C., 2000). The nitrogen

was converted to protein by multiplying the nitrogen level with a factor of 6.25.

**Growth and Feed Utilization Parameters** Determination of parameters such as Specific Growth Rate (SGR), Feed Conversion Ratio (FCR), Protein Efficiency Ratio (PER) were carried out. Parameters determined and their formulae include:

1. Weight gain = W1 - W0

78 2. Relative Weight Gain (RWG%)= $(W1 - W0) / W0 \times 100$ 

- 79 3. Specific Growth Rate (SGR %)= {(In W1 In W0)/T}  $\times$  100
- 80 W0: mean initial weight (g) W1: mean final weight (g) T: time in 7 days between 81 weightings
- 4. Feed conversion ratio (FCR) = feed intake (g) / weight gain (g)
- 83 5. Protein efficiency ratio (PER) = weight gain (g) / protein intake (g)
- 84 6. Net protein utilization (NPU) =  $\{(BP1 BP0)/CP\} \times 100$

Where; BP0: Initial body protein content (g) BP1: Final body protein content (g) CP: Protein
intake (g) Sampling was carried out weekly by weighing the whole fish in each replicate.
Statistical Analysis: All analyzed data were tested for significant differences using analysis
of variance (ANOVA) test and the means were compared using Genstat 2012 version, all at
5% level of significance.

- 90
- 91 **Result**

# Table 2: Growth Performance and Feed Utilization of Heteroclarias Fingerlings Fed Blood Meal Based Diets.

Parameters	TRT1	TRT2	TRT3	TRT4	TRT5	SEM
	10%BM	15%BM	20%BM	25%BM	30%BM	
Weight Gain(g)	1.60 <sup>a</sup>	0.96 <sup>b</sup>	0.90 <sup>b</sup>	0.36 <sup>c</sup>	0.24 <sup>c</sup>	0.32
Specific Growth Rate	1.58 <sup>a</sup>	1.12 <sup>a</sup>	1.10 <sup>a</sup>	0.14 <sup>b</sup>	0.30 <sup>b</sup>	0.58
(%/day)						
Relative Weight Gain (%)	15.66 <sup>a</sup>	10.19 <sup>b</sup>	10.08 <sup>b</sup>	6.30 <sup>c</sup>	5.72 <sup>c</sup>	3.82
Protein Efficiency Ratio	83.23 <sup>b</sup>	74.41 <sup>ab</sup>	79.62 <sup>ab</sup>	36.81 <sup>a</sup>	21.30 <sup>a</sup>	3.86
Feed Intake(g)	2.53 <sup>a</sup>	2.14 <sup>a</sup>	2.07 <sup>a</sup>	1.22 <sup>b</sup>	1.02 <sup>b</sup>	0.33
Feed Conversion Ratio	1.58	2.23	2.30	3.39	4.25 <sup>NS</sup>	2.68

94 *N/B: Mean Values with the same superscript on the same row are not significantly different, (P> 0.05) SEM* 95 = standard error of mean. *NS* = *No significant difference* 

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97 Result from the growth performance and feed utilization of Heteroclarias (Table 2) showed 98 that Weight gained by Heteroclarias fingerlings after ten weeks of culture was significantly 99 higher (P<0.05) in Diet 1 (1.60) while Diet 5 (0.24) recorded the least value.

The Specific growth rate for Diet 1 (1.58), Diet 2 (1.12) and Diet 3 (1.10) were not significantly different (P>0.05) from each other while Diet 5 (0.30) and Diet 4 (0.14) were also not significantly different (P>0.05).

The Relative Weight in Diet 2 (10.19) and Diet 3 (10.08) showed no significant difference
(P>0.05) from each other as well as Diet 4 (6.30) and Diet 5 (5.72) showed no significant
differences (P>0.05). The Relative Weight gain for Diet 1 (15.66) was significantly superior

- 106 (P < 0.05) than all the other treatments.
- 107 The Protein Efficiency Ratio for Diet 2 (74.41) and Diet 3 (79.62) were not significantly 108 different (p>0.05) from each other. The Protein Efficiency Ratio for Diet 1 (83.23) was 109 significantly superior (P<0.05) than all the other treatments.

110 Feed intake in Diet 1, 2 and 3 were not significantly different (P > 0.05) from each other but

- they were significantly different from Diet 4 and 5 meaning that feed were consumed at
- different levels within each treatment. Diet 1 had the highest feed intake value of 2.53g while
- the lowest feed intake value was recorded in Diet 5 (1.02g).

Feed conversion ratio showed no significant difference (P > 0.05) between all treatments as this was similar with Diet 5 (4.25) being the highest and Diet 1 (1.58) being the lowest.

# 116117 Discussion

The weight gain was significantly higher (p < 0.05) for fish fed 10% blood meal. As the level 118 119 of blood meal inclusion increased, the response of the fish to the diet became poor. This 120 finding indicate that fishmeal in the diet of *Heteroclarias* can only be efficiently replaced with 10% blood meal and this observation agree with the result of Otubusin (2001) who 121 122 reported that the replacement of fish meal with 10% blood meal in pelleted feeds was 123 adequate for *Oreochromis niloticus* production in floating bamboo net-cages. However, this 124 finding is in contrast to that of Agbebi et al. (2009) who reported that a 25% blood meal 125 substitution of fishmeal in diets gave the best growth performance.

126 The specific growth rate and feed conversion ratio was significantly higher in Diet 1; this 127 result is different from that indicated by Adejoke (2012) on the use of bovine blood and rumen digest in catfish diet to replace fish meal at 0%, 25%, 50%, 75 and 100% where he 128 reported that the best growth performance was recorded in fish with inclusion level of 25% 129 bovine blood and ruminant digest meal. The results obtained from this study showed that fish 130 meal can be replaced partially with blood meal at 10% inclusion level which differs from the 131 132 report of Agbebi et al. (2009) who stated that fish meal can be replaced completely by blood 133 meal at 100% with no adverse effects on the growth, survival and feed conversion ratio of 134 Clarias gariepinus juveniles.

135 The relative weight gain varied with the different inclusion level of bloodmeal, it was highest in treatment 1 (15.66%) and decreased variably as the inclusion level of bloodmeal increased. 136 137 This variation in growth rate that was highest in diet 1 can be attributed to the use of high 138 level of fishmeal as the major animal protein source. Fish meal is known to have balanced amino acid profile, high digestibility and palatability which promote good growth of fish 139 (Hardy and Tacon, 2002). The poor performance of the high blood meal based diets on the 140 141 fishes agrees with the study carried out by Otubusin (1987) in which the feed containing the highest amount of bloodmeal gave the poorest performance in terms of growth and feed 142 conversion ratio. This could be due to the imbalance nature of the essential amino acid 143 144 composition in the bloodmeal which was translated into the diet (Fasakin, et al., 2005; Kirimi 145 et al., 2016).

146 In general, performance of the fish reduced as the level of fishmeal inclusion was reduced. The replacement of the fishmeal with bloodmeal not only changed the nutritional profile of 147 148 the diet (generally reduced methionine levels) but it also affected the palatability. Poor 149 performance was attributed to shifts in palatability as fish meal was replaced with other protein sources (Davis et al., 1995; Meilahn et al., 1996). Palatability is defined as the degree 150 of acceptability of feed materials by a particular animal. Based on visual observations of the 151 152 fish in this experiment as well as subjective ranking of the quantity of feed remaining in the tanks after feeding, it was clear that palatability was reduced as fish meal was reduced. The 153 154 acceptability of the bloodmeal based diets was very low and this also resulted to low growth 155 rate.

The lower the FCR of a feed, the higher the efficiency of the feed and vice versa. The lowest FCR in this study was recorded from the fish fed 10% bloodmeal. This is in accordance with MacDonald *et al.* (1994) who reported an increase in feed conversion ratio of fish as the inclusion level of LabLab bean meal increased across the treatment.

161 Conclusion

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The results of this present study showed that fish fed on diet with high inclusion level of fishmeal performed better in terms of growth than those fed on feed with high inclusion level of bloodmeal. However, although blood meal cannot compete with fishmeal in terms of growth performance, the economics of using it to replace fish meal is positive in the long run.

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