## **Research Article**



# Growth performance and body composition of the African catfish (*Clarias gariepinus*) juveniles fed diets supplemented with *Afrostyrax lepidophyllus* fruit powder

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#### Abstract

To minimize the dependence of pisciculture on imported feed and improve the production of fish through the valorization of local feeds, this study was conducted to evaluate the effects of dietary inclusion of Afrostyrax lepidophyllus fruit powder on the growth and chemical composition of *Clarias gariepinus* juveniles. The juveniles weighing 13±3.1 g were divided into four treatments in triplicates and fed with diets containing 0 (T<sub>0</sub> or control), 2 (T<sub>1</sub>), 4 (T<sub>2</sub>), and 6 (T<sub>3</sub>) g/kg A. lepidophyllus fruit powder for 63 days. The feeding rate was 5% of their body weight. The results showed that adding different levels of A. lepidophyllus fruit powder to the diets improved growth, feed utilization, and carcass quality attributes. The African catfish fed with T<sub>3</sub> diet recorded higher final weight (43.44±0.23 g), weight gain (WG, 31.43±0.41 g), feed conversion ratio (FCR, 0.91±0.05), and protein efficiency ratio (PER, 2.67±0.14) than those recorded in the control group (40.20 g final weight, 27.20±0.52 g WG, 1.1±0.05 FCR, and 2.24±0.10 PER). The same trend was observed in the body macro-nutrient retention (32.33±0.58 % crude protein, 25.33±0.58 % lipid, and 23.33±0.58 % ash) compared to the control group (20.67±0.58 % crude protein, 16.00±0.00 % lipid, and 17.33±0.58 % ash). Finally, this study clearly showed that dietary inclusion of A. lepidophyllus fruit powder at 6 g/kg improves growth performance, feed nutrient utilization, and body composition in juvenile C. gariepinus.

**Keywords**: *Afrostyrax lepidophyllus*, Carcass quality, *Clarias gariepinus*, Zootechnical performance

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## Introduction

The amount of wild-caught fish has not increased since 1980; aquaculture has mainly contributed to the supply of fish for human consumption in the world (FAO, 2016). In Africa, fish production is hampered by the unavailability of aqua-feed which is mostly imported and very expensive (Atangana et al., 2019). However, in Africa and particularly in Cameroon, there is a wide range of products and agricultural by-products rich in nutrients and the addition of some feed additives or dietary supplements can keep fish healthy and enhance growth rate (Sayed et al., 2011; Adegbesan et al., 2019). Research shows that herbs and spices (phytobiotics) are used as supplements or feed additives in animal diets (Muneendra et al., 2014; Mohsen et al., 2018).

Phytobiotics contain active secondary metabolites belonging to the classes of isoprene derivates, flavonoids, and phenols compounds which have been suggested to act as antioxidants (Frankič et al., 2009; Sumczynski et al., 2015). Beneficial effects of phytobiotics in farm animals result from feed intake improvement and secretion of digestive enzymes, immune stimulation, antibacterial, anthelmintic, antiviral, antiinflammatory, and antioxidant properties (Muneendra et al., 2014). Previous studies revealed that ginger (Zingiber officinale), garlic (Allium sativum), and scent leaf (Ocimum gratissimum) contain active substances such as alkaloids, tannins, flavonoids, saponins, and phenols compounds (Muhammad et al., 2009; Abdou et al., 2010). Those compounds are known to improve animal performances through their anti-oxidative and anti-microbial actions and improve feed palatability (Muneendra *et al.*, 2014). An example of a plant possessing secondary metabolites is *A. lepidophyllus* fruit (Moukette *et al.*, 2015; Namkona *et al.*, 2017; Sokamte *et al.*, 2018).

A. lepidophyllus is a plant of the Huaceae family and is commonly found in Equatorial and Tropical Africa (Cronquist, 1981; Moukette et al., 2015; Namkona et al., 2017). This plant is used in Congo as an antiseptic and in traditional medicine in treating gastroenteric diseases (Bouquet, 1969). In Cameroon, the seeds of this plant are traditionally used as a spice. Moreover, pharmacological studies showed that the extracts of A. lepidophyllus seed possess very interesting properties. Ngono (1999) demonstrated the antifungal activity and identified Afrostyraxthioside A. Afrostyraxthioside B. and Afrostyraxthioside C. Agbor et al. (2005) and Boufack et al. (2021) showed that the seed extracts possess antioxidant properties. In addition, Toumnou et al. (2012) reported the insecticidal activity of the extract from the seeds of A. lepidophyllus. Qualitative analysis of A. *lepidophyllus* fruit showed the presence pholyphenols, flavonoids, of total tannins, and anthocyanins (Namkona et al., 2017). These compounds possess antimicrobial significant activity (Sokamte et al., 2018) and antioxidant properties (Oben et al., 2010; Fogang et al., 2014; Sokamte et al., 2018) with

# many other attributes including digestive enzyme stimulation, lipid metabolism, and microbial load modulation (Muneendra *et al.*, 2014).

In the view of the chemical composition of A. lepidophyllus fruit, its ability to balance the gut microbial load stimulate digestive enzymes and (Frankič et al., 2009; Oben et al., 2010), this phyto-additive can positively affect performance the growth of  $C_{-}$ gariepinus. This study aims to determine the effects of A. lepidophyllus fruit powder on growth performance, feed utilization, and body composition of the African catfish (*Clarias gariepinus*) juveniles.

#### Materials and methods

#### Site of study

This study was conducted from April to May 2021 in the research and applied farm in fishery and aquaculture of Kanhé-Moyo (FRAPAIK) in Baham, the western region of Cameroon. This farm is located at 5°19′ North and 10°24′ East and an altitude of 1728 m above sea level. The annual average temperature is 23°C. Rainfall ranges between 1500-2000 mm per annum, over 9 months of the rainy season (March to November).

#### Experimental design

360 *C. gariepinus* juveniles were distributed following a completely randomized design to four treatments of 90 fish each, the average weight was  $13\pm3.10g$ . Each experimental treatment had three (03) experimental units (hapas) of 30 juveniles each placed in a non-fertilized pond of 80 m<sup>2</sup>.

Origin of A. lepidophyllus

*A. lepidophyllus* fruit were bought from a local market then ground, sieved and the powder was incorporated at different levels in locally produced feed.

#### Animal material

Three hundred and sixty C. gariepinus (360) juveniles, were purchased from a renowned fish farm in Douala. These juveniles were acclimatized for two weeks in 03 hapas of  $1 \text{ m}^3$  each, they were fed during the first three days with a commercial imported feed (Gouessant, France; 2 mm) with 46% crude protein, 10 % crude fat, 2.83 % crude fiber, 8.1 % crude ash, 1 % calcium, 1.4% phosphorus, and 0.4 sodium as declared by the company. From the fourth day, 25% of their ration was substituted with local feed without A. lepidophyllus. From the seventh to the tenth day respectively 50 and 75 % of the imported feed was substituted. On the thirteenth and fourteenth days, the animals were entirely fed with local feed (100%). At the end of this acclimatization period, ten С. gariepinus juveniles were randomly sampled for the determination of the carcass quality characteristics.

#### Housing and equipment

After acclimatization, juveniles were randomly transferred to 12 experimental hapas of 0.64 m<sup>2</sup>. The different feed rations were distributed manually at a frequency of three times per day (6:00 a.m., 12:00, and 5:00 p.m.); at a rate of 5% of the fish biomass. In order to monitor growth and adjust the quantities of feed distributed, using a landing net, control fishing was done after 21 days and during cool hours of the day (6:00 a.m.). The different individual weights and lengths were respectively measured using a 1g precision electronic scale balance and an ichthyometer. A 50cm long Secchi disc, JBL Test Kits (JBL GmbH & Co. KG–Dieselstraβe 3 -67141 Neuhofen – Germany).

Maximum-minimum thermometer and pH meter were respectively used daily to measure water transparency, dissolved oxygen, temperature, and pH. The values of the physico-chemical parameters of water recorded during the experiment are resumed in Table 1.

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|------------------------|----------------------------------------|------------------------|------------------------|
| Table 1: water quality | parameters (Mean ± SD)                 | ) during the experimen | ital period (65 days). |

| Experimental period (days) |                                              |                                                                                                                                                                       |                                                        |
|----------------------------|----------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------|--------------------------------------------------------|
| 0-21                       | 21-42                                        | 42-63                                                                                                                                                                 | Optimum values                                         |
| 22.55±1.41                 | 20.63±1.60                                   | 22.2±1.66                                                                                                                                                             | 20-30                                                  |
| $7.03\pm0.08$              | 6.81±0.49                                    | 7.07±0,73                                                                                                                                                             | 6.5-8                                                  |
| $5.85 \pm 0.02$            | $7,89\pm0,55$                                | 7.63±0,59                                                                                                                                                             | 7.5-9.1                                                |
| 30±0                       | 23±2.64                                      | 25±3                                                                                                                                                                  | <50                                                    |
| 45.23±6.62                 | 43.04±7.20                                   | 42.17±7.04                                                                                                                                                            | <60                                                    |
|                            | 22.55±1.41<br>7.03±0.08<br>5.85±0.02<br>30±0 | 0-21         21-42           22.55±1.41         20.63±1.60           7.03±0.08         6.81±0.49           5.85±0.02         7,89±0,55           30±0         23±2.64 | $\begin{array}{c c c c c c c c c c c c c c c c c c c $ |

Temperature (T°C); Hydrogen potential pH; dissolved oxygen (D.O); nitrate (NO<sub>3</sub><sup>-</sup>)

#### Experimental diet

In addition to the locally prepared diet  $(T_0 \text{ or control})$ , three isoprotein, isolipidic and isoenergetic diets including  $T_1$ ,  $T_2$ , and  $T_3$  were made by incorporating 2, 4, and 6g/kg *A*. *lepidophyllus* fruit powder, respectively. The basal diet  $(T_0)$  ingredients and proximate composition are summarized in Table 2.

### Preparation of diets

The raw ingredient was finely ground in a mill. *A. lepidophyllus* previously crushed was incorporated in the mixtures while respecting the different proportions specified in Table 2. In each treatment, cold water (20% per treatment) was mixed, then stirred delicately before extrusion in a DGP60-C brand extruding machine, with a capacity of 120-150kg/hour, at a 105°C temperature equipped with a die of 2.5mm. The extrudates were sun-dried for 20 hours then placed in plastic bags, labeled, and kept away from humidity until use.

| Table 2: Composition of the basal diet. |                      |  |  |  |
|-----------------------------------------|----------------------|--|--|--|
| Ingredients                             | % in dry matter (DM) |  |  |  |
| Fishmeal                                | 27                   |  |  |  |
| Soybean meal                            | 15                   |  |  |  |
| Peanut meal                             | 20                   |  |  |  |
| Cotton meal                             | 8                    |  |  |  |
| Wheat bran                              | 8                    |  |  |  |
| Maize meal                              | 16                   |  |  |  |
| *Premix 5%                              | 5                    |  |  |  |
| Palm oil                                | 1                    |  |  |  |
| Total                                   | 100                  |  |  |  |
| Chemical composit                       | ion                  |  |  |  |
| Protein (% DM)                          | 41.136±3.68          |  |  |  |
| Energy (kcal/kg<br>DM)                  | 2759.24±6.04         |  |  |  |
| Lipid (% DM)                            | 11.18±0.5            |  |  |  |
| Ash (% DM)                              | 11.101±0.95          |  |  |  |
| Moisture (% DM)                         | $5.5 \pm 0.57$       |  |  |  |
| Fiber (% DM)                            | 6.31±0.63            |  |  |  |
| Dry matter (%)                          | 94.5±0.57            |  |  |  |

\*Premix 5%: Crude protein =40%; Lysine =3.30; Methionine= 2.40; Calcium= 8; Phosphorus= 2.05; Metabolized Energy = 2078 kcal/kg. The formulated diet samples (10g) was analyzed following the procedures of AOAC (1990). Moisture content was determined by drying the sample in an oven at 105°C overnight. Crude protein was analyzed in a KJELTEC SYSTEM 1002 Distilling Unit made in Belgium following the Kjeldahl method while lipid content was determined by extraction with hexane using the Soxhlet method. The ash content was determined by combustion in a muffle furnace at 500°C for 6 h.

# Growth, survival rate, and feed utilization parameters

Growth performance, survival rate, feed utilization, and nutrient retention were assessed per treatment by determining the weight gain (WG), specific growth rate (SGR), survival rate (SR), condition factor (K), feed intake (FI), feed conversion ratio (FCR), feed efficiency ratio (FER), protein efficiency ratio (PER) and nutrient retention (NR). The calculations were done using the following formulae:

WG (g) = Wf – Wi  
SGR (%/day) = 
$$\frac{\ln Wf - \ln Wi}{T} \times 100$$
  
Where, Wf = final weight; Wi = initial  
weight; T = experimental period (days);  
SR (%) =  $\frac{final number of fish}{initial number of fish} \times 100$   
K =  $\frac{Weight}{Length^3} \times 100$   
FI (g/fish)  
=  $\frac{total dry feed distributed}{number of fish}$   
FCR =  $\frac{feed intake}{Fish weight gained}$   
FER =  $\frac{Fish weight gained}{feed intake}$   
PER =  $\frac{Fish weight gained}{Protein consumed}$ 

 $Protein consumed = \frac{Total feed consumed x Crude protein in feed}{100}$  $NR (\%) = \frac{Final carcass composition - Initial carcass composition}{Amount of nutrient fed} \times 100$ 

#### Statistical analysis

Results on growth performance, survival rate, feed utilization, and nutrient retention obtained from each replicate were used for statistical analysis. Data collected were submitted to a one-way Analysis of Variance test by the General Linear Model's procedure of Statistical Package for Social Science (SPSS 20.0) software. Where there was a significant difference between treatments, their means were separated using Duncan's multiple range test and probability values less than 0.05 were considered significant.

#### Results

#### Growth performance

Independent of the incorporation level of *A. lepidophyllus* fruit powder in different levels, Figure 1 shows an inflection in the body weight and weight gain curves on the 42<sup>nd</sup> day of the experiment. After this point, the curves have a linear and increasing evolution until the end of the experiment. Moreover, the evolution of

weight and weight gain of C. gariepinus juveniles is proportional to the increasing incorporation level of A. lepidophyllus powder in the diets. Thus, after the  $21^{st}$  day, the curve of C. gariepinus juveniles fed diet supplemented with 6 g/kg Α. lepidophyllus (T<sub>3</sub>) remained above all the other curves. At the end of the trial, the body weight  $(44.43\pm0.23 \text{ g})$  was significantly higher (p < 0.05) by 9.52%,

4.82%, and 2.99% compared to those fed respectively  $T_0$  (40.20±0.30 g),  $T_1$ (42.29±0.14 g) and  $T_2$  (43.10±0.2 g). On the other hand, weight gain (31.43±0.41 g) was significantly higher (*p*<0.05) by 13.46%, 6.78%, and 4.23% compared to those fed with  $T_0$  (27.20±0.52 g),  $T_1$ (29.30±0.24 g), and  $T_2$  (30.10±0.34 g), respectively.

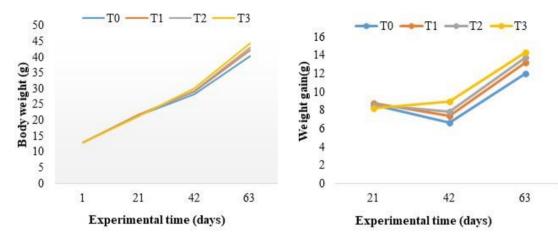


Figure 1: Body weight and weight gain of *Clarias gariepinus* juvenile fed different levels of *Afrostyrax lepidophyllus*. T<sub>0</sub>= control ration, T<sub>1</sub>= T<sub>0</sub>+2g/kg *A. lepidophyllus*, T<sub>2</sub>= T<sub>0</sub>+4 g/kg *A. lepidophyllus*, T<sub>3</sub>= T<sub>0</sub>+6 g/kg *A. lepidophyllus*.

The relationship between fish growth and dietary inclusion level of *A*. *lepidophyllus* powder is illustrated by the linear regression curve (Figure 2) below. *C. gariepinus* juvenile growth rate increased with increasing inclusion level of *A. lepidophyllus* in the diet ( $R^2$ =0.9671).

The specific growth rate of C. gariepinus juveniles fed with different experimental diets for 63 days is illustrated in Figure 3. It is noted that irrespective of the treatments, the growth was faster during the first week of feeding and gradually decreased with the increasing experimental period. At the end of the experimental period, fish fed 6 g/kg A. *lepidophyllus* powder (T<sub>3</sub>) had a specific growth rate of  $1.95\pm0.06\%$ g/d, significantly higher (p<0.05) compared to the other treatments T<sub>0</sub> ( $1.79\pm0.02\%$  g/d), T<sub>1</sub> ( $1.87 \pm 0.009\%$ g/d), T<sub>2</sub> ( $1.90 \pm 0.01\%$  g/d).

# Survival Rate and Feed Nutrient Utilization

Table 3 resumes the survival rate and feed nutrient utilization of *C. gariepinus* juveniles at the end of the feeding period. It is noticed that, though no

significant difference was observed amongst treatments. Mortalities occurred in two treatments  $T_1$  and  $T_2$ with a survival rate of 93.33  $\pm$  0.00%. Determination of the feed nutrients utilization parameters (feed conversion ratio, feed efficiency ratio, and protein efficiency ratio) indicated that the fish fed diets supplemented with *A*. *lepidophyllus* had better feed utilization characteristics (p<0.05) compared to the control diet. On the other hand, we notice a significant increase in the value of condition factor K concerning the dietary inclusion level of *A*. *lepidophyllus*.

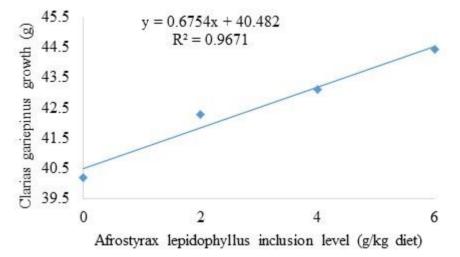


Figure 2: Regression curve between *Clarias gariepinus* growth and dietary inclusion levels of *Afrostyrax lepidophyllus* fruit powder.

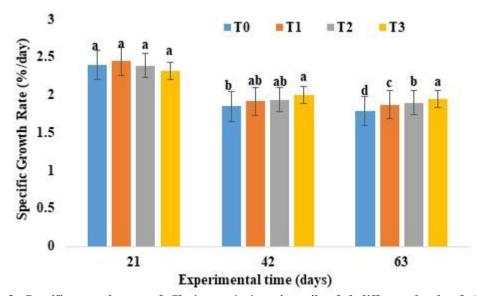


Figure 3: Specific growth rate of *Clarias gariepinus* juveniles fed different levels of *Afrostyrax lepidophyllus* fruit powder. T<sub>0</sub>=control ration, T<sub>1</sub>=T<sub>0</sub>+2g/kg *A. lepidophyllus*, T<sub>2</sub>=T<sub>0</sub>+4 g/kg *A. lepidophyllus*, T<sub>3</sub>=T<sub>0</sub>+ 6 g/kg *A. lepidophyllus*. Means on the same rearing period carrying different superscripts were significantly different at p<0.05.

Chemical (nutrient) Composition and Nutrient Retention of C. gariepinus Nutrient composition of C. gariepinus juveniles produced at the end of the experimental period (63 days) is recorded in Table 4. As compared to fish nutrient composition at the beginning of the study, it is noted that apart from the moisture content that dropped significantly (p<0.05) in the juveniles fed diets supplemented with Α. lepidophyllus powder, other macroelements like the ash, protein, and lipid significantly (p<0.05) levels were enhanced by increasing the incorporation levels of A. lepidophyllus powder in the diets. With the exception of the energy level that was significantly (p < 0.05) higher in the fish fed T<sub>2</sub> diet.

 Table 3: Survival rate and feed utilization of Clarias gariepinus fed diets supplemented with Afrostyrax lepidophyllus fruit powder at the end of the feeding period.

| <u>v                                </u> | Treatments                                  |                        |                               |                           |                  |  |
|------------------------------------------|---------------------------------------------|------------------------|-------------------------------|---------------------------|------------------|--|
| Parameters                               | <b>T</b> <sub>0</sub> <b>T</b> <sub>1</sub> |                        | $T_2$                         | <b>T</b> <sub>3</sub>     | <i>p</i> - value |  |
| Ni                                       | 90                                          | 90                     | 90                            | 90                        | -                |  |
| Nf                                       | 90                                          | 88                     | 88                            | 90                        | -                |  |
| SR                                       | $100.00\pm0.00$                             | $93.333 \pm 0.00$      | $93.33 \pm 0.00$              | $100.00\pm0.00$           | -                |  |
| FI (g/fish)                              | $39\pm0.00$                                 | $38.409 \pm 0.00$      | $38.116\pm0.00$               | $39\pm0.00$               | -                |  |
| Wi (g)                                   | $13,02 \pm 0.01^{a}$                        | $13.01\pm0.02^{a}$     | $13.02\pm0.01^{a}$            | $13.01\pm0.1^{\rm a}$     | 0.977            |  |
| Wf (g)                                   | $40.20 \pm 0.30^{d}$                        | $42.29\pm0.14^{\rm c}$ | $43.10\pm0.2^{b}$             | $44.43\pm0.23^{\rm a}$    | 0.000            |  |
| WG (g)                                   | $27.20 \pm 0.52^{\ d}$                      | $29.29 \pm 0.24^{c}$   | $30.10 \pm 0.34$ <sup>b</sup> | $31.43\pm0.41^{a}$        | 0.000            |  |
| Lf (cm)                                  | $17.38\pm0.02^{\text{d}}$                   | $17.59\pm0.05^{\rm c}$ | $17.94\pm0.02^{a}$            | $17.72\pm0.02^{\text{b}}$ | 0.000            |  |
| Κ                                        | $0.76\pm0.01^{bc}$                          | $0.78\pm0.01^{ab}$     | $0.75\pm0.004^{\rm c}$        | $0.80\pm0.002^{a}$        | 0.006            |  |
| FCR                                      | $1.1\pm0.05^{\rm a}$                        | $0.94\pm0.02^{b}$      | $0.90\pm0.05^{\text{b}}$      | $0.91\pm0.05^{b}$         | 0.004            |  |
| FER                                      | $0.92 \pm 0.04^{b}$                         | $1.06 \pm 0.03^{a}$    | $1.11\pm0.06^{a}$             | $1.11\pm0.06^{a}$         | 0.007            |  |
| PER                                      | $2.24 \pm 0.10^{b}$                         | $2.59{\pm}0.07^{a}$    | 2.69±0.15 <sup>a</sup>        | 2.67±0.14 <sup>a</sup>    | 0.007            |  |

 $T_0$ = control ration,  $T_1=T_0+2g/kg$  *A. lepidophyllus*,  $T_2=T_0+4$  g/kg *A. lepidophyllus*,  $T_3=T_0+6$  g/kg *A. lepidophyllus*.

Values are mean±standard deviation of three replicates of 30 fish each. Mean within the row with different superscripts are significantly different from other at p<0.05. Ni, initial number of fish; Nf, final number of fish; FI, feed intake; Lf, final length of fish; K, condition factor; FCR, feed conversion ratio; FER, feed efficiency ratio; PER, protein efficiency ratio.

 Table 4: Proximate composition (% or kJ/g WW) of carcass and nutrient retention in Clarias gariepinus juveniles fed different inclusion levels of Afrostyrax lepidophyllus fruit powder.

| Parameters      | Initial                         | T <sub>0</sub>       | $T_1$                         | $T_2$                   | T3                      | <i>p</i> - value |
|-----------------|---------------------------------|----------------------|-------------------------------|-------------------------|-------------------------|------------------|
| 1 al alletel S  |                                 | Chem                 | ical compositio               | on (% or KJ/g V         | VW)                     |                  |
| Ash             | $2.47 \pm 0.38^{d}$             | 2.66±0.34°           | 2.76±0.24 <sup>b</sup>        | 2.76±0.26 <sup>b</sup>  | 2.99±0.84ª              | 0.000            |
| Crude protein   | 11.34±0.22 <sup>e</sup>         | $11.96 \pm 1.09^{d}$ | 14.32±0.40°                   | 14.72±0.18 <sup>b</sup> | $15.08 \pm 1.05^{a}$    | 0.000            |
| Lipid           | 2.36±0.28 <sup>e</sup>          | $2.50\pm0.32^{d}$    | 2.55±0.45°                    | 3.10±0.26 <sup>b</sup>  | $3.17 \pm 0.34^{a}$     | 0.000            |
| Energy          | 7.31±0.58 <sup>e</sup>          | $7.43 \pm 0.40^{d}$  | 8.84±0.35°                    | $9.22 \pm 0.46^{a}$     | 9.16±0.22 <sup>b</sup>  | 0.000            |
| Moisture        | 81±0.01 <sup>a</sup>            | 81±0.01 <sup>a</sup> | 77±0.01 <sup>b</sup>          | 77±0.01 <sup>b</sup>    | 77±0.01 <sup>b</sup>    | 0.000            |
|                 | Nutrient retention (% dry feed) |                      |                               |                         |                         |                  |
| Ash             |                                 | 17.33±0.58°          | 19.67±0.58 <sup>b</sup>       | $20.00 \pm 0.00^{b}$    | 23.33±0.58 <sup>a</sup> | 0.000            |
| Crude protein   |                                 | $20.67{\pm}0.58^{d}$ | $28.67 \pm 0.58^{\circ}$      | $30.33 \pm 0.58^{b}$    | 32.33±0.58 <sup>a</sup> | 0.000            |
| Lipid           |                                 | $16.00 \pm 0.00^{d}$ | 18.00±0.00°                   | $23.67 \pm 0.58^{b}$    | 25.33±0.58ª             | 0.000            |
| Energy          |                                 | $19.00 \pm 0.00^{d}$ | 26.00±0.00°                   | $28.00 \pm 0.00^{b}$    | $29.00 \pm 0.00^{a}$    | 0.000            |
| Dry matter      |                                 | 14.00±0.00°          | $20.00 \pm 0.00^{b}$          | 20.33±0.58 <sup>b</sup> | $21.00\pm0.00^{a}$      | 0.000            |
| T = control rat | ion $T = T + 2a$                | leg A lanidon        | $h_{\rm M} H_{\rm M}$ T – T – | 1 alea A lanida         | nhullus T2-T            | 6 a/2 a 1        |

 $T_0$ = control ration,  $T_1$ =  $T_0$ +2g/kg *A. lepidophyllus*,  $T_2$ =  $T_0$ +4 g/kg *A. lepidophyllus*, T3=  $T_0$ + 6 g/kg *A. lepidophyllus*; WW, wet weight.

Mean within the same row with different superscripts are significantly different at p < 0.05. p= probability

Results on nutrient retention of C. gariepinus carcass (Table 4) after 63 days of feeding revealed that incorporating A. lepidophyllus powder in fish diet affects protein, lipid, ash, energy, and dry matter contents. The fish fed  $T_3$  diet had protein (32.33±0.58%), (25.33±0.58%), lipid ash  $(23.33\pm0.58\%)$ , energy  $(29.00\pm0.00\%)$ , and dry matter (21.00±0.00%) contents retention significantly higher compared to those fed the other experimental diets. Figure 4 illustrates the linear regression between nutrient retention in the body of fish and the dietary inclusion level of *A*. *lepidophyllus* powder. Regardless of the nutrient, there is a relationship between nutrient retention and dietary inclusion level of *A*. *lepidophyllus* powder. This is illustrated by the regression coefficient which indicates the relationship between lipid retention ( $R^2$ =0.95) and the inclusion level of *A*. *lepidophyllus* powder followed by ash retention ( $R^2$ =0.91) then by protein retention ( $R^2$ =0.86) and finally energy retention ( $R^2$ =0.84).

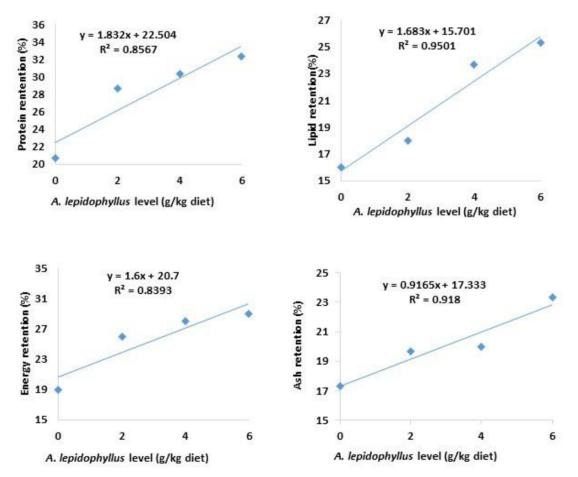


Figure 4: Regression curves between nutrient retention of *Clarias gariepinus* juveniles and dietary inclusion levels of *Afrostyrax lepidophyllus* fruit powder.

#### Discussion

Water quality parameters had a great influence on fish survival rate as well as

on their growth. Moreover, the recorded physicochemical parameters such as temperature, dissolved oxygen, pH, and nitrate  $(NO_3)$ were within the recommended range for freshwater fish breeding. According to the above observations, the mortalities observed in the fish fed with 2 g/kg ( $T_1$ ) and 4 g/kg (T<sub>2</sub>) dietary inclusion of A. lepidophyllus fruit powder could be probably attributed to stress-induced during the fish rearing. The application of phytochemical agents (herbal components) in aquaculture has greatly increased as they have been reported to prevent diseases and reduce the use of hazardous antibiotics (Sakai, 1999). Results from this study showed positive effects of dietary inclusion Α. lepidophyllus on growth performance utilization. and feed nutrients Incorporating A. lepidophyllus fruit powder in C. gariepinus feed markedly enhanced body weight and body weight gain compared to those fed the control diet. The highest response was observed in fish fed with the highest level of phyto-additive. These results agree with Soosean et al. (2010) and Abbasi et al. (2017), who showed that the highest final weight and weight gain was recorded when the African catfish was fed the highest level of Garcinia mangostana and Zingiber officinale powder in common carp, Cyprinus carpio diets. This result can be explained by the presence of anti-oxidative and antimicrobial substances such as flavonoids and phenolic compounds (Fogang et al., 2014; Namkona et al., 2017; Sokamte et al., 2018) which are known to maintain gut equilibrium and improve growth performance of animal (Frankič et al., 2009). The highest improvement in body weight was recorded with C. gariepinus fed 6 g/kg A. lepidophyllus powder. This could be attributed to the antimicrobial properties of their specific active compounds and their impact on gut function. Increasing feed inclusion levels of A. lepidophyllus increased flavonoids and phenolic compounds. which are known to improve animal performances by changing the intestinal ecosystem of the animal through their antimicrobial action (Odoemelam et al., 2013). These compounds act by forming complexes with many proteins, causing the destruction of the bacterial membranes (Frankič et al., 2009). The changes in the intestinal ecosystem due to their antimicrobial action could lead to greater availability of some nutriments for the host and consequently improve body weight gain and feed utilization. This is in agreement with Frankič et al. (2009), who noticed the growthpromoting effect of most herbs and spices that act by killing parasites that hinder the digestibility and growth performance of animals. On the other hand, the improvement in body weight of C. gariepinus juveniles obtained with high a level of incorporation of A. lepidophyllus could also be attributed to the potent antioxidant properties of their major components as reported by Fogang et al. (2014) and Moukette et al. (2015).

Some phyto-additives have been reported to improve specific growth rate (Dada and Ikuerowo, 2009; Nyadjeu *et al.*, 2021). In this study, the fish fed diet containing 6 g/kg *A. lepidophyllus*  powder had the specific growth rate value significantly higher than all the other treatments. This result contradicts those obtained using dietary Zingiber officinale extract in beluga juvenile (Huso huso) diet (Vahedi et al., 2017). The conflicting results of the present study with those of Vahedi et al. (2017) could be attributed to the different species, feeding programs, and farming conditions. On the other hand, it was noted that regardless of the treatment, juvenile growth was faster during the first week of feeding and gradually decreased with the increase in the period. This experimental was comparable to that obtained by Gnikpo et al. (2014) who reported that the specific growth rate decreases while the muscle mass increases.

The improvement in feed nutrient utilization in C. gariepinus juveniles induced by A. lepidophyllus inclusion compared to those fed the control diet could have emanated from the presence of bioactive compounds contained in the phytobiotic. These results agree with the findings of Kana et al. (2017) where the incorporation of A. lepidophyllus fruits in broilers diet enhanced body weight and body weight gain and tended to reduce feed conversion ratio compared to birds fed the control diet. The improvement in growth and feed nutrient utilization in animals could be attributed to gallic acid which was the most abundant phenolic acid present in A. lepidophyllus fruit (Sokamte et al., 2018). According to Kang Yang et al. (2020), gallic acid influences the gut microbiome and modulates the immune

response to maintain intestinal health. The improvement of intestinal health reduced the animal's exposure to microbial toxins and other undesired microbial metabolites such as ammonia and biogenic amines (Windisch et al., 2008). Thereby, animals are relieved from immune defense stress during critical situations and there is increased availability of essential nutrients for absorption. This is in accordance with the results from this study where there was an increase in nutrient retention (protein, lipid, ash, energy, and dry matter) in the fish fed diets supplemented with A. lepidophyllus compared to those in the control diet. Moreover, the fish fed diet containing 6 g/kg A. lepidophyllus powder registered the highest nutrient retention of these macro-elements. This result indicates that increasing A. lepidophyllus level in C. gariepinus diet increases the presence of bioactive compounds present in the phyto-additive such as flavonoids and phenolics compounds (Sokamte et al., 2018) These secondary metabolites are known to maintain gut equilibrium (Frankič et al., 2009) of the microbiome that plays a crucial role in the maintenance of physiological homeostasis within the gastrointestinal tract (Kang Yang et al., 2020). Hence, they help the animals to grow better within the framework of their genetic potential.

The results of the present study revealed that supplementing *C*. *gariepinus* diet with *A. lepidophyllus* improved growth performance, feed nutrient utilization, and nutrient retention in the African catfish juveniles. The highest effect was obtained with the highest level of inclusion. These observed beneficial effects have been attributed to the bioactive contents of *A*. *lepidophyllus* than to their nutritional properties. The present findings provided a ray of hope for dietary supplements that are widely and intensively used in aquaculture.

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