

**Studies on the Breeding Biology And Behaviour
of Barbus Pituitora (Mahseer)**

C-QU / BIO (84)
FINAL REPORT
(January 1978 – December 1978)

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PAKISTAN SCIENCE FOUNDATION

ANNUAL RESEARCH REPORT

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17. Designation: Assistant Professor
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FINAL REPORT

NAME & ADDRESS OF THE
INSTITUTION:

DEPARTMENT OF BIOLOGICAL SCIENCES
QUAID-I-AZAM UNIVERSITY, ISLAMABAD.

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PROJECT NO:

C-QU/BIO (84)

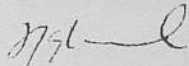
PROJECT TITLE:

STUDIES ON THE BREEDING BIOLOGY
AND BEHAVIOUR OF BARBUS PITUITORA
(MAHSEER)

REPORT PERIOD

JANUARY, 1978 TO DECEMBER 1978.

VICE-CHANCELLOR
QUAID-I-AZAM UNIVERSITY
ISLAMABAD


(DR. QAZI JAVED IQBAL)
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S U M M A R Y

The present investigations were aimed initially at the primary productivity of the Ramli stream used for supplying running water to experimental ponds employed to study the breeding biology and behaviour of Mahseer (Barbus Pituitora). These investigations are important as Mahseer is an economically important fish but very difficult to breed. It is of utmost importance to know A-biotic and biotic factors operating and controlling a particular fish fauna. Our investigations show that a Temperature of 16^o- 34^oC and dissolved Oxygen (2.7-7.9 ppl) are important controlling factors in the breeding of this fish. Besides, phyto- and zooplankton, of the stream are described with reference to those one on which the fish predate upon, such as Cyclops magnus, Diaptomus sp. Daphina sp. Moina brachitata Chydeus ovalis. Other a-biotic factors like pH and phosphate content are described for reference.

Growth rate of the fish under different dietary conditions have been worked out, using length-weight relationships. Two running water ponds and four aquariums were employed in the study. The investigations show that 35% proteins in the fish diet are best suited for adequate growth rate and breeding of B. pituitora.

The length-weight relationships show that $W = aL^b$
(-3.7590227 + 2.4026106 log L).

Feeding behaviour is initiated by visual stimuli and oriented by orthokineses, though telotaxis and dorsal light reaction play important role. Food orientation is impressed by pavlovian classical conditioning.

Diurnal short length and long length migration are supported by protean behaviour which acts as energy conservation mechanism. Both zigzagging pattern and looping is observable.

Sexual Behaviour breeding start in the end of April and last upto middle of Sept.

Spawning was induced in the fish by giving injection of preserved pituitary gland in a ratio of 5 mg/Kg of the body weight of the fish. Spawning occurred after 7-7½ hrs. of the injection.

INTRODUCTION

Fresh water fishes of Pakistan have a significant economic importance due to their nutritive value especially animal proteins. Taxonomic status of most of these fresh water fishes has been established and catalogued (Ahmad, 1961). Attempts have also been made by the fisheries deptt; to update fresh water fisheries management programme for the proliferation, survival, feed, breeding and economic use of some of these fishes. It need be emphasized here that for successful breeding and propagation of fishes, controlled mechanism involved are to be related to the ecological complexes in which these fishes thrive. In this regard little work has been done (Baqai and Zubairi, 1974, Azra and Iqbal, 1975). Another important factor operating on the fish biology is fish feed and food preference and the fish growth in relation to feed and ecological factors such as dissolved oxygen, temperature, phosphates, nitrates and chlorides. Little work is reported on the food habits and feed of carps (Menon, 1955; Hussain; 1955; Vas, 1957; Alikunbi, 1958; Arshad, 1965; Desai, 1968; Ahmad; 1969; Sukmaran, 1969; Hephher, 1969; Szunmiec; 1969, Khandker, 1970; Javed, 1970; Chinkich, et al, 1976). Lastly, investigations are needed to breed the fish artificially in the fish pond, Here also we find scanty reports on the experimental breeding of the carps (Hussain, 1965; Wlodek, 1968; Choudhury 1968, 1969; Konradt, 1968; Jafri, 1973; 1977; Doha and Dowan, 1967; Qazi, 1965).

The present investigation were aimed at describing environmental factors such as dissolved oxygen and organic phosphate pH and temperature, which play an important role in controlling survival growth and successful breeding of the fish.

The investigation also describe food formulation, food preference and length weight relationship of the fish. Lastly investigation briefly described experimentation with induced breeding.

PLAN OF WORK

1. Limnological Studies of the Supply Stream.
2. Limnological Studies of the Experimental ponds.
3. Collection of Fish
4. Studies on the Environmental factors controlling survival and breeding of the fish.
5. Food formulation.
6. Studies on the feeding behaviour.
7. Studies on the weight-length relationship.
8. Studies on Energy conservation..
9. Breeding season
10. Induced Breeding.

METHODOLOGY

Primary productivity of the stream and ponds.

Water samples were collected from two stations selected at the stream and were then compared with samples collected from the two experimental ponds, supplied with stream water through 1/2" and 2" diameter pipes at different flow rate. Day and night samples were taken from the ponds. Samples were collected simultaneously upstream downstream and from the ponds. Ponds were of two different dimensions 45'x25' & 6' high & 35' & 5' high, water could either flow independently of each pond or could be run from pond I to pond II.

Oxygen curves were constructed by wrinkler's method as given below:

Estimation of dissolved oxygen by Wrinkler's method

Reagents:- Chemicals used for fixation

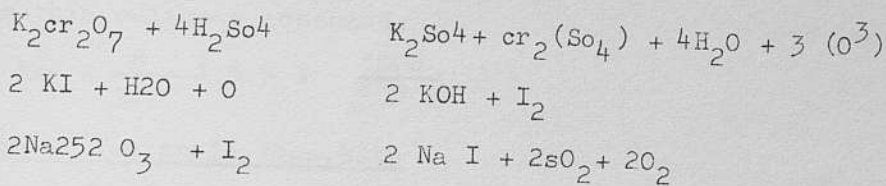
1. Manganese sulphate:- 367 gms. of $MnSO_4$ was dissolved in distilled H_2O and diluted to one liter.
2. Alkaline Iodide:- 360 gms. of $NaOH$ was dissolved in distilled & then 150 gms of KI was added to the solute, which was then diluted to one liter. crystals were dissolved by shaking.
3. Sulfuric Acid 50%

Chemicals used for titration:-

1. .01 N sodium thio sulphate:- 2.5 gms of $\text{Na}_2\text{S}_2\text{O}_3$ was weighed and dissolved in one liter of distilled H_2O .

Standardization of sodium thio-sulphate:- Stock solution 1.225 gms = .01 N of $\text{K}_2\text{Cr}_2\text{O}_7$ was dissolved in 1000 cc of distilled water chemically equivalent 0.025 N. To 10 c c of $\text{K}_2\text{Cr}_2\text{O}_7$ solution of .01 N, was added approx. 2.3 cc. of concentrated H_2SO_4 and 5 ml of 15% KI solution so that an equal equality of Iodine was liberated.

Reaction as follows



Procedure:- $\text{Na}_2\text{S}_2\text{O}_3$ was draw in a burette, and the level marked. 10 ml of prepared $\text{K}_2\text{Cr}_2\text{O}_7$ solution was taken in a conical flask, a few drops of Phenolphthelin was added as indicator showing blue colour. This was filtrated against $\text{Na}_2\text{S}_2\text{O}_3$ till, the solution became colourless.

Result:-

$$N \text{ of } \text{Na}_2\text{S}_2\text{O}_3 = N \text{ of } \text{K}_2\text{Cr}_2\text{O}_7 \times \frac{\text{vol of } \text{K}_2\text{Cr}_2\text{O}_7}{\text{vol of } \text{Na}_2\text{S}_2\text{O}_3}$$

2. Starch Solution:- 2%

Procedure:- Fixation of water samples:- 250 ml brown bottle was filled with water, so that no air bubble was trapped in. 2 ml of M_nSO_4 was added to it followed by 2 ml of alkaline solution. After shaking, a dense cloud of precipitation appeared at the bottom. The precipitate was allowed

to settle down. After half an hour 2 ml of H_2SO_4 was added. The precipitate dissolved and the sample became yellow. This colour is due to the formation of iodine.

Titration

The burette was filled with sodium thio-sulphate, the initial reading was noted. 20 ml of fixed sample was drawn by pipette in a titration flask, 5-6 drops of starch solution were added as indicator, the solution became blue, then titrated against sodium thio-sulphate drop by drop, till the sample became colourless. Final reading was noted. Difference of the two readings gave the amount of sodium-thio sulphate consumed.

Results: $Nxv \times \frac{B}{B-2} \times 5.6 \times \frac{1000}{5}$

Estimation of Inorganic Phosphate

Fiske & Subbarow Method (1925)

Reagents:-

1. Standard phosphate Solution:- 0.351 gms of pure dry mono-potassium phosphate was dissolved in water and transferred quantitatively to a 1-liter volumetric flask. 10 ml of 10 N sulfuric acid was added, and diluted to the mark with water, and mixed. This solution contained 0.4 mg of phosphorus in 5 ml. This made the solution stable indefinitely.
2. Molybdate Solution:- 25 gms of reagent grade ammonium molybdate was dissolved in approximately 200 ml of water. To a 1-liter volumetric flask containing 300 ml of 10 N sulphuric acid, the molybdate solution was added and diluted with washing to a 1 liter and mixed to render it stable indefinitely.

3. Aminonaphtholsulfonic Acid Reagent:- To 195 ml 15% sodium-bisulfite solution contained in a glass-stoppered cylinder, 0.5 gms of 1,2,4-aminonaphtholsulfonic acid was added and then 5 ml of 20% sodium sulfite. This was shaken until the powder was dissolved. The solution was transferred to a brown glass bottle and stored in the cold incubator. This solution was useable for about four weeks.

Procedure:-

Standard was prepared by taking 3 ml of stock standard, adding to it 12 ml of distilled water, so that it became 5 times dilute. From this 6 different concentrations were prepared as follow:-

- S₁ = 0.5 ml of 5 time dilute solution + 4.5 ml distilled water.
- S₂ = 1.0 " " " " " " " " 4.0 " " "
- S₃ = 1.5 " " " " " " " " 3.5 " " "
- S₄ = 2.0 " " " " " " " " 3.0 " " "
- S₅ = 2.5 " " " " " " " " 2.5 " " "
- S₆ = 3.0 " " " " " " " " 2.0 " " "

To each standard i.e. S₁ S₂.....S₆, 0.4 ml ANSA and 1 ml Molybdate solution was added.

Samples were prepared as:

T₁ = 5 ml of pure sample + 0.4 ml ANSA + 1 ml molybdate solution.

T₂ = 2.5 ml of sample + 2.5 ml distilled water + 0.4 ml ANSA + 1 ml molybdate solution.

The solution was left for 10-15 minutes and then transferred a portion of the coloured solution into a container and readings were taken with photometer at 660. The photometer was set at zero density with a blank, prepared by treating 5 ml of distilled water with 1 ml of molybdate solution and 0.4 ml of ANSA reagent. Densities of standard and samples were noted from photometer.

Results:-

$$\frac{\text{Density of Sample}}{\text{Density of standard}} \times \frac{\text{Standard's concentration}}{\text{Volume of sample}} \times 100$$

"= mg pi/100 ml of sample.

Temperature was noted using electronic thermister and pH. with a portable pH meter.

Observations were made over a period of six months. Approximate variables of dissolved Oxygen and temperature were also monitored using aquariums (4) 5'x3' & 4' high placed alongside the ponds and by supplying oxygen with air pump and different levels of warm water with heated water flow into the aquariums at different flow intensities (Fig.1).

COLLECTION OF FISH

Fishes were collected from Hassan Abdal fish farm Bara K h Stream, Chattar Bagh Stream and Outlet stream of Rawal Dam. Fishes were brought in fish containers.

Measurement of Fishes

Length: Fishes were measured from tip of the snout to the tip of the gill and from tip of the gill to the tip of tail fin and then from the tip of the tail fin to the tail fin end.

Weight. A container was half filled with water and weighed as (a), then the fish was put into it and again weighed as (b). The actual weight (W), was worked out as $b-a = W$.

Length-Weight relationship

The length-weight relationship was expressed as follows:
 $W = a L^b$, Where W is weight, L is length and a and b empirically derived constants determined by standard regression analysis. This was further expressed in Logarithmic form; $\log W = \log a + b \log L$ (Grover, 1976).

Each fish was tagged before release and bi-monthly length-weight measurements were taken to observe growth-food relationship.

Plankton fauna on which the fish were observed to predate was noted and identified.

Observations on protean Behaviour were made at the inlet stream of Rawal Dam. Observation on the maturity of male and female fish were made by (a) squeezing out spermatazoa and ova from the fish (b) by dissecting the fish and then scanning the testes and ovary under a Binocular microscope.

The breeding behaviour of the fish was observed in the experimental ponds.

Induced spawning was effected by injecting pituitary in the male and female fish as given below.

The pit. gland of the several fishes were taken out by dissecting the fish. It was then placed in a polythene bag and preserved in cooled incubator at -10°C .

The fishes of known weight and length were given injections of pituitary to affect spawning behaviour in April. Experiments on induced breeding were carried on in aquariums.

OBSERVATIONS & RESULTS

Limnological Studies

pH: Hydrogen ion concentration ranged between 7.2-8.6, which is slightly alkaline in nature.

Phosphate: Inorganic phosphates were maximum .03-.04 mg/ml in June-July, while in winter, the concentration was low being .003-.001 mg/ml.

Oxygen: Dissolved oxygen was minimum in June, ranging between 4.5-4.7/pp1 and was maximum in January ranging between 6.8-7.9/pp1.

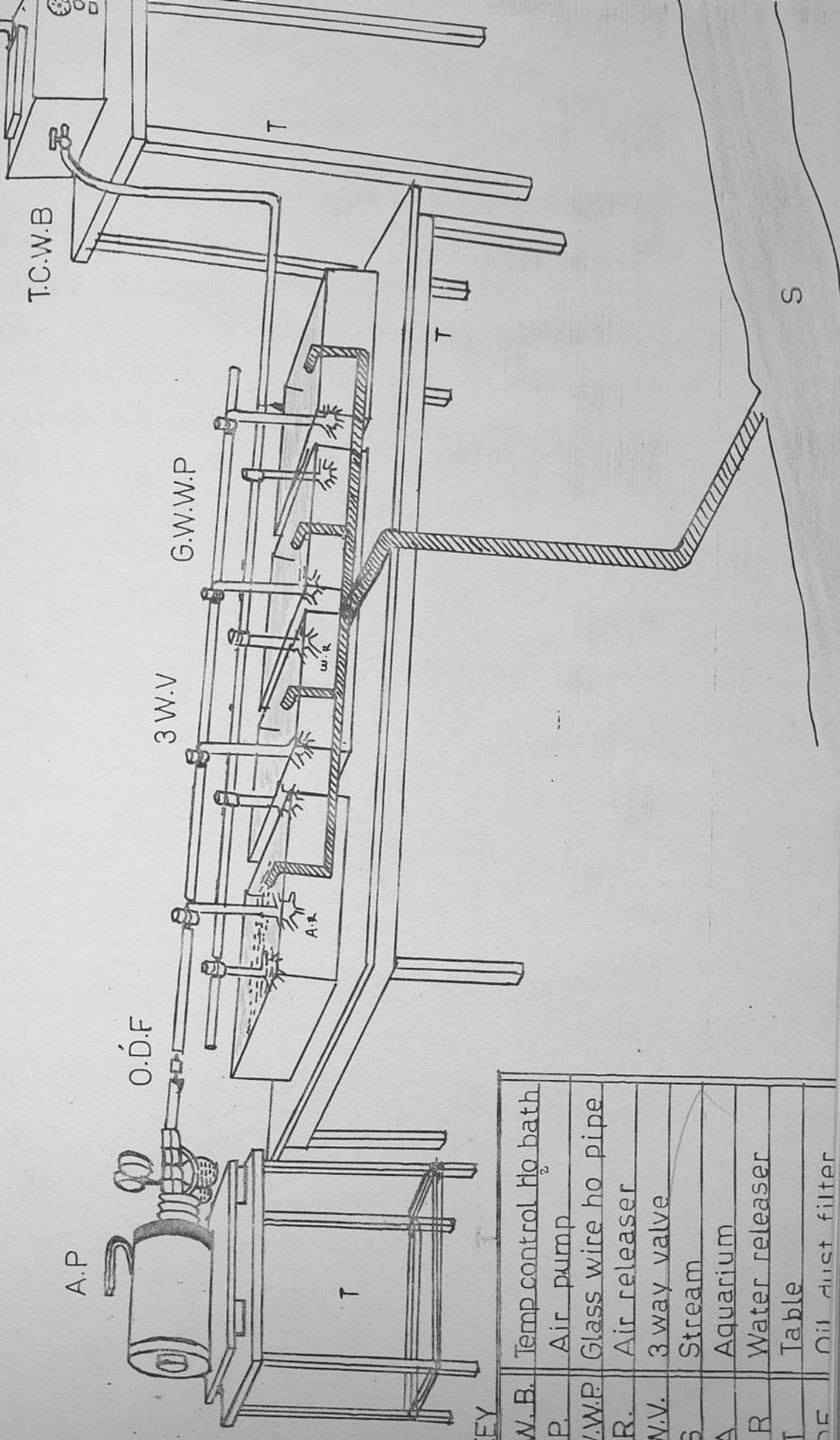
When water flow into the ponds was controlled and brought to a minimum, the dissolved oxygen content went down to a minimum and toxicity caused 100% mortality to the fish.

Experimental observations were made, using aquaria placed alongside the ponds (fig.I). Results indicated that a low concentration of oxygen (2.7/pp1) caused mortality to fish.

Temperature: Minimum temperature ranged between 13-15°C in Jan, and max. ranged between 20-36°C in June. Experiments on temperature effects were conducted in the aquaria (Fig.I) The results show that temperature is also an important factor in the breeding and survival of the fish and that a temperature ranging between 16-34°C is favourable for the fish.

Zooplanktons such as Cyclops magnus, Diatomus sp. Daphnia sp. Moina brachitata Chydorus ovalis are usually predated upon by the fish Barbus pituitora.

Fig -1



KEY	
T.C.W.B.	Temp control H ₂ bath
A.P.	Air pump
G.W.W.P.	Glass water pipe
A.R.	Air releaser
3 W.V.	3 way valve
S	Stream
A	Aquarium
R	Water releaser
T	Table
O.D.F.	Oil dust filter

CONCLUSIONS

Limnological studies indicate that dissolved oxygen is a critical factor in the survival and successful breeding of the fish Barbus pituitora and that the fish may only be bred and propagated in ponds, lakes or streams having a safe range of dissolved oxygen level as indicated in the results. It is also shown that temperature also effects successful maintenance and breeding of the fish.

Table No. I

Formulation of feed containing different level of protein.

Ingredients	0% Protein	25% Protein	30% Protein	35.0 gms.
Casein	0.0 gms	25.0 gms.	30.0 gms	35.0 gms.
Starch	20.0 "	10.0 "	15.0 "	10.0 "
Cellulose	30.0 "	25.0 "	18.0 "	19.0 "
Minerals	4.0 "	2.0 "	2.0 "	2.0 "
Vitamins	2.0 "	4.0 "	4.0 "	4.0 "
Oil	25.0 "	19.0 "	16.0 "	15.0 "
Glucose	20.0 "	15.0 "	15.0 "	15.0 "
	100.0 = 385 calories value	100.0 = 371 calories value	100.0 = 384 calories value	100.0 = 375 calories value

FEDDING BEHAVIOUR

Observations and Feeding Behaviour.

Feeding Behaviour is initiated by visual stimuli of the food and orientation as by telotaxis and orthokinesis, through dorsal light reaction play important role in the intensity of orientation Food orientation may be impressed by Pavlovian conditioning..

Food Relationships.

Results:- Four dietary experiments were conducted on the fish, alongwith a control groups. Each group was of four fishes, each fish was measured weighed, tagged and then released into the experimental aquariums. Results are shown in tables 2-10. For comparison see graphs of weight relationship and length relationship(see graph 1-4).

The fish food was prepared is given in table I.

The results indicate that 35% protein in the diet are best suited for adequate growth of the fish.

Observation:-

On feeding behaviour show that stimulation orientation mechanism play a supporting factor in artificial feeding of the fish and that the fish may be conditioned to feed an farmulated feed.

Conclusions:-

It may be concluded from the experiments on A-biotic and biotic factors favourable as shown in liminological section, a diet containing 35% proteins is best suited for the beneficial breeding and propagation of the fish Barbus pituitora

Table No II

Initial ~~consent~~ weight and length of the Mahseer (Barbus pituitora) before treating with any specific diet.

No of Species	Total weight of the Species <u>Barbus pituitora</u> (gms) W	Length from mouth to the end of gill cover (mm)	Length from end of gill cover upto end of body (mm)	Length of tail fin (mm)	Total length from mouth upto tail end (mm)	Total Increasing length of the species <u>B. pituitora</u> (mm)	Total Increasing weight of the <u>B. pituitora</u> (gms) W
1.	4.0	10	50	15	75	x	x
2.	4.0	10	50	15	75	x	x
3.	4.0	10	50	15	75	x	x
4.	4.0	10	50	15	75	x	x

Table No. III Group A.

Result of the feeding Experiments with Mahseer (Barbus. pituitera) group A.

No of Species	Total weight of the Species Barbus pituitera (gms) W	Length from mouth the end of gill cover (mm)	Length from end of gill cover upto end of body (mm)	Length of tail fin (mm)	Total length from mouth upto tail end (mm)	Total Increasing length of the species B. pituitera (mm)	Total Increasing weight of the B. pituitera (gms)w
.	4.8	10	54	15	79	4	0.8
.	4.8	10	54	15	79	4	0.8
.	4.8	10	54	15	79	4	0.8
.	4.8	10	54	15	79	4	0.8

Mahseer (Barbus pituitera) were fed experimental diet containing (0% protein) for fifteen days at 25-27°C of water temperature.

No mortality occurred during the experimental period.

Table No. IV (Group A)

Result of the feeding experiment with Mahseer (Barbus pituitora) group A.

No of Species <u>Barbus pituitora</u>	Total weight of the Species <u>Barbus pituitora</u> (gms) W	Length from mouth to the end of gill cover (mm)	Length from end of gill cover upto end of body (mm)	Length of tail fin (mm)	Total length from mouth upto tail end (mm)	Total Increasing length of the species <u>B. pituitora</u> (mm)	Total Increasing weight of the <u>B. pituitora</u> (gms)w
1.	5.7	16	58	20	94	19	1.7
2.	5.7	16	58	20	94	19	1.7
3.	5.7	16	58	20	94	19	1.7
4.	5.7	16	58	20	94	19	1.7

Mahseer (Barbus pituitora) were fed experimental diet containing (0% protein) for thirty day at 25-27°C of water temperature.

No mortality occurred during the experimental period.

0% Protien.

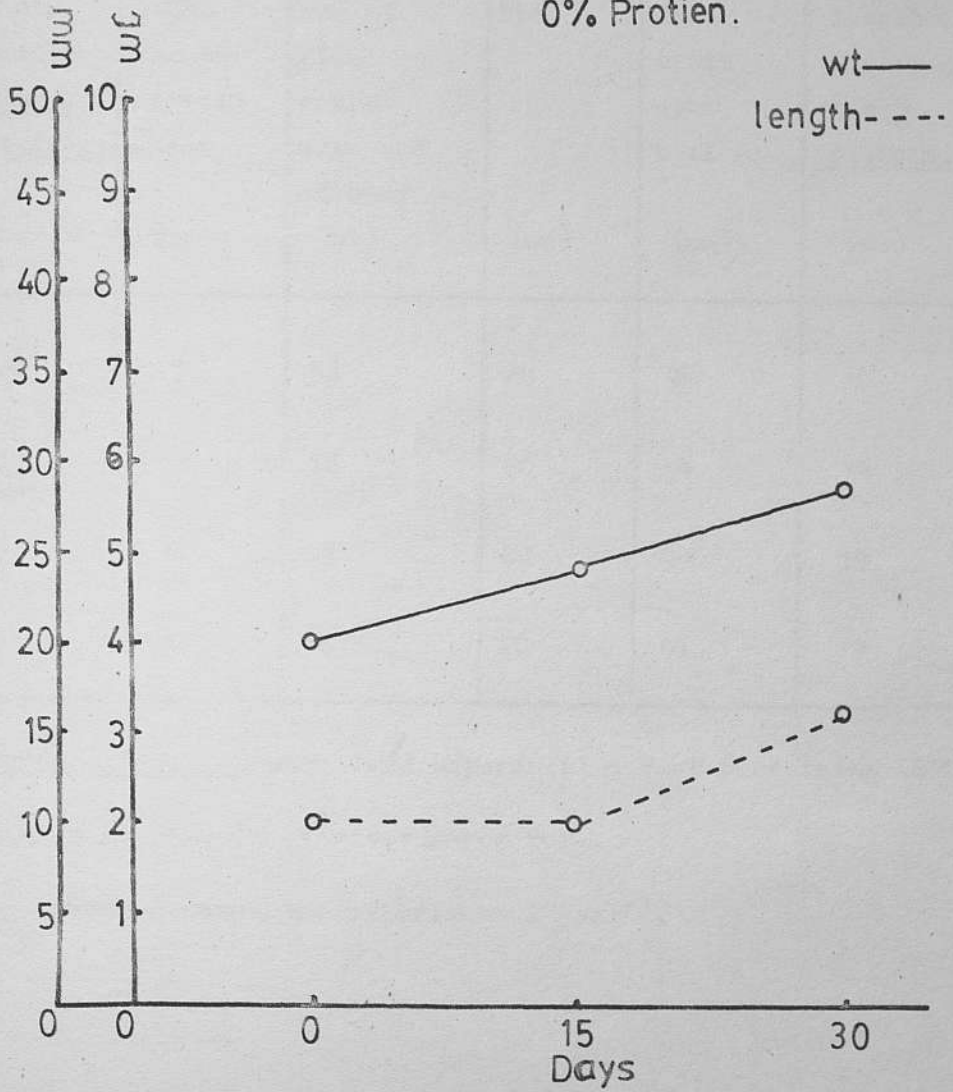


Table No. V (Group B)

Result of the feeding experiment with Mahseer (Barbus pituitera) group B.

No. of fish	Total weight of the Species <u>Barbus</u> <u>pituitera</u> (gms) W	Length from mouth to the end of gill cover (mm)	Length from end of gill cover upto end of body (mm)	Length of tail fin (mm)	Total length from mouth upto tail end (mm)	Total Increasing length of the spec- ies <u>B.</u> <u>pituitera</u> (mm)	Total Increasing weight of the <u>B.</u> <u>pituitera</u> (gms) w
	6.0	16	58	20	94	19	2.0
	6.0	16	58	20	94	19	2.0
	6.0	16	58	20	94	19	2.0
	6.0	16	58	20	94	19	2.0

Mahseer (Barbus pituitera) were fed experimental diet containing (25% protein) for fifteen days at 25-27°C of water temperature.

No mortality occurred during the experimental period.

Table No VI (Group B)

Result of the feeding experiment with Mahseer (Barbus pituitora) Group B.

Species <u>Barbus pituitora</u>	Total weight of the Species <u>Barbus pituitora</u> (gms) w	Length from mouth the end of gill cover (mm)	Length from end of gill cover upto end of body (mm)	Length of tail fin (mm)	Total length from mouth upto tail end (mm)	Total Increasing length of the species <u>B. pituitora</u> . (mm)	Total Increasing weight of the <u>B. pituitora</u> . (gms) w
	8.4	18	60	21	99	24	4.4
	8.4	18	60	21	99	24	4.4
	8.4	18	60	21	99	24	4.4
	8.4	18	60	21	99	24	4.4

Mahseer (Barbus pituitora) were fed experimental diet containing (25% protein) for thirty days at 25-27°C of water temperature.

No mortality occurred during the experimental period.

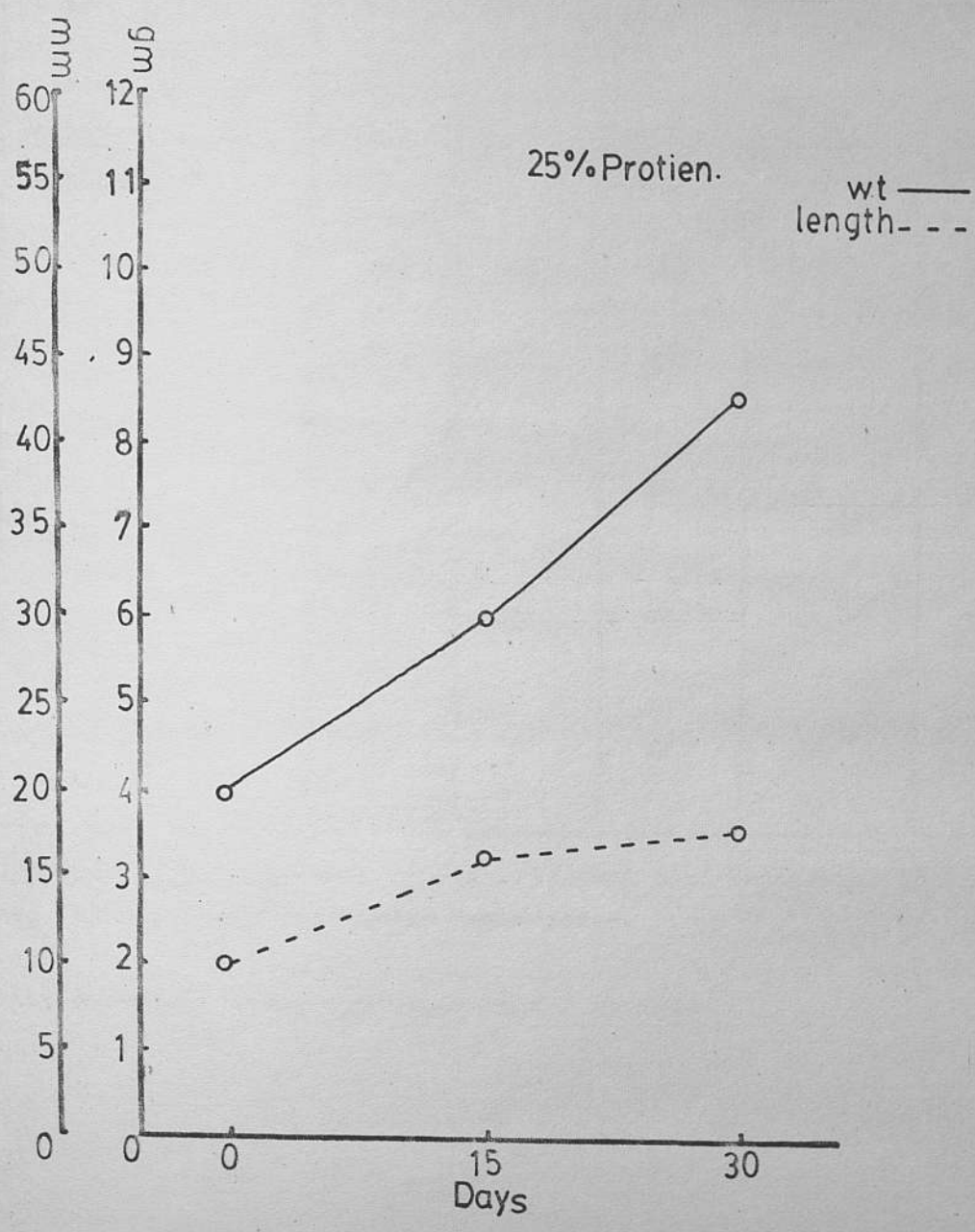


Table No.VII(Group C)

Result of the feeding experiment with Mahseer (Barbus. pituitora) group C.

Species <u>Barbus. pituitora</u>	Total weight of the Species <u>Barbus. pituitora</u> (gms) W	Length from mouth to the end of gill cover (mm)	Length from end of gill cover upto end of body (mm)	Length of tail fin (mm)	Total length from mouth upto tail end (mm)	Total Increasing length of the species <u>B. pituitora</u> . (mm)	Total Increasing weight of the <u>B. pituitora</u> . (gms) w
	7.1	18	60	21	99	24	3.1
	7.1	18	60	21	99	24	3.1
	7.1	18	60	21	99	24	3.1
	7.1	18	60	21	99	24	3.1

Mahseer (Barbus. pituitora) were fed experimental diet containing (30% protein) for fifteen days at 25-27°C of water temperature.

No mortality occurred during the experimental period.

Table No. VII (Group C)

Result of the feeding experiment with Mahseer (Barbus. pituitora) group C.

Species	Total Weight of the Species (gms) W	Length from mouth to the end of gill cover (mm)	Length from end of gill cover upto end of body (mm)	Length of tail fin (mm)	Total length from mouth upto tail end (mm)	Total Increasing length of the species <u>B. pituitora</u> . (mm)	Total Increasing weight of the <u>B. pituitora</u> . (gms) w
<u>Barbus. pituitora</u>	10.5	20	71	22	113	38	6.5
<u>Barbus. pituitora</u>	10.5	20	71	22	113	38	6.5
<u>Barbus. pituitora</u>	10.5	20	71	22	113	38	6.5
<u>Barbus. pituitora</u>	10.4	20	71	22	113	38	6.4

Mahseer (Barbus. pituitora) were fed experimental diet containing (30% protein) for thirty days at 25-27°C of water temperature.

No mortality occurred during the experimental period.

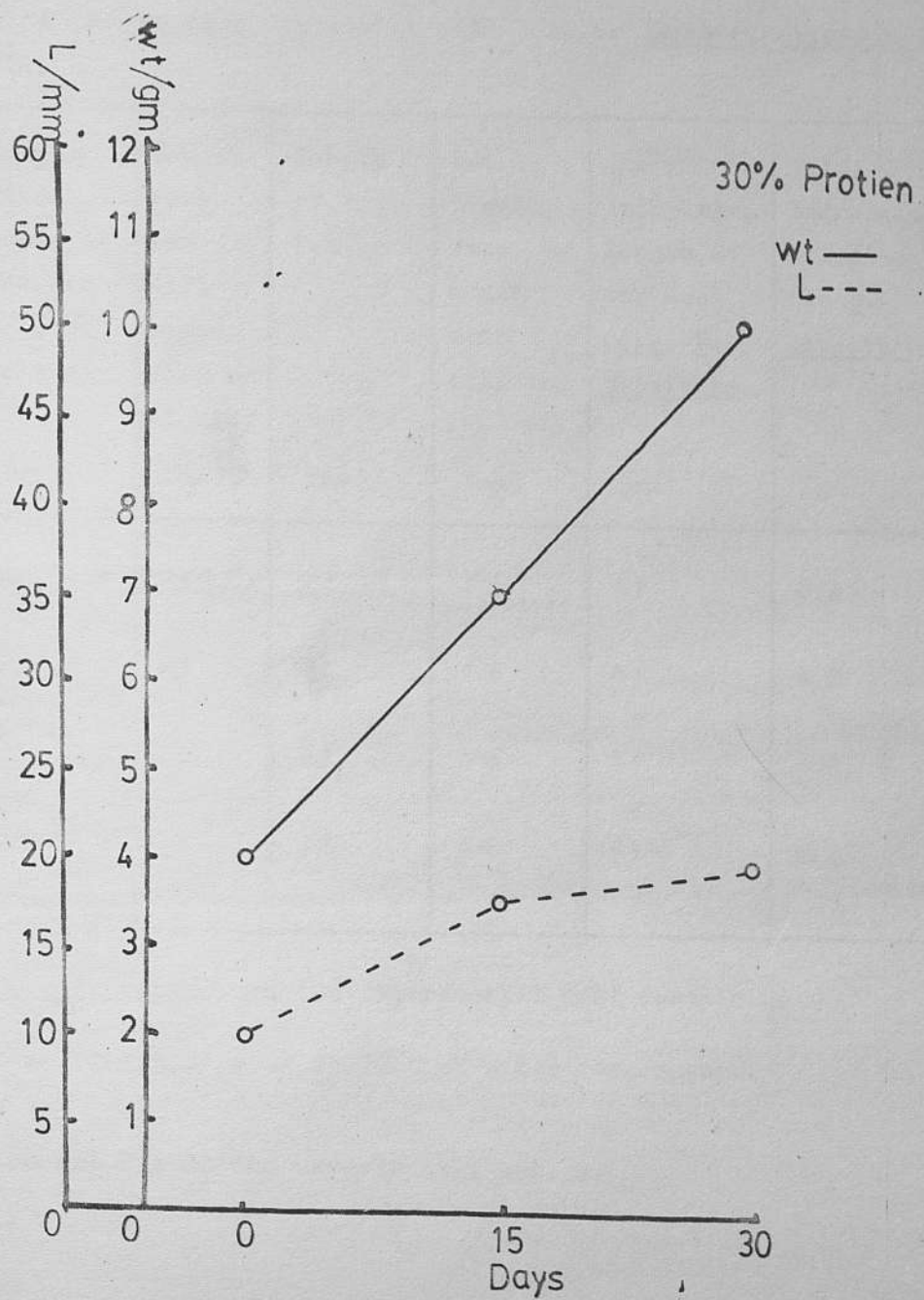


Table No. IX (Group D)

Result of the feeding experiment with Mahseer (Barbus. pituitora) group D.

No. of species <u>Barbus. pituitora</u>	Total weight of the Species <u>Barbus. pituitora</u> (gms) w	Length from mouth to the end of gill cover (mm)	Length from end of gill cover upto end of body (mm)	Length of tail fin (mm)	Total length from mouth upto tail end (mm)	Total Increasing length of the species <u>B. pituitora.</u> (mm)	Total Increasing weight of the <u>B. pituitora</u> (gms) w
	12.2	21	73	22	116	41	8.2
	12.2	21	73	22	116	41	8.2
	12.2	21	73	22	116	41	8.2
	12.2	21	73	22	116	41	8.2

Mahseer (Barbus. pituitora) were fed experimental diet containing (35% protein) for fifteen days at 25-27°C of water temperature.

No mortality occurred during the experimental period.

Table No. X (Group D)

Result of the feeding experiment with Mahseer (Barbus. pituitora) group D.

No. of Species	Total weight of the Species	Length from mouth to the end of gill cover	Length from end of gill cover upto end of body	Length of tail fin	Total length from mouth upto tail end	Total Increasing length of the species	Total Increasing weight of the <u>B. pituitora</u> .
<u>Barbus. pituitora</u>	<u>Barbus. pituitora</u>					<u>Barbus. pituitora</u>	
	(gms) W	(mm)	(mm)	(mm)	(mm)	(mm)	(gms) w
1.	25.2	28	90	29	147	72	21.2
2.	25.2	28	90	29	147	72	21.2
3.	25.2	28	90	29	147	72	21.2
4.	25.2	28	90	29	147	72	21.2

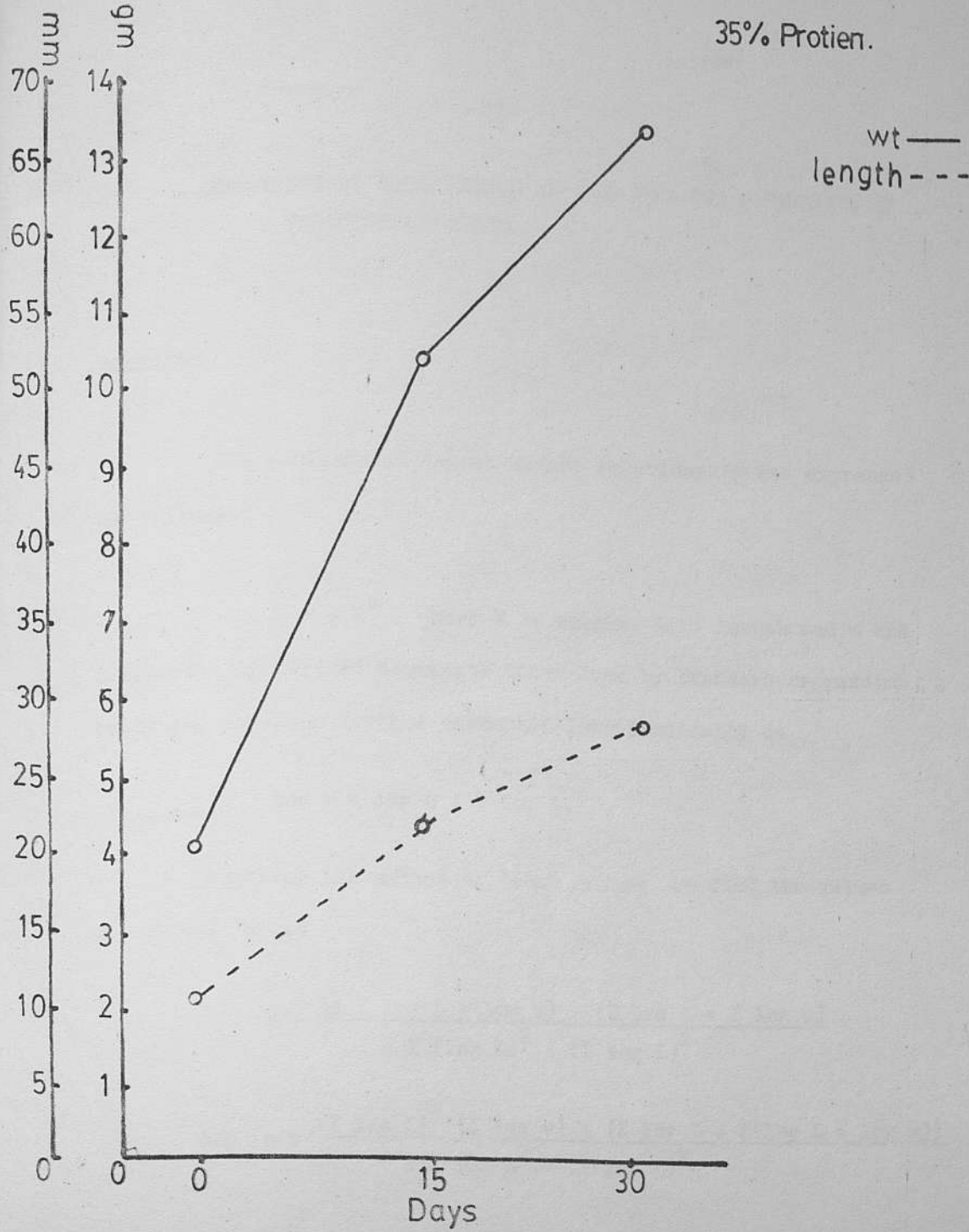
Mahseer (Barbus. pituitora) were fed experimental diet containing (35% protein) for thirty days at 25-27°C of water temperature.

No mortality occurred during the experimental period.

GRAPH NO. 4

WEIGHT LENGTH RELATIONSHIP OF THE FISH (BARBUS- PITUITORA)
FED ON 35% PROTEIN.

35% Protien.



LENGTH-WEIGHT RELATIONSHIP OF THE FISH FOR ESTIMATION OF
PRODUCTIVITY LEVEL.

RESULTS:

The analysis of length-weight relationship was expressed as follows:

$W = a L^b$, where W is weight, L is length and a and b empirically derived constants determined by standard regression analysis. This was further expressed logarithmically as

$$\log W = \log a + b \log L.$$

Applying the method of least square, we find the values of 'a' and 'b' as

$$b = \frac{(n \sum \log L * \log w) - (\sum \log L * \sum \log w)}{n \sum (\log L)^2 - (\sum \log L)^2}$$

$$\log a = \frac{(\sum \log L)^2 (\sum \log w) - [\sum \log L * \sum (\log L * \log w)]}{n \sum \log L^2 - (\sum \log L)^2}$$

Now putting the values from table in the equation for finding the values of b and log a.

$$\begin{aligned} b &= \frac{(n \Sigma (\log L * \log w) - (\Sigma \log L * \Sigma \log w))}{n \Sigma (\log L)^2 - (\Sigma \log L)^2} \\ &= \frac{(23 * 91.301445) - (39.158889 * 52.934678)}{23 * 122.32554 - (52.934678)^2} \\ &= \frac{27.0701}{11.4073} = 2.4026106 \end{aligned}$$

$$b = 2.4026106$$

$$\begin{aligned} \text{For log a} &= \frac{(n \Sigma \log L * \log w) - (\Sigma \log L * \Sigma \log w)}{n \Sigma (\log L)^2 - (\Sigma \log L)^2} \\ \log a &= \frac{[\Sigma (\log L)^2 (\Sigma \log w)] - [(\Sigma \log L * \Sigma (\log L * \log w))]}{n \Sigma (\log L)^2 - (\Sigma \log L)^2} \\ &= \frac{(122.32554 * 39.158889) - (52.934678 * 91.301445)}{(23 * 122.32554) - (52.934678)^2} \\ &= -\frac{42.8803}{11.4073} = -3.7590227 \end{aligned}$$

$$b = 2.4026106$$

$$\log a = -3.7590227$$

so equation becomes

$$\log w = -3.7590227 + 2.4026106 \log L$$

$$w = \text{antilog} (-3.7590227 + 2.4026106 \log L)$$

consult Table 11.

CONCLUSION:

The empirically derived results show that any experiments to be conducted on food, feeding behaviour and propagation of the fish should be deduced using length-weight relationships, so that comparable results may be obtained.

No. of species	W gram	L m.m.	Logw	LogL	LogL*logw	logL ²
1.	7.0	99	.84509	1.995635	1.6864911	3.982559
2.	11.0	113	1.041392	2.053078	2.138059	4.21512292
3.	15.0	121	1.176091	2.082785	2.4497520	4.3379933
4.	18.0	132	1.255272	2.120573	2.6618959	4.4963298
5.	25.0	147	1.39794	2.167317	3.0297791	4.6972629
6.	28.0	152	1.447158	2.18184	3.1574497	4.7604257
7.	35.0	165	1.544063	2.21748	3.4239399	4.9172175
8.	40.0	177	1.602059	2.2479732	3.6013856	5.0533835
9.	46.0	184	1.662757	2.264817	3.7658403	5.129396
10.	50.0	189	1.69897	2.2364618	3.7996815	5.0017613
11.	62.0	213	1.79239	2.328379	4.1733632	5.4213487
12.	66.5	212	1.82282	2.326335	4.2444899	5.4118345
13.	72.0	225	1.857332	2.35218	4.36878815	5.53227507
14.	75.6	230	1.87852	2.36172	4.4365382	5.5777213
15.	79.5	239	1.900367	2.378397	4.5193271	5.6567722
16.	82.0	245	1.913813	2.389166	4.5724169	5.7081141
17.	88.0	254	1.94448	2.404833	4.6761496	5.7832217
18.	93.5	261	1.970811	2.4166405	4.7627416	5.8401513
19.	105.0	280	2.021189	2.4477158	4.9461683	5.9835822
20.	113.0	288	2.05307	2.459392	5.0493039	6.048609
21.	114.5	296	2.058805	2.4712917	5.0879077	6.1072826
22.	132.5	317	2.122215	2.501059	5.3077849	6.2552961
23.	142.0	339	2.15228	2.530199	5.4456967	6.4019069
			39.158889	52.934678	91.301445	122.32554

OBSERVATION ON PROTEAN BEHAVIOUR

The fish show two type of diurnal movements; Short-length movements in slow water. These donnot exceed more than 60 yards up-stream and down-stream lures. Long-length movements during which the fish and the school travel up-stream in the morning (6.00-6.30 am) travelling several hundred yards and then travel back to the lake at approx. 11.30 am to 12.00 noon.

PROTEAN MOVEMENTS

The fish show single erratic movements by zig-zagging or looping in the stream. This protean behaviour is shown during down-stream and up-stream journey, fish do not travel parallel to water flow but show protean behaviour by zig-zagging so that when they travel down-stream the water flow is at 120° to the right or left of the front region of the fish. Similarly, while travelling up-stream, the water flow hits the tail at 60° . This zig-zagging protean behaviour is used as a postural device by the fish to offer least resistance in water and at the same time, movements at an angle to water flow facilitate swift journey using least possible energy. Whereas, when showing movement-pause the fish always orientate parallel to water flow.

CONCLUSION

The fish Barbus pituitora uses protean behaviour as a means of energy conservation during up-stream a down-stream for energy.

INDUCED BREEDING

Two males weighing $1\frac{1}{2}$ kg and one female weighing $1\frac{1}{2}$ kg were used in the induced spawning experiment.

The preserved pituitary (male and female) of the fishes was homogenized and diluted with a drop of distilled water. The sample was then centrifuged. Intramuscular injection of the female pituitary was then given to the female fish at a ratio of 3 mg/kg of the body weight. After four hours another injection was given at a ratio of 5 mg/kg. The female was then freed into the hapae net. The net was spread in the middle of the aquarium water, so that it did not touch the bottom.

Same process of injecting male pituitary was simultaneously done with the two males, which were also released into the same net. After some time the female showed splashing movements in water. After seven and half hours the female secreted the spawn and the male discharged milt over the eggs.

Hatching of the eggs occurred after fourteen hours of the fertilization. Two nets were used, one was large and the other one small. The larger hapae contained the smaller one inside it. When the eggs hatched the larvae came out of the meshes of the smaller one and went to the larger net mesh, through which the fingerlings could not pass down.

After the completion of this procedure, all the fingerlings were transferred to the pond. The cage was not poured at once into the pond because the abrupt change of temperature may affect the fingerlings. The cage was gradually poured so that the fingerlings could acclimatize to the temperature.

GENERAL DISCUSSION

Present studies were conducted to study the possibility of successful breeding and propagation of the fish Mahseer Barbus so that it may be utilized as an economic crop.

A-biotic factors play an important role in the productivity of a given. Initial studies on the stream supplying water to experimental tanks, and later experimental studies conducted in the aquariums indicate that temperature and Oxygen are controlling factors in the successful propagation and survival of the fish in fish tanks. Similarly, pH also plays an important role in the survival of the fish.

It is important to know a balanced diet if one intends to use fish for economic purposes. Previously, studies have been made on carps with respect to their dietary energy. But these were mainly concerned on the utilization of carbohydrates (G. Jayachandran et al., 1969; Chinkichi et al., 1976). Present studies indicate that for economic growth of the fish a diet consisting 35% protein is most efficient as shown in length-weight relationship viz $W = \text{antilog} (-3.7590227 + 2.4026106 \log L)$.

Failure of the major carps to breed in captivity, is in part due to the refractiveness to induced spawning, though successful studies have been made in some carps (Doha and Dewan 1964; Qazi 1965; Mitra, 1969; Choudhri 1969; Singh, 1969; Jafri, 1977.

Present studies indicate a break through by successfully inducing spawning with hormone injection.

Occurance of protean behaviour in prey organisms has been reviewed and discussed by Humphries and Driver (1967), and Humphries and Driver (1970). No evidence is available on the eco-behavioural aspects of protean displays involved in other behavioural patterns except for courtship displays (Iqbal, personal observation). Present studies indicate that protean displays are used as postural device for energy conservation. Such postural erratic devices are comparable to sudden jet blasts employed for propulsion.

1. NEED FOR ADDITIONAL RESEARCH:

Further studies are recommended on the feeding behaviour and dietary formulation. These studies will be made on weight length relationships, so that a model for economic breeding of the fish may be formulated.

2. Studies on the sexual behaviour are also recommended as only initial observations were made during the present study.

3. Studies on the growth rate of the fish are also recommended for economic purposes.

4. Lastly detailed studies on the breeding biology, using modern induced breeding techniques should be made.

It is recommended that the above mentioned studies should be conducted for a period of three years.

Note:- An extension scheme is being submitted under separate cover.

LIST OF PUBLICATIONS:

Papers under preparations.

- i) Environmental factors controlling the survival and breeding of the fish Barbus pituitora. (Mahseer)
- ii) Feeding behaviour of the fish Barbus pituitora.
- iii) Length weight relationship of the fish Barbus pituitora
- iv) Protean behaviour exhibited by the fish Barbus pituitora as a postural device for energy conservation.

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HEAD WISE EXPENDITURE DURING THE PERIOD

JANUARY 1978 TO DECEMBER 1978

RECURRING

Heads	Provision	Expenditure	Out Standing Liabilities
1. Chemicals & Glassware	4000/-	4000/-	-
2. Conveyance	1000/-	1000/-	-
3. Films	4000/-	4000/-	-
4. Books & Journals	1000/-	1000/-	-
TOTAL	10,000/-	10,000/-	-

DETAILS OF FISCAL REPORT

Pay and allowance paid to Research Officer during the period
January 1978 to December 1978.

Name of Research Officer

Mr. Fazile Subhan

Period	Pay	M.A.	C.A.	L.C.A.	H.R.	Net Payment
January	900/-	100/-	60/-	90/-	630/-	1780/-
February	900/-	100/-	60/-	90/-	630/-	1780/-
March	900/-	100/-	60/-	90/-	630/-	1780/-
April	900/-	100/-	60/-	90/-	630/-	1780/-
May	900/-	100/-	60/-	90/-	630/-	1780/-
June	900/-	100/-	60/-	90/-	630/-	1780/-
July	900/-	100/-	60/-	90/-	630/-	1780/-
August	900/-	100/-	60/-	90/-	630/-	1780/-
September	900/-	100/-	60/-	90/-	630/-	1780/-
October	900/-	100/-	60/-	90/-	630/-	1780/-
November	900/-	100/-	60/-	90/-	630/-	1780/-
December	900/-	100/-	60/-	90/-	630/-	1780/-
TOTAL	10800/-	1200/-	720/-	1080/-	7560/-	21,360/-
