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Note

# Amino-acid and biochemical composition of whole body and molt of the Indian white shrimp

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#### ABSTRACT

The amino acid and biochemical composition of molt (exuvia) and whole body of the shrimp, *Fenneropenaeus indicus* were analysed along with morphometric relations. The wet weight of exuvia  $(0.22 \pm 0.07 \text{ g})$  is nearly 4.8% of the whole shrimp weight  $(4.50 \pm 1.42 \text{ g})$ . The goodness fit statistics R<sup>2</sup> revealed highly significant relationship in weights of wet exuvia- wet shrimp, dry exuvia- wet shrimp and dry exuvia- dry shrimp with values 0.90, 0.89 and 0.88, respectively indicating that more than 88% of variation of the data explained by the models. The crude protein and crude fat contents of whole shrimp expressed on dry matter basis were 3-5 times higher than those found in exuvia. However, the exuvia of the shrimp had more than twice the dry matter and about four times ash contents when compared to that of whole shrimp body. The amino-acid composition of the whole shrimp body and the exuvia expressed on dry matter basis were 63.37 and 11.94 per cent, respectively. Glutamic acid, leucine and aspartic acids in the whole shrimp and glutamic acid, glycine and alanine in the exuvia were the predominant amino acids.

The integument of crustaceans consists of basement membrane, epidermis and cuticle. The cuticle of crustacea acts as an exoskeleton providing mechanical support for their body preventing from physical injury and protecting them against extreme ionic and osmotic changes (Travis, 1960). This exoskeleton is periodically shed through the process of molting. Molting is one of the most important physiological processes for crustacean growth since increase in body size occurs in a series of steps associated with the castings of the old exoskeleton (Passano, 1960). Frequency of molting depends not only on species but also on the growth stages. Penaeids molt more frequently, once every few days or weeks while, large

crustaceans such as crabs and lobsters may molt only once or twice a year in their adult stages (Dall *et al.*, 1990).

The molt of crustacea consists of both organic and inorganic materials such as protein, chitin and minerals (Travis, 1960) and their composition varies greatly from species to species (Welinder, 1974). As a result of molting, crustaceans loose considerable amounts of these nutrients and these should be replenished.

Crustaceans are having the habit of eating their exuvia after ecdysis (Forster, 1976; Thomas *et al.*, 1984). If this nutrient source is not accounted for, it could lead to errors in nutritional studies, especially those relating to

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dietary requirements of protein, amino acids and minerals. If the composition of this exuvia is known, the amount of these nutrients provided by ingested molts can be determined. Most of the works on these aspects are limited to relationships between exuvia and the total body length and weight only (Dall *et al.*, 1990). The present study is aimed at determining the biochemical composition along with amino acid profiles of exuvia and the whole body in the Indian white shrimp, *F. indicus*.

Juveniles of *F. indicus* weighing 4.5 g  $\pm$  1.42 (mean  $\pm$  S.D.) were obtained from the wild and maintained in a 1000 l fibre reinforced plastic tank. One shrimp per tank were randomly distributed in twelve 50 l fibre reinforced plastic tanks. During the experimental period, the water temperature was 29  $\pm$  1°C, salinity at 33 °/ 00 and dissolved oxygen 7 mg/l. Eighty per cent of the water in the tanks was exchanged daily. The animals were fed *ad libitum*, twice a day with a standard formulated feed having 39.5 % crude protein.

The molting of the shrimp was closely monitored and the exuvia and freshly molted shrimp were removed from the tanks. Empty tanks were cleaned and restocked with new, randomly selected shrimps from holding tank. The trial was continued for 30 days to collect exuvia and molted shrimp. Three shrimps from holding tank and exuvia samples were analyzed for biochemical composition (AOAC, 1990). Amino acids were analyzed after sealed tube hydrolysis with 6 N HCl (Spackman et al., 1958; Finlayson, 1964) in Shimadzu HPLC system using dye as fluorescent detector post-column derivatization with O-Pthaladehyde. Each analysis was done in duplicate.

The functional relationship between

shrimp weight and exuvia weight was determined by the following regression equation

Where  $Y_i$  is the response variable,  $X_i$  is a regressor variable,  $\beta_0$  and are unknown parameters to be estimated, and is an error term (Gomez and Gomez, 1984). As mentioned above, regression function is fitted to the data. The following stochastic regression equations are considered.

$$DEW = aWSW + \varepsilon$$
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$$DEW = aDSW + \varepsilon$$

Where WEW is wet exuvia weight (g), WSW is wet shrimp weight (g), DEW is dry exuvia weight (g), DSW is Dry shrimp weight (g), 'a' is parameter to be estimated and  $\varepsilon$  is error term. The constant term is not included in all the equations 2, 3 and 4 because if input equals to zero then one would expect the output to be zero as well.

The average weight of shrimp and exuvia were 4.50  $\pm$  1.42 and 0.22  $\pm$  0.07 g, respectively indicating that the wet weight of exuvia was  $4.8 \pm 0.46\%$  of the wet weight of the whole shrimp body. The dry exuvia was  $11.74 \pm 1.13\%$  of the dry weight of the whole shrimp body. Kibra (1993) reported that the loss of weight immediately following ecdysis averaged 4.9% in Penaeus monodon, which is linked with the loss of the exuvia. The morphometric relationships (Fig.1 and Fig.2) between wet and dry weight exuvia and wet shrimp weight are highly significant ( $R^2 = 0.90$ , P < 0.001) indicating strongest relationship in both the cases. Regression calculations

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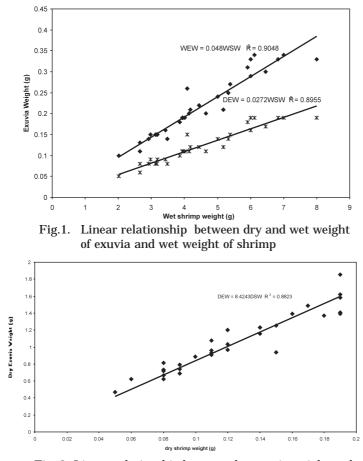


Fig. 2. Linear relationship between dry exuvia weight and Dry shrimp weight

reported earlier (Sarac *et al.*, 1994) in *P. monodon* on wet exuvia weight-wet body weight, dry exuvia weight-dry body weight and dry exuvia weight-wet body weight were 0.75, 0.78 and 0.83, respectively. The relationship between dry weight of exuvia and dry shrimp weight shown similar results ( $R^2 = 0.88$ , P < 0.001). Comparable relationship between the dry weight of whole prawn body and exuvia ( $R^2 = 0.97$ , P < 0.001) was reported by Hewitt (1984) in the freshwater prawn, *Macrobrachium australiense*.

The dry matter content of whole shrimp body was  $23.15 \pm 0.16$  per cent.

Alava and Pascual (1987) and Sarac et al. (1993) also found that the dry matter was in the range of 23.4 - 26.7 per cent for juvenile P. monodon. The exuvia had more than half (56.48%) dry matter. Similarly, high dry matter content was reported in other crustacean species, Cancer magister (Allen, 1971). The crude protein value observed (69.94%) in *F. indicus* in this study was within the range of crude protein reported for juveniles of P. indicus (59.9-64.0), P. monodon (61.8 - 75.81) and Metapenaeus monoceros (62.1-67.81) (Vattheeswaran and Ahamad Ali, 1986; Ahamad Ali, 1996; Alava and Pascual, 1987; Sarac et al., 1993). Alava and Lim (1983) reported that crude protein levels in the body of juvenile P.

*monodon* were directly related to dietary protein levels. Variations in dietary energy levels significantly influenced the total body lipid levels (5.1-10.7%) of juvenile *M. monoceros* (Kanazawa *et al.*, 1981).

The exuvia of the shrimp had more than twice the dry matter and about four times ash contents when compared to that of whole shrimp body (Table1). However, crude protein and crude fat contents of whole shrimps expressed on dry matter basis were 3-5 times higher than those found in exuvia. The crude protein recorded in exuvia (23.11%) in *F. indicus* is within the range (20 – 40%)

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| Composition(%) | Shrimp body      | Exuvia            | Ratio * (%)      |  |
|----------------|------------------|-------------------|------------------|--|
| Dry matter     | $23.15 \pm 0.16$ | $56.48 \pm 0.84$  | $11.74 \pm 1.12$ |  |
| Crude protein  | $69.94 \pm 0.14$ | $23.11 \pm 0.218$ | $3.87 \pm 0.37$  |  |
| Ether extract  | $5.48\pm0.08$    | $1.09 \pm 0.14$   | $2.34 \pm 0.22$  |  |
| Ash            | $16.68 \pm 1.01$ | $63.01 \pm 1.48$  | $44.34 \pm 4.26$ |  |

# On Dry matter basis except for dry matter

\* Component in wet exuvia (g) \_\_\_\_\_ X 100

Component in wet shrimp (g)

 $T_{ABLE} \ 2: \ Amino \ acid \ composition \ of \ whole \ body \ and \ exuvia \ F. \ indicus.$ 

| Amino acids | Whole shrimp body        |                             | Exuvia                   |                             | Ratio <sup>*</sup> of AA<br>in wet exuvia<br>to AA in wet<br>shrimp body |
|-------------|--------------------------|-----------------------------|--------------------------|-----------------------------|--|
|             | % Dry<br>matter<br>basis | Ratio to<br>total<br>AA (%) | % Dry<br>matter<br>basis | Ratio to<br>total<br>AA (%) | 1 3  |
| ASP         | 5.83                     | 9.20                        | 1.01                     | 8.46                        | 2.04   |
| THR         | 2.66                     | 4.2                         | 0.6                      | 5.03                        | 2.65   |
| SER         | 2.51                     | 3.96                        | 0.57                     | 4.77                        | 2.67   |
| GLU         | 10.11                    | 15.95                       | 2.51                     | 21.02                       | 2.92   |
| PRO         | 3.91                     | 6.17                        | 0.75                     | 6.28                        | 2.25   |
| GLY         | 5.33                     | 8.41                        | 1.1                      | 9.21                        | 2.42   |
| ALA         | 4.02                     | 6.34                        | 1.05                     | 8.79                        | 3.07   |
| CYS         | 0.6                      | 0.95                        | 0.08                     | 0.67                        | 1.57   |
| VAL         | 2.94                     | 4.64                        | 0.66                     | 5.53                        | 2.64   |
| MET         | 1.66                     | 2.62                        | 0.11                     | 0.92                        | 0.78   |
| ISO         | 3.08                     | 4.86                        | 0.28                     | 2.35                        | 1.07   |
| LEU         | 6.14                     | 9.69                        | 0.6                      | 5.03                        | 1.15   |
| TYR         | 2.29                     | 3.61                        | 0.53                     | 4.44                        | 2.72   |
| PHE         | 2.66                     | 4.20                        | 0.8                      | 6.70                        | 3.53   |
| HIS         | 2.01                     | 3.17                        | 0.11                     | 0.92                        | 0.64   |
| LYS         | 4.21                     | 6.64                        | 0.33                     | 2.76                        | 0.92   |
| ARG         | 3.41                     | 5.38                        | 0.85                     | 7.12                        | 2.93   |
| TOTAL       | 63.37                    | 100                         | 11.94                    | 100                         | 2.21   |

\* Amino acid in wet exuvia (g)

Amino acid in wet shrimp (g)

reported earlier (Dall et al., 1990). The crude protein of exoskeleton decreased from 53.5%, just after molting to 30.8%, after14 days in Astacus fluviatilis (Welinder, 1975). Exuvia contain up to 20% chitin (Welinder, 1975) which is a non- protein compound containing 7% N (Muzzarelli, 1977). Further, the exuvia also contains other non-protein nitrogen compounds such as chitosan. Because of this true protein content in exuvia is very less as represented by low amino acid content (Table 2). The ash content of dry exuvia is 63.01%, which is mainly present in the form of calcium salts (Welinder, 1975). Since calcium is a major part of the exuvia, shrimps require high levels of calcium intake, especially after molting. This requirement seems to be met mainly by absorbing Ca available in the surrounding water (Davis and Gatlin, 1991).

The amino-acid composition of the whole shrimp body and the exuvia expressed on dry matter basis is presented in Table 2. Glutamic acid was the predominant amino acid in whole shrimp body (10.11%) and in exuvia (2.51%). This was followed by leucine (6.14%) and aspartic acid (5.83%) in shrimp body and glycine (1.10%) and alanine (1.05%) in exuvia. The amounts of sulphur-containing amino acids, cystine and methionine, were low in the exuvia (0.08 and 0.11%, respectively). The essential and non-essential amino acids constituted 28.72 & 34.65 and 4.34 & 7.6 per cent, respectively in the whole body and in the exuvia. The total amino acids in exuvia had 2.21% of that of the whole shrimp. Glutamic acid, leucine and aspartic acids were also the same three amino acids predominant in Penaeus aztectus (Shewbert et al., 1972). The exoskeleton amino acid reported in Cancer magister by Allen (1971) were valine (1.84%), aspartic acid (1.14%), glutamic acid (1.03%), glycine (0.91%) and alanine (0.69%). The amino acid composition of the exuvia of F. indicus is similar to those reported for other penaeid shrimps (Sameshima et al., 1973; Welinder, 1974). The loss of essential and non-essential amino acids through ecdysis in *F. indicus* is 1.81% and 2.45%, respectively. Loss of these nutrients in *P. monodon* was 1.51% and 1.92%, respectively (Sarac et al., 1994). The study revealed that there is a strong relationship between the weights of exuvia and shrimp body weight. The proportion of dry matter and ash were more in exuvia compared to whole shrimp body, whereas it is vice - versa in case of protein and lipid proportions. The exuvia contains considerable amounts of crude protein and 2.21% of amino acids when compared to that of shrimp. Nutrient consumption from exuvia could be estimated using the regression equations calculated in the present study especially in dietary nutrient requirement studies.

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