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 Aquafeeds

A look at protease enzymes in crustacean nutrition

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A better understanding of enzyme supplementation in aquafeeds is essential



The enzyme composition and their interaction with the environment and diets can significantly influence the digestive capacity and immune response of shrimp.

Digestion, a complex physiological process, depends on molecular activation and recognition and hydrolysis of food in specific conditions. The process involves digestive enzymes for breaking down polymeric macromolecules into smaller building blocks facilitating nutrient absorption.

In crustaceans, life stage, age, molting stage, circadian cycle, feed composition and environmental conditions all affect digestive enzyme production. Among these enzymes, proteases are the most important ones responsible for hydrolyzing dietary proteins and breaking them into smaller peptides and free amino acids. The protease activities in the digestive tract of an animal are induced by feeding, and are usually affected by genetic features of the individual, age, life stage and the culture environment.

Effect of environment and diet on digestive enzymes

Environment, diets and their interactions appeared to affect the enzyme production in all animals. The species-specific differences in the optimum pH and temperature range of proteolytic enzymes of aquatic animals can be attributable to their habitat, food and genotypic and phenotypic differences. For example, the optimum pH range in crustaceans usually ranges between 7.0 and 8.0 for trypsin-like enzymes, and for chymotrypsin-like enzymes, from 7.5-8.0 for the São Paulo shrimp (*Farfantepenaeus paulensis*). Whereas, in the marine, edible crab (*Cancer pagurus*) it ranges from 8.0 to 10.0.

Differences in proteolytic activity can be significant for shrimp reared in culture environments with large variations in quantity and quality of natural food. For example, Moss et al. (2001) reported significantly higher activity of serine protease (71648-unit vs 30200-unit/mg soluble protein), collagenase (653-unit vs 312-unit/mg soluble protein), and acid phosphatase (189-unit vs 83-unit/mg soluble protein) in Pacific white shrimp cultured in pond water than those in well water. Higher enzymatic activity in shrimps in pond water indicates the presence of natural food in the system. Interestingly, among



Shrimp have about 14 different types of digestive proteolytic enzymes, with varying degrees of temperature and pH optima.

digestive enzymes, only the chitinase activity was higher in shrimp reared in clean water, indicating that the animals were feeding on molted shells, high in chitin.

Dietary protease, a solution in hand

High cost and limited availability of fishmeal and other balanced protein sources are frequently forcing aquafeed manufacturers in many regions to rely on low-cost but poorly digestible alternatives. At the same time, there is increasing pressure to lower environmental and economic costs, putting a serious onus on feed manufacturers to improve nutrient digestibility of aquafeeds while lowering the manufacturing cost.

Better nutrient digestibility

Davis et al. (1998) tested the apparent digestibility of dry matter (ADCDM) and crude protein (ADCCP) in Pacific white shrimp (*Litopenaeus vannamei*) fed diets supplemented with graded levels (0, 0.2 or 0.4 percent) of a protease, and reported that the ADC of CP (74.3 percent) in shrimp fed 0.4 percent protease diets was higher than for those animals fed the control (65.3 percent) or other diets. In a subsequent trial, these authors reported poor growth performance (weight gain, feed efficiency, and protein efficiency ratio) in shrimp fed diets with the protease.

Better performance and immune response

Recently, Li et al. (2015) reported significant increases in weight gain (288 percent) and feed conversion (1.29) for Pacific white shrimp fed low-fishmeal diets (38 percent CP, 22 percent fishmeal) supplemented with 175 mg/kg of a commercial multi-component protease (Jefo Nutrition Inc., Canada) compared to those animals fed the same diet without the supplementation (275 percent and 1.42, respectively). The authors suggested the possible use of protease in aquafeed to reduce or replace expensive protein sources with poorly digestible and less-expensive ingredients without affecting the performance of the cultured animals (Table 1).

Chowdhury, Table 1

Parameters	Positive control (PC) 40% CP, 25% FM	Negative control (NC) 38% CP, 22.5% FM	NC + 175 mg/kg protease 38% CP, 22.5% FM
FBW, g	12.9 +/- 0.6b	11.5 +/- 0.5a	12.4 +/- 0.2b
FCR	1.14 +/- 0.05a	1.28 +/- 0.05b	1.16 +/- 0.04a
HP_PROT, U/g tissue	3665 +/- 203b	3257 +/- 192a	3788 +/- 167b
IN_PROT, U/g tissue	520 +/- 57b	499 +/- 30a	527 +/- 37b

Final body weight (FBW), feed conversion ratio (FCR) and digestive protease activities in hepatopancreas (HP_PROT) and intestine (IN_PROT) in Pacific white shrimp fed high and low fishmeal (FM) diets where the low-

FM diet was supplemented with a protease [from Li et al. (2015)].

Song et al. (2016) also reported significantly better weight gain and FCR in shrimp fed low-fishmeal (LFM, 10 percent) diets supplemented with the protease, compared to the same diet without supplementation. They also reported significantly higher serum phenol oxidase concentration and lower malonaldehyde concentration in both serum and hepatopancreas in shrimp fed the protease supplemented diets. The authors also conducted a disease challenge test with a pathogenic strain of *Vibrio parahaemolyticus* and reported a linear decrease in 96 hours cumulative mortality of shrimp fed diets with 0, 125, 150 and 175 ppm of the protease (Table 2).

Chowdhury, Table 2

Parameters	Positive control (PC) 20% FM	Negative control (NC) 10% FM	NC + 125 mg/kg protease	NC + 150 mg/kg protease	NC + 175 mg/kg protease
FBW, g	12.3 +/- 0.3d	11.0 +/- 0.3a	11.3 +/- 0.3ab	11.7 +/- 0.5bc	12.0 +/- 0.5cd
Trypsin, U mg ⁻¹ PROT	4927 +/- 21e	2106 +/- 59a	3419 +/- 192b	4048 +/- 137c	4576 +/- 141d
PO_S, U mL ⁻¹	672 +/- 11bc	525 +/- 14a	605 +/- 55b	691 +/- 4c	890 +/- 63d
MDA_S, nmol/mL	12 +/- 1.1b	16 +/- 0.2d	14 +/- 0.1d	13 +/- 0.6c	9 +/- 0.2a
PO_HP, U/mL	1028 +/- 13c	701 +/- 30a	739 +/- 52a	833 +/- 11b	1032 +/- 35c
CMR_96h	0.31 +/- 0.09a	0.54 +/- 0.04b	0.46 +/- 0.05b	0.40 +/- 0.04ab	0.32 +/- 0.06a

Final body weight (FBW), endogenous trypsin activity, phenol oxidase activity in serum (PO_S) and hepatopancreas (PO_P), malon-dealdehyde activity in serum (MDA_S), and 96-hour cumulative mortality (CMR_96h) under a disease challenge test (a pathogenic strain of *Vibrio parahaemolyticus*) of Pacific white shrimp fed a high-fishmeal (FM) diet and a low-FM diet supplemented with graded level (0, 125, 150 and 175 mg/kg diet) of a protease [from Song et al. (2016)].

Perspectives

Ingredients, diets, age, size, molting stage and culture environment (such as pH, temperature and dissolved oxygen levels) can significantly affect digestive enzyme production in crustaceans and other animals. The complex nature of diet and environment, and their effects on digestive enzyme production, requires a better understanding of these enzymes. There are about 14 different types of digestive proteolytic enzymes reported in shrimp, with varying degrees of temperature and pH optima. The composition of these enzymes and their interaction with environment and diets can significantly influence the digestive capacity and immune response of the shrimp.

Ingredients, diets, age, size, molting stage and culture environment (such as pH, temperature and dissolved oxygen levels) can significantly affect digestive enzyme production in shrimp, and other crustaceans and animals.

Dietary enzymes are a major alternative to improve feed quality and for better nutrient digestibility, and also to reduce organic wastes. Additionally, enzyme supplementation can improve gut health, compensate digestive enzymes when needed, and may also improve immune responses.

However, a large majority of the stakeholders remains skeptical because of issues with heat stability, interactions with other enzymes, safety of recommended dosages and insufficient knowledge on suitable substrates for enzymes. Our lack of understanding on suitable substrates often leads to unnecessary or improper use of some enzymes under various brands of multi-enzyme complexes, where proteases, carbohydrases, phytase – and sometime even lipases – are combined.

It is important to understand that each specific enzyme usually attacks specific active sites in a complex molecule. For example, an enzyme containing a single protease enzyme may be only suitable for specific substrates or ingredients. On the other hand, an enzyme containing multiple protease activities can be suitable for multiple substrates and can react positively with a wide range of raw materials. With the omnipresence crisis for quality raw materials, an industry-wide understanding of dietary enzyme application is essential, not only for shrimp aquafeeds but also for aquafeeds in general.

Further research is needed to understand the effects of different dietary proteases on the quality of proteins in feed, their interactions with various ingredients, gut health and nutrient utilization. The appropriate application of the dietary proteases available in the market has also been an issue because of the heat stability concerns and fear of destruction during manufacturing. Some better heat-stable enzymes are available in dry powder form, can be readily added to the mixer during aquafeed manufacturing, and can keep most of their activity intact. Other enzymes are either coated or available in liquid form for post-pelleting application requiring costly mechanical tools.

We also need a better understanding of the advantages and disadvantages of the various application methods when selecting an enzyme. The ability to apply enzymes directly to the mixer may have several advantages. In addition to the ease of handling, the protease may also alter the quality of feed proteins during cooking or conditioning, resulting in better quality aquafeeds. And a partial coating could provide a better solution to improve heat stability.

As the cost and availability of quality protein sources are increasingly becoming an issue, the use of dietary proteases could provide a better solution to the sustainability of the aquaculture industry. However, we are still in the infancy of our understanding of the complexity of the protease world. A renewed and targeted focus in our research agenda is, therefore, strongly recommended to fully utilize the potential of protease enzymes.

All references available from author.

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