

Chemoattraction of Low-Molecular-Weight Compounds in Shrimp Feeds

Summary:

The methodology used in a Texas Agricultural Experiment Station study provided a useful tool for evaluating the attractiveness of individual ingredients in prepared shrimp feeds. Although not as effective an attractant as the squid muscle meal control, putrescine was the highest rated of eight chemoattractant compounds tested.

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Marine shrimp possess the ability to detect and differentiate among a multitude of environmental chemical stimuli. Whether olfactory, gustatory, or both, this ability is referred to as chemoreception. Chemoreception enables shrimp to identify potential predators, mates, symbionts, changes in the chemical makeup of their environments, and sources of food.

Due to the high cost of feed, rapid food identification and ingestion by cultured shrimp is of major economic importance. Further, since 50-100% of the low-molecular-weight, water-soluble compounds in feeds can leach out within one hour after the feed is added to culture water, it is critical that feed is quickly ingested for greater nutrient quality and reduced potential water pollution.

Low-Molecular-Weight Compounds

Most chemoattractants for shrimp are metabolites with low molecular weights below 1,000 daltons. They include amino acids, quaternary ammonium compounds, nucleotides, nucleosides, and organic acids. As a group, low-molecular-weight substances are water-soluble and occur in tissues at concentrations many orders of magnitude higher than those in the aquatic environment.

The introduction of these substances into the aquatic environment from the tissues of prey species can be caused by leakage, excretion, tissue damage, or decomposition. Assimilation of these low-molecular-weight compounds by bacteria as well as higher organisms serves to keep ambient concentrations low.

Evaluating Chemoattractants

The development of methodology to evaluate the effectiveness of one chemoattractant relative to another is an important step in reducing the amount of feed used in

commercial aquaculture. The process of ranking chemoattractants can be achieved by assigning data to an ordinal scale or by transforming discrete data to a hypothesized underlying scale. Although ordinal methods can be very useful with regards to ranking, they lack the ability to measure the magnitude of difference between the objects ranked.

Furthermore, depending on experimental design, ordinal ranking scales require assessors to learn specific scoring systems that can be dependent on previous questions or observations, thus placing demands on the assessors' experience and training. In contrast, paired comparison tests only require that assessors choose which of two objects is greater with respect to a predefined attribute.

Counting the number of shrimp present on a feeding tray at each observation is a useful paired comparison. Transforming this discrete data to a log-linear scale provides an estimated magnitude of difference between the two objects compared.

Individual ingredients can also be indirectly ranked relative to one another within an experiment by conducting pairwise comparisons of all possible choice combinations. This can be very useful when evaluating potential chemoattractants and their levels of inclusion with respect to cost as well as efficacy.

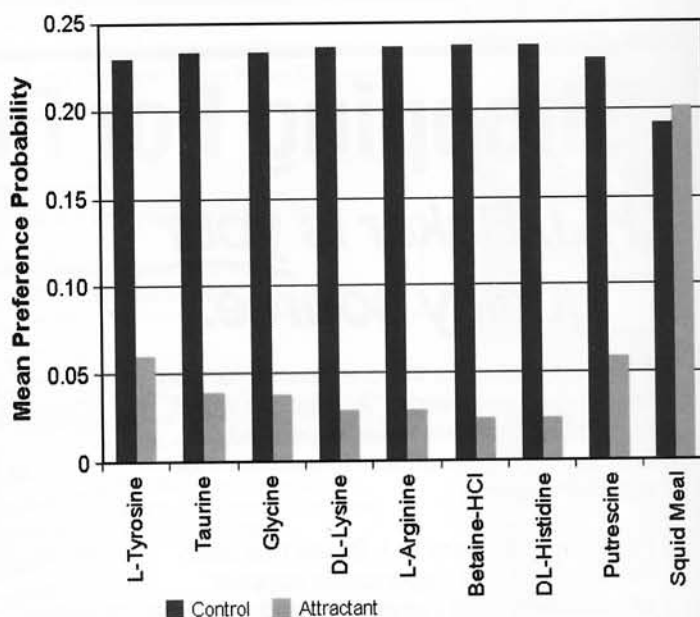


Figure 1. Mean preference probabilities of low-molecular-weight compounds compared to a common control containing squid muscle meal.

TAES Study

In May 2003, a 19-day study of chemoattractants at the Texas Agricultural Experiment Station of Texas A & M University in Port Aransas, Texas, USA, used an experimental system consisting of nine fiberglass tanks 3.4 m in diameter. Each tank was stocked with 50 pacific white shrimp, *Litopenaeus vannamei*, with initial weights of about 7.6 g.

None of the prepared feeds containing low-molecular-weight chemoattractants were as attractive to *L. vannamei* as the control feed, which contained squid muscle meal.

Rankings indicate most attractive to least attractive. Based on pairwise comparisons of all possible combinations of exponentiated differences in mean log-relative strength. Chemoattractants that share identifiers are not significantly different.

Table 1. Numerical ranking of compounds evaluated as chemoattractants.

Rank	Identifier	Chemoattractant	a = 0.05
1	a	Squid muscle meal	a
2	b	Putrescine	b, c, d, e
3	c	L-tyrosine	b, c, d, e
4	d	Taurine	b, c, d, e, f, g
5	e	Glycine	b, c, d, e, f, g
6	f	DL-lysine	d, e, f, g
7	g	L-arginine	d, e, f, g, h
8	h	Betaine-HCl	g, h, i
9	i	DL-histidine	g, h, i

Each tank contained two sets of two 0.6-m² black, gel-coated fiberglass feed trays, with each set placed against opposite tank walls. Experimental feeds were formulated to contain 10% crude protein and 4% low-molecular-weight compound on a dry-weight basis. The compounds tested were L-tyrosine, taurine, glycine, DL-lysine, L-arginine, betaine-HCl, DL-histidine, and putrescine.

Each experimental feed was added to the feed trays at a rate of 6% of shrimp body weight. The number of shrimp on each set of trays was observed at 15-minute intervals over two hours. Applied every other day, experimental feeds were rotated through all nine tanks, and tray positions were alternated each trial day to reduce within-system variability.

On days opposite the chemoattractant trials, a single feeding of commercial shrimp feed was hand-broadcast into the tanks at a rate of 5% of the shrimp wet body weight. The average shrimp growth during the experiment was about 1.05 g/week.

For this experiment, the basic Bradley-Terry model of paired comparisons was extended to 3-choices to account for the entire population of each experimental tank at each 15-minute observation time.

Results

The study showed that none of the prepared feeds containing low-molecular-weight chemoattractants were as attractive to *L. vannamei* as the control feed, which contained squid muscle meal (Figure 1). However, some of the compounds were significantly more attractive than others (Table 1).

For example, putrescine was not significantly more attractive than L-tyrosine. However, it was significantly more attractive than DL-lysine, L-arginine, betaine-HCl, and DL-histidine.

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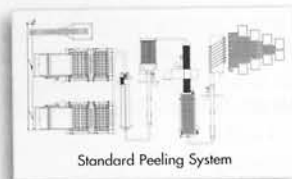


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