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Research Article

Effects of Crab Shellmeal Inclusions to Fishmeal Replacement on the Survival, Growth, and Feed Utilization of Mangrove Crab *Scylla serrata* (Forsskal 1775)

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Abstract: Mangrove crab *Scylla serrata* is associated with mangroves throughout the Indian and Pacific oceans. This species is crucial to aquaculture and fishing. As wild stocks decline and demand rises, mangrove crab aquaculture has become increasingly popular. However, feed development research and its quality are still meager in the industry. This study examined the interactive effects of different levels of crab shellmeal (CSM) to fishmeal (FM) replacement on proximate composition, feed utilization, carcass composition, growth, and survival performance of mangrove crab *S. serrata*. Four formulated diets were prepared, and one for chopped trash fish (TF) supplement: 30% FM and 0% CSM (Diet 1) as a negative control, 20% FM and 10% CSM (Diet 2), 10% FM and 20% CSM (Diet 3), 0% FM and 30% CSM (Diet 4), and TF as a positive control (Diet 5). Experiments were conducted in each group for 30 days with ten replicates. Results revealed that formulated diets using different levels of CSM and FM did not significantly affect mangrove crabs' growth and survival rates as well as feed utilization. However, the proximate composition of Diet 4 was significantly higher among other experimental diets. Moreover, the crab's whole body composition (ash, moisture, carbohydrates, crude protein, crude fat, and calories) with different levels of CSM and FM was significantly improved. Hence, it is possible to enhance the carcass composition and proximate composition by supplementing CSM; however, it has no effect on feed utilization, as well as the growth and survival rates of mangrove crab *S. serrata*.

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1. Introduction

Mangrove crab *S. serrata* (Forsskal, 1775), also known as green crab or mud crab, is one of the largest portunid species that lives along the coast and in mangrove swamps all along the Indo-Pacific coastline (Barnes et al., 2002; Le Vay et al., 2007; Shelley and Lovatelli, 2011). It has become increasingly popular in aquaculture due to its economic importance and is currently admired seafood all over the world because of its savory sweet flesh, larger size, and nutritional value (Sathiadhas and Najmudeen, 2004; Ali et al., 2011). As a matter of fact, mangrove crab is one of the most commonly

used ingredients in a number of dishes (Mirera, 2011). Together with shrimp and mollusks, it already ranks among the top most in-demand aquaculture products globally (Sathiadhas and Najmudeen, 2004). In 2020, crustacean production accounted for nearly 11 237 metric tons in world aquaculture, of which 159.4 metric tons came from crabs (FAO, 2022). Furthermore, aside from the delicious and nutritious crab meat, its shells are also demanded in the pharmaceutical industry for its high levels of chitin, which has a variety of applications such as water-repellent adhesive, oral mucoadhesive liquid, and wound dressings (Kato et al., 2003). Aquaculture is also exploring the possibility of using shells as a formulated feed ingredient because they contain chitin like krill (Ringo et al., 2012).

Mangroves' crabs are initially harvested in the wild (Agbayani, 2011; Mirera, 2011; Qunitio, 2017; Hasanuzzaman et al., 2014; Basu and Roy, 2018). Mangroves provide a high level of salinity and temperature and protect young against cannibalism and predators, which is why these species thrive in these areas (Baylon, 2010; Alberts-Hurbatsch et al., 2014). In addition to its undeniable importance to society, it is an integral part of the mangrove ecosystem since it is critical in nutrient cycling and energy transfer (Mirera, 2011). Thus, wild mangrove crabs are increasingly being collected, leading to increased concern that a decrease in their abundance might negatively affect their ecosystems (Alberts-Hurbatsch et al., 2016). Furthermore, habitat destruction has been a serious concern resulting from the hunting of these wild species (Shelley, 2008). Due to high demand in the market, even juveniles of mangrove crabs (crab farming's main source of crab seeds) are still extracted despite rising concerns (Lindner, 2005). Due to its higher survival rate and resistance to disease than shrimps, the mangrove crabs trade is a reliable source of income and is also a good alternative for shrimp farmers (Marichamy and Rajapackiam, 2001). As a result, commercial mangrove crab culture initiatives are being developed (Williams and Field, 1999). Different techniques have been developed to achieve the best possible method for fattened crabs to survive and grow (Shelley, 2008; Agbayani, 2011; Petersen et al., 2011; Qunitio, 2017). In addition, there is an ongoing effort to develop alternative feeds that would produce better quality mangrove crabs, both in size and consistency (Alber, 2003).

Mangrove crabs consume fish, mollusks, crustaceans, sand, and detritus (Viswanathan and Raffi, 2015). However, as the culture of this product has grown, the supply of feeds has become a major problem (Alber, 2003). Particularly, trash fish, one of the fishmeal feeds to mangrove crabs, have competing demands with the growing human food consumption (Gasco et al., 2018). Therefore, mangrove crab culture may not be worthwhile due to the high prices of trash fish in response to the competing demand over the years. As a result of this problem, researchers have worked on formulating alternative feeds to fatten mangrove crabs, such as plant-based, algae, animal-based, insects, and crustacean feeds (Alber, 2003; Krogdahl, 2016; Gasco et al., 2018). Crab shellmeal has been used as a replacement for fishmeal in poultry and livestock (Vijayalingam and Rajesh, 2020). Due to the high chitin content of crustacean by-products such as shrimp heads and shells and crab shells, the use of these products in feed has gained an increasing amount of attention in recent years (Gasco et al., 2018). In food and pharmaceutical manufacturing, chitin is a biopolymer with various uses (Kato et al., 2003). Toppe et al. (2006) stated that crab by-products could be used as a feed in diets for the fish Atlantic cod, *Gadus morhua*. In crustaceans, such as prawn *Macrobrachium rosenbergii*, chitin diets are shown to advance growth (Kumar et al., 2006). In addition, increasing survival with chitin diets has been linked to removing potentially pathogenic bacteria from adult male shore crab *Carcinus maenas* (Powell and Rowley, 2007). However, researchers are still investigating whether crab shells can be used as a feed ingredient in aquaculture (Ringo et al., 2012). Considering this, the study examined the effectiveness of formulated feed incorporated with crab shells as a meal for mangrove crab *S. serrata* on survival, growth, proximate composition, feed utilization, and carcass composition.

2. Material and Methods

2.1. Study site and duration

This study was conducted at the Multi-Species Hatchery of the College of Fisheries, Mindanao State University Tawi-Tawi College of Technology and Oceanography (MSU-TCTO), Sanga-Sanga, Bongao Tawi-Tawi, Philippines (Figure 1) for a duration of 30 days.

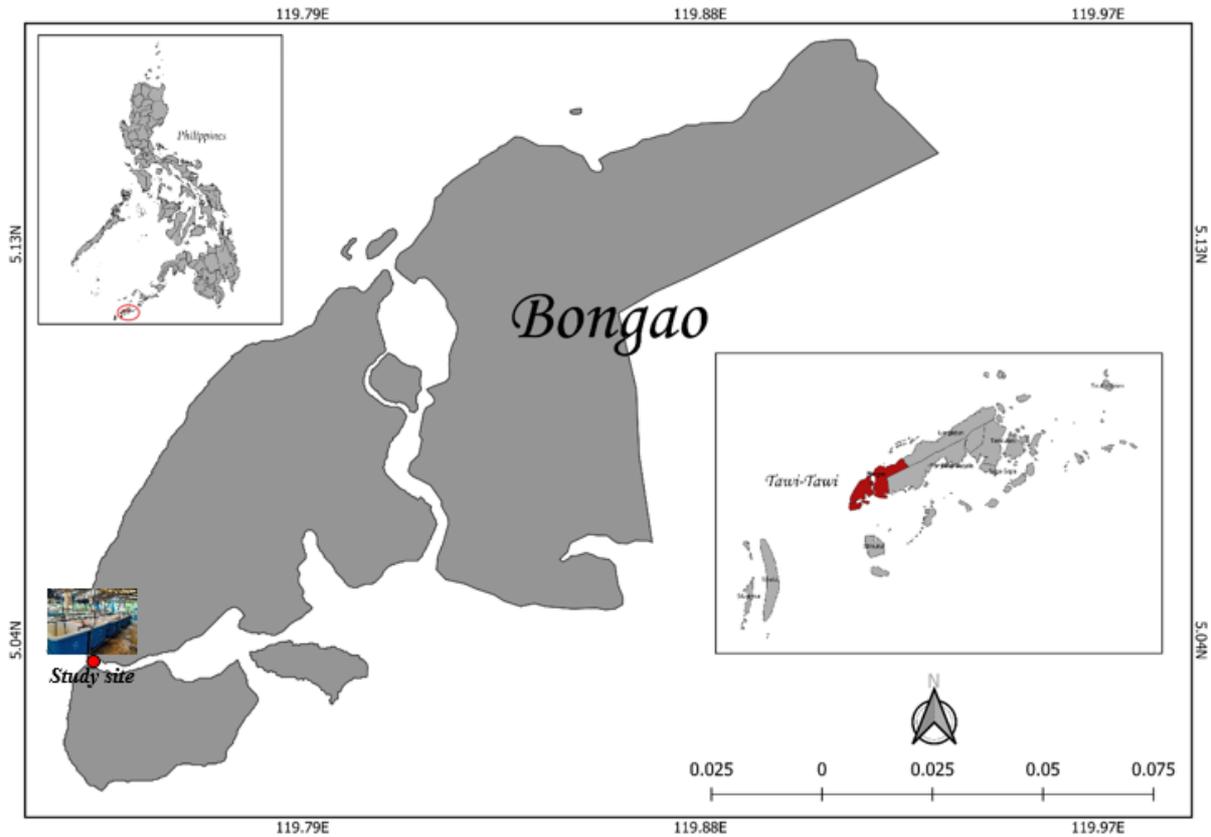


Figure 1. Map of the study site.

2.2. Source of crab shells and feed ingredients

Blue swimming crab (*Portunus pelagicus*) shells were used as the source of powdered crab shells from the carapace and pincers only because these parts of crab shells could be cleaned and powdered easily. These were collected from Crab Meat Processing Plant in Tubig Boh, Bongao, Tawi-Tawi, Philippines. Roughly 1.4 tons (wet weight) of crab shells (Figure 2) were discarded monthly by this processing plant (Pers. Com.). In addition, the feed ingredients used in the experiment were purchased from Bongao Public Market, as well as the fresh *Sardinella* sp. that was used for the positive control diet.



Figure 2. Discarded crab shell in Crab Meat Processing Plant, Bongao, Tawi-Tawi, Philippines.

2.3. Experimental Species

Mangrove crab *S. serrata* was used as an experimental animal, purchased from fisherfolk in Panglima Sugala, Tawi-Tawi, which was previously collected from mangrove areas. The crabs were transported by pump boat to Bongao, Tawi-Tawi, and landed at the MSU-TCTO Multi-purpose Hatchery.



Figure 3. Experimental animals.

2.4. Preparation of powdered crab shells

Blue swimming crab (*P. pelagicus*) shells were cleaned thoroughly and air-dried within five days until they could be crushed easily and ground using a grinder. The ground crab shells were sieved, packed in ziplock plastic, and stored at room temperature to be used to formulate experimental diets.

2.5. Preparation of treatment diets

Table 1 presents the ingredients of the experimental diet. Using locally available ingredients, crab shells were gradually substituted for fishmeal in a feed formulation. They were dried and processed, along with other ingredients for animal feed. The ingredients were powdered in an osteorizer and sieved through a 0.5 mm mesh sieve. In an electric blender, with the binder dissolved in 500 mL of purified water, the ingredients were mixed according to the formula. The mixture was kneaded into a dough and formed into a spherical ball approximately 2.0 cm in diameter by hand (Ali et al., 2011). The spherical balls were steamed at atmospheric pressure for five minutes and dried in an electric oven at 60 °C for 12 hours to remove excess water. The dried feeds were stored in sealed plastic containers until used. There were five diets prepared, four experimental diets incorporated with 0% (negative control), 10%, 20%, and 30% powdered crab shells kg^{-1} , respectively, and positive control (chopped *Sardinella* sp.). All treatment diets were evaluated using proximate analysis, which was done at Davao Analytical Laboratories Inc. (DALI) in Davao City and the Department of Science and Technology (DOST) in Region IX, Zamboanga City, Philippines.

Table 1. Composition of the treatment diets in crab shellmeal as replacement of fishmeal (g kg^{-1})

Ingredients	Inclusion level			
	0% Crab shell	10% Crab shell	20% Crab shell	30% Crab shell
Crab shell meal (g)	0	100	200	300
Fishmeal (g)	300	200	100	0
Copra meal (g)	300	300	300	300
Rice Bran (g)	300	300	300	300
Corn starch (g)	40	40	40	40
Palm Oil (mL)	40	40	40	40
NaCl (g)	19.5	19.5	19.5	19.5
Vitamin Premix (g)	0.5	0.5	0.5	0.5
Total	1 000	1 000	1 000	1 000

2.6. Stocking of crabs

Experimental animals to be considered were a young adult male and female mangrove crabs that have already reached sexual maturity (approximately 150 – 350 grams in weight). Prior to stocking, 50 crabs were used and acclimatized for one week with salinity ranging from 30 to 33 ppt and temperature from 26 to 30 °C. The crabs were subjected to an intensive environment where feeds were provided, the water was replenished, and waste was removed regularly. During the experiments, the crabs were randomly weighed ($207.24 \pm 7.66\text{g}$), assigned, and stocked in 30-L plastic containers (one crab per container). Each treatment had ten replicates.

2.7. Water maintenance

The temperature and salinity of the water were monitored using a thermometer and refractometer. Prior to feeding each day, both parameters were checked before and after changing the water in the plastic containers. Every week, approximately 80% of the water was changed. Siphoning the container every early in the morning was done before changing the water to remove uneaten feeds.

2.8. Feeding

There were five dietary treatments used in the experiment: Diet 1 (30% FM and 0% CSM) as the negative control, Diet 2 (20% FM and 10% CSM), Diet 3 (10% FM and 20% CSM), Diet 4 (0% FM and 30% CSM crab), and Diet 5 (chopped *Sardinella* sp.) as the positive control. The experimental feeds and chopped *Sardinella* sp. were weighed and supplemented based on the weight of each crab per plastic container. Feeding was done once a day in the afternoon with 5% of the average body weight.

2.9. Experimental design

This study used a Completely Randomized Design (CRD) with five experimental diets and ten replicates for each diet (Figure 4). The experiment was conducted using a 50 rectangular clear plastic container (30 L volume capacity) half-filled with seawater with holes on its cover to enable air passage and another hole on its side that serve as excess water passage from the container.

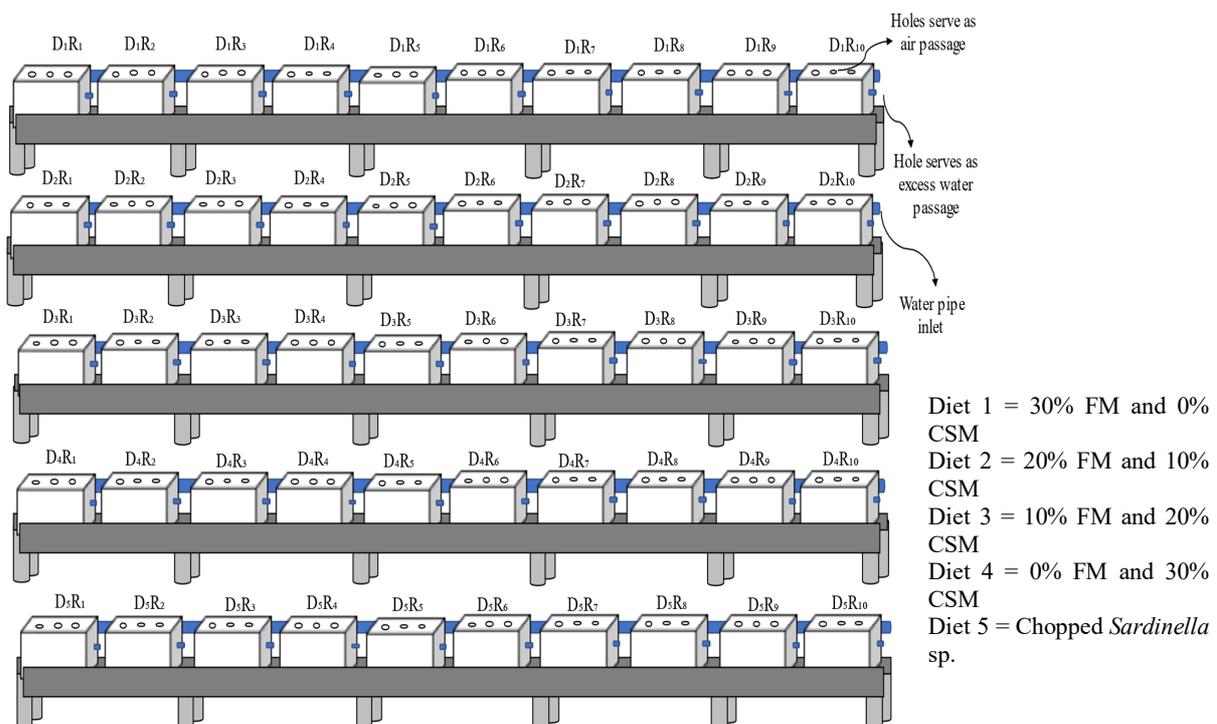


Figure 4. Experimental set-up of the study.

2.9. Sampling

Sampling was done every 15 days to determine the survival and growth rates. Analysis of the body composition was done at the end of the culture period. In order to avoid scaping the crab, a plastic bowl was used, which was weighed using a weighing scale, then tared to zero weight before placing the crab inside. Samples of treatment diets and body composition of experimental animals were sent to DOST, Zamboanga City, and DALI, Davao City, for proximate and carcass analysis such as ash, moisture, carbohydrates, crude fat, crude protein, crude fiber, and calorie. Proximate and carcass analysis was conducted after 30 days of the culture period. The survival rate, specific growth rate (SGR), and feed utilization were computed using the following formula (Kader et al., 2017):

A. Survival rate

$$\text{Survival rate} = \frac{\text{Final number of stocks}}{\text{Initial number of stocks}} \times 100 \quad (1)$$

B. Growth rate

$$\text{WG (g)} = \text{ABWf} - \text{ABWi} \quad (2)$$

$$\text{SGR (\% Day)} = \frac{\ln(\text{ABWf}) - \ln(\text{ABWi})}{\text{DOC}} \times 100 \quad (3)$$

Where: WG = weight gain

ABWf = average body weight final

ABWi = average body weight initial

DOC = days of culture

C. Feed utilization

$$\text{FCR} = \frac{\text{FI}}{\text{WG}} \quad (4)$$

$$\text{PER} = \frac{\text{WG}}{\text{FI}} \times \text{Diet CP} \quad (5)$$

Where: FCR = feed conversion ratio

FI = feed intake

WG = weight gain

PER = protein efficiency ratio

2.10. Statistical analysis

The collected data were presented as mean \pm standard error of the mean (SEM) and subjected to a one-way analysis of variance (ANOVA) using IBM SPSS 20.0 software package to identify significant differences in the different treatments in terms of growth, survival rate, and feed utilization at $\alpha = 0.05$. Levene's Test was used to test for homogeneity of variance, and Duncan's Post-Hoc Test was used to rank the mean.

3. Results

3.1. Proximate composition treatment diets

The proximate composition of treatment diets is shown in Table 2. Levels of ash content and carbohydrates varied from $5.57 \pm 0.12\%$ – $15.23 \pm 0.13\%$ and $1.16 \pm 0.87\%$ – $38.23 \pm 0.27\%$,

respectively, indicating that Diet 4, which contains 30 % crab shellmeal inclusion significantly higher ($p < 0.05$) among all diets. Moisture content ranged from 15.09 ± 0.04 % – 20.38 ± 0.18 %, where Diet 2, Diet 3, and Diet 4 were significantly lower ($p < 0.05$) than Diet 1 and Diet 5. Crude fat levels varied from 9.60 ± 0.06 % – 28.47 ± 0.23 %. Diet 1 was significantly greater ($p < 0.05$) in all diets in terms of crude fat. Levels of crude protein in all experimental diets ranged from 8.93 ± 0.22 % – 57.60 ± 0.70 %, where Diet 5 was significantly higher ($p < 0.05$) among all diets. Crude fiber levels varied from 0.46 ± 0.04 % – 9.79 ± 0.41 %, and Diets 3 and 4 were significantly higher ($p < 0.05$) in all diets.

Table 2. Proximate composition of experimental diets (%)

¹ Diet	Ash	Moisture	Carbohydrates	Crude fat	Crude protein	Crude fiber
Diet 1	5.57 ± 0.12^c	17.30 ± 0.10^b	28.63 ± 0.58^d	28.47 ± 0.23^a	20.03 ± 0.18^b	3.91 ± 0.27^d
Diet 2	9.33 ± 0.03^d	15.25 ± 0.05^d	31.73 ± 0.26^c	25.90 ± 0.17^b	17.77 ± 0.09^c	6.62 ± 0.44^c
Diet 3	11.67 ± 0.03^b	16.90 ± 0.08^c	35.80 ± 0.44^b	24.27 ± 0.43^c	11.30 ± 0.06^d	9.50 ± 0.59^{ba}
Diet 4	15.23 ± 0.13^a	15.09 ± 0.04^d	38.23 ± 0.27^a	22.53 ± 0.12^d	8.93 ± 0.22^c	9.79 ± 0.41^a
Diet 5	11.27 ± 0.12^c	20.38 ± 0.18^a	1.16 ± 0.87^c	9.60 ± 0.06^c	57.60 ± 0.70^a	0.46 ± 0.04^c

¹Values are measures of triplicates. Means with the same letters within a row do not differ significantly ($P > 0.05$), $n = 15$.

3.2. Growth and survival performance and feed utilization

The gain weight and specific growth rate (SGR, % day⁻¹) of *S. serrata* are shown in Figures 4 and 5. The gain weight of Diet 1, Diet 2, Diet 3, Diet 4, and Diet 5 groups were 19.08 ± 7.48 g, 16.04 ± 2.93 g, 15.96 ± 1.91 g, 21.75 ± 7.91 g, and 23.87 ± 8.99 g, respectively. All experimental diets were not significantly different ($p > 0.05$) in terms of weight gain after 30 days of the culture period. However, mangrove crab fed with Diet 4 was higher than Diet 2, Diet 3, and the negative control diet. Moreover, Diet 1, Diet 2, Diet 3, Diet 4, and Diet 5 achieved SGR of 0.26 ± 0.09 % day⁻¹, 0.26 ± 0.07 % day⁻¹, 0.26 ± 0.05 % day⁻¹, 0.35 ± 0.11 % day⁻¹, and 0.42 ± 0.18 % day⁻¹, respectively. All experimental diets were not significantly different ($p > 0.05$). Nevertheless, Diet 4, which contains a 30 % diet of crab shellmeal inclusion fed on mangrove crab *S. serrata* obtained higher SGR, although there was no significant difference.

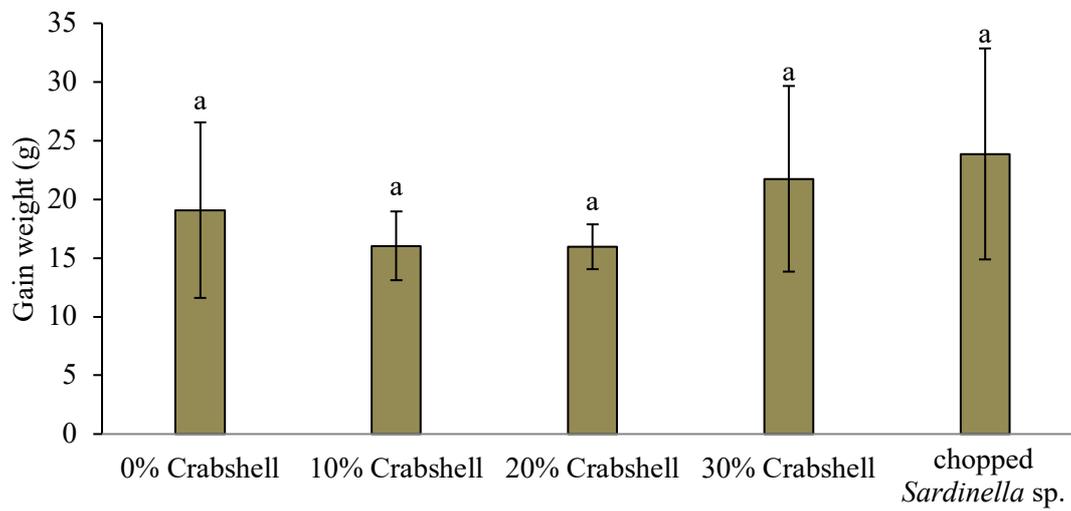


Figure 4. Gain weight performance of mangrove crab *S. serrata* fed with different diets after 30 days of culture. Diet 1 = 0% Crabshell, Diet 2 = 10% Crabshell, Diet 3 = 20% Crabshell, Diet 4 = 30% Crabshell, Diet 5 = Copped *Sardinella* sp. Bars with the same letters do not differ significantly ($P > 0.05$). Values are means \pm S.E.M (standard error mean), $n = 6 - 40$.

The survival rate (SR, %) of Diet 1, Diet 2, Diet 3, Diet 4, and Diet 5 groups were 80.00 ± 13.33 %, 80.00 ± 13.33 %, 60.00 ± 16.33 %, 80 ± 13.33 %, and 100.00 ± 0.00 % (Figure 6). Inclusions of all levels of crab shellmeal fed on mangrove crab *S. serrata* were not significantly different among experimental diets in terms of SR. Moreover, Table 3 shows the feed intake (FI), feed conversion ratio

(FCR), and protein efficiency ratio (PER) values of mangrove crab *S. serrata*. The combination of fishmeal and crab shellmeal had no significant effects in terms of FI, FCR, and PER. However, the FI and FCR of all experimental diets were significantly higher ($p < 0.05$) than those of trash fish (chopped *Sardinella* sp.), whereas the PER of trash fish was significantly higher ($p < 0.05$) than that of all experimental diets.

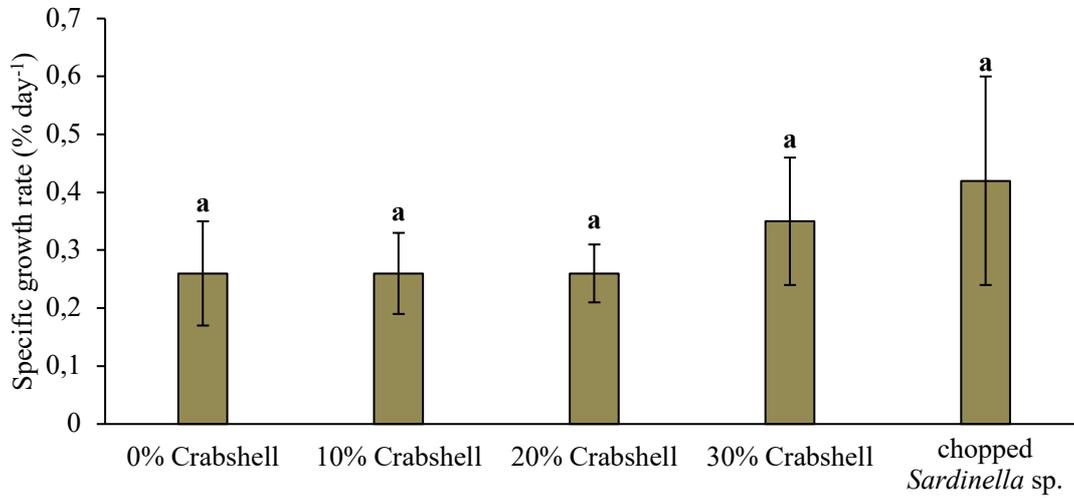


Figure 5. Growth performance (SGR) of mangrove crab *S. serrata* fed with different diets after 30 days of culture. Diet 1 = 0% Crabshell, Diet 2 = 10% Crabshell, Diet 3 = 20% Crabshell, Diet 4 = 30% Crabshell, Diet 5 = Copped *Sardinella* sp. Bars with the same letters do not differ significantly ($P > 0.05$). Values are means \pm S.E.M (standard error mean), $n = 6 - 40$.

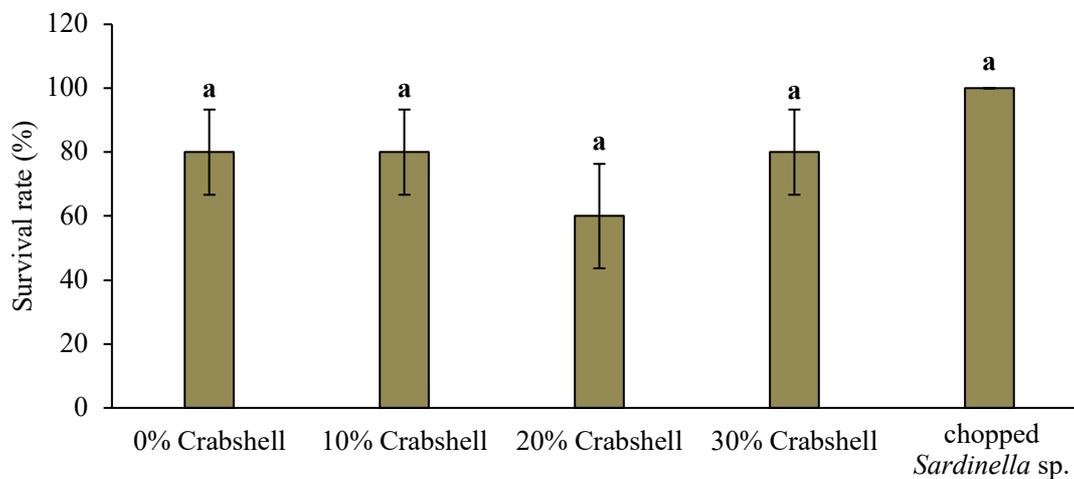


Figure 6. Survival rate of mangrove crab *S. serrata* fed with different diets after 30 days of culture. Diet 1 = 0% Crab shellmeal, Diet 2 = 10% Crab shellmeal, Diet 3 = 20% Crab shellmeal, Diet 4 = 30% Crab shellmeal, Diet 5 = Copped *Sardinella* sp. Bars with the same letters do not differ significantly ($P > 0.05$). Values are means \pm S.E.M. (standard error mean), $n = 10 - 50$.

Table 3. Feed utilization of mangrove crab *S. serrata*

¹ Diets	FI	FCR	PER
Diet 1	261.94 \pm 34.19 ^a	22.17 \pm 3.73 ^a	1.41 \pm 0.49 ^b
Diet 2	285.12 \pm 27.81 ^a	22.23 \pm 4.12 ^a	1.15 \pm 0.33 ^b
Diet 3	265.43 \pm 29.13 ^a	18.31 \pm 3.40 ^a	0.75 \pm 0.16 ^b
Diet 4	253.45 \pm 14.17 ^a	21.30 \pm 6.43 ^a	0.75 \pm 0.25 ^b
Diet 5	115.71 \pm 21.75 ^b	7.71 \pm 1.99 ^b	161.81 \pm 140.44 ^a

¹Values are means \pm S.E.M. (standard error mean). Means with the same letters within a row do not differ significantly ($P > 0.05$), $n = 40$.

3.3. Whole body proximate composition of mangrove crab

Proximate compositions of mangrove crabs are shown in Table 4. Ash content and moisture content varied from $21.53 \pm 0.22\%$ – $31.60 \pm 0.20\%$ and $7.69 \pm 0.04\%$ – $21.48 \pm 0.05\%$, respectively, where crab fed with Diet 3 was significantly ($P < 0.05$) greater among all diets. Carbohydrates ranged from $13.00 \pm 0.10\%$ – $34.07 \pm 0.87\%$, indicating that crab fed with Diet 4 was significantly higher ($P < 0.05$) in all experimental diets. Levels of crude fat ranged from $1.73 \pm 0.18\%$ – $5.03 \pm 0.07\%$, where crab fed with Diet 2 and Diet 4 were significantly lower ($P < 0.05$) than crab fed with Diet 1, Diet 3, and Diet 5. In addition, crude protein varied from $21.83 \pm 0.73\%$ – $37.70 \pm 0.12\%$, where crab fed with Diet 2 was significantly greater ($P < 0.05$) among all diets. Levels of calories ranged from $218.33 \pm 1.45\%$ – $275.00 \pm 1.15\%$, indicating that crab fed with Diet 5 was significantly higher ($P < 0.05$) among all experimental diets.

Table 4. Whole body composition of mangrove crab *S. serrata* (%)

¹ Diet	Ash	Moisture	Carbohydrates	Crude Fat	Crude Protein	Calorie
Diet 1	26.67 ± 0.20^c	16.33 ± 0.09^b	26.57 ± 0.35^c	3.00 ± 0.15^b	27.43 ± 0.24^d	243.00 ± 0.58^e
Diet 2	26.10 ± 0.15^d	21.48 ± 0.05^a	13.00 ± 0.10^e	1.73 ± 0.18^d	37.70 ± 0.12^a	218.33 ± 1.45^e
Diet 3	31.60 ± 0.20^a	7.69 ± 0.04^e	28.77 ± 0.20^b	2.47 ± 0.09^c	29.47 ± 0.19^c	255.33 ± 0.33^b
Diet 4	29.40 ± 0.23^b	12.79 ± 0.13^d	34.07 ± 0.87^a	1.90 ± 0.26^d	21.83 ± 0.73^e	241.00 ± 2.52^d
Diet 5	21.53 ± 0.22^e	16.09 ± 0.07^{cb}	25.07 ± 0.78^d	5.03 ± 0.07^a	32.30 ± 0.60^b	275.00 ± 1.15^a

¹Values are means \pm S.E.M. (standard error mean). Means with the same letters within a row do not differ significantly ($P > 0.05$), $n = 15$.

4. Discussion

CSM supplementation can enhance the proximate composition of experimental diets, thereby improving mangrove crab's whole body composition. However, CSM supplementation had no significant effect on feed utilization, as well as the growth and survival rates of mangrove crabs. In mangrove crab farming, shrimp diet and trash fish are commonly used. Therefore, developing an appropriate mangrove crab feed formulation is a priority area of research (Say and Ikhwanuddin, 1999; Zhao et al., 2016; Dayal et al., 2019; Aaqillah-Amr et al., 2022). A farmer can further enhance the quality of the feed by understanding the composition and reformulating the feed if the nutrients are not adequate to sustain the crabs. The crab's body is made of claws and legs, and its back shell is made of hard, opaque tissue called chitin (Cornwall, 2014). Chitin is a polymer that is found in arthropods' exoskeletons, such as crabs (Guerrero, 2000; Brock et al., 2003). Among the most abundant organic compounds in nature, chitin is made up of sugar molecules arranged in a network with a variety of uses, such as agriculture, aquaculture, and biotechnology (Beundia, 1999; Shamshina et al., 2020). About 10% of the shell of the mangrove crab *S. tranquebarica* contains chitin (Thirunavukkarasu and Shanmugam, 2009). Using underutilized by-products in the seafood industry as a substitute for costly FM in mangrove crab diets, the present study aims to develop a more cost-effective diet. By substituting FM with 30% CSM, it is evident that the survival and growth of mangrove crab *S. serrata* can be boosted, although not significantly different. In aquaculture nutrition, the replacement of FM with alternative protein sources has long been well-established for many crustacean and fish species, and research on crab species is meager. Species of crabs such as *Eriocheir sinensis* (Luo et al., 2011; Jiang et al., 2013), *S. serrata* (Nguyen et al., 2014), and *S. paramamosain* (Suwirya et al., 2009) have been studied to replace FM with different vegetable sources of protein. In addition, the fish bone meal (FBM) is regarded as a significantly improved source of dietary protein that can be used to replace 45% of FM in Atlantic cod diets. Luo et al. (2011) stated that it is possible to replace 40% of FM with rapeseed meal and soybean meal in the diet of *E. sinensis*. In addition, *S. paramamosain*, soybean meal, and corn gluten meal could replace 20 – 40% of FM (Suwirya et al., 2009). In our study, 10 – 20% FM could be replaced with CSM for *S. serrata*. The higher level of FM could be replaced by animal protein sources, possibly

due to higher protein content, fatty acids, balanced amino acids, palatability, and less toxin or antinutrients (Kader et al., 2017).

FBM tended to increase dietary phosphorus, calcium, and ash contents, which helped to enhance Atlantic cod growth and feed consumption (Toppe et al., 2006). A crabstick (farm-made feed) developed for crab fattening increased moisture, crude protein, crude lipids, crude fiber, and total ash, thereby enhancing FI and FCR in blue swimming crab *P. pelagicus* (Kalidas et al., 2020). In the present study, increasing supplementation of CSM significantly affects dietary ash, moisture content, carbohydrates, crude fat, crude protein, and crude fiber. The nutrient composition of feed, such as ash, moisture content, carbohydrates, fat, protein, and fiber, can control the metabolism of crustaceans, shrimps, and other fishes, provide energy, and contribute to a living cell's functioning (Krogdahl et al., 2005; Ndome et al., 2010). For aquafeed ingredients, the PER is an effective parameter for measuring protein quality (Goytortua-Bores et al., 2006; Luo et al., 2011). In the present study, the FCR, PER, and FI showed no effect with different levels of CSM supplementation; hence, the feed utilization, growth, and survival of the mangrove crab *S. serrata* were not affected. In other studies, different levels of phospholipids enhanced feed utilization for kuruma shrimp (Michael et al., 2008), blue swimming crab (Li et al., 2014), and blunt snout bream (Li et al., 2015), and black tiger shrimp (Kumaraguruvasagam et al., 2005).

CSM replacement changed the body composition of the *S. serrata*, such that ash, carbohydrates, moisture, crude fat, crude protein, and calories were significantly different ($p < 0.05$). Kader et al. (2017) stated that there were no significant effects of replacing FM with FBM on the composition of whole bodies of *S. paramamosain*. However, the inclusion of phospholipids significantly improved crude lipids and crude protein content. Moreover, researchers have observed that dietary phospholipids positively affect protein retention in *Megalobrama amblycephala* (Li et al., 2015) and *Seriolla dumerilli* (Uyan et al., 2009). As shown in the body composition analysis, diets based on CSM (Diet 2, Diet 3, and Diet 4) contain higher levels of ash. Conversely, dietary FBM did not influence the whole body ash composition of *S. paramamosain* (Kader et al., 2017), *Megalobrama amblycephala* (Li et al., 2015), *Penaeus monodon* (Kumaraguruvasagam et al., 2005), *Litopenaeus vannamei* (Gonzalez-Felix et al., 2002) and *Gadus morhua* (Toppe et al., 2006).

Conclusion

The results of this study suggest that fishmeal may be replaced with crab shellmeal from mangrove crab diet formulations. Supplementing crab shellmeal to the diets of mangrove crabs can enhance their whole body composition; however, it did not affect feed utilization, growth performance, and survival rate of the mangrove crab. By utilizing the by-products and waste products of crab meat processing plants, crab shellmeal can be the most cost-effective and efficient option for aquatic animal feed as it contains chitin and protein.

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