

The Siphonal Mantle Morphology of *Tridacna crocea*

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Tridacna crocea is the smallest among the eight species of Family Tridacnidae and the most abundant tridacnids in reefs around the Philippine archipelago. This investigation describes the characteristic mantle pattern and color of *T. crocea* and correlates them to genetic structure; verifies the characteristic mantle color and pattern of *T. crocea* underwater and differentiates it from its closely related species, *T. maxima* in the field. Tissue samples of *Tridacna crocea* and *T. maxima* were collected and preserved in 95% alcohol. Prior to any mantle collection, each clam was photographed.

Thirteen mantle patterns were identified from the 174 *Tridacna crocea* individuals of six reef areas: Pamilacan, Tanon Strait, Carbin, Camiguin, Southeastern Samar and Spratlys. Results revealed that *Tridacna crocea* can be distinguished from *T. maxima* in the field by the appearance and arrangement of their hyaline organs. Moreover, analysis on genotype-phenotype correlation using the *T. crocea* mantle morphology/color, found no significant relationship between the mantle morphs and genetic structure of the individuals.

KEYWORDS: *Tridacna crocea*, *T. maxima*, mantle morphology, mantle color, genetic structure.

INTRODUCTION

T*ridacna crocea* is the smallest among the eight species of Family Tridacnidae and the most abundant tridacnids in reefs around the Philippine archipelago despite the existence of its commercial harvesting (Ravago-Gotanco et al., 2007; Juinio-Menez, et al., 2003;

Calumpong & Cadiz, 1993; Gomez & Alcala, 1988; Junio, et al., 1988; Alcala, 1986). As described by Rosewater (1965), this species, in its natural environment, can be recognized by its habit of being completely imprisoned in coral pockets with free margins of its valves nearly flush with the substrate. This habit distinguishes *Tridacna crocea* from its closely related species, *T. maxima* which lives in relatively shallower burrows in coral, hence its shell protrudes halfway from the coral rubble, but which is also tightly fastened to the substrate by a byssus.

The widely accepted taxonomic classification of the family Tridacnidae is by Rosewater (1965) and was based on shell morphology. He described *Tridacna crocea* and *T. maxima* as having opposed valves with a well-defined byssal orifice without tightly fitting teeth. However, the valves of *T. crocea* are usually quite smooth and stouter with depressed sculpture and are more triangularly ovate in shape (Figure 1A); *T. maxima*, although sometimes nearly scaleless, usually does not have the shell sculpture so reduced and its valves tend to be more triangularly elongate in shape (Figure 4B). Other differences that Rosewater (1965) noted between the two include the shapes of the adductor-retractor muscle scar complexes; interdigitating projections of the dorsal margins of the valves, the number of riblets on the radial folds and the relative lengths of byssal orifices.

However, there is a considerable overlap in mantle coloration and pattern between these two species. The mantle is an extension of the inhalant and exhalant siphons and is also referred to as siphonal tissue. It contains the majority of the zooxanthellae as well as the fixed cells called iridophores that contain pigments which mainly protect the clam against excessive light and UV radiation. These pigments have color range of blue to brown or green to yellow. These pigments and their combinations are the reason for the wide range of colors and patterns that are found in these clams.

Both species are exceedingly variable in these characters, but this may be due to convergence where these characters are subject to no strong selective pressures (Rosewater, 1965). Like *T. maxima*, mantle color in *T. crocea* runs an array of brilliant green, blue, purple and brown with great pattern variation. According to Rosewater (1965), color brilliancy in *T. maxima* may vary in widely separated geographic regions. At Eniwetok, marshal Islands, the mantle colors were observed to be extremely bright, as well as in the Great Barrier Reef. However, the colors were observed to be more subdued like in *T. crocea* at Andaman Sea in Malaysia and Thailand and off southern Sumatra.

This phenomenon may be due to differences in the conditions of the animals at the time the observations were made, to a real geographic variation based on genetic difference, or to environmental factors. Rosewater (1965) emphasized the difference between the two species in terms of their hyaline organs which tend to be concentrated along the edge of the mantle on papillae in *T. maxima* while in *T. crocea* they are more diffuse on the mantle surface. However, this character may be confusing underwater especially when the two species occur together in reef flats and coral rubble.

Based on the amendment done by Lucas et al. (1991) on Rosewater's taxonomy using morphologic characters, Family Tridacnidae has eight extant recognized species with two genera and three subgenera (*Tridacna*: subgenus *T. sensu stricto* (*T. gigas* Linnaeus, 1758); subgenus *Chametrachea*: *T. maxima* Roding, 1798, *T. squamosa* Lamarck, 1819, and *T. crocea* Lamarck, 1819; subgenus *Persikima*: *T. derasa* Roding, 1798 and *T. tevoroa* Lucas, Ledua & Braley, 1991); and *Hippopus*: *H. hippopus* Linnaeus, 1758 and *H. porcellanus* Rosewater, 1982) (Rosewater, 1965, 1982; Lucas et al., 1991). The resulting major groups were confirmed by Benzie and Williams (1998) using protein electrophoresis, however, they suggested that some characters, such as, the lack of attachment to the substrate as adults, the lack of boring into coral, the small byssal orifice, and the dorsally extended ctenidia, are not reliably diagnostic of the genus, subgenus or species levels. Many of the characters shared by pairs of subgenera within *Tridacna* are also found in *Hippopus*, and are likely to be primitive and therefore give little clue to the evolutionary relationships of the taxa concerned. The first three mentioned above are also part of a set of ecologically plastic and functionally inter-related characters.

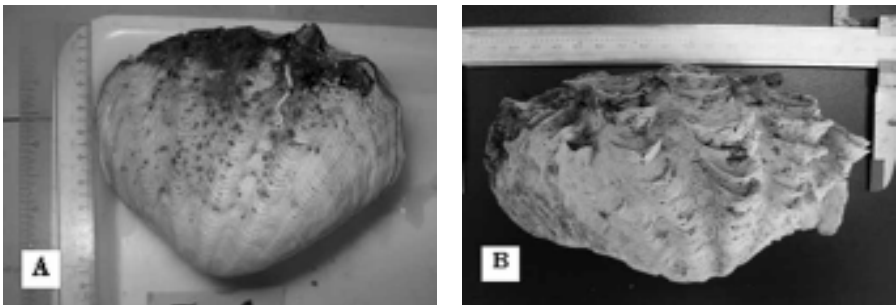


Figure 1. Lateral view of the shell valves of [A] *Tridacna crocea* and [B] *T. maxima*.

This investigation describes the characteristic mantle pattern and color of *T. crocea* and differentiates it from its closely related

species, *T. maxima*. This study verifies the characteristic mantle color and pattern of both species using genetic analysis. Furthermore, an attempt to correlate the clams' mantle morphology to their genetic structure was also done.

METHODS

Phenotypic Differentiation (Mantle Color and Morphology)

Fifty clams from each of the following reefs were sampled: Pamilacan (Bohol), Bolisong (Tanon Strait), Carbin (Sagay, Negros Occidental), Spratlys (South China Sea), Camiguin Island, and Camanga (Southeastern Samar). Using an underwater digital camera, each clam was photographed before a piece of mantle was cut from each clam. Pictures were downloaded and sorted according to mantle color and pattern. Samples were then grouped according to categories (Table 1).

Relating Mantle Morph to Genetic Structure

To analyze the genetic structure that may be associated with the mantle pattern and color of the *Tridacna crocea*, only nine mantle characters (morphs 1 to 9) were considered. This was based on the individuals with DNA that were successfully sequenced. The DNA sequences were converted into a FASTA format and analyzed for genetic variation using the analysis of molecular variance (AMOVA) implemented in the ARLEQUIN version 3.

Differentiating *Tridacna crocea* from *T. maxima*

Tissue samples of *Tridacna crocea* and *T. maxima* were collected and preserved in 95% alcohol. Ten clams were picked up and identified as *Tridacna crocea* (5 individuals) and *T. maxima* (5 individuals) by a panel of three giant clam experts from the underwater pictures obtained in the above-mentioned sampling. Selection and identification were based on mantle pattern and color. These were labeled TcP1-5 and TmP1-5, respectively. Another five *T. crocea* and six *T. maxima* that were positively identified by their shell and mantle morphology (collected and maintained in the hatchery for spawning purposes) were also sampled for comparison and were labeled TcL1-5 and TmL1-6. Six more samples that were randomly collected from the field as *T. crocea*

were also included.

Genomic DNA was extracted using standard phenol-chloroform extraction method utilizing TNES-urea digestion buffer (6 M urea, 1M Tris-HCl pH 7.5, 5 M NaCl, 0.5 M EDTA, and 1% SDS or sodium dodecyl sulfate) as described in Wasko et al. (2003) and Proteinase K treatment. Partial sequences (500 bp) of the mitochondrial *cytochrome c oxidase 1* (COI) gene were amplified with a specific primer for *Tridacna crocea* (Tridacna 1F 5'- ACC CTT TAY TTT TTA TTA GCA Y- 3'; Tridacna 3R 5'- CAA TGC TGT AAT CGC CAA TGA C-3') designed by Barber (2006). PCR products were visualized using 1% (w/v) agarose gel electrophoresis. Clones (forward and reverse stands) were sequenced on an ABI 377 or an ABI 3730 automated sequencer using Big Dye (Applied Biosystems, Foster City, CA) terminator chemistry. Nucleic acid sequences were subjected to BLAST/N (Altschul et al., 1990) searches at the National Center for Biotechnology Information (NCBI). ChromasPro version 1.33 available at <http://www.technelysium.com.au/ChromasPro.html>, sequences were downloaded and subjected to BLAST search at <http://www.ncbi.nlm.nih.gov.csulib.ctstateu.edu/blast/blastFAQs.html>.

Table 1.

Mantle color and patterns generated from the field and picture observations used as criteria in grouping the samples

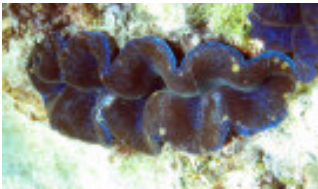

MANTLE COLORMORPH	DESCRIPTION	PHOTO
1	Dark blue to chocolate brown overall mantle color. Eyespots or hyaline organs bounded by Iridescent blue circles. Mantle margins are also iridescent blue.	
2	Light brown with thin cream/white specks on central part of siphonal mantle. Hyaline organs/eyes are black in color bounded by thinner cream/tan margins. Mantle margins are usually light green.	

Table 1. Continued...





MANTLE COLORMORPH	DESCRIPTION	PHOTO
3	Light brown/tan with specks of dark brown, tan and cream/white. Hyaline organs black bounded by cream to white margins. Occasional warty protuberances are found on the lateral side of siphonal mantle folds. Mantle margins are generally yellow green and prominent.	
4	Tan with specks of white with blue mantle edge/margin.	
5	Dark brown with iridescent yellow or tan eye margins. Central siphonal mantle appeared to be plain.	
6	Dark brown with cream or white fine lines (horizontal) along siphonal mantle outer fold.	

Table 1. Continued...








MANTLE COLORMORPH	DESCRIPTION	PHOTO
7	Dark brown either with white specks horizontally spread on the siphonal mantle or white or cream rays regularly spaced on the outer mantle fold. Yellow green mantle margins.	
8	Tan to olive green with green or cream irregular spots scattered on the siphonal mantle. White stripes maybe present along the lateral side. Mantle margins are usually yellow green.	
9	Light to dark brown with scattered blue specks. Mantle margin yellow green.	
10*	Light brown on lateral side of siphonal mantle with warty protuberance. Dark brown on central portion with mint green specks.	
11*	Olive green mantle. Hyaline organs are black bounded by white margins. They occur in two or three layers or are scattered.	

Table 1. Continued...

MANTLE COLORMORPH	DESCRIPTION	PHOTO
12*	Dark brown with specks of blue on inner mantle fold. Mantle margins yellow green.	
13*	Siphonal mantle plain without obvious pattern. Sky blue to deep blue mantle margins. Hyaline organ margin same color with mantle margin.	

Photographs are found in Figure 1
 * not included in the AMOVA

RESULTS

Phenotypic Differentiation (Mantle Color and Morphology)

Thirteen mantle patterns (Table 1) were identified from the 174 *Tridacna crocea* individuals of six reef areas: Pamilacan, Tañon Strait, Carbin, Camiguin, Southeastern Samar, and Spratly. A look at the occurrence and distribution of mantle patterns in each population (Figure 2) shows Pamilacan with nine mantle patterns (1, 2, 3, 4, 5, 6, 7, 8 and 11), Spratly, only three (1, 2 and 4), Tañon Strait with 10 (1, 3, 5, 6, 7, 8, 9, 10, 12 and 13), Carbin has six (1, 2, 3, 6, 7 and 13), Camiguin, five (1, 2, 3, 7 and 8) and Southeastern Samar, nine (1, 2, 3, 4, 5, 6, 7, 8 and 13). Mantle morph 1 was the most common in all populations. Among the percentages of each population exhibiting each mantle pattern (Figure 3), all patterns except for morphs 9, 10, 11 and 12 were found in all six sites.

Relating Mantle Morph to Genetic Structure

A total of 64 clam DNA sequences from five populations: Pamilacan, Carbin, Bolisong, Spratly and Southeastern Samar (Table 1) were analyzed. Pamilacan samples displayed six mantle patterns (morphs 1, 2, 4, 6, 7 and 8), Guiuan and Bolisong have five (morphs 1, 2, 3, 4 and

5 and morphs 1, 3, 5, 6 and 9, respectively), Carbin has three (morphs 1, 2 and 6) and Spratly, three (morphs 1, 2 and 4).

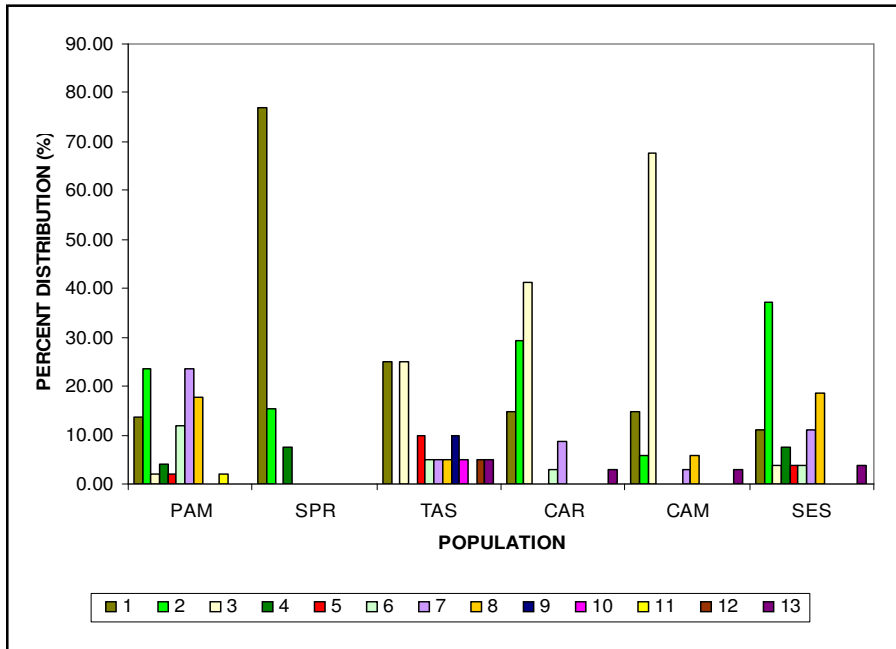


Figure 2. Percent distribution of mantle patterns in each of the six populations.

Note: PAM (Pamilacan), SPR (Spratly), TAS (Tañon Strait), CAR (Carbin), CAM (Camiguin), and SES (South-Eastern Samar)

The 64 DNA sequences were grouped according to the nine mantle patterns and subjected to analysis of molecular variance (AMOVA). Results revealed no significant genetic variation among the morphs ($F_{ST} = -0.03737$, $p = 0.93842$) at 1023 permutations. As indicated in Table 2, different mantle morphs may even share the same haplotypes like in the case of haplotype 7. This result is in conformity with the results obtained by Laurent et al. (2002) on *Tridacna maxima* using allozymes. They found no relationship between color of the mantle and genetic structure of *T. maxima*, with individuals of different patterns showing similar genetic structures. Rosewater (1965) had reported variation in mantle color of giant clams in which he described several morphs *Tridacna maxima*. According to McMichael (1974), such variation in color can be due to genetic variation in the clam. However, Rosewater (1965) emphasized the role of zooxanthellae variation.

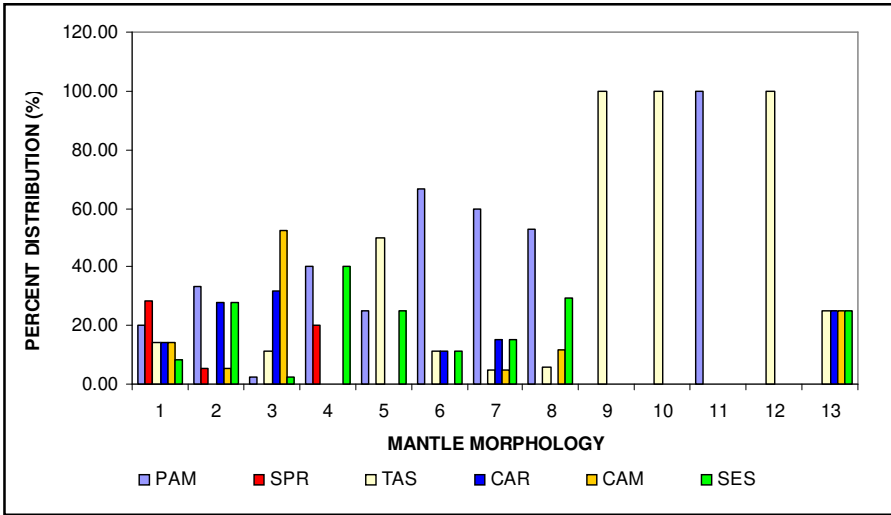


Figure 3. Percentage of population exhibiting each mantle pattern.

Codes are given in Figure 2.

Differentiating *Tridacna crocea* from *T. maxima*

Sequences were subjected to BLAST search (BLASTN 2.2.18, Zheng Zhang et al., 2000). Among the photograph-based samples, all probable *Tridacna crocea* were verified exactly as *T. crocea*, but, the probable *T. maxima* turned out to be *T. crocea*. For the *T. maxima* (TmL1-6) and *T. crocea* (TcL1-6) samples identified by their mantle and shell morphology were exactly *T. maxima* and *T. crocea*, except for TmL5 which failed to match with any of the tridacnid CO1 sequences in GeneBank using the NCBI basic Tool Alignment Search Tool (BLAST). On the other hand, sequences of the seven “*T. crocea*” samples randomly collected from the field all matched with *T. maxima cytochrome c oxidase I* sequence during the BLAST search.

DISCUSSION

The only description Rosewater (1965) has given to differentiate *T. maxima* from *T. crocea* was in terms of their hyaline organs arrangement, wherein in the former, they tend to be concentrated along the edge of the mantle on papillae (Figure 4A) while in the latter, they are more diffuse on the mantle surface (Figure 4B & 4C). This difference has been observed in the present study. Specifically, these “dark spots” or “eyes” appeared as a distinct continuous line along the siphonal mantle

margin in *T. maxima*. Moreover, these eye-like structures as called by Stasek (1966), typically were bounded by lighter margins usually white, cream and blue in *Tridacna crocea*, whereas in *T. maxima*, were not (Fig 4B & 4C). This feature was consistently observed in the 64 samples of *T. crocea* with BLAST results matching with the *T. crocea* sequences in GeneBank. Likewise, 94.4% of the sequenced *T. maxima* samples exhibited the unbounded hyaline organs.

Table 2.

Haplotypes of Tridacna crocea and their mantle morphology

HAPLO-TYPE	MANTLE MORPHOLOGY								
	1	2	3	4	5	6	7	8	9
2			1						
3									1
4			2				1		
5					1				
6			1						
7	2	4	2	1		3	4	2	1
9							1		
10	1								
11	3			1	1				
12	1								
13		1	1					1	
19							1		
20			1						
28	1								
29	2		1			1			
30		1							
31			1						
32	1								
33							1		
34		1							
36								1	
37	2							1	
41	1								
42	1								
43	1								
44	1								
50	2	1							
51					1				
52				1					
53	1								
54			1						

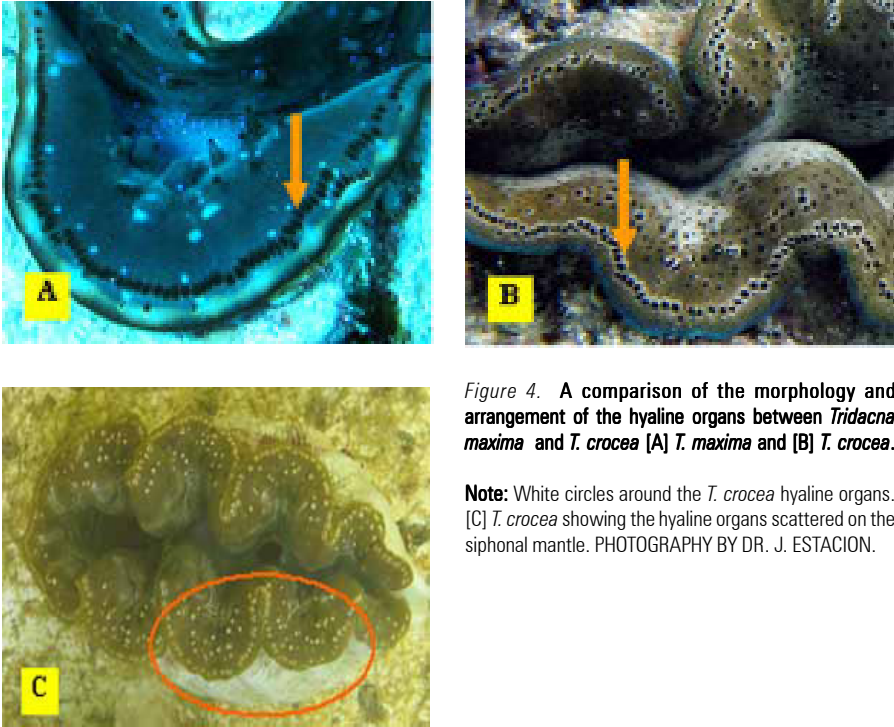


Figure 4. A comparison of the morphology and arrangement of the hyaline organs between *Tridacna maxima* and *T. crocea* [A] *T. maxima* and [B] *T. crocea*.

Note: White circles around the *T. crocea* hyaline organs. [C] *T. crocea* showing the hyaline organs scattered on the siphonal mantle. PHOTOGRAPHY BY DR. J. ESTACION.

Relating genetic structure to mantle morphology in the present study does not provide new insights because no significant genetic difference according to mantle morphology was observed. In summary, the absence of difference is not conclusive; it can be interpreted as genetic homogeneity of *Tridacna crocea* and mantle pattern like color may be a result of zooxanthellae and iridophores variation or local adaptation (Laurent et al., 2002). Moreover, the non-significant variation may be simply due to the reason that the genetic marker (mitochondrial *cytochrome c oxidase 1*) used in the study is not linked to color pattern regulation. Moreover, *Tridacna crocea* can be distinguished from *T. maxima* in the field by the appearance and arrangement of their hyaline organs. These organs are bounded by white, iridescent blue or yellow circles. *T. crocea* has distinct mantle patterns but colors are overlapping with *T. maxima*.

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REFERENCES CITED

- Alcala, A.C. 1986. Distribution and abundance of giant clams mollusks (Family Tridacnidae) in the south-central Philippines. *Silliman Journal*, 33, 1-4.
- Alcazar, S. 1988. Spawning and larval rearing of tridacnid clams in the Philippines. In Copland, J. & Lucas, J., eds. *Giant Clams in Asia and the Pacific*. ACIAR Monograph No. 9, Canberra, Australia, 125-128.
- Benzie, J. A. H & Williams, S. T. 1998. Phylogenetic relationships among giant clam species (Mollusca: Tridacnidae) determined by protein electrophoresis. *Marine Biology*, 132, 123-133.
- Brale, R. 1992. *The giant clam: hatchery and nursery culture manual*. ACIAR Monograph No. 15, Canberra. 144pp.
- Calumpong, H.P., Ablan, M.C., Macaranas, J, Solis-Duran, E., Alcazar, S., Abdon-Naguit, R. 1993. Biochemical evidence of self-fertilization in *Hippopus* species. In Fitt, W.K., ed. *Biology and mariculture of giant clams*. Australian Centre for International Agricultural Research, Canberra, Australia, 103-110 (Proc.No. 47, ACIAR).
- Calumpong, H.P. & Cadiz, P. 1993. Observations on the distribution of giant clams in protected areas. *Silliman Journal* 36(2), 107-116.
- Heslinga, G.A., & Watson, T.C. 1990. *Giant clam farming*. Honolulu, HI: Pacific Fisheries Development Foundation (NMFS/NOAA). 179pp.
- <http://www.technelysium.com.au/ChromasPro.html>
- Juinio-Meñez, M.A., Magsino, R.M., Ravago-Gotanco, R., Yu, E.T., 2003. Genetic structure of *Linckia laevigata* and *Tridacna crocea* populations in the Palawan shelf and shoal reefs. *Marine Biology*, 142, 717–726.
- Laurent V, Planes S, Salvat B (2002) High variability of genetic pattern in giant clam (*Tridacna maxima*) populations within French Polynesia. *Biological Journal of the Linnaean Society*, 77, 221-231.
- Lucas, J.S., Ledua, E., & Braley, R.D. 1991. *Tridacna tevoroa* Lucas, Ledua and Braley: A recently described species of giant clam (Bivalvia; Tridacnidae) from Fiji and Tonga. *Nautilus*, 105, 92-103.
- McMichael, D.F. 1974. Growth rate, population size and mantle coloration in the small giant clam *Tridacna maxima* (Roding), at One Tree Island, Capricorn Group, Queensland. Proceedings of the Second international Coral Reef Symposium 1, 241-254.

- Rosewater, J. 1965. The family Tridacnidae in the Indo-Pacific. *Mollusca*, 1, 347- 394.
- Schneider, J.A. & Foighil, D.O.1999. Phylogeny of giant clams (Cardiidae: Tridacninae based on partial mitochondrial 16S rDNA gene sequences. *Molecular Phylogenetics and Evolution*, 13, 59–66.
- Schneider J. 1992. Preliminary cladistic analysis of the bivalve family Cardiidae. *American Malacological Bulletin*, 9, 145-155.
- Stasek, C. R. 1966. The eye of the giant clam (*Tridacna maxima*). *California Academy of Science*, 58, 9.
- Zheng Zhang, Schwartz, S., Wagner, L., & Miller, W. 2000. A greedy algorithm for aligning DNA sequences. *Journal of Computational Biology*, 7(1-2), 203-214.