



Short communication

Mendelian inheritance of golden shell color in the Pacific oyster *Crassostrea gigas*



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ARTICLE INFO

Article history:

Received 1 October 2014

Received in revised form 31 December 2014

Accepted 31 January 2015

Available online 7 February 2015

Keywords:

Pacific oyster

Crassostrea gigas

Shell color

Inheritance

Epistatic

ABSTRACT

Shell coloration in many molluscs is highly variable. A shell color segregation study with progenies from a full factorial cross generated among Pacific oysters exhibiting distinct shell colors (golden, white and black) was conducted to investigate the inheritance of the golden shell color and its correlation with dark pigmentation. Random samples from twenty-three full-sib families were obtained and the shell coloration of offspring within each family was recorded. Results revealed that golden coloration was inherited in a different pattern from dark pigmentation, indicating its different genetic basis. Dark pigmentation was identified as a foreground color while golden or white color were background ones. The locus controlling background colors has two alleles with the allele for golden background being dominant to the allele for white background. In addition, the overlying foreground pigmentation of shells with a golden background was significantly lighter than that of shells with a white background, which suggested an epistatic effect of background color on shell foreground pigmentation. All these findings will facilitate the selection of elite oyster lines with desired shell coloration for aquaculture.

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1. Introduction

Molluscs are a large group of invertebrate animals presenting variable shell shapes and colorations, which have attracted many naturalists and collectors. Moreover, consumers tend to choose live seafood in the market according to their colors, which endows the coloration with product value of great importance (Alfnes et al., 2006). Pacific oyster (*Crassostrea gigas*) has the largest production among all cultured aquatic animals (FAO, 2011) and its coloration is of interest to the whole oyster industry (Brake et al., 2004; Kang et al., 2013). The oysters with golden shell coloration are rarely seen in the market and are sold at much higher prices than others (Nell, 2001). In our selective breeding practice of Pacific oyster, we obtained a number of golden shell variants, and established a set of full-sib families using these golden shell oysters as parents (Cong et al., 2014). These provided great materials to unravel the genetic basis of the golden coloration that remains unknown at present.

Mollusca shell color is known to be inheritable and affected by environmental factors, such as lights, salinity and substratum (Heath, 1975; Lindberg and Pearse, 1990; Sokolova and Berger, 2000). Several studies investigating the genetic basis of shell coloration with limited data from experimental crosses reported one or two loci with dominance to

control shell coloration (Adamkewicz and Castagna, 1988; Cole, 1975; Innes and Haley, 1977; Kobayashi et al., 2004; Liu et al., 2009; Wada and Komaru, 1990; Winkler et al., 2001; Zheng et al., 2013).

Pacific oyster shell pigmentation exhibits a continuous variation from near-white, pigment-free shells to near-black, fully pigmented shells, which was defined as foreground pigmentation. The shell pigmentation has been considered as a quantitative trait that is controlled by many genes with small-effects (Brake et al., 2004; Evans et al., 2009; Imai and Sakai, 1961). However, in some cases, only a small amount of major genes were identified to control shell coloration (Evans et al., 2009; Hedgecock et al., 2006). Our recent study also identified a SCAR marker which was well correlated with Pacific oyster shell pigmentation of black and white, suggesting a major locus that is responsible for foreground pigmentation (Ge et al., 2014). However, the genetic basis and mechanism of shell coloration in Pacific oyster are still inconclusive, especially for specific colors, such as golden coloration, which is clearly different from those dark pigmentation phenotypically. Does golden coloration possess a same genetic basis as the black pigmentation? If not, what are the genetic loci controlling golden coloration and is there any correlation between golden coloration and typical dark pigmentation?

In the present study, we performed controlled crosses between oysters showing different shell colors to determine the inheritance of golden coloration in *C. gigas* and to examine the relationship between golden coloration and typical dark pigmentation, which will facilitate the selection of elite oyster lines with desired coloration patterns.

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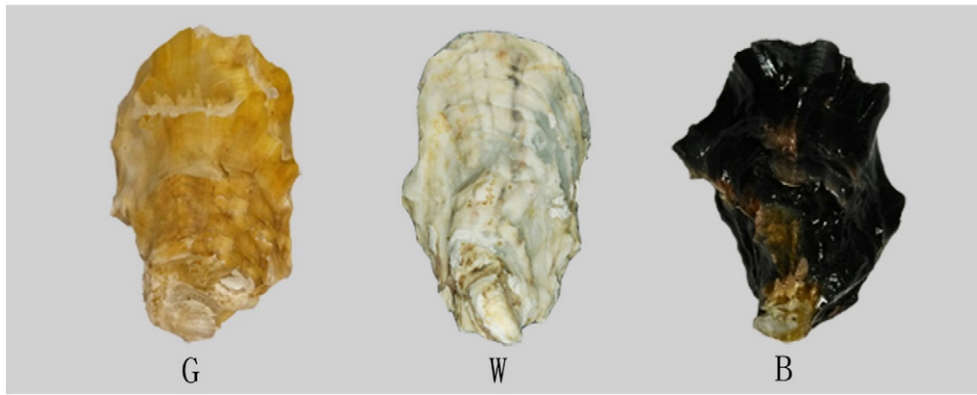


Fig. 1. Representative parents of three shell coloration patterns. G = Golden; W = White; B = Black.

2. Materials and methods

2.1. Parental source

One-year-old oysters with desired shell coloration were selected from nine full-sib families to conduct cross mating. All families were the first-generation selective families and were produced using wild oysters with specific shell colors in Rushan, China (Cong et al., 2014). Oysters with three shell coloration patterns were used as parents in

this study (Fig. 1), including those with golden shell (G) and two extreme foreground pigmentation patterns of solid white shell (W) and solid black shell (B).

2.2. Mating experiment and rearing

A full factorial cross among oysters with the three shell color patterns was performed to generate 27 families with 9 cross-mating groups

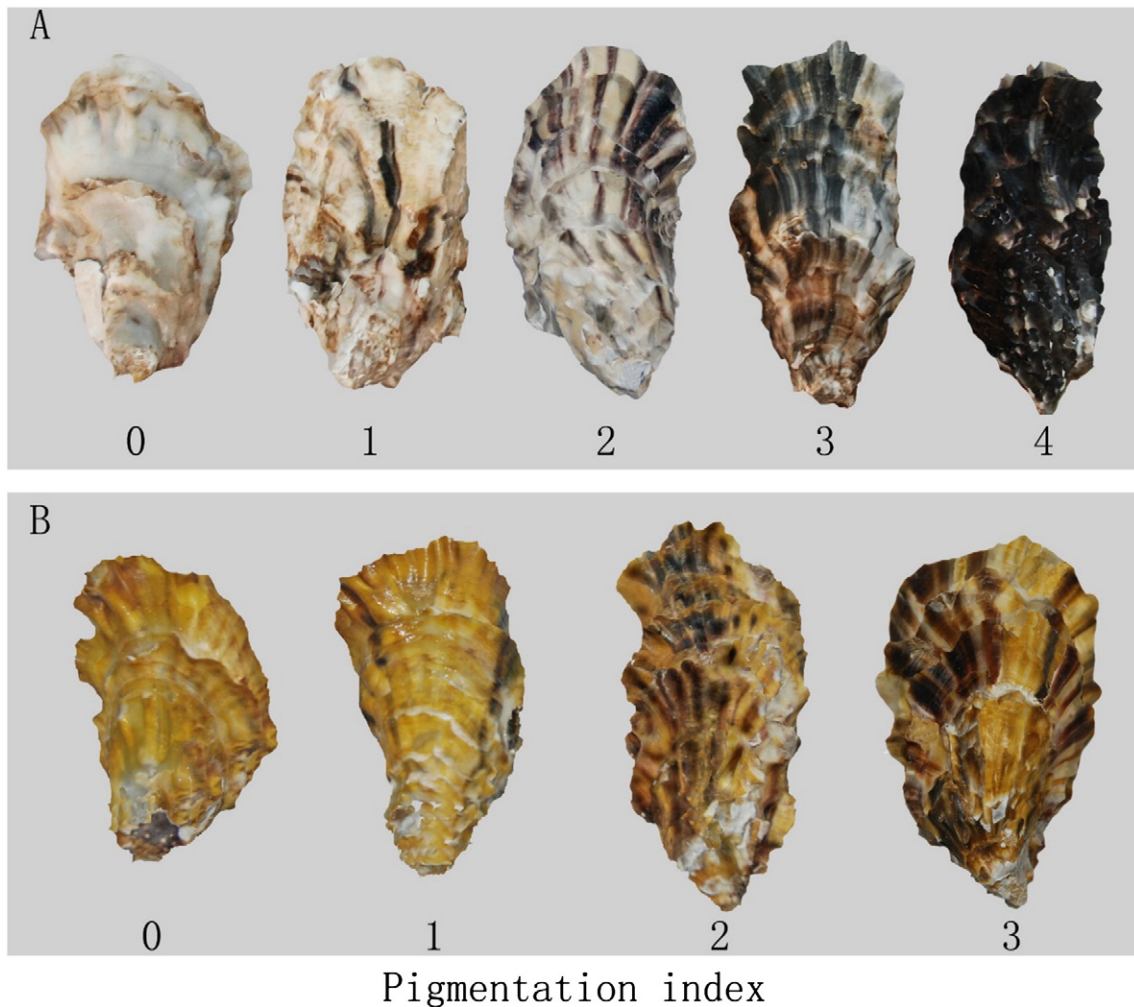


Fig. 2. Representative shell color morphs of offspring. (A) Offspring with white background and different foreground pigmentation (pigmentation index: 0–4); (B) offspring with golden background and different foreground pigmentation (pigmentation index: 0–3).

Table 1
Shell background color of the offspring from 23 crosses among three shell color patterns of *C. gigas*.

Family no.	Parents ^a		Background color of offspring					χ^2 (P value)
	Female	Male	Golden	White	Total	Expected ratio		
1	G1	G4	42	16	58	3:1	0.649	
2	G2	G5	92	30	122	3:1	0.917	
3	G3	G6	51	20	71	3:1	0.537	
4	W1	W4	0	45	45	0:1	NA	
5	W2	W5	0	44	44	0:1	NA	
6	W3	W6	0	70	70	0:1	NA	
7	B1	B4	0	53	53	0:1	NA	
8	B2	B5	0	45	45	0:1	NA	
9	B3	B6	0	38	38	0:1	NA	
10	G1	W4	61	49	110	1:1	0.253	
11	W1	G4	35	34	69	1:1	0.904	
12	G3	W6	29	28	57	1:1	0.895	
13	W3	G6	19	20	39	1:1	0.873	
14	G1	B4	26	28	54	1:1	0.700	
15	G2	B5	25	23	48	1:1	0.773	
16	B2	G5	26	34	60	1:1	0.302	
17	G3	B6	33	30	63	1:1	0.705	
18	B3	G6	46	42	88	1:1	0.670	
19	W1	B4	0	61	61	0:1	NA	
20	B1	W4	0	60	60	0:1	NA	
21	W2	B5	0	58	58	0:1	NA	
22	B2	W5	0	68	68	0:1	NA	
23	W3	B6	0	39	39	0:1	NA	

^a G Golden shell; W White shell; B Black shell.

and three replicates. To guarantee the survival of offspring, inbreeding between siblings was avoided.

Gametes of oysters with each color were rinsed into separate buckets by stripping the gonad. A suspension of eggs from each dam was divided into three equal portions and fertilized by sperm from the three sires of each color, respectively. Fertilized eggs of all cross groups were hatched at temperature of 24 °C and salinity of 30 psu. After 24-h incubation, D-stage larvae of each family were collected and transferred into a 100-L plastic bucket with an initial stocking

density of 8 larvae/mL. The larvae stocking density was decreased along with larval growth, and half volume of water was replaced with fresh water twice per day. Veligers were fed with daily rations of *Isochrysis galbana* at early stage (shell length <120 μm), and were fed with *Platymonas helgolandica* and *Chaetoceros calcitrans* at later stage. When oyster eye spots appeared, strings of scallop shells were placed into the bucket for larvae to attach. Within one week, all eyed larvae metamorphosed to spat and then were transferred to an outdoor nursery tank. All spats were spread into lantern nets after 40 days and then deployed to grow-out areas in Rushan, China.

2.3. Shell coloration record

Offspring samples were collected randomly from all families at 410 days post-fertilization in summer 2012. All sampled oysters were shucked, with shells washed and brushed thoroughly with freshwater. The shell background color was recorded as golden or not (i.e., white), while the foreground pigmentation was recorded as five categories with five index numbers, according to Imai and Sakai (1961). The categories were described as follow:

- 0—Whole surface has no dark coloration;
- 1—A small part of shell is dark colored;
- 2—Nearly half part of shell is dark colored;
- 3—Most part of shell is dark colored;
- 4—All part is dark colored.

2.4. Statistical analysis

The proportions of shell background color in the progeny were analyzed with the assumption that the single locus-two-allele model is the major determinant for shell color. The fit of the phenotypic frequencies to the expected frequencies under the proposed model of inheritance was tested using the Chi-squared test for goodness-of-fit. Fisher's exact test of independence was used to test the association of shell foreground pigmentation and shell background color.

Table 2
Independence tests of the association of shell foreground pigmentation (pigmentation index: 0–4) and background color (ie. golden and white) within 12 families.

Family no.	Parents		Background color type	Foreground pigmentation					Total	P value [†]
	♀	♂		0	1	2	3	4		
1	G1	G4	Golden	9	17	8	8	0	42	0.001
			White	0	2	12	2	0	16	
2	G2	G5	Golden	38	52	2	0	0	92	0.000
			White	5	14	9	2	0	30	
3	G3	G6	Golden	21	21	7	2	0	51	0.000
			White	1	6	6	6	1	20	
10	G1	W4	Golden	26	31	4	0	0	61	0.067
			White	12	28	7	2	0	49	
11	W1	G4	Golden	17	11	7	0	0	35	0.005
			White	5	13	12	4	0	34	
12	G3	W6	Golden	9	10	10	0	0	29	0.011
			White	6	5	9	8	0	28	
13	W3	G6	Golden	6	9	4	0	0	19	0.000
			White	1	2	12	5	0	20	
14	G1	B4	Golden	0	4	10	12	0	26	0.001
			White	0	0	6	13	9	28	
15	G2	B5	Golden	2	6	14	3	0	25	0.000
			White	0	5	3	7	8	23	
16	B2	G5	Golden	3	8	11	4	0	26	0.250
			White	0	10	14	8	2	34	
17	G3	B6	Golden	1	12	15	5	0	33	0.000
			White	0	1	8	10	12	31	
18	B3	G6	Golden	2	22	17	5	0	46	0.000
			White	0	1	22	13	6	42	
Total			Golden	134	203	109	39	0	485	0.000 [‡]
			White	30	87	120	80	38	355	

[†] P values were calculated with the use of Fisher's exact test.

[‡] The P value was based on Chi-squared test.

3. Results and discussion

Twenty-three full-sib families were obtained eventually, with four families lost during the grow-out stage. Several representative shell color morphs in the offspring were shown in Fig. 2. It is clear that dark pigmentation present in the surface of both white and golden shells. Golden coloration tends to be distributed all over the shell, while dark pigmentation is only mottled some area or banded radially in the surface (Fig. 2). All these observations support the hypothesis that golden coloration follows a different pattern from black pigmentation, which is named as the background color (golden or white) to distinguish the foreground dark pigmentation.

As shown in Table 1, the presence of golden background color was dominant over its absence (white). Each of the three golden families (families 1–3) produced both golden and white offspring in a ratio closely approaching 3:1. In contrast, none of the white background families (families 4–9, 19–23) produced golden offspring. Families 10–18, cross matings between golden oysters and different overlying pigmentation oysters, resulted in background color of offspring at ratios that were not different from 1:1. These observations clearly support the hypothesis that the two background colors are controlled by one locus, with one allele for golden background being dominant to the other allele for white. This result is consistent with the observation of Nell (2001) that true breeding oysters with golden shell and mantle were obtained after a few generations of selection. Similar inheritance model has also been reported in the bay scallop (*Argopecten irradians*) that yellow background color versus its absence appears to be inherited as a single gene with absence (white) being recessive (Adamkewicz and Castagna, 1988).

Independence tests indicated that shell foreground pigmentation was significantly related to shell background color in families 1–3 and 10–18, except families 10 and 16 (Table 2). In general, there were more golden offspring with lighter foreground pigmentation (pigmentation index: 0 and 1), whilst more white offspring with darker foreground pigmentation (pigmentation index: 2–4; Table 2). These all point to a tentative hypothesis that the background locus may be epistatic to one or more pigmentation genes which only are permitted to express when the allele is for white background. This has also been reported in the bay scallop as all the offspring with white background color had dark mottling on their shells, whereas none of the orange offspring followed this pattern in that study (Adamkewicz and Castagna, 1988).

In conclusion, our results demonstrated that the shell background color of Pacific oyster is controlled by one locus with two alleles and that the allele for golden background is dominant to the allele for white. Additionally, the oyster shell background color may have an epistatic effect on its foreground pigmentation. These results suggested major implications for the selection of oyster lines with desired shell coloration.

Acknowledgments

This study was supported by grants from the National High Technology Research and Development Program (2012AA10A405-6), the National Natural Science Foundation of China (31372524), and the Special Fund for Independent Innovation of Shandong Province (2013CX80202).

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