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Recommended Citation
Masaru, H; Komaru, A; De Sousa, Joana Teixeira; and Allen, Standish K. Jr., "Meiotic And Early Zygotic Development In Crassostrea Virginica Observed Through Confocal Microscopy" (2017). VIMS Articles. 301.
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MEIOTIC AND EARLY ZYGOTIC DEVELOPMENT IN CRASSOSTREA VIRGINICA OBSERVED THROUGH CONFOCAL MICROSCOPY

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ABSTRACT Several studies have examined early development in oysters, but few have reported the normal (expected) behavior of meiosis and early mitosis or of the organization of the spindle by the centrosomes. To establish this baseline, normal development of fertilized eggs in Crassostrea virginica from meiosis until the four–cell stage using confocal microscopy was documented. To visualize the centrosomes and microtubules, eggs were stained with antibodies for γ- and α-tubulin. In addition to establishing normal development, two additional features of early development were documented. First, centrosome behavior was documented. The centrosome associated with the male pronucleus disappeared after polar body formation. The single centrosome associated with the male pronucleus duplicated by late anaphase of meiosis II to eventually form the spindle for mitosis I, thus the two centrosomes in the first cleavage were derived from the sperm. Second, linkage between the cell and centrosome cycles was established. In the first and second mitosis, centrosome duplication began at early anaphase. At late anaphase, the centrosomes duplicated and were observed as two distinct units at each pole. Asynchrony was also observed at the four–cell stage between the large and small blastomeres, with chromatin and centrosome duplication showing more advanced stages in the large blastomere. This documentation will serve as a baseline for future studies on chromosome loss in polyploid oysters.

KEY WORDS: oyster, Crassostrea virginica, centrosome, microtubule, meiosis, cleavage

INTRODUCTION

Chromosome loss in Crassostrea species had been documented for both diploids (Thiriot-Quivreux et al. 1992, Leitão et al. 2001, de Sousa et al. 2011) and polyploids (Allen et al. 1996, McCombie et al. 2005, Zhang et al. 2010, Zhang et al. 2014). In a previous study on the subject, de Sousa et al. (2016) hypothesized the possible involvement of supernumerary centrosomes in early development and possibly in later ontology in the loss of chromosomes. To know what is unusual in (possibly) abnormal chromosome behavior in polyploids, it is critical to understand normal chromosome behavior. Surprisingly, little information is available on microtubule and centrosome behavior in early development of oysters and especially Crassostrea virginica. The objectives of this work were to document early development starting with meiosis in newly fertilized eggs through the first and second cleavage divisions, including centrosome inheritance and duplication observed by confocal microscopy.

Several studies have made cytological observations on early development in bivalves and can be roughly divided into two purposes. The first type concerns genetic improvement or chromosome manipulation, including polyploid induction, for aquaculture. Especially, for triploid production, which has high commercial value because of reproductive sterility, numerous studies have demonstrated meiotic and early zygotic development (Komaru et al. 1990b, Longo et al. 1993, Li et al. 2000) during chromosome set manipulation. Chromosome manipulation techniques include the induction of polyploidy with chemical reagents, such as, cytochalasin B and inactivation of chromosomes using UV irradiation (Longo et al. 1993, Li et al. 2000). The main target of chromosome manipulation in these studies was inhibition of polar body (PB) formation or first cleavage division. Most of these studies observed and described chromosome movements using normal light (Crassostrea virginica: Longwell & Stiles 1968), electron microscopy (Crassostrea gigas: Longo et al. 1993), or epifluorescence microscopy (Pinctada fucata: Komaru et al. 1990a, 1990b; Chlamys nobilis: Komaru & Wada 1991; C. gigas: Li et al. 2000). Work of this nature has contributed to the production of triploid scallops, clams, and mussels—some on at least a semicommercial scale (Beaumont & Fairbrother 1991, Nell 2002).

The second type of cytological study of early development in bivalves concerns investigations in developmental biology and/or cytology. Marine bivalves are excellent candidates for meiotic and early zygotic studies (Guo & Allen 1997) as reported by Longo (1972), Guo et al. (1992a, 1992b), and Longo et al. (1993). The main advantage of this system, compared with insect and mammalian ones, is that marine bivalves have high fecundities and practice external fertilization. For example, a 2-y-old female Pacific oyster, Crassostrea gigas, may produce 20–50 million eggs (Guo et al. 2009). Eggs from bivalves, in general, are smaller and contain less yolk material than eggs from fish and amphibians. Moreover, because meiosis is arrested at prophase I in the follicle, meiosis I and II can both be visualized in mature bivalve eggs after controlled fertilization. For example, Kuriyama et al. (1986) showed the microtubule cycle in oocytes of Spisula solidissima using fluorescence microscopy and immunological methods. Surprisingly, few studies have been done on the meiotic and mitotic chromosome behavior in Crassostrea spp. Longwell and Stiles (1968, 1973) reported chromosome behavior stained by aceto-orcein at fertilization, meiosis, and early cleavage in Crassostrea virginica using optical microscopy. Li et al. (2000) compared nuclear behavior between normal and gynogenetic.
eggs of *Crassostrea gigas* using DAPI staining and epifluorescence microscopy, Longo et al. (1993) showed the course of fertilization and meiosis in CB-treated and untreated *C. gigas* eggs using electron microscopy and immunofluorescent techniques for the observation of DNA and microtubule organization, only from meiosis through metaphase of mitosis I. They discussed the behavior of centrosomes based on the organization of microtubules in the chromosome assemblages. Actual centrosomal behavior, however, has not been reported and there is little information for meiosis and even less for early cleavage. More contemporary techniques allow detailed visualization of complete meiotic and mitotic cytological structure and organelles. For example, Pielak et al. (2003, 2004, 2005) were able to elucidate the interaction between microtubule asters and cortical actin in *Spisula solidissima* oocytes using confocal microscopy. Several studies showed localization of centrosomes and nucleolus in *S. solidissima* oocytes using confocal microscopy with *in situ* hybridization (Alliegro & Alliegro 2005, 2008, 2013, Alliegro et al. 2006, 2010, 2012). Confocal microscopy affords exceptional clarity into the cytol-ogy of development. In this article, meiotic and early zygotic developments were documented in normal diploid *C. virginica* with confocal microscopy to document normal development.

**MATERIALS AND METHODS**

**Crosses Experiment and Sampling**

Two sets of pair matings were produced in the hatchery on August 9 (Cohort 1) and August 16 (Cohort 2), 2016. Ripe oysters from line DEBY.14 and hANA.14 held by Aquaculture Genetics and Breeding Technology Center were shucked and gonad biopsies were examined microscopically to determine the sex. The best specimens (maturity and fecundity) were used to make experimental crosses. Eggs for cohort 1 and 2 were stripped-spawned (Allen & Bushek 1992) from hANA females. Males were from the DEBY.14 line for cross 1 and hANA.14 for the cohort 2. Eggs were counted before fertilization to obtain appropriate densities of eggs in the beakers. Both cohorts were performed at 24.5°C in 14.1 psu-filtered seawater.

Samples were fixed at 0, 10, 30, 60, 80, and 100 min postfertilization, with the first three samples corresponding to periods of meiosis and the last three—mitosis. Oyster eggs are arrested before meiosis I and commence after fertilization. An appropriate volume of eggs was taken to assure at least 20,000 eggs were fixed.

**Sample Preparation for Confocal Microscopy**

**Fixation of Eggs and Staining**

Eggs sampled from beakers were screened through a 20-μm Nytex screen to remove cell debris and placed into a 15-mL Falcon tube containing 12 mL of fixative consisting of 0.1% TritonX-100, 4% formaldehyde in PEM (100-mM PIPES, 5-mM EGTA, 1-mM MgCl₂, pH 6.8, using NaOH). Eggs were simultaneously permeabilized and fixed in this fixative for 2 h. After fixation, the samples were rinsed three times with 10-mM glycine-phosphatase buffered saline (glycine-PBS pH7.4) and then washed with 10-mM glycine-PBS twice. The samples were stored in 10-mM PBS (without glycine) in 15-mL falcon tube and stored at 4°C until they were used for cytoskeletal labeling.

For microtubules and centrosomes, before staining, fixed eggs were incubated for 1 h in 3% bovine serum albumin (BSA)/PBS. Eggs were then washed once with glycine-PBS. Afterward, eggs were washed again, this time with 0.1% BSA/PBS (containing 0.1% TritonX). For staining, either of two primary antibodies was used, specific for either microtubules or centrosomes, in combination with a secondary antibody containing the fluorochrome. Microtubules were labeled with mouse monoclonal anti-α tubulin (Sigma-T9026) diluted to 0.5 μL/mL by 0.1% BSA/PBS (containing 0.1% TritonX) by incubating for 24 h at room temperature. Centrosomes were labeled with mouse monoclonal anti-γ tubulin (Sigma-T6557) diluted to 0.5 μL/mL by 1% BSA/PBS (containing 0.1% TritonX) as the primary antibody by incubating for 48 h at room temperature. After incubation, the eggs were washed three times with 0.1% BSA/PBS (containing 0.1% TritonX).

Eggs stained for microtubules or for centrosomes were labeled with the goat antimouse IgG (H + L) secondary antibody pretagged with Alexa Fluor 488 conjugate diluted to 5 μL/mL by 1% BSA/PBS (containing 0.1% TritonX). The eggs were washed three times with 0.1% BSA/PBS (containing 0.1% TritonX).

For chromosomes and nuclei, eggs labeled for either microtubules or for centrosomes were then counterstained with DAPI stock solution (DAPI 5 μg/mL, 2-mercapt-ethylamine hydrochloride 10 mM, Tris 10 mM, EDTA-2Na 100 mM, NaCl 100 mM) diluted to 0.5 μg/mL by 1% BSA/PBS (containing 0.1% TritonX) and incubated for 20 min to visualize the chromosomes. The labeled samples were washed twice with 0.1% BSA/PBS (containing 0.1% TritonX).

For mounting slides permanently, after removal of most of the supernatant, a suspension of eggs was added to about 60 μL of mounting medium (ProLong Gold Antifade Mountant) on the slide glass (Matunami Glass Ind., Ltd). The specific slide glasses were coated around the edges and uncoated in the center to avoid crushing the eggs when the cover slip was mounted (Alliegro et al. 2010). Eggs and mounting medium were aspirated gently to mix the eggs and medium. A cover slip was placed over the sample/mounting medium mixture and the edges sealed with nail polish.

Samples were viewed with a standard Olympus epifluorescence microscope and by confocal microscopy using an Olympus Laser Scanning Fluoview FV1200 with image processing via FV10-ASW 4.2 Viewer (Olympus). Images are maximum intensity projections of the entire z-stack. In addition, an optical microscope BX-51(Olympus) using cellSens (Olympus) image analysis software was used for estimating egg diameter.

**Measurements**

Egg diameter was determined under 100× magnification in fixed eggs by taking four diameter readings at each egg. Eggs in which the maximum egg diameter differed from a minimum egg diameter by more than 10 μm were not analyzed because of their probable immaturity. Average egg diameter from each diploid female was statistically analyzed with a Welch’s t-test at P = 0.05.

Spindle size was defined as the distance between two centrosomes. Measurement of spindle size was determined on a photograph and measured using imaging software FV10-ASW 4.2 Viewer. In meiosis I, the images were taken with the following three conditions: (1) Developmental stage was...
meiosis I metaphase. (2) The peripheral centrosome was attached to the egg cortex. (3) The images from the confocal were taken with the 2D-mode and the two centrosomes had to be in the same section. The thickness of single optical section in the 2D-mode was theoretically 1.34 μm. For measuring the first mitotic division (MT1), the distance between centrosomes was determined under following two conditions: (1) Developmental stage was metaphase of MT1. (2) The images were taken by the 2D-mode with confocal microscopy and two centrosomes were in the same section.

RESULTS

Egg Morphology

The overall morphology of eggs obtained from stripped follicles after incubation in seawater for 30 min took two forms: (1) Eggs were round (Fig. 1A) or (2) eggs were teardrop shaped (Fig. 1B). Uneven distribution of egg yolk was observed in both forms with optical microscopy. Egg diameter was 47.0 ± 1.9 μm (n = 73) for the first cohort and 49.1 ± 1.8 μm (n = 93) for the second, which was significantly different (P < 0.001) between the two diploid females used in the cohort. Teardrop-shaped eggs rounded up after fertilization.

Meiosis

Microtubules and centrosomes were visualized by immunostaining of the eggs and maternal and paternal chromosomes with DAPI staining. A spindle was formed by two centrosomes and microtubules spread radially from each centrosome. The meiotic spindles were located near the animal pole. Peripheral astral microtubules were oriented near the animal pole and spread along the cell cortex (Fig. 1C). Central astral microtubules were spread radially at the center of the egg. In most eggs, one centrosome was attached to the cortex of oocytes and the other centrosome was located centrally. The average distance between the two centrosomes was 14.4 ± 1.3 μm (n = 50) for the first cross and 13.9 ± 1.2 μm (n = 50) for the second. There was a significant difference between the two crosses, i.e., egg source (P = 0.031). Chromosomes derived from the egg were arranged along the equatorial plane of the meiotic spindle. Karyokinesis was observed to proceed perpendicular to the egg surface (Fig. 1D, E). After extrusion of the first polar body (PB1), the second meiotic spindle appeared. During the transition from the first to the second meiotic division, the maternal chromosomes never appeared in a decondensed state. At sampling periods immediately after cytokinesis, chromosomes were observed at the start of second meiosis. When the second meiotic spindle appeared, it was not always perpendicular to the egg surface, but later was observed perpendicular to the egg surface for the release of second polar body (PB2) (Fig. 1F). Extruded PB1 and PB2 were very close together. In some cases, the number of polar bodies on the egg surface could not be counted accurately.

Male Pronucleus Transformation and Centrosome Inheritance

Just after insemination, the sperm nuclei in the cytoplasm of eggs were observed still in a condensed (Fig. 2A) state. Sperm could be seen at various locations around the perimeter of the egg. The centrosome was not observed at the periphery of the sperm nucleus at early metaphase II (Fig. 2A). At meiosis II, one centrosome was observed beside the sperm nucleus (Fig. 2B). The centrosome originating from the sperm duplicated by late anaphase of meiosis II (Fig. 2C). When the release of the PB2 was complete, the sperm nucleus began to swell (Fig. 2D), with two centrosomes still associated with it. The female pronucleus also swelled, but the centrosomes associated with the female pronucleus during meiosis disappeared after PB2 formation. At this time, duplicated centrosomes migrated to opposite sides of the male pronucleus (Fig. 2E). The network of microtubules began to build in the egg cytoplasm (not shown, but could be observed with microtubule staining). Syngamy occurred as the swollen male and female pronuclei combined in the center of the egg, with the two centrosomes derived from the male pronuclei (Fig. 2F). Intermingled chromosomes began to condense and line up on the equatorial plane for first mitosis (Figs. 2G and 3H).

Mitosis—First Cleavage and Mitotic Centrosome Duplication

Chromosomes from the male and female pronuclei gathered and aligned on the equatorial plane of the spindle organized by

![Figure 1. Oocytes 30 min after strip spawning (A, B) observed by light microscopy and newly fertilized eggs of Crassostrea virginica by fluorescence microscopy (C–F). Microtubules (C, E, F) and centrosomes (D) were labeled with α- or γ-tubulin antibody, respectively. Chromosomes were located on the metaphase plate of first meiosis (C, D). (E) Early anaphase of first meiosis. (F) Metaphase of second meiosis. The first polar body was on the egg surface. Scale bar = 10 μm.]
the two centrosomes (Fig. 3A). The spindle was organized in the center of the egg. The spindle axis of the first cleavage formed perpendicular to the axis of the animal–vegetal pole. At metaphase of MT1, the average distance between the two centrosomes was 17.56 \pm 1.9 \mu m (n = 10) for the first cohort and 15.9 \pm 1.0 \mu m (n = 11) for the second, with a significant difference (P = 0.045) between the two cohorts. Beginning at anaphase of MT1, the fluorescent signal of the centrosome diminished, probably because the centrosome was undergoing duplication. Two fluorescent signals were recognized at each pole of the spindle (Figs. 3B and 4C). Also, at late anaphase of MT1, the polar lobe (PL) formed at the vegetal pole (Figs. 3C and 4D) and was later absorbed into one of the two cells of the first cleavage to form large and small blastomeres. Cytokinesis was observed at times
synchronous to the absorption of the PL. The cleavage furrow was colocated with the position of the extruded polar bodies at the animal pole. Also, the midbody between the cells became apparent at telophase (Fig. 3D), apparently decreasing with the contraction of the contractile ring.

At early anaphase of MT1 (Fig. 3E), centrosomes were distinct, one at each pole of the spindle. The centrosomes were elongated. But by late anaphase, they had begun to duplicate and were observed as two distinct signals at each pole (Fig. 3F). By telophase, segregated chromosomes had started to decondense (Fig. 3G) with duplicated centrosomes.

Two–Cell

After the formation of two cells (Fig. 3H), nuclei swelled and chromatin in the nuclei was observed in a more condensed state. A network of microtubules was evident throughout the egg cytoplasm at this stage (Fig. 4A). The astral microtubules that had nucleated from each centrosome were duplicated in late anaphase of MT1 and had migrated to opposite poles of their respective nuclear masses and started to form the spindle poles (Fig. 4B). At metaphase of MT1, chromosomes segregated after aligning on the equatorial plane of each spindle. This seemed to occur simultaneously in both blastomeres (Fig. 4C). The second karyokinesis of the two blastomeres, however, was asynchronous in the vast majority of observations (Fig. 4D). In the eggs observed that were asynchronous, chromosome segregation in the large blastomere was seen before those in the small blastomere. The PL again formed and was then absorbed in the large blastomere at anaphase. At the telophase of the second mitotic division (MT2), the width of the midbody that formed between each blastomere decreased with the contraction of the contractile ring as in the first cleavage.

Four–Cell

The blastomere that absorbed the PL became the large blastomere (Fig. 4E). The other cells appeared to be of equal size. Chromosome decondensation occurred simultaneously in all four cells (Fig. 4F), whereas a network of microtubules was built in the cytoplasm, as observed in former stages. The large blastomere was the first to form the next mitotic spindle in the four–cell stage (Fig. 4G). As the mitotic apparatus of the large blastomere became organized with chromosomes at the metaphase plate, the chromosomes of the three small blastomeres were less condensed and the aster not completely segregated. Figure 4D shows the four–cell stage. In the large blastomere, centrosomes were already duplicated at the poles for the next division. In the small blastomeres, the centrosomes were observed as one unit located on both sides of the condensed chromosomes, but not on the metaphase plates. Thus, at the four cell stage, asynchrony was obvious.

DISCUSSION

Egg Structure at Fertilization

In this study, eggs were obtained by strip spawning (Allen & Bushek 1992), which is an increasingly common way for research and commercial spawning of oyster species. Other studies have documented early development of oyster eggs obtained by strip spawning. Longo et al. (1993) documented two morphologies of Crassostrea gigas fertilized egg from strip spawning. The significance of these two forms with respect to gamete maturation, fertilizability, and potential for development was unclear. It was assumed that teardrop-shaped eggs were immature, having been torn from their follicle walls where they were still attached. Such eggs generally round up after time and are equally fertilizable, but this has not been tested directly. Nonetheless, egg diameter was measured only on round eggs.
On average between the cohorts, the diameter of eggs from diploid *Crassostrea virginica* was 48.2 μm. The egg size differing significantly from each other (female from first cohort versus female from second) was surprising because the egg size of *C. virginica* is highly regulated, undergoing constant balancing selection between fecundity and nourishment (Powell et al. 2011). At the same time, eggs from the two cohorts came from two different highly selected lines and the effect of domestication on the egg size is unknown.

**Meiosis**

Meiotic maturation in molluscan species is characterized by two general types: eggs with a secondary arrest in the first meiotic metaphase or eggs with no secondary arrest. In some bivalves, such as *Spisula or Barnea*, spawned oocytes are arrested at prophase I. Fertilization occurs at this stage leading to reinitiation of meiosis (Dubé & Guerrier 1982, Colas & Dubé 1998). By contrast, bivalves, such as *Ruditapes* and *Crassostrea* (Kuraishi & Osanai 1988), undergo germinal vesicle breakdown after spawning and are then arrested at metaphase of meiosis I. The *Crassostrea gigas* eggs in this study were incubated for at least 30 min after strip spawning, and most of the eggs were arrested at metaphase of meiosis I. In some cases, the meiotic spindle pole was unattached to the cortex of the animal pole (Fig. 2A). It seemed clear that the spindle migrates to the animal pole eventually. Kuriyama et al. (1986) and Pielak et al. (2003) described similar meiotic spindle behavior using *Spisula* oocytes. Longo et al. (1993) also documented spindle formation during meiosis in *Crassostrea gigas*. Interestingly, Longo et al. (1993) proposed that this spindle migration in bivalves can occur without microfilaments.

The male pronucleus condensed and remained quiescent until PB2 extrusion in the egg cytoplasm, at which point it began to swell. Similar findings were reported in previous work (*Crassostrea gigas*: Longo et al. 1993, Stephano & Gould 2000, *Asterina miniatata*: Stephano & Gould 2000; *Corbicula sandai and Corbicula fluminea*: Obata et al. 2006). On the other hand, duplicated centrosomes were observed at late anaphase of meiosis II (Fig. 3C). The paternal centrosome may have been duplicated at the same time as PB2 extrusion or before. Stephano and Gould (2000) proposed that MAP kinase is involved in sperm centrosome suppression based on the observation for microtubule organization in which MAP kinase is inhibited during meiosis in oyster eggs.

**Pronuclei and Centrosome Inheritance**

Longo et al. (1993) investigated the course of fertilization events in CB-treated *Crassostrea gigas* eggs using electron microscopy and immunofluorescent techniques for the observation of DNA and microtubule organization. Based on microtubule formation, they suggest that centrosomes of the embryo of *C. gigas* are paternally inherited (Longo et al. 1993, Longo 1997). Actual centrosome behavior was not reported, however. Observations from this study followed the behavior of centrosomes from metaphase of meiosis I through metaphase I and clearly showed that centrosomes are inherited from sperm whereas maternal centrosomes disappear (Fig. 2D). Therefore, it is evident from this study that two centrosomes derived from spermatozoa act as microtubules organization center of MT1. More correctly, the one centrosome derived from sperm duplicated just after PB1 extrusion. If oocytes were to retain a fully functional maternal centrosome capable of replication, introduction of a paternal centrosome would result in the presence of two replication-competent centrosomes and lead to the formation of tetrapolar spindles and disruption of chromosome segregation or aneuploidy (Palazzo et al. 1999). Thus, the ability of the maternal centrosome to replicate seems lost during the completion of meiosis, leaving the paternal centrosome to replicate as needed for coordinated development (Sluder et al. 1989, Schatten 1994).

In oocytes of starfish, *Asterina pectinifera* and the clams, *Spisula solidissima* and *Corbicula sandai*, the ability of the maternal centrosome to replicate is selectively destroyed during meiosis II (Kuriyama et al. 1986, Sluder et al. 1989, 1993, Obata et al. 2006, Borrego-Pinto et al. 2016). Importantly, in both *Asterina* and *Spisula*, the male centrosome is present in the egg cytoplasm when the ability of the maternal centrosome to replicate and to nucleate microtubules was destroyed (Sluder et al. 1989, 1993, Wu & Palazzo 1999). Yet, despite the deactivation of the maternal centrosome, maternal components are still required for restoration of the zygotic centrosome during organization of microtubules (Schatten 1994).

**Mitosis**

Normal cell division proceeds through the linkage of the cell cycle, the nuclear cycle, and the centrosome cycle in animal cells (Doxsey et al. 2005). The first two mitotic divisions were observed in this study. Although cell number was different, centrosomes and cytoplasmic structure exhibited similar behavior in both MT1 and MT2 in accordance with the nuclear cycle. Furthermore, the microtubule cycle was closely related to the centrosome cycle because centrosomes coordinate microtubule organization. The following description presents the linkage of nuclear cycle, cell cycle, and centrosome cycle in first two mitotic divisions of *Crassostrea virginica* zygotes.

**Metaphase to Anaphase**

The duplicated genome must be divided into two daughter cells by the spindle apparatus that is organized by centrosomes. Centrosomes are composed of two orthogonally arranged centrioles surrounded by an amorphous mass of protein, termed the pericentriolar material (Nigg 2002). One centrosome with two centrioles performs as a single microtubule organization center (Nigg 2002, 2007). In the initial step for *Crassostrea virginica*, chromosomes condensed and aligned on the equatorial plane of the spindle that was formed by microtubules radiating from the two centrosomes at metaphase. Chromosome segregation began after all chromosomes were aligned on the equatorial plane. Normally, the process is regulated by a spindle assembly checkpoint that monitors attachment of the spindle microtubules to kinetochores of chromosomes (Doxsey et al. 2005). Changes in cell structure and centrosome number during the metaphase to anaphase stage were not observed by us.

**Late Anaphase**

There were several changes in both cell structure and centrosome behavior in late anaphase of the two mitotic divisions. For the centrosome, although a single centrosome was observed at each pole at early anaphase of MT1, two distinct signals of γ-tubulin were recognized in each spindle pole at late anaphase. In more detail, these duplicated signals of γ-tubulin were
Allen, S. K. & D. Bushek. 1992. Large-scale production of triploid crassostrea gigas: Luetjens & Dorresteijn 1998; Corbicula sandat: Obata et al. 2006). On the contrary, the spindle of the first cleavage with or without PL formation. Polar lobe formation has been observed in other species during early development (e.g., Crassostrea virginica, centrosome duplication began at late anaphase of MT1. Furthermore, change of cell structure also started at late anaphase of early mitotic division. For MT1, the PL protruded from vegetal pole. For the second cleavage, the PL extruded from the large blastomere (data not shown). Borrego-Pinto, J., K. Smogyi, M. A. Karreman, J. König, T. Müller-Reichert, M. Bettencourt-Dias, P. Gönczy, Y. Schwab & P. Lénárt. 2016. Distinct mechanisms eliminate mother and daughter centrosomes in meiosis of starfish oocytes. J. Cell Biol. 212:815–827.

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BEATTIE, J. & A. BUSHEK. 1992. Large-scale production of triploid crassostrea gigas: Luetjens & Dorresteijn 1998; Corbicula sandat: Obata et al. 2006). On the contrary, the spindle of the first cleavage with or without PL formation. Polar lobe formation has been observed in other species during early development (e.g., Crassostrea gigas: Deno 1998; Crepidula fornicata; Henry et al. 2006). In embryos without a PL, the first mitotic division shifts to one side and one spindle pole attaches to the cortex (e.g., Dreissenia polymorpha: Luetjens & Dorresteijn 1998; Corbicula sandat: Obata et al. 2006). On the contrary, the spindle of the first cleavage is oriented in the center of the zygote. This is the typical pattern of the first cleavage with PL formation.

Telophase to Prophase

In Crassostrea virginica, the centrosomes organized the microtubules from metaphase to late anaphase to establish the spindle apparatus. The fluorescence of the astral formation diminished when the chromosomes decondensed and formed the nucleus. This loss of γ-tubulin signal is probably associated with disassembly of microtubules at late anaphase and seemed to be related to the centrosome cycle. γ-Tubulins are essential for nucleation of microtubules (Palazzo et al. 1999). The main feature of telophase was cytoplasmic division. The cleavage furrow was observed when segregated chromosomes decondensed and formed nuclei. Moreover, the midbody was observed between the two dividing cells at the end of cytoplasmic division. The midbody is common and synchronous with the formation of the contractile ring in animal cells (Steigemann & Gerlich 2009). At this time, synchrony was observed between cytoplasmic division and PL reabsorption. In contrast to this dynamic cell structural change, the two duplicated centrosomes apparently kept their position and structure. Important centrosomal behavior was observed when cell division was completed: two adjacent centrosomes separated and acted as microtubule organizing centers. Centrosome separation is known to be caused by the action of microtubules-dependent motor proteins, such as Eg5 (Smith et al. 2011). The two separated centrosomes formed a bipolar spindle and contributed to chromosome segregation. Asynchrony of the cell cycle in each cell at four-cell stage was also observed. Even though cell number and cell structure are different between first and second cell division, the cell cycle, nuclear cycle, and centrosome cycle are coordinated in both cell divisions.

In this article, detailed observations on meiotic and early zygotic development were presented. The results, obtained by double staining nuclear and microtubules or centrosomes clearly showed linkage of cell cycle, nuclear cycle, and centrosome cycle. This fundamental knowledge will be useful not only to understand the detailed mechanism of meiosis and early mitosis in the oyster, but also for future investigations into meiotic and mitotic abnormalities in polyploids of Crassostrea virginica.
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