

A cleavage clock regulates features of lineage-specific differentiation in the development of a basal branching metazoan, the ctenophore *Mnemiopsis leidyi*

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A cleavage clock regulates features of lineage-specific differentiation in the development of a basal branching metazoan, the ctenophore *Mnemiopsis leidyi*

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Abstract

Background: An important question in experimental embryology is to understand how the developmental potential responsible for the generation of distinct cell types is spatially segregated over developmental time. Classical embryological work showed that ctenophores, a group of gelatinous marine invertebrates that arose early in animal evolution, display a highly stereotyped pattern of early development and a precocious specification of blastomere fates. Here we investigate the role of autonomous cell specification and the developmental timing of two distinct ctenophore cell types (motile compound comb-plate-like cilia and light-emitting photocytes) in embryos of the lobate ctenophore, *Mnemiopsis leidyi*.

Results: In *Mnemiopsis*, 9 h after fertilization, comb plate cilia differentiate into derivatives of the E lineage, while the bioluminescent capability begins in derivatives of the M lineage. Arresting cleavage with cytochalasin B at the 1-, 2- or 4-cell stage does not result in blastomere death; however, no visible differentiation of the comb-plate-like cilia or bioluminescence was observed. Cleavage arrest at the 8- or 16-cell stage, in contrast, results in the expression of both differentiation products. Fate-mapping experiments indicate that only the lineages of cells that normally express these markers in an autonomous fashion during normal development express these traits in cleavage-arrested 8- and 16-cell stage embryos. Lineages that form comb plates in a non-autonomous fashion (derivatives of the M lineage) do not. Timed actinomycin D and puromycin treatments show that transcription and translation are required for comb formation and suggest that the segregated material might be necessary for activation of the appropriate genes. Interestingly, even in the absence of cytokinesis, differentiation markers appear to be activated at the correct times. Treatments with a DNA synthesis inhibitor, aphidicolin, show that the number of nuclear divisions, and perhaps the DNA to cytoplasmic ratio, are critical for the appearance of lineage-specific differentiation.

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Conclusion: Our work corroborates previous studies demonstrating that the cleavage program is causally involved in the spatial segregation and/or activation of factors that give rise to distinct cell types in ctenophore development. These factors are segregated independently to the appropriate lineage at the 8- and the 16-cell stages and have features of a clock, such that comb-plate-like cilia and light-emitting photoproteins appear at roughly the same developmental time in cleavage-arrested embryos as they do in untreated embryos. Nuclear division, which possibly affects DNA-cytoplasmic ratios, appears to be important in the timing of differentiation markers. Evidence suggests that the 60-cell stage, just prior to gastrulation, is the time of zygotic gene activation. Such cleavage-clock-regulated phenomena appear to be widespread amongst the Metazoa and these cellular and molecular developmental mechanisms probably evolved early in metazoan evolution.

Keywords: Ctenophore, Comb jelly, Photocyte, Comb cell, Cytochalasin B, Puromycin, Actinomycin, Cell cycle arrest, Cell lineage

Background

Ctenophores are a monophyletic group of seemingly simple marine animals with distinct features, such as unique comb rows, body symmetry and stereotypic cleavage program [1-11]. The name Ctenophore means ‘comb-bearing’ and refers to eight rows of comb or ctene plates, each possessing thousands of motile cilia arranged in linear arrays to form small paddles [12]. Each comb plate in a ctene row beats in a highly synchronized manner and they are used to move the animal through the water column [12]. The oral-aboral axis is the major body axis of ctenophores and it is characterized by the mouth at one pole and the statocyst-containing apical organ at the opposite (aboral) pole [3,4,12] (Figure 1A, B). The view from the aboral pole shows that ctenophores comprise four highly similar quadrants, which are separated by two orthogonal planes – the tentacular plane and the esophageal or sagittal plane (Figure 1A) [4,13-15]. These planes are defined by axes of rotational symmetry around the oral-aboral axis because diagonally opposed quadrants are more identical to one another than adjacent quadrants [3-6,10,13,15].

Many ctenophores, including the lobate ctenophore *Mnemiopsis leidyi*, generate distinct light-producing photocytes during development, which migrate to locations adjacent to each of the eight comb rows. Genome sequencing of *Mnemiopsis* has revealed two genomic clusters containing ten distinct copies of photoproteins closely resembling the luciferase-type photoproteins found at the base of the Metazoa. *In situ* hybridization studies have shown that at least subsets of these mRNAs are expressed in photocytes prior to when these embryos are bioluminescent [16]. Thus, the presence or absence of bioluminescence is a strong indicator of the developmental fate of a differentiated photocyte.

Ctenophore development is highly stereotypic and unique within the animal kingdom [1,4,5]. In the lobate ctenophore *Mnemiopsis leidyi*, embryos are generally fertilized at the time of spawning and cleavages are unipolar and

holoblastic (Figure 1, and 2 Additional file 1) [4-6,10,17]. Subsequent rounds of division occur every 15 to 20 min at room temperature and the juvenile cydippid stage hatches from the egg membrane within 24 h after the first cleavage. The second cleavage, which is perpendicular to the first cleavage, gives rise to four similar-sized EM blastomeres [1,4,10,18]. The third cleavage is oblique and results in the formation of four E cells (end cells) and four M cells (middle cells) [1,4,10,14,18] (Figure 2 and Additional file 1). Each M blastomere undergoes two rounds of asymmetric division, giving rise to two small m micromeres (m_1 and m_2) towards the aboral pole and one M macromere towards the oral pole [1,4,14]. E blastomeres undergo three asymmetric cell divisions each, producing three small e micromeres (e_1 , e_2 and e_3) at the aboral pole and an E macromere at the oral pole [1,4,14]. The micromeres proliferate further and begin to envelop the macromeres during gastrulation via epiboly at approximately 3 to 4 hours post fertilization (hpf) [4,6] (Figure 2 and Additional file 1). Later during development, the macromeres, which are located at the oral pole, generate another set of oral micromeres [4]. Gastrulation is complete around 5 to 6 hpf. Ciliated comb cells appear by 9 hpf at the same time bioluminescence is detected. Cydippid stage juveniles hatch after 18 to 24 hpf [1,4,6,18].

The stereotyped cleavage program in ctenophores allows each blastomere to be identified and its fate followed by the injection of intracellular lineage tracers [1,4,7] (summarized in Figure 1C). For example, the mesoderm, including muscle, mesenchymal cells and photocytes, is generated by the micromeres born from endodermal precursors at the future oral pole [4] (summarized in Figure 1C). Early labeling experiments identified that the e_1 micromeres give rise to the comb plate cilia [18,19]; however, later fate-mapping experiments were able to detect the m_1 micromere's contributions to the formation of comb plates (summarized in Figure 1C). Interestingly, when e_1 micromeres are deleted, no comb plate cilia form,

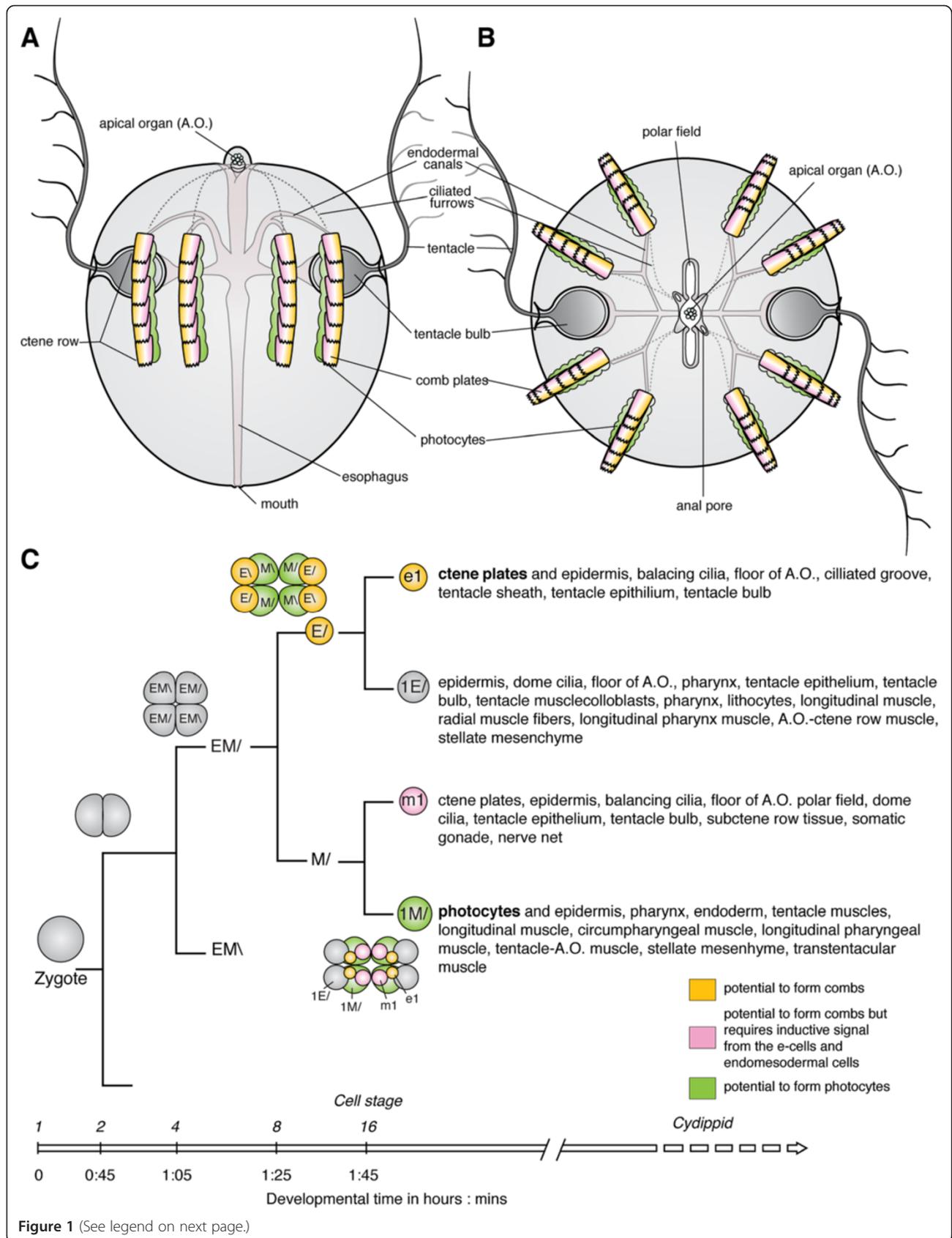


Figure 1 (See legend on next page.)

(See figure on previous page.)

Figure 1 Ctenophore development and cell lineage. Ctenophore body plan, with (A) lateral view, aboral pole up and (B) aboral view. The comb plates are shown in yellow to indicate the contribution from e1 micromeres (yellow) and m1 micromeres (pink). The photocytes, are a derivative of the 2 M cells and associated with the endodermal canals underlying the comb rows [4,16]. (C) Simplified cell lineage fate map showing the e₁, m₁, 1E/ and 1 M/ micromere sublineages. For each of the different early cleavage stages there are diagrams showing the embryo from the aboral pole. Modified after [1,4,14].

indicating that e₁ micromeres are autonomously specified to give rise to comb plates while m₁ micromeres require inductive signals [18,20].

Here, we investigate the role of the cleavage program in the segregation and expression of the developmental potential of two distinct cell types (motile comb-plate-like cilia and light-emitting photocytes) during the development of the ctenophore *Mnemiopsis*. Using an inhibitor of filamentous actin polymerization (cytochalasin B) to block cytokinesis (but not nuclear division), we show that the formation of differentiation markers associated with each of these cell types appears only after they have segregated into their own distinct lineages and they are never co-expressed in the same cells. Furthermore, using inhibitors of protein synthesis, transcription and DNA synthesis, we provide evidence for the existence of a cleavage clock that is based on the number of rounds of DNA synthesis (and possibly controlled by the nuclear-to-cytoplasmic ratio), which determines the temporal appearance of differentiation products in cleavage-arrested embryos.

Methods

Mnemiopsis collection and embryos

Mnemiopsis leidyi adults were collected from the National Oceanic and Atmospheric Administration jetty and Eel Pond in Woods Hole, MA, during June and July and from 13660 Deering Bay Dr, Coral Gables, FL 33158, during December. Adults were brought into the lab and induced to spawn as described previously [21]. Eggs were washed

multiple times with 0.2 μm filtered seawater to remove any jelly or debris. Only batches in which a high percentage of embryos developed normally were used in the study. Fertilization in these animals occurs at spawning, so for developmental timing purposes this was designated as 0 hpf.

Pharmaceutical inhibitors

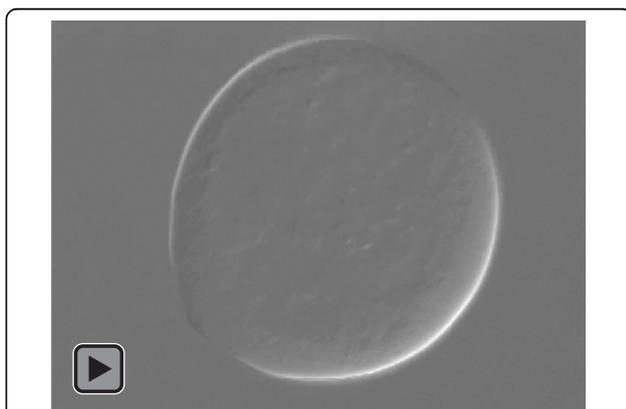
After the embryos were collected, they were transferred by pipet to 24-well culture dishes for drug treatments. The following drugs were used: cytochalasin B (Sigma, St. Louis, MO, USA, C6762), actinomycin D (Sigma, St. Louis, MO, USA, A1410), puromycin (Sigma, St. Louis, MO, USA, P7255) and aphidicolin (Sigma, St. Louis, MO, USA, A0781). Stock solutions of cytochalasin (1 mg/ml), actinomycin (1 mg/ml) and aphidicolin (1 mg/ml) were made up in dimethyl sulfoxide, with aliquots stored at -20°C. A puromycin stock solution (12.5 mg/ml) was made up in distilled water and stored at -20°C. Freshly thawed aliquots were used in each experiment. Working solutions were made by diluting stock solutions with filtered seawater. Approximately 1 ml of solution was added to each well containing embryos.

Hoechst staining and immunohistochemistry

The embryos were fixed for antibody staining in 4% paraformaldehyde and 0.02% glutaraldehyde, as previously described by Pang and Martindale [21]. Following fixation, the embryos were removed from their membranes by gentle pipetting, washed with PBS plus 0.2% Triton (PBT), blocked in 5% goat serum for 1 h and then incubated in anti-tyrosine tubulin (Sigma, T9028) overnight at 4°C. The embryos were washed six times for 30 min in PBT and incubated with the secondary antibody, goat anti-mouse conjugated to Alexa-594 (Invitrogen, Molecular Probes, Carlsbad, CA, USA), Alexa-488 phalloidin (Invitrogen, Molecular Probes, Carlsbad, CA, USA) and Hoechst 33342 (Invitrogen, Molecular Probes, Carlsbad, CA, USA) overnight at 4°C. Afterwards the embryos were washed twice for 5 min in PBS and mounted for imaging.

Lineage tracing

After the embryos were collected, the vitelline membranes were mechanically removed using sharpened forceps. The de-membrated embryos were allowed to develop to the correct cleavage stage prior to injection with DiI, (Catalog No. D-282; Molecular Probes, Inc.,



Additional file 1 Normal embryonic development of *Mnemiopsis leidyi*. DIC microscopy time-lapse movie of developing *Mnemiopsis* embryos beginning with a zygote until 12 hpf. The aboral side is up. The animal shown is the same as in Figure 2A-P.

Eugene, OR) a lipophilic membrane stain that diffuses laterally to stain the entire cell, as described in [4]. Following DiI injection, the embryos were carefully transferred into individual wells of a Terasaki plate (Nunc, Roskilde, Denmark) so that each embryo remained separate. Each well of the plate contained 10 μ l of filtered seawater (controls) or cytochalasin B solution (1 μ g/ml). The Terasaki plate was then stored in a humidified chamber to minimize evaporation.

Imaging

The embryos were scored for the presence or absence of comb cells using a Stereo Discovery (Zeiss, Inc) or Axio Scope (Zeiss, Inc) under transmitted light. Live comb plate cells were imaged using time-lapse microscopy with time intervals of five to ten images per second. Normal *Mnemiopsis* development was recorded at three images per minute using DIC (Differential interference contrast). Visualization of bioluminescence was performed on an

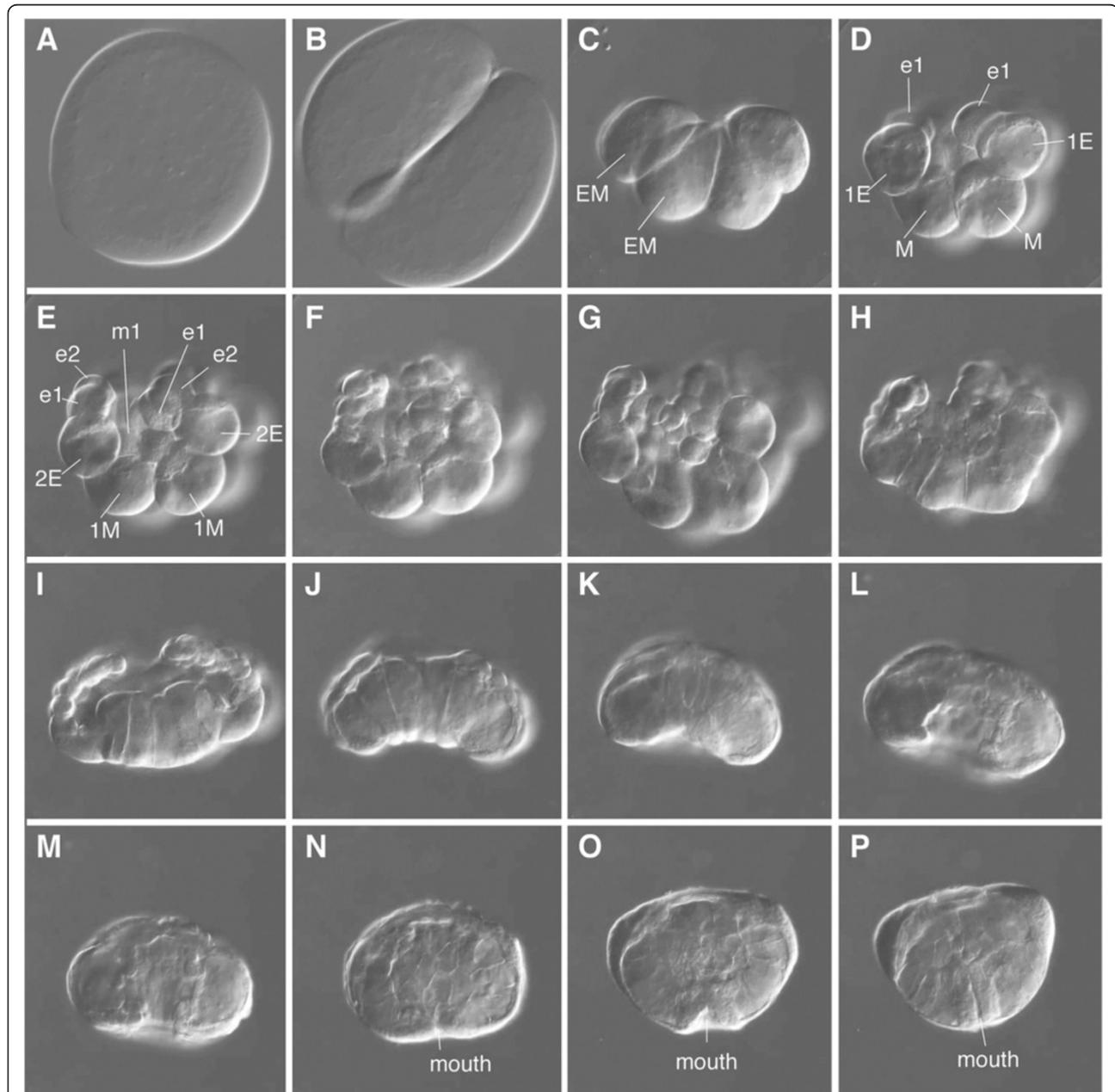


Figure 2 Embryonic development of *Mnemiopsis leidyi*. DIC images of developing *Mnemiopsis* embryos at various stages, beginning with a zygote in (A) until 9 hpf in (P). (A) Zygote. (B) 2-cell stage. (C) 4-cell stage. (D) 8-cell stage. (E) 16-cell stage. (F, G, H) 32- to 60-cell stages. (I, J, K, L) Gastrulation. (M, N, O, P) Post-gastrulation. The aboral side is up. hpf, hours post fertilization. Additional file 1 shows the embryonic development of the same specimen in a time-lapse movie.

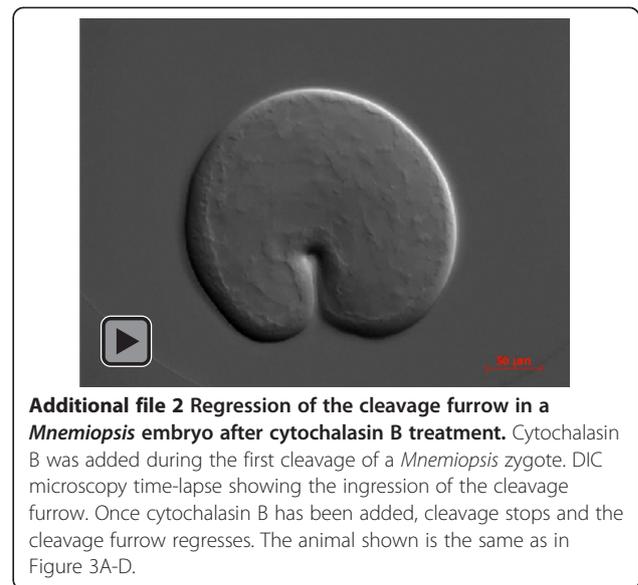
Axio Scope, using the GFP filter (38 HE Green Fluorescent Prot. filter, excitation BP 470/40 nm, emission BP 525/50 nm) to stimulate photoprotein emission and with 0.5 to 1 sec exposure times with an AxioCam Mr to capture faint signals. Prior to visualization, an embryo was kept undisturbed in darkness for 10 to 20 min to prevent activation and depletion of the photoprotein.

Confocal imaging after immunohistochemical staining was performed using a Zeiss 710 confocal microscope. Images were processed using Zen software (Zeiss, Inc) and Volocity (Improvision, Inc) to create 3D image reconstructions of confocal sections.

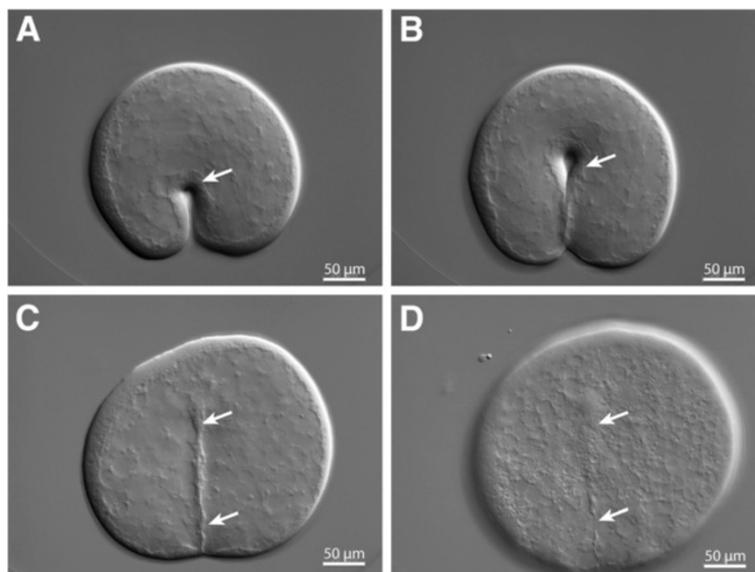
Results

Cytochalasin B inhibits the polymerization of actin and thus prevents cytokinesis. During embryonic development, individual cleavages and specific developmental stages can be arrested by cytochalasin B, and thus the developmental potential of cells that were present at the time of arrest can be analyzed. We used cytochalasin B on zygotes, the 2-, 4-, 8- and 16-cell stages and at later time points up to the 60-cell stage (at 3, 3.5, 4, 5, 6, 7, 8 and 9 hpf). An overview of *Mnemiopsis* development is shown in Figure 2 and Additional file 1.

Cytochalasin B treatment of *Mnemiopsis* embryos led to the immediate arrest of cytokinesis. If the embryo was undergoing cell division at the time of treatment, the cleavage furrow immediately began to regress and the blastomeres rounded up (Figure 3, Additional file 2). Throughout the cytochalasin B treatment, the blastomeres



did not die and cell nuclei continued to divide as karyokinesis does not require actin polymerization [22]. For the first 4 to 5 h of treatment, the nuclei divided at the normal rate and remained in the periphery of the cell (Figure 4A), but after approximately 5 hpf, individual nuclei fused together into one or several large nuclei (Figure 4B, C, D). Cleavage-arrested blastomeres adhered tightly to each other until 7 to 8 hpf, after which individual cells tended to lose contact with one another. Although the normal configuration of the blastomeres was lost, they



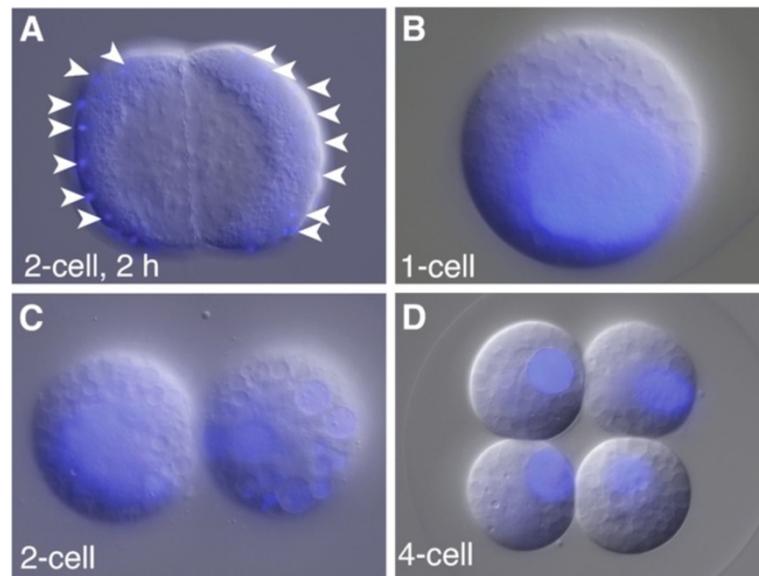


Figure 4 Nuclear divisions in cleavage-arrested embryos continue. Nuclei continue to divide after cytochalasin B treatment resulting in multiple nuclei in each cell, which later fuse into one or several large nuclei per cell. DNA staining with Hoechst (blue) merged with DIC images. **(A)** Embryo inhibited with cytochalasin B at the 2-cell stage. Multiple nuclei are visible along the periphery of the cells after 2 h of treatment. **(B)** A zygote arrested with cytochalasin B with one overly large nucleus. **(C)** Embryo inhibited with cytochalasin B at the 2-cell stage, with one large nucleus in the left cell and several large nuclei in the right cell. **(D)** Embryo inhibited with cytochalasin B at the 4-cell stage. Each cell has one large nucleus. Since all the embryos were inhibited prior to the 8-cell stage, none of them have differentiated comb cells.

were still contained within the vitelline membrane, allowing for the identification of all descendants from an individually arrested embryo.

While cytochalasin-treated zygotes and 2- and 4-cell stages did not show any visible sign of cell differentiation even after 24 hpf (Figure 4), surprisingly, 70% (31/44) of arrested embryos treated at the 16-cell stage (Figure 5F) and 83% (19/23) of embryos treated at the 32-cell stage had ctene-like cilia around 9 hpf. When treated at the 60-cell stage and onwards, nearly 100% of all arrested embryos formed ctene-like cilia. And when treated with cytochalasin B at the 8-cell stage, 26% (9/35) of the arrested embryos had two or more cells that formed motile comb-plate-like compound cilia (Figure 5, Additional file 3, Additional file 4, Additional file 5). These cilia were approximately the same size and formed at the same time as for untreated control embryos (Figure 5G, Additional file 6). A closer look at the ctene cells in treated embryos revealed that individual comb-plate-like cilia appeared morphologically normal with rows of cilia beating back and forth (Figure 5A, B, C, H, H', I, I', Additional file 3, Additional file 4, Additional file 5). Ctenophores make other kinds of ciliated structures (for example, dome cilia, sensory pegs, ciliated grooves and balancing cilia) but most of these are not motile, and none of them comprise compound cilia such as those made by the cleavage-arrested cells. Many individual cleavage-arrested cells formed multiple motile comb-plate-like cilia;

these combs were not arranged in discrete rows as is the case during normal development [23] and their beating was not coordinated and appeared random with respect to one another and to neighboring cells. The cilia of the comb-plate-like generating cells were seen beating for up to 18 hpf and these cells quickly separated from the other cells in the embryos.

Ctene cells in cleavage-arrested embryos are derived from e_1 micromeres

During normal *Mnemiopsis* development, ctene cells are formed by e_1 micromeres and m_1 micromeres [4]. However, while e_1 micromeres can differentiate into comb-bearing cells autonomously, m_1 micromeres require an inductive signal from the E lineage [1]. To determine which cell lineages were forming the comb cells in cleavage-arrested embryos, we utilized DiI labeling. Labeling of all four E blastomeres at the 8-cell stage and arresting the embryo at the 8- or 16-cell stage led to embryos in which all cells producing comb-plate-like cilia were labeled (Table 1, Figure 6A, B, C, E, F, F"). The same was true after labeling all four E macromeres at the 8-cell stage, allowing them to undergo one more cell division (to form DiI-labeled 1E macromeres and e_1 micromeres) before arresting them at the 16-cell stage (Table 1, Figure 6F, F', F"). These experiments show that all cells that gave rise to comb-plate-like cilia in cleavage-arrested embryos were derived from the E lineage.

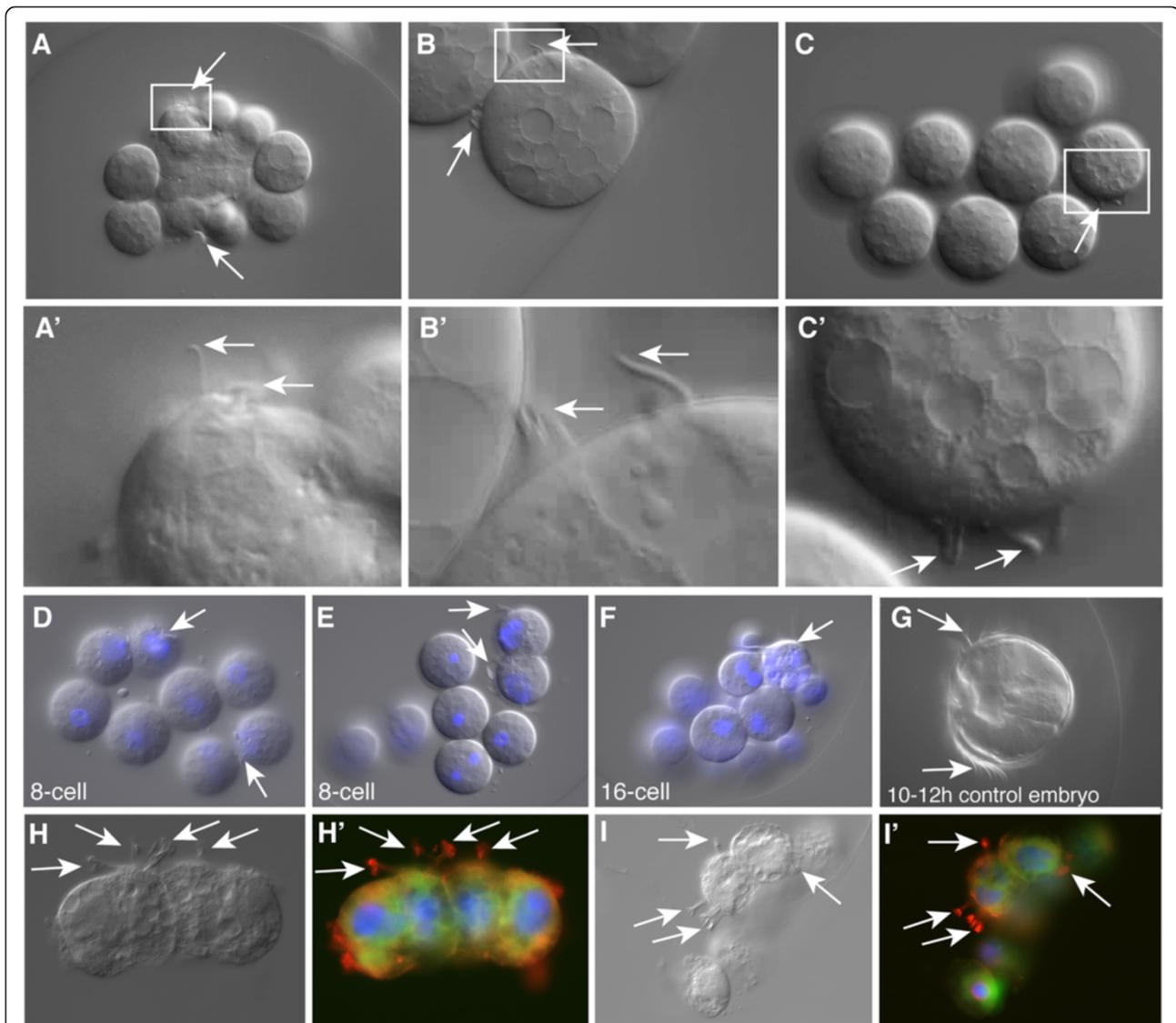


Figure 5 Comb cells in cleavage-arrested embryos. The arrows point to the cilia of the differentiated comb cells. **(A, B, C)** DIC images of cleavage-arrested embryos. **(A', B', C')** Close-up views of the rectangles in **(A, B, C)**. **(D, E, F)** DNA staining with Hoechst (blue) merged with DIC images of embryos inhibited with cytochalasin B at the 8-cell stage **(D, E)** or 16-cell stage **(F)**. **(G)** Control embryo at 10 to 12 hpf. DIC images **(H, I)** and z-projections of confocal images **(H', I')** of embryos stained with anti-tyrosinated tubulin (red) showing the cilia, Alexa-488 phalloidin (green) showing cell borders and Hoechst 33342 (blue) showing the nuclei. hpf, hours post fertilization. Additional file 3, Additional file 4, Additional file 5 are movies showing the motility of the cilia in the specimens shown in **A, B** and **C** respectively. Additional file 6 shows the movement of the cilia in the comb rows of the specimen shown in **G**.

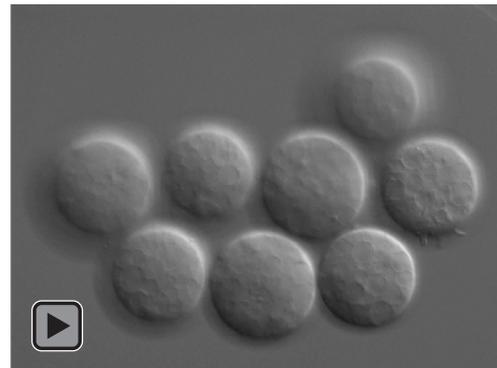
To rule out the possibility that DiI labeling could induce the cell fate of the comb-plate-like cilia, we labeled all four M blastomeres at the 8-cell stage or all four 1M macromeres and m_1 micromeres at the 16-cell stage and arrested the embryos with cytochalasin B. In only one case (which was likely to have been generated from a misinjected cell), were the cells producing the comb-plate-like cilia labeled (Figure 6D, G).

To determine whether e_1 cells specifically gave rise to the comb plates, we labeled all four e_1 micromeres at the 16-cell stage and immediately afterwards treated the

embryos with cytochalasin B. All of the resulting embryos formed cells producing comb-plate-like cilia that were labeled with DiI (Figure 6E, Table 1, Additional file 7). We also labeled all four 1E macromeres after the division of the E blastomeres into e_1 and 1E at the 16-cell stage and cleavage arrested the embryos with cytochalasin B. None of the cells producing comb-plate-like cilia that formed were labeled (Table 1). These data demonstrate that only E lineage blastomeres at the 8-cell stage and e_1 micromeres at the 16-cell stage give rise to comb-bearing cells. 1E macromeres, but not M lineage



Additional file 3 Cleavage-arrested embryo with beating cilia at 2 comb cells. DIC microscopy movie showing the beating cilia on comb-cells in a cleavage-arrested embryo. The embryo shown is the same one as in figure 5A and 5A'.



Additional file 5 Cleavage-arrested embryo with beating cilia at a comb cell. DIC microscopy movie showing the beating cilia on comb-cells in a cleavage-arrested embryo. The embryo shown is the same one as in figure 5C and 5C'.

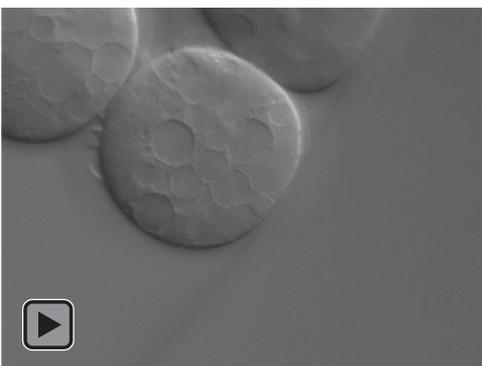
descendants, appear to be able to contribute to the formation of comb-plate-like cilia.

Photocytes are specified by the M lineage in cleavage-arrested embryos

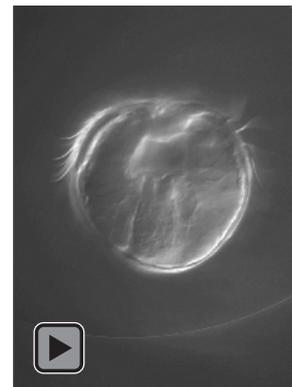
To determine whether the M lineage also gives rise to products specific to differentiated cells in cleavage-arrested *Mnemiopsis* embryos, we looked for the presence of bioluminescence or light production. In *Mnemiopsis*, bioluminescence is generated by the expression of luciferase-like photoproteins in a specific cell type, the photocytes [11,16]. Fate-mapping experiments have shown that during development, the 2M macromeres give rise to photocytes [4,16,24,25]. Photocytes are normally located in the endodermal canals underneath the comb rows [11,26] (Figure 7A, A', A'', B). To visualize the photoprotein, embryos were acclimatized to the dark and imaged under a GFP filter set on a Zeiss compound

microscope (Additional file 8). The blue fluorescent light excites the photoprotein in these cells causing them to emit light at around 485 nm to 496 nm [16,27]. In living cydippids, the luminescence appeared as green specks and faded away within 1 to 3 sec, which can be seen by the naked eye and was recorded by a CCD camera.

In embryos that were treated with cytochalasin B at the 8- or 16-cell stage, we observed green bioluminescence in the cleavage-arrested blastomeres. This began at the same time as in control embryos at around 9 hpf (Additional file 9). Because DiI labeling of M blastomeres masked the weak fluorescence of the photoproteins in injected embryos, we labeled the E blastomeres at the 8-cell stage with DiI. Observation of DiI-injected E lineage cleavage-arrested embryos at the 8- or 16-cell stage revealed that the photoprotein was located in the unlabeled M (8-cell) and 1M macromeres (precursors to the 2M lineage) at the 16-cell stage (Figure 7C, D, E, F).



Additional file 4 Close-up on one a comb cell with beating cilia in a cleavage-arrested embryo. DIC microscopy movie showing the beating cilia on comb-cells in a cleavage-arrested embryo. The embryo shown is the same one as in figure 5B and 5B'.



Additional file 6 Control embryo at around 10-12 hpf. DIC microscopy movie showing the beating cilia of the comb cells, which are well visible on both sides of the embryo. The embryo shown is the same one as in figure 5G.

Table 1 Overview of the cytochalasin treatment experiments on *Mnemiopsis leidyi* embryos

Developmental stage	Treatment	Observation	Conclusion
8-cell stage	Cytokinesis arrested with cytochalasin B	50% have combs	All combs that were formed were labeled
16-cell stage		75% have combs	
32-cell stage		83% have combs	
8-cell stage	Cytokinesis arrested with cytochalasin B, labeling of all four E blastomeres	62% (8/13) have combs that are labeled 38% (5/13) did not form combs	
16-cell stage	Cytokinesis arrested with cytochalasin B at 16-cell stage, labeling of all four E blastomeres at the 8-cell stage	95/128 (74%) formed combs that are labeled 32/128 (25%) did not form combs 1/128 (0.8%) formed combs that are not labeled	99% of embryos that formed combs were labeled, E cell lineage forms combs
16-cell stage	Cytokinesis arrested with cytochalasin B at the 16-cell stage, labeling of all e ₁ micromeres at the 16-cell stage	27/36 (75%) formed combs that are labeled 9/36 (25%) did not form combs	100% of embryos that formed combs were labeled, e ₁ cells form combs
16-cell stage	Cytokinesis arrested with cytochalasin B at the 16-cell stage, labeling of all 1E macromeres at the 16-cell stage	No labeled combs	Only e ₁ cells at 16-cell stage form combs and E1 cells do not form combs
8-cell stage	Cytokinesis arrested with cytochalasin B at the 8-cell stage, labeling of all M blastomeres	19/27 (70%) formed combs that are not labeled 7/27 (26%) did not form combs 1/27 (4%) formed combs that are labeled	95% of the embryos formed combs that were not labeled and combs do not arise from M lineage cells
8-cell stage and 16-cell stage	Cytokinesis arrested with cytochalasin B at the 8-cell stage or 16-cell stage, all E blastomeres were labeled	Unlabeled cells have photoprotein	M cells can make photoprotein even when they are cell-cycle arrested

Transcription and translation are required for the formation of comb-plate-like cilia in cleavage-arrested embryos

The differentiation of comb-plate-like cilia in E lineage cells and photoproteins in M lineage descendants in cleavage-arrested embryos suggests that developmental potential is faithfully segregated to the correct spatial lineages by the early cleavage program. However, it is unclear what key factors may be segregated into the lineages. To gain insight into the molecular nature of the determinants for the formation of comb-plate-like cilia, we used inhibitors of mRNA and protein synthesis to see if these classes of molecules could be responsible for the expression of these structures.

We treated embryos at various stages, beginning with zygotes up to the time that comb-plate-like cilia are normally generated in intact embryos (9 hpf), with 10 µg/ml of actinomycin D, an inhibitor of transcription [28,29]. Embryos that were treated up to 6 hpf, did not form comb plates (Table 2), suggesting that the localized

determinant is not solely an mRNA (possibly maternal in nature) that is required for comb-plate formation. These results suggest that additional transcription is necessary for the comb plates to form in descendants of the E lineage.

It is interesting to note that embryos that were treated at the 1-, 2-, 4-, 8- and 16-cell stages continued to divide normally and arrested only prior to gastrulation at about the 60-cell stage. Embryos that were treated with actinomycin D at the 32- or 60-cell stage divided only a few more times and then became arrested (Table 2). This suggests that if actinomycin D is inhibiting the bulk of mRNA synthesis, then zygotic transcription appears to begin at about the 32- or 60-cell stage and there are sufficient stores of maternal mRNAs to allow the embryo to reach these stages of development.

To block protein synthesis, we treated embryos at various stages with puromycin (25 to 125 µg/ml) [30]. Comb formation was only observed in embryos that were treated after 6 hpf (at 25 µg/ml puromycin), suggesting that the

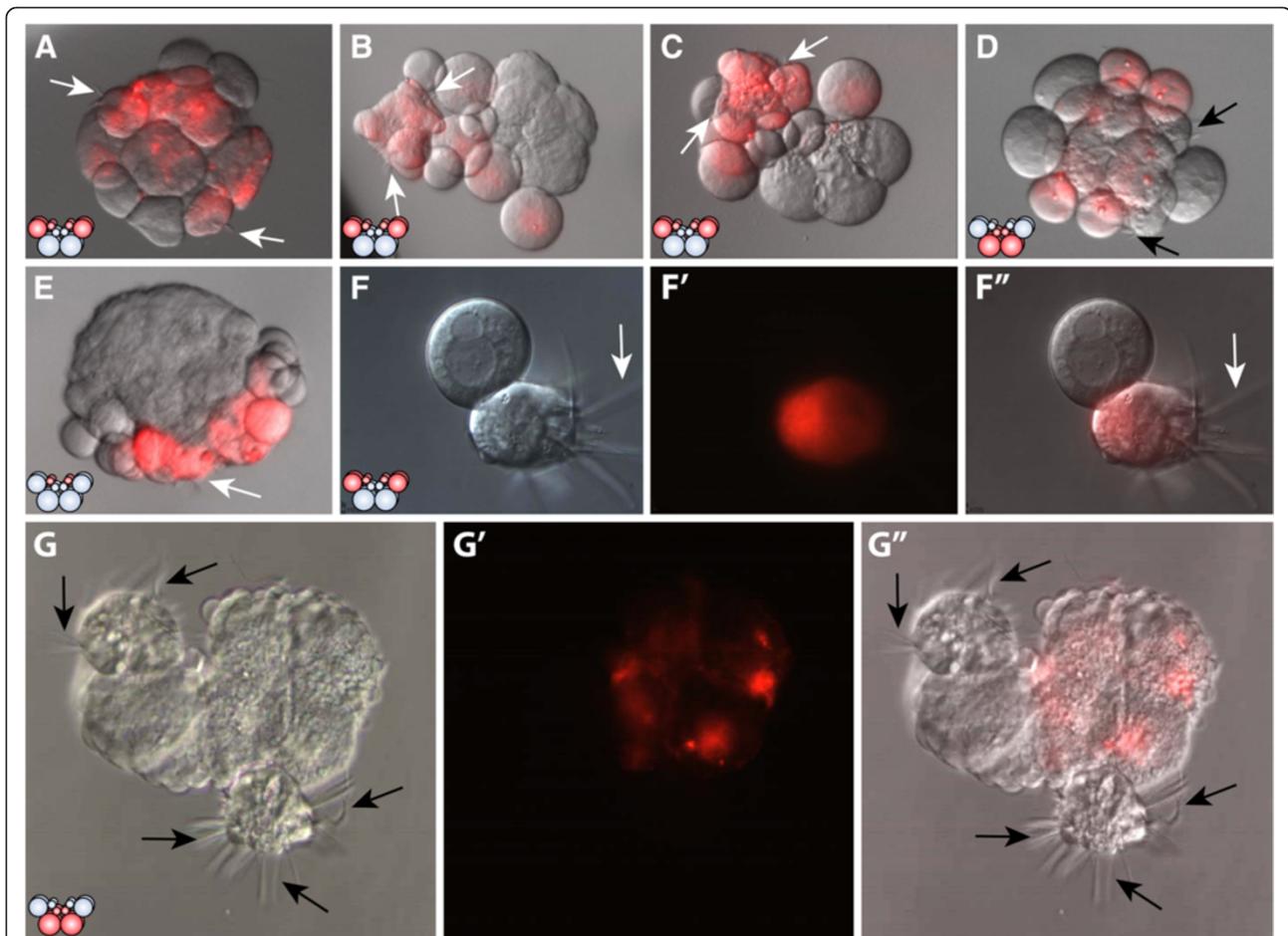


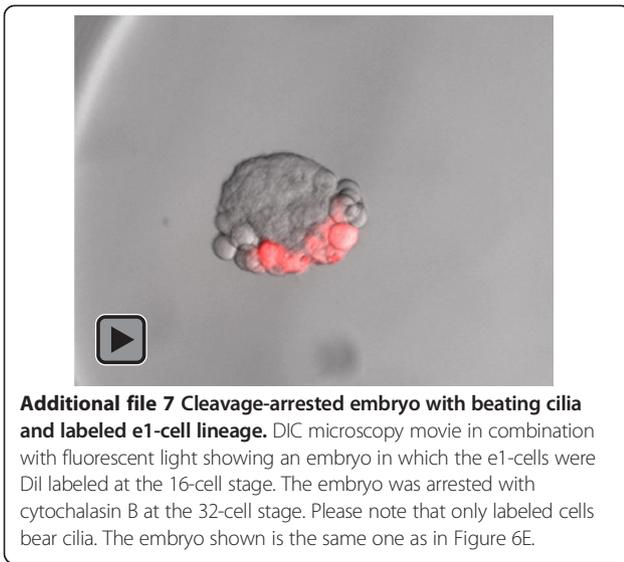
Figure 6 Lineage tracing experiments in cleavage-arrested embryos show that e1 micromeres give rise to the comb-cells. DIC images of cleavage-arrested embryos with DiI-labeling shown in red. White arrows point to labeled comb plates and black arrows point to unlabeled comb plates. The diagrams in the lower left corners show the stage when the embryo was inhibited with cytochalasin B and the appropriate DiI-labeled cells. **(A, B, C)** All four E macromeres were labeled at the 8-cell stage and the embryos were arrested at the 16-cell stage. Only labeled cells have combs. **(D)** All four M macromeres were labeled at the 8-cell stage and the embryos were arrested at the 16-cell stage. Only unlabeled cells form combs. **(E)** All four e1 micromeres were labeled and the embryos were arrested at the 16-cell stage. All comb cells are labeled. **(F, F'')** Part of an embryo that was arrested at the 16-cell stage. All E macromeres are labeled. **(F)** is a DIC image. **(F')** is the red channel and **(F'')** shows both images merged. The comb cell is clearly labeled. **(G, G', G'')** The embryo was arrested at the 16-cell stage. All four M macromeres are labeled. **(G)** is the DIC image and **(G')** the red channel. **(G'')** shows both images merged. All comb cells are unlabeled.

key determinant is not solely that a protein is required for comb formation, but that proteins must be translated between 5 and 6 hpf (Table 2).

The differentiation of combs is regulated by the number of nuclear divisions

Cells producing cteno-like cilia and photocytes are formed at the same time in control embryos and in embryos treated with cytochalasin B at the 8-cell stage or later, suggesting that the embryo has a molecular timing mechanism that operates in the absence of cytokinesis. Because nuclear divisions appear to proceed normally in embryos treated with cytochalasin B (Figure 4), the number of nuclear divisions may be an important component

of the timing mechanism. To test whether the timing of the formation of the comb-plate-like cilia is dependent on the number of nuclear divisions, we treated embryos with the DNA synthesis inhibitor aphidicolin (2 µg/ml) [31] at various cleavage stages up through 5 hpf (Table 2). When treated during early cleavage, up to the 32-cell stage, we did not observe the formation of any comb-plate-like cilia. When treated at the 60-cell stage, 4 hpf, and 5 hpf, we did observe comb formation, although it was delayed by approximately 2 to 3 h (12 hpf as opposed to 9 hpf). Embryos that were treated with aphidicolin at 5 hpf for 2 h and then removed from the drug, formed comb-plate-like cilia with a delay corresponding approximately to the length of the drug treatment. These results are consistent



Additional file 7 Cleavage-arrested embryo with beating cilia and labeled e1-cell lineage. DIC microscopy movie in combination with fluorescent light showing an embryo in which the e1-cells were Dil labeled at the 16-cell stage. The embryo was arrested with cytochalasin B at the 32-cell stage. Please note that only labeled cells bear cilia. The embryo shown is the same one as in Figure 6E.

with the fact that the aphidicolin treatment is reversible, that DNA synthesis is able to resume following the rescue (for example, the drug is not lethal) and that the correct total lineage-specific DNA content is important for the timing of the formation of comb-plate-like cilia. We did not see any comb formation in embryos that were treated with both cytochalasin B and aphidicolin, even after 6 hpf. The double treatment with cytochalasin B and aphidicolin is not reversible and embryos that were treated for 2, 3 or 4 hpf did not recover and did not form comb cells. These results showed that the combined effects of blocking DNA synthesis and cytokinesis are more detrimental than

either drug alone and indicate that a critical number of nuclear divisions is essential.

Discussion

We used cytochalasin B to inhibit cytokinesis in early ctenophore embryos to examine how developmental potential is partitioned into two distinct cell lineages: those giving rise to comb plates and those forming the photocytes. When cytokinesis is arrested in early *Mnemiopsis leidyi* embryos, treated embryos do not undergo apoptosis, but rather nuclear divisions continue and give rise to multinucleated cells. Cytochalasin B was used to study the segregation of developmental potential in *Mnemiopsis* by Freeman [18]. Freeman [18] inhibited certain divisions including the first, second or third cleavage or the second and third cleavage by short treatments with cytochalasin B. When only the first cleavage was inhibited by a brief exposure, most embryos developed like the control embryos but were delayed by one cleavage cycle [18]. However, when the second, third or second and third cleavages were reversibly blocked, the inhibited cleavages were skipped and the embryos continued the cleavage program with a reduced number of cells [18].

Surprisingly, as we demonstrate here, if cytokinesis is arrested at the 8-cell stage or later and is permitted to remain arrested for the following hours of development, comb-plate-like cilia and the photoprotein-mediated bioluminescence characteristic of photocytes form at the correct time in development compared with control embryos. Lineage-tracing experiments reveal that comb-plate-like cells are derived from E cells at the 8-cell stage and from the e₁ micromeres at the 16-cell stage. These are the

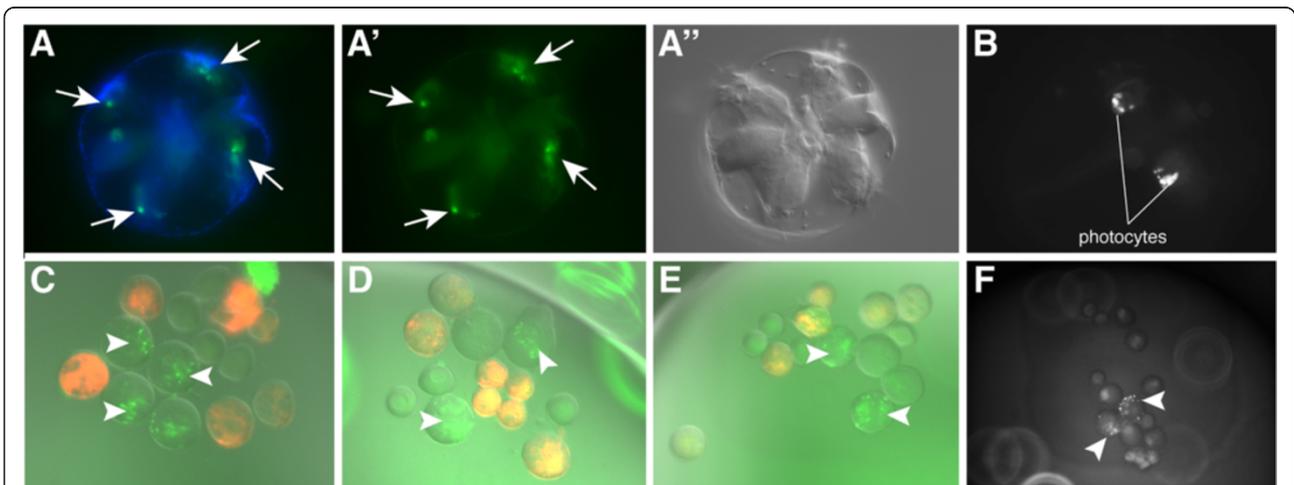
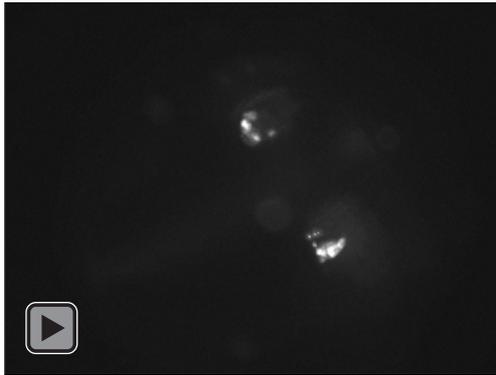
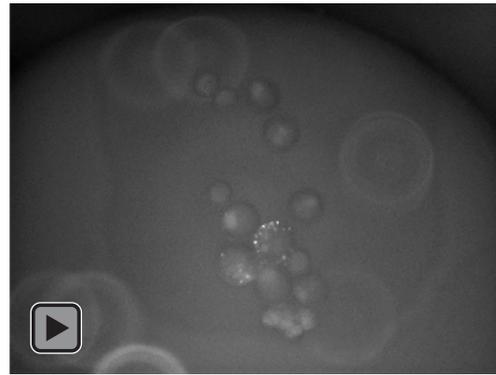


Figure 7 Photocytes in control embryos and cleavage-arrested embryos. (A, A', A'') In normal embryos, photocytes (white arrows) are visible at 9 to 10 hpf in the endoderm beneath the comb rows. In this live animal, viewed from the aboral pole, (A) shows nuclei in blue (Hoechst 33342 staining) overlaid with photoprotein fluorescence (green). (A') Photoprotein fluorescence as viewed under the GFP channel. (A'') DIC image of the same embryo shown in (A) and (A'). (B) Photoprotein fluorescence of a control animal. (C, D, E) E macromeres were labeled with Dil at the 8-cell stage and embryos were cleavage arrested at the 16-cell stage. Photoprotein is visible as green dots (arrowheads), in cells that are not labeled. (F) Photoprotein in an embryo that was cleavage arrested at the 16-cell stage. hpf, hours post fertilization. Additional files 8 and 9 show videos of the light-emitting photocytes of the specimens shown in figures B and F.



Additional file 8 Bioluminescence in photocytes of a control embryos. Light microscopy movie showing light emission from the photocytes that was stimulated by a brief illumination with fluorescent light. The photocytes are well visible on either side of the embryo. The animal shown is the same one as in Figure 7B.



Additional file 9 Bioluminescence in photocytes of a cleavage arrested embryos. Light microscopy movie showing light emission from the photocytes that was stimulated by a brief illumination with fluorescent light. Those cells that contain the bioluminescent protein are photocytes. The embryo shown is the same one as in Figure 7C.

same cell lineages that autonomously generate comb-plate-like cilia when isolated from the rest of the embryo [1]. Previous intracellular fate-mapping experiments showed that m_1 micromere descendants also have the capacity to make comb-plate-like cilia [4] but M lineage descendants were not found to give rise to comb-plate-like cilia in cleavage-

arrested embryos. Likewise, the bioluminescence characteristic of M-lineage-derived photocytes was only seen in M-lineage-derived cells in cleavage-arrested embryos. Earlier experiments showed that the m_1 micromeres require inductive signals first from the e_{11} or e_{12} micromeres and later from endomesodermal cells derived from E and M

Table 2 Overview of the aphidicolin, actinomycin D and puromycin treatment experiments on *Mnemiopsis leidyi* embryos

Developmental stage	Treatment	Observation	Conclusion
2-cell stage	Arrest of DNA synthesis for 2 h with aphidicolin, 2 $\mu\text{g}/\text{ml}$	No combs formed (0/30)	DNA synthesis is essential for comb formation.
16-cell stage			
32-cell stage			
60-cell stage		27/30 (90%) formed combs, comb formation was delayed by 2 h	
4 hpf		29/30 (97%) formed combs, comb formation was delayed by 2 h	
5 hpf	29/30 (97%) formed combs, comb formation was delayed by 2 h		
60-cell stage and 4 hpf	Arrest of DNA synthesis and cytokinesis with aphidicolin, 2 $\mu\text{g}/\text{ml}$, plus cytochalasin B, 1 to 2 $\mu\text{g}/\text{ml}$	Immediately arrested, single nuclei observed, neither combs nor photoprotein were formed in any embryo (0/30)	DNA synthesis is essential for comb formation even in cleavage-arrested embryos.
1- to 2-cell stage	Blocking of transcription with actinomycin D, 10 $\mu\text{g}/\text{ml}$	0/30 gastrulated, development arrests at 32- to 60-cell stage	Zygotic transcription begins around the 32- to 60-cell stage and the specification of the combs requires more than only maternal RNAs.
4- to 8-cell stage		0/30 gastrulated, development arrests at 32- to 60-cell stage	
16- to 32-cell stage		10/30 (33%) gastrulate, 20/30 (67%) arrest at 32- to 60-cell stage	
60-cell stage		28/30 (93%) gastrulate but no pharynx is formed, 2/30 (7%) do not gastrulate	
5 hpf (post-gastrulation)		No pharynx is formed	
2-cell stage	Arrest of translation with puromycin, 125 $\mu\text{g}/\text{ml}$, 25 $\mu\text{g}/\text{ml}$	0/30 formed combs	Comb protein is translated prior to 6 hpf, comb formation requires protein that is translated in the embryo and not maternally loaded.
16-cell stage		0/30 formed combs	
60-cell stage		0/30 formed combs	
6 hpf (post-gastrulation)		32/30 (97%) formed combs	

macromeres before they form combs [14,20]. Our results support the finding that m_1 micromeres require additional inductive signals from endomesodermal cells for comb formation, since no combs were formed by m_1 micromeres in cleavage-arrested embryos. This supports the finding that the presence of e_1 micromeres is not sufficient to induce comb formation in m_1 micromeres [14,20].

Interestingly, neither comb-plate-like cilia nor bioluminescence was detected in cleavage-arrested zygotes or 2- or 4-cell stage embryos, suggesting that the segregation of distinct E and M lineages is crucial for the expression of the lineage-specific differentiation products. These results suggest that factors that are required for comb-plate and photocyte determination are already present and localized at the 8- and 16-cell stages and may have mutually exclusive activity.

The segregation of developmental determinants

One of the fundamental processes of development is the localization of factors that are required to establish cell polarity, break symmetry and drive the asymmetric specification of cell fates. Experimental embryologists have provided evidence for the segregation of factors that determine specific cell fates in a diverse variety of different organisms (examples are summarized in [32-34]). In some embryos, segregated factors are asymmetrically localized maternally to the cell cortex such as the developmental potential required for gastrulation in echinoderm embryos [35-39] and for the establishment of dorso-ventral polarity in many embryos that undergo spiral cleavage [40-43], or the maternal gradients in insect eggs [44-46]. Other cytoplasmic factors are segregated actively at the time of fertilization as in ascidians [47-49], amphibians [50,51] and soil nematodes [52,53].

In ctenophore embryos, all evidence suggests that the asymmetrical localization of developmental potential is actively segregated by the cleavage process itself. Each of the early cell divisions leads to a definite asymmetric cell fate. The site of first cleavage gives rise to the oral-aboral axis [10]. Interestingly, a similar correlation between the site of the first cleavage and the site of gastrulation, and thus the formation of the oral-aboral axis, has been observed in the hydrozoan *Clytia hemisphaerica* (previously called *Phialidium gregarium*) [54]. Freeman [54] showed that the site of the first cleavage can be altered from the side of the polar body formation experimentally in ctenophores as well as in hydrozoans [10,54]. In contrast, the eggs of several sea urchin species are already polarized along the animal-vegetal axis at the time of fertilization (summarized in [55]). Starfish eggs and even ascidian eggs show a similar polarization [56,57]. Based on currently available data, it could be assumed that an irrevocable polarization of the unfertilized egg was an evolutionary novelty of bilaterians whereas in basally

branching taxa, such as ctenophores and cnidarians, the egg is polarized by the position of the female pronucleus, which determines the site of the first cleavage, after fertilization [17]. However, a broader taxon sampling is desired to further support this hypothesis.

In ctenophores, the second division gives rise to the anal axis [4,15,19]. The third division separates the E and M lineages [4,18], and the fourth division segregates fates between micromere lineages (for example, comb rows) and macromere lineages (for example, photocyte cells) [4,18]. Consequently, when cytokinesis is inhibited, the factors remain associated with the proper lineages that would have generated the descendants during normal development. Interestingly, photoprotein formation and comb-plate formation never appeared to occur in the same cell. It is possible that the developmental determinants, which are required for comb-plate and photoprotein differentiation, inhibit each other and only after these factors have been spatially segregated can they be activated.

Although we have little information on the cell biological nature of the segregation process in ctenophores, centrosomes have been shown to be a causal factor in the segregation of factors that specify cell fates in other systems. In *Caenorhabditis elegans*, the sperm-derived centrosome, a complex of several proteins that acts as the microtubule-organizing center, breaks the symmetry of the oocyte and sets off a series of events that relies on maternally deposited proteins and eventually leads to the establishment of the anterior-posterior, dorsal-ventral and left-right body axes [58,59]. A surprisingly large number of mRNAs are localized to specific cell lineages in the snail *Ilyanassa* [60]. These mRNAs are associated with the centrosomes, and as the cells divide the mRNAs are subsequently distributed in an asymmetric manner to specific daughter cells [61]. Experimental evidence shows that these factors play an active role in the unique development of these cells [60,61]. It would be interesting to investigate whether a centrosome-dependent mechanism is also involved in the segregation of developmental factors in *Mnemiopsis*, where distinct developmental fates are decided at each of the early divisions.

What is the nature of segregated developmental determinants?

Ctenophore development is rapid, with cleavages occurring every 15 to 20 min. The asymmetrical localization of proteins or mRNAs into distinct lineages during each division could be a simple way to distribute components quickly so as to distinguish different cell fates. Freeman [18] showed that the developmental determinants that are required for comb formation begin to be localized to the aboral region already during the 2-cell stage. Although crude and not gene specific, our experiments that inhibit transcription and translation suggest that the distributed determinants are not simply represented by the full set of

comb-plate protein or comb-plate mRNA. Instead subsequent transcription and translation are required for comb formation. The nature of the components that are asymmetrically localized during the early cleavage stages in *Mnemiopsis* remains unknown.

In other systems, RNAs, proteins, protein complexes or a mix of these are the developmental determinants that are distributed unevenly during development and which break the symmetries. A well-known example is the localization of the protein Dishevelled – a key player of the Wnt/ β -catenin pathway – to one side of an unfertilized egg. This is one of the first symmetry-breaking events in many bilaterians [17,62,63]. Other well-known examples of proteins with asymmetric localizations are Miranda, Prospero and Staufen during *Drosophila* neuroblast division [64] and the Par proteins during *C. elegans* development [65].

Besides the localization of proteins or protein complexes, RNAs are often asymmetrically stored and thus establish cell polarity and determine the different daughter cell fates. One of the most famous examples of RNA localization is the localization of the *Oskar* mRNA, which is one of the key components used to establish polarity in the *Drosophila* oocyte during *Drosophila* oogenesis [66-69]. Other examples of RNA localization are *ASH1* in budding yeast, *bicoid* in *Drosophila* embryos, *Vg1* in *Xenopus* oocytes and *CamKIIa* in distal dendrites in mammalian neurons (summarized in [70]).

Future experiments will be needed to reveal the nature of the developmental determinants that are segregated during early ctenophore development.

Temporal regulation of comb-plate formation: counting cell divisions

The regulation of growth and differentiation are key to developmental processes. Moreover, the timing of developmental events is a particularly important aspect of development. We showed that the differentiation of comb cells in *Mnemiopsis* embryos, which are cleavage arrested at the 8-cell stage or later, subsequently occurs at the same time as in untreated control embryos. Furthermore, our results show that comb-plate formation is blocked if DNA replication is inhibited by aphidicolin in addition to the inhibition of cytokinesis by cytochalasin B. This indicates that the number of nuclear division cycles or the amount of DNA (the nuclear-to-cytoplasmic ratio) appear to be important factors for comb-cell differentiation and not simply the amount of time that has elapsed since the third cleavage. As previously shown, after the second and/or third cleavage is blocked, *Mnemiopsis* embryos skip the respective cleavages and continue in a timely manner with their subsequent cleavage program, suggesting that a timing system determines the orientation of each division [18]. Once this process is activated, the cleavage plane is

determined by this timing mechanism and cleavages do not follow a set order with respect to each prior cleavage division [18]. Since the inhibition of the first cleavage only causes a delay of the program, which otherwise occurs normally, Freeman [18] concluded that the timing mechanism is initiated with the completion of the first cleavage.

Embryos that undergo exact numbers of cell divisions are known from a broad range of developmental systems; however, little is understood about the underlying mechanisms [71-75]. A stunning example is the highly stereotypic cell division pattern of the ventral germ band in malacostracan crustaceans. In *Cherax destructor*, for example, all ectoteloblasts undergo exactly 15 rounds of asymmetric cell division and give birth to ectodermal blast cells, which undergo two distinct divisions each [74].

Fundamental experiments by Whittaker showed that ascidian embryos developed muscle-specific acetylcholinesterase and brain pigment cell tyrosinase in specific blastomeres even in cleavage-arrested embryos and concluded that specific positional information is differentially segregated during early development [76]. Since then several studies have employed cytochalasin B to investigate the cell fate specification and cell lineage of ascidian embryos (for example, [77-80]). Satoh and Ikegami [75] performed a series of experiments combining cytochalasin B and aphidicolin in ascidian embryos, showing that future muscle cells must undergo eight cell-division cycles before they start expressing the muscle cell lineage marker acetylcholinesterase. Nevertheless, the question of how cells 'count' the number of cell cycles remains unanswered.

A number of counting mechanisms have been suggested for this phenomenon. Based on the observation that genomic DNA is highly methylated in zygotes and gradually demethylated during development, which tends to de-repress transcriptional activity in mammals [81], Kataoka et al. [73] suggested that changes in DNA methylation could be involved in keeping count of the number cell cycles in ascidians. However, their own results do not support this hypothesis as they were unable to detect any changes in DNA methylation [73]. Another possible explanation is that a specific ratio of DNA to cytoplasm is required. This ratio is crucial during early *Xenopus* development, where zygotic transcription is initiated after 11 to 12 rounds of cell division [82]. The required ratio of DNA-to-cytoplasmic volume can also be acquired when cytokinesis is blocked with cytochalasin B but DNA replication is allowed to continue further [82]. In contrast, in *C. elegans*, the timing of gut marker gene expression only depends on an early period of DNA synthesis until the 8-cell stage when the gut is clonally established [83]. Subsequent rounds of cell divisions, which usually occur after the 8-cell stage, can be inhibited without preventing the expression of the marker gene at a later stage, thus this is independent of the DNA-to-cytoplasmic ratio or the number of DNA synthesis rounds [83].

Future studies are required to unravel the mechanism that regulates the timing of cell differentiation in ctenophores, whether this is the ratio of DNA to cytoplasm, DNA methylation, protein degradation, telomeric alterations, a combination of different factors or an entirely different mechanism that is currently unknown.

The onset of zygotic transcription

Early developmental events are regulated by gene products, which may be provided maternally [84]. The activation of the zygotic genome marks the maternal-to-zygotic transition (MZT), which was first described for *Xenopus* [82] and has since been studied in many model organisms [84]. It is estimated that around 40% (in mice) to 75% (in sea urchins) of all protein coding genes are represented as maternal mRNAs and are degraded throughout early development [85,86].

Other than bilaterians, the onset of zygotic transcription has only been described in cnidarians. In *Clytia hemisphaerica*, zygotic transcription starts at the late

blastula to early gastrula stage [63,87]. The sea anemone *Nematostella vectensis* exhibits an upregulation of gene expression around 10 hpf [88], at a time in development when cell division is asynchronous [89].

We used actinomycin D to inhibit transcription and we showed that treatment prior to the 60-cell stage did not visibly affect development until the animal reached the 60-cell stage, suggesting that maternally loaded transcripts are sufficient to support the early cleavage program up to this stage but not subsequent development or differentiation. These results suggest that the onset of zygotic transcription starts at about the 60-cell stage, just prior to gastrulation. The rapid early development of ctenophores also suggests that there may be little time for the transcription of new messages until the cell cycle began to slow down. Future experiments using labeled nucleotides will help to determine the precise onset of zygotic transcription in *Mnemiopsis*.

The transition from maternal to zygotic transcription is most likely an ancient feature in animal evolution and

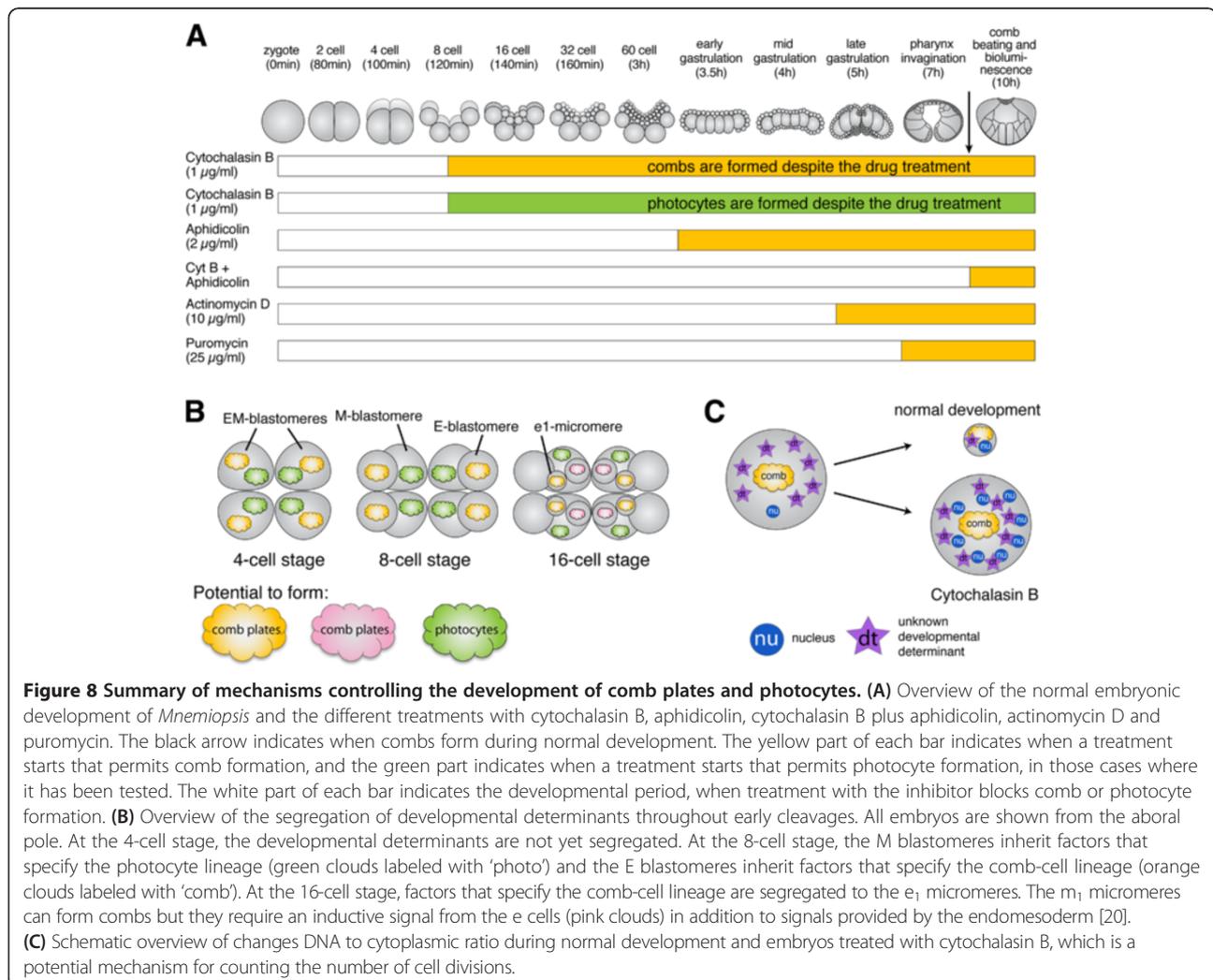


Figure 8 Summary of mechanisms controlling the development of comb plates and photocytes. (A) Overview of the normal embryonic development of *Mnemiopsis* and the different treatments with cytochalasin B, aphidicolin, cytochalasin B plus aphidicolin, actinomycin D and puromycin. The black arrow indicates when combs form during normal development. The yellow part of each bar indicates when a treatment starts that permits comb formation, and the green part indicates when a treatment starts that permits photocyte formation, in those cases where it has been tested. The white part of each bar indicates the developmental period, when treatment with the inhibitor blocks comb or photocyte formation. **(B)** Overview of the segregation of developmental determinants throughout early cleavages. All embryos are shown from the aboral pole. At the 4-cell stage, the developmental determinants are not yet segregated. At the 8-cell stage, the M blastomeres inherit factors that specify the photocyte lineage (green clouds labeled with 'photo') and the E blastomeres inherit factors that specify the comb-cell lineage (orange clouds labeled with 'comb'). At the 16-cell stage, factors that specify the comb-cell lineage are segregated to the e₁ micromeres. The m₁ micromeres can form combs but they require an inductive signal from the e cells (pink clouds) in addition to signals provided by the endomesoderm [20]. **(C)** Schematic overview of changes DNA to cytoplasmic ratio during normal development and embryos treated with cytochalasin B, which is a potential mechanism for counting the number of cell divisions.

might be tightly linked with the evolution of multicellularity and sexual reproduction. If ctenophores are the sister group of all metazoans, as suggested by recent studies [90,91], one could conclude that the inclusion of the MZT is part of the metazoan ground pattern. However, so far data about the MZT in sponges are not available and the phylogenetic position of ctenophores is still very much under debate [92]. There are currently several contradictory hypotheses, including: (a) ctenophores and cnidarians are sister groups [93-95] and (b) ctenophores are the sister group to all bilaterians [96]. Recent developmental studies support the idea that ctenophores lack several characters that are shared between cnidarians and bilaterians [13,97-101] and thus they support the idea that (c) ctenophores are the sister group to all remaining eumetazoans [102] or even, as mentioned above, that (d) ctenophores are the sister group to all metazoans [90,91].

A phylogenetic comparison of maternally loaded proteins and the molecular regulation of the maternal-to-zygotic transition in bilaterian and non-bilaterian taxa might provide further insight into the evolution of sexual reproduction and life history.

Conclusions

As in other animals with a mosaic development, embryos of the comb jelly *Mnemiopsis leidyi* show differentiation of selected cell fates even when cytokinesis is arrested using cytochalasin B during the course of early embryogenesis [48,77,78,103,104]. An overview is given in Figure 8. We demonstrated that embryos arrested prior to the 8-cell stage do not show visible signs of cell fate differentiation from either the E or M lineages. However, if cytokinesis is blocked at the 8-cell stage or later, comb-plate-like cilia, which are derived autonomously from the E lineage, and the bioluminescence characteristic of photocytes form at the correct time compared with control embryos. Treatment with the DNA synthesis inhibitor aphidicolin reveals that the number of nuclear divisions is essential for the proper timing of the differentiation of the comb-plate-like cilia. Lineage tracing experiments show that cells producing comb-plate-like cilia are formed autonomously from the E lineage, particularly the e_1 micromeres, and not any of the M or m_1 lineages. Likewise, bioluminescence is only observed in the M lineage and specifically in M macromeres. In addition, we showed that developmental determinants that are required to generate motile comb-plate-like cilia and photoprotein expression are already present and localized at the 8-cell stage and further localize to the appropriate cells in the subsequent cell divisions. Further studies are required to unravel the nature of the developmental determinants and which mechanisms are used to segregate them. The timed inhibition of transcription and translation with actinomycin D and puromycin, respectively, shows that both transcription and translation are

required for comb cells to form, suggesting that not all the necessary factors are maternally deposited. While these inhibitor studies are not definitive gene-specific approaches, they do provide insight into the potential nature of materials that are differentially segregated into different embryonic lineages. Finally, we presented evidence that suggests that zygotic transcription in *Mnemiopsis* begins around the 60-cell stage, just before the onset of gastrulation.

Abbreviations

hpf: hours post fertilization; MZT: Maternal-to-zygotic transition; PBS: Phosphate-buffered saline; PBT: PBS plus 0.2% Triton; DIC: Differential interference contrast; AO: Apical organ.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

AHLF, KP, JQH and MQM conceived the study and analyzed the data. AHLF and KP conducted and analyzed the drug treatments and produced the figures. KP analyzed the immunohistochemistry, produced the confocal micrographs and recorded the time-lapse images. JQH and MQM performed the microinjection. KP, JQH and MQM imaged the injected embryos. AHLF wrote the first draft of the manuscript. All other authors helped to complete and edit the manuscript. All authors read and approved the final manuscript.

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