Yolk syncytial layer formation is a failure of cytokinesis mediated by Rock1 function in the early zebrafish embryo

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Summary
The yolk syncytial layer (YSL) performs multiple critical roles during zebrafish development. However, little is known about the cellular and molecular mechanisms that underlie the formation of this important extraembryonic structure. Here, we demonstrate by timelapse confocal microscopy of a transgenic line expressing membrane-targeted GFP that the YSL forms as a result of the absence of cytokinesis between daughter nuclei at the tenth mitotic division and the regression of pre-existing marginal cell membranes, thus converting the former margin of the blastoderm into a syncytium. We show that disruption of components of the cytoskeleton induces the formation of an expanded YSL, and identify Rock1 as the regulator of cytoskeletal dynamics that lead to YSL formation.

Our results suggest that the YSL forms as a result of controlled cytokinesis failure in the marginal blastomeres, and Rock1 function is necessary for this process to occur. Uncovering the cellular and molecular mechanisms underlying zebrafish YSL formation offers significant insight into syncytial development in other tissues as well as in pathological conditions.

Key words: Yolk syncytial layer, Cytoskeleton, Cytokinesis, Rock1

Introduction
The yolk syncytial layer (YSL) is a multinucleate layer of non-yolky cytoplasm immediately beneath the cellular blastoderm. It is the first lineage-restricted extraembryonic structure to form in the zebrafish embryo (Kimmel et al., 1995). Although it does not contribute cells or nuclei to the embryo, it is nevertheless important for embryonic development (Carvalho and Heisenberg, 2010). It is crucial for induction of the embryonic organizer (Mizuno et al., 1999), early patterning of mesoderm and endoderm (Mizuno et al., 1996; Rodaway et al., 1999), epiboly (Trinkaus, 1993; Solnica-Krezel and Driever, 1994) and cardiac morphogenesis (Sakaguchi et al., 2006). Additionally, the YSL plays a nutritive role during embryonic and larval stages (Walzer and Schönberger, 1979a; Walzer and Schönberger, 1979b).

Our understanding of YSL formation is largely limited to descriptive studies. The YSL forms in a similar manner in a number of teleosts: marginal blastomeres undergo a morphogenetic “collapse” upon the yolk to form the YSL (Kimmel and Law, 1985; Trinkaus, 1993). In the zebrafish embryo, this occurs at the ninth or tenth cell division (Kimmel and Law, 1985). The final step of cell division is cytokinesis – the division of the cytoplasm into two daughter cells following karyokinesis. Cytokinesis is a highly regulated multistep process (reviewed by Eggert et al., 2006), which involves continuous remodelling of the cytoskeleton and plasma membranes to effect the necessary changes in cytoarchitecture. Small GTPases of the Rho family, which are molecular switches that control many cytoskeleton-dependent cell functions, have been implicated in this process (Etienne-Manneville and Hall, 2002). Rho kinases (Rocks) are downstream effectors of Rho signalling – activated GTP-bound Rho binds to Rock at its Rho-binding domain, thereby relieving the autoinhibitory loop formed by interaction between the N- and C-termini of Rock (Riento and Ridley, 2003).

Although the molecular mechanism underlying cytokinesis has been a matter of intense investigation, the molecular mechanism underlying YSL formation is only just beginning to be elucidated (Takesono et al., 2012). This report demonstrated that solute carrier family 3 member 2 (Slc3a2) inhibits the RhoA/Rock pathway via phosphorylation of c-Src to modulate YSL microtubule dynamics and YSL formation. Here, we show using timelapse confocal microscopy of a transgenic zebrafish line expressing membrane-targeted GFP that the YSL forms as a result of the absence of cytokinesis between marginal daughter nuclei at the tenth mitotic division and the regression of pre-existing marginal cell membranes. Disruption of components of the cytoskeleton by various means, including nocodazole, cytochalasin D and attenuation of Rock function by small molecule inhibitors led to YSL expansion. To define which Rho kinase is responsible for this effect we employed the dominant-negative loss-of-function (LOF) strategy and demonstrated that...
LOF of Rock1, but not Rock2, led to YSL expansion. Taken together, these results indicate that the YSL forms as a result of regulated cytokinesis failure of the marginal blastomeress, and attenuation of Rock1 activity is necessary for this process to occur.

Results and Discussion
Microtubule and membrane dynamics in the early zebrafish embryo
The distribution and organization of microtubules in the yolk cell of the early zebrafish embryo have been analysed by immunolabelling (Strähle and Jesuthasan, 1993; Solnica-Krezel and Driever, 1994). In this study, late cleavage and blastula stage embryos were examined by whole-mount immunofluorescence to visualize changes in the organization of yolk cortical microtubules for the period leading to and spanning YSL formation (Fig. 1). From the 32- to 512-cell stage, the microtubule arrays of the marginal blastomeres extended into the yolk cortex along the animal-vegetal axis (Fig. 1Aii–Dii). Cell borders were evident between marginal blastomeress from the 32- to the 256-cell stage (Fig. 1Ai–C, arrowheads). At the 256-cell stage, the yolk cortical microtubules emanated mainly from the central region of the base of the marginal blastomeress (Fig. 1Ciii, inset). At the 512-cell stage (2.75 hpf), just prior to YSL formation, the marginal blastomeress flattened towards the yolk cortex (note difference in curvature of marginal blastomeress between Fig. 1Dii and Fig. 1Cii). Cell boundaries were detected between some marginal blastomeress (Fig. 1Di, arrowheads), but not others (Fig. 1Di, arrow).

Midblastula transition (MBT) begins at this stage (Kane and Kimmel, 1993). MBT is characterized by initiation of zygotic transcription, cell cycle lengthening and loss of cell cycle synchrony (note some cells in interphase [Fig. 1Diii inset, arrow] and others in mitosis [Fig. 1Diii inset, arrowhead]). The YSL is formed at the end of the 512-cell stage, after the tenth mitosis (Kimmel and Law, 1985). At the 1k-cell stage, just after YSL formation, most marginal blastomeress have lost their boundaries, suggesting that pre-existing cell membranes recede. However, remnants of a few boundaries could still be detected (Fig. 1Ei, arrowheads). The microtubules of the newly formed YSL radiate from the yolk syncytial nuclei (YSN), forming large interdigitating aster-like arrays (Fig. 1Eii, arrowheads) (Solnica-Krezel and Driever, 1994). The yolk cortical microtubules now emanate from the YSL (i.e., the former marginal blastomeress) instead of the current marginal blastomeress. By the oblong stage (3.7 hpf), the YSN have divided several times (Kimmel et al., 1995) and spread vegetally (Fig. 1Fi, arrowhead), as well as beneath the blastoderm to form the internal-YSL (I-YSL) (Korzh et al., 1989; Solnica-Krezel and Driever, 1994). The YSL microtubule network now consists of a dense mat of intercrossing microtubules adjacent to the blastoderm and large astral-like assemblies radiating from the vegetal-most YSN (Fig. 1Fiii, dashed circle). The yolk cytoplasmic layer (YCL) microtubule array extends from these astral-like arrays of the vegetal YSN.

To better understand the membrane dynamics during YSL formation, time-lapse confocal microscopy was employed to visualize events at the blastoderm margin using the mGFP (membrane-targeted GFP) transgenic line, which expresses GFP fused with the last 20 amino acids of c-Ha-Ras, a sequence that provides farnesylation and palmitoylation signals for plasma membrane targeting, driven by the medaka β-actin promoter (Cooper et al., 2005). At the 512-cell stage, cell membranes of the marginal blastomeress are still evident (Fig. 2A, white arrowhead; supplementary material Movie 1). During the next division cycle, no new cleavage furrows form at the margin (Fig. 2G–N, arrow). Pre-existing cell membranes of the marginal blastomeress regress over the next cycle (Fig. 2M–R; supplementary material Movie 1), and nuclei reappear after this mitosis sharing a common cytoplasm, thus forming the YSL. These events occur very quickly, giving the appearance of marginal blastomeress collapsing onto the yolk cell.

Disruption of the cytoskeleton induces YSL formation
Microtubules and microfilaments are dynamic components of the cell cytoskeleton. Both cytoskeletal elements are constantly being remodelled and redistributed within the cell. This dynamism is vital to the function of the cytoskeleton especially in the control
of cell division, cell shape and motility. To characterize the role of these elements in YSL formation, we tested the effects of the microtubule-depolymerizing drug, nocodazole, and the inhibitor of actin polymerization, cytochalasin D, on this process. Consistent with a previous study (Jesuthasan and Strähle, 1997), nocodazole-treated embryos developed an enlarged syncytium (Fig. 3Bi, arrow) with small asters (Fig. 3Biii, arrowheads) instead of the large asters characteristic of normal YSL (Fig. 3Aiii, broken circle). Note a pre-existing membrane at the margin regressing as the YSL develops (red arrowhead in M–R). Nuclei are unstained and appear as dark circles.

Cytochalasin D-treated embryos also developed an enlarged acellular region of cytoplasm (arrow in Fig. 3Ci), which contained dense disorganized microtubules without asters (Fig. 3Ci, Ci). Some regions in the YCL were devoid of microtubules (Fig. 3Ci, asterisks). Again, the enlarged syncytium expressed mtx1 (Fig. 3Civ) and induced ntl in the overlying blastoderm (Fig. 3Cv).

Y27632 is a specific inhibitor of the ROCK family of kinases. Its affinity for ROCKI and ROCKII as determined by Ki values was at least 20 to 30 times higher than those for two other Rho effector kinases, citron kinase and protein kinase PKN (Ishizaki et al., 2000). In order to determine whether it plays a role upstream of actin in this context, embryos at the 1-cell stage were injected with Y27632. Embryos injected with 2 pmol Y27632 developed enlarged syncytia similar to cytochalasin-treated embryos (Fig. 3Di, arrow) with dense disordered microtubules.
(Fig. 3Dii,iii). In contrast, the YCL microtubules in Y27632-injected embryos appeared sparse in comparison to control YCL (compare Fig. 3Diii with Fig. 3Aiii). The enlarged syncytium in Y27632-injected embryos expressed mtx1 (Fig. 3Div) and could induce ntl endogenously in the overlying blastoderm (Fig. 3Dv) as well as in animal caps (Fig. 3E) confirming the identity of this layer as the YSL. Together, the data suggest that disruption of cytoskeletal elements induces YSL formation, and the Rho-Rock signalling pathway is involved in this process.

Rock1, but not Rock2, is involved in YSL formation

Two ROCK isoforms have been identified in mammals – ROCKI/ROKI and ROCKII/ROKα (Riento and Ridley, 2003). In zebrafish, only rock2 paralogues have been cloned thus far: rock2a, which has been shown to be required for mediolateral cell elongation in convergence-extension movements during gastrulation (Marlow et al., 2002) and rock2b, which is necessary for establishing anteroposterior asymmetry of the Kupffer’s vesicle (Wang et al., 2011). Unlike rock2b, rock2a is expressed maternally; hence, it may have a role prior to gastrulation. To determine whether zebrafish Rock2a is involved in YSL formation, dominant-negative Rock2a (dn-Rock2a) was generated by deletion of the amino terminus containing the kinase domain (Marlow et al., 2002) (Fig. 4A). Overexpression of dn-rock2a resulted in dose-dependent enlargement of cells with single enlarged nuclei in each cell (Fig. 4Cii,iii, arrowheads), which indicates failure of both karyokinesis and cytokinesis. Since cytokinesis and karyokinesis are temporally coordinated such that furrow ingression closely follows the segregation of chromosomes in anaphase, failure of cytokinesis in dn-Rock2a expressing cells could be a secondary consequence of karyokinesis failure. The YSL, however, appeared to have formed normally suggesting that Rock2a is not responsible for regulating the cytoskeletal dynamics leading to YSL formation.

Hence, we proceeded to clone zebrafish rock1. The predicted protein shares 80–90% identity with Rock1 from other teleosts (Fig. 5A). Interestingly, amino acid sequence identity between human ROCKI and ROCKII is 65.9%, whereas that between zebrafish Rock1 and Rock2a and Rock2b is slightly less (61.8% and 54.5%, respectively), implying a faster rate of divergence between the zebrafish Rocks. Transcripts of rock1 are ubiquitously expressed during cleavage blastula and gastrula stages (Fig. 5Bi–iii). By 36hpf, expression is restricted to anterior structures, including the forebrain and hindbrain (Fig. 5Biv).

Similar to Y27632-injected embryos, dn-Rock1 expressing embryos developed an enlarged YSL with a disorganized microtubule network (Fig. 4Cii,iii). Time-lapse confocal microscopy of mGFP embryos expressing dn-Rock1 revealed that nuclear division proceeds normally in the absence of cell membrane formation in the affected cells (Fig. 6; supplementary material Movie 2) (n=7). This occurs predominantly in the marginal cells (Fig. 6A–C; supplementary material Movie 2, white arrows) whereas cells near the animal pole are not affected as much (Fig. 6H–M; supplementary material Movie 2, arrowheads). Cell membranes of non-marginal cells that eventually syncytialize appear to regress after several rounds of nuclear division (Fig. 6D–R; supplementary material Movie 2, red arrow), suggesting that Rock1 activity is also required for prevention of furrow regression in addition to cleavage furrow formation during cytokinesis in the early zebrafish embryo. This is consistent with observations in other systems. In the C. elegans hypomorphemic Rho-binding kinase mutant, let-502 (sb106), some cells fail to complete cytokinesis: furrows initiate and ingress slightly but later regress (Pieknay and Mains, 2002). In Drosophila, Rok has been shown to promote cleavage furrow ingression (Hickson et al., 2006); and in mammalian cells, preventing ROCK activation of myosin II by dissociating the septin scaffold that physically links the kinase to its substrate results in cleavage furrow regression (Joo et al., 2007).

Our data suggest a model of YSL formation involving specific attenuation of Rock1 function at the blastoderm margin leading to localized and controlled cytokinesis failure. This could occur in response to a maternally expressed signal localized only to the blastoderm margin. Since Rock is inactivated directly by Gem (Ward et al., 2002) and indirectly by Rho-GAP (Riento and Ridley, 2003).
2003), they are potential targets for this maternal signal. Another possible model of YSL formation was recently proposed by Takesono and colleagues, in which solute carrier family 3 member 2 (Slc3a2) inhibits RhoA/Rock signalling via c-Src activation, thereby regulating microtubule dynamics and YSL organization (Takesono et al., 2012). They reported that Slc3a2 knockdown caused YSL expansion, which was rescued by treatment with Y27632. The likely explanation for the apparent discrepancy between our results is that the level of Rock activity at the margin needs to be within an acceptable range for normal YSL formation – any deviation from this range in either direction results in YSL expansion. Our hypothesis that attenuation of Rock1 activity is necessary for YSL formation is consistent with the proposed Slc3a2-mediated signalling pathway, and implicates Rock1, but not Rock2, as the effector of RhoA signalling in this pathway.

Maternal-effect mutants that have defects in the cytokinetic machinery such as cellular island (cei), which affects Aurora-B kinase (Yabe et al., 2009), and cellular atoll (cea), which affects the centriolar protein Sas-6 (Yabe et al., 2007), also develop an expanded YSL (supplementary material Fig. S1), lending support to the hypothesis that YSL formation is a result of cytokinesis failure. Indeed, this mechanism of controlled cytokinesis failure or abortive cytokinesis has been employed in mammalian developmental programs such as cardiomycyte binucleation (Engel et al., 2006), hepatocyte tetraploidization (Margall-Ducos et al., 2007), and megakaryocyte polyploidization (Lordier et al., 2008). Understanding how cytokinesis is aborted in these physiological settings would expand our understanding of how cytokinesis fails in pathological states such as cancer.

Materials and Methods
Zebrafish husbandry
Wild-type zebrafish (AB strain), transgenic and mutant lines were maintained as described (Westerfield, 2000) in accordance with IACUC regulations (Biopolis IACUC application NO. 050096). All embryos were generated by pair-wise matings and staged according to Kimmel and colleagues (Kimmel et al., 1995).

Embryo stainings
Whole-mount in situ hybridization, immunohistochemistry and cryosectioning were performed as described (Korzh et al., 1993; Korzh et al., 1998). KMX-1 (Solnica-Krezel and Driever, 1994) (Chemicon) and secondary goat anti-mouse IgG-Alexa-488/Alexa-633 (Molecular Probes) were used to detect β-tubulin. Alexa-546-phalloidin (Molecular Probes) was used at 66 nM for fluorescent labelling of actin. DNA was labelled by incubating embryos in 0.3 μM DAPI (Sigma).

Drug treatments
Embryos in their chorions were incubated in 0.1 μg/ml nocodazole (10 mg/ml stock in DMSO) from the 16-cell stage to 4 hpf. For cytochalasin D, embryos in their chorions were incubated in 15 μg/ml (5 mg/ml stock in DMSO) for 35 minutes from the 2–4 cell stage, then washed and left to develop until stage 24 common embryo. Y27632 (Calbiochem) was prepared as a 30 mM stock in DMSO. All drugs were diluted to the appropriate concentrations in embryo medium prior to use. Controls were performed by treating sibling embryos in the same manner with the corresponding concentration of solvent in embryo medium.

Cloning of full-length rock1
RNA was isolated from 2 hpf embryos using RNA Purification Kit (QIAGEN). First-strand cDNA was synthesized by oligo(dT) priming using PowerScript reverse transcriptase (Clontech). To determine the 5′-end of rock1, we performed

Fig. 5. Cloning of rock1 and its expression pattern during development. (A) Multiple amino acid sequences were aligned using ClustalW algorithm. Calculation of sequence similarity and distance as well as phylogeny reconstruction were done using MegAlign from DNASTAR Lasergene v8.0.2. Full-length amino acid sequences were retrieved either from Ensembl database (Zv9, release 63), ENSORLP00000022079 (medaka Rock1), ENSGACP00000022893 (stickleback Rock1), ENSTNIP000000910 (tetraodon Rock1), FBpp0074061 (Drosophila Rock) or from NCBI database, NP_053597.1 (human ROCK1), NP_03097.1 (mouse ROCK1), NP_008481.1 (human ROCK2), NP_03098.1 (mouse ROCK2), AAK97854.1 (zebrafish Rock2a), CAI21196.1 (zebrafish Rock2b). (B) Early rock1 mRNA is of maternal origin. It is expressed ubiquitously at (Bi) 64-cell, (Bii) 512-cell, and (Biii) 6 hpf. (Biv) At 36 hpf, rock1 is detected in anterior structures, including forebrain and hindbrain.
Fig. 6. In vivo membrane dynamics at the blastoderm edge during YSL formation upon injection of dn-rock1. Selected frames from supplementary material Movie 2 showing the sequence of membrane failure events in a representative dn-Rock1 expressing embryo. Note the lack of membrane deposition between marginal nuclei (white arrows in A–C). Most cells near the animal pole undergo normal nuclear and cell division (arrowheads in H–M). Note the regression of cell membranes in a syncytializing non-marginal cell after multiple rounds of nuclear division (red arrow in D–R).

5'-RACE PCR (SMART RACE cDNA Amplification Kit from Clontech) with gene-specific primers (5'-ACA ACC TCC GCA GTG TAG AAG CGT G-3' and 5'-TGC TGA GCA GTC TCA TGG GTG AGA C-3'). Full-length cDNA of rock1 was amplified using Expand Long Template PCR System (Roche). Gene-specific primers (5'-GAA GGC GGA CAT GGT CGG GCA G-3' and 5'-TGG TGC GTC GTG CGA ACA GAA CAG CTG A-3') were designed based on sequence derived from 5'-RACE and Ensembl database (v8). The PCR product of approximately 5.2 kb was cloned into pGEM-T Easy vector (Promega). The sequence information related to defects in the contractile ring and Rho/Rock signaling.

RNA injections
5'-capped sense RNA was synthesized with the mMessage mMachine T7 kit (Ambion). RNA was injected into a single blastomere at the 2-cell stage. Doses injected ranged from 93–1800 pg of dn-rock2a and 200–400 pg of dn-rock1.

Generation of yolk cell-animal cap conjugates
The mesoderm induction assay was performed as described by Ober and Schulte-Mmerker (Ober and Schulte-Mmerker, 1999).

Microscopy
Observation of live embryos was performed using either SZX12 dissecting microscope (Olympus) or Fluoview FV1000 confocal system attached to an inverted microscope (Olympus IX81). Immunofluorescence-stained embryos were imaged by confocal microscopy (Fluoview FV1000 attached to upright Olympus BX61 microscope).

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Competing Interests
The authors declare that there are no competing interests.

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