Molecular characteristics of the edge cells responsible for expansion of the chick embryo on the vitelline membrane

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During early avian development, only a narrow band of cells (the edge cells, also called ‘margin of overgrowth’) at the rim of the embryo is responsible for blastoderm expansion by crawling over the vitelline membrane (VM) to cover the whole egg yolk in just 4 days (a process called epiboly). Surprisingly, this has not yet been studied in detail. Here we explore the edge cells of the chick embryo using in situ hybridization, immunohistochemistry and live imaging. Morphological and molecular properties reveal that the edge has a distinctive structure, being subdivided into sub-regions, including at least four distinct zones (which we name as leading, trailing, deep and stalk zones). This allows us to study reorganization of the edge region that accompanies reattachment of an explanted blastoderm to the VM. Immunohistochemistry uncovers distinct polarized cellular features resembling the process of collective cell migration described in other systems. Live imaging reveals dynamic lamellipodial and filopodial activity at the leading edge of the outermost cells. Our data provide evidence that edge cells are a distinct tissue. We propose that edge cells may be a useful model system for the study of wound healing and other closure events in epithelial cell sheets.

1. Introduction

During the first few days of development of the chick embryo, the extra-embryonic tissue (initially called the ‘area opaca’) at the periphery of the flat blastoderm gradually expands until, about 4 days’ incubation after egg laying, it covers the entire surface of the spherical yolk mass. This represents more than a 200-fold increase in surface area and a rate of expansion of 200–550 µm h\(^{-1}\) [1] (figure 1a). This expansion is driven by specialized cells located at the extreme edge of the embryo, which are the only cells that adhere to the inner (yolk-facing) surface of the vitelline membrane (VM). This narrow ring of cells has been called ‘margin of overgrowth’ [2] and spans approximately 40–60 µm [1,3–5]. Several requirements need to be met for proper expansion of the blastoderm and normal embryo development. First, the attachment of this edge is required for blastoderm expansion and their outgoing movement generates and maintains tension in the embryo [4–6], which rises to a peak at around 20–24 h’ incubation [1]. Second, tension in the acellular VM that constitutes the substrate for this expansion is also important for normal embryo development [4–6]. In addition, the orientation of both the embryo and the VM is important. For example, on an inverted VM, the embryo becomes abnormally strongly attached to the outer VM and does not expand, while an inverted blastoderm develops into a hollow vesicle probably by folding of the edge, allowing attachment of the dorsal side of the edge cells to the VM, on which they crawl to seal to form a sphere [5].

Cells at the edge of the embryo change their shape from initially cuboidal multiple layers to later a squamous and flattened monolayer [1,3]. The extreme
edge region has larger cells with larger nuclei, while the immediately proximal region has densely packed cells [3]. Perhaps paradoxically given the rapid expansion of the circumference at this time, the extreme edge cells do not appear to proliferate [3,7]. It has been shown that proximal cells close to the edge contribute to the edge region; the...
Table 1. List of marker genes for the edge region.

| gene       | NCBI gene ID | full name                          | marker of           |
|------------|--------------|------------------------------------|---------------------|
| RNH1       | 423111       | ribonuclease/angiogenin inhibitor 1 | leading/trailing/deep |
| SNAI2      | 432368       | snail family transcriptional repressor 2 | leading/trailing/deep |
| DACT2      | 421561       | dishevelled binding antagonist of beta catenin 2 | leading/trailing/deep |
| MSX1       | 396484       | msh homeobox 1                     | stalk               |
| TGM4       | 420706       | transglutaminase 4                 | leading/trailing/deep/stalk |
| GATA2      | 416018       | GATA binding protein 2             | trailing            |
| LHX1       | 396381       | LIM homeobox 1                     | trailing            |
| DKK1 (DKK4) | 374156     | dikkopf WNT signalling pathway inhibitor 4 | trailing/stalk      |

same study also investigated some molecular markers expressed in this region and described specific expression of the intermediate filament protein, vimentin [7].

The edge cells have unique cellular morphologies that are related to their adhesiveness and migration. At their distal face, broad and flattened cell-processes adhere to the inner VM [8,9]. Incubation of embryos in small peptide inhibitors of adhesion to fibronectin causes transient detachment of the embryo from the VM and arrest of expansion for a short time, suggesting that fibronectin is involved in the adhesion of the embryo to the VM [10]. Chemical inhibition of microtubules also affects blastoderm expansion [11,12]. Apart from this, we know very little about the mechanisms of chick blastoderm expansion.

In this study, we revisit chick embryo expansion and study its molecular characteristics. We reveal molecular markers expressed in different sub-domains of the edge region. We show that when an embryo is removed from its membrane and then replaced onto it, reattachment is accompanied by an increase in the number of cells with edge cell properties (including molecular markers), suggesting that reattachment involves reorganization of the tissue. We use immunohistochemistry to explore various cellular properties including cell polarity, proliferation and proteins involved in cell adhesion—these are strongly reminiscent of similar properties described for cells undergoing collective cell migration in other systems. Finally, high-resolution live imaging reveals dynamic morphological changes of the most distal, leading-edge cells.

2. Results and discussion

2.1. Markers for the edge cells

No specific markers for edge cells have yet been described in any avian species. To find candidate genes that are specifically expressed in the edge region, we first examined whole-mount in situ hybridization data from previous studies (including [13] and data for a large number of genes from our laboratory). This revealed eight genes that are specifically expressed or highly enriched at the rim of the embryo: RNH1, SNAI2, DACT2, MSX1, TGM4, GATA2, LHX1 and DKK1 (table 1). To investigate these in more detail, we performed in situ hybridization on embryos incubated for 24 h (approximately HH stage 5–7 embryos) [14]. In dorsal view, all the genes showed localized expression in the outermost 2–3 cells all around the periphery of the embryo (figure 1c–f and k–n); in the case of MSX1 and TGM4, expression was also seen in non-edge proximal cells (figure 1k,l). Histological sections of these embryos revealed distinctive expression patterns of the markers for cells located at slightly different positions within the edge region. This allows classification of four distinct regions, which we name leading (at the leading/free edge, and adjacent to the VM), trailing (also adjacent to the VM but just behind the leading cells), deep (underlying the trailing zone, towards the yolk) and stalk (also deep, towards the yolk but further removed from the leading edge) (figure 1b). The cells in all four zones have flattened morphology and much larger nuclei and cell size compared to other area opaca cells more distant from the edge (non-edge cells), but the four zones differ from each other. Extensive membrane ruffles are seen particularly in the leading zone, extending mostly towards the direction of migration (figure 1b). Cells in the stalk zone provide a connection between the edge region and non-edge cells (figure 1b).

This classification reveals quite specific expression patterns for the eight markers (table 2). Figure 1c–r shows representative images in dorsal and sectioned view. RNH1 and SNAI2 are expressed in all edge cells but more weakly in the leading zone (figure 1c,d,g,h). DACT2 is expressed specifically in upper cells (leading and trailing zones, adjacent to the VM) but not in cells in the deep zone (figure 1c,i). MSX1 is specifically expressed in the stalk zone (as well as in neighbouring non-edge cells), but absent in all the other edge sub-regions (figure 1f,j). TGM4 is expressed strongly in the trailing zone but weakly in the other edge zones and non-edge cells (figure 1k,o). Both GATA2 and LHX1 are expressed exclusively in the trailing zone furthest from the leading edge (1 cell thick) (figure 1l,m,p,q), while the latter showed mosaic (heterogeneous) expression in the trailing zone furthest from the leading edge (figure 1m). DKK1 is expressed in the trailing zone and also occasionally in stalk zone (figure 1n,r). To confirm these results, we used double in situ hybridization to visualize the expression of either RNH1 with MSX1, or of DACT with LHX1, in the same embryos. RNH1 and MSX1 are expressed in separate regions: RNH1 in all edge cells except in the stalk zone, while MSX1 is expressed exclusively in the stalk zone (figure 1s,t,u,v,w,y). The expression of DACT2 and LHX1 overlaps, the latter being heterogeneous (mosaic) in the rearmost cells of the trailing zone (figure 1z,1,2).

Next, to investigate if these patterns of expression are maintained throughout the process of expansion, two more stages, 12 h (approx. HH stage 2–3, an early stage of
expansion) and 3 days (approx. HH stage 19, when the edge has passed the equator and is now close to the South pole of the yolk), were investigated for expression of RNH1, SNAI2, DACT2 and MSX1. At 12 h, three of the markers (RNH1, SNAI2 and DACT2) are expressed strongly in edge cells, while MSX1 is expressed in the stalk zone and non-edge cells (electronic supplementary material, figure S1A–L), which is comparable to 24 h embryos in figure 1. Unlike 24 h embryos, expression of RNH1, SNAI2 and DACT2 in non-edge cells is also observed (electronic supplementary material, figure S1I–J). Also, the edge region as a whole at 12 h contains fewer cells in thickness (electronic supplementary material, figure S1E–L) compared to 24 h (figure 1). On the other hand, at 3 days (electronic supplementary material, figure S1M), the edge shows fairly regular, marked infoldings, suggesting that this may be a mechanism for reducing the circumference of the blastoderm edge as embryo expansion reaches its end (electronic supplementary material, figure S1N). All four markers are specifically expressed in the edge cells, but MSX1 is only expressed in the stalk (electronic supplementary material, figure S1O–Z). At 3 days of incubation, the edge has thickened, involving more cell layers, in comparison with a very thin, one-cell layer of non-edge cells (electronic supplementary material, figure S1S–T). This could be an additional mechanism contributing to reducing the circumference of the edge as expansion around the yolk nears completion. These results reveal that, throughout embryo expansion, edge cells maintain expression of the markers while they change their cellular composition, suggesting that some of these genes may have a role in the migration of the edge cells.

Together, our data reveal that the edge region of the embryo can be subdivided into at least four regions using different markers (table 1), implying different biological properties of edge cells in each zone and defining a distinctive structure for the edge region as a whole.

### 2.2. Reorganization of the edge region during reattachment process

Next, we investigated the behaviour of the edge region that accompanies attachment to the VM, using the newly found markers. Embryos were detached from the VM, kept in saline for 30 min before replacing them on the VM and incubation for different periods of time, followed by in situ hybridization to check the expression of edge cell markers (figure 2). RNH1 and SNAI2 were selected as they are expressed in all four zones, while DACT2 and MSX1 were selected because they mark upper edge cells and cells in the stalk zone, respectively.

Table 2. Details of expression of markers in the edge region. Here, the rear zone includes the trailing zone and the deep zone. Numbers: (number of sections with expression) / (total number of sections). n/e: not expressed.

| sub-regions of rear zone | leading zone | rear zone | trailing zone | deep zone | stalk zone |
|--------------------------|-------------|-----------|---------------|-----------|-----------|
| RNH1                     | 33/33       | 33/33     | 33/33         | 29/33     | 0/33      |
| SNAI2                    | 10/14       | 13/14     | 14/14         | 13/14     | 0/33      |
| DACT2                    | 20/30       | 30/30     | 30/30         | 5/30      | 0/33      |
| MSX1                     | 0/21        | 0/21      | n/e           | n/e       | 21/21     |
| TGM4                     | 11/25       | 23/25     | 23/25         | 15/25     | 13/25     |
| GATA2                    | 0/19        | 19/19     | 19/19         | 0/19      | 0/19      |
| LHX1                     | 0/20        | 20/20     | 20/20         | 0/20      | 0/20      |
| DKK1                     | 0/26        | 19/26     | 14/26         | 7/26      | 10/26     |

Several distinct features were observed during the reattachment process. First, three of the four markers (RNH1, SNAI2 and DACT2) are upregulated in non-edge cells just proximal to the edge soon (30 min) after detachment from the VM (figure 2a–g, m–y–an). Second, all four genes gradually expand their expression domains during culture, from a width of 2–3 cells before culture to a maximum of 10 cell diameters from the edge as seen in dorsal view, and 4–6 cells in depth as shown by sections after 2 h (figure 2a–x). The greatest rate of expansion occurs by 2 h (figure 2l, j, p, v). This expansion of the expression domains depends on attachment to the VM, as embryos kept in saline for 2 h showed only weak expression and no expansion of the expression domains (figure 2f, l, r, x). Finally, during reattachment, the marker genes maintain their expression pattern (spatial expression in different zones) regardless of the extra thickness due to additional cell layers (figure 2ab–ac). For example, RNH1 and SNAI2, which are initially expressed in all four zones, are still expressed throughout the edge after reattachment (figure 2ab–ac). DACT2 maintains its expression, limited to the upper cell layer (figure 2ai), and MSX1 is expressed in rear cells (like the stalk zone) but is absent from distal cells (figure 2ae).

The cell morphology, including nuclear shape, of edge cells quickly becomes rounded after detachment (figure 2y–an) and does not fully recover even several hours after reattachment (figure 2ab–ae). Some of the leading cells start to flatten again at 1 h and most of them have done so 2 h after replacing the embryo on the VM (figure 2ab–ae). Figure 2af presents a schematic summary of the cellular changes seen during the reattachment process.

Additionally, the increased adhesion to the VM was also apparent by the embryo retaining its association with the membrane even after the harsh in situ hybridization process from 2 h onwards. Attachment of non-edge cells was frequently observed at multiple points (electronic supplementary material, figure S2A,C). Occasionally, the edge folded back on itself, and the ventral face of the edge region became attached to the VM (electronic supplementary material, figure S2B).
Cells that migrate collectively have unique cellular properties: the cells in the leading zone exhibit filopodial and lamellipodial activity at their leading face while keeping adhesion at their lateral and rear faces to cells that follow them, which in some cases defines a distinct structure comprising a group of cells that migrate collectively as a unit, which has been considered as a ‘giant supracell’ [15,16]. To investigate this in chick edge cells, immunohistochemistry was conducted to visualize the spatial distribution of a variety of cellular components (figures 3 and 4).

First, we checked the localization of cell polarity markers in the edge cells. PKCζ is localized apically in epithelialized cells in the early chick embryo [13]. In non-edge cells of the area opaca, it is also localized apically, but is not detectable in the edge cells (figure 3a), suggesting that the latter lack apical-basal polarity. RHOA and RAC1, members of the Rho family of small GTPases that regulate actin dynamics, show localized expression in migrating cells [17,18]. In the edge cells, both of them show strong localization in the ventral-facing protrusions and elsewhere in the cytoplasm (figure 3b,c). Similar protrusions can be seen in a subset of non-edge cells in the outer area opaca (figure 3b,c). E-cadherin and CTNNB are components of adherens junctions [19]; they are absent from the leading face of the edge cells but strongly expressed at junctions between non-edge cells (figure 3d,e), as

Figure 2. Reorganization of the edge during reattachment. Embryos at HH stage 6 were detached from VM and then cultured by replacing them onto the VM for different time periods before assessing the expression of edge cell markers RNH1, SNAI2, DACT2 and MSX1. As controls, (f, j, r, x) show the expression of the same four genes in embryos incubated in suspension in saline for 2 h. After detachment, the former three makers are induced weakly in non-edge cells proximal to the edge region (arrowheads in (y–ao)). During reattachment, all genes show a gradual increase in the width of the expression domain, especially from 2 h after culture (d, j, p, v). During reattachment, there is an increase in cell layers in the edge, including expression of RNH1, SNAI2 and DACT2 (ab–ad). From 2 h, the front-most edge cells flatten (arrows in (ab, ac)). DACT2 expression is limited to upper edge cells (ad). The expression of MSX1 is absent from the front edge cells (bracket, (ae)). Dashed magenta lines: ventral side of VM. Black arrows: attachment points of the embryo to the VM. White arrows, flattening of the cell in the leading zone. Scale bars: 100 µm. (af) Schematic summary of cellular changes that accompany the reattachment process.
previously shown [7] Notably, in the case of PKCζ, E-cadherin and CTNNB, supracellular cables perpendicular to the direction of migration can be seen at the border of the edge cells (arrowhead). Arrow, no localization at the leading edge of the leading cell. (b,c) Both RHOA (b) and RAC1 (c) show expression in the cytoplasm of the edge cells. Strong localization is seen in the ventral-facing irregular shaped protrusions of the edge cells (arrow), also seen in some non-edge cells (arrowhead in (c)). (d,e) Both E-cadherin and CTNNB are strongly expressed at the junctions between non-edge cells, but both are absent from the leading front of the edge cells (arrow). Supracellular cables perpendicular to the migration direction are seen at the border of the edge cells (arrowhead) (f) No phospho-H3-positive cells are observed in the edge cell regions indicating no proliferating cells. The bottom images under each panel show a slice-view from the confocal image stacks. Left-to-right: proximal-to-distal. Blue colour: DAPI stained nuclei. Brackets: edge cells. Scale bar: 100 µm.

Next, we investigated markers of the ECM and for cellular components that participate in cell-substrate adhesion (figure 4a–f). Fibronectin, normally expressed basally in epithelial cells as a continuous sheet [20], is expressed as a sheet under non-edge cells (figure 4a,b). However, in sections through edge cells, staining is sparse, consistent with organization as cables oriented parallel to the direction of migration [21] (figure 4a,c). Another ECM molecule, laminin, shows punctate staining in edge cells (figure 4f) as previously reported [7]. The ECM receptors ITGA6 and ITGB1 are strongly and specifically localized to the edge cells (figure 4d, e). As most migrating cells exhibit unique cytoskeletal dynamics [15], we investigated markers of some of these. α-tubulin staining reveals a mesh-like structure in the lamellipodia of the leading-edge cells, while supracellular cables (2–3 cells thick) perpendicular to the migration direction can be observed at the border between edge cells and non-edge cells (figure 4g). Phospho-MYL2 and F-actin (phalloidin) staining reveals 2–3 layers of supracellular cables (actomyosin cables) in the edge cells (figure 4h,i), suggesting tension perpendicular to the direction of migration [15]. Punctate localization of phospho-MYL2 was also observed at the free margin of the leading-edge cells (arrowhead, figure 4h). The expression of F-actin is strong in the edge cells, mostly in the cytoplasm (figure 4i). Together, these immunohistochemistry data demonstrate unique molecular features of the edge cells compared to neighbouring non-edge cells.

2.4. Live imaging of edge cells during epiboly
To study the migratory behaviour of edge cells, we used two different approaches to observe different aspects of edge cell

Figure 3. The expression of markers of cell polarity (PKCζ, RHOA and RAC1), cell-to-cell adhesion (E-cadherin and CTNNB), and mitosis (phospho-H3) in edge cells. (a) PKCζ shows apical localization in non-edge cells but is absent from the edge cells. Supracellular cables perpendicular to the migration direction are seen at the border of the edge cells (arrowhead). Arrow, no localization at the leading edge of the leading cell. (b,c) Both RHOA (b) and RAC1 (c) show expression in the cytoplasm of the edge cells. Strong localization is seen in the ventral-facing irregular shaped protrusions of the edge cells (arrow), also seen in some non-edge cells (arrowhead in (c)). (d,e) Both E-cadherin and CTNNB are strongly expressed at the junctions between non-edge cells, but both are absent from the leading front of the edge cells (arrow). Supracellular cables perpendicular to the migration direction are seen at the border of the edge cells (arrowhead) (f) No phospho-H3-positive cells are observed in the edge cell regions indicating no proliferating cells. The bottom images under each panel show a slice-view from the confocal image stacks. Left-to-right: proximal-to-distal. Blue colour: DAPI stained nuclei. Brackets: edge cells. Scale bar: 100 µm.
behaviour. First, we immersed the entire embryo in a lipophilic dye, CM-DiI (CellTracker) to label all cell membranes; this revealed details of lamellipodial and filopodial actions at the membranes of edge cells (figure 5a–d; electronic supplementary material, movie S1). Lamellipodia were constantly generated and resorbed at the migratory front of the outermost edge cell (figure 5a–d; electronic supplementary material, movie S1). Numerous very thin filopodia grew out from the periphery of the lamellipodia (figure 5a,b,d; electronic supplementary material, movie S1), as previously reported [10]. Rapid changes in local fluorescence intensity were observed in these lamellipodia, resembling the focal adhesion points of migrating cells [22,23] (figure 5b; electronic supplementary material, movie S1). As the edge cell moves forward, the lamellipodia retract away from the direction of migration, while filopodia keep their attachment to the VM and do not retract. This suggests a possible combinatorial action of the lamellipodia and filopodia of the edge cells for their migration, as shown in a model of rapid wound healing [24]. Next, we stained the embryo with Calcein-AM (figure 5e–g). This revealed details of lamellipodial action and clearly marks the width of the edge region (red brackets in figure 5c–g; electronic supplementary material, movie S2). Our data confirm that very dynamic changes in morphology take place at the outermost front of the edge cells during chick embryo expansion.

In conclusion, we describe some cellular and molecular characteristics of the edge region, specialized for blastoderm expansion on the VM. This reveals that the region is made up of at least four molecularly and anatomically distinct zones. The molecular markers hint at possible roles of WNT signalling. We propose that the edge region could be a useful model to study other general processes, such as wound healing and cancer cell metastasis, which appear to involve similar migratory behaviours.

3. Materials and methods

3.1. Embryo harvest, fixation and whole-mount in situ hybridization

Fertilized White Leghorn hens' eggs were obtained from Henry Stewart, UK, and incubated for 12 h or 24 h to obtain embryos at HH stage 2–3 or 6–7 [14], respectively, or
3 days (about HH stage 19) at 38°C. To keep the VM attached to the embryo, fixation was carried out as a series of sequential steps during embryo harvest. After opening the egg shell, the embryo was brought to the top by rolling the yolk with a spatula. Two to three drops of 4% paraformaldehyde (PFA) in calcium-magnesium free phosphate-buffered saline (PBS) (pH 7.4) were applied to cover the embryo and the adjacent VM. After 5–10 s, an area of VM slightly larger than the embryo was cut and lifted out with a spoon-spatula. Two to three drops of 4% PFA were applied directly to the embryo on the spatula for another 5–10 s. Embryos still attached to their membrane were cleared of adherent yolk while submerged in Pannett-Compton saline [25] using streams of saline from a glass Pasteur pipette. They were then fixed in 4% PFA either at 4°C overnight prior to whole-mount in situ hybridization, or at room temperature for 1 h prior to immunohistochemistry. Whole-mount in situ hybridization was performed as previously described [26,27]. The probes used were RNH1 (ChEST73n20), SNAI2 [28], DACT2 [29], MSX1 [30], TGM4 (ChEST698a23), GATA2 [31], LHX1 (ChEST389n6) and DKK1 [32]. For double in situ hybridization, the first probe was labelled with fluorescein isothiocyanate and cyan colour development was done with 5-bromo-4-chloro-3-indolyl phosphate (BCIP) only. The second probe was labelled with digoxigenin and dark purple colour development was done with nitro blue tetrazolium plus BCIP. After imaging as whole mounts, embryos were embedded in paraffin and sectioned at 10 µm in a Zeiss Microm microtome. Sections were mounted with Vectashield Plus antifade mounting medium which contains 4',6-diamidino-2-phenylindole (DAPI) (H-2000, Vector Laboratories) as a nuclear stain. Photographs of sections and whole mounts

Figure 5. Dynamics of edge cell migration using different staining methods. Frames from a time-lapse sequence showing membrane activity of edge cells during migration using Dil staining (a–d) and Calcein-AM staining (e–g). (a–d): Dil staining). Retraction (bracket in (a)) and de novo generation (bracket in (b)) of lamellipodia are observed. Simultaneously, many highly active filopodia (white arrows) are observed protruding beyond the edge of the lamellipodia. In the lamellipodia, rapid appearance and disappearance of fluorescent localization (arrowheads) is observed, which may correspond to focal adhesions. During cell crawling (dashes in (c) and dashed arrows in (d)), the lamellipodia retract while the filopodia keep their attachments (c,d). The pseudo-colour scale reflects fluorescence intensity. Black arrows: lamellipodia. (e–g) Calcein-AM staining. Active lamellipodial activity of the membrane of edge cells is observed. The staining clearly marks the extent of the edge region (red brackets). Grey arrow in (a,e): direction of edge cell migration. Scale bar: 20 µm in (a), 100 µm in (e).
were obtained with an Olympus Vanox-T microscope with a QImaging Retiga 2000R camera.

3.2. Live imaging

Two different dyes were used for live staining of embryos and imaging; CM-Dil (C7001, Invitrogen) and Calcein-AM (17783, Sigma-Aldrich). For CM-Dil staining, embryos were incubated in PBS containing 4 µM CM-Dil in 0.2% dimethyl formamide in Pannett-Compton saline at 38°C for 30 min and then washed with PBS three times. The embryo was then cultured using a modification of the New culture method [33,34] at 38°C for at least 1 h. For Calcein-AM staining, the embryos were first set-up for culture, and then a few drops of 10 µM Calcein-AM in PBS were applied to the edge region of the embryos. The embryo was then imaged with a Leica SPE inverted confocal microscope at 38°C.

3.3. Immunohistochemistry and antibodies

For whole-mount immunostaining, fixed embryos were dehydrated with ice-cold methanol and then rehydrated gradually in steps by adding PBS containing 1% Triton X-100 (PBST). After further washing three times with PBST, they were blocked with blocking buffer (PBST containing 5% normal goat serum and 0.02% thimerosal) for 2–6 h at room temperature on a rocker. The embryos were then incubated at 4°C for 2–3 days with primary antibodies: PKCζ (SC-216, Santa Cruz), RHOA (SC-179, Santa Cruz), RAC1 (05-389, Millipore), E-cadherin (610182, BD Biosciences), β-catenin (C7207, Sigma), phospho-H3 (06-570, Millipore), laminin (31 or 31-2, Developmental Studies Hybridoma Bank, DSHB), α-tubulin (2125S, Cell Signalling Technology) and phospho-MYL2 (3674S, Cell Signalling Technology), all diluted 1 : 40 except the DSHB antibodies which were diluted 1 : 10. After washing three times with PBST, the embryos were incubated at 4°C for 1 day with fluoresecently labelled secondary antibodies: Alexa Fluor 488-conjugated goat anti-mouse IgG (A21202, Invitrogen), Alexa Fluor 488-conjugated goat anti-rabbit IgG (A11008, Invitrogen) or Cy3-conjugated anti-mouse IgG (115-165-073, Jackson), all diluted 1 : 200. For filamentous actin staining, embryos were incubated with rhodamine-phalloidin (Invitrogen R415). For fibronectin staining (VA1, DSHB), integrin-α6 (P2C62C4, DSHB) and integrin-β1 (V2E9, DSHB), fixed embryos were embedded in paraffin and sectioned, then antigen retrieval was conducted in 10 mM sodium citrate (pH 6) for 10 min at 98°C. Then, sectioned tissues were incubated for 1 day at 4°C with primary antibodies, and for 1 h at room temperature with secondary antibodies. For nuclear staining, 2.5 µg ml−1 of DAPI was applied to the embryos or sections for 10 min and then washed thoroughly. For whole embryo mounting, the embryo was transferred onto a slide dorsal side-up and flattened carefully, then the excessive solution was removed using a paper tissue. Twenty microlitres of mounting medium (FH-1000-10, VECTASHIELD) were applied before covering with a coverslip. The stained embryos and sectioned tissues were imaged with a Leica SPE1 confocal microscope. The images were processed using Fiji software [35] to obtain a maximum projection or a slice view.

Data accessibility. The data are provided in the electronic supplementary material [36].

Authors’ contributions. H.C.L.: conceptualization, formal analysis, investigation, methodology, validation, visualization, writing—original draft and writing—review and editing; Y.F.: investigation and writing—review and editing; C.D.S.: funding acquisition, project administration, supervision, writing—original draft and writing—review and editing.

All authors gave final approval for publication and agreed to be held accountable for the work performed therein.

Conflict of interest declaration. The authors declare no competing interests.

Funding. This research was funded by a Wellcome Trust Investigator Award (grant no. 107055/Z/15/Z) to C.D.S. H.C.L. was supported by the above Wellcome Trust award and by a Basic Science Research Program through the National Research Foundation of Korea (NRF) funded by the Ministry of Education (grant no. 2014R1A6A3A0303468). Y.F. was supported by a scholarship from the Custodian of The Two Holy Mosques’ External Scholarship Program from the Ministry of Education of Saudi Arabia (grant no. SE-12267).

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