Fgfr-Ras-MAPK signaling is required for apical constriction via apical positioning of Rho-associated kinase during mechanosensory organ formation

Molly J. Harding1,2 and Alex V. Nechiporuk1,*

SUMMARY
Many morphogenetic movements during development require the formation of transient intermediates called rosettes. Within rosettes, cells are polarized with apical ends constricted towards the rosette center and nuclei basally displaced. Whereas the polarity and cytoskeletal machinery establishing these structures has been extensively studied, the extracellular cues and intracellular signaling cascades that promote their formation are not well understood. We examined how extracellular Fibroblast growth factor (Fgf) signals regulate rosette formation in the zebrafish posterior lateral line primordium (pLLp), a group of ~100 cells that migrate along the trunk during embryonic development to form the lateral line mechanosensory system. During migration, the pLLp deposits rosettes from the trailing edge, while cells are polarized and incorporated into nascent rosettes in the leading region. Fgf signaling was previously shown to be crucial for rosette formation in the pLLp. We demonstrate that activation of Fgfr receptor (Fgfr) induces intracellular Ras-MAPK, which is required for apical constriction and rosette formation in the pLLp. Inhibiting Fgfr-Ras-MAPK leads to loss of apically localized Rho-associated kinase (Rock) 2a, which results in failed actomyosin cytoskeleton activation. Using mosaic analyses, we show that a cell-autonomous Ras-MAPK signal is required for apical constriction and Rock2a localization. We propose a model whereby activated Fgfr signals through Ras-MAPK to induce apical localization of Rock2a in a cell-autonomous manner, activating the actomyosin network to promote apical constriction and rosette formation in the pLLp. This mechanism presents a novel cellular strategy for driving cell shape changes.

KEY WORDS: Fgfr-Ras-MAPK, Rho-kinase, Apical constriction, Lateral line primordium

INTRODUCTION
In many species, apical constriction drives cell shape changes that are important for diverse developmental processes including gastrulation and neural tube closure (Sawyer et al., 2010). This cellular behavior is typically dependent on contraction of an actomyosin network composed of non-muscle myosin (Myosin II) and an apical meshwork of filamentous actin (F-actin). To form this actomyosin network, an apical domain is established through Par protein activity (Suzuki and Ohno, 2006) and the microtubule organizer γ-tubulin is apically localized. Subsequently, cytoskeletal components including F-actin, Myosin II and adherens junctions accumulate apically. Finally, the molecular motor Myosin II is activated by phosphorylation of Myosin regulatory light chain (MRLC), relieving auto-inhibition and promoting constriction (reviewed by Sawyer et al., 2010; Somlyo and Somlyo, 2003). Activation of MRLC is crucial for myosin activity and is mediated by multiple kinases, including Rho-associated kinase (Rock) (Vicente-Manzanares et al., 2009). Whereas the cytoskeletal network driving constriction is fairly well understood, how extracellular cues regulate this process during development is largely unknown. Here, we examine how extracellular signals regulate apical constriction during rosette formation in the zebrafish posterior lateral line primordium (pLLp).

The pLLp is a group of ~100 cells that migrate along the developing embryonic trunk between 22 and 48 hours postfertilization (hpf). The pLLp is organized into rosettes (Ghysen and Dambly-Chaudière, 2007), within which cells are apically constricted, with basally displaced nuclei (Lecaudey et al., 2008). During migration, the pLLp deposits mature rosettes from the trailing edge that give rise to mechanosensory neuromasts. Concurrently, new cells are generated, polarized and assembled into nascent rosettes in the leading edge (Nechiporuk and Raible, 2008). We and others have previously demonstrated that rosette renewal in the pLLp is dependent on Fibroblast growth factor (Fgf) signaling (Lecaudey et al., 2008; Nechiporuk and Raible, 2008). However, how extracellular Fgf is interpreted intracellularly to control cell shape is not understood.

Fgfs are secreted molecules that promote diverse cellular processes including survival and fate specification. Upon binding to transmembrane receptors (Fgfrs), Fgfs activate multiple intracellular cascades, including the Ras-MAPK (ERK) pathway (Tsang and Dawid, 2004). Ras-MAPK signaling mediates multiple processes ranging from transcription to cytoskeletal remodeling (Tsang and Dawid, 2004). However, whether Ras-MAPK mediates Fgf signaling and cell shape changes in the pLLp is unknown.

We show that the Ras-MAPK pathway is activated in the pLLp in an Fgf-dependent manner, and that Fgfr-Ras-MAPK signaling is required cell-autonomously for apical constriction and apical positioning of Rock, two necessary processes for rosette formation. This previously unrecognized mechanism is crucial for proper pLLp development and might reveal a novel cellular strategy for driving cell shape changes.
MATERIALS AND METHODS
Fish strains, heat shock, pharmacological treatments and live imaging
Adult zebrafish were maintained under standard conditions (Westerfield, 1995). The pLLp was visualized using Tg(–8.0claudinB:lynEGFP)p^106 (Haas and Gilmour, 2006), referred to here as claudinB:EGFP. Ras signaling was conditionally inhibited using Tg(hsp70:dnHRAS.cryaa:EGFP)p^167 (also called hsp70:dn-Ras) embryos (Lee et al., 2009) heat shocked at 28 hpf for 40 minutes at 38°C, unless otherwise indicated. Inhibitors were diluted in embryo medium containing 1% DMSO and used at the following concentrations unless indicated otherwise: 100 µM SU5402 (Calbiochem), 7 µM PD0325901 (Stemgent) and 50 µM Rockout (Calbiochem). Live imaging was performed as described (Nechiporuk and Raible, 2008).

Plasmids and injections
par3-EGFP plasmid (von Trotha et al., 2006) was modified to express a Par3-TagRFP fusion. par3-TagRFP mRNA was synthesized using the mMessage mMachine Kit (Life Technologies) and microinjected at 500 pg/embryo.

Immunolabeling and in situ hybridization
Immunolabeling followed established protocols (Ungos et al., 2003), with the following exceptions: embryos stained with anti-Rock2a and anti-pMAPK were fixed in Glyo-Fixx (Thermo Scientfic) and embryos stained with anti-pMRCLC were fixed in Bouin’s fixative (Ortiz-Hidalgo, 1992). Antibodies used: rabbit anti-GFP (1:1000; Invitrogen), mouse anti-GFP (1:100; Roche), rabbit anti-pMAPK (1:150; Cell Signaling), rabbit anti-Rock2a (1:50; Anaspec), rabbit anti-pMRCLC (1:20; Cell Signaling), rabbit anti-γ-tubulin (1:1000; Sigma), rabbit anti-Myosin IIA (1:500; Sigma); and Alexa Fluor 568 phalloidin (1:500; Invitrogen). In situ hybridization was performed as described (Andermann et al., 2002). Digoxigenin-labeled antisense RNA probes were generated for fgfr1 (Scholpp et al., 2004), pe3 (Raible and Brand, 2001) and fgfr10a (Grandel et al., 2000). Fluorescently labeled embryos were imaged using an FV1000 confocal microscope (Olympus). Images were processed in ImageJ (Abramoff, 2004); brightness and contrast were adjusted in Adobe Photoshop.

Cell shape and fluorescence intensity analysis
Three-dimensional reconstructions were generated from images of claudinB:EGFP pLLp using Imaris software (Bitplane). Apical constriction indexes (ACIs) were generated from manually collected measurements. Measurements were taken from the surface of each cell oriented toward the midline. Apical width measurements were made 1 µm below the apical surface. Height measurements represent a straight line between the apical and basal surfaces. Rock2a apical intensity was measured from the average surface. Height measurements represent a straight line between the apical midline. Apical width measurements were made 1 µm below the apical surface.

Statistical analysis
We used JMP (SAS) to perform one-way ANOVA, Wilcoxon and Tukey-Kramer post-hoc tests. Student’s t-tests were performed in Microsoft Excel. Data are presented as mean ± s.e.m.

RESULTS AND DISCUSSION
Rosette formation is dependent on the Fgfr-Ras-MAPK signaling cascade
To address how Fgf signals are intracellularly interpreted within the pLLp, we tested whether the MAP kinase (MAPK) signaling pathway, a common mediator of Fgfr signaling, is active in the pLLp. Activation of the MAPK signaling cascade, as detected by phosphorylated MAPK (pMAPK) immunostaining, was observed in pLLp cells and surrounding tissues (Fig. 1A). Although the MAPK expression pattern was somewhat variable, the majority of cells with high levels of pMAPK were positioned in forming or existing rosettes (Fig. 1A). pMAPK was not detectable in the leading-most progenitor cells of the pLLp (McGraw et al., 2011). Blocking Fgfr signaling with the chemical inhibitor SU5402 (Mohammadi et al., 1997) resulted in pMAPK loss, indicating that Fgf activity is required for MAPK cascade activation (Fig. 1B). Consistent with previous observations (Lecaudey et al., 2008; Nechiporuk and Raible, 2008), inhibiting Fgfr activity also interfered with rosette formation in the leading zone (Fig. 1A,B, yellow bracket). Blocking Ras-MAPK signaling using hsp70:dn-Ras heat-shock-inducible dominant-negative transgensics or the MAPK kinase (MAPKK) inhibitor PD0325901 resulted in loss of the pMAPK signal and a failure of leading rosette formation, phenotypes similar to those observed following Fgfr inhibition (Fig. 1C,D). Additionally, prolonged treatment with PD0325901 caused a progressive expansion of the rosette-free region (data not shown). We thus conclude that Fgfr-Ras-MAPK signaling is necessary for MAPK activation, which is in turn required for rosette formation.

Similar to Fgfr inhibition, inhibiting MAPKK depressed Fgf target gene expression (fgfr1 and pe3), suggesting that MAPK mediates Fgf-dependent patterning in the pLLp (supplementary material Fig. S1) (Nechiporuk and Raible, 2008). Combinatorial treatment with suboptimal doses of Fgfr and MAPKK inhibitors between 30 and 48 hpf arrested pLLp migration, implying that Fgf and MAPK function in the same pathway (supplementary material Fig. S2). Together, these data show that MAPK is the intracellular transducer for Fgfr signaling in the pLLp.

The Fgfr-Ras-MAPK signaling cascade is required for apical constriction in leading pLLp cells
Apical constriction has been suggested to underlie pLLp rosette formation (Hava et al., 2009; Chitnis et al., 2012). Cell shape analysis during live imaging of rosette formation revealed that, over a 1-hour period, columnar cells become apically constricted as nascent rosettes form (Fig. 1E-G), confirming that apical constriction promotes rosette formation.

Because blocking Fgfr-Ras-MAPK signaling caused a failure of leading rosette formation, we examined whether Fgfr-Ras-MAPK inhibition disrupts apical constriction. We limited our analyses to the leading 30 pLLp cells, as our live imaging indicated that this population gives rise to new rosettes. We measured apical constriction indices (ACIs; ratios of lateral height to apical width) of pLLp cells after 2 hours of Fgfr, Ras or MAPKK inhibition, and compared them with DMSO-treated controls. Control cells had ACIs of pLLp cells after 2 hours of Fgfr, Ras or MAPKK inhibition, and compared them with DMSO-treated controls. Control cells had ACIs of pLLp cells after 2 hours of Fgfr, Ras or MAPKK inhibition, and compared them with DMSO-treated controls. Control cells had ACIs of pLLp cells after 2 hours of Fgfr, Ras or MAPKK inhibition, and compared them with DMSO-treated controls.

Transplantation experiments
Transplantation experiments were carried out as previously described (Nechiporuk and Raible, 2008). Host embryos expressed claudinB:EGFP pLLp using Imaris software (Bitplane). Apical constriction indexes (ACIs) were generated from manually collected measurements. Measurements were taken from the surface of each cell oriented toward the midline. Apical width measurements were made 1 µm below the apical surface. Height measurements represent a straight line between the apical and basal surfaces. Rock2a apical intensity was measured from the average fluorescence intensity in the apical domain (apical one-third of the cell) compared with the rest of the cell (the basal two-thirds).

Statistical analysis
We used JMP (SAS) to perform one-way ANOVA, Wilcoxon and Tukey-Kramer post-hoc tests. Student’s t-tests were performed in Microsoft Excel. Data are presented as mean ± s.e.m.

Fgfr-Ras-MAPK signaling is not required for subcellular distribution of apical polarity and cytoskeletal components
Analyses in other models have shown that apical polarity establishment and actomyosin network assembly are important for apical constriction (Sawyer et al., 2010). Because Fgfr-Ras-MAPK signaling underlies apical constriction in the pLLp, we...
examined whether MAPK activation modulates the localization of polarity [Par3 (Pard3) and γ-tubulin] or cytoskeletal (Cadherin 2, F-actin and Myosin II) components. We found that Par3 and γ-tubulin were apically localized in the leading cells, caudal to the first rosette, demonstrating that cells are polarized prior to apical constriction and rosette formation (supplementary material Fig. S4A-H and Table S1). Inhibiting Fgfr or MAPKK did not disrupt the apical localization of these polarity components (supplementary material Fig. S4A-H). Cadherin 2, F-actin and Myosin II were also localized to apical domains, but slightly more rostral than to the polarity markers (supplementary material Fig. S4I-T). The localization of these components was unchanged following inhibition of Fgfr or MAPKK (supplementary material Fig. S4I-T). These data indicate that Fgfr-Ras-MAPK signaling is not required for the localization or assembly of cytoskeletal polarity machinery in the pLLp.

**Rock is required for apical constriction and activation of MRLC**

In order for Myosin II to generate force, it must be activated through MRLC phosphorylation (Vicente-Manzanares et al., 2009). Because Rho-associated kinase (Rock) is known to phosphorylate MRLC in other contexts (Ishiuchi and Takeichi, 2011; Plageman et al., 2011), we examined whether Rock is required for myosin activation during rosette formation in the pLLp. We treated embryos with the Rock inhibitor Rockout (Weiser et al., 2007) for 2 hours beginning at 28 hpf, and assayed constriction and MRLC phosphorylation. We observed apical constriction failures in leading-edge cells following Rockout treatment (Fig. 2A-E). Rock inhibition also resulted in failure of MRLC phosphorylation in the leading edge (Fig. 2F-H). From these data, we conclude that Rock activity is required to activate MRLC in the pLLp and to drive apical constriction during rosette formation.

**Fgfr-Ras-MAPK signaling is required for subcellular distribution of Rock2a**

Next, we investigated whether Fgfr-Ras-MAPK regulates the subcellular localization of Rock during pLLp apical constriction. We focused on Rock2a as it is the predominant Rho kinase in neuronal tissues (Amano et al., 2010) and is expressed in the pLLp (supplementary material Fig. S5). A second isoform, Rock2b, does not appear to be expressed in the pLLp (Wang et al., 2011). In control embryos, Rock2a was apically localized in leading cells. However, following Fgfr or MAPKK inhibition, Rock2a failed to segregate apically and appeared dispersed throughout the cell (Fig. 3A-C). rock2a expression was unaffected by MAPKK inhibition, suggesting that MAPK does not regulate rock2a transcription (supplementary material Fig. S5).

As Rock is required for phosphorylation of MRLC, we examined whether Fgfr-Ras-MAPK activity is also necessary for MRLC phosphorylation. Indeed, pMRLC was lost from leading pLLp cells in embryos treated with Fgfr or MAPKK inhibitors (Fig. 3D-F). This implies that proper Rock2a localization is required for the activation of MRLC. Interestingly, the loss of apical Rock2a and of MRLC activation were unchanged in trailing rosettes following Fgfr-Ras-MAPK inhibition, indicating that this pathway is required for the initiation of Rock2a apical positioning but not its maintenance. Finally, combining suboptimal inhibition...
of Ras-MAPK signaling with suboptimal inhibition of Rockout resulted in a failure of MRLC activation in the leading pLLp cells (supplementary material Fig. S6 and Table S2), suggesting that Ras-MAPK and Rock2a act in the same pathway.

**Ras-MAPK signaling is required cell-autonomously for Rock2a localization and apical constriction**

We next investigated whether individual pLLp cells lacking Ras-MAPK signaling are capable of apically constricting in the context of wild-type neighbors. We generated mosaic embryos that contained a small number of hsp70:dn-Ras or wild-type (control) donor cells. Mosaic embryos were heat shocked at 28 hpf to inhibit Ras activity, and pMAPK and apical constriction were assayed at 30 hpf. As expected, induction of dn-Ras in donor cells resulted in loss of pMAPK (Fig. 4A, B). Reconstructed cell shapes from before and after heat shock showed that wild-type cells constricted over the 2-hour period (Fig. 4C). By contrast, the average ACIs of Ras-deficient cells did not change, indicating constriction failures (Fig. 4D-E). Loss of Ras-MAPK activity also corresponded to cell-autonomous loss of apical Rock2a in dn-Ras cells (Fig. 4H-I). However, this loss had no obvious effect on the overall distribution of Rock2a (Fig. 4F-G'). Our data show that individual cells constrict in response to the apical accumulation of Rock2a initiated by intracellular Ras-MAPK, and not as a result of neighbor interactions.
An intriguing question raised by this work is that of how Fgf-Ras-MAPK acts to regulate the apical localization of Rock2a. Although Rock does not contain a consensus sequence for MAPK phosphorylation, MAPK might regulate Rock via phosphorylation and suppression of p190A RhoGAP activity and subsequent promotion of RhoA/Rock activity (Pullikuth and Catling, 2010). Alternatively, Fgf signals have been shown to regulate transcription of shroom3 (Chitnis et al., 2012), which encodes a scaffolding molecule that can anchor Rock and promote apical Rock localization (Nishimura and Takeichi, 2008).

**Conclusions**

Our observations reveal a novel role for Fgf signaling during morphogenesis that might represent a general strategy for controlling apical constriction. Fgf signaling has previously been implicated in apical constriction via activation of basally localized Myosin II during inner ear morphogenesis (Sai and Ladher, 2008). However, our data provide the first example that Fgfr activation, through the Ras-MAPK pathway, can regulate the localization of Rock without affecting other polarity or cytoskeletal components. When combined with the role of Fgf in pLp hair cell specification (Nechiporuk and Raible, 2008), this model elucidates the elegant manner in which Fgf couples morphogenesis and patterning of the pLp.

**Acknowledgements**

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**Competing interests statement**

The authors declare no competing financial interests.

**Supplementary material**

Supplementary material available online at http://dev.biologists.org/lookup/suppl/doi:10.1242/dev.082271/-/DC1

**References**


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DMSO control

7 μM PD0325901

0.5 μM PD0325901 + 20 μM SU5402

36ºC heat-shock dn-Ras

20 μM SU5402

0.5 μM PD0325901 + 20 μM SU5402

0.5 μM PD0325901 + 36ºHS dn-Ras
**A** Absolute cell # from the pLLp tip

**A'**

**B** Apical constriction index (ACI) for DMSO control

**B'**

**C** Apical constriction index (ACI) for hsp70:dn-Ras

**C'**

**D** Apical constriction index (ACI) for 7 μM PD0325901

**D'**

**E** Graph showing the relationship between absolute cell # from the pLLp tip and apical constriction index (ACI). Lines represent different treatments:
- Red: DMSO control
- Purple: SU5402
- Green: hsp70:dn-Ras
- Blue: PD0325901

**F** Magnified images showing the difference in cell size and shape due to different treatments.
30 hpf rock2a

A

DMSO control

B

2 hours PD0325901

C

4 hours PD0325901
Supplemental figure 6

DMSO control

A

A'

36°C heat-shock hsp70:dn-Ras

B

B'

36°C hs hsp70:dn-Ras/10 μM Rockout

E

E'

0.5 μM PD0325901 / 10 μM Rockout

F

F'

10 μM Rockout

C

C'

10 μM Rockout

D

D'

0.5 μM PD0325901

G

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<th>Treatment</th>
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<tr>
<td>10 μM Rockout</td>
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<tr>
<td>10 μM Rockout+ dn-Ras 36°C</td>
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n.s.  **
Table S1. Quantification of cytoskeletal recruitment in the leading zone of the pLLp Par3-tagRFP

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<th>PD0325901</th>
<th>DMSO</th>
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<tr>
<td>Mean</td>
<td>11.8%</td>
<td>13.5%</td>
<td>11%</td>
<td>10.2%</td>
<td>10.1%</td>
<td>11.3%</td>
<td>19.2%</td>
<td>15.36%</td>
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<tr>
<td>s.e.m.</td>
<td>0.52%</td>
<td>0.97%</td>
<td>0.88%</td>
<td>0.62%</td>
<td>0.7%</td>
<td>0.71%</td>
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<tr>
<td>P-value</td>
<td>0.67</td>
<td>0.88</td>
<td>0.88</td>
<td>0.67</td>
<td>0.71</td>
<td>0.71</td>
<td>0.67</td>
<td>0.71</td>
</tr>
</tbody>
</table>

Mean 17.7% 18.9% 16.1% 17.6% 19% 16.5%
s.e.m. 0.51% 0.45% 1.2% 1.13% 0.55% 0.93%
P-value 0.369 0.407

Myosin II

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<td>Mean</td>
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<td>P-value</td>
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To calculate means, cells were counted from n=6 pLLps in each condition. Data were analyzed using one-way ANOVA.
Table S2. Pairwise comparisons between Rockout and Ras-MAPK synergy conditions

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<td>23.3±6.6%</td>
<td>24.5±4.5%</td>
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<td>33.4±6.0%</td>
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<td>DMSO</td>
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<td>36°C HS/10 μM</td>
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<tr>
<td>Rockout</td>
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<td>10 μM Rockout/0.5 μM PD0325901</td>
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To calculate means, cells were counted from n=10 pLLps in each condition. P-values were derived from Tukey-Kramer post-hoc tests. Heat shock (HS) conditions were 40 minutes at 36°C.