FGF8 coordinates tissue elongation and cell epithelialization during early kidney tubulogenesis.

Author(s)
Atsuta, Yuji; Takahashi, Yoshiko

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ABSTRACT
When a tubular structure forms during early embryogenesis, tubular elongation and lumen formation (epithelialization) proceed simultaneously in a spatiotemporally coordinated manner. We here demonstrate, using the Wolffian duct (WD) of early chicken embryos, that this coordination is regulated by the expression of FGF8, which shifts posteriorly during body axis elongation. FGF8 acts as a chemoattractant on the leader cells of the elongating WD and prevents them from epithelialization, whereas static (‘rear’) cells that receive progressively less FGF8 undergo epithelialization to form a lumen. Thus, FGF8 acts as a binary switch that distinguishes tubular elongation from lumen formation. The posteriorly shifting FGF8 is also known to regulate somite segmentation, suggesting that multiple types of tissue morphogenesis are coordinately regulated by macroscopic changes in body growth.

KEY WORDS: Wolffian duct/nephric duct, Chemoattraction, Live imaging, Body axis

INTRODUCTION
In many organs, tubular epithelia perform key physiological functions. For example, they convey food in the gut, exocrine factors from the pancreas and wastes from the kidney. For these functions, tubular integrity is important because its failure would cause pathological defects (Andrew and Ewald, 2010). To understand how such tubular integrity is established, it is necessary to delineate the mechanisms underlying epithelial tubule formation.

Tubule formation involves two major, coordinated steps: tubular elongation, a process that controls the length of the tube; and cell epithelialization, a process that forms an internal lumen. These two events must take place in a spatiotemporally coordinated manner. However, how such coordination is achieved at the cellular and molecular levels has remained largely unexplored. This lack of knowledge is mainly due to the lack of experimental models that can simultaneously address the integrated questions regarding tubule elongation and epithelialization.

We have recently developed a novel technique of in ovo electroporation that enables gene manipulation of the forming Wolffian duct (WD; also called the nephric duct) in chicken embryos (Atsuta et al., 2013). The WD emerges in the anterior intermediate mesoderm (IMM) of the pronephric region, and subsequently extends caudally as a straight cord along a stereotypic path in between the presomitic mesoderm (PSM) and lateral plate (Obara-Ishihara et al., 1999; Saxén and Sariola, 1987). During WD elongation, the mesenchymal cord progressively hollows to form a single-layered epithelial tube through the process of mesenchymal-epithelial transition (MET). Importantly, cells located at the leader of the elongating WD (‘leader’ cells) are mesenchymal in shape and highly motile, as previously reported in chickens (Atsuta et al., 2013) and mice (Chia et al., 2011; Soofi et al., 2012), whereas ‘rear’ cells are epithelial and less motile (static).

Here, we studied how the mesenchymal and epithelial states are coordinately regulated in both time and space during WD elongation. We asked three questions: (1) what regulates the behavior of leader cells; (2) what determines the relative locations of the leader and static rear cells; and (3) what triggers epithelialization/lumenization? We found that FGF8, which is produced in a caudal region of the embryo (Dubrulle and Pourquie, 2004), plays crucial roles in these processes. FGF8 not only maintains the mesenchymal state of the leader cells, but also acts as a direct chemoattractant for their path finding. Since the FGF8-positive domain shifts caudally as the tail region elongates, the anteriorly positioned WD cells (i.e. rear cells) receive progressively less FGF8 signal, leading to their epithelialization and concomitant lumenization. Thus, tubule formation is harmonized with the growth rate of the embryo via FGF signals: mesenchymal and epithelial cells coordinately participate in elongation and lumenization, allowing tubule formation at the same rate as body axis elongation. Coordinated morphogenesis between the body axis elongation, WD elongation and somite segmentation is also discussed. Our results are in part consistent with those reported recently by Attia et al. (2015), who also showed the importance of FGF signals for WD elongation.

RESULTS
Tissue elongation is coordinated with cell epithelialization during WD formation
It is known that the WD emerges from the anteriorly located pronephric region of HH10 chick embryos, spanning the sixth to twelfth somite levels (Hiruma and Nakamura, 2003). Subsequently, the WD extends posteriorly as a simple straight cord, and this elongation is in register with somitic segmentation: the leader of the extending WD is constantly located in the PSM (unsegmented) at the level of one to two presumptive somites posterior to the most recently formed somite [somite level (sm) –1 to –2] (Atsuta et al., 2013; Saxén and Sariola, 1987). We found in HH13 embryos that the cells at the leader of the WD were mesenchymal with no tubular structure, whereas those located anterior to sm V (the fifth somite anterior to the forming somite) were part of an epithelial tubule. In a transverse view, WD cells at sm V were enclosed by the basal marker laminin 1, a component of the extracellular matrix (ECM), and exhibited apicobasal polarity as revealed by the tight junction marker ZO-1.
and E-cadherin (Fig. 1A-C; n=10). By contrast, in the leader cells at sm –1, signals for laminin 1 and ZO-1 were sparse, with no obvious apicobasal polarity of E-cadherin signals (Fig. 1D,E; n=10). These observations were also supported by observation of sagittally sectioned specimens (Fig. 1F,G; n=10). In this study, we defined leader cells as those located between sm –2 and I of the WD and rear cells as those between sm V and VII. Although leader cells exhibit sparse and weak laminin 1 and ZO-1 signals, we describe the leader cells as ‘mesenchyme’, so that morphological differences between leader cells and rear cells (epithelial) can be clearly highlighted.

To examine how the leader and rear cells behave in embryos, we performed time-lapse live imaging confocal microscopy at high resolution. During WD elongation, which progresses in register with somite segmentation (Fig. 1H; supplementary material Movie 1; n=3 embryos), leader cells were elongated in shape and possessed numerous cellular processes that repeatedly extended and retracted in the direction of WD elongation (Fig. 1J; supplementary material Movie 3). By contrast, static cells were polygonal in shape with few cellular processes, as is characteristic of epithelial cells (Fig. 1I; supplementary material Movie 2). In addition, long-term (270 min) tracing of WD cells demonstrated that leader cells were highly motile compared with rear cells (Fig. 1KL; n=3 embryos).

The spatial relationship between the mesenchymal (leader) and epithelial (rear) cells was maintained regardless of the number of somites (embryonic stages), at least until HH16 (data not shown). Thus, these two types of cells were coordinated during WD elongation. Since the mesenchymal cells at the leader and the epithelial cells in the rear would serve for tubule elongation and lumenization, respectively, we reasoned that mesenchymal-epithelial coordination between the leader and rear would be crucial for correct WD formation in terms of size and shape.

Leader cells are attracted to their adjacent intermediate mesoderm

We explored the cellular and molecular mechanisms by which spatiotemporal coordination is achieved between the motile leader cells and epithelial static cells during WD elongation. To specifically address the differential regulation of these two cell types, we analyzed
how the motility of leader cells is regulated in the context of interactions between adjacent tissues. Unlike the situation reported in axolotl embryos (Morris et al., 2003), we found that the surface ectoderm appears to be dispensable, blockage of ectodermal signals by removing this tissue (supplementary material Fig. S1A-C; n=12/15) or by inserting a piece of aluminium foil underneath this tissue (supplementary material Fig. S1D-E′; n=7/9) did not affect elongation of the WD, at least until HH15.

To determine which tissue(s) regulates WD elongation, we ectopically placed a piece of leader WD into the lateral plate region (Fig. 2A). The grafted WD, which was labeled by PKH26, was attracted medially toward the IMM of host embryos (Fig. 2B; n=17/23). This attraction was also observed in a similar experiment in which a host WD had been ablated (Fig. 2C,D; n=7/9), suggesting that the IMM might be a primary source of attracting signals for the leader cells. To locate precisely the leader cell-attracting site in the IMM along the anteroposterior (AP) axis, leader cells dissected from embryos were PKH26 labeled and placed into five different positions along the AP axis in the lateral plate of the host embryo (Fig. 2E). We found that only the two posterior transplants moved to the IMM (Fig. 2F; n=4/4). Thus, during normal development the IMM near the leader cells, but not the IMM near the rear, provides cues for WD elongation. Of note, even the rear WD cells reacted to the leader-attracting signal: when relocated to the leader position of WD, the rear cells resumed migration (Fig. 2G,I; n=5), whereas this phenomenon was not observed when they were placed back in original position as a control (Fig. 2G,H; n=5). Thus, even quiescent epithelial WD (rear) cells can resume migration if they encounter attractive signals. Hereafter, we refer to the IMM region that surrounds the leader cells as ‘leader IMM’ (L-IMM) (Fig. 2J).

To further examine the leader cell-attracting activity exerted by L-IMM, we grafted a piece of L-IMM into a region adjacent to the leader WD of a host embryo (Fig. 2K). This resulted in ectopic branching of the host WD, which migrated toward the graft (Fig. 2L; n=12/27). Furthermore, the grafted L-IMM also induced ectopic branching from quiescent epithelial tube from the rear WD (approximately sm VI) (Fig. 2M,N; n=6/12). Thus, the L-IMM exerts attracting activities to both leader and rear WD. Such activity was not seen when lateral plate mesoderm (LPM) was used as a control (Fig. 2Q,R; n=0/9).

Together, these observations suggest that WD elongation is regulated by interactions between the WD leader and its adjacent L-IMM. The L-IMM exerts activities that might induce or promote the motility and migration of leader cells, and such activities must be located specifically to the region surrounding the leader cells.

FGF signaling is activated in the elongating WD

To identify the WD-attracting signaling molecule(s) elicited by L-IMM, we focused on fibroblast growth factor (FGF) signals as candidates for the following reasons. First, FGF8 is known to be expressed in PSM (Dubrulle et al., 2001; Dubrulle and Pourquie, 2004). Second, a tail bud, also known to express FGF8 (Dubrulle and Pourquie, 2004), was found to induce ectopic deviation of the WD (Fig. 2O,P). Together, these observations suggest that WD deviates due to the FGF8 signaling, which is known to be expressed in the PSM. Therefore, we hypothesized that FGF signaling is activated in the elongating WD.
when placed into the lateral plate (Fig. 20.P; n=5/7). Third, FGF signals are known to be essential for tubular branching of trachea in *Drosophila* (Sutherland et al., 1996) and lung and mammary glands in mammals (Andrew and Ewald, 2010; Lu and Werb, 2008).

We found that mRNAs of FGF receptor 1 (FGFR1), FGFR2 (supplementary material Fig. S2A-D) and FGFR3 (Fig. 3C,D) were localized in the elongating WD. In addition, as previously reported (Dubrulle et al., 2001; Dubrulle and Pourquié, 2004), FGF8 was markedly expressed in the posterior region of the body, and this expression shifted posteriorly as the body axis elongated. Importantly, this *FGF8*-positive area included not only PSM, but also IMM (Fig. 3E-H). Thus, the posterior shift of the *FGF8*-positive zone was in register with WD elongation (Fig. 3P).

To determine whether FGF signaling was activated in the WD leader, we examined the expression patterns of *PEA3* and *SEF*, which are downstream factors of FGF signaling. We found that they were expressed in WD at HH10 (Fig. 3i; supplementary material Fig. S2E) (Lunn et al., 2007), and also at HH13 (Fig. 3j; supplementary material Fig. S2F). We also detected phosphorylation of ERK1/2 (pERK), a prominent marker of FGF signaling, in the elongating WD (Fig. 3k,l). Notably, pERK signals were more intense in the leader cells than in rear cells (Fig. 3m-o). These observations suggested that the leader cells, but not the rear cells, receive FGF ligands, and also raised the possibility that the posteriorly shifting *FGF8* would direct WD elongation by activating the leader cells (Fig. 3p).

**FGF8 is sufficient to attract leader cells of the WD**

To determine whether *FGF8* would act as a chemoattractant for WD cells, we implanted an aggregate of *FGF8*-producing DF1 cells (a chicken fibroblast cell line) into LPM. This induced ectopic deviation of the WD toward the implant (Fig. 4a-c; n=38/56). Similar manipulations using EGFP-transfected DF1 cells as a control did not affect WD elongation (Fig. 4d,e; supplementary material Fig. S3e). Ectopic deviation was also observed when FGF8-soaked beads were placed in a similar way (supplementary material Fig. S3f). In these embryos, implants (cell aggregates or beads) were placed in between the somatopleural (lateral) mesoderm and surface ectoderm (Fig. 4f-h).

Since FGFs are known to promote cell proliferation, we asked if the ectopic deviation of WD that we observed was achieved by cell proliferation or cell migration. *In vivo* time-lapse analyses spanning 180 min revealed that a majority of WD cells migrated toward the implant of FGF8/DF1 without significant cell division (Fig. 4j; supplementary material Movies 4 and 5). Thus, the FGF8-induced deviation of the WD is achieved by chemoattractive activity with little contribution from cell proliferation.

We further tested whether the WD migration was mediated by a direct effect by FGF8. We conducted *in vitro* time-lapse imaging analyses in which leader cells were dissected from E2/HH13 embryos and co-cultured with FGF/DF1 cells (Fig. 4k). As shown in supplementary material Movie 6, WD cells migrated toward FGF/DF1 cells, which were also motile. Similarly, WD cells were attracted to an FGF8-soaked bead (Fig. 4l; supplementary material Movie 7). By contrast, there was no directional migration toward a control PBS-soaked bead, although the cells were motile (Fig. 4m; supplementary material Movie 8). Furthermore, the FGF8-induced attraction was inhibited in the presence of SU5402, an inhibitor of FGF signaling (Fig. 4n; supplementary material Movie 9). Together, we concluded that FGF8 acts as a chemoattractant for WD leader cells.

We also examined the effects of FGF4 and FGF10, transcripts of which were found in the tail bud and PSM, respectively (supplementary material Fig. S3a-d) (Karabagli et al., 2002). A bead soaked with FGF4 protein and FGF10-producing DF1 cells were implanted in between the surface ectoderm and somatopleure (in a similar manner to Fig. 4a and supplementary material...
Fig. S3F). Whereas FGF4-soaked beads induced deviation of the WD leader cells, no effect was seen with FGF10-producing cells (supplementary material Fig. S3G,H). These observations raise the possibility that FGF8 and FGF4 may act redundantly, which would also account for the previously reported phenomenon that WD elongation is unaffected in mice lacking FGF8 (Perantoni et al., 2005).

Although glial cell line-derived neurotrophic factor (GDNF) has been shown to direct WD cells in axolotl (Drawbridge et al., 2000), we found no effects in chick of GDNF/cells implanted near the WD (supplementary material Fig. S3I). This is consistent with previous reports for mice showing that GDNF is not involved in early WD elongation (Chi et al., 2009; Chia et al., 2011).

FGF signals in the elongating WD through a Ras/ERK cascade

To determine whether FGF signals are essential for WD elongation, an SU5402-soaked bead was implanted into the lateral region. This treatment markedly disturbed WD elongation, whereas the control bead soaked with DMSO had no effect (Fig. 5A,B; \( n = 22/30 \) for SU5402, \( n = 0/30 \) for control). The expression of LIM1, an early marker for WD differentiation (Kobayashi et al., 2004; Tsang et al., 2000), was unaffected by SU5402 (supplementary material Fig. S3I). This is consistent with previous reports for mice showing that GDNF is not involved in early WD elongation (Chi et al., 2009; Chia et al., 2011).

FGF-deprived WD cells undergo epithelialization to form a tubular structure

We observed that SU5402-treated WD leader cells not only cease migration, but also undergo precocious epithelialization to form a tubular structure. An elongating WD, electroporated with GAP-EGFP (plasma membrane) and H2B-mCherry (nucleus), was treated with an SU5402-soaked bead (Fig. 6A-C). SU5402 abolished this attraction activity (N). Scale bars: 100 μm.
staining as a marker for basement membrane, which showed that SU5402-treated leader cells were apicobasally polarized (Fig. 6D,E). Thus, FGFs coordinate leader migration with tubular epithelialization during WD formation by acting as a molecular switch: in the presence of FGFs, WD cells are endowed with migratory and motile properties, whereas when deprived of FGFs they undergo tubular epithelialization.

Epithelialized cells convert to migratory cells when placed in the leader region

As shown above, rear WD cells (tubular epithelium) can be converted to leader-like migratory cells when relocated to the leader region (Fig. 2G) or provided with a leader environment (Fig. 2M). To further examine the potential of tubular-to-migratory conversion, we exploited SU5402-treated leader cells, which underwent precocious epithelialization, since these cells are amenable to more detailed analyses than is possible with rear cells. We carried out time-lapse imaging of these cells (GAP-EGFP electroporated) when transplanted into either the leader or rear positions of elongating WD (Fig. 6F). When placed into the leader, SU5402-treated cells resumed migration with mesenchymal character (Fig. 6H,H′; supplementary material Movie 10). These cells also exhibited intense pERK signal (Fig. 6K,L). By contrast, when transplanted into the rear region, the cells retained their epithelial morphology with little motility (Fig. 6G,G′; supplementary material Movie 11), and exhibited little pERK signal (Fig. 6I,J).

Together, the FGF signal acts as a binary switch between migratory leader cells and quiescent epithelial tubular cells. Thus, the progressive posterior shift of FGF8 expression coordinates leader elongation with tubule formation during WD formation.

DISCUSSION

By exploiting the simple structure of the WD and the amenability of tissue manipulations in chicken embryos, we have demonstrated that the WD is formed through a remarkably coordinated morphogenesis of tubule elongation and cell epithelialization. This coordination is mediated by a graded activation of FGF8 signals: posteriorly high signals chemoattract the leader cells, whereas anteriorly low signals enhance the tubular epithelialization of rear cells (Fig. 7). The posterior-to-anterior gradient of FGF8 is initiated by FGF8 transcription in the tail bud, and, as the body axis elongates posteriorly, the FGF8-positive zone is likewise progressively...
We have demonstrated that FGF8 acts as a chemoattractant and FGF8 signaling in migrating WD cells.

FGF8 signaling in migrating WD cells

We have demonstrated that FGF8 acts as a chemoattractant and induces directional migration of WD cells both in embryos and in primary cultured WD cells in vitro (Fig. 4; supplementary material Movies 6 and 7). Notably, it seems that leader cells attracted to FGF8 are not fully determined, and other cells can become leader cells (supplementary material Movie 6; Y. A., unpublished). These phenomena are reminiscent of other leader/tip cells reported for blood vessel sprouting (Jakobsson et al., 2010) and neural crest migration (Wynn et al., 2013). Furthermore, epithelialized rear cells in the WD can be converted to motile leader cells when relocated into the leader environment without any substantial effects on cell proliferation, and this conversion is likely to be mediated by FGF8. Thus, the newly epithelialized tubular structure is naïve and susceptible to an FGF8-mediated EMT action. A few examples of the chemoattractive activities of FGFs have been shown previously, including cell migration along the primitive streak controlled by FGF4 (not by FGF8) (Yang et al., 2002) and neurite extension of motor neurons by FGF8 (Shirasaki et al., 2006).

It is likely that the FGF8 signaling is transmitted via FGFR3, transcripts of which are detected in the elongating WD. This might also account for the selective attraction of the leader cells to FGF8, as FGF8 binds to FGFR3 with higher specificity than to FGFR1 and FGFR2 (Zhang et al., 2006). Furthermore, it is likely that FGF4 acts redundantly with FGF8 during WD elongation, consistent with mouse genetic studies showing cooperative actions between FGF4 and FGF8 in limb and body axial outgrowth (Boulet and Capecchi, 2012; Boulet et al., 2004; Naiche et al., 2011; Sun et al., 2002). Intracelularly, the FGF8 signal during WD elongation is transmitted via the Ras/ERK pathway, whereas PLCγ and the PI3K pathways are dispensable (Fig. 5). Of note, sprouty 1 appears to be expressed in chicken WD (Geisha, http://geisha.arizona.edu/geisha/) (Antin et al., 2014). Since sprouty 1 is known to inhibit the PLCγ but not Ras/ERK pathway during mesodermal specification in frogs (Nutt et al., 2001; Sivak et al., 2005), similar inhibition of PLCγ might occur during WD elongation in chickens.

Recently, Attia et al. reported that FGF signals are crucial for WD elongation (Attia et al., 2015). However, they did not detect attractive activity by FGF8. The reason for the discrepancy between their results and ours is not known, although several possibilities are conceivable. The first relates to how the activities of FGF8 protein and drug inhibitors were validated: for the FGF8 transfection into cultured cells, we used the bidirectional promoter (pBI) (Watanabe et al., 2007), which drives EGFP and FGF8 cDNAs simultaneously, allowing us to confirm high transfection efficiency by EGFP signal. The second possibility relates to how FGF8-producing cells and drug-soaked beads were implanted into embryos: in our experience, implants should be placed precisely in between the surface ectoderm and somatopleural mesoderm (as shown in Fig. 4E-H). If embedded too deep in the coelomic cavity or splanchnopleura, the implant would not influence WD elongation.

Coordination between tissue morphogenesis and body growth

In this study, the FGF gradient has been shown to coordinate the leader and rear cells during WD elongation. This gradient is initiated by FGF8 produced in the tail bud, which is essential for the body elongation (Bénazéraf et al., 2010; Dubrulle and Pourquie, 2004). Thus, the body growth of the embryo coordinates the WD elongation, and this is mediated by FGF8.

In addition to the coordinated elongations between the body axis and WD, it is also known that the FGF8 gradient regulates the progression of somite segmentation (Dubrulle et al., 2001). Thus, the growth of the embryo body coordinates at least two types of morphogenesis—segmentation and WD elongation—in which FGF8 operates as a master coordinator. The coordination within this trinity can also explain the phenomenon at the molecular level that the WD leader is always located at sm –1 to –2, with the relative positions unchanged during body elongation. Importantly, our findings have revealed the coordination not only between multiple tissues, but also...
between different morphogenetic cells within the WD. It is remarkable that a macroscopic change, such as body axis elongation, is coordinately translated to morphogenetic changes at the microscopic level, such as the chemoeattraction of leader cells and tubule formation of rear cells, both of which progress posteriorly with a regular interval between them. Of note, WD formation, in turn, appears to regulate Müllerian duct elongation, which also involves coordination between exploring front cells and epithelializing tubular cells (Orvis and Behringer, 2007). Along this line, an elegant study on mouse lung development should be mentioned in which branching morphogenesis antagonizes alveolar differentiation, so that both events progress in a coordinated manner as the developing lung grows (Cunningham et al., 2013), and it awaits to be seen whether such morphogenetic coordination within the lung would also be controlled by body growth. From this viewpoint, it should be highlighted that diseases seen at the microscopic level (i.e. cellular dysfunction) might link to a macroscopic malformation (i.e. body growth), providing a new avenue for therapeutic treatments.

MATERIALS AND METHODS

Experimental animals
Fertilized chicken eggs were obtained from Shiroyama-Keien (Sagamihara, Japan). Embryos were staged according to Hamburger and Hamilton (1992) (HH) or somite number. All animal experiments were conducted with the ethical approval of Kyoto University (No. H2620).

PKH labeling
The presumptive pronephric region of 10-somite embryos (10 sm) was labeled with PKH26 or PKH67 fluorescent cell linker (Sigma) as previously reported (Atsuta et al., 2013).

Expression vectors
pCAGGS-GAP43-EGFP and pCAGGS-tTA were described previously (Atsuta et al., 2013). For pCAGGS-tdTomato, tdTomato was subcloned into the EcoRI-BglII sites of the pCAGGS plasmid (Niwa et al., 1991). For pCAGGS-GAP-P2A, GAP43-EGFP and oligonucleotides encoding P2A (Kim et al., 2011) were subcloned into the MluI-Xhol and Xhol-EcoRI sites, respectively, of the pCAGGS plasmid. To obtain pCAGGS-GAP-EGFP, 2A-H2B-mCherry, H2B-mCherry was ligated into the RV site of pBI-TRE-EGFP. cDNAs for chicken FGF8 and FGF10 were gifts of Drs H. Nakamura (Tohoku University) and H. Ohuchi (Okayama University), respectively. Each was subcloned into the Midt-Nhel sites of pB1-TRE-EGFP (Watanabe et al., 2007) to obtain pB1-TRE-FGFR8-EGFP and pB1-TRE-FGFR10-EGFP. cDNA for chicken GFN4 was isolated by PCR using primers (5′-3′): GFN4-F, ATGAGGTGACCTGACATGGTCAAGGCT; GFN4-R, ATAGCTGTCGACGACATCCACCA. An amplified fragment was subcloned into the Midt-Nhel sites of pB1-TRE-EGFP to obtain pB1-TRE-GDNF-EGFP.

In ovo electroporation
In ovo DNA electroporation into WD cells was performed using CUY21 EX (BEX) as described (Atsuta et al., 2013).

DNA transfection and preparation of DF1 cell aggregate
Chicken fibroblast-derived DF1 cells were maintained at 39°C with Dulbecco’s modified Eagle’s medium (Nissui) containing 10% fetal bovine serum (FBS/DMEM) and Penicillin-Streptomycin (Gibco). pB1-TRE-EGFP, pB1-TRE-FGFR8-EGFP, pB1-TRE-FGFR10-EGFP or pB1-TRE-GDNF-EGFP was co-transfected with pCAGGS-tTA into DF1 cells using Lipofectamine 2000 (Invitrogen) as previously described (Saito et al., 2012). Twenty-four hours after transfection, cells were transferred to a dish coated with 1% agarose to obtain cell aggregates (Tonogawa et al., 1997).

Embryological manipulation
Surgical manipulations were performed with E2/HH13 embryos (about 20 sm). Transplantation of a donor tissue, cell aggregate or bead into a host embryo was performed using a tungsten needle. FGFs were delivered using heparin acrylic beads (Sigma) soaked overnight with 250 μg/ml human FGF4 (WAKO) or 250 μg/ml human FGF8 (WAKO) at 4°C. For chemical-soaked bead experiments, AG1X2 ion-exchange beads (formate form; BioRad) were incubated in 10 mM SU5402 (Calbiochem), 20 mM PD184352 (Sigma), 20 mM LY294002 (Sigma), 10 mM U73122 (Sigma), 20 mM SP600125 (Sigma) or 20 mM SB203580 (Sigma) for 2 h at room temperature before grafting.

Immunohistochemistry
For immunohistological staining on histological sections, anti-laminin mouse monoclonal antibody (3H11; DSHB), anti-E-cadherin mouse monoclonal antibody (BD Biosciences, 610182) and anti-ZO-1 rabbit monoclonal antibody (Zymed, 40-2200) were used as described (Atsuta et al., 2013). Sections treated with antibodies were also exposed to DAPI (Sigma).

To detect diphosphorylated ERK (pERK), sections were treated with 3% H2O2 in TNT (0.1 M Tris-HCl, 0.1 M NaCl, 0.1% Tween 20, pH 7.5) for 30 min, and pre-blocked with 1% Blocking Reagent (Roche) in TNT for 1 h. The sections were incubated overnight at 4°C with 1:300 anti-pERK rabbit polyclonal antibody (Cell Signaling, 4370). They were then washed in TNT and incubated with 1:1000 anti-rabbit IgG horseradish peroxidase (HRP)-conjugated donkey antibody (GE HealthCare, NA9340-1ML) for 30 min. After washing in TNT, sections were reacted with the TSA Plus Cy3 system (PerkinElmer) for 5 min at room temperature. The reaction was terminated by washing in TNT, and sealed with FluorSave Reagent (Calbiochem) containing DAPI. Fluorescent images were obtained using an Axioplan 2 microscope with Apotome system (Carl Zeiss).

For whole-mount detection of pERK, embryos were fixed with 4% paraformaldehyde (PFA) in PBS overnight at 4°C. Fixed embryos were dehydrated with methanol and treated with 3% H2O2 in methanol for 4 h. After rehydration with PBS containing 0.1% Tween 20 (PBST), embryos were washed in PBS containing 0.5% Tween 20. After fixation with 5% FBS in PBST, they were incubated with 1:200 anti-pERK antibody for 48 h at 4°C. After washing in TBST (0.1 M Tris-HCl, 0.15 M NaCl, 0.5% Triton X-100, pH 7.5), they were incubated with 1:200 anti-rabbit IgG HRP-conjugated donkey antibody.

For DAB (3,3′-diaminobenzidine; WAKO) staining, specimens were washed with TBST and incubated with DAB/PBST for 20 min, then reacted with a 1:2000 dilution of 30% H2O2 for 5 min. For double staining of LIM1 protein and pERK, 1:200 anti-LIM1 mouse monoclonal antibody (4F2; DSHB) was used along with the anti-pERK antibody.

Probes and in situ hybridization
Chicken cDNA fragments for FGFR2 and FGFR3 were isolated by PCR using primers (5′-3′): FGFR2-F, TTTACGCGTGAGAGCGTAGTCCCA- TCCGA; FGFR2-R, TGAGCTAGCTCATTTTGGATCCTCTGGCAC- TAGC; FGFR3-F, ATAGATATCATGTTCAAAACG- ATTTTTC; FGFR3-R, ATAGCTGTCGACGACATCCACCA. An amplified fragment was subcloned into the Midt-Nhel sites of pB1-TRE-EGFP to obtain pB1-TRE-GDNF-EGFP.

Whole-embryo culture and time-lapse imaging
Whole-embryo culture was performed as described (Atsuta et al., 2013). For cell tracking analyses, images were processed with the Manual Tracking Tool of ImageJ (NIH).

In vitro migration assay
WD cells were dissected using tungsten needles and placed with FGF8-soaked beads or FGF8-expressing cells in a 35-mm glass-bottom dish (MATSUNAMI) coated with 10 ng/ml fibronectin (WAKO). The specimens were embedded in Matrigel (BD Biosciences) diluted with Opti-MEM (Invitrogen), and cultured for 6 h with FBS/DMEM.
TUNEL assay
The TUNEL assay was performed with the In Situ Cell Death Detection Kit (Roche) according to the manufacturer’s instructions.

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Competing interests
The authors declare no competing or financial interests.

Author contributions
Y.A. and Y.T. designed the study, analyzed the data and wrote the manuscript. Y.A. performed the experiments.

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Supplementary material
Supplementary material available online at http://dev.biologists.org/lookup/suppl/doi:10.1242/dev.122408/-/DC1

References
**Figure S1.** Signals from the surface ectoderm are dispensable for WD elongation. (A) Schematic diagrams showing that the surface ectoderm overlying leader cells was removed from an E2/HH13 embryo. (B, C) Caudal migration of WD on the treated side was comparable to the untreated side (black arrowheads). (D, E) A piece of aluminum foil was inserted in between the WD and surface ectoderm, resulting in no disturbed elongation of WD (black arrowheads).
Figure S2. Expression patterns of FGFR1, FGFR2 and SEF. Whole-mount in situ hybridization to show transcripts of FGFR1 (A, B), FGFR2 (C, D) and SEF (E, F) in E1.5/HH10 and E2/HH13 embryos. FGFR1 and SEF were expressed in the WD at both stages, and FGFR2 was expressed in E2/HH13 embryo (white arrowheads).
Figure S3. FGF8, FGF4, but not FGF10, attracted leader cells. Whole-mount in situ hybridization to show expression patterns of *FGF4* (A, B) and *FGF10* (C, D) in E1.5/HH10 and E2/HH13 embryos. An implanted FGF8-soaked or FGF4-soaked bead attracted leader cells ([F, G], arrows) whereas EGFP-expressing cells yielded no effect (E). (H, I) Neither FGF10-producing nor Glial-derived growth factor (GDNF)-producing DF1 cells showed attracting activity.
Figure S4. Inhibition of FGF signaling did not affect *LIM1* expression or cell survival. (A, B) Expression of *LIM1* mRNA was not downregulated by SU5402 treatment (arrowheads). (C-F) SU5402 did not excessively induce cell deaths. (D, E) Transverse sections indicated in (C) were double-stained with antibody for Laminin1 (red) and TUNEL (green). (F) Quantification of apoptotic cells within DMSO-treated (n = 63 cells in 8 embryos) or SU5402-treated leader cells (n = 121 cells in 8 embryos). Error bars represent SEM. Scale bars: 20 μm for (D).
Movie 1. Time-lapse analyses using a cultured whole embryo, the WD of which was labeled with a red fluorescent dye PKH26. Frames were taken every 3 min with a 10 × Plan-Apochromat objective lens. Total movie length: 270 min.
Movies 2, 3. In vivo time-lapse analyses for rear cells (Movie 2) and leader cells of WD (Movie 3), which were electroporated with pCAGGS-GAP-EGFP-P2A-H2B-mCherry. Frames were taken every 1 min with a 40× Plan-Achromat objective lens. Total movie length: 30 min.
**Movies 4, 5.** Time-lapse analyses using cultured whole embryos where pCAGGS-tdTomato (red) was electroporated into WD. When FGF8-producing DF1 cells (green) were implanted, leader cells migrated toward the FGF8/DF1 (Movie 4). 10 cells in each movie were traced by digitally processed colors. Movie 5 is of a normal embryo. Frames were taken every 3 min with a 10× Plan-Achromat objective lens. Total movie length: 180 min.
**Movie 6.** *In vitro* live-imaging and chemoattraction assay using leader cells (red) taken from PKH26-labeled WD co-cultured with FGF8-EGFP-producing DF1 cells (green). Frames were taken every 5 min with a 20× Plan-Achromat objective lens. The length of each movie: 300 min.
Movie 7-9. *In vitro* chemoattraction assay using leader cells co-cultured in Matrigel with an FGF8-soaked bead (Movie 7), DMSO-soaked bead (Movie 8) and FGF8-soaked bead in the presence of 2μM SU5402 (Movie 9). pCAGGS-tdTomato (red) was electroporated into WD to visualize individual cells prior to the culture. Frames were taken every 5 min with a 20× Plan-Achromat objective lens. The length of each movie: 360 min.
Movies 10, 11. Time-lapse analyses showing that forced-epithelialized cells by SU5402 could be transformed into motile cells when transplanted into the leader region (Movie 10), whereas cells transplanted into the rear region remained static (Movie 11). Frames were taken every 1 min with a 20× Plan-Achromat objective lens. The length of each movie: 60 min.