Notch signaling controls the differentiation of transporting epithelia and multiciliated cells in the zebrafish pronephros

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Epithelial tubules consist of multiple cell types that are specialized for specific aspects of organ function. In the zebrafish pronephros, multiciliated cells (MCCs) are specialized for fluid propulsion, whereas transporting epithelial cells recover filtered-blood solutes. These cell types are distributed in a ‘salt-and-pepper’ fashion in the pronephros, suggesting that a lateral inhibition mechanism may play a role in their differentiation. We find that the Notch ligand Jagged 2 is expressed in MCCs and that notch3 is expressed in pronephric epithelial cells. Morpholino knockdown of either jagged 2 or notch3, or mutation in mind bomb (in which Notch signaling is impaired), dramatically expands ciliogenic gene expression, whereas ion transporter expression is lost, indicating that pronephric cells are transfigured to MCCs. Conversely, ectopic expression of the Notch1a intracellular domain represses MCC differentiation. Gamma-secretase inhibition using DAPT demonstrated a requirement for Notch signaling early in pronephric development, before the pattern of MCC differentiation is apparent. Strikingly, we find that jagged 2 knockdown generates extra cilia and is sufficient to rescue the kidney cilia mutant double bubble. Our results indicate that Jagged 2/Notch signaling modulates the number of multiciliated versus transporting epithelial cells in the pronephros by way of a genetic pathway involving repression of rfx2, a key transcriptional regulator of the ciliogenesis program.

KEY WORDS: Pronephros, Jagged 2, Multiciliated cell, Notch3, double bubble, mind bomb, Zebrafish

INTRODUCTION

The generation of cell diversity during organ development is a crucial event in defining the proportion of specialized cell types that coordinate organ function. How epithelial organ primordia are patterned to give rise to specialized epithelial cells is largely unknown. In the kidney, the nephron is the functional unit of organ physiology. The mature nephron is populated by multiple different cell types that are specialized for blood filtration, and water and ion homeostasis (Seldin and Giebisch, 1992; Vize et al., 2002). Functionally similar cell types in the nephron are typically arranged in discrete tubule segments that process filtered blood sequentially as it passes through the nephron. In other parts of the nephron, functionally distinct cells are intermingled or ‘intercalated’ between non-like cells. For instance, in the collecting system, intercalated cells that regulate pH homeostasis are morphologically and functionally distinct from adjacent principal cells, whose primary function is water transport (Schuster, 1993; Seldin and Giebisch, 1992). Similar intermingling of functionally distinct cell types is found in other epithelial organs; for example, in the lung, intestine and stomach. In the lung, neuroendocrine cells, which have specialized sensory and secretory functions, are dispersed among airway epithelial cells (Ito et al., 2000). In the intestinal epithelium, mucous-secreting goblet cells are interspersed among vilius epithelial cells (Zecchini et al., 2005). In the stomach of the chick, glandular epithelial cells differentiate from an initially homogeneous layer of epithelium and lie interspersed with lumenal epithelial cells (Matsuda et al., 2005). The existence of adjacent non-like cells in these organs suggests that a lateral inhibition mechanism, similar to well-defined mechanisms underlying the development of proneural clusters in Drosophila (Artavanis-Tsakonas et al., 1995), may play a role in vertebrate organogenesis. In peripheral neural development in Drosophila, clusters of cells destined to become nerves and supporting cells are partitioned by the expression of the Notch ligand Delta in future nerve cells. Delta, acting on Notch receptors in adjacent cells, prevents them from adopting a neural fate (Artavanis-Tsakonas et al., 1995). Using loss- and gain-of-function approaches targeting various elements of vertebrate Notch signaling pathways, lateral inhibition mechanisms have in fact been demonstrated to control the differentiation of neuroendocrine cells in the lung (Ito et al., 2000), goblet cells in the intestine (Crosnier et al., 2005; Zecchini et al., 2005) and gland cells in the stomach (Matsuda et al., 2005).

In Notch-mediated lateral inhibition, a specialized cell type expresses a transmembrane DSL (Delta/Serrate/Lag-2) ligand of Notch which binds a Notch receptor in neighboring cells (Kadesch, 2004). The Notch receptor undergoes regulated intramembrane cleavage to generate an active intracellular domain peptide that transits to the nucleus to participate in transcriptional regulation (Mumm and Kopan, 2000). By binding to and activating the CSL [CBF-1, Suppressor of Hairless (Rbpsuh), Lag-1] transcriptional complex, the Notch intracellular domain (NICD) initiates a transcriptional cascade involving the expression of Hes and HRT/HER/Hey transcriptional repressors (Kadesch, 2004). These, in turn, repress genes that are expressed in Notch-ligand-expressing cells, often including the ligand gene itself, and thereby direct the neighboring cell along a different developmental pathway (Kadesch, 2004). In kidney organogenesis, Notch signaling has been implicated in the development of glomerular vasculature and in segmentation of the nephron (Cheng et al., 2003; McCright, 2003). Expression of the Notch ligand Jagged 1 in endothelial cells and Notch2 in glomerular epithelial cells is required for proper development of the glomerular capillary tuft (McCright et al., 2001). Broadly activating or inhibiting Notch signaling by the expression of activated or dominant-negative Suppressor of hairless proteins have been shown to perturb proximal versus distal nephron fate in Xenopus embryos (McLaughlin et al., 2000). Mouse kidney explant...
cultures treated with the gamma-secretase inhibitor DAPT to inhibit Notch cleavage also show a loss of proximal versus distal nephron segments (Cheng et al., 2003). These experiments demonstrate a role for Notch signaling in the specification of relatively large nephron domains. However, there is currently no evidence implicating Notch signaling in the generation of 'salt-and-pepper' patterns of interdigitating cell types within the kidney nephron.

The zebrafish pronephros is an accessible model for nephron cell specification and patterning because it consists of typical vertebrate kidney cell types, and is amenable to both forward and reverse genetic manipulation (Drummond, 2003). Expression of jagged and notch genes in the pronephros suggests that they could play an important role in patterning the pronephric nephron (Zecchin et al., 2005). Evidence for nephron patterning in the pronephros is seen in the segment-specific expression of ion transporters (Nichane et al., 2006; Shmukler et al., 2005), in the expression of receptors and transcription factors in discrete segments (Bisgrove et al., 1997; Marcos-Gutierrez et al., 1997; Van Campenhout et al., 2006), and in the disruption of segment-boundary formation in the pax2.1 (also known as pax2a – Zebrafish Information Network) mutant no isthmus (Majumdar et al., 2000). In our examination of cilia function in the kidney (Kramer-Zucker et al., 2005) we have found that two types of ciliated cells exist in the pronephros: single ciliated cells that have the morphology of typical transporting epithelial cells, and multiciliated cells that, by several criteria, appear specialized for fluid propulsion in the pronephros. We show here that multiciliated cells are isolated, specialized cells surrounded by non-like transporting epithelia in the distal segment of the pronephros. We present evidence that Notch signaling via Jagged 2-Notch interactions, as well as the downstream transcriptional regulation of ciliogenic genes, is essential in the generation of the 'salt-and-pepper' pattern of multiciliated and transporting epithelial cells in the pronephros. We also demonstrate that transfecting cells to the MCC lineage is sufficient to overcome defects in ciliogenesis in a zebrafish cyst mutant and prevent the formation of pronephric cysts.

MATERIALS AND METHODS
Zebrafish lines
Wild-type TL or TUAB, double bubble (m468) and mind bomb (m132) zebrafish were maintained and raised as described (Westerfield, 1995). Dechorionated embryos were kept at 28.5°C in E3 solution with or without 0.003% PTU (1-phenyl-2-thiourea, Sigma) to suppress pigmentation and were staged according to somite number (som) or hours post-fertilization (hpf) (Westerfield, 1995). The Na,K-ATPase alpha1A4:GFP transgenic line was created by amplifying an 8112 base-pair fragment of the Na,K-ATPase alpha1A4-subunit promoter (cDNA accession: NM_131689) from genomic DNA using nested PCR with the following primers: LNaK-F1: 5'-GGTGGCCTGGGCTAAGTCACAGAC-3' 
LNaK-F2: 5'-GCGCTGAATGTGGGCCCCCTTA-3' 
LNaK-R: 5'-TCCACACTCTCAGGCCCCATT-3' 
NaKB-R2: 5'-TCGGGTAATCAGGCCCCATT-3' 

The 8112 bp fragment was cloned in PG1 upstream of EGFP and injected as a circular plasmid into TuAB wild-type embryos at the 1-cell stage. Founders were screened by GFP fluorescence in the pronephros and were out crossed to create a stable transgenic line. For notch1a founders were screened by GFP fluorescence in the pronephros and were crossed to create a stable transgenic line. For notch1a founders were screened by GFP fluorescence in the pronephros and were crossed to create a stable transgenic line. For notch1a founders were screened by GFP fluorescence in the pronephros and were crossed to create a stable transgenic line.

In situ hybridization
Whole-mount in situ hybridization was performed as previously described (Thiess and Thiess, 1998). For two-color in situ hybridization, probes were synthesized using DIG-UTP or FITC-UTP. Stained embryos were cleared in Benzyl alcohol and photographed on a Leica MZ12 or Nikon E800 microscope equipped with a Spot Image digital camera. Two-color stained embryos were cleared in glycerol and photographed after removal of the yolk extension. Two-color fluorescent in situ hybridization (S. Holley, Yale University, New Haven, CT, personal communication) was performed using DIG-UTP and FITC-UTP-labeled RNA probes followed by sequential detection using HRP labeled anti-DIG and HRP-labeled anti-FITC antibodies (Boehringer Mannheim) and fluorescent tyramide amplification. Following RNA-probe hybridization, embryos were blocked in 150 mM maleic acid, 100 mM NaCl pH 7.5 with 2% BSA and then incubated with HRP anti-FITC at 1:500 dilution for 4 hours in blocking solution. After washing, anti-FITC antibody was visualized with TSA Plus Fluorescein Solution (Perkin Elmer) for 45 minutes, dehydrated in methanol and quenched with 1% H2O2 in methanol for 30 minutes. After rehydrating, embryos were blocked again and incubated with HRP anti-DIG at 1:1000 dilution for 4 hours, then washed and reacted with TSA Plus Cy3 solution (Perkin Elmer) for 45 minutes. For combined fluorescent shippedo1 DIG in situ anti-acetylated tubulin staining, embryos were incubated with both anti-acetylated tubulin (clone 6-1B1-1; Sigma) and HRP anti-DIG (Boehringer Mannheim) after hybridization. After complete fluorescent in situ hybridization, embryos were refixed with 4% PFA, and were then washed and blocked in 150 mM maleic acid, 100 mM NaCl pH 7.5 with 10% NGS for 2 hours at room temperature. Acetylated tubulin was visualized with Alexa Fluor 546 goat anti-mouse IgG (Molecular Probes) in blocking solution. Embryos were cleared in glycerol, mounted and viewed on a Zeiss LSM5 PASCAL confocal fluorescence microscope.

Morpholino antisense oligonucleotides
Morpholino antisense oligonucleotides (Gene Tools, Philomath, OR) were designed to target either the translation start or an exon splice-donor site causing splicing defects of the mRNA. The following morpholinos were used:
j2exon 20: 5'-TCTTTGAGATACCTACTGATAGCCG-3' 
j2exon 20 control: 5'-CGCGCATAGCTCATCAGAGTTCT-3' 
r2exon 6: 5'-GGGTTAGTCGACCCTGTT-3' 
r2exon 6 control: 5'-CATGTGCTAGCTGATG-3' 
notch3exon 27: 5'-TGACCAACTCTCCATGCCCAGTG-3' 
jagged 2: 5'-TCTGGTACATTGCGCTGC-3' 
jagged 1b: 5'-TCTGGTACATTGCGCTGC-3'

Morpholino oligos were diluted in 100 mM KCl, 10 mM HEPES, 0.1% phenol red (Sigma) and injections were performed using a nanoliter2000 microinjection (World Precision Instruments). Injection concentrations were 0.2-0.5 mM and injection volume was 4.6 nl/embryo (7-16 ng morpholino/embryo). The molecular defect caused by splice-donor morpholinos was verified by reverse-transcriptase PCR from total RNA from a single morphant embryo with nested primers in flanking exons.

Histochemistry and immunohistochemistry
For histology, embryos were fixed with 4% formaldehyde in PBS at 4°C overnight followed by dehydration and embedding in JB-4 (Polysciences), and then sectioned at 5-7 μm. Slides were stained in methylene blue/azure II (Humphrey and Pritman, 1974) and examined using a Nikon E800 microscope. For acetylated tubulin staining, the embryos were fixed in Dent’s Fix (80% methanol/20% DMSO) at 4°C overnight. After rehydration they were washed in PBS plus 0.5% Tween20 and blocked in PBS-DTB (PBS plus 1% DMSO, 1% BSA and 0.5% Tween20) with 10% stringency.
MULTICILIALED AND TRANSPORTING EPITHELIAL CELLS ARE DISTINCT, SPECIALIZED CELL TYPES IN THE PRONEPHROS

We have shown previously that motile cilia in the zebrafish pronephros are required for the normal high rate of fluid output from this organ and that defects in cilia function are sufficient to cause cyst formation in the proximal pronephric tubules (Kramer-Zucker et al., 2005). More-detailed examination of ciliated cells in the pronephros showed that the majority of cells possessed a single cilium, while a subset of cells were multiciliated and extended up to 20 cilia from their apical surface into the pronephric lumen (Fig. 1). Acetylated tubulin immunofluorescence of whole-mount zebrafish embryos at 52 hpf revealed bright bundles of cilia in the pronephric lumen, which were interspersed among cells with a single cilium (Fig. 1A,B). Ultrastructurally, these bundles of cilia originated from individual cells characterized by multiple apical basal bodies and apically oriented mitochondria (Fig. 1C,F). By contrast, the surrounding single-ciliated cells possessed a microvillar brush border and basolaterally oriented mitochondria, a morphology typical of transporting epithelia. The distribution of multiciliated cells (MCCs) as isolated individual cells along the pronephros was also apparent in high-speed micro video images of the pronephros (see Movie 1 in the supplementary material). Bundles of cilia emanating from MCCs beating in a coordinated wave pattern, in

**Fig. 1. Distribution of multiciliated cells and transporting epithelia containing single cilia in the pronephros.** (A) Whole-mount immunofluorescence of the trunk region of a 48 hpf embryo stained with anti-acetylated tubulin reveals bright cilia bundles (arrowheads) in the pronephric lumens, as well as single ciliated cells (arrows) in a more caudal nephron segment. (B) At higher magnification, condensed bundles of cilia (arrowheads) and single cilia (arrows) can be observed in the pronephric lumen. (C) Bundles of cilia emanating from individual cells project into a distented lumen of a mechanically obstructed pronephros. Dotted lines in B and C outline the pronephric tubules. (D-F) Electron micrographs of the pronephros show isolated multiciliated cells (MCC; false-colored in red) interspersed among transporting epithelial cells (false-colored in light blue). Arrowheads show apical basal bodies. (D) MCCs are distinguished by multiple apical basal bodies, by the lack of a brush border (bb), by a small basal cell surface and by multiple apical cilia (asterisks). n, nucleus. (E) Example of an MCC with multiple apical cilia basal bodies, and bundles of cilia in the lumen. (F) Cross section of a pronephric tubule in a 7 dpf larva, showing a single MCC with multiple basal bodies, apical mitochondria (asterisks) and bundles of cilia fitting tightly into the pronephric lumen.
contrast to the single cilia on neighboring transporting epithelial cells, which appeared to beat autonomously (see Movie 1 in the supplementary material).

Consistent with the abundance of cilia on MCCs, we found that genes for axonemal structural proteins and regulators of cilia formation were highly expressed in a pattern similar to MCCs in the pronephros (Fig. 2). shippo1 (zgc:63985 – Zebrafish Information Network) encodes an axonemal protein originally isolated from male germ cells (Egydio de Carvalho et al., 2002) (Fig. 2A) and rfx2 belongs to the family of X-box-binding transcription factors, which act as master regulators of the ciliogenesis program (Bonnafe et al., 2004; Efimenko et al., 2005; Swoboda et al., 2000). rfx2 was initially expressed throughout the pronephros, implying a role for this transcription factor in the development of both single- and multiciliated cells (data not shown). By 34 hpf, however, rfx2 was concentrated in a spotted pattern similar to the distribution of multiciliated cells (Fig. 2B). shippo1- and rfx2-positive cells are present throughout the distal segment of the pronephric nephron (Nichane et al., 2006) but were not present in the most-caudal nephron segment (Fig. 2A,B). Because cells directly adjacent to MCCs appear by electron microscopy to be more typical of transporting epithelia (Fig. 1D-F), we examined ion channel gene expression in similar regions of the pronephros. Cells in the ‘early distal’ segment of the pronephric nephron expressed the cation transporter trpM7 (Fig. 2C) and the sodium-sulfate co-transporter slc13a1 (Fig. 2D). Whereas epithelial cells expressing these two channel genes appeared to coexist with shippo1- and rfx2-positive cells in the early distal segment, the distribution of shippo1- and rfx2-positive cells further extends distally into the ‘late distal’ segment of the pronephros (Nichane et al., 2006).

To confirm that shippo1/rfx2 and trpM7/slc13a1 marked non-equivalent, but adjacent, cell types in the pronephros, we performed two-color whole-mount in situ hybridization and double fluorescent in situ hybridization with these probes. Two-color whole-mount in situ hybridization verified that trpM7/slc13a1-expressing epithelial cells and shippo1-expressing cells are indeed two distinct cell types that exist side by side in the pronephros (Fig. 2E,F). To further distinguish MCCs from transporting epithelia, we created a transgenic line of zebrafish that expresses GFP from the Na,K-ATPase alpha A4 gene promoter. This Na,K-ATPase subunit is highly expressed in the pronephros, where, by analogy to mammalian kidney epithelia, it is likely to power a variety of ion-transport activities (Seldin and Giebisch, 1992). In the distal segment of the pronephros, the Na,K-ATPase::GFP transgene was expressed in a subset of cells (Fig. 2G) and, based on co-staining for cilia bundles with anti-acetylated tubulin, was excluded from MCCs (Fig. 2H,I). The complementary expression pattern of axonemal genes/cilia bundles with genes for ion transporters suggests that MCCs and transporting epithelia are distinctly different cell types, intermixed in the distal segment of the pronephros.

Using a combination of shippo1 fluorescent in situ hybridization and anti-acetylated tubulin immunofluorescence, we found that shippo1-positive cells were multiciliated (Fig. 3A,B). Neighboring cells with a single cilia were shippo1-negative (Fig. 3A,B). Using shippo1 as a reference probe for MCCs in double fluorescent in situ hybridization with ciliogenic or ion channel gene probes, we confirmed that other ciliogenic genes – including fleer (flr), a tetratricopeptide-repeat protein required for ciliogenesis (N.P. and I.A.D., unpublished) (Fig. 3C-E), and rfx2 (Fig. 3F-H) – were co-expressed with shippo1 in MCCs. Conversely, the ion channel genes for slc13a1 (Fig. 3I-K) and trpM7 (Fig. 3L-N) were expressed in a non-overlapping, adjacent set of cells in the pronephros.

The Notch ligand Jagged 2 is expressed in multiciliated cells

The apposition of different epithelial cell types in the pronephros suggested that a lateral inhibition mechanism, possibly involving Notch signaling, might play a role in pronephric cell patterning. Because the Notch receptor notch3 and the Notch ligands jagged 2 and jagged 1b (previously jagged 3) have recently been reported to be expressed in the pronephros (Lawson et al., 2001; Zecchin et al., 2005), we examined the expression of Notch signaling genes in relation to MCCs and transporting epithelial cells. Interestingly, jagged 2 was expressed in the pronephric mesoderm during early somitogenesis (Fig. 4A) and was later expressed in a subset of cells in the pronephros in a pattern similar to that of the MCC-marker gene expression (Fig. 4B). Double staining by jagged 2 in situ hybridization and anti-acetylated tubulin immunohistochemistry demonstrated that jagged 2-positive cells were indeed multiciliated (Fig. 4C). A bundle of cilia could be seen originating from a single jagged 2-expressing cell in a cross section of the pronephros (Fig. 4D). To confirm further the identity of jagged 2-expressing cells, we performed two-color and double fluorescent in situ hybridization...
with *jagged 2* and *shippo1* probes. By both methods, we found that *shippo1* was co-expressed with *jagged 2* in a distinct subset of cells (Fig. 4D,E-G), confirming that *jagged 2* is specifically expressed in MCCs.

**Jagged 2 represses MCC fate**

Expression of *jagged 2* mRNA in MCCs suggested that it could be required in some way for MCC cell differentiation or, by analogy to Notch signaling in other contexts, it could give rise to the intercalated pattern of MCCs by repressing the multiciliated cell fate in neighboring cells. To distinguish between these possibilities, we disrupted the expression of *jagged 2* with antisense morpholino oligos that targeted either the *jagged 2* ATG initiation codon (*J2atgMO*; protein knockdown) or the *jagged 2* exon 20 splice donor (*J2ex20MO*; mRNA mis-splicing). The *J2atgMO* has previously been shown to block *jagged 2* protein translation (Lorent et al., 2004) and the *J2ex20MO* was found to delete exon 20 and generate an in-frame stop codon immediately following the coding sequence of exon 19, with the resultant mRNA predicted to encode a truncated *jagged 2* extracellular domain peptide (see Fig. S1 in the supplementary material). Injection of either morpholino oligo gave rise to the same phenotype: the number of MCCs was dramatically expanded in the pronephros (Fig. 5A-I). Injection of a control invert sense morpholino had no effect on the expression pattern of the MCC markers *shippo1*, *flr* or *rfx2* (Fig. 5). Sections of whole-mount stained embryos confirmed that the single-cell pattern of MCC-marker expression (Fig. 5J) expanded to include all tubule cells in the section in *jagged 2* morphants (Fig. 5K,L). Interestingly, expression of *jagged 2* itself was also dramatically expanded in *jagged 2* morphants (Fig. 5R,U).

One interpretation of the enhanced expression of MCC markers in *jagged 2* morphants would be that MCC cell number was increased by the transfating of transporting epithelial cells to MCCs. Alternatively, it is possible that MCC and transporting-cell phenotypes could co-exist in the same cells and that *jagged 2* loss of function may simply be derepressing axonemal gene expression. We therefore examined *jagged 2* morphants for expression of the transporters *trpM7*, *slc13a1* and Na,K-ATPase alpha1A4:GFP. Strikingly, *jagged 2* morphants showed a complete loss of all mid-segment ion-transporter gene expression (Fig. 5M-Q) as well as a segment-specific reduction in Na,K-ATPase alpha1A4:GFP expression (Fig. 5S,T). As a further test for specificity, we also knocked down *jagged 1b* expression and assessed MCC and transporter gene expression. *jagged 1b* knockdown had no effect on marker gene expression, indicating that the observed effects were specific to *jagged 2* (data not shown). The results suggest that *jagged 2* acts as a repressor of a genetic pathway leading to MCC differentiation and that part of this patterning mechanism involves...
jagged 2 repression of its own mRNA synthesis in a subset of pronephric cells. Additionally, the fates of MCCs and transporting epithelial cells appear to be mutually exclusive, with jagged 2 repression of MCC fate being necessary to allow neighboring cells to acquire a secondary, transporting-epithelial-cell fate.

A role for Notch3 in mediating Jagged 2 signaling in the pronephros

Notch3 has previously been shown to be expressed in the pronephros (Lawson et al., 2001) and was a reasonable candidate for mediating the effects of Jagged 2. We confirmed that notch3 mRNA was expressed strongly in the proximal and distal pronephros, and is diffusely expressed in the pronephric midsegment (Fig. 6A,B). Knockdown of notch3 with a morpholino oligo targeting notch3 exon 27 produced a morphant ‘allele’ truncated after exon 26 (stop codon following 20 mis-sense amino acids; see Fig. S1 in the supplementary material) and lacking both the transmembrane domain and the notch3 intracellular domain required for nuclear signaling. Similar to jagged 2 morphants, both shippo1-positive (Fig. 6C,D) and flr-positive (Fig. 6E,F) cells were significantly increased in number in notch3 morphants, whereas the expression of slc13a1 (Fig. 5G,H) and trpM7 (Fig. 5LJ) was nearly completely abolished in the early-distal pronephros.

Regulation of pronephric cell patterning requires mind bomb activity

mind bomb mutants lack an E3 ubiquitin ligase that is required to process Notch ligands and to facilitate Notch signaling (Itoh et al., 2003). To test whether Jagged 2 signaling required mind bomb activity, we examined MCC and transporting-cell markers in mind bomb homozygotes. Consistent with the jagged 2- and notch3-morphant phenotypes, expression of the MCC markers shippo1 and jagged 2 was dramatically expanded in mind bomb homozygotes (Fig. 7A-D). Shippo1 expression in single cells of the distal segment (Fig. 7A) was expanded to include nearly all cells of this segment (Fig. 7B, ds). In addition, the number of shippo1-positive cells was significantly increased in the proximal segment of the pronephros (Fig. 7B, ds). Expression of the transporting-cell markers trpM7 and slc13a1 was nearly completely abolished (Fig. 7E-H). These results indicate that mind bomb-processing activity is required for jagged 2/notch3 patterning of the pronephros.

Timing of Jagged 2/Notch3 signaling during pronephric development

jagged 2 mRNA expression is initiated early in the intermediate mesoderm and is maintained in fully differentiated MCCs (Fig. 4A,B). To determine when Jagged 2/Notch3 signaling acts to restrict the MCC fate and to assess whether continued Jagged 2/Notch3 signaling is required to generate the mature pattern of cell types in the pronephros, we used the gamma-secretase inhibitor DAPT to block notch cleavage (Geling et al., 2002) at different time points during pronephric development. Expansion of MCC fate (increased shippo1 expression; Fig. 8A-C) and the loss of trpM7-expressing transporter cells (Fig. 8G-I) occurred when DAPT was added to egg water at 9.5 hpf, but not if it was added later, at 24 hpf (Fig. 8). Interestingly, slc13a1 expression was sensitive to inhibition of Notch signaling at later developmental stages (24+ hpf; Fig. 8J-L), indicating that, although expressed in the same cells, trpM7 and slc13a1 may respond to Notch signaling in different ways. The effect of DAPT on MCC differentiation suggested that cell fate choice is decided during late gastrulation and early somitogenesis, and that continued Notch3 signaling after 24 hpf may not be required for its expression.
to limit the number of MCCs. To further refine the time window in which Jagged 2/Notch signaling acts, we treated embryos with DAPT starting at progressively later time points and assayed the number of MCCs at 34 hpf by counting the number of DAPT-induced MCCs at 34 hpf (Fig. 9A). As shown in Fig. 9A, control pronephric tubules contained, on average, 26±4 (±s.d., n=8) MCCs, and treatment with 100 μM DAPT at 9.5 hpf increased this number to 67±14 (n=8; although this is possibly an underestimate of MCC number given the difficulty in counting overlapping shippo1-positive cells). MCC differentiation was sensitive to DAPT only during a narrow time window (around 9-10 hpf), and later treatment at 12 or 15 hpf did not significantly affect the number of MCCs (Fig. 9A).

**Ectopic expression of the Notch1a intracellular domain blocks MCC differentiation**

Gamma-secretase cleavage of Notch proteins liberates the NICD, which is then transported to the nucleus where it acts as a transcriptional regulator. Our results on Jagged/Notch loss of function indicate that Notch signaling acts as a repressor of the MCC fate. To directly test this, we expressed the Notch1a ICD under heat-shock control and assayed shippo1 expression in the pronephros. Heat-shock induction of the NICD at 8 hpf dramatically reduced the number of shippo1-positive MCCs at 34 hpf (Fig. 8E); when the NICD was induced by heat shock at 9 hpf (Fig. 8F), MCC number was also reduced, but to a lesser extent. shippo1-positive cells were counted in each case and the results quantified (Fig. 9B). Control heat-shocked siblings had 27±2 (±s.d.; n=10) MCCs per tubule, whereas those in which the NICD was induced at 8 hpf and 9 hpf had 9±3 (±s.d.; n=35) and 17±4 (±s.d.; n=7) MCCs per tubule, respectively. These results confirm that Notch signaling acts as a repressor of MCC fate early in the development of the pronephros.

**Enhanced ciliogenesis and rescue of the double bubble kidney cyst mutant by jagged 2 knockdown**

The strong expression of ciliogenic genes and the absence of transporter gene expression suggested that loss of Jagged 2/Notch signaling could cause cells to fully transfate to the MCC cell type. To assess whether jagged 2-morphant pronephric cells acquired the functional character of MCCs, we examined the abundance and motility of cilia in morphant embryos. Immunostaining with anti-acetylated tubulin revealed that the loss of jagged 2 function increased the abundance of luminal cilia in the pronephros (Fig. 10A,B). Moreover, the newly formed cilia in jagged 2-morphant pronephroi all beat rapidly (see Movie 2 in the supplementary material), similar to the behavior of wild-type pronephric cilia (see Movie 1 in the supplementary material). To better visualize beating cilia, the pronephric opening of these embryos was mechanically obstructed to expand the pronephric lumen. Notch3 morphants also showed an increase in cilia bundles in the pronephros (data not shown). These results indicate that loss of Jagged 2/Notch signaling not only enhanced axonemal gene expression, but was sufficient to transfate pronephric cells to fully functional MCCs.

Motile cilia in the pronephros are required to maintain high rates of fluid output and prevent backpressure, which can lead to pronephric cyst formation (Kramer-Zucker et al., 2005). The transfating of epithelial cells to MCCs in jagged 2 morphants raised the possibility that the enhancement of the ciliogenic program might be sufficient to compensate for deficiencies in ciliogenesis that are observed in a class of pronephric cyst mutants. We therefore tested whether jagged 2 knockdown in pronephric cyst mutant

![Fig. 6. Notch3 mediates Jagged 2 signaling in the pronephros.](Image)

(A) Expression of notch3 in the pronephros (arrowheads). (B) Expression of notch3 in the proximal pronephric tubules (arrowhead). (C,D) shippo1 expression in single cells of control embryos at 34 hpf (C) is expanded by notch3 exon 27-morpholino injection (D). (E,F) Similarly, expression of flr that is observed in control embryos (E) is expanded in notch3 exon 27 morphants (F). (G,I) Transporting-cell markers slc13a1 (G) and trpM7 (I) are greatly reduced in notch3 exon 27 morphants (H and J, respectively).

![Fig. 7. Transfating of pronephric cells in the mind bomb mutant.](Image)

(A, B) shippo1 expression in control wild-type embryos is restricted to single cells (A), whereas, in mind bomb homozygotes (B), pronephric distal segment cells (ds) are uniformly positive and there is an increase in shippo1-positive cells in the proximal segment (ps). (C,D) Wild-type jagged 2 expression (C) is also expanded in mind bomb homozygotes (D). (E,G) trpM7 (E) and slc13a1 (G) expression is almost completely lost in mind bomb homozygotes (F and H, respectively).
embryos would have any effect on cilia structure, cilia motility and/or pronephric cyst formation. We screened five different pronephric cyst mutants [flr, oval (ij888), tg238a, schmalhans and double bubble] for potentially beneficial effects of jagged-2-morpholino injection on ciliogenesis. Based on an apparent improvement in phenotype (absence of cysts and edema), one of these mutants, double bubble (m468) (Drummond et al., 1998), was chosen for further analysis. The double bubble (dbb) mutant is characterized by a loss of cilia structure (Fig. 10E,G) with consequent lumen dilatation and pronephric cyst formation (Drummond et al., 1998). dbb homozygotes can also be identified by a characteristic ventral axis curvature (Drummond et al., 1998).

High-speed micro videos showed that dbb homozygotes lack detectable motile cilia (see Movie 4 in the supplementary material) in comparison to similar pronephric segments in wild-type embryos (see Movie 3 in the supplementary material). Significantly, jagged-2 knockdown rescued normal cilia formation in double bubble-mutant embryos (Fig. 10C,D). Rescued cilia bundles in dbb homozygotes were actively motile (see Movie 5 in the supplementary material), similar to wild-type cilia (see Movie 3 in the supplementary material). Remarkably, jagged-2 knockdown completely prevented pronephric cyst formation in dbb homozygotes and restored wild-type pronephric structure to mutant embryos (Fig. 10E-H). In three separate experiments, injection of jagged-2 exon 20 donor-morpholino into embryos from pair matings of dbb heterozygotes prevented cyst formation (embryos with cysts/total embryos in clutch: 0/42, 0/46 and 0/41). Control morpholino-injected embryos developed cysts at standard Mendelian frequencies (25%) and injection of the same concentration of jagged-1b morpholino or control morpholinos had no effect on the dbb-mutant phenotype (data not shown). We also tested whether DAPT treatments that significantly increased MCC expression (Fig. 8B, Fig. 9A) would affect cyst formation in double bubble mutants. Treating dbb-mutant embryos with 100 μM DAPT at 9.5 hpf significantly reduced cyst formation (Fig. 10I,J). These results strengthen our previous findings that motile cilia are key organelles in fluid transport in the pronephros (Kramer-Zucker et al., 2005) and further suggest that the capacity of the kidney for fluid transport can be modulated by Notch-mediated control of the number of multiciliated versus transporting cells during a crucial stage of kidney-nephron patterning.

DISCUSSION

Like all epithelial organs, the physiological activity of the zebrafish pronephros is dependent on a partitioning of epithelial tubules into segments and cell types with unique activities that combine to achieve overall organ function. Kidney cells are specialized for osmotic homeostasis and electrolyte balance, which, in a freshwater vertebrate, translates to excretion of water and avid recovery of filtered ions and metabolites (Hickman and Trump, 1969). In the pronephros, multiciliated cells form the biological equivalent of a ‘screw pump’ by beating in a helical wave pattern in a tight-fitting tubule lumen to propel fluid out of the pronephric opening (Kramer-Zucker et al., 2005), while transporting epithelial cells recover specific ions from filtered blood depending on the spectrum of ion transporters that they express. Very little is known about how kidney-tubule cells achieve their different identities. What we have shown is that a patterning mechanism underlying cell-type specification in the pronephros involves Jagged-2/Notch signaling. Our findings point to a lateral inhibition mechanism in generating a ‘salt-and-pepper’ pattern of multiciliated cells intercalated among transporting cells.
Notch signaling and kidney development

Previous studies on mouse kidney development have demonstrated roles for Notch signaling in glomerulus formation and in specifying the fates of proximal versus distal nephron tubule segments. Notch2 expression in glomerular endothelial cells and Jagged 1 expression in endothelial cells is required for development of the glomerular vasculature (McCright, 2003). Defects in glomerular vasculature structure have also been reported in combined jagged 2/jagged 1b morphants in zebrafish, suggesting that similar developmental mechanisms may underlie zebrafish pronephric development (Lorent et al., 2004). General inhibition of Notch signaling by gamma-secretase inhibition or by mutations in mouse presenilin genes results in a loss of both glomerular and proximal tubule cells, and in the preservation of the distal nephron fate (Cheng et al., 2003). Jagged 1 signaling has also been shown to regulate branching morphogenesis of the renal collecting system (Kuure et al., 2005). In pronephric development in Xenopus, Notch signaling also regulates the patterning of proximal versus distal tubule types (McLaughlin et al., 2000). Together, these studies point to a role for Notch signaling in nephron precursor structures (S-shaped bodies or their equivalents) in broadly defining major nephron segments and structures.

Our focus on Jagged 2/Notch3 signaling in the zebrafish introduces an additional role for Notch signaling in the generation of intercalated cell patterns in the nephron. In the mammalian collecting system, ‘intercalated cells’ exist interspersed among principal cells, and function in pH homeostasis (Seldin and Giebisch, 1992). Although mechanisms regulating intercalated cell differentiation are currently unknown, the expression of multiple Notch and jagged genes in the nephron suggests that notch signaling could play a role (Challen et al., 2006). In the zebrafish larval intestine, secretory and absorptive cells exist intermixed, and this balance of cell types is controlled by a Notch-mediated lateral inhibition mechanism involving deltaD expression in secretory cells (Crosnier et al., 2005). In the zebrafish liver, the hepatocyte/cholangiocyte cell-fate choice also appears to be under the control of Notch signaling (Lorent et al., 2004). In the pancreas, disruption of Notch signaling by mutation in mind bomb or by expression of a dominant-negative Suppressor of Hairless construct accelerates differentiation of exocrine acinar cells (Esn et al., 2004). The picture that emerges from these studies is that Notch signaling may act at multiple stages of organogenesis to first regulate the timing and overall pattern of organ progenitor cell differentiation, and may later serve to modify cell-type specification by superimposing a repression mechanism that regulates the proportion of different cell types in the mature organ.

Notch signaling pathways and the emergence of multiciliated cells

The work presented here suggests that, early in pronephric cell differentiation (8–10 hpf), cells expressing high levels of Jagged 2 interact with neighboring Notch-expressing cells (summarized in Fig. 11). Jagged 2/Notch signaling requires the E3 ligase mind bomb, and initiates proteolysis of Notch3 and the liberation of the transcriptionally active NICD. The NICD activates expression of unknown repressors that downregulate rfx2 (and downstream cilia genes) and jagged 2. Later in development, a subset of cells becomes multiciliated while neighboring cells acquire a transporting-epithelial-cell fate and express ion channel genes.

In other Notch signaling contexts, the hairy/enhancer of split family of genes exert the transcriptional-repressor function of Notch signaling (Iso et al., 2003). We have found that her9 is expressed in the pronephros (see Fig. S2 in the supplementary material); however, it is clear that other factors must act to repress ciliogenic gene expression, because the expression of rfx2 and shippo1 was unaffected by a her9-translation-blocking morpholino (Y.L. and I.A.D., unpublished). Whatever the identity of Notch-induced transcriptional repressors may be, it is likely that the transcription factor rfx2 may be a primary target of repression, because it belongs to a gene family known to serve as master regulators of ciliogenic...
Although we have characterized several elements of Notch signaling in pronephric patterning, aspects of our data indicate that the framework is incomplete. In embryos in which Notch signaling has been generally inhibited (i.e. mind bomb mutants and DAPT-treated embryos), the transfecting of cells to MCCs is observed not only in the pronephric distal segment but also in the proximal segment, where MCC cells are not as frequently seen, suggesting that additional notch genes or Notch ligands may regulate MCC differentiation. These factors are unlikely to include jagged 1b, because single knockdown of jagged 1b or double knockdown of jagged 1b and jagged 2 did not replicate the mind bomb phenotype (data not shown).

Development and functions of cilia

The importance of cilia in normal organ function is highlighted by the pathology observed in organisms with non-functional cilia: kidney cystic disease, retinal degeneration, organ laterality defects and hydrocephalus (Pazour, 2004). Mutations in a variety of cilia-associated proteins cause kidney tubules to become cystic (Barr et al., 2001; Blacque et al., 2004; Fan et al., 2004; Haycraft et al., 2001; Kim et al., 2004; Kramer-Zucker et al., 2005; Morgan et al., 2002; Murcia et al., 2000; Myktyyn et al., 2004; Otto et al., 2003; Pazour et al., 2000; Pazour and Rosenbaum, 2002; Qin et al., 2001; Sun et al., 2004; Yoder et al., 2002). In cultured mammalian epithelial cells, cilia are proposed to function as non-motile fluid-flow sensors that regulate epithelial responses to flow via calcium signaling (Nauli et al., 2003; Prætorius and Spring, 2001). We have demonstrated a role for motile cilia in cystic kidney pathology that affects the zebrafish pronephros (Kramer-Zucker et al., 2005). Multiciliated cells similar to what we describe here in the zebrafish have also been shown to exist intercalated between proximal tubule cells in the human kidney (Duffy and Suzuki, 1968; Hassan and Subramanyan, 1995; Katz and Morgan, 1984; Ong and Wagner, 2005). The function of these cells is currently unknown. Although there is no evidence for Notch signaling in the development of these cells, studies of epidermal development in Xenopus have highlighted a role for Notch signaling in the context of ciliated-cell development. The regularly spaced pattern of MCC differentiation in the Xenopus epidermis has been shown to be regulated by X-Delta-1 expression in a subset of epidermal cells, which later abundantly express alpha-tubulin and become multiciliated (Deblandre et al., 1999). Taken together with our work, the results suggest that Notch signaling might be a general feature of specialized ciliated cell development.

Our finding that jagged 2 knockdown can rescue cilia assembly in the double bubble mutant and prevent cyst formation strengthens our previous results that cilia-driven fluid flow in the pronephros is a key factor in cystic pathology. This result also raises questions concerning the process of ciliogenesis and the nature of the dbb mutation. It is not clear at present why only dbb and not other ciliogenesis mutants are rescued by jagged 2 knockdown. The dbb allele we are studying (m468) could be a hypomorphic allele or, alternatively, the dbb protein may not be absolutely essential for ciliogenesis. Other intraflagellar transport (IFT) proteins, if expressed in high enough amounts, may be sufficient to compensate for the loss of dbb. Further characterization of the dbb mutation and its interactions with other proteins involved in IFT will be necessary to resolve these issues.

The authors thank Mike Pack for jagged 2 and jagged 3 ATG morpholinos; Ajay Chitnis for jagged cDNA clones, mind bomb-mutant embryos and advice on the manuscript; Alan Davidson for mind bomb embryos; Calum MacRae for the notch3 exon 27 morpholino; Anirudh Majumdar for Fig. 1c; Mary McKee and Dennis Brown for microscopy support; Scott Halley for the
fluorescent in situ protocol; and members of the Drummond laboratory for feedback on this work. This work was supported by NIH RO1 DK53903 to I.A.D.

Supplementary material
Supplementary material for this article is available at http://dev.biologists.org/cgi/content/full/134/6/1111/DC1

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