Control of hair follicle cell fate by underlying mesenchyme through a CSL–Wnt5a–FoxN1 regulatory axis

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Epithelial–mesenchymal interactions are key to skin morphogenesis and homeostasis. We report that maintenance of the hair follicle keratinocyte cell fate is defective in mice with mesenchymal deletion of the CSL/RBP-Jk gene, the effector of “canonical” Notch signaling. Hair follicle reconstitution assays demonstrate that this can be attributed to an intrinsic defect of dermal papilla cells. Similar consequences on hair follicle differentiation result from deletion of Wnt5a, a specific dermal papilla signature gene that we found to be under direct Notch/CSL control in these cells. Functional rescue experiments establish Wnt5a as an essential downstream mediator of Notch–CSL signaling, impinging on expression in the keratinocyte compartment of FoxN1, a gene with a key hair follicle regulatory function. Thus, Notch/CSL signaling plays a unique function in control of hair follicle differentiation by the underlying mesenchyme, with Wnt5a signaling and FoxN1 as mediators.

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The skin provides a well-studied example of epithelial cell differentiation and morphogenesis [Botchkarev and Paus 2003]. In the interfollicular epidermis, keratinocytes terminally differentiate along a single lineage, while in hair follicles they undergo a complex differentiation program that gives rise to several concentric layers of cells with separate fates. Hair follicle morphogenesis and cycling are critically dependent on an interplay between keratinocytes and cues originating from the underlying mesenchyme [Botchkarev and Paus 2003]. Several key developmental pathways are involved in establishing and/or maintaining the hair follicle keratinocyte cell fate, in concert with keratinocyte-specific genes like FoxN1, a transcription factor whose mutation results in the nude [mouse, rat, or human] phenotype [Mecklenburg et al. 2005].

Notch signaling is an important form of cell–cell communication that plays a key role in control of cell fate, stem cell potential, and differentiation [Bray 2006; Hurlbut et al. 2007]. The Notch gene family [Notch1, Notch2, Notch3, and Notch4] consists of evolutionarily conserved transmembrane receptors. Interactions with ligands of the Delta and Jagged families, present on the surface of neighboring cells, trigger proteolytic cleavage of these receptors with release of the intracellular domain of Notch receptors (NICD). The best characterized “canonical” pathway of Notch activation involves translocation of NICD to the nucleus, where it associates with the DNA-binding protein RBP-Jk (CBF-1 or CSL), converting it from a repressor into an activator of transcription [Kopan and Ilagan 2009].

This pathway is highly context-dependent, and can exert opposite roles in growth/differentiation of different cell types, at different developmental stages, and/or under normal versus pathological conditions. While great attention has been paid to the role of Notch signaling in the epithelial compartment of the skin, for both interfollicular and hair follicle keratinocyte differentiation [Dotto 2008; Watt et al. 2008], a possible role of this pathway in the adjacent mesenchyme has not been investigated. However, an indication of Notch activity in both dermal fibroblasts and dermal papilla cells has been provided by the analysis of transgenic mice with a Notch-responsive GFP reporter, immunostaining for activated Notch1 and/or expression of “canonical” Notch target genes of the Hes/HERP family [Okuyama et al. 2004; Estrach et al. 2006; Moriyama et al. 2006; Lee et al. 2007]. We report...
here that CSL/RBP-Jk, the effector of “canonical” Notch signaling, plays an as yet unsuspected role in control of hair follicle cell fate by the underlying mesenchyme. This is linked to a unique nonredundant function of a specific Wnt family member, Wnt5a, as a direct Notch/CSL target in dermal papilla cells, with FoxN1 as a downstream target in keratinocytes.

Results

Compromised hair follicle differentiation in mice with mesenchymal deletion of the RBP-Jk gene

Little is known about the function of Notch signaling in cells of the mesenchymal lineage. “Canonical” Notch signaling is critically dependent on the DNA-binding protein CSL/RBP-Jk (Bray 2006). To delete this gene specifically in the mesenchyme, transgenic mice with Cre expression driven by the promoter of the collagen type I [α2 chain] gene [Col-I-Cre] (Florin et al. 2004) were crossed with mice with the RBP-Jk gene flanked by loxP sites (Han et al. 2002). Further crossing of mice into a ROSA26R background confirmed that, as reported previously [Florin et al. 2004], Cre activity in the skin was confined to dermal papilla cells and dermal fibroblasts [Supplemental Fig. 1]. Specificity of deletion of the RBP-Jk gene in the mesenchymal compartment of the skin [dermal papilla cells and dermal fibroblasts] was further verified by physical separation of various skin components, followed by real-time PCR analysis of genomic DNA [Supplemental Fig. 2]. Analysis of embryonic skin showed specific deletion of the RBP-Jk gene in the mesenchymal compartment as early as embryonic day 15.5 (E15.5) [data not shown].

Phenotypically, homozygous [Col-I-Cre × RBP-JkloxP/loxp] mice were recognizable after birth by their “kinky” tail and smaller size, with areas of decreased hair density (Fig. 1A). They also exhibited a skeletal phenotype that will be presented elsewhere. Histologically, hair follicles were structurally normal at birth [Supplemental Fig. 3], but, by 4–10 d, a significant percentage started to show structural alterations, with disrupted inner root sheath [IRS] and cuticle layer organization, followed by hair shaft/cortex alteration and signs of cystic degeneration (Fig.

Figure 1. Defects in hair follicle differentiation in mice with mesenchymal deletion of the RBP-Jk gene. (A) Mice with the RBP-Jk deletion [−/−] and age-matched controls [+/+] at P14. (B) Histological images of back skin sagittal sections from a mouse at 10 d after birth with mesenchymal deletion of the RBP-Jk gene [−/−] versus an age-matched control [+/+], for analysis of hair follicles at earlier times, see Supplemental Figures 3 and 4. (C) Higher-magnification image of an abnormal hair follicle from a mouse with mesenchymal deletion of the RBP-Jk gene [−/−] [marked by a square in B] versus a hair follicle from an age-matched control [+/+]. (D, E) Immunofluorescence analysis of hair follicles of mice with mesenchymal deletion of the RBP-Jk gene [−/−] versus controls [+/+] at 10 d of age with antibodies against 44-kDa/46-kDa acidic hair keratins [AE13], trichohyalin [AE15], and cytokeratins 1 and 10. (F) Real-time RT–PCR analysis of physically separated hair follicles from mice with [−/−] and without [+/] deletion of the RBP-Jk gene at P10, with primers specific for the indicated genes. Hair follicles from mice with deletion of the RBP-Jk gene were analyzed separately, depending on their relatively normal [RN] and abnormal [AN] structure, and were isolated as shown in Supplemental Figure 2C. Results are expressed as relative expression levels, using β-actin for internal normalization. (G) Histochemical staining of alkaline phosphatase activity in hair follicles of mice with mesenchymal deletion of the RBP-Jk gene [−/−] versus controls at 10 d of age. (H) Immunofluorescence analysis of similar hair follicle sections [left two panels] and of cysts that develop in RBP-Jk−/− mice by 3 wk of age (right panel) with antibodies against versican. (I) Immunofluorescence analysis of hair follicles of control and RBP-Jk mutant mice with antibodies against Sox2. Bars, 40 µm.
1B,C; Supplemental Fig. 4). Immunoﬂuorescence analysis showed substantially decreased expression of 44-kDa/46-kDa acidic hair keratins, a marker of cortex and cuticle layers of the hair shaft, and trichohyalin, a marker of medulla and IRS [Lynch et al. 1986], while, conversely, expression of markers normally conﬁned to the interfollicular epidermis (cytokeratins 1 and 10) was detectable in hair follicles [Fig. 1D,E]. This “switch” between hair follicle and interfollicular marker expression was further documented and quantiﬁed by real-time RT–PCR analysis of physically isolated hair follicles from the [ColI-Cre × RBP-JkloxP/loxP] mice and age-matched controls [Fig. 1F]. Degeneration of hair follicles into cysts became widespread by 2–3 wk, with the mature cysts exhibiting high levels of interfollicular marker expression [Supplemental Fig. 5A–C]. Vibriasse hair follicles were also normal after birth, but were then lost in the mutant mice by 3–4 wk, undergoing similar cystic degeneration as pelage hair follicles [Supplemental Fig. 5D,E].

Versican and alkaline phosphatase staining showed that dermal papillae remained in contact with hair follicles as well as the cysts that formed over time [Fig. 1G,H]. Expression of another dermal papilla cell marker, Sox2, was also similar in control and mutant mice [Fig. 1I]. Sox2-positive dermal papilla cell populations were shown recently to be essential for production of auchen and awl hair ﬁbers [Driskell et al. 2009]. Consistent with our expression data, plucked hair analysis showed that mutant mice produced all four types of hair in an approximately normal ratio [Supplemental Fig. 6A]. Reflecting the hair follicle structural defects, a large percentage of all hair types showed abnormal packing and irregular ﬁber proﬁle, with aberrant restrictions and bending [Supplemental Fig. 6B].

As with many other mutations, mice with keratinocyte-speciﬁc deletion of the RBP-Jk gene exhibit also hair follicle degeneration and cyst formation [Blanpain et al. 2006]. We therefore compared in detail alterations resulting from deletion of the RBP-Jk gene by the Cre transgene driven by a Coll versus keratin 5 [K5] promoter, using mice of a young age and at an early stage of hair follicle degeneration [Supplemental Fig. 7]. In both cases, deletion of the RBP-Jk gene resulted in markedly decreased expression of acidic hair keratins and trichohyalin, and K6 up-regulation. However, while in the [ColI-Cre × RBP-JkloxP/loxP] mice there was pronounced hair follicle expression of suprabasal interfollicular cytokeratins 1 and 10 (K1 and K10), no such expression was found in the [K5-Cre × RBP-JkloxP/loxP] mice [Supplemental Fig. 7B]. As reported previously [Blanpain et al. 2006], K1/K10 expression was decreased even in the interfollicular epidermis of mice with keratinocyte-speciﬁc RBP-Jk deletion, while in mice with the mesenchymal deletion, expression of these genes was increased [Supplemental Fig. 7C]. This was paralleled by the expected decrease of direct Notch target genes Hes1 and Hey1 in mice with the ﬁrst genotype, but not in mice with the second genotype [Supplemental Fig. 7C]. Besides differentiation marker expression, dermal papilla cells, as identiﬁed by versican and Sox2 staining, were normally present in the skin of mice with the Coll-Cre-driven deletion but not K5-Cre-driven deletion [Supplemental Fig. 7D], consistent with distinct phenotypes resulting from RBP-Jk deletion in the two skin compartments.

**Intrinsically defective hair-inducing capabilities of dermal papilla cells with RBP-Jk deletion**

To assess whether the phenotype of the [ColI-Cre × RBP-JkloxP/loxP] mice is skin intrinsic or secondary to systemic alterations, we resorted to full-thickness skin grafting onto nude mice. Grafts of RBP-Jk mutant skin showed aberrant hair follicle differentiation and cyst formation, as in the original mice with this mutation [Fig. 2A].

The interesting possibility exists that RBP-Jk deletion in extrafollicle dermal cells contributes to the observed phenotype. To speciﬁcally establish whether the hair follicle alterations can be attributed to a defect in dermal papilla cells, we performed hair follicle reconstitution assays [Zheng et al. 2005] using freshly isolated dermal papilla cells with or without RBP-Jk deletion in combination with wild-type keratinocytes. The latter were obtained from mice with a GFP transgene driven by an actin promoter [Okabe et al. 1997], which allowed unequivocal identiﬁcation of hair follicles reconstituted by donor cells relative to endogenous hair follicles of the recipient mice.

Efficient hair follicle formation was observed with GFP-positive keratinocytes admixed with control dermal papilla cells, while only few hair follicles (~10-fold difference) were formed by the same keratinocytes admixed with RBP-Jk-deﬁcient papilla cells [Fig. 2B]. Instead, the latter combination of cells resulted in widespread formation of cysts, which were rarely observed in the graft controls [Fig. 2C]. Immunoﬂuorescence analysis showed that the aberrantly reconstituted hair follicles and cysts formed by GFP-positive keratinocytes in combination with RBP-Jk−/− dermal papilla cells had—as in the [ColI-Cre × RBP-JkloxP/loxP] mice—reduced acidic hair keratins and trichohyalin expression, and were positive for cytokeratins 1 and 10 [Fig. 2D].

**The Wnt5a gene is a direct Notch/CSL target in dermal papilla cells**

Gene expression studies were undertaken to probe into the molecular defects resulting from the RBP-Jk deletion in dermal papilla cells. As expected, real-time RT–PCR analysis showed that the “canonical” Notch–RBP-Jk target genes Hes1, Hes5, and Hey1 were down-modulated in dermal papilla cells isolated from mice with the deleted RBP-Jk gene [Fig. 3A]. Expression of signaling molecules that make part of the “dermal papilla-speciﬁc signature”—Wnt5a, Fgf7, Fgf10, and Noggin—was confirmed at the protein level by immunoblot analysis of dermal papilla cell extracts [Fig. 3B] and immunoﬂuorescence of corresponding skin.
sections [Fig. 3C]. Modulation of these genes, similar to that observed in vivo, was found with cultured dermal papilla cells from (ColI-Cre × RBP-J\(^{loxp/loxp}\)) mice [Fig. 3D], as well as with dermal papilla cells with the floxed RBP-J\(_k\) gene after “acute” deletion by adenovirally mediated Cre expression [Fig. 3E].

The Wnt5a, Noggin, and Fgf7/10 genes code for diffusible factors that may be under parallel RBP-J\(_k\) control, or regulation of one of them might play a primary role on the others. To address this question, dermal papilla cells with the RBP-J\(_k\) deletion were treated with each of these factors individually, in a purified form, followed by analysis of expression of the others. As shown in Figure 4A, treatment of the RBP-J\(_k\)^\(-/-\) dermal papilla cells with Wnt5a rescued, to a partial extent, expression of Noggin, Fgf7, and Fgf10 expression. The converse treatment with Noggin, Fgf7, or Fgf10 had no effect on Wnt5a levels [Fig. 4B], suggesting that Wnt5a is a primary target of RBP-J\(_k\) function.

Consistent with this conclusion, activated Notch1 expression by adenoviral vector infection, or tamoxifen-dependent activation of a retrovirally transduced Notch1–ER fusion protein [Schroeder and Just 2000], induced Wnt5a in wild-type dermal papilla cells, but not in RBP-J\(_k\)-deficient cells [Fig. 4C]. Induction of Wnt5a expression was similarly observed upon endogenous Notch activation by treatment of dermal papilla cells with purified Jagged 1 ligand [Fig. 4D]. Nucleotide sequence analysis of the Wnt5a gene promoter and first intronic regions showed the presence of “canonical” RBP-J\(_k\)-binding sites, to which the endogenous Notch1 protein was found to bind by chromatin immunoprecipitation (ChiP) assays with control dermal papilla cells but not cells with the RBP-J\(_k\) deletion [Fig. 4E]. In parallel, luciferase reporter assays showed that activated Notch1 expression induces activity of a Wnt5a promoter construct containing RBP-J\(_k\)-binding sites, but not one with the sites deleted [Fig. 4F].

Compromised hair follicle differentiation in mice with deletion of the Wnt5a gene

Mice with homozygous deletion of the Wnt5a gene die at birth due to a number of developmental defects [Yamaguchi et al. 1999; Yang et al. 2003]. So far, the consequences of Wnt5a deletion in the skin have not been analyzed. Histological analysis of E18.5 embryos showed that loss of Wnt5a, as the mesenchymal RBP-J\(_k\) deletion, causes no detectable defects in hair follicle morphogenesis and differentiation at this time (data not shown).
To assess the role of Wnt5a after birth, E17.5 embryo skins were grafted onto nude mice. Hair follicle abnormalities started to be seen by 1 wk after grafting of the Wnt5a mutant skin, becoming more substantial by 2 and 4 wk (Fig. 5A, B, Supplemental Fig. 8). In parallel with these alterations, immunofluorescence analysis showed consistent down-modulation of acidic hair keratins and inappropriate cytokeratin 10 expression, with dermal papilla cells (as detected by versican staining) still present (Fig. 5C). In the latter, however, there was strong reduction of Noggin and Fgf10 expression, consistent with our other results pointing to Wnt5a as an upstream regulator of these molecules (Fig. 5D).

While Wnt5a expression is limited to dermal papilla cells during development, after birth it also extends to the keratinocyte compartment (Reddy et al. 2001; Rendl et al. 2005). To determine to what extent the hair follicle alterations of the Wnt5a−/− grafted skin can be attributed to defective dermal papilla cell function, hair follicle reconstitution assays were undertaken with dermal papilla cells isolated from Wnt5a−/− versus control littermates in combination with keratinocytes harboring an intact Wnt5a gene and expressing GFP. Wnt5a-deficient dermal papilla cells, like those with the RBP-Jk deletion, were similarly impaired in their hair follicle-inductive properties, inducing instead extensive cyst formation.

Figure 3. Changes of dermal papilla signature genes by mesenchymal deletion of the RBP-Jk gene. (A) Real-time RT–PCR analysis for expression of the indicated “canonical” Notch target genes [left panel] and dermal papilla signature genes [right panel] (Rendl et al. 2005) in freshly isolated dermal papilla cells (by the method illustrated in Supplemental Fig. 2A) from mice with mesenchymal deletion of the RBP-Jk gene (−/−) and age-matched controls (+/+). Expression levels are expressed as relative values after internal normalization for β-actin. (B) Immunoblot analysis of dermal papilla cell extracts from mice with (−/−) and without (+/+) deletion of the RBP-Jk gene with antibodies against the Wnt5a, Noggin, and Fgf7 proteins, using β-actin for equal loading control. The position of molecular size markers is indicated. The immunoblot for Wnt5a resulted in multiple bands, as reported in previous publications (Blumenthal et al. 2006). (C) Immunofluorescence analysis of Wnt5a, Noggin, and Fgf10 expression in anagen hair follicles of mice with and without RBP-Jk deletion, at 10 d of age. Hair follicle boundaries and dermal papillae are indicated by dotted lines and arrows, respectively. (D) Real-time RT–PCR analysis of expression of the indicated genes in dermal papilla cells that were cultured from mice with (−/−) and without (+/+) deletion of the RBP-Jk gene, at 4 d after plating. (E) Real-time RT–PCR analysis of cultured dermal papilla cells from mice with the RBP-Jk gene flanked by loxP sites, 24 h after infection of these cells with a Cre-expressing adenovirus [Adeno-Cre] or Adeno-GFP control. Besides the indicated genes, real-time PCR analysis with primers specific for exons 6 and 7 of the RBP-Jk gene showed >80% deletion of this portion of the gene in the Cre-expressing cells (data not shown).
according to the following formula: percentage total = \(2^{\Delta Ct} \times 5\), where \(\Delta Ct = Ct\) (input) – \(Ct\) (immunoprecipitation). \(Ct\) (cycle threshold) of the real-time RT–PCR. (Bottom panel) Gel electrophoresis of the PCR products at the end of the real-time RT–PCR, showing, in each case, single-amplified DNA bands of the expected size. [I.P.] input DNA; [N1] Notch1 immunoprecipitate; [IgG] nonimmune control. The primers used to amplify these Wnt5a promoter regions are listed in Supplemental Table S1. (I.P.) input DNA; (N1) Notch1 immunoprecipitate; (IgG) nonimmune control. The amount of precipitated DNA was calculated relative to the input chromatin. Mutually precipitated DNA was similarly analyzed and used as “input DNA” control. The amount of precipitated DNA was calculated relative to the total input chromatin, and was expressed as percentage of the total

[Fig. 5E,F]. Even in this case, immunofluorescence analysis showed down-modulation of hair follicle differentiation markers and inappropriate cytokeratin 1 expression (Fig. 5G).

Functional rescue of defective RBP-Jk\(^{-/-}\) hair follicles by Wnt5a

To assess whether the defects caused by deletion of RBP-Jk can be attributed to decreased Wnt5a expression, intact skin of postnatal day 4 (P4) mice was placed in culture and incubated for 4 d in either the presence or absence of exogenous Wnt5a. The fraction of hair follicles undergoing cystic degeneration over a 4-d incubation period was significantly decreased as a consequence of Wnt5a treatment (Fig. 6A,B). In parallel, hair follicle differentiation-related genes were up-regulated and interfollicular markers were down-modulated in hair follicles isolated from the Wnt5a-treated skins (Fig. 6C). Similar rescue of gene expression defects was observed when purified follicles from the mutant mice, instead of skin explants, were placed in culture and directly treated with Wnt5a (Fig. 6D).

Figure 4. The Wnt5a gene as a primary target of Notch/RBP-Jk-dependent transcription in dermal papilla cells. (A) Cultured dermal papilla cells from (ColI-Cre \(\times\) RBP-Jk\(^{loxP/loxP}\)) mice \((-/-\)) were treated with purified recombinant Wnt5a (200 ng/ml) or BSA control for 24 h. Cells were analyzed in parallel with control cells from (RBP-Jk\(^{loxP/loxP}\)) mice \((+/+\) for levels of Noggin, Fgf7, and Fgf10 expression by real-time RT–PCR. \(\beta\)-Actin mRNA levels were used for internal normalization. (B) Parallel cultures as in A were treated with purified Noggin (200 ng/ml), Fgf7 (200 ng/ml), and Fgf10 (200 ng/ml), followed by real-time RT–PCR analysis of Wnt5a expression. (C) Real-time RT–PCR analysis of Wnt5a expression levels in cultured RBP-Jk\(^{-/-}\) versus RBP-Jk\(^{+/+}\) dermal papilla cells, infected with a retrovirus expressing an activated Notch1–ER fusion protein, after a 24-h 4-OHT treatment (left panel) (Schroeder and Just 2000), or with an activated Notch1-expressing adenovirus (right panel), along with corresponding controls. (D) Primary passage wild-type dermal papilla cells were plated on dishes coated with BSA or purified Jagged 1 protein, in the indicated concentrations of coating solution, for 24 h, followed by real-time RT–PCR analysis of the “canonical” Notch target gene Hey1 and Wnt5a. (E) Cultured dermal papilla cells from mice with \((-/-\)) and without \((+/+\) deletion of the RBP-Jk gene were processed for ChIP assays with an antibody against the proteolytically cleaved active form of Notch1 and purified rabbit IgG as nonimmune control. Real-time PCR of three distinct regions of the Wnt5a gene containing conserved RBP-Jk-binding sites at positions \(-3.87\) kb (RBP-Jk-A), \(-1.63\) kb (RBP-Jk-B), and \(+1.81\) kb (RBP-Jk-C) from the transcription start site (see map above) were performed along with PCR of an upstream region (C) devoid of such sites. Unprecipitated chromatin preparations were similarly analyzed and used as “input DNA” control. The amount of precipitated DNA was calculated relative to the total input chromatin, and was expressed as percentage of the total
Figure 5. Defective hair follicle differentiation as a consequence of Wnt5a deletion in grafted skin and hair follicle reconstitution assays. (A) Full-thickness skin of E17.5 mouse embryos with homozygous deletion of the Wnt5a gene was grafted in parallel with skin of wild-type littermates onto the backs of nude mice. Photographs are representative of three grafted animals at 1 wk after grafting. Results of grafts at 2 and 4 wk are shown in Supplemental Figure 8. (B) Histological analysis of the corresponding skin samples illustrated in A. Hair follicle density, assessed by counting two grafts per group, was 16 ± 4 and 13 ± 2 follicles per millimeter for wild-type and mutant grafts, respectively \( (P < 0.01 \text{ by Student's } t\text{-test}) \). Bar, 160 µm. (C) Immunofluorescence analysis of mutant versus control skin at 7 d after grafting with antibodies against the indicated proteins. Bar, 40 µm. (D) Immunofluorescence analysis of Wnt5a, Noggin, and Fgf10 expression in hair follicle and dermal papillae of mutant versus control skin at 7 d after grafting. Hair follicle boundaries and dermal papillae are indicated by dotted lines and arrows, respectively. Images were taken under the same exposure and capture conditions, and are representative of multiple hair follicles of similar developmental stages in two grafts per condition. Bar, 30 µm. (E) Hair follicle reconstitution assays with primary keratinocytes from EGFP transgenic mice in combination with freshly isolated dermal papilla cells from E18.5 Wnt5a mutant embryos \((-/-)\) versus wild-type littermates \((+/+)\). Stereomicrographs are representative of four independent assays per combination of cells, at 2 wk after grafting. Average number of reconstituted hair follicles was 150–200 in the control grafts, and 20–40 in the grafts with mutant dermal papilla cells. Bars, 400 µm. (F) Histological analysis of hair follicle reconstitution assays described in E. Bar, 100 µm. (G) Immunofluorescence analysis of hair follicle reconstitution assays with antibody against the indicated proteins. Epithelial-mesenchymal borders are indicated by dotted lines. Bar, 40 µm.
FoxN1 as a target and mediator of RBP-Jk/Wnt5a signaling in hair follicles

Wnt5a has been implicated as an activator of noncanonical Wnt signaling, although, in some contexts, it can also induce the canonical pathway (Mikels and Nusse 2006). In the functional rescue experiments above, real-time RT–PCR analysis of isolated hair follicles from cultured skin showed that Wnt5a treatment caused no modulation of canonical Wnt target genes like Lef1, Axin2, or DKK1. In contrast, expression of FoxN1, a keratinocyte-specific transcription factor reported previously to be under Wnt control (Balciunaite et al. 2002), was substantially induced (Fig. 7A). Immunofluorescence analysis confirmed that Wnt5a treatment caused a significant increase of FoxN1 expression in hair follicles of cultured (Coll-Cre × RBP-JkloxP/loxP) skin (Fig. 7B). Levels of FoxN1 expression were similarly analyzed in hair follicle reconstitution assays with dermal papilla cells with or without deletion of the RBP-Jk and Wnt5a genes. The results indicated that sustained FoxN1 expression is under dermal papilla cell control through a RBP-Jk- and Wnt5a-dependent mechanism (Fig. 7C,D).

A number of transcriptional targets of FoxN1 have been reported previously, including Phospholipase C-δ1 (Plcd1) (Janes et al. 2004; Nakamura et al. 2008) and desmocollin 2 (DSC2) (Johns et al. 2005). Similar down-modulation of these genes, as well as trichohyalin, was found in cultured hair follicles with siRNA-mediated knockdown of FoxN1 expression as in hair follicles derived from mice with the RBP-Jk deletion (Fig. 7E,F). Conversely, Wnt5a treatment led to induction of these genes in wild-type hair follicles, with no such induction occurring in similarly treated hair follicles derived from FoxN1 mutant mice (Fig. 7G).

Involvement of the Wnt5a receptors ROR2 and Fzd6 in control of FoxN1 expression

Wnt5a can function through multiple receptors, like the Frizzled family member Fzd6 and ROR2, a transmembrane tyrosine kinase with an extracellular Fzd-like cystein-rich domain (van Amerongen et al. 2008). Real-time RT–PCR analysis showed that expression of both receptors is reduced in hair follicles as a consequence of RBP-Jk or Wnt5a deletion (Supplemental Fig. 9A), and is induced by Wnt5a treatment (Supplemental Fig. 9B), consistent with a positive feedback regulatory mechanism reported in other systems (O’Connell et al. 2010). Functionally, knockdown of ROR2 in hair follicles led to down-modulation of FoxN1 (Supplemental Fig. 9C) and suppressed induction of this gene, as well as its downstream targets, by Wnt5a treatment.
Similar results were observed after Fzd6 knockdown (Supplemental Fig. 9E,F). We complemented the above findings by assessing the consequences of loss of ROR2 function in vivo. Mice with homozygous disruption of the ROR2 gene die at birth due to breathing defects (Takeuchi et al. 2000). Immunofluorescence analysis of the skin of these mice after full-thickness grafting onto nude mice revealed that expression of FoxN1 as well as trichohyalin was reduced consistently in the hair follicles of these mice (Supplemental Fig. 10A,B). However, this was associated with only slight alterations in hair follicle structures [in IRS and/or hair shaft organization] that did not proceed to overt cyst formation, consistent with ROR2 functioning in concert with Fzd6 and possibly other Wnt5a receptors.

**Discussion**

Epithelial–mesenchymal interactions play a key role in organ morphogenesis and homeostasis. Hair follicle development and cycling provide a model paradigm for these interactions (Botchkarev and Paus 2003). In this system, the function of Notch signaling in the epithelial compartment has been studied extensively, while its role in the neighboring inductive mesenchyme has not been investigated. We unveiled an as yet unsuspected mechanism for
maintenance of the hair follicle keratinocyte cell fate by underlying dermal papilla cells through the CSL/RBP-Jk gene, the mediator of “canonical” Notch signaling. Wnt5a functions as a primary CSL effector in these cells, with FoxN1 as one of its targets in keratinocytes.

Significant attention has been given to the role of Notch signaling in control of keratinocyte stem cell potential and differentiation. Deletions of the RBP-Jk (Yamamoto et al. 2003; Blanpain et al. 2006; Demehri and Kopan 2009), Notch1 and Notch2 (Pan et al. 2004), and jagged 1 (Estrach et al. 2006) genes in keratinocytes cause loss of hair follicles, with replacement of these structures with cysts. We found similar alterations in mice with mesenchymal deletion of the RBP-Jk gene. However, comparative analysis of these mice with those with keratinocyte-specific deletion of the same gene pointed to distinct molecular defects, with likely different underlying mechanisms. In mice with the Col1-Cre deletion of RBP-Jk, there was inappropriate hair follicle expression of suprabasal interfollicular keratins (K1 and K10), which were not found in hair follicles of mice with the K5-Cre deletion. While dermal papilla cells (identified by versican, alkaline phosphatase, and Sox2 staining) were normally present in mice with the first genotype, they were hard to find in the others. The impact that loss of RBP-Jk function in keratinocytes has on underlying dermal papilla cells will have to be assessed in future studies. Here, by hair follicle reconstitution assays, we showed that the RBP-Jk gene plays an intrinsic essential role in dermal papilla cells. Deletion of this gene did not compromise cell fate identity, as assessed by a panel of dermal papilla signature genes. Rather, it resulted in decreased expression of diffusible molecules that act as cues to overlying keratinocytes. Analysis of late gestation embryos and mice of early age indicates that dermal papilla cells lacking the RBP-Jk gene are not impaired in their overall ability to contribute to hair follicle morphogenesis, but in subsequent steps required for hair follicle maturation and/or differentiation.

Wnt5a is a specific Wnt family member whose expression during early skin development is limited to dermal papillae (Rendl et al. 2005), with only later expression in the hair follicle compartment [Reddy et al. 2001]. Wnt5a expression in the developing skin is dependent on Shh function, and Wnt5a has been proposed as a mediator of the “second dermal signal” required for hair follicle proliferation downstream from Shh expression in the overlying epithelium [Reddy et al. 2001]. Wnt5a expression has also been connected with the site-specific function of dermal fibroblasts in the body, under transcription control of the distal site-specific homeobox gene HOX13 [Rinn et al. 2008]. Our present findings point to Wnt5a as a likely direct target of Notch/RBP-Jk function in dermal papilla cells, as the promoter and first intronic region of the Wnt5a gene contain RBP-Jk-binding sites to which endogenous Notch1 binds, and which are required for Notch1-dependent promoter activation.

Hair follicle reconstitution assays with dermal papilla cells from mice with mesenchymal RBP-Jk deletion or deletion of the Wnt5a gene point to a common function of the two genes in these cells. In fact, functional rescue experiments indicate that Wnt5a is a primary mediator of RBP-Jk function in this context, with other diffusible molecules involved in dermal papilla signaling, like Noggin and Fgf7/10, being under more indirect RBP-Jk/Wnt5a control. Interestingly, the demonstrated ability of Wnt5a to induce JNK/c-Jun activity [Pukrop et al. 2006; Nomachi et al. 2008] could underlie its regulation of Fgf7 expression, which was found to be under c-Jun control in the skin [Szabowski et al. 2000]. Future studies will have to assess to what extent this and other factors synergize or antagonize with Wnt5a function in this context.

Downstream from RBP-Jk/Wnt5a, we found that expression of the FoxN1 transcription factor is compromised in the keratinocyte compartment. FoxN1 plays an important role in control of hair follicle keratinocyte differentiation [Mecklenburg et al. 2001]. Genes under FoxN1 control are down-modulated as a consequence of the RBP-Jk/Wnt5a deletions, and are up-regulated in hair follicles by Wnt5a treatment in a FoxN1-dependent manner, implicating this transcription factor as a mediator of RBP-Jk/Wnt5a activity in hair follicles. FoxN1 has also been implicated in providing the necessary cues for specific transfer of pigment from melanocytes to keratinocytes of the hair shaft cortex [Weiner et al. 2007]. Reduced pigment deposition in the hair cortex was found in hair follicles of RBP-Jk or Wnt5a mutant mice or grafts, as well as in hair follicle reconstitution assays with corresponding dermal papilla cells [Supplemental Fig. 11], consistent with our other findings of decreased FoxN1 expression and activity.

Little is known about control of this gene. In thymic epithelia, FoxN1 expression is positively modulated by Wnt4 and/or Wnt5b [Baliunaite et al. 2002], whereas it is negatively regulated by ERK1/2 activity in keratinocytes [Baxter and Brissette 2002; Mandinova et al. 2009]. The data we presented indicate that FoxN1 expression in hair follicles is under positive Wnt5a control, as it is down-regulated in the skin of mice with deletion of the Wnt5a gene and in reconstituted hair follicles with Wnt5a-null dermal papilla cells, while it is induced in wild-type hair follicles by Wnt5a treatment. Wnt5a has been most frequently linked to “noncanonical” Wnt signaling, through multiple receptors leading to activation of calcium- and Rho/JNK-dependent pathways [van Amerongen et al. 2008]. Knockdown experiments indicate that two of these receptors, Fzd6 [Wang and Nathans 2007] and ROR2 [Oishi et al. 1999], participate in control of FoxN1 expression by Wnt5a. Studies from other systems indicate that these proteins have the potential to functionally and physically interact [Yamamoto et al. 2008]. Little or nothing is known about the signaling function of these receptors in keratinocytes, and an exciting line of future work will involve the elucidation of their role in the regulatory axis that we uncovered.

Materials and methods

Mice

RBP-Jk<sup>loxp/loxp</sup> mice [Han et al. 2002] were crossed with transgenic mice expressing a Cre recombinase under control of the
promoter/enhancer unit of the α2 chain of the collagen type 1 gene (Florin et al. 2004) or the keratin 5 promoter (Berton et al. 2000) to generate [ColII-Cre × RBP-Jk^loxP/loxP^] and [K5-Cre × RBP-Jk^loxP/loxP^] mice, respectively. [ColII-Cre × RBP-Jk^loxP/loxP^] mice were also crossed into the ROSA 26R reporter strain, an in vivo reporter of Cre activity (Soriano 1999). Transgenic mice with EGF expression driven by a ubiquitous β-actin promoter (CS7BL/6-Tg[β-actinEGFP]) were described (Okabe et al. 1997). B6;129S7-Wnt5atm1Amc/J mice (Yamaguchi et al. 1999) were described previously (Oishi et al. 1999). All mouse work except for ROR2 mice was performed according to the Swiss guidelines and regulations for the care and use of laboratory animals. For ROR2 mice, work was performed according to International Animal Care and Use Committee guidelines and regulations of Tulane University.

Skin cell preparation and purification

The whole procedure is illustrated in Supplemental Figure 2A, along with a description of various markers to assess the effectiveness of the separation procedure. For separation of interfollicular epidermis from the dermis, back skin from P1 mice was treated with 1% Dispase II (Roche) overnight at 4°C. For hair follicle separation, the dermis was further digested by 0.3% type I collagenase (Sigma) in HBSS (Gibco) for 1 h at 37°C. After centrifugation in a tabletop centrifuge (1000 rpm for 5 min, followed by 400 rpm for 5 min twice), the sedimented material was subjected to two rounds of density gradient centrifugation—first in 10% Percoll (d = 1.01), and the second in 33% Percoll (d = 1.06)—collecting the bottom fraction each time. Microscopic examination showed >90% hair follicle purification by this approach. The recovered hair follicles were further dissociated into cell suspension by mechanical disruption (20 strokes through a 23-gauge and then a 26-gauge needle). Cells were then centrifuged and resuspended into 500 μL of culture medium and incubated with rat anti-mouse c-Kit (eBioscience) and rat anti-mouse CD133 (eBioscience) antibodies (at a 1:100 dilution) for 1 h at room temperature. Cells were then washed twice in culture medium, followed by incubation with goat anti-rat microbeads (Miltenyi Biotech) for 30 min at 4°C. Magnetic cell sorting, using a MACS midi separator and LS columns (Miltenyi Biotech), was then used to deplete cells positive for c-Kit (melanocytes) and CD34-negative fraction (dermal fibroblasts). Cells recovered after this depletion procedure were further incubated with rat anti-mouse CD133 antibodies, and magnetic cell sorting was used to separate them in into a CD133-positive fraction (dermal papilla cells) and CD133-negative fraction (dermal fibroblasts). For hair follicle reconstitution and dermal papilla cell culture assays, dermal papilla cells were directly isolated from dermal cell preparations by magnetic cell sorting with anti-CD133 antibodies.

For the isolation of mature hair follicles [from P10 mice] and cysts [from 3-wk-old mice], the total skin was at first digested in 1% type I collagenase for 2 h at 37°C. Hair follicles and cysts were then purified by the two-step density gradient centrifugation described above. Hair follicles with relatively normal versus abnormal structures from the [ColII-Cre × RBP-Jk^loxP/loxP^] mice, as well as the cysts, were further manually isolated under a dissecting stereomicroscope (Leica MZ 16F).

Cell culture, viral infection, and plasmids

For culturing, the sorted dermal papilla cells from newborn mice were plated at an initial density of 2 × 10⁵ per 10-cm dish and cultured in DMEM (Gibco) and 10% fetal bovine serum [FBS] (Sigma) for 3 d (to 60%–70% confluence) prior to further analysis. For in vitro deletion of the RBP-Jk gene, cultured dermal papilla cells from [RBP-Jk^loxP/loxP^] mice were infected with an adenovirus expressing activated Notch1 or the Cre recombinase together with GFP, in parallel with a control virus expressing GFP alone, at a multiplicity of infection of 100, which resulted in >99% of GFP-positive infected cells. Adenoviruses and our conditions for adenovirus production and infection were as reported previously (Rangarajan et al. 2001). As an alternative approach, dermal papilla cells were infected with a retrovirus expressing constitutively active Notch1 fused to an estrogen receptor domain (rNERT-neo) (Schroeder and Just 2000) and empty vector control (rneo), followed 4 d later by treatment with 1 μM 4-hydroxy-moxifen (OHT) for 48 h to induce nuclear translocation and activation of Notch-dependent transcription.

For treatment of dermal papilla cells with Jagged 1 ligand, 2 mL of goat anti-rat IgG (R&D Systems) (diluted in PBS) were added to nontreated 10-cm plastic dishes at a concentration of 10 μg/mL and incubated for 30 min at 37°C. Dishes were washed five times with PBS, followed by blocking with 2 mL of 2% BSA in PBS for 1 h at 4°C. The blocking medium was then replaced with 2 mL of ligand solution. For this, recombinant rat Jagged 1 protein (R&D Systems) was diluted in 2% BSA in PBS to a final concentration of 10 or 20 μg/mL. Dishes were incubated with the Jagged 1 ligand solution for 2 h at 37°C, followed by five washes with PBS. Freshly isolated dermal papilla cells from P1 wild-type mice were then seeded at a concentration of 200,000 cells per dish and cultured in DMEM (Gibco) and 10% FBS [Sigma] for 24 h.

For Wnt5a promoter activity assays, the mouse Wnt5a promoter and first intronic region (3.943 kb, spanning from −1.66 to +2.29 kb relative to the transcription start site) was amplified by PCR of genomic DNA using the following primers: 5′-CTTT CTACCCCTTGAAGCTTG-3′ (Forward) and 5′-TTGAAGGACT GGGACTCAGA-3′ (Reverse). The product of the PCR was cloned into the pGL4.1 vector [Promega] at the BglII/HindIII sites to generate the Wnt5a-luc reporter plasmid. For in vitro deletion of the Wnt5a promoter (Berton et al. 2001), two RBP-Jk-binding sites, located at positions −1.64 and +1.81 kb relative to the transcription start site, the Wnt5a-luc plasmid DNA was digested by BglII/Smal to remove 368 base pairs [bp] of 5′ upstream sequence, enclosing the first site, and by Sbal/MluI, for an internal deletion of 233 bp (+1.78 to +2.01 kb), enclosing the second site. The Wnt5a-luc plasmid and its Wnt51ΔRBP-Jk-luc derivative were cotransfected into Saos-2 cells [in 12-well plates; 1 μg of DNA per well] together with an expression vector for activated Notch1 [2 μg of DNA per well] (Rangarajan et al. 2001) or empty vector control and the renilla phRL-TK reporter [Promega] for internal normalization. Cells were harvested 48 h after transfection, and were assayed with Dual Luciferase Reporter Assay System [Promega] according to the manufacturer’s instructions. All experiments were performed in triplicate.

Skin and hair follicle organ cultures

Full-thickness back skin from mice of the indicated age and background was placed in semisolid agarose medium [Sigma] in DMEM + 10% FBS, with the upper surface of

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Hu et al.

the skin exposed to the air interface. For Wnt5a treatment experiments, two halves of the same skin samples were placed in culture in separate wells, followed, after 24 h of adaptation to culture conditions, by transfer to fresh semisolid medium containing either Wnt5a (200 ng/mL) or BSA alone. Samples were further transferred to freshly supplemented semisolid medium 48 h later. Recombinant Wnt5a protein was purchased from a commercial source (R&D Systems) and dissolved in 1% BSA-PBS to achieve 100 μg/mL stock solution.

For hair follicle cultures, hair follicles physically separated from the skin of P0 mice of the indicated background were placed onto 12 wells with organotypic inserts (Millipore) (~1,000 follicles per well), with 2 mL of DMEM +10% FBS per well and 0.5 mL of the same added to the insert. Samples were adapted to culture conditions for 24 h prior to treatment with Wnt5a (200 ng/mL final concentration) or BSA alone.

For siRNA-mediated knockdown experiments, P0 hair follicles, isolated as described above, were resuspended in HBSS (without Ca²⁺ and Mg²⁺) (Gibco) and transferred to six-well plates (Ultra-Low attachment microplates, Corning). About 1000 hair follicles were added per well, in 1.8 mL of culture medium, and kept on ice. siRNA transfection was performed with INTERFERin reagent (Polyplus) according to the manufacturer’s recommendations. For each sample, 20 μL of siRNAs (from 10 μM stock) was added to 80 μL of 5% glucose and mixed with 12 μL of INTERFERin diluted into 88 μL of 5% glucose by gentle vortex for 10 sec. The mix was then incubated for 10 min at room temperature before adding to the hair follicle HBSS suspension, with a final siRNA concentration of 100 nM. The transfected hair follicle suspension was kept for 24 h at 4°C, followed by replacing HBSS with 2 mL of low-calcium mouse keratinocyte culture medium (Dotto et al. 1988) and culturing for 4 d at 34°C in 7.5% CO₂ condition. Medium was changed after 48 h (by hair follicle centrifugation at 450 rpm for 5 min at 4°C). For Wnt5a treatment, culture medium was replaced for the last 24 h of the experiment with 2 mL of fresh culture medium containing recombinant Wnt5a (200 ng/mL final concentration) or BSA alone (0.05%).

Hair follicle reconstitution assays

For each injection, 1 × 10⁶ freshly isolated CD133⁺ dermal papilla cells from P1 mice (for [ColI-Cre × RBP-JkloxP/loxP] and [RBP-JkloxP/loxP]) and E18.5 embryos (for Wnt5a control and knockout mice) were admixed with 0.5 × 10⁶ freshly isolated P1 keratinocytes from actin-EGFP transgenic mice. The mixed cells were resuspended in 50 μL of DMEM +10% FBS, and were injected at the dermal-hypodermal junction into NMRI nude mice as described previously (Zheng et al. 2005). Each mouse was injected in parallel, on the left and right flank, with keratinocytes admixed with either [ColI-Cre × RBP-JkloxP/loxP] or [RBP-JkloxP/loxP] dermal papilla cells, or with Wnt5a−/− versus Wnt5a+/+ dermal papilla cells. Eight to 12 mice were injected, being sacrificed at various times after injection as indicated.

Full-thickness skin grafting

Back skin of Wnt5a−/− and littermate wild-type control mice at E17.5, and (ColI-Cre × RBP-JkloxP/loxP) and (RBP-JkloxP/loxP) at P0, were cut into 5 × 5-mm pieces, washed thoroughly in PBS, and placed onto the recipient graft bed of nude mice [prepared by removing epidermis and superficial dermis with curved surgical scissors]. To minimize individual animal variability, control and mutant skin were grafted in parallel on the left and right back flank, respectively, of each recipient mouse. Grafts were covered with a 16 × 57-mm wound bandage and further secured with adhesive tape. Bandage and adhesive tape were changed every day. Three days later, grafts were exposed to air until harvest.

Immunofluorescence staining and confocal analysis

The following primary antibodies were used: AE13 and AE15 (1:25; kindly provided by Dr. T.T. Sun), CD133 (1:200; R&D Systems), Cre (1:200; Novagen), cytokeratin 1 (1:500; Covance), cytokeratin 10 (1:500; Covance), cytokeratin 14 (1:1000; Covance), cytokeratin 6 (1:1000; Covance), FoxN1 (1:50; Santa Cruz Biotechnology), GFP (1:500; Abcam), Sox2 (1:50; R&D Systems), Versican (1:200; Chemicon), and Wnt5a (1:100; R&D Systems). The secondary antibodies were conjugated with Alexa Fluor 488 or 568 (Invitrogen). Counterstaining of nuclei was performed with DAPI.

Embryos or newborn pups were cryosectioned in sagittal direction at 20 μm at the midline of the body. Sections were fixed in precooled acetone for 20 min at −20°C, permuted for 20 min in PBST (PBS with 0.1% Triton X-100), and incubated in blocking solution (PBS with 2.5% donkey serum, 2.5% goat serum, 0.5% cold water fish gelatin, 0.5% BSA) for 60 min. Incubation with primary antibodies was for either 2 h at room temperature or 12 h at 4°C. For immunostaining of Cre and CD133, enzymatically dissociated total dermal cells were fixed in 4% PFA for 20 min on ice and then stained with the above protocol.

Images were acquired with a Zeiss LSM510 meta laser-scanning microscope (Carl Zeiss MicroImaging). For histological sections and whole-mount samples, Z stacks (1 μm) of six to eight and 20 planes, respectively, were acquired.

Quantitative real-time RT–PCR, immunodetection, and ChIP techniques

Conditions for real-time RT–PCR and immunoblotting were as described previously (Lefort et al. 2007). Immunoblotting was performed with the following antibodies: Wnt5a (R&D Systems), Fg10 (Santa Cruz Biotechnology), Noggin (Santa Cruz Biotechnology), and β-actin (Santa Cruz Biotechnology). ChIP assays were performed as described previously [Lefort et al. 2007]. The list of gene-specific primers is provided in Supplemental Table S1.

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Control of hair follicle cell fate by underlying mesenchyme through a CSL–Wnt5a–FoxN1 regulatory axis

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