

RESEARCH COMMUNICATION

A dermal *HOX* transcriptional program regulates site-specific epidermal fate

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Reciprocal epithelial–mesenchymal interactions shape site-specific development of skin. Here we show that site-specific *HOX* expression in fibroblasts is cell-autonomous and epigenetically maintained. The distal-specific gene *HOXA13* is continually required to maintain the distal-specific transcriptional program in adult fibroblasts, including expression of *WNT5A*, a morphogen required for distal development. The ability of distal fibroblasts to induce epidermal keratin 9, a distal-specific gene, is abrogated by depletion of *HOXA13*, but rescued by addition of *WNT5A*. Thus, maintenance of appropriate *HOX* transcriptional program in adult fibroblasts may serve as a source of positional memory to differentially pattern the epithelia during homeostasis and regeneration.

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Epithelial tissues such as skin demonstrate remarkable anatomic diversity in their structure and function. For instance, while long terminal hairs decorate the scalp, palmo-plantar skin lacks hairs but possesses thickened epidermal barriers for mechanical stress. These anatomic differences lead to many body site-specific manifestations of diseases and guide their appropriate treatments (Bologna et al. 2003). Because the epidermis is continually turned over, the apparent stability of site-specific features raises the fundamental question of how

positional identity is acquired and maintained in the skin. Classic heterotypic recombination experiments in chick suggested that a primary dermal signal may dictate the positional identity of epithelial differentiation (Dhouailly 1984). We and others have used a genomic approach to examine the anatomic diversity of gene expression of cell types in the dermis and identified fibroblasts as a cell type that demonstrates consistent and large-scale differences of the expression of cell–cell signaling proteins in a site-specific manner (Chang et al. 2002; Chi et al. 2003; Rinn et al. 2006). Systematic comparison of the global transcriptional programs of fibroblasts from anatomic sites that finely map the body suggest that fibroblasts are differentiated based on their position along three anatomic divisions: anterior–posterior, proximal–distal, and dermal–nondermal (Rinn et al. 2006). Moreover, primary adult fibroblasts retained many features of the embryonic pattern of expression of *HOX* genes (Chang et al. 2002; Rinn et al. 2006), a family of homeodomain transcription factors that act to specify positional identity during development. These data support the idea, first proposed by Chuong (1993), that a “*HOX* code” may dictate the positional identity of skin and thus influence site-specific epidermal differentiation. However, fundamental elements of this hypothesis, such as the stability of *HOX* expression, gene regulatory function, and inductive activities governed by *HOX* genes in adult fibroblasts remain poorly understood. Here we show that expression of *HOXA13* is required and the transcriptional targets are sufficient to maintain epidermal inductive properties of distal fibroblasts.

Results and Discussion

To test the hypothesis that ongoing *HOX* expression in dermal fibroblasts contributes to skin positional identity, we first examined the stability of *HOX* gene expression. Microarray analysis of serially passaged primary fibroblasts derived from foot or thigh showed that expression levels of all *HOX* genes are stable for at least 35 passages in vitro (Fig. 1A). We next tested whether the site-specific gene expression patterns were influenced by secreted factors in the media by culturing foot fibroblasts in conditioned media from thigh fibroblasts and vice versa (proximal vs. distal), followed by microarray analysis. Although the addition of conditioned media from any source predictably altered fibroblast gene expression, unsupervised hierarchical clustering of the gene expression patterns show a clear separation of thigh fibroblasts from foot fibroblasts, irrespective of the source of conditioned media (Fig. 1B). Their repertoire of *HOX* genes also remained distinctive and site-specific. Reciprocal, heterotypic conditioned media treatment of foreskin and lung fibroblasts also showed that their site-specific gene expression programs were essentially unaltered (Supplemental Fig. 1). These experiments suggest that secreted factors from adult fibroblasts are unable to reprogram site-specific gene expression of other fibroblasts, at least in the assay conditions used.

To address whether site-specific gene expression of fibroblasts may be reprogrammed by coculture and physical contact between heterotypic fibroblasts, we adapted a genetically encoded method of RNA labeling and capture (Fig. 1C–E; Cleary et al. 2005). Briefly, introduction of

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Rinn et al.

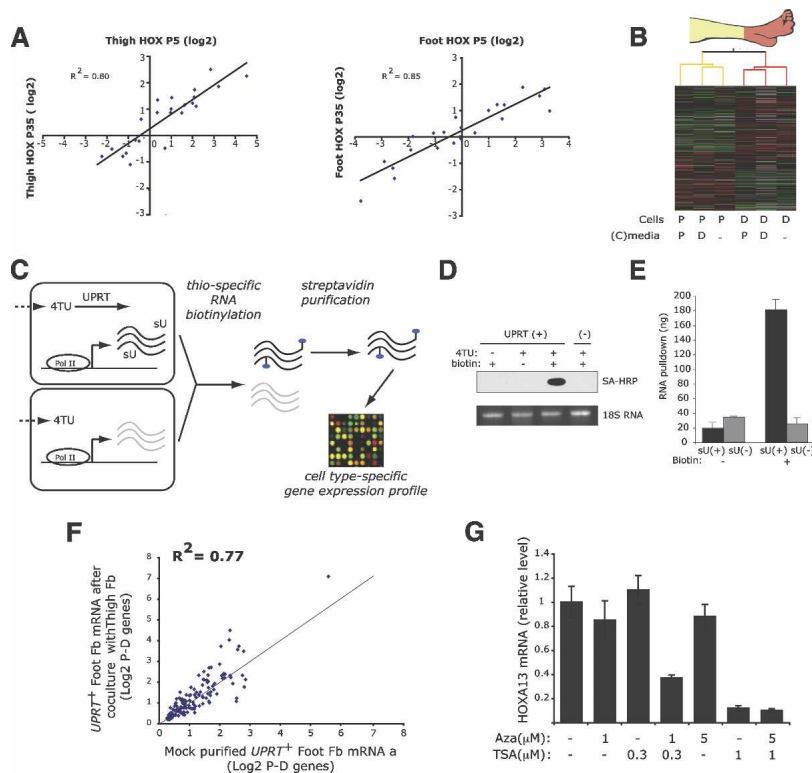


Figure 1. Stability of site-specific gene expression in fibroblasts. (A) Stability of *HOX* gene expression after serial passage. The expression level of *HOX* genes plotted for passage 5 (X-axis) and passage 35 (Y-axis) for thigh (left) and foot fibroblasts (right). (B) Stability of site-specific gene expression patterns with heterotypic conditioned media. Unsupervised hierarchical clustering of thigh fibroblasts (P) without coculture (-), cultured with their self-conditioned media (P, P) or with conditioned media from foot fibroblasts (P, D) and vice versa for foot fibroblasts. Expression level of each gene above (red) or below (green) the global median (black) is denoted by the color scale (fourfold to 0.25-fold on linear scale or +2 to -2 on \log_2 scale). (C) UPRT-mediated RNA capture. Only UPRT⁺ cells can incorporate 4TU into thio-uracil (sU) in RNA, enabling retrieval of cell-type-specific RNA by thio-directed RNA biotinylation and purification. (D) UPRT-mediated RNA biotinylation. The 18S RNA band is shown by Northern blot probed with streptavidin-horseradish peroxidase (top) or ethidium bromide (bottom). (E) Verification of cell-type-specific RNA capture from 1:1 mixture of UPRT⁺ and UPRT⁻ RNA. In the absence of the biotinylation reaction, little RNA was retrieved from either population. With biotin-mediated RNA pull down, RNA from UPRT⁺ cells was specifically enriched by eightfold to 10-fold compared with that of the mixed population. (F) Little change in the expression of genes diagnostic of proximal-distal limb origin in human foot fibroblasts after coculture with thigh fibroblasts. (G) Synergistic effects of trichostatin A (TSA) and Aza-C (Aza) on *HOXA13* mRNA expression.

the *Toxoplasma gondii* UPRT gene allows all RNAs in the UPRT⁺ cells to be specifically biotinylated and captured from a pool of UPRT⁺ and UPRT⁻ cells; we could therefore study the cell-type-specific consequences of coculture on gene expression without artifacts introduced by cell suspension or cell sorting. Using UPRT-mediated RNA capture and microarray analysis, we found that the global gene expression profiles of UPRT⁺ foot fibroblasts are not appreciably affected by coculture with thigh fibroblasts when compared with UPRT⁻ foot fibroblasts (Fig. 1F). This result suggests that the majority of the site-specific gene expression in fibroblasts is controlled by a cell-autonomous mechanism. Importantly, site-specific expression of *HOX* genes in fibroblasts can be altered by treatment with trichostatin, a histone deacetylase inhibitor, and is potentiated by Aza-C, a DNA methylation inhibitor (Fig. 1G). These results are consistent with data indicating that site-specific *HOX* expression is

epigenetically maintained by chromatin modifications in fibroblasts (Bernstein et al. 2005; Rinn et al. 2007).

We next addressed the requirement of ongoing *HOX* activity for site-specific gene expression in fibroblasts. By comparing global gene expression profiles of primary fibroblasts from 43 anatomic sites, we previously defined a “distal gene expression neighborhood” that comprises 52 probes representing 44 unique genes specifically induced in cells from distal anatomic sites such as fingers, feet, and foreskin (Fig. 2A). *HOXA13* is member of this distal neighborhood; interestingly, *HOXA13* mutations in human patients produce malformations in these exact same anatomic sites and is termed hand-foot-genital syndrome (Mortlock and Innis 1997). We hypothesized that *HOXA13* positively regulates transcriptional activation of this distal module, as these 44 genes are highly correlated in their expression across all 43 fibroblast sites. We also suspected that *HOXA13* may negatively regulate the transcription of genes that are anti-correlated to the express level of *HOXA13* across our fibroblast compendium (Fig. 2A, bottom right). Interestingly, the most highly anti-correlated genes to *HOXA13* include *HOXB2*, *HOXB4*, *HOXB5*, *HOXB6*, and *HOXB7*, suggesting a negative interaction between two *HOX* loci located on separate chromosomes, similar to what has been seen for the *HOXC* and *HOXD* loci (Rinn et al. 2007). siRNA-mediated depletion of *HOXA13* in foreskin fibroblasts followed by microarray analysis showed that almost all genes in the distal neighborhood are substantially reduced in expression upon acute loss of *HOXA13* ($P < 0.00011$, hypergeometric distribution) (Fig. 2B,C; Supplemental Table 1). The 32 genes anti-correlated with *HOXA13* did not exhibit coordinate regulation upon depletion of *HOXA13*; however, *HOXB2*, *HOXB5*, and *HOXB6* showed a trend toward increased expression. These data suggest that

ongoing *HOXA13* expression is required for the activation of a distal-specific pattern of gene expression in fibroblasts.

To determine which of these transcriptional changes are directly regulated by *HOXA13*, we performed chromatin immunoprecipitation (ChIP) of *HOXA13*, followed by hybridization to microarrays tiling over 30,000 human promoters (“ChIP-Chip” analysis) (Fig. 2C; Supplemental Table 2). The promoters of seven of the 44 genes in the distal neighborhood were occupied by *HOXA13*, which is significantly more enriched than expectation by chance alone ($P < 1 \times 10^{-3}$, hypergeometric distribution). *HOXA13*-occupied promoters in fibroblasts include *HOXA13* itself; *HOXB6* the most anti-correlated gene to *HOXA13*; *BMP2*, a *HOXA13* target gene important in distal limb development (Knosp et al. 2004); and *MAB21L1*, a cell fate determining protein. Together, these results identify both direct and indirect targets of

HOXA13 in the distal-specific gene expression program of adult human fibroblasts (Supplemental Tables 2, 3).

To begin to understand how *HOX*-dependent gene expression in fibroblasts may impact epithelial–mesenchymal interactions, we focused our analysis on *WNT5A*, a member of the distal neighborhood whose expression requires ongoing *HOXA13* function, but is not a direct transcriptional target of *HOXA13*, at least for the genomic region of *WNT5A* promoter that was interrogated. *WNT5A* is a member of the WNT family of extracellular cell–cell signaling proteins, and it has been considered an atypical WNT because it can either activate or inhibit a canonical WNT signaling cascade, depending on the presence of its coreceptor ROR2 kinase (Mikels and Nusse 2006). Importantly, murine *Wnt5a* is highly expressed in distal limbs, external genitalia, and distal snout during development, and deletion of *Wnt5a* in mice leads to distal truncation of all these structures (Yamaguchi et al. 1999). Despite the clear evidence implicating *Wnt5a* as a mediator distal outgrowth, the mechanism that specifies distal-specific expression of *Wnt5a* has not been identified. Our microarray analysis thus provides the first experimental evidence that *HoxA13* may activate *Wnt5a* expression, leading to distal patterning. To confirm this hypothesis, first we performed whole-mount in situ hybridization of *HoxA13* and *Wnt5a* during murine development and showed that these two genes are indeed spatially and temporally coexpressed in the distal limb bud (Fig. 3A). Second, we examined the effect of *HoxA13* loss of function on *Wnt5a* expression by using *HoxA13*^{GFP/GFP} mice, where a *GFP* sequence was inserted into a *HoxA13* coding sequence, thereby disrupting *HoxA13* function (Stadler et al. 2001). *Wnt5a* mRNA was detected in the cartilage condensate, scattered cells in the dermis, and in the epidermis of wild-type distal embryonic limb buds; this pattern was similar to the *HoxA13* expression domain, as visualized by GFP expression in *HoxA13*^{GFP/GFP} limb buds (Fig. 3B). Importantly, *Wnt5a* expression was substantially reduced in *HoxA13*^{GFP/GFP} limb buds, indicating that *HoxA13* is required for *Wnt5a* expression (Fig. 3B). In contrast, microarray expression of *HOXA13*-depleted fibroblasts showed a 2.5-fold induction of WNT receptor *FZD8*, and in situ hybridization confirmed the overexpression of murine *FZD8* mRNA in *HoxA13*^{GFP/GFP} limb buds (Fig. 3B). This finding suggests that loss of *Wnt5a* expression in *HoxA13* mutant limbs is unlikely due to cell death or general tissue dysfunction, but is more likely a specific gene regulatory function of *HoxA13*.

Finally, we examined whether *HOXA13*-induced expression of *WNT5A* in adult human fibroblasts continue to provide a distal-specific function. In postnatal human skin from an anatomically distal site, *HOXA13* protein is expressed in the dermis but not in the epidermis (Rinn et al. 2006). Previous studies have shown that coculture

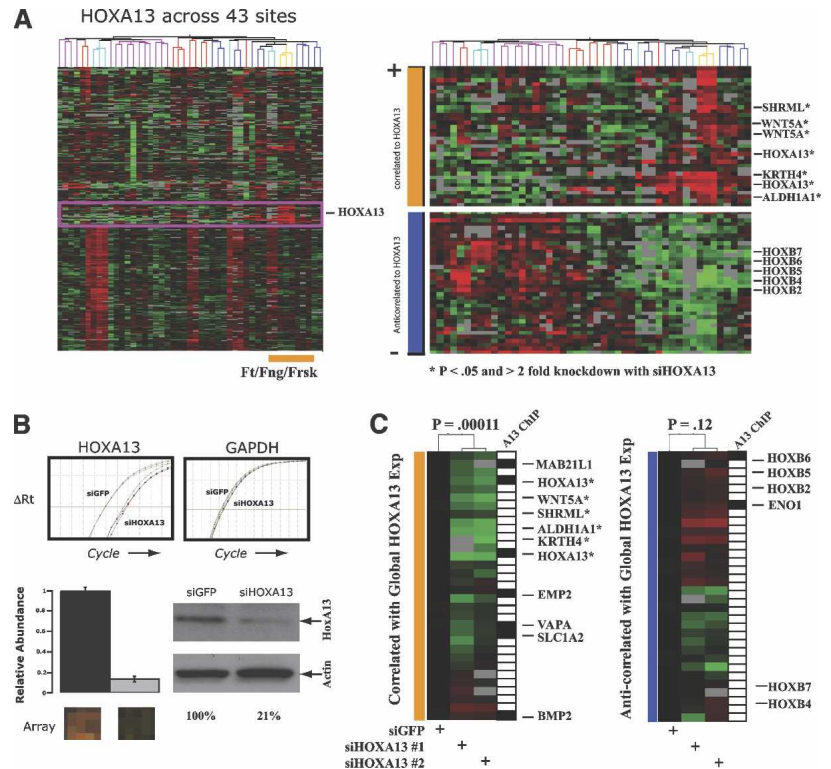


Figure 2. HOXA13 regulates a distal gene expression program. (A, top right) Identification of genes most similar in expression pattern to HOXA13 across fibroblasts from 43 anatomic sites, which comprise a distal-specific neighborhood. (Bottom right) We also identified a set of genes that show the most opposite pattern of expression to HOXA13. (B) siRNA-mediated depletion of HOXA13 as measured by qRT-PCR (top), microarray (bottom left), and immunoblot (bottom right). (C) Effect of HOXA13 depletion on distal gene expression. Expression level of each gene above (red) or below (green) the average mock siRNA treatment (black) is denoted by the color scale (fourfold to 0.25-fold on linear scale or +2 to -2 on log₂ scale). Genes differing significantly (twofold and $P < 0.05$) are denoted by asterisk. The significance of the entire distal neighborhood being coordinately repressed or induced is 0.00011 versus 0.12 (hypergeometric distribution), respectively. Direct binding of HOXA13 to a given promoter as determined by ChIP-chip is denoted by a black box.

of primary dermal fibroblast from palms or soles can reprogram keratinocytes from other sites to express acral-specific differentiation genes, such as keratin 9 (K9) (Fig. 4A; Yamaguchi et al. 1999). Mutations in K9 cause epidermolytic palmoplantar keratoderma (Reis et al. 1994), underscoring the importance of proper regulation of K9. Therefore, we tested whether such a distal-specific inductive function involved the HOXA13–WNT5A pathway. Consistent with previous reports (Yamaguchi et al. 1999), we found that plantar fibroblasts could induce K9 expression in cocultured keratinocytes, albeit a modest induction compared with K9 level in vivo. This effect required the coculture of fibroblasts and keratinocytes, and is more efficient using fibroblasts of palmo-plantar origin (Fig. 4B). This inductive effect appears to be regulated at the transcriptional level as induction of K9 mRNA paralleled protein accumulation (Fig. 4C). Depletion of HOXA13 in plantar fibroblasts before coculture, but not treatment with control siRNAs against GFP or HOXB4, substantially reduced the ability of these cells to induce epidermal K9 transcription, and this defect can be fully rescued by addition of recombinant purified WNT5A protein to the coculture (Fig. 4C). In fact, direct addition of WNT5A protein to keratinocytes in the ab-

Rinn et al.

sence of cocultured fibroblasts could also induce K9 transcription (Fig. 4C), and this effect occurred without perturbing the general differentiation program of keratinocytes (Supplemental Fig. 2). Microarray analysis of the transcriptional response of keratinocytes to WNT5A confirmed that WNT5A predominantly regulates genes involved in cell differentiation (Supplemental Table 4). These results suggest that HOXA13 control of WNT5A in fibroblasts is one of the mechanisms that can regulate site-specific epidermal differentiation.

Here we provide several lines of evidence supporting a fibroblast *HOX* code that contributes to the positional identity of skin. Consistent with heterotopic epithelial-mesenchymal recombination experiments, where the mesenchyme can reprogram heterotopic epithelia, we showed that the positional-specific *HOX* code in fibroblasts is stably maintained in the face of heterotypic signals. Ongoing HOXA13 activity is required to maintain a distal-specific gene expression program in adult fibroblasts. As illustrated by WNT5A, the same HOXA13-dependent gene is used to pattern distal outgrowth early in development and can be used again to specify distal epidermal differentiation in adulthood (Fig. 4D). Our analysis also gave some of the first insights into the human HOXA13 transcriptional networks in fibroblasts, revealing the potential autoregulation of *HOXA13*, *trans*-repression of *HOXB6*, and continued regulation of embryonic target genes such as *BMP2*, which is required for footpad epidermal differentiation (Plikus et al. 2004).

We found that the HOXA13 protein occupied the *HOXB6* promoter, but not the *WNT5A* promoter. Inter-

estingly, HoxB5 and HoxB6 demonstrate an antagonistic pattern of expression as *Wnt5a* expression in lung development (Sakiyama et al. 2000). Therefore, we speculate that HOXA13 activation of WNT5A may occur directly via more distant enhancer sequences or indirectly via HOXB6. *Wnt5a* expression is substantially reduced, but not completely absent in *HoxA13* knockout limb. *Wnt5a* is also expressed more proximally than the *HoxA13* expression domain, suggesting that other transcription factors also induce *Wnt5a* expression. These additional layers of regulation may explain the more severe distal truncation phenotype in *Wnt5a* knockouts.

Previous studies have shown that DKK1, an antagonist of canonical Wnt signaling, is also induced in palmoplantar fibroblasts (Yamaguchi et al. 2004). The combination of DKK1, WNT5A, and repression of FZD8 in palmoplantar skin could block canonical WNT-dependent appendage formation and promote site-specific epidermal differentiation, such as induction of K9. The specialized fibroblasts that support hair follicle differentiation and cycling, termed dermal papilla cells, control the site-specific identity of the hairs and also express *HOX* genes (Jahoda 1992; Rendl et al. 2005), suggesting a possible role of *HOX* genes in patterning epidermal appendage development.

Because the proper position-specific expression of *HOX* genes would appear to have critical importance in dictating skin positional identity, mechanisms that ensure the fidelity of *HOX* expression in skin should be investigated in future studies.

In summary, human adult fibroblasts maintain features of the embryonic *HOX* pattern over decades in vivo and over numerous cell divisions in vitro, which suggests that they are a valuable system to investigate the mechanisms of transcriptional memory. The maintenance of region-specific *HOX* gene expression in adult fibroblasts may serve as a source of positional memory for skin during homeostasis and regeneration. These data are consonant with emerging evidence in other model organisms that *HOX* genes not only orchestrate segmental patterning in development, but continue to act as micromanagers of differentiation in subsequent developmental stages (Akam 1998). Whether ectopic expression of *HOX* genes in adult skin may suffice to reprogram positional identity will be examined in future studies. Finally, additional cell types likely also contribute to skin positional identity, and their roles also await clarification in future studies.

Materials and methods

Heterotypic cell interactions

Primary fibroblasts (Rinn et al. 2006) and *UPRT*-mediated RNA capture (Cleary et al. 2005) were as described. Media conditioned for 4 d were applied to fresh cells for 4 d. Cells were treated with 5-AzaC for 5 d, replacing the drug every 48 h, or with trichostatin A for 48 h.

HOXA13 transcriptional network

After siRNA-mediated depletion of HOXA13, RNA was isolated and hybridized to Stanford cDNA microarrays (Rinn et al. 2006). ChIP using HOXA13 antibody (Knosp et al. 2004) was hybridized to human HG18 promoter tiling arrays (Nimblegen, Inc.). Candidate ChIP peaks of different enrichment values were validated by ChIP followed by qPCR (Supplemental Fig. 2). Candidate HOXA13 targets were compared with genes altered by HOXA13 RNAi or HOXA13 overexpression (Williams et al. 2005), yielding a HOXA13 transcriptional network supported by independent lines of evidence. In situ hybridization of wild-type and *HOXA13^{GFP/GFP}* mice (Stadler et al. 2001) were as described (Albrecht et al. 1997). Array data are available at GEO (GSE9922) and Stanford Microarray Database.

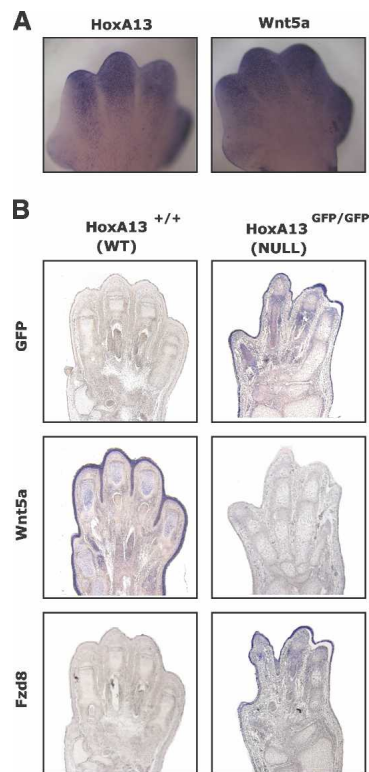


Figure 3. HoxA13 regulates *Wnt5a* expression in vivo. (A) Whole-mount in situ hybridization of E10.5 mouse limb buds for *HoxA13* or *Wnt5a* mRNA expression. (B) Expression of *GFP*, *Wnt5a*, or *Fzd8* in wild-type or *HoxA13^{GFP/GFP}* (null) E16.5 limb bud sections.

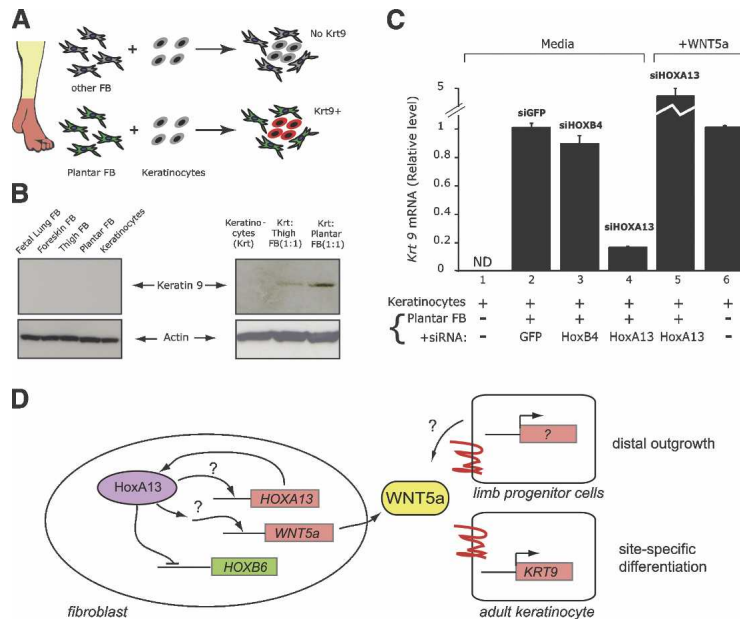


Figure 4. HOXA13 is required for fibroblast induction of site-specific keratin. Epidermal induction of plantar-specific K9. (A) Schematic of experiment. K9, a palmo-plantar differentiation marker, is induced in keratinocytes by coculture with plantar fibroblasts, but not more proximal fibroblasts. (B) Induction of K9 protein by coculture with plantar fibroblasts. (C) Relative induction of K9 mRNA in coculture assay upon depletion of HOXA13 in fibroblasts or addition of WNT5A. (ND) Not detectable. (D) Model of HOXA13 transcriptional network. In the developing limb, cells that make WNT5a may also respond to WNT5a themselves. The ectodermal cells in limb bud also express WNT5A in a HOXA13-dependent manner (Fig. 3), suggesting that the distal transcriptional program may occur in both mesenchymal and ectodermal cells of the distal limb bud.

K9 induction

KRT9 qPCR was performed 24 h after coculture of keratinocytes and fibroblasts in KGM media or addition of 300 ng of recombinant WNT5A.

Full methods are available in the Supplemental Material.

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