

# Diverse regulation of mammary epithelial growth and branching morphogenesis through noncanonical Wnt signaling

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The mammary gland consists of an adipose tissue that, in a process called branching morphogenesis, is invaded by a ductal epithelial network comprising basal and luminal epithelial cells. Stem and progenitor cells drive mammary growth, and their proliferation is regulated by multiple extracellular cues. One of the key regulatory pathways for these cells is the  $\beta$ -catenin–dependent, canonical wingless-type MMTV integration site family (WNT) signaling pathway; however, the role of noncanonical WNT signaling within the mammary stem/progenitor system remains elusive. Here, we focused on the noncanonical WNT receptors receptor tyrosine kinase-like orphan receptor 2 (ROR2) and receptor-like tyrosine kinase (RYK) and their activation by WNT5A, one of the hallmark noncanonical WNT ligands, during mammary epithelial growth and branching morphogenesis. We found that WNT5A inhibits mammary branching morphogenesis *in vitro* and *in vivo* through the receptor tyrosine kinase ROR2. Unexpectedly, WNT5A was able to enhance mammary epithelial growth, which is in contrast to its next closest relative WNT5B, which potently inhibits mammary stem/progenitor proliferation. We found that RYK, but not ROR2, is necessary for WNT5A-mediated promotion of mammary growth. These findings provide important insight into the biology of noncanonical WNT signaling in adult stem/progenitor cell regulation and development. Future research will determine how these interactions go awry in diseases such as breast cancer.

mammary stem cells | noncanonical Wnt signaling | receptor tyrosine kinase | epithelial morphogenesis

The mammary gland is composed of a highly dynamic epithelial structure that undergoes multiple rounds of remodeling during puberty, pregnancy, lactation, and involution (1). At puberty, the mammary gland forms a branching ductal network, which connects the nipple to the milk-producing lobuloalveolar structures that arise during pregnancy (2). Development and growth of the mammary gland depends on the function of adult mammary stem cells (MaSCs) (1). These MaSCs are capable of reconstituting a complete mammary epithelial ductal structure when implanted as a single cell into a cleared fat pad *in vivo* (3–5). The morphogenetic changes of the breast epithelium are closely coordinated within the context of its microenvironment, which consists of a variety of stromal cells such as adipocytes, macrophages, and fibroblasts (6, 7). MaSCs respond to extracellular signals such as wingless-type MMTV integration site family (WNT) ligands provided by stromal cells of the microenvironment. For example, hyperactivation of the canonical WNT/ $\beta$ -catenin signaling pathway in the mammary gland expands the MaSC population by sixfold (4), and WNT ligands are necessary for self-renewal properties of MaSCs (8). Constitutive overexpression of the gene encoding the canonical *Wnt1* ligand in this organ ultimately gives rise to tumors, suggesting a direct link between MaSC accumulation and tumor susceptibility (9, 10).

The role of the noncanonical WNT signaling pathway in the regulation of mammary gland development and breast cancer function is obscure. WNT5A and WNT5B represent two noncanonical WNT ligands expressed in the mammary gland (11–14). Within the mammary epithelium, expression of both *Wnt5a* and *Wnt5b* is restricted to the more differentiated luminal epithelial cell lineage (14). Several receptors have been implicated in mediating the function of WNT5A and WNT5B. These include the noncanonical receptors, receptor tyrosine kinase-like orphan receptor 1 (ROR1) and ROR2, which are expressed in both the basal and luminal compartments (14, 15), whereas the expression of the receptor RYK remains less well defined. Although several studies have reported inhibitory roles for noncanonical WNT ligands, WNT5A during branching morphogenesis (16) and WNT5B in mammary stem and progenitor outgrowth (16, 17), the receptors mediating these inhibitory functions remain poorly characterized.

Even though the actions of WNT5A and WNT5B have been associated with events occurring during breast cancer initiation and progression, the role of noncanonical WNT signaling in breast cancer remains elusive. Although there is evidence suggesting that secretion of WNT5A by stromal cells may inhibit the

## Significance

The extracellular signals that regulate cell growth and tissue rearrangements during organogenesis are still poorly understood. One large family of secreted signaling molecules, called wingless-type MMTV integration site family member (WNTs), governs many of these processes by controlling cell-cell interactions. Here, we show that two closely related members of the WNT superfamily influence the development of the mammary gland by signaling through different tyrosine kinase receptors. WNT5A enhances the growth of mammary stem/progenitor cells while restricting the process of branching morphogenesis. In contrast, WNT5B represses mammary stem/progenitor cell proliferation through a different receptor. Together, the distinct actions of these highly homologous extracellular cues regulate the developmental processes that supply cells and shape tissues during periods of rapid growth.

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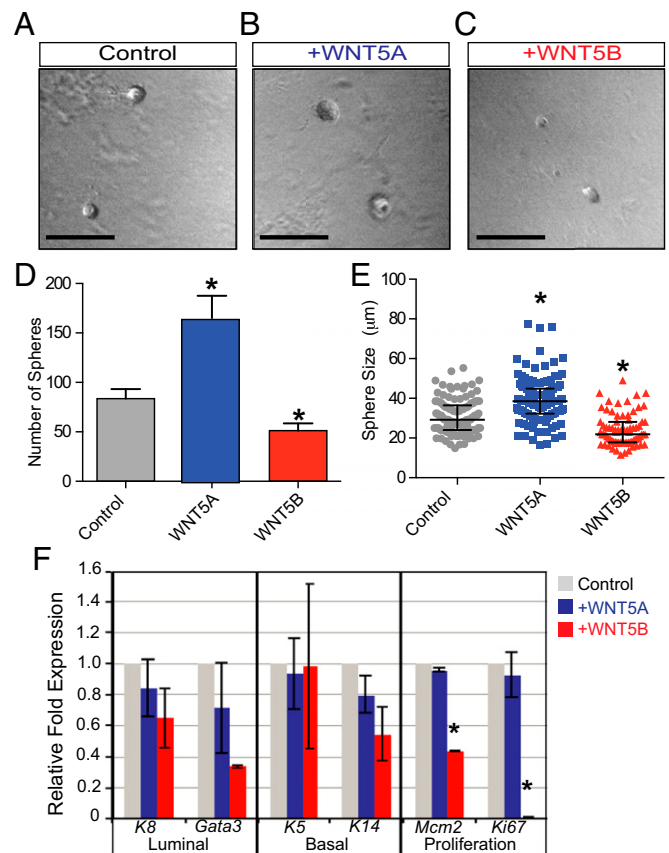
actions of tumor-initiating cells in breast cancer (18), other studies show that WNT5A/B may promote epithelial to mesenchymal transition and metastatic progression in breast and other cancers through a noncanonical *Frizzled2* pathway (19). Overall, the role of noncanonical WNT signaling in breast cancer appears to be particularly context-dependent, and the diverse effects of noncanonical WNT ligands on cell and developmental pathways remain largely unexplored.

Here, we focused on the role of WNT5A and WNT5B as two of the main mediators of noncanonical WNT signaling in the mammary gland. In particular, we sought to understand how noncanonical WNT signaling is involved in the regulation of MaSC function and branching morphogenesis. Our results show that despite their high degree of similarity, WNT5A and WNT5B may work in a distinct manner involving different receptor molecules. These findings further shed light on our understanding of the intricacies of the WNT signaling pathway and provide crucial insights to better understand their role in diseases such as breast cancer.

## Results

**WNT5A and WNT5B Differentially Regulate Mammary Growth and Progenitor Cell Proliferation.** We previously observed that WNT5B is capable of inhibiting mammary epithelial stem and progenitor cell growth capacity in vitro and in vivo, using lentiviral-mediated overexpression in transplanted MaSCs (17). WNT5A and WNT5B show a high degree of amino acid sequence similarity, at 83% (Fig. S1A), which initially suggested that these two WNT ligands may function in a redundant manner. To test whether these two noncanonical WNT ligands are indeed functionally redundant in the context of MaSC/progenitor regulation, we first used the clonal mammosphere assay as a proliferative analysis tool in which single mammary epithelial cells (MECs) are embedded into a 3-dimensional Matrigel-based culture. Using this assay, we then compared the effects of exogenous recombinant WNT5A and WNT5B to vehicle-treated mammosphere formation. Interestingly, we observed a pronounced increase in the number, as well as the size, of mammospheres treated with WNT5A compared with vehicle-treated MECs (Fig. 1A–E). In contrast, WNT5B demonstrated a strong inhibitory effect on mammosphere formation, which is consistent with our previous findings (17). These distinct effects of WNT5A and WNT5B were dose-dependent in a range from 0.1 to 1  $\mu\text{g}/\text{mL}$  (Fig. S1B). Subsequent quantitative PCR (qPCR) analysis revealed that in contrast to WNT5A, WNT5B-treated cells down-regulate expression of genes associated with proliferation and luminal differentiation (Fig. 1F). This suggests that WNT5B blocks mammosphere formation by inducing cell cycle arrest, presumably at an earlier basal MaSC state. Also, despite the high degree of amino acid sequence homology shared by WNT5A and WNT5B (Fig. S1A), these hallmark noncanonical WNT ligands may work in a distinct fashion in the regulation of MEC biology.

**Differential Expression of Noncanonical WNT Receptors in Mammary Epithelial Subpopulations.** To investigate which noncanonical WNT receptors are involved in mediating WNT5A and WNT5B signaling in breast epithelium, we used qPCR expression analysis on isolated basal and luminal MECs. To this end, we used FACS to separate basal, luminal progenitor, and mature luminal cells harvested from 8-wk-old FvB/N mice (Fig. S2A–C) and performed qPCR analysis with primers targeting *Ror1*, *Ror2*, and *Ryk*. We also confirmed isolation of mammary subpopulations, using epithelial lineage marker expression (Fig. S2D). We found all three receptors expressed in the mammary gland (Fig. S2E). Although *Ror1* and *Ryk* show expression in all epithelial cell populations, *Ror2* showed a differential expression pattern, with high abundance in basal cells, low abundance in mature luminal cells, and undetectable expression in luminal progenitor cells. We also confirmed the basal-specific

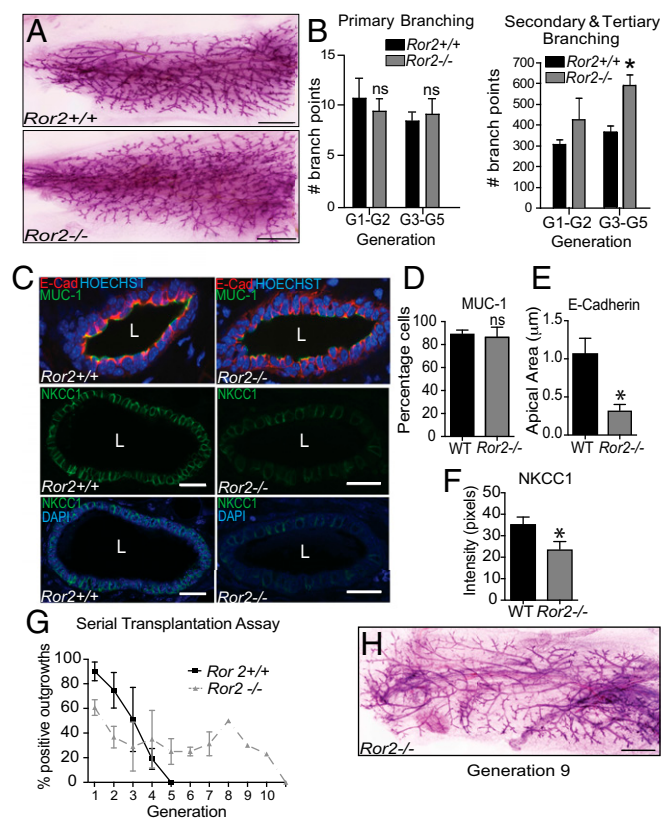


**Fig. 1.** Differential effects of WNT5A and WNT5B on mammosphere outgrowth. Primary WT MECs (2,000 cells per well) were used as single cells in the mammosphere assay and treated with vehicle control or medium containing either WNT5A (0.5  $\mu\text{g}/\text{mL}$ ) or WNT5B (0.5  $\mu\text{g}/\text{mL}$ ) recombinant protein. Representative images show mammosphere formation after 7 d. Compared with control conditions (A), WNT5A-treatment promotes mammosphere formation (B), whereas WNT5B treatment inhibits mammosphere formation (C). For each condition, quantification of sphere number (D) and size (E) was carried out. The effects of WNT5A and WNT5B on cell fate and proliferation were measured by qPCR, using cDNA from MECs cultured in a low-adhesion plate (200,000 cells per well) and treated with control medium (gray, Left) or medium containing either WNT5A (blue, Middle) or WNT5B (red, Right) recombinant protein (F). Error bars show SD ( $n = 3$ ); data representative of three independent experiments. WNT5A-treated cells show an expression profile that was similar to control-treated cells, whereas WNT5B-treated cells showed a significant decrease in the expression of proliferation markers (*Mcm2*, *Ki67*) and a trending decrease in luminal differentiation markers (*K8*, *Gata3*). Error bars show average with SD (D–F) or median measurement with interquartile range (F). (Scale bars, 100  $\mu\text{m}$  in A–C.) Asterisks represent  $*P \leq 0.05$ , Student's *t* test or relative to control.

expression of ROR2 in situ (Fig. S2F) by immunostaining in wild-type (WT) tissue, observing strong staining in the basal compartment (15). Taken together, our results suggest that ROR2 could be mediating a cross-talk between basal and luminal cells by functioning as a receptor on basal cells for luminal-derived WNT5A and WNT5B.

**Ror2-Deficient Mammary Glands Show Increased Branching Morphogenesis.** The basal-prominent expression pattern of ROR2 prompted us to study its role in the regulation of mammary gland development in more detail. *Ror2*-deficient (*Ror2*<sup>−/−</sup>) mice have severe developmental defects leading to death of the animal at age embryonic day 18.5 (E18.5) (20). To analyze the postnatal development of the mammary gland, we generated *Ror2*<sup>−/−</sup> glands in vivo by using standard protocols to rescue the anlage from E16.5 embryos and transplanted them into precleared fat pads of

immunocompromised mice (21). To ensure only epithelial tissue was propagated, we allowed the anlage transplant to develop 12 wk before performing serial transplants with tissue fragments taken from the previous outgrowth. Whole-mounted generation 1 (G1) *Ror2*<sup>-/-</sup> outgrowths were morphologically similar to littermate WT outgrowths generated in the contralateral fat pad (Fig. S2G). By G4, however, *Ror2*<sup>-/-</sup> outgrowths appeared hyperbranched compared with WT glands (Fig. 2A). We performed branching analysis on four independently derived lines of WT and *Ror2*<sup>-/-</sup> tissue (Fig. 2B and Fig. S2H and I). We found no significant differences in primary branch number between *Ror2*<sup>-/-</sup> and WT tissue, even with successive transplantation (G1–G5), indicating that terminal end bud bifurcation is unaffected by the lack of *Ror2*. In contrast, there was a significant increase in secondary and tertiary branching in *Ror2*<sup>-/-</sup> tissue that had been serially transplanted for three to five generations, indicating a progressive misregulation in mechanisms that constrain side-branch formation (Fig. 2B). Cross-sections through WT G4

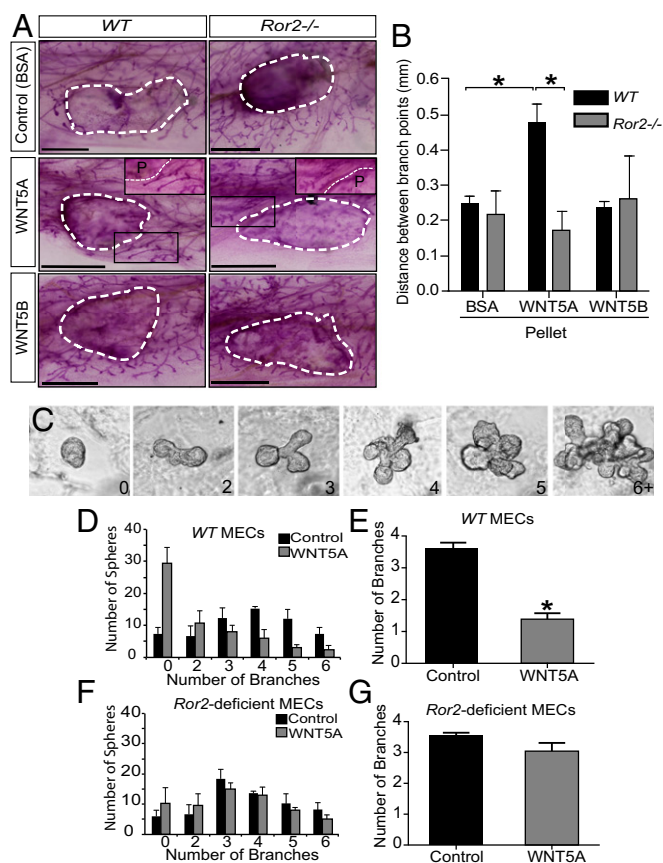


**Fig. 2.** *Ror2*-deficient mammary glands show increased secondary and tertiary branching morphogenesis in vivo. Anlage were rescued from E16.5 embryos and transplanted into precleared fat pads of athymic nude mice. After 12 wk, epithelial tissue fragments were used to propagate the line. Whole-mounts of G4 tissue reveal a hyperbranched phenotype (A) that was quantified by tracing the primary and secondary/tertiary branches (B) ( $n = 4$ ). Immunohistochemistry in cross-sections of G4 *Ror2*<sup>-/-</sup> and WT tissue, using E-cadherin (red) MUC-1 (green), NKCC1 (green) antibodies, and Hoechst (blue) for nuclei, reveal disorganization in the *Ror2*<sup>-/-</sup> tissue ( $n = 3$ ) (C). Quantification of MUC-1 (D), E-cadherin (E), and NKCC1 immunostaining (F) in WT and *Ror2*<sup>-/-</sup> luminal cells ( $n = 3$ ). Serial transplantation of tissue fragments show that WT tissue ( $n = 3$ ) senesces at G5, whereas *Ror2*<sup>-/-</sup> tissue displays enhanced transplantability ( $n = 2$ ).  $n$  is the number of independently derived lines, with five contralaterally transplanted mice analyzed per line. (G) Whole-mount of G9 *Ror2*<sup>-/-</sup> outgrowth shows epithelial tissue filling the fat pad (H). (Scale bars, 1.5 mm for A and H, 10 μm for C.) \* $P < 0.05$ , Student's  $t$  test. Error bars represent  $\pm$  SE.

tissue revealed a normal bilayered epithelial architecture (Fig. 2C), with luminal cells lining up in register to create a smooth apical surface. In contrast, luminal epithelial cells are slightly out of register in the *Ror2*<sup>-/-</sup> G4 tissue, but there is no significant difference in apical MUC1 immunostaining, suggesting *Ror2*<sup>-/-</sup> luminal cells are properly polarized (Fig. 2C and D). However, E-cadherin immunostaining in *Ror2*<sup>-/-</sup> tissue is reduced at the adherens junctions, which reside just below the tight junctions that demarcate the apical surfaces of luminal cells (Fig. 2C and E). Instead of the organized belt of E-cadherin staining observed in WT tissue, staining in *Ror2*<sup>-/-</sup> tissue extends along the lateral and basal membranes. We also examined the localization of the Na/K/Cl cotransporter, NKCC1, which marks the basolateral surface (22), and found an overall decrease in staining in *Ror2*<sup>-/-</sup> tissue (Fig. 2C and F). Taken together, the data suggest that luminal epithelial cells maintain their proper orientation with regard to the lumen in *Ror2*<sup>-/-</sup> tissue, but contacts between cells appear compromised. Previous studies showed that WT tissue fragments can be serially transplanted up to five times before no further outgrowths are obtained (23). In our studies, all three WT lines senesced at G5 (Fig. 2G and H). In contrast, two of four *Ror2*<sup>-/-</sup> transplanted lines displayed increased transplantability, with one senescing at G9 and the second at G11, suggesting that loss of *Ror2* regulates either the number of MaSC/progenitor cells or their self-renewal.

**WNT5A Inhibits Mammary Epithelial Branching Through ROR2.** Next, we focused on the interaction of ROR2 with noncanonical WNT ligands in the context of mammary branching morphogenesis. Both WNT5A and WNT5B have previously been shown to inhibit mammary branching morphogenesis in vivo (16, 17). To test whether ROR2 is involved in this process, we used our *Ror2*<sup>-/-</sup> transplantation model in combination with Elvax slow-release pellets to locally deliver WNT ligands into the tissue adjacent to the epithelial outgrowths. To quantify the effect of WNT5A and WNT5B on branching morphogenesis in vivo, we measured the distance between branch points in close proximity to the transplanted pellets. We observed significantly increased distance between branches near the WNT5A pellets, an indication of reduced branching compared with BSA-containing control pellets. In contrast, WNT5B had no measurable effect on mammary side branching under these conditions, which further underlines that WNT5A and WNT5B may possess different functional properties. Importantly, in the absence of *Ror2*, the inhibitory effects of WNT5A on side branching were completely abolished (Fig. 3A and B), indicating that ROR2 functions as the primary receptor mediating this regulatory role of WNT5A during mammary branching morphogenesis. We further corroborated these findings using an established ex vivo surrogate assay for mammary branching morphogenesis (Fig. 3C). After 7 d of WNT5A treatment, WT mammary epithelial organoids showed significantly fewer branches compared with mock treated samples (Fig. 3D and E). In contrast, *Ror2*<sup>-/-</sup> organoids treated with WNT5A showed no change in branching after 7 d (Fig. 3F and G), demonstrating that WNT5A inhibits mammary branching morphogenesis through ROR2.

**WNT5A Promotes Mammosphere Formation Through RYK.** We initially observed distinct effects of WNT5A and WNT5B on mammosphere formation, which may be driven primarily by different basal stem and progenitor cell functions, rather than the process of branching morphogenesis. We therefore explored the role of ROR2 as it relates to its basal epithelial expression pattern, and our results suggest that the loss of *Ror2* promotes the longevity of stem cells and their potential to serially undergo mammary morphogenesis in vivo (Figs. 1 and 2). To this end, we isolated MECs from WT and *Ror2*<sup>-/-</sup> outgrowths harvested from contralaterally transplanted animals and subjected these to the

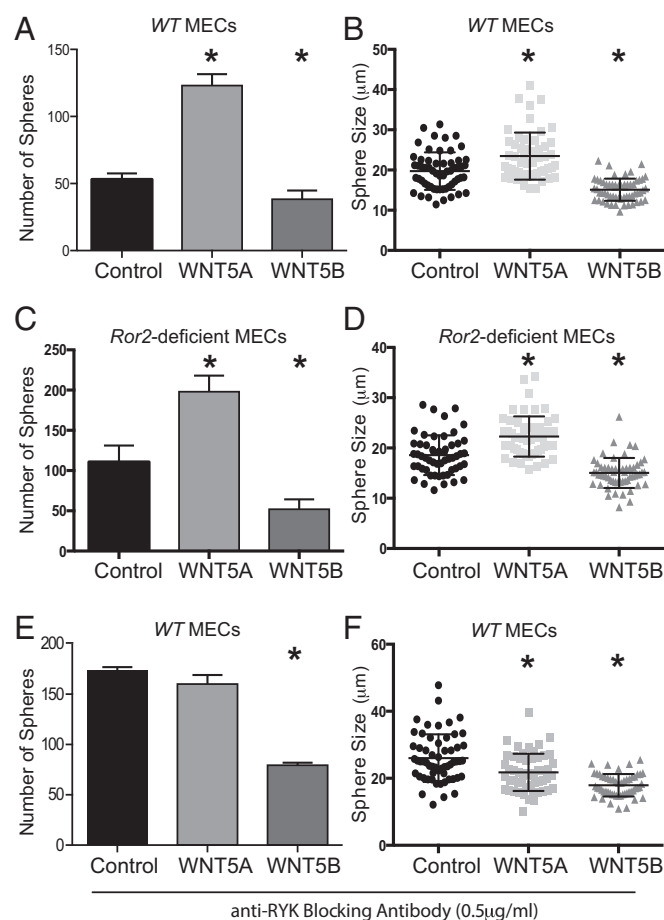


**Fig. 3.** WNT5A inhibits mammary branching morphogenesis through ROR2. WT and *Ror2*<sup>-/-</sup> fragments were contralaterally transplanted into precleared mammary fat pads. Elvax slow-release pellets containing BSA, WNT5A, or WNT5B were implanted bilaterally 3 wk posttransplant. The tissue was harvested and carmine stained for whole-mount analysis 1.5 wk after the Elvax implantation (A). The distance between branch points was quantified by tracing the ductal structure in proximity of the pellet and measuring the length between branch points ( $n = 3$  lines per five contralaterally transplanted mice per line) (B). Ex vivo branching morphogenesis of reaggregated primary MECs in the presence of FGF2 was carried out over the course of 7 d, and the number of branches per organoid was determined. Example images for organoids with 0–6+ branches are shown (C). WT MECs show significant reduction of organoid branching in the presence of WNT5A compared with control conditions (D and E) ( $n = 3$ ). Organoids derived from *Ror2*-deficient MECs show no decrease in the number of branches per organoid when treated with WNT5A (F and G) ( $n = 3$ ). \* $P < 0.05$ , Student's *t* test. Error bars represent  $\pm$  SE. (Scale bars, 1.5 mm for A.)

mammosphere formation assay in the presence or absence of WNT5A and WNT5B (Fig. 4). We observed no difference in the effects of WNT5A or WNT5B on sphere forming capacity in WT and *ROR2*<sup>-/-</sup> samples (Fig. 4 A–D), demonstrating these Wnt ligands act independent of ROR2 in this context. In treated samples, we found that WNT5A promoted and WNT5B reduced the number and size of mammospheres independent of ROR2 (Fig. 4 A–D), indicating that a different noncanonical receptor is governing the effects of WNT5A and WNT5B on mammosphere formation.

We next focused on the receptor tyrosine kinase RYK, which has been implicated as a receptor for WNT5A (24). To interfere with RYK function in the presence or absence of WNT ligand, we used a blocking antibody approach that has been previously used to inhibit RYK function during neuron specification (25). Although WNT5B inhibited mammosphere formation independent of RYK blockade, we found that the increase in number and size of mammospheres treated with WNT5A could be abolished by the presence of anti-RYK blocking antibody (Fig. 4 E and F),

demonstrating that WNT5A may increase MaSC capacity and proliferation by activating RYK-dependent, noncanonical WNT signaling. Importantly, control experiments under the same conditions, but treated with nonspecific IgG instead of anti-RYK treatment, did not alter mammosphere capacity (Fig. S2J). Interestingly, we noticed that in the presence of RYK-blocking antibody, WNT5A changed its function from promoting to inhibiting mammosphere growth, as measured by sphere size, whereas the number of spheres remained comparable to control (Fig. 4 E and F). This could suggest that in the absence of RYK, WNT5A engages different WNT coreceptors, leading to inhibition of progenitor cell proliferation or promotion of cell differentiation. Together, these findings show that WNT5A may activate the receptor tyrosine kinases ROR2 and RYK, presumably on different stem or progenitor populations that contribute to distinct morphogenetic processes during mammary branching and MaSC outgrowth (Figs. 3 and 4). To further corroborate this notion, we analyzed the expression pattern for ROR2 and RYK on a single-cell level, using a single-cell RNAseq dataset generated in a currently ongoing study, which revealed that ROR2 and RYK are



**Fig. 4.** The role of ROR2 and RYK in WNT5A- and WNT5B-mediated regulation of mammosphere formation. MECs from contralateral WT (A and B) or *Ror2*<sup>-/-</sup> outgrowths (C and D) (2,000 cells per well) were used as single cells in mammosphere assays and treated with control medium or medium containing either WNT5A (0.5  $\mu$ g/ml) or WNT5B (0.5  $\mu$ g/ml) recombinant protein ( $n = 3$ ). The experiment was also performed in the presence of anti-RYK blocking antibody, using WT cells (E and F) ( $n = 3$ ). After 7 d in WNT containing culture medium, sphere number (A, C, and E) and size (B, D, and F) were determined for the WT, *Ror2*<sup>-/-</sup>, and anti-RYK-treated spheres. Error bar shows SD (A, C, and E) or median measurement with interquartile range (B, D, and F). \* $P \leq 0.05$ , Student's *t* test.

indeed expressed in a mutually exclusive manner in the vast majority of MECs (Fig. 5 *A* and *B*).

## Discussion

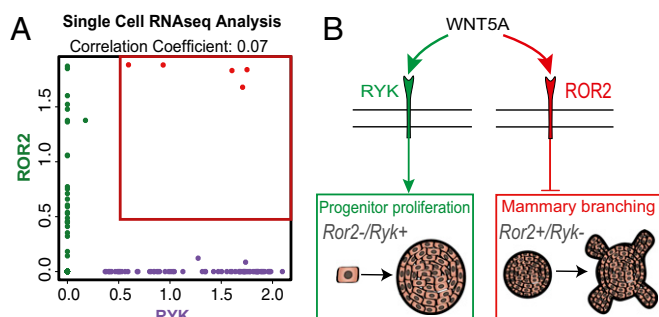
Although canonical WNT signaling has been extensively studied in the context of mammary gland biology, MaSC regulation, and cancer, the noncanonical WNT pathway remains elusive. To learn more about the role of noncanonical WNT signaling in mammary development, we focused here on the hallmark noncanonical ligands WNT5A and WNT5B because they are prominently expressed in the mammary gland. By investigating the alternate WNT receptors, ROR2 and RYK receptor tyrosine kinases, our studies provide insight into noncanonical WNT signaling pathways during mammary development. Whereas previous studies have shown that loss of either *Wnt5a* or *Ror2* results in a hyperbranching phenotype (15), we demonstrate a ligand/receptor relationship by showing that ROR2 is required for the inhibitory action of WNT5A on branch formation. Despite being highly homologous, WNT5B has no such effect. We further distinguished the activities of these two WNTs in mammosphere assays by demonstrating that WNT5A enhances mammosphere formation through the RYK receptor. This stimulatory activity is surprising in light of studies on human tumors that showed tumor-suppressive activity of WNT5A, with loss of *Wnt5a* expression correlating with more aggressive tumor subtypes, earlier relapse, and significantly shorter overall patient survival (26, 27). WNT5A was shown to induce the formation of a RYK/TGF $\beta$ R1 complex in MCF10A cells and activate SMAD2 in mammary epithelial cells harvested from preneoplastic MMTV-*ErbB2*/*Wnt5a*<sup>+/-</sup> mice (27). However, in studies on nonimmortalized cells, we and others (28) find that WNT5A promotes mammosphere formation, a positive effect on stem/progenitors consistent with WNT5A's ability to support the self-renewal of mouse spermatogonial stem cells by inhibiting apoptosis (29). In two studies, pharmacological inhibition of the Jun N-terminal kinase abrogated the WNT5A-mediated outcome (28, 29), and in the studies on mammospheres, the effect was also reversed by knocking down *Ror2* (28). In contrast, our studies show that WNT5A enhances the formation of mammospheres through the RYK receptor, independent of ROR2, perhaps reflecting the difference between knocking down and knocking out a receptor. Our studies also demonstrate that WNT5B blocks mammosphere formation, possibly by inhibiting cell cycle progression because markers for cell cycle entry, *Mcm-2* and *Ki67*, are significantly down-regulated in response to WNT5B. These results are consistent with our previous studies demonstrating an

inhibitory effect associated with WNT5B treatment on MaSC proliferation in vitro and in vivo (17). Although we have not yet identified the receptor mediating this effect, we have ruled out ROR2. Taken together, our data show that noncanonical WNT5A and WNT5B have distinct functions despite a high degree of sequence homology. Our study reveals that WNT5A signals through two different receptors: ROR2 to inhibit branching morphogenesis and RYK to promote mammary MaSC/progenitor proliferation. ROR2 also has a separate function in enhancing MaSC/progenitor self-renewal, but the ligand mediating this effect is unknown.

Multiple potential noncanonical WNT receptors are expressed in the mammary gland. Similar to a previous study (16), our results show that *Ror2* is predominantly expressed in the basal population with little or no expression in luminal progenitors and low expression in mature luminal cells. In contrast, the expression of *Ryk* and *Ror1* is more evenly distributed across populations. Because the basal compartment contains a subpopulation of MaSCs (1), we further explored the function of ROR2 in this compartment by examining the phenotype of *Ror2*<sup>-/-</sup> glands. Our immunostaining data revealed strong basal expression of ROR2 in myoepithelial cells along the duct (15). We further explored ROR2 function by examining its knock-out phenotype, using the technique of transplantation to circumvent the perinatal lethality of the mutation. Loss of either *Wnt5a* or *Ror2* has been associated with a hyperbranching phenotype in the mammary gland, but the molecular mechanism behind this phenotype remains elusive. Although *Ror2*<sup>-/-</sup> mammary glands exhibited no defects in primary branch formation, excessive secondary and tertiary branching was observed after three successive rounds of transplantation. In contrast, others have demonstrated an initial branching defect in mammary outgrowths generated from *Ror2* knockdown cells (15), suggesting there may be activation of compensatory signaling pathways caused by the germline deletion of this tyrosine kinase receptor, which complicates the detection of branching defects in *Ror2*<sup>-/-</sup> tissue. By implanting Elvax pellets in vivo into fat pads containing WT and *Ror2*<sup>-/-</sup> outgrowths, we demonstrate that WNT5A, but not WNT5B, has the capacity to reduce branch formation in a *Ror2*-dependent manner. Histological analysis revealed disorganized epithelial architecture that is consistent with a low level of ROR2 expression in mature luminal cells and may be a result of defects in the actin cytoskeleton, which is disrupted in the absence of ROR2 (15). Using an in vitro branching assay, our studies further revealed that WNT5A negatively regulates branch formation through the ROR2 axis, supporting a crucial developmental role for WNT5A/ROR2 signaling in restricting mammary branching morphogenesis.

Despite their high degree of sequence similarity, WNT5A showed substantially distinct functional properties compared with WNT5B, resulting in enhanced MaSC/progenitor-mediated mammosphere formation in a RYK-dependent manner. By using an anti-RYK, function-blocking antibody, we determined that the RYK pathway is responsible for the WNT5A-mediated increase in mammosphere formation. It is tempting to speculate that high local levels of WNT5A may be important during the formation of terminal end buds invading into the mammary fat pad, when enhanced proliferation is required and premature formation of side branches has to be suppressed. The distinct effects of WNT5A inhibiting branching morphogenesis through ROR2, and activation of MaSC/progenitor proliferation through RYK, also indicate that there are distinct stem and progenitor cell populations associated with these aspects of mammary gland development.

Taken together, our studies provide important insight into the biology of noncanonical WNT signaling in adult stem cell regulation and development. We discovered that WNT5A and WNT5B may have biologically distinct functions during mammary development. Although WNT5B blocks MaSC growth and has no measurable effect on branching morphogenesis, WNT5A enhances MaSC outgrowth and negatively regulates mammary branching. These effects are mediated through diverse engagement of RYK and ROR2,



**Fig. 5.** Single-cell analysis and functional model summarizing the role of WNT5A in mammary development. Scatter plot of single-cell correlation analysis of ROR2 and RYK expression in mouse MECs shows very low correlation and almost mutually exclusive expression (*A*), demonstrating that ROR2 and RYK are expressed in different populations of cells. Schematic illustrating the differential expression patterns of noncanonical WNT receptors in MEC populations and the signaling capacities of WNT5A acting either through ROR2 to block branching morphogenesis or through RYK to promote stem/progenitor proliferation (*B*).

which highlights the complexity and context dependency of WNT signaling in general. Future research will determine how these interactions go awry in diseases such as breast cancer.

## Experimental Procedures

**Mice.** FVB/N mice were purchased from Charles River Laboratories. KO mice were generated and genotyped as described; *Ror2*<sup>-/-</sup> (20). Mice were maintained in a pathogen-free facility. All mouse procedures were approved by the University of California, San Francisco, or University of California, Santa Cruz, Institutional Animal Care and Use committees.

**Recombinant Proteins and Antibodies.** The list of recombinant proteins and antibodies is provided in the *SI Experimental Procedures*.

**Mammosphere Formation and Branching Assay.** Mammosphere assays were carried out as described previously (4, 5), and as described in *SI Experimental Procedures*.

**Mammary Gland Transplantation, in Vivo Branch Quantification, and Elvax Implantation.** Mammary anlage were rescued from *Ror2*<sup>-/-</sup> embryos and

transplanted into precleared fat pads of *Foxn1*<sup>nu</sup> mice, as previously reported (21) and described in the *SI Experimental Procedures*.

**Immunohistochemistry.** Immunohistochemistry was performed as previously described with ref. 30. Images were collected on a Leica SP5 confocal microscope. Brightfield imaging was performed on a Bioevo BZ-9000 Digital Microscope (Keyence).

**Statistical Analysis.** All statistical analyses (Student's *t* test) were conducted on Graphpad Prism; *P* values of less than 0.05 were considered significant.

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