WNT5a-ROR Signaling Is Essential for Alveologenesis

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Abstract: WNT5a is a mainly “non-canonical” WNT ligand whose dysregulation is observed in lung diseases such as idiopathic pulmonary fibrosis (IPF), chronic obstructive pulmonary disease (COPD) and asthma. Germline deletion of Wnt5a disrupts embryonic lung development. However, the temporal-specific function of WNT5a remains unknown. In this study, we generated a conditional loss-of-function mouse model (Wnt5aCAG) and examined the specific role of Wnt5a during the saccular and alveolar phases of lung development. The lack of Wnt5a in the saccular phase blocked distal airway expansion and attenuated differentiation of endothelial and alveolar epithelial type I (AT1) cells and myofibroblasts. Postnatal Wnt5a inactivation disrupted alveologenesis, producing a phenotype resembling human bronchopulmonary dysplasia (BPD). Mutant lungs showed hypoalveolization, but endothelial and epithelial differentiation was unaffected. The major impact of Wnt5a inactivation on alveologenesis was on myofibroblast differentiation and migration, with reduced expression of key regulatory genes. These findings were validated in vitro using isolated lung fibroblasts. Conditional inactivation of the WNT5a receptors Ror1 and Ror2 in alveolar myofibroblasts recapitulated the Wnt5aCAG phenotype, demonstrating that myofibroblast defects are the major cause of arrested alveologenesis in Wnt5aCAG lungs. Finally, we show that WNT5a is reduced in human BPD lung samples, indicating the clinical relevance and potential role for WNT5a in pathogenesis of BPD.

Keywords: WNT5a; ROR; lung; alveologenesis; secondary crest myofibroblast; migration

1. Introduction

Structurally and functionally, the mammalian lung represents a remarkably complex and cellularly diverse organ. The primary architecture of the mouse lung as a highly branched structure is laid down during early embryonic days (E) 9.5 through E16.5 by branching morphogenesis. Subsequently, from E17.5 to postnatal day (PN) 4, structures known as saccules are formed that are
predecessors of the final respiratory units or alveoli. Alveolar formation, or alveologenesis in the mouse lung occurs entirely in the postnatal period and is estimated to be complete by PN30 [1]. Biological events spanning the late saccular stage to postnatal alveologenesis are required for full maturation of the lung as an efficient gas exchange organ. While a wealth of information is currently available on mechanisms of early lung morphogenesis, our understanding of late embryonic development and postnatal alveologenesis remain surprisingly limited.

During the saccular phase of mouse lung development, distal airways in the lung undergo "expansion" to form saccules. Subsequently, the progenitors of AT1 (NKX2.1<sup>pos</sup>HOPX<sup>pos</sup>) and AT2 (NKX2.1<sup>pos</sup>SFTPC<sup>pos</sup>) cells begin to give rise to mature and functional AT1 (AQP5<sup>pos</sup>PDPN<sup>pos</sup>) and AT2 (SFTPB<sup>pos</sup>SFTPC<sup>pos</sup>) cells [2]. Concurrently, the saccular walls undergo thinning as the capillary networks are established and surfactant production commences. A major event is the formation of structures known as secondary crests (SC), which on 2-dimensional images appear as protrusions into the saccular space [3]. Formation of SC is driven by mesodermally derived secondary crest myofibroblasts (SCMF), also known as alveolar myofibroblasts. Progenitors of SCMF express Pdgfra in the saccular phase and Atca2 in the alveolar phase of lung development [4–6]. As they differentiate, SCMF undergo directed migration to form the SC. Disruption of SCMF differentiation or migration leads to arrested alveologenesis [4,5]. Differentiation of the latter cell types during late stage of lung development or lung maturation is associated with expression of several cell-type-specific genes, as listed in Table S1.

WNT signaling is a critical regulator of normal lung morphogenesis, homeostasis and injury repair. WNT ligands can activate the beta-catenin-dependent (the canonical) or the alternative beta-catenin-independent (the non-canonical) pathways [7]. WNT5a, a predominantly non-canonical WNT ligand, is increasingly recognized as an important regulator of stem-cell renewal, cell migration, cell polarity and inflammatory responses [8,9]. WNT5a expression is present in both epithelial and mesenchymal compartments during embryonic stages and is mainly in fibroblasts and endothelial cells in adult lungs [9–11]. Dysregulated WNT5a signaling is observed in many lung diseases, ranging from chronic obstructive pulmonary disease (COPD) [12] and idiopathic pulmonary fibrosis (UIP/IPF) [13] to asthma [14]. There are reliable data indicating that exquisitely regulated, temporal and cell-type-specific WNT5a signaling is a strict requirement for normal lung development. We and others have shown that lack of WNT5a activity is associated with abnormal branching of distal airways together with defects in capillaries and alveolar airspaces [10,15]. Conversely, overexpression of Wnt5a in Sftpc-Wnt5a transgenic mice disrupts epithelial branching and lobe formation but the mice are postnatally viable, indicating no impact on lung function [16]. Importantly however, to date the role of Wnt5a in alveologenesis remains unclear.

RO1R and ROR2 are tyrosine kinase receptors with functional redundancy [17]. Germline deletion of either Ror1 or Ror2 disrupts lung development [17]. Compound Ror1/Ror2 mutant mice present with a more severe, abnormal phenotype compared to single gene mutants, and are similar in phenotype to Wnt5a knockout mice [17,18]. Both ROR1 and ROR2 mediate WNT5a signaling [18–21]. In compound transgenic lungs, inactivation of Ror1 and Ror2 blocks the lung phenotype caused by overexpression of Wnt5a in Sftpc-Wnt5a mice [21].

In this study, we generated a number of conditional Wnt5a loss-of-function genetic models to elucidate the precise function of Wnt5a during specific late stages of lung development. We found that differentiation of multiple cell types was disrupted by conditional Wnt5a inactivation in the saccular phase. In contrast, conditional Wnt5a inactivation in the alveolar phase had no impact on epithelial cell differentiation, but interrupted differentiation and migration of the mesodermally derived SCMF, leading to defective alveolar formation. Finally, conditional inactivation of the WNT5a receptors Ror1 and Ror2 in SCMF also resulted in a similar arrested alveologenesis phenotype, validating our findings on the role of WNT5a signaling during the critical phase of alveolar formation.
2. Materials and Methods

2.1. Mouse Breeding and Genotyping

All animals were maintained and housed in pathogen-free conditions at constant room temperature (20–22 °C), with a 12 h light/dark cycle, and free access to water and food in the animal facility of University of Southern California, according to a protocol approved by the USC Institutional Animal Care and Use Committee (IACUC) (Los Angeles, CA, USA). CAG-creER;Rosa26mTmG;Wnt5a/f (Wnt5aCAG) mice were generated by breeding CAG-creER mice (Tg(CAG-cre/Esr1*)5Amc/J, The Jackson Laboratory) and Rosa26mTmG (Gt(ROSA)26Sor tm4(ACTB-tdTomato,-EGFP)Luo/J, The Jackson Laboratory) mice with Wnt5af/f mice [22]. Gli1-creERT2;Rosa26mTmG mice (mTmGGli) were generated by breeding Gli1-creERT2 [23] and Rosa26mTmG mice. Gli1-creERT2;Rosa26mTmG;Ror1f/f;Ror2f/f (RorGli;GFP) mice were generated by breeding mTmG mice with the Ror1f/f;Ror2f/f mice [18].

Genotyping of the transgenic mice was performed by PCR with genomic DNA isolated from mouse tails. The forward (F) and reverse primers (R) for transgenic mouse genotyping are listed below.

CAG-creER: (forward) 5’-TAAAGATATCTCACGTACTGACGGTG-3’ and (reverse) 5’-TCTCTGACCAGAGTCATCCTAGC-3’.

Gli1-creERT2: (forward) 5’-TAA AGA TAT CTC ACG TAC TGA CGG TG-3’ and (reverse) 5’-TCT CTG ACCAGAGTCACTCATTAGC-3’.

Wnt5a/f: (Forward) 5’-TGAGGGACTGGAAGTTG CAGG-3’, and (Reverse) 5’-TTCCAATGGGCTTCTGGAGAG-3’.

Ror1/f: (Forward) 5’-CGTTTTCCTCCTGCTCACAGG-3’, and (Reverse) 5’-GCAGACCTGGTGAATTCTACCTCAG-3’.

Ror2/f: (Forward) 5’-CATGGGCCACATCCACATCCCGAG-3’, and (Reverse) 5’-CCGGCGACGTGCCTTGAATGACC-3’.

2.2. Tamoxifen Administration

For embryonic studies, tamoxifen (Sigma, 40mg/mL in peanut oil) was administered by oral gavage to pregnant females from E14 to E17 (2.4 mg each mouse, one dose per day). Embryonic lungs were collected at E18. For neonatal studies, tamoxifen (8mg/mL in peanut oil) was administered by oral gavage to neonates at postnatal day 2 (P2, 400 μg each pup) with a plastic feeding needle (Institute Laboratories, PA). Neonatal lungs were collected between P12 and P13 for morphological, immunohistochemical and molecular biological analyses.

2.3. Human Neonatal Lung Samples

Human lung tissue samples were obtained postmortem by expedited autopsy of preterm infants having the diagnoses of mild RDS, evolving and established BPD as well as term infants as controls (no lung disease). Consented collection and processing of the lung samples as from the deceased were approved by the University of Rochester Institutional Review Board. Selected clinical and recovery details have been previously published [24].

2.4. Immunofluorescent Staining

Immunofluorescent staining was conducted as previously described [25] using paraffin-embedded lung sections. In brief, five micrometer (μm) tissue sections were deparaffinized, rehydrated and subjected to antigen retrieval. After blocking with normal serum, the sections were probed with primary antibodies at 4 °C overnight. Combinations of fluorescein anti-mouse and Cy3 anti-rabbit or anti-goat IgG secondary antibodies (Jackson ImmunoResearch Laboratories, ING) were applied to detect specific primary antibodies. Nuclei were counterstained with 4′,6-diamidino-2-phenylindole (DAPI). Primary antibodies used are listed in Table S3.
2.5. Neonatal Lung Fibroblast Isolation, Treatment and Migration Assay

P5 neonatal lungs of Wnt5aCAG mice and littermate controls were dissected in HBSS (GIBCO24020-117), inflated with Dispase and digested by continuous shaking in Dispase at 37 °C for 15 min. Lung lobes were then isolated, cut into small pieces, transferred into Miltenyi tubes in 5 mL HBSS and dissociated with a gentle MACS dissociator (Miltenyi Biotec. Inc., San Diego, CA, USA). After filtering through 40 μm cell strainers, the dissociated cells were collected by centrifugation at 1200 rpm for 5 min and resuspended in DMEM containing 10% FBS. The cells were then plated in cell-culture plates and incubated at 37 °C with 5% CO2 for 1 h. After removing floating cells, the attached fibroblasts were washed with PBS and cultured in fresh medium. Loosely attached cells were removed by repeated washes within 20 h of culture. The purity of the fibroblasts isolated under this condition was examined by using the Pdgfra-EGFP model, which expresses enhanced GFP under the control of an endogenous Pdgfra promoter [26]. Approximately 90% of the attached cells were GFP positive 20 h after cell isolation. When the cells grew to near confluence, they were trypsinized and stored in 10% DMSO (DMEM + 10% FBS + 10% DMSO) in liquid nitrogen for future use. For the cell migration assay, cells grown in culture flasks (ThermoFisher Scientific, Waltham, MA, USA) were treated with tamoxifen (1.5 μg/mL) for 24 h, and then re-plated (1 × 10^5 cells per Transwell) onto 12 μm Transwell membranes (MiliporeSigma, Burlington, MA, USA) and cultured in DMEM containing 1% FBS inside and 10% FBS outside the Transwell insert. After 24 h, the cells were fixed in 4% formaldehyde, washed with PBS and stained with crystal violet. The cells on the upper side of the membrane were cleaned with a cotton swab and the cells that migrated to the lower side of the Transwell membrane were imaged and counted.

2.6. Real-time Quantitative Polymerase Chain Reaction (real-time RT-PCR)

Neonatal mouse lung fibroblasts were isolated and cultured as described above (Section 2.5). 1.5x10^5 cells were plated onto a 12-well culture plate until they grew to near confluence. The cells were treated with tamoxifen (1.5 μg/mL) for 24 h and then collected for RNA isolation with TRIzol reagent (ThermoFisher Scientific, MA, USA). cDNA was synthesized with an EasyScript Plus cDNA synthesis kit according to the manufacturer’s protocol (Lamda Biotech Inc., Ballwin, MO, USA). Expression of selected genes was quantified by real-time RT-PCR using a LightCycler with LightCycler Fast Start DNA Master SYBR Green I Kit (Roche Applied Sciences, Indianapolis, IN, USA) as previously described [16]. Each reaction contained 3 μL of FastStart SYBR green reaction mix plus enzyme, 2.5 μM of primers and 0.5 μL of cDNA in a total volume of 20 μL. The running protocol consisted of four steps, including 1) denaturation at 95 °C for 6 min; 2) amplification and quantification under 95 °C for 0–10 s, 62 °C for 15 s and 72 °C for 20 s for 45 cycles; 3) melting curve program, where the reaction temperature rapidly increases to 95 °C, then decreases to 60 °C for 15 s, followed by a slow increase to 98 °C at a rate of 0.1 °C per s with continuous fluorescence monitoring; and 4) cooling to 40 °C. Reaction conditions were optimized to create a one-peak melting curve. The ΔΔCt method was used to calculate relative ratios of a target gene mRNA in mutant lungs compared to littermate control lungs. TBP (TATA-Box Binding Protein) was used as the reference gene. Primers for real-time RT-PCR were designed using the program of the Universal ProbeLibrary Assay Design Center from Roche Applied Sciences (IN). Sequences of the primers are listed in Table S4.

2.7. Statistics Analysis

At least three biologically independent control and mutant lungs were used for each morphometric analysis and real-time RT-PCR analyses. Five to six images (10× magnification) from each lung were used to manually count MLI. Quantitative data are presented as mean values ± the standard error of the mean. p values were calculated by two-tailed Student’s t tests.
3. Results

3.1. WNT5a is Required for Differentiation of Multiple Cell Types During Saccular Stage Lung Development

We previously reported that germline deletion of Wnt5a (Wnt5a−/−) disrupts development of saccular stage lungs [10]. However, whether the lung saccular defects (E18) were secondary consequences of defects that may have occurred during branching morphogenesis was not addressed. In the present study, we generated a regulatable (conditional) triple transgenic mouse model, CAG-creER;Wnt5aff;Rosa26mTmG, referred to as Wnt5aCAG (see Materials and Methods) to target Wnt5a globally upon activation of CAG-creER by administration of tamoxifen. To assess the impact of Wnt5a inactivation during the saccular stage of lung development, pregnant Wnt5aff;Rosa26mTmG female mice, mated with Wnt5aCAG males, received tamoxifen orally from E14 to E17 (Figure 1). The embryos and the isolated lungs were analyzed at E18. As shown in Figure 1, Wnt5aff;Rosa26mTmG control (Panels A and B) embryos showed a normal morphology, and Wnt5aCAG littermate embryos (Panels C and D) were similar to control embryos with the exception of minor craniofacial differences (i.e., mouth) (Figure S1). In contrast, the germline Wnt5a−/− embryos (Figure 1, Panel E) are profoundly abnormal with truncation of multiple structures, including limbs, tail and mouth [27]. Further analysis showed that compared to Wnt5aff;Rosa26mTmG controls (Panels F and G), gross morphology of the littermate Wnt5aCAG lungs was unaffected by a lack of WNT5a signaling during the saccular phase of lung development (Panels H and I). In contrast, germline Wnt5a−/− lung lobes were clearly deformed (Panel J) [10]. Histological analysis using hematoxylin and eosin (H&E)-stained multiple lung sections showed that Wnt5aff;Rosa26mTmG control lungs (Panel K) are composed of uniformly organized and expanded saccular structures, while the Wnt5aCAG lungs are heterogenous and consist of both areas of expanded (i.e., sacculi) as well as unexpanded airspace (Figure 1, Panel L). The germline Wnt5a−/− lungs contained nearly uniformly unexpanded airspace (Figure 1, Panel M). The phenotypic differences between the control and Wnt5aCAG lungs were further illustrated by immunostaining for GFP and AT1 cell marker PDPN (Figure 1, Panels N–S).
Our previous analysis found that AT1 cell differentiation was reduced in Wnt5a−/− lungs [28]. The saccular phase is the critical time period during which major decisions on establishment of AT1 and AT2 cell lineages as well as differentiation of various mesodermally derived cells take place. We therefore used our conditional Wnt5aCAG mice in combination with lineage-specific markers for AT1, AT2 and endothelial cells, as well as for myofibroblasts (Table S1) to determine the role of WNT5a specifically during the saccular phase of lung development. As shown in Figure 2, conditional Wnt5a inactivation in Wnt5aCAG embryos during the E14→E17 saccular transition period decreased expression of multiple AT1 (Cav, Emp2, Hopx, Rage and Rtkn2), but not AT2 cell markers. This
indicates that differentiation of AT1 cells, which are derived from NKX2.1posHOPXpos AT1 progenitors is exquisitely sensitive to loss of WNT5a signaling. In the endothelial lineage, multiple markers were decreased in the mutant lungs (Cd31, Emcn and Tek). In addition, many genes related to myofibroblast cell differentiation and function were significantly decreased in Wnt5aCAG lungs. Therefore, Wnt5a signaling is a critical regulator of differentiation for multiple cell types during the saccular phase of lung development.

Figure 2. Wnt5a inactivation attenuates differentiation of AT1 and endothelial cells and myofibroblasts during the saccular stage of lung development. A–D: Relative mRNA for AT1 (A), AT2 (B), endothelial (C) and myofibroblast (D) markers in E18 Wnt5aCAG lungs (Wnt5a CAG) compared to control lungs by real-time RT-PCR analysis. Data represent the mean ± standard error of the mean (SEM). * indicates *p* < 0.05. *n* = 3.

3.2. WNT5a is Required for Myofibroblast Differentiation and Migration During Alveolar Stage Lung Development

Because germline Wnt5a−/− mice do not survive postnatally, the role of WNT5a in alveolar formation has remained unknown. To address this issue, we conditionally inactivated Wnt5a globally at the onset of postnatal alveolar development in Wnt5aCAG mice by administering tamoxifen on postnatal day 2 (P2). P2 inactivation of Wnt5a disrupted alveologenesis resulting in enlarged alveolar structures, reminiscent of a phenotype characteristic of the human neonatal chronic lung disease known as bronchopulmonary dysplasia (BPD). Morphometric quantification showed a significantly increased mean linear intercept, MLI, a measure of alveolar dimensions [29,30]. Analysis of cell differentiation showed no changes in expression of endothelial markers (Figure 3), nor in AT1 and AT2 cell markers (Figure 4). To determine whether the relative proportion of AT1 and AT2 cells was altered, we analyzed lung tissue immunostained for AT1 (HOPX) and AT2 (SFTPC) cell markers (Figure 4). The relative ratio of HOPXpos cells over total DAPIpos cells was not significantly altered in the Wnt5aCAG lungs (Control: 6.5% ± 1.1%; Mutant: 5.9% ± 1.0%, *p* = 0.69). Similarly, no significant
difference was observed in the ratio of SFTPC\textsuperscript{pos} cells (Control: 7.9% ± 0.8%; Mutant: 9.5% ± 1.3%, \( p = 0.30 \)).

**Figure 3.** Postnatal inactivation of Wnt5a attenuates alveologenesis. Top: Schematic of the experimental plan. A–D: Gross morphology of P12 control (A–B) and Wnt5a\textsuperscript{CAG} (C–D) lungs. Panels A and C show bright-field images. Panels B and D show green fluorescent images of control and Wnt5a\textsuperscript{CAG} embryos, respectively. Scale bars (D): 4 mm for A–D. E–F: H&E staining of P12 control (E) and Wnt5a\textsuperscript{d-AC} (F) lungs. E1 and F1 show higher magnification views of boxed areas (rotated 90 degree clockwise in E1) in E and F, respectively. Please note that distal alveolar units (*) surrounding the dividing alveolar space (a) are profoundly enlarged in the mutant lungs. Arrows indicate 2nd crests. Scale bars (F): 100 \( \mu \)m for E and F; 25 \( \mu \)m for E1 and F1. G: Mean linear intercept (MLI) of control and Wnt5a\textsuperscript{CAG} lungs (CKO). H: Relative mRNA for endothelial markers in Wnt5a\textsuperscript{CAG} lungs (Wnt5a CAG, P12-13) compared to the littermate control lungs by real-time RT-PCR analysis. Data represent the mean ± SEM. * indicates \( p < 0.05 \). \( n = 4 \).
Figure 4. No significant changes in AT1 and AT2 cell markers in P12 Wnt5aCAG lungs. A–B: Relative mRNA for AT1 (A) and AT2 (B) cell markers in Wnt5aCAG lungs (Wnt5a CAG, P12-13) compared to littermate control lungs by real-time RT-PCR analysis. C–H: Immunostaining of GFP (green) and HOPX (red) in P12 control (C, D, D1 and E) and Wnt5aCAG (F, G, G1 and H) lungs. D1 and G1 show higher magnification views of boxed areas in D and G, respectively. Arrows indicate HOPX-positive cells. Scale bar (H): 50 μm for C, D, E, F, G and H; 25 μm for D1 and G1. I–N: Immunostaining of GFP (green) and SFTPC (red) in P12 control (I, J, J1 and K) and Wnt5aCAG (L, M, M1 and N) lungs. J1 and M1 show higher magnification views of boxed areas (rotated 90 degree clockwise in M1) in J and M, respectively. Arrows indicate SFTPC-positive cells. Scale bar (N): 50 μm for I, J, K, L, M and N; 25 μm for J1 and M1. Data represent the mean ± SEM. * indicates p < 0.05. n = 4.

In contrast, multiple markers related to myofibroblast differentiation were reduced (Des: 0.81 ± 0.04, p < 0.05; Dbn1: 0.58 ± 0.05, p < 0.05; Tgfbi: 0.74 ± 0.10, p = 0.07; Fbln1: 0.60 ± 0.08, p < 0.05). Consistent with the latter data, immunostaining with antibodies against TAGLN and ACTA2, two markers of myofibroblast differentiation, revealed the number of TAGLNpos and ACTA2pos cells in the alveolar
region was decreased in the Wnt5a mutant lungs (Figure 5). In control lungs, ACTA2^pos^ cells were elongated, having a stretched cell shape. In contrast, ACTA2 staining was condensed in Wnt5a^CAG^ lungs. Both ACTA2 and TAGLN are expressed in alveolar myofibroblasts. We used TAGLN, which shows stronger perinuclear staining, to evaluate the abundance of alveolar myofibroblasts and found that the ratio of TAGLN^pos^ cells is significantly decreased in Wnt5a^CAG^ lungs (control: 12.7% ± 0.7%, mutant: 8.0% ± 0.7%, p < 0.05). However, TAGLN and ACTA2 staining around the blood vessels and airways appeared similar between the control and mutant lungs.

**Figure 5.** Postnatal inactivation of Wnt5a decreases expression of genes enriched in myofibroblasts. A: Relative mRNA for genes enriched in myofibroblasts in Wnt5a^CAG^ lungs (Wnt5a CAG, P12-13) compared to the littermate control lungs by real-time RT-PCR analysis. Data represent the mean ± SEM. * indicates p < 0.05. n = 4. B–G: Immunostaining of TAGLN (B and C) and ACTA2 (D–G) in P12 control (B, D and F) and Wnt5a^CAG^ (C, E and G) lungs. Panels F and G show higher magnification views of boxed areas in D and E, respectively. Nuclei are counterstained with DAPI. Arrowheads in B and C indicate TAGLN-positive perivascular smooth muscle cells. Block arrows in D and E indicate ACTA2 positive parabronchial smooth muscle cells. Arrows in F and G indicate ACTA2 positive alveolar myofibroblast cells. Scale bars (G): 50 μm for B and C, 100 μm for D and E, 25 μm for F and G.

The above data indicate that WNT5a signaling during the alveolar phase is required for myofibroblast differentiation. To confirm this finding, we conducted in vitro analyses using fibroblasts isolated from P5 neonatal Wnt5a^CAG^ and littermate Wnt5a^f/f^;Rosa26^tm1G^ (control) lungs. We activated CAG-cre^ER^ by treating cultured cells with tamoxifen for 24 hrs. As shown in Figure 6, several myofibroblast genes that were decreased in vivo were also reduced in cultured Wnt5a mutant cells, compared to the controls. These included Acta2 (0.73 ± 0.06, p < 0.05), Tagln (0.74 ± 0.03, p < 0.05) and...
These data support a necessary role for WNT5a signaling in directing differentiation of myofibroblast progenitors in the early postnatal period.

Figure 6. Inactivation of Wnt5a in vitro disrupts differentiation and migration of cultured fibroblasts. Fibroblasts from P5 neonatal lungs were isolated, cultured and treated with tamoxifen (1.5 μg/mL) for 24 h before RNA isolation. A: Relative mRNA for myofibroblasts-enriched genes in Wnt5aCAG fibroblasts compared to control lung fibroblasts determined by real-time RT-PCR analysis. B: Relative mRNA of cell migration genes in Wnt5aCAG fibroblasts compared to control lung fibroblasts determined by real-time RT-PCR analysis. Data represent the mean ± SEM. * indicates p < 0.05. n = 4. C–D: Transwell migration assay. Control (C) and Wnt5aCAG (D) fibroblasts treated with tamoxifen (1.5 μg/mL, 24 h) were plated on 12-micron Transwell membranes and cultured for 24 h. Cells migrated to the lower side of the Transwell were stained with crystal violet and counted. E: Quantification of the number of cells that migrated to the lower side in unit area. Four areas from each experiment (each Transwell) were imaged, counted and the means calculated. Data represent the mean of three independent experiments ± SEM. * indicates p < 0.05.

We recently showed that conditional targeting of Pdgfra in SCMF during alveologenesis also disrupts alveolar formation resulting in a phenotype similar to that of Wnt5aCAG lungs [25]. RNA-seq analysis of the Pdgfra mutant SCMF revealed cell migration as one of the top affected cellular functions, reflected as altered expression of multiple cell migration genes [25]. In the present study, we selected several cell migration genes that were disrupted in the Pdgfra mutant SCMF (Table S2) and analyzed their expression in fibroblasts isolated from P5 control and Wnt5aCAG lungs. As shown in Figure 6, we found decreased expression of Actc1 (0.80 ± 0.05, p < 0.05), Lum1 (0.64 ± 0.09, p < 0.05), Cyr61 (0.53 ± 0.18, p = 0.08) and Cnn1 (0.72 ± 0.03, p < 0.05) in Wnt5a mutant cells, suggesting WNT5a plays a role in lung myofibroblast migration. To further test this possibility, we examined the migration of control and Wnt5aCAG fibroblasts in vitro using the Transwell cell culture. As shown in
Figure 6, many control fibroblasts migrated to the lower side of the Transwell after 24 h of culture. In contrast, we noted only a few trans-migrated mutant fibroblasts indicating defective migration of \( Wnt5a^{\text{CAG}} \) myofibroblasts (control: 56.2 ± 15.5, mutant: 9.2 ± 2.9 in unit area).

3.3. RORs are Essential for Myofibroblast Migration During Alveolar Stage Lung Development

The sum of the above observations indicates that myofibroblasts are a major target of WNT5a signaling and suggest that defects in this cell lineage may be the major cause of abnormal alveologenesis in \( Wnt5a^{\text{CAG}} \) lungs. To test this potential mechanism, we blocked WNT5a signaling by targeting its receptors \( \text{Ror1} \) and \( \text{Ror2} \) in SCMF using \( \text{Gli1-cre}^{\text{ERT2}} \). Previously we showed that \( \text{Gli1-cre}^{\text{ERT2}} \) targets SCMF during mouse alveologenesis [6,25,31]. We therefore generated mice with the compound genotype, \( \text{Gli1-cre}^{\text{ERT2}};\text{Rosa26}^{\text{mTmG}};\text{Ror1}^{\text{f/f}};\text{Ror2}^{\text{f/f}} \) (\( \text{RorGli;GFP} \)) and activated the \( \text{Gli1-cre}^{\text{ERT2}} \) by tamoxifen on P2. Lungs from \( \text{RorGli;GFP} \) and \( \text{Rosa26}^{\text{mTmG}};\text{Ror1}^{\text{f/f}};\text{Ror2}^{\text{f/f}} \) control neonatal mice were characterized on P13 (Figure 7). Postnatal inactivation of \( \text{Ror1} \) and \( \text{Ror2} \) in SCMF resulted in a phenotype nearly identical to that of postnatal \( Wnt5a \) inactivation in \( Wnt5a^{\text{CAG}} \) mice. Characterization of \( \text{RorGli;GFP} \) lungs showed profound enlargement of alveoli, indicating disruption of alveolar formation. To determine the impact of \( \text{Ror} \) inactivation in the \( \text{Gli1-cre}^{\text{ERT2}} \) targeted cells, we conducted double immunostaining for GFP and \( \text{ACTA2} \) in \( \text{RorGli;GFP} \) and \( \text{Gli1-cre}^{\text{ERT2}};\text{Rosa26}^{\text{mTmG}} \) (\( \text{mTmGGli} \), control) lungs. Consistent with our previous observations, \( \text{Gli1-cre}^{\text{ERT2}} \) labeled mainly SCMF that are scattered within the alveolar region and are \( \text{ACTA2}^{\text{pos}} \) at this time point. The majority of GFP\(^{\text{pos}} \) cells in the control lungs were localized to the tip of the SC. In \( \text{RorGli;GFP} \) lungs, the majority of GFP\(^{\text{pos}} \) cells were still localized within the primary septa, indicating that lack of \( \text{Ror1} \) and \( \text{Ror2} \) decreased myofibroblast cell migration in the \( \text{RorGli;GFP} \) lungs. Consistent with this conclusion, expression of the myofibroblast genes \( \text{Des} \) (0.76 ± 0.02, \( p < 0.05 \)) and \( \text{Fbln1} \) (0.74 ± 0.06, \( p = 0.05 \)) and cell migration genes \( \text{Actc1} \) (0.66 ± 0.10, \( p = 0.07 \)), \( \text{Bmp2} \) (0.68 ± 0.12, \( p = 0.11 \)) and \( \text{Lum} \) (0.68 ± 0.03, \( p < 0.05 \)) were collectively decreased in the \( \text{RorGli;GFP} \) lungs. Thus, signaling thru ROR1/ROR2, the cognate WNT5a receptors is required for normal migration of SCMF and the process of alveologenesis.
Figure 7. Postnatal inactivation of Ror1 and Ror2 attenuated alveologenesis. Top: Schematic of the experimental plan. A–B: H&E staining of P13 control (A) and RorGli;GFP (B) lungs. C: Mean linear intercept (MLI) of P13 control and RorGli;GFP (CKO) lungs. D–K: Immunostaining of GFP (green) and ACTA2 (red) in P13 mTmGli (D–G, control) and RorGli;GFP (H–K) lungs. Panels G and K show higher magnification views of boxed areas in F and J, respectively. Nuclei are counterstained with DAPI. Scale bars (K): 200 μm for A and B, 50 μm for D–F and H–J, 11 μm for G and K. L–M: Relative mRNA for myofibroblast enriched genes and cell migration genes in P13 RorGli;GFP (Ror Gli) lungs compared to control lungs by real-time RT-PCR analysis. Data represent the mean ± SEM. * indicates p < 0.05. n = 3.

3.4. Relevance to Human Neonatal Lung Disease

Both Wnt5aCAG and RorGli;GFP lungs represent a phenotype of arrested alveologenesis similar to the histological characteristic of human BPD. Expression or disruption of WNT5a in BPD lungs has not been previously reported. Therefore, to ascertain the relevance of our findings in the mouse to human BPD, we determined the expression of WNT5a in lungs of human patients who died with BPD. As shown in Figure 8, WNT5a was significantly lower in the BPD lungs (# 11, #14, #17, #18, #44)
as compared to lungs from the non-BPD patients (#30, #50, #52, #56) who died due to non-pulmonary causes. No significant difference was observed in ROR1 and ROR2 expression.

**Figure 8.** Expression of WNT5a, ROR1 and ROR2 in human bronchopulmonary dysplasia (BPD) lungs. Human lung samples from deceased donors were collected and processed at the University of Rochester following the protocol approved by University of Rochester Institutional Review Board. Relative mRNA for the examined genes is presented in a 2-D column chart with levels of #56 arbitrarily set as “1”. Data distribution of the non-BPD (#30, #50, #52, #56) and the BPD (#11, #14, #17, #18, #44) samples is shown by the boxplot. * indicates \( p < 0.05 \).

4. Discussions

Late lung development that encompasses the periods of saccular and alveolar development are especially important as they include the cellular and molecular processes required for lung functional maturation as an efficient gas exchange organ at birth and throughout life. The role of WNT5a, a non-canonical WNT, in late lung development is unknown. In the present study, we generated and utilized conditional loss-of-function genetic models to examine the temporal-specific role of WNT5a in the saccular and alveolar phases of lung development. We demonstrate that finely tuned regulation of WNT5a is critical for both structural and cellular differentiation in the lung. In the saccular phase, WNT5a regulates differentiation of both mesodermal and epithelial lineages. Most importantly, we found that in the alveologenesis phase, inactivation of Wnt5a interrupts alveolar formation by disrupting differentiation and migration of the key mesodermally derived SCMF. The significance and relevance of these findings are highlighted by the fact that alveolar defects are a hallmark of many human neonatal and adult lung diseases.

The key findings of the present study are that WNT5a regulation of late lung development is stage and cell-type specific. Conditional inactivation of Wnt5a in the saccular phase resulted in defects in structural development of the lung that appeared to be caused by failure in distal airway expansion (Figure 1). This is reminiscent of our previously reported results using a conventional germline Wnt5a loss-of-function model [10]. In the saccular model, we also found significantly reduced AT1 cell differentiation. Five of the seven tested AT1 cell markers were significantly decreased in the Wnt5aCAG lungs (Figure 2), reflecting an important role of WNT5a in AT1 cell differentiation during the saccular phase of lung development. Interestingly, expression of most AT1 cell markers was not significantly affected by Wnt5a inactivation during alveologenesis (Figure 4), suggesting that the role of WNT5a in AT1 cell differentiation may be stage dependent. During the saccular phase of lung development, AT1 progenitors (NKX2.1posHOPXpos) and AT2 progenitors (NKX2.1posSFTPCpos) begin to give rise to mature AT1 and AT2 cells, respectively [2]. As lung development enters the alveolar phase, a subpopulation of AT2 cells maintains progenitor properties. These alveolar epithelial progenitors (AEP) are SFTPCposAXIN2pos and are capable of self-renewal or differentiation into mature AT1 and
AT2 cells [32]. Results from the current study demonstrate that it is the saccular phase events, which are exquisitely sensitive to lack of WNT5a signaling (Figure 9). Interestingly, in adult lungs Wnt5a is expressed in the AT2 stem-cell niche (Axin2pos AT2 cells), indicating a potential role for WNT5a in AT2 stem-cell maintenance [9]. Deletion of Wnt5a with Tbx4LME-cre was shown to moderately reduce AT2 stem cells but the reduction was not significant [9]. Recently Wu and colleagues showed that WNT5a represses the growth of alveolar epithelial progenitors in both lung-slice and organoid cultures [33]. Collectively, these data demonstrate a stage-dependent functional importance of WNT5a in lung development and stem-cell maintenance.

**Figure 9.** A simplified model indicating the role of WNT5a in AT1 and AT2 cell differentiation. During the saccular stage, WNT5a activity is required for differentiation of AT1 cells from NKX2.1posHOPXpos AT1 progenitors. During alveologenesis, WNT5a activity does not appear to be required for AT1 and AT2 cell differentiation. AEP: alveolar epithelial progenitors.

The present study revealed that differentiation and morphogenesis of mesodermally derived cells are also sensitive to lack of WNT5a signaling in a temporally specific manner. Wnt5a inactivation reduced endothelial markers only in the saccular stage, but not during alveologenesis. Formation of capillary networks, which commences during the saccular phase, is regulated by the interaction between endothelial cells and pericytes. A recent study demonstrated that WNT5a from endothelial cells is important for pericyte recruitment [11]. Interestingly, our present study showed that expression of the pericyte marker Pdgfrb was reduced by Wnt5a inactivation during both the saccular and alveolar phases of lung development. Abnormal capillary distribution was also found in the germline deletion model of Wnt5a [10]. A number of previous studies have demonstrated that WNT5a induces endothelial cell migration through non-canonical WNT signaling pathways [15,34,35].

The major impact of Wnt5a inactivation during alveologenesis is on myofibroblast differentiation. As shown in Figure 5, multiple myofibroblast markers that included Acta2, Tagln, Des, Dbn1, Tgfbi and Fbln1 were reduced in neonatal Wnt5aCAG lungs. In contrast, markers for other fibroblast cell types, such as lipofibroblasts (Adrp), was not significantly altered. This indicates that myofibroblasts are a major target of WNT5a signaling during alveologenesis, and Wnt5a deficiency disrupts their differentiation and migration. A key myofibroblast cell type with a critical role in alveologenesis is SCMF. In support of our findings, we found that blocking WNT5a signaling via targeting its receptors, Ror and Ror2, in SCMF (Figure 7) arrested alveologenesis. While the expression of myofibroblast markers was only moderately reduced in the RorsGli;GFP lungs (Figure 7), this is likely because Gli1creERT2 does not target all ACTA2pos myofibroblasts (Figure 7, Panels D-F). Targeting Rors in SCMF also did not completely inhibit their differentiation, as some of the GFPpos cells were still
positive for ACTA2. Nevertheless, because of the critical role of SCMF, this level of gene-targeting was sufficient to cause a significant defect in alveologenesis, indicating the importance of WNT5a-ROR signaling in SCMF function during neonatal lung development.

Our recent study demonstrated that PDGFRA-mediated elastogenesis is important for lung maturation [25]. However, we found no evidence by RNA-seq analysis for alterations in Wnt5a expression in Pdgfra mutant SCMF. In addition, Pdgfra expression was not reduced in the Wnt5αCAG lungs (Figure 5). This indicates that PDGFRA signaling and WNT5a signaling may regulate alveologenesis via separate pathways. Interestingly, inactivation of Wnt5a, Ror1 and Ror2, as well as Pdgfra all decreased the elastogenic gene Fbln1 (Figures 5 and 7) [25], indicating that the two pathways may cross-talk over key mediators of alveologenesis, such as Fbln1.

BPD is a morbid and sometimes lethal chronic lung disease in human prematurely born neonates [36]. Histological analysis of lung tissue from infants who died with BPD is consistent with a morphology characteristic of arrested alveolar formation [37]. While a number of molecules have been associated with BPD, causative factors in its pathogenesis remain unknown. A key reason is paucity of data on mechanisms of alveolar formation. The findings of the present study show that WNT5a is causally related to alveologenesis, the process that is arrested in human BPD. To examine the clinical relevance of the mouse findings, we show that WNT5a is also reduced in human BPD lung samples. While expression or the potential role of WNT5a in BPD has not been directly elucidated, multiple studies reported elevated nuclear β-catenin in both rodent hyperoxia BPD-phenocopy models and human BPD tissue [38,39]. Moreover, it has been shown that neonatal hyperoxia increases nuclear β-catenin and decreases AT2 to AT1 cell differentiation [40]. WNT5a inhibits β-catenin-mediated WNT signaling [6,12,28,33]. Thus, it is likely that reduced levels of WNT5a may contribute to increased canonical WNT signaling and pathogenesis of BPD.

Supplementary Materials: Figure S1: Craniofacial defects of E18 Wnt5aCAG embryos. Table S1: Expression of cell type specific markers during saccular and alveolar phases of lung development (References: LungGENS: https://research.cchmc.org/pbge/lunggens; [32,41,42]). Table S2: Changes of cell migration genes in Pdgfra deficient SCMF identified by RNAseq analysis (GSE126457, [25]. Table S3: Antibody information. Table S4: Sequences of real-time RT-PCR primers.

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