

Research Article

Role of *HIF-1 α -miR30a-Snai1* Axis in Neonatal Hyperoxic Lung Injury

Yuhao Zhang, Xiaoyu Dong, and Krithika Lingappan 

Department of Pediatrics, Section of Neonatology, Texas Children's Hospital, Baylor College of Medicine, Houston, Texas, USA

Correspondence should be addressed to Krithika Lingappan; lingappa@bcm.edu

Received 21 April 2019; Revised 5 September 2019; Accepted 19 September 2019; Published 22 October 2019

Guest Editor: Lynette K. Rogers

Copyright © 2019 Yuhao Zhang et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Bronchopulmonary dysplasia (BPD) is characterized by a severe impairment in lung alveolarization and vascular development. We have previously shown that pulmonary angiogenesis is preserved in hyperoxia-exposed female mice accompanied by increased *miR-30a* expression, which is a proangiogenic miRNA. Also, *miR-30a* expression is decreased in human BPD. HIF-1 α plays an essential role in postnatal lung development, especially in recovery from hyperoxic injury. *Snai1* activation promotes pathological fibrosis through many mechanisms including Endo-MT, which may in turn adversely impact lung vascular development. Our objective was to test the hypothesis that higher *miR-30a* expression through *HIF-1 α* decreases *Snai1* expression in females and attenuates injury in the developing lung. Neonatal male and female mice (C57BL/6) were exposed to hyperoxia (P1-5, 0.95 FiO₂) and euthanized on P21. Neonatal human pulmonary microvascular endothelial cells (HPMECs; 18-24-week gestation donors; 3/group either sex) were subjected to hyperoxia (95% O₂ and 5% CO₂) or normoxia (air and 5% CO₂) up to 72 h. *Snai1* expression was measured in HPMECs *in vitro* and in neonatal mouse lungs *in vivo*. Also, *Snai1* expression was measured in HPMECs after *miR-30a* mimic and *miR-30a* inhibitor treatment. To further establish the potential regulation of *miR-30a* by *Hif-1 α* , *miR-30a* expression after *Hif-1 α* inhibition was measured in HPMECs. *In vivo*, *Snai1* expression was decreased in neonatal female lungs compared to males at P7. Increased *Snai1* expression was seen in male HPMECs upon exposure to hyperoxia *in vitro*. Treatment with the *miR-30a* mimic decreased *Snai1* expression in HPMECs, while *miR-30a* inhibition significantly increased *Snai1* expression in HPMECs. siRNA-mediated loss of *Hif-1 α* expression in HPMECs decreased *miR-30a* expression. *Hif-1 α* may lead to differential sex-specific *miR-30a* expression and may contribute to protection from hyperoxic lung injury in female neonatal mice through decreased *Snai1* expression.

1. Introduction

With the increasing survival of extremely premature babies, the incidence of bronchopulmonary dysplasia (BPD) has remained steady, despite the advances in neonatal intensive care. BPD is the cause of significant morbidity and leads to prolonged impairment in lung function in this group of patients. Many prenatal and postnatal factors are involved in the pathogenesis of BPD, and exposure to supraphysiological concentrations of oxygen contributes to its development. Murine models have utilized varying durations of postnatal hyperoxia exposure to simulate the human disease in neonatal mice [1, 2].

Bronchopulmonary dysplasia (BPD) is characterized mainly by an arrest in lung development with severe impair-

ment of alveolar septation and vascular development, and pathological fibrosis in severe cases [3–6]. Histopathological analysis in neonates with “new BPD” shows evidence of variable interstitial fibroproliferation compared to extensive fibroproliferation in “old BPD” [7]. BPD disproportionately affects male infants compared to females. The molecular mechanisms underlying this sex bias are not known. We have previously shown that alveolarization and pulmonary angiogenesis is preserved in hyperoxia-exposed female neonatal mice compared to males [8]. The protective effect in females is accompanied by an increase in pulmonary *miR-30a* expression in females [9]. *miR-30a* has proangiogenic, anti-inflammatory, and antifibrotic effects in many diseases processes [10–17], including those affecting the lung. Significantly, *miR-30a* expression is decreased in human patients

with BPD [9]. The upstream regulation of *miR-30a* leading to differential expression in males and females has not been elucidated. Previously published work has shown that *Hif-1 α* may increase *miR-30a* expression [18]. HIF-1 α plays a vital role in postnatal lung development [19], especially in recovery from hyperoxic injury [20] and acute lung injury [21]. Overexpression of *Hif-1 α* in hyperoxia-exposed neonatal mice attenuates lung injury [20]. *HIF-1 α* binding to its target genes is higher in female lungs after hyperoxia exposure [22]. Whether *HIF-1 α* increases *miR-30a* expression in the pulmonary microvascular endothelium is not known. *Snail* is a transcriptional repressor and a profibrotic molecule [23–28]. Studies have shown that *miR-30a* downregulates *Snail* [10, 14, 17].

However, the mechanistic role of the intersection between HIF-1 α , miR-30a, and *Snail* in neonatal hyperoxic lung injury has not been studied. Our objective was to test the hypothesis that HIF-1 α increases the miR-30a expression in females and decreases the *Snail* expression in the developing lung.

2. Methods

2.1. Animals. All experiments were performed per relevant guidelines and regulations. The care of animals was as per the 8th edition of the guide for the care and use of laboratory animals and other IACUC protocols. Timed pregnant C57BL/6J WT mice were obtained from Charles River Laboratories (Wilmington). The sex in neonatal mouse pups was determined as described before and with PCR analysis for the *Sry* gene [8].

2.1.1. Mouse Model of BPD. Mouse pups were exposed to normoxia (21% O₂) or hyperoxia (95% O₂), within 12 h of birth for five days. Neonatal mice are at the sacular stage of lung development during this period, equivalent to 26–36 weeks in human gestation. The dams were rotated between air- and hyperoxia-exposed litters every 24 hours to prevent oxygen toxicity in the dams. Oxygen exposure was conducted in plexiglass chambers as previously described [8]. Mice were euthanized on P7 and P21 (after recovery in room air). The mice were euthanized with sodium pentobarbital, 100 mg/kg, i.p. For the *in vivo* experiments, neonatal mice from different litters were randomly allocated to room air or hyperoxia. The experimental unit was a single neonatal mouse pup and data from 4–6 animals/group were used for analysis.

2.1.2. Cell Culture and Hyperoxia Treatment. Neonatal human pulmonary microvascular endothelial cells (HPMEC; male: #10885, #11367, and #10899; female: #5016, #10160, and #10169; 6 biological replicates in total) were purchased from ScienCell and maintained in Endothelial Cell Medium (Cat # 1001, ScienCell) at 37°C in 5% CO₂. The gestational age of the donors varied from 18 to 24 weeks. Male and female HPMECs were used from passages 3 to 5 to ensure their endothelial characteristics. Male or female HPMEC cells (1 × 10⁵) were seeded in a 6 mm dish. 24 h later, these cells were incubated at 37°C in room air condition (21% O₂, 5% CO₂) or in hyperoxia (95% O₂, 5% CO₂) as described

before for 72 h. For measuring *SNAIL* expression, both male and female HPMECs were used. For the rest of the *in vitro* experiments, female HPMECs were used as the miR-30a expression was increased only in female HPMECs [9].

2.1.3. miRNA and Plasmid Transfection in HPMECs. miR30a-5p mimic (HMI0454), miR30a-5p inhibitor (HSTUD0454), Hif-1 α -siRNA (3017635209), and negative control siRNA (NCSTUD001) were purchased from MilliporeSigma. Hif-1 α overexpression plasmid (#18948) was purchased from Addgene. pcDNA3.1 (V79020) plasmid was purchased from Thermo Fisher. MicroRNA negative control (30 pmol), miR30a-5p mimic (30 pmol), or miR30a-5p inhibitor (30 pmol) were used for HPMEC transfection separately using a Lipofectamine 2000 reagent (Cat #11668019; 6 μ l per well of a 6-well plate). Hif-1 α plasmid (2 μ g) with or without miR30a-5p inhibitor (60 pmol) was transfected using a Lipofectamine 2000 reagent (6 μ l per well of a 6-well plate). pcDNA3.1 plasmid (2 μ g/well) was transfected as a negative control of Hif-1 α . After six hours, fresh medium was added to the plate. After 48 h incubation in room air (21% O₂, 5% CO₂), cells were harvested for later use.

2.1.4. Quantitative PCR. Total RNA and microRNA was extracted from the lung tissues and cell lines using TRIzol (Cat #15596026, Thermo Fisher) and chloroform (Cat #C2432, Millipore-Sigma) and then treated with DNase I (Cat #K1622, Invitrogen). cDNA was prepared using a RevertAid Reverse Transcriptase (Thermo Fisher). MicroRNA cDNA was generated using a MystiCq microRNA cDNA Synthesis Mix (MilliporeSigma). Quantitative PCR was performed using the QuantStudio 7 Flex Real-Time PCR Detection System (Thermo Fisher) and SYBR Green (Cat #1725274, Bio-Rad). The thermal cycling conditions used were as follows: one cycle at 95°C for 1 min, 40 cycles at 95°C for 15 s, and one cycle at 60°C for 30 s. The primers used in the real-time PCR test were listed as follows: *Snail* (NM_005985, human) forward primer: GCGAGCTGCAG GACTCTAAT, reverse primer: GGACAGAGTCCCAGAT GAGC; *Snail* (NM_011427, mice) forward primer: CACC CTCATCTGGGACTCTC, reverse primer: GAGCTTTTG CCACTGTCCCTC; Hif-1 α (NM_001243084, human) forward primer: GATGTAATGCTCCCCTCACC, reverse primer: CTTGATTGAGTGCAGGGTCA; Hif-1 α (NM_001313919, mice) forward primer: ATTCTCCAAGCCCT CCAAGT, reverse primer: TCATCAGTGGTGGCAGTTG T; β -actin (NM_001101, human) forward primer: CATC GAGCACGGCATCGTCA, reverse primer: TAGCACAGC CTGGATAGCAAC; β -actin (NM_007393, mice) forward primer: GATCTGGCACCACACCTTCT, reverse primer: GGGGTGTTGAAGGTCTCAA; beta 2-microglobulin (NM_009735.3, mouse) forward primer: CTGACCGGCCT GTATGCTAT, reverse primer: CCGTTCTTACGATTT GGAT; and beta 2-microglobulin (NM_00408.3, human) forward primer: TGCTGTCTCCATGTTTGATGTATCT, reverse primer: TCTCTGCTCCCCACCTCTAAGT. miR30a-5p (MIRAP00079), MystiCq Universal PCR (MIRUP), and U6 (MIRCP00001) primers were purchased

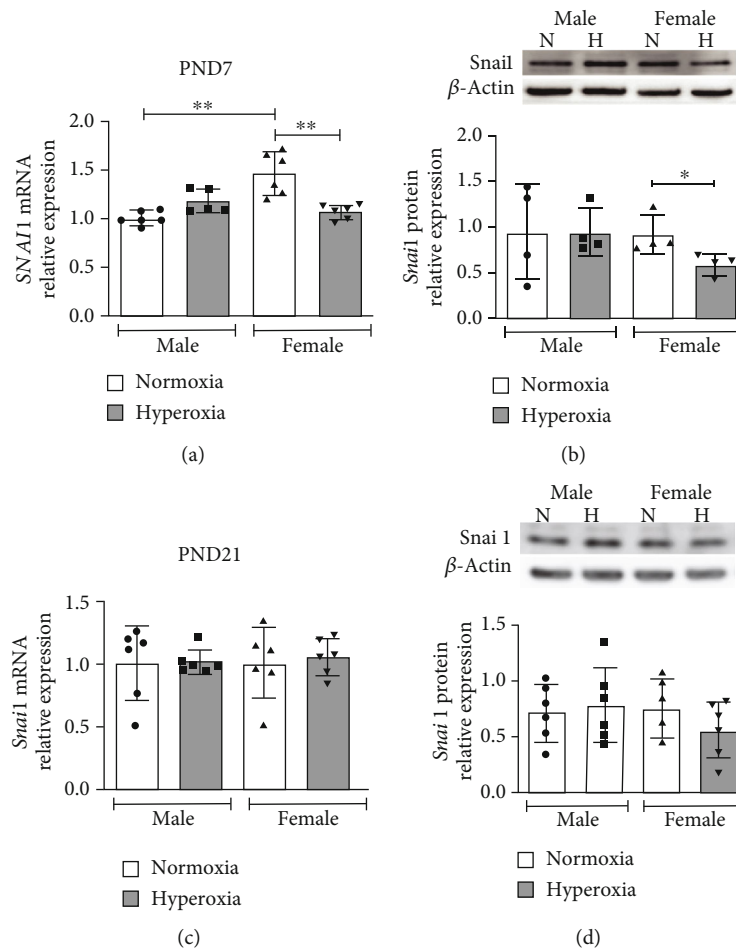


FIGURE 1: Differential sex-specific expression of *Snail* in neonatal mice after postnatal hyperoxia exposure: *Snail* mRNA (β -actin as housekeeping gene) and protein expression neonatal male and female mice exposed to hyperoxia (P1-5, 0.95 FiO₂) during the saccular stage of lung development. *Snail* mRNA (a, c) and protein expression (b, d) were measured in the lungs at P7 (a, b) immediately after hyperoxia exposure and at P21 (c, d) after recovery in normoxia. Values are means \pm SD. $n = 5 - 6$ mice per group. Significant differences between groups indicated by * $P < 0.05$ and ** $P < 0.01$.

from MilliporeSigma. Relative mRNA levels were calculated using the $2^{-\Delta\Delta CT}$ method and normalized by β -actin or beta 2-microglobulin in the same sample.

2.1.5. Western Immunoblotting. Protein was isolated from murine lung tissue and HPMECs using RIPA buffer (Thermo Fisher) containing protease mixture inhibitors (Thermo Fisher). Proteins were separated by 4–12% sodium dodecyl sulfate polyacrylamide gel electrophoresis, then transferred to a PVDF membrane using a Mini-PROTEAN Tetra Cell system (Bio-Rad). The following primary antibodies were used: goat anti-Snail (1:1000, Abcam, ab53519), rabbit anti- β -actin (1:5000, Cell Signaling, 4970), and rabbit anti-vinculin (1:1000, Cat#4650, Cell Signaling). Pierce ECL plus a Western blotting substrate (Cat #32132, Thermo Fisher) was used for visualizing immunoreactive protein bands. The protein bands were normalized by β -actin or vinculin on the same membrane.

2.1.6. Statistical Analysis. GraphPad version 7 was used for the analysis of our data. Data are expressed as means \pm SD.

Data were analyzed by two-way ANOVA to test for the independent effects of sex and hyperoxia and to look for any interaction (sex \times hyperoxia) or by Mann-Whitney U test. Multiple-comparison testing (Bonferroni) was performed if statistical significance ($P < 0.05$) was noted by ANOVA.

3. Results

3.1. Differential Sex-Specific Expression of *Snail* in Neonatal Mice after Postnatal Hyperoxia Exposure. WT male and female neonatal mice were exposed to hyperoxia (0.95 FiO₂ from P1-5) during the saccular stage of lung development, and *Snail* mRNA and protein expression were measured at P7 (early) and P21 (during recovery in normoxia) and compared to respective normoxic controls. *Snail* mRNA expression (β -actin as housekeeping gene) was decreased in female neonatal mice at P7 after hyperoxia exposure compared to female normoxic controls. No difference in expression was seen in males (Figure 1(a)). The results were similar with another housekeeping gene (β 2-microglobulin) as shown in Supp. Figure 1. Similarly, Snail

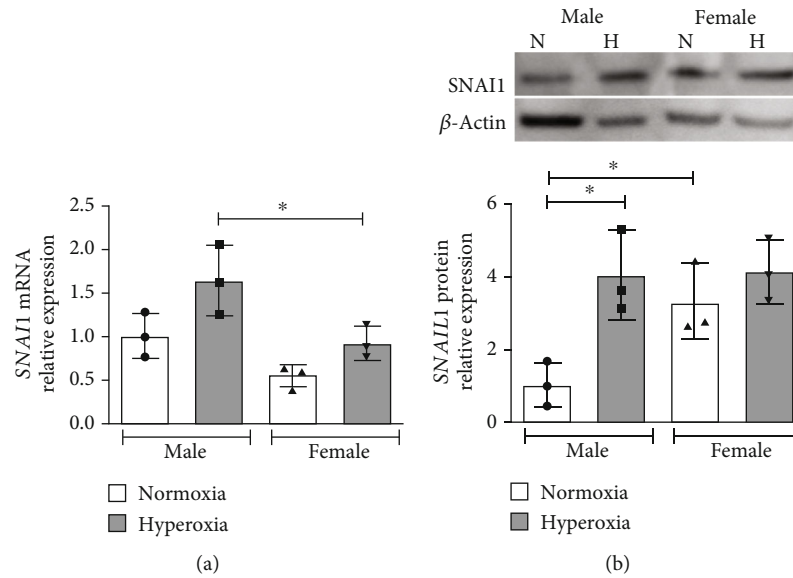


FIGURE 2: *Snail* mRNA and protein expression is increased in human pulmonary microvascular endothelial cells upon exposure to hyperoxia: *SNAIL* mRNA (a) and protein expression (b) were measured in neonatal human pulmonary microvascular endothelial cells exposed to normoxia or hyperoxia (0.95 FiO₂ up to 72 hrs). Values are mean \pm SD. $n = 3$ per group. Significant differences between indicated groups are indicated by * $P < 0.05$.

protein expression in the lungs was also decreased in females compared to female normoxic controls and compared to hyperoxia-exposed male mice (Figure 1(b)). *Snail* mRNA and protein expression did not show statistically different expression levels between any of the groups at P21 (Figures 1(c) and 1(d)). Protein expression was repeated with vinculin as the loading control; female mice tended to have lower *Snail* protein in the lungs at P7 compared to males, and in HPMECs, males tended to have a greater increase in *Snail* expression. These results however, were not statistically significant (Supp. Figure 1).

3.2. *Snail* mRNA and Protein Expression Are Increased in Male Human Pulmonary Microvascular Endothelial Cells upon Exposure to Hyperoxia. Next, we wanted to elucidate the response to hyperoxia in neonatal human pulmonary microvascular endothelial cells (HPMECs). These cells were exposed to hyperoxia (0.95 FiO₂) for up to 72 hours, and *SNAIL* mRNA (Figure 2(a)) and protein expression (Figure 2(b)) were measured. *SNAIL* mRNA expression was significantly higher in male HPMECs compared to female. Male HPMECs showed increased *SNAIL*1 protein upon exposure to hyperoxia, while female HPMECs did not show this change.

3.3. *Snail* Regulation by *miR-30a*. TargetScan reported *miR-30a* binding sites 630-637 bp in the 3'UTR of *SNAIL* (Figure 3(a)). *miR30a-5p* expression was increased in female HPMECs using the *miR30a-5p* mimic (Figure 3(b)). Following treatment with the mimic, *miR30a-5p* expression was increased in female HPMECs. *SNAIL* mRNA (Figure 3(c)) and protein expression (Figure 3(d)) were decreased following *miR30a-5p* overexpression. In contrast, *SNAIL* protein expression was increased following treatment with a *miR30a-5p* inhibitor (Figure 3(e)).

3.3.1. Regulation of *miR30a-5p* by *HIF-1 α* . We have previously shown that *Hif-1 α* expression was increased in female mice at P7 and decreased in male mice at P21 after hyperoxia exposure [29] and that *HIF-1 α* binding to its target genes is more significant in female lungs after hyperoxia exposure [22]. *HIF-1 α* protein expression was decreased in both male and female HUVECs upon exposure to hyperoxia [29]. Motif analysis revealed a hypoxia response element (HRE) [AG]CGTG site 428 base pairs upstream of *miR-30a* (Figure 4(a)). To further establish the potential regulation of *miR-30a* by *Hif-1 α* , we used the siRNA-mediated loss of the *Hif-1 α* expression in female human pulmonary microvascular endothelial cells and its effect on *miR30a-5p* expression. Treatment of female HPMECs with *HIF-1 α* siRNA significantly decreased *HIF-1 α* expression (Figure 4(b)). Also, the expression of *miR30a-5p* was decreased in pulmonary microvascular endothelial cells transfected with *HIF-1 α* siRNA compared to cells treated with scrambled siRNA (Figure 4(c)).

3.3.2. *SNAIL* Expression following *HIF-1 α* Silencing in Female Human Pulmonary Microvascular Endothelial Cells. We next wanted to see if *HIF-1 α* played a role in regulating *SNAIL* expression in human pulmonary microvascular endothelial cells. Following the silencing of *HIF-1 α* in female HPMECs, *SNAIL* mRNA (Figure 5(a)) and protein expression (Figure 5(b)) were decreased in female HPMECs.

3.3.3. *SNAIL* Expression following *miR30a-5p* Inhibition, *HIF-1 α* Overexpression, or Both in Female HPMECs. Since both *miR30a-5p* and *HIF-1 α* had opposing effects on *SNAIL* expression, we wanted to determine the independent and additive effects of *miR30a-5p* inhibition and *HIF-1 α* overexpression in female HPMECs. *HIF-1 α* overexpression and *miR30a-5p* inhibition both independently increased *SNAIL* mRNA expression in female HPMECs (Figure 6(a)).

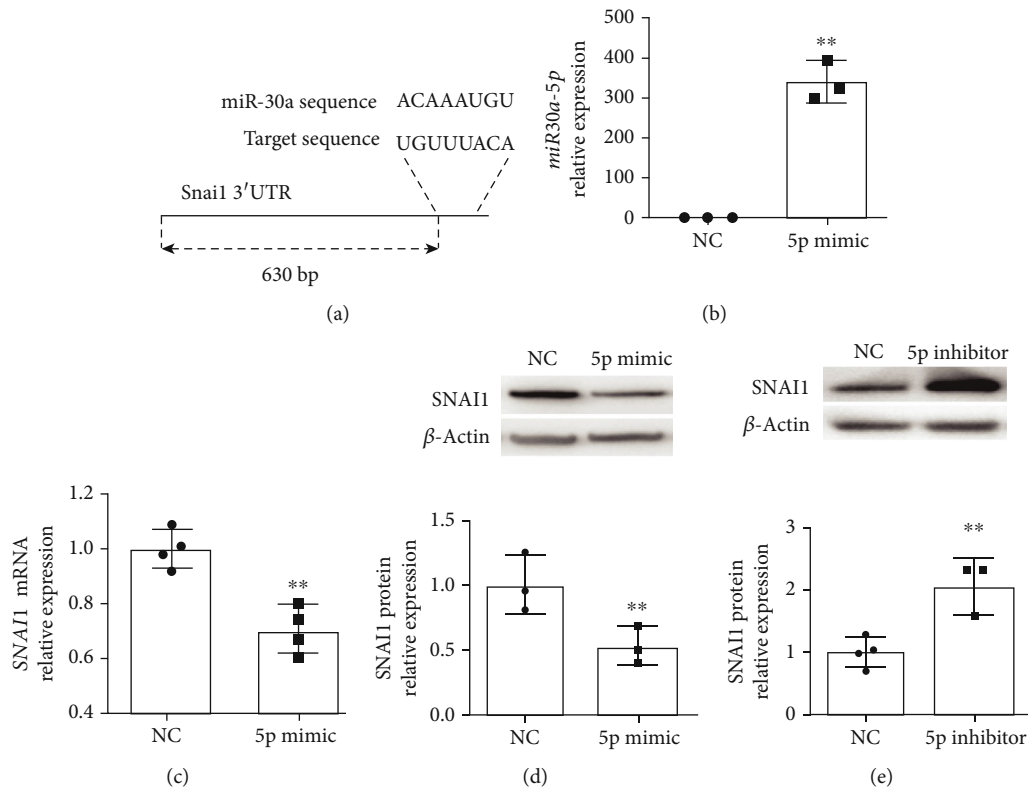


FIGURE 3: *miR30a-5p* decreases SNAIL expression in female neonatal human pulmonary microvascular endothelial cells: (a) TargetScan reported *miR-30a* binding sites 630-637 bp in the 3'UTR of SNAIL. Treatment with the *miR30a-5p* mimic increased the *miR30a-5p* expression (b) in HPMECs. SNAIL mRNA (c) and protein (d) expressions were decreased in neonatal HPMECs after *miR30a-5p* overexpression. In contrast, treatment with the *miR30a-5p* inhibitor increased SNAIL protein (e) expression ($n = 3 - 4$ experimental replicates/group). Values are means \pm SD. Significant differences between nontreated controls are indicated by * $P < 0.05$ and ** $P < 0.01$.

HIF-1 α overexpression had a more significant effect on SNAIL expression compared to *miR30a-5p* inhibition. Combined *HIF-1 α* overexpression and *miR30a-5p* inhibition increased SNAIL expression compared to controls, but there was no additive effect (Figure 6(a)). SNAIL protein was increased by *miR30a-5p* inhibition or *HIF-1 α* overexpression independently and by the combined treatment with *miR30a-5p* inhibition and *HIF-1 α* overexpression compared to controls. No statistically significant changes in protein levels were noted between the individual treatment groups (Figure 6(b)).

4. Discussion

Sex-specific differences exist in many neonatal morbidities including bronchopulmonary dysplasia (BPD), which adversely affects a significant number of extremely premature newborns. Elucidation of mechanisms responsible for the sex-specific differences in the development of this disease is critical for the development of individualized therapies. We identified miR-30a as a potential modulator of many of the genes differentially expressed in the angiogenesis pathway [9]. Interestingly, miR-30a expression was increased in the female lung after exposure to hyperoxia. These findings were replicated in neonatal human pulmonary microvascular endothelial cells (HPMECs), with higher miR-30a expression in female HPMECs upon exposure to hyperoxia. miR-30a

modulates many biological processes and has proangiogenic and antifibrotic properties. A miRNA can have many putative mRNA targets. *Snail* (a profibrotic gene) is a miR-30a target. We show that *Snail* expression is decreased in female neonatal mice *in vivo*. On the other hand, male HPMECs show increased SNAIL expression *in vitro* after exposure to hyperoxia. Also, we show that *Hif-1 α* could modulate miR-30a expression in HPMECs. Interestingly, *Hif-1 α* also increases *Snail* expression, and the net protective effect in females could be mediated through a *Hif-1 α* -mediated increase in the *miR-30a* expression which keeps the *Snail* expression in check during exposure to hyperoxia (Figure 7).

Snail (Snai Family Transcriptional Repressor 1) is a zinc finger transcriptional repressor protein and plays a central role in both epithelial to mesenchymal (EMT) [30] and endothelial to mesenchymal (Endo-MT) in many disease processes. Exposure to hyperoxia leads to a profibrotic phenotype leading to an increase in myofibroblasts [31-34]. Endo-MT is a complex biological process in which endothelial cells lose their surface expression of endothelial-specific markers (e.g., CD31 and vWF) and acquire a mesenchymal phenotype (elongated and fusiform) and express mesenchymal markers (like α -SMA and vimentin) [35]. The role of Endo-MT is known in lung diseases such as idiopathic pulmonary fibrosis (significant contribution to fibrosis and vascular regression) and pulmonary arterial hypertension (by

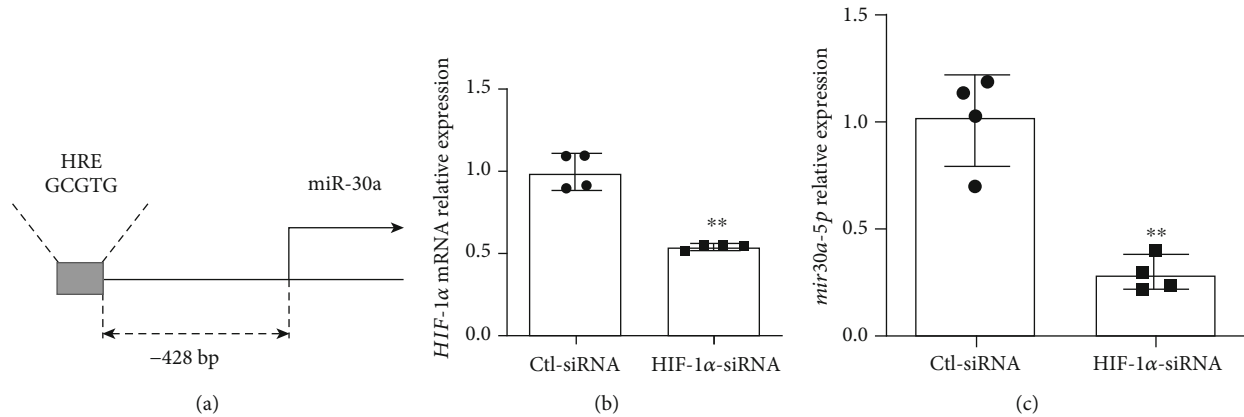


FIGURE 4: *miR-30a* regulation by HIF-1 α . (a) Motif search revealed a hypoxia response element [AG]CGTG site 428 base pairs upstream of *miR-30a*, based on the mm10 genome build. (b) *HIF-1α* mRNA expression in female HPMECs after *HIF-1α* knockdown using siRNA (Ctl-siRNA: control siRNA). (c) *mir30a-5p* expression in female HPMECs after *HIF-1α* knockdown. Values are expressed as mean \pm SD. $N = 4$ experimental replicates for each group in each test. Significant differences from baseline are indicated by ** $P < 0.01$.

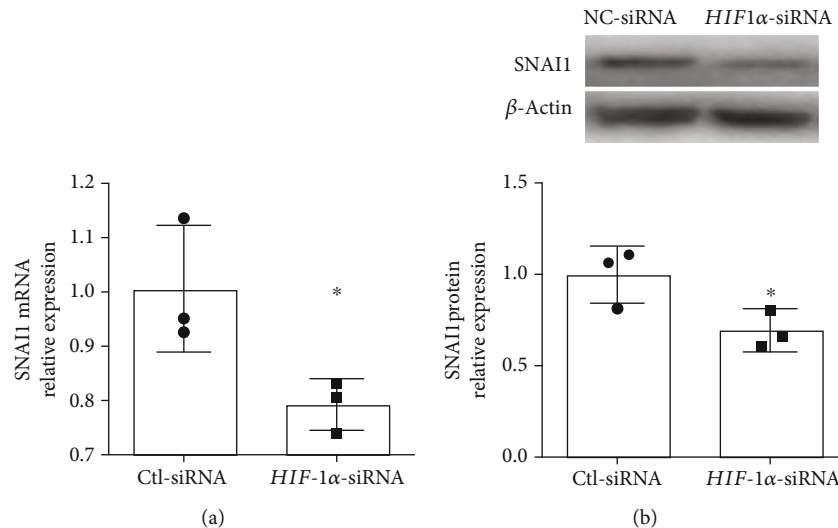


FIGURE 5: *HIF-1α* knockdown decreased *SNAIL1* mRNA and protein expression in neonatal female human pulmonary microvascular endothelial cells. *SNAIL1* mRNA (a) and protein (b) expression in HPMECs after *HIF-1α* knockdown using siRNA (Ctl-siRNA: control siRNA). Values are expressed as mean \pm SD. $N = 3$ experimental replicates for each group in each test. Significant differences from baseline are indicated by * $P < 0.05$.

causing pulmonary vascular remodeling and endothelial dysfunction) [36–38]. We have previously shown that male HPMECs show higher expression of α -SMA upon exposure to hyperoxia accompanied by a decrease in CD31 expression [39]. In this study, *SNAIL1* mRNA expression was significantly higher in male HPMECs compared to female. Male HPMECs showed increased *SNAIL1* protein upon exposure to hyperoxia, while female HPMECs did not show this change. In vivo, *Snail* expression was decreased in females immediately after (P7) hyperoxia exposure.

Since our group had discovered sex-specific differences in *miR-30a* expression with females showing higher expression than males [9], we wanted to elucidate the possible regulation of *SNAIL1* expression by *miR-30a* in female HPMECs. Other studies have shown *miR-30a* mediated downregulation of *Snail* [10, 14, 17]. Zhang et al. reported that *miR-30a*

targeted and downregulated *SNAIL1* in a diabetic cataract model *in vitro*. Similar antifibrotic effects were reported in non-small cell lung cancer [40], atrial fibrillation-induced myocardial fibrosis [14], and TGF- β 1-induced peritoneal fibrosis [10]. In female HPMECs, in the present study, we show decreased expression of *SNAIL1* with *miR-30a* overexpression and increased expression with *miR-30a* inhibition. To our knowledge, this is the first study to show that *miR-30a* targets *SNAIL1* in neonatal HPMECs.

Even though we had reported sex-specific differences in *miR-30a* expression [9], upstream modulation of *miR-30a* was not elucidated. *Hif-1α* is expressed in early lung development and slowly decreases expression as the lung matures, with a second transient peak at the alveolar stage of lung development around PND10 [20, 41]. Use of prolyl hydroxylase domain (PHD) inhibitors to increase HIF protein levels

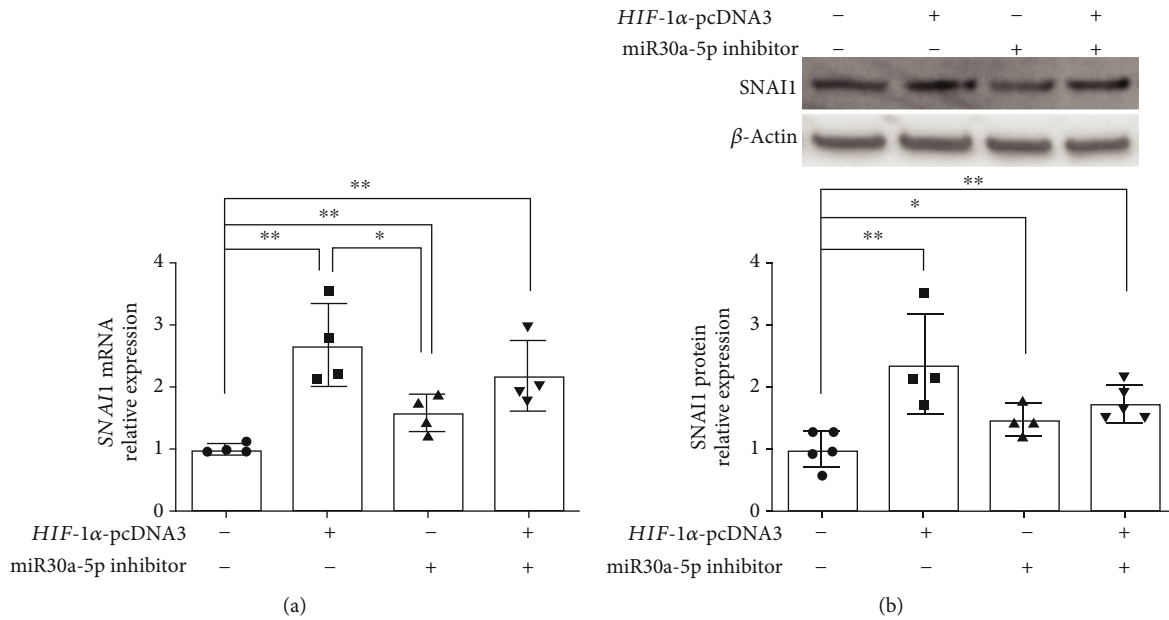


FIGURE 6: *SNAI1* expression following miR30a-5p inhibition, HIF-1 α overexpression, or both in female HPMECs: *SNAI1* mRNA (a) and protein (b) expression in female HPMECs transfected with HIF-1 α overexpression plasmid (*HIF-1 α -pcDNA3*), miR30a-5p inhibitor, or both. Values are expressed as mean \pm SD. $N = 4$ experimental replicates for each group. Significant differences between the indicated groups are shown by * $P < 0.05$ and ** $P < 0.01$.

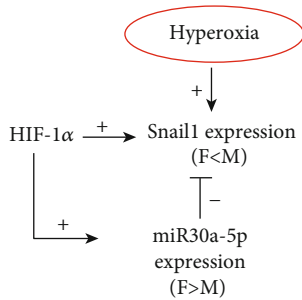
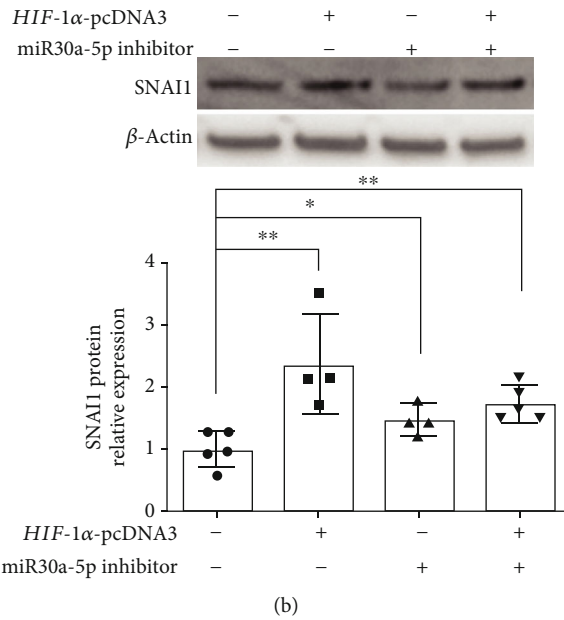


FIGURE 7: Overall schematic. Hyperoxia increases *SNAI1* expression *in vitro*, but females show decreased expression *in vivo*. miR-30a decreases *Snail* expression, and females show higher expression. HIF-1 α increases both the miR-30a and *Snail* expressions, but HIF-1 α -mediated miR-30a expression in females keeps the *Snail* expression in check upon exposure to hyperoxia.

improved lung growth in a preterm baboon model of BPD [42] Interestingly, *Hif-1 α* activity is increased, when human umbilical vein endothelial cells are returned to normoxic conditions after being exposed to hyperoxia [43]. We had previously reported that *HIF-1 α* binding to its target genes is more significant in female lungs after hyperoxia exposure [22]. In contrast to male mice, females did not show a decrease in *Hif-1 α* expression at PND21 after early hyperoxia exposure. Yang et al. showed that HIF-1 α inhibition abrogated *miR-30a* upregulation in cardiomyocytes [18]. In this study, we were able to show that the silencing of *HIF-1 α* in HPMECs significantly decreased *miR-30a* expression in HPMECs, thus postulating a role for *HIF-1 α* in increasing *miR-30a* expression in the hyperoxia model, especially during the normoxic recovery phase after exposure to hyperoxia.



Many factors including HIF-1 α , NFKB, STAT3, and NICD can induce *Snail* expression [30]. Xu et al. showed that *HIF-1 α* could induce *Snail*, independent of *TGF- β* , and that *SNAI1* is a direct target of *HIF-1 α* in human coronary endothelial cells [23]. We wanted to see if *HIF-1 α* modulated *SNAI1* expression in HPMECs. *HIF-1 α* knock-down in HPMECs decreased *SNAI1* mRNA and protein expression. Similar results were reported by Zhu et al. in pancreatic cancer cells [44]. Hypoxia-induced Endo-MT in a model of radiation-induced pulmonary fibrosis was mediated through *HIF-1 α* [45]. *HIF-1 α* may be thus inducing both profibrotic (*SNAI1*) and antifibrotic (*miR-30a*) pathways in HPMECs. The balance of expression between the two mediators may eventually predispose or protect the lung from pathologic fibrosis. In our study, *HIF-1 α* overexpression and *miR30a-5p* inhibition either separately or in combination increased *SNAI1* expression, but the combined treatment did not produce an additive effect in *SNAI1* expression.

We recognize the following limitations in this study. Our study only evaluated two time points *in vivo* after hyperoxia exposure during the saccular stage of lung development in mice. Later time points may have sustained differences in *Snail* expression, or these differences may be lost. Also, we used high oxygen concentration (0.95 FiO₂) for our hyperoxia exposure protocol based on the reproducible BPD-like phenotype produced in mice. Even though this limits hyperoxia exposure to the saccular stage of lung development, which equates to 26-36 weeks in human neonates, this oxygen concentration is much higher than what is used in the human NICU. Also, these results may be different from other models, where hyperoxia exposure is continued for longer durations [2]. We are also aware that findings in the murine

model may be strain-specific [46]. Our *in vivo* data is from whole lung tissue and does not show endothelial-specific expression. Our future studies will seek to validate the cell-specific expression of *Snail* in primary mouse endothelial cells and BPD and non-BPD patient samples. In conclusion, we show that the intersection between *HIF-1 α* , *miR-30a*, and *Snail* may play a role in driving sex-specific differences in after neonatal hyperoxic lung injury.

Data Availability

The data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest

The authors declare that there is no conflict of interest regarding the publication of this paper.

Acknowledgments

This work was supported by grants from NIH (K08-HL127103, R03-HL141572, and R01-HL144775 to KL).

Supplementary Materials

Supplementary Figure 1: *Snail* expression in vivo and in vitro: *Snail* mRNA (A–C) and protein (D–F) expression in neonatal male and female mice exposed to hyperoxia (P1-5, 0.95 FiO₂) during the saccular stage of lung development at P7 (A, D) and P21 (B, E) and in male and female neonatal human pulmonary microvascular endothelial cells (HPMECs) (C, F). Significant differences between the indicated groups is shown by **P* < 0.05. (Supplementary Materials)

References

- [1] C. Nardiello, I. Mižiková, D. M. Silva et al., “Standardisation of oxygen exposure in the development of mouse models for bronchopulmonary dysplasia,” *Disease Models & Mechanisms*, vol. 10, pp. 185–196, 2017.
- [2] J. Berger and V. Bhandari, “Animal models of bronchopulmonary dysplasia. I: the term mouse models,” *American Journal of Physiology-Lung Cellular and Molecular Physiology*, vol. 307, pp. L936–L947, 2014.
- [3] A. J. Jobe, “The new BPD: an arrest of lung development,” *Pediatric Research*, vol. 46, pp. 641–643, 1999.
- [4] C. D. Baker and C. M. Alvira, “Disrupted lung development and bronchopulmonary dysplasia: opportunities for lung repair and regeneration,” *Current Opinion in Pediatrics*, vol. 26, pp. 306–314, 2014.
- [5] C. D. Baker and S. H. Abman, “Impaired pulmonary vascular development in bronchopulmonary dysplasia,” *Neonatology*, vol. 107, pp. 344–351, 2015.
- [6] C. D. Baker, S. H. Abman, and P. M. Mourani, “Pulmonary hypertension in preterm infants with bronchopulmonary dysplasia,” *Pediatric Allergy, Immunology and Pulmonology*, vol. 27, pp. 8–16, 2014.
- [7] J. J. Coalson, “Pathology of new bronchopulmonary dysplasia,” *Seminars in Neonatology*, vol. 8, pp. 73–81, 2003.
- [8] K. Lingappan, W. Jiang, L. Wang, and B. Moorthy, “Sex-specific differences in neonatal hyperoxic lung injury,” *American Journal of Physiology-Lung Cellular and Molecular Physiology*, vol. 311, pp. L481–L493, 2016.
- [9] Y. Zhang, C. Coarfa, X. Dong et al., “MicroRNA-30a as a candidate underlying sex-specific differences in neonatal hyperoxic lung injury: implications for BPD,” *American Journal of Physiology-Lung Cellular and Molecular Physiology*, vol. 316, pp. L144–L156, 2018.
- [10] Q. Zhou, M. Yang, H. Lan, and X. Yu, “miR-30a negatively regulates TGF- β 1-induced epithelial-mesenchymal transition and peritoneal fibrosis by targeting *Snail*,” *The American Journal of Pathology*, vol. 183, pp. 808–819, 2013.
- [11] Y.-H. Chung, S. C. Li, Y. H. Kao et al., “miR-30a-5p inhibits epithelial-to-mesenchymal transition and upregulates expression of tight junction protein claudin-5 in human upper tract urothelial carcinoma cells,” *International Journal of Molecular Sciences*, vol. 18, p. 1826, 2017.
- [12] Z. Liu, K. Tu, and Q. Liu, “Effects of microRNA-30a on migration, invasion and prognosis of hepatocellular carcinoma,” *FEBS Letters*, vol. 588, pp. 3089–3097, 2014.
- [13] R. Peng, L. Zhou, Y. Zhou et al., “miR-30a inhibits the epithelial–mesenchymal transition of podocytes through downregulation of NFATc3,” *International Journal of Molecular Sciences*, vol. 16, pp. 24032–24047, 2015.
- [14] C.-T. Yuan, X. X. Li, Q. J. Cheng, Y. H. Wang, J. H. Wang, and C. L. Liu, “MiR-30a regulates the atrial fibrillation-induced myocardial fibrosis by targeting *snail 1*,” *International Journal of Clinical and Experimental Pathology*, vol. 8, pp. 15527–15536, 2015.
- [15] J. Chen, Y. Yu, S. Li et al., “MicroRNA-30a ameliorates hepatic fibrosis by inhibiting beclin1-mediated autophagy,” *Journal of Cellular and Molecular Medicine*, vol. 21, pp. 3679–3692, 2017.
- [16] C. Mao, J. Zhang, S. Lin et al., “miRNA-30a inhibits AECs-II apoptosis by blocking mitochondrial fission dependent on Drp-1,” *Journal of Cellular and Molecular Medicine*, vol. 18, pp. 2404–2416, 2014.
- [17] L. Zhang, Y. Wang, W. Li et al., “MicroRNA-30a regulation of epithelial-mesenchymal transition in diabetic cataracts through targeting *SNAIL1*,” *Scientific Reports*, vol. 7, article 1117, 2017.
- [18] Y. Yang, Y. Li, X. Chen, X. Cheng, Y. Liao, and X. Yu, “Exosomal transfer of miR-30a between cardiomyocytes regulates autophagy after hypoxia,” *Journal of Molecular Medicine*, vol. 94, pp. 711–724, 2016.
- [19] L. A. Shimoda and G. L. Semenza, “HIF and the lung: role of hypoxia-inducible factors in pulmonary development and disease,” *American Journal of Respiratory and Critical Care Medicine*, vol. 183, pp. 152–156, 2011.
- [20] A. Vadivel, R. S. Alphonse, N. Etches et al., “Hypoxia inducible factors promote alveolar development and regeneration,” *American Journal of Respiratory Cell and Molecular Biology*, vol. 50, pp. 96–105, 2014.
- [21] T. Eckle, K. Brodsky, M. Bonney et al., “HIF1A reduces acute lung injury by optimizing carbohydrate metabolism in the alveolar epithelium,” *PLoS Biology*, vol. 11, article e1001665, 2013.
- [22] C. Coarfa, Y. Zhang, S. Maity et al., “Sexual dimorphism of the pulmonary transcriptome in neonatal hyperoxic lung injury: identification of angiogenesis as a key pathway,” *American*

- Journal of Physiology-Lung Cellular and Molecular Physiology*, vol. 313, no. 6, pp. L991–L1005, 2017.
- [23] X. Xu, X. Tan, B. Tampe, E. Sanchez, M. Zeisberg, and E. M. Zeisberg, “Snail is a direct target of hypoxia-inducible factor 1 α (HIF1 α) in hypoxia-induced endothelial to mesenchymal transition of human coronary endothelial cells,” *Journal of Biological Chemistry*, vol. 290, pp. 16653–16664, 2015.
- [24] T. Kokudo, Y. Suzuki, Y. Yoshimatsu, T. Yamazaki, T. Watabe, and K. Miyazono, “Snail is required for TGF β -induced endothelial-mesenchymal transition of embryonic stem cell-derived endothelial cells,” *Journal of Cell Science*, vol. 121, pp. 3317–3324, 2008.
- [25] M. M. Mahmoud, J. Serbanovic-Canic, S. Feng et al., “Shear stress induces endothelial-to-mesenchymal transition via the transcription factor Snail,” *Scientific Reports*, vol. 7, p. 3375, 2017.
- [26] J.-X. Sun, T. F. Chang, M. H. Li et al., “SNAI1, an endothelial-mesenchymal transition transcription factor, promotes the early phase of ocular neovascularization,” *Angiogenesis*, vol. 21, pp. 635–652, 2018.
- [27] C. Sahlgren, M. V. Gustafsson, S. Jin, L. Poellinger, and U. Lendahl, “Notch signaling mediates hypoxia-induced tumor cell migration and invasion,” *Proceedings of the National Academy of Sciences*, vol. 105, pp. 6392–6397, 2008.
- [28] Y. Matsuno, A. L. Coelho, G. Jarai, J. Westwick, and C. M. Hogaboam, “Notch signaling mediates TGF- β 1-induced epithelial-mesenchymal transition through the induction of Snail,” *The International Journal of Biochemistry & Cell Biology*, vol. 44, pp. 776–789, 2012.
- [29] Y. Zhang, W. Jiang, L. Wang, and K. Lingappan, “Sex-specific differences in the modulation of growth differentiation factor 15 (GDF15) by hyperoxia in vivo and in vitro: role of Hif-1 α ,” *Toxicology and Applied Pharmacology*, vol. 332, pp. 8–14, 2017.
- [30] S. Kaufhold and B. Bonavida, “Central role of Snail1 in the regulation of EMT and resistance in cancer: a target for therapeutic intervention,” *Journal of Experimental & Clinical Cancer Research*, vol. 33, p. 62, 2014.
- [31] L. G. Boros, J. S. Torday, W.-N. Paul Lee, and V. K. Rehan, “Oxygen-induced metabolic changes and transdifferentiation in immature fetal rat lung lipofibroblasts,” *Molecular Genetics and Metabolism*, vol. 77, pp. 230–236, 2002.
- [32] J. Ni, Z. Dong, W. Han, D. Kondrikov, and Y. Su, “The role of RhoA and cytoskeleton in myofibroblast transformation in hyperoxic lung fibrosis,” *Free Radical Biology & Medicine*, vol. 61, pp. 26–39, 2013.
- [33] D. Kondrikov, R. B. Caldwell, Z. Dong, and Y. Su, “Reactive oxygen species-dependent RhoA activation mediates collagen synthesis in hyperoxic lung fibrosis,” *Free Radical Biology & Medicine*, vol. 50, pp. 1689–1698, 2011.
- [34] J. M. S. Sucre, P. Vijayaraj, C. J. Aros et al., “Posttranslational modification of β -catenin is associated with pathogenic fibroblastic changes in bronchopulmonary dysplasia,” *American Journal of Physiology-Lung Cellular and Molecular Physiology*, vol. 312, pp. L186–L195, 2017.
- [35] S. Piera-Velazquez, F. A. Mendoza, and S. A. Jimenez, “Endothelial to mesenchymal transition (EndoMT) in the pathogenesis of human fibrotic diseases,” *Journal of Clinical Medicine*, vol. 5, p. 45, 2016.
- [36] Y. Li, K. O. Lui, and B. Zhou, “Reassessing endothelial-to-mesenchymal transition in cardiovascular diseases,” *Nature Reviews Cardiology*, vol. 15, pp. 445–456, 2018.
- [37] R. B. Good, A. J. Gilbane, S. L. Trinder et al., “Endothelial to mesenchymal transition contributes to endothelial dysfunction in pulmonary arterial hypertension,” *The American Journal of Pathology*, vol. 185, pp. 1850–1858, 2015.
- [38] N. Hashimoto, S. H. Phan, K. Imaizumi et al., “Endothelial-mesenchymal transition in bleomycin-induced pulmonary fibrosis,” *American Journal of Respiratory Cell and Molecular Biology*, vol. 43, pp. 161–172, 2010.
- [39] Y. Zhang, X. Dong, J. Shirazi, J. P. Gleghorn, and K. Lingappan, “Pulmonary endothelial cells exhibit sexual dimorphism in their response to hyperoxia,” *American Journal of Physiology-Heart and Circulatory Physiology*, vol. 315, pp. H1287–H1292, 2018.
- [40] R. Kumarswamy, G. Mudduluru, P. Ceppi et al., “MicroRNA-30a inhibits epithelial-to-mesenchymal transition by targeting Snail and is downregulated in non-small cell lung cancer,” *International Journal of Cancer*, vol. 130, pp. 2044–2053, 2012.
- [41] J. P. Bridges, S. Lin, M. Ikegami, and J. M. Shannon, “Conditional hypoxia inducible factor-1 α induction in embryonic pulmonary epithelium impairs maturation and augments lymphangiogenesis,” *Developmental Biology*, vol. 362, pp. 24–41, 2012.
- [42] T. M. Asikainen, L. Y. Chang, J. J. Coalson et al., “Improved lung growth and function through hypoxia-inducible factor in primate chronic lung disease of prematurity,” *The FASEB Journal*, vol. 20, pp. 1698–1700, 2006.
- [43] F. Cimino, C. Balestra, P. Germonpre et al., “Pulsed high oxygen induces a hypoxic-like response in human umbilical endothelial cells and in humans,” *Journal of Applied Physiology*, vol. 113, pp. 1684–1689, 2012.
- [44] G.-H. Zhu, C. Huang, Z.-Z. Feng, X.-H. Lv, and Z.-J. Qiu, “Hypoxia-induced snail expression through transcriptional regulation by HIF-1 α in pancreatic cancer cells,” *Digestive Diseases and Sciences*, vol. 58, pp. 3503–3515, 2013.
- [45] S.-H. Choi, Z.-Y. Hong, J.-K. Nam et al., “A hypoxia-induced vascular endothelial-to-mesenchymal transition in development of radiation-induced pulmonary fibrosis,” *Clinical Cancer Research*, vol. 21, pp. 3716–3726, 2015.
- [46] S. Leary, P. Das, D. Ponnalagu, H. Singh, and V. Bhandari, “Genetic strain and sex differences in a hyperoxia-induced mouse model of varying severity of bronchopulmonary dysplasia,” *The American Journal of Pathology*, vol. 189, pp. 999–1014, 2019.