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Sheng Xia Children's Mercy Hospital

Heather Menden Children's Mercy Hospital

Nicholas Townley MD Children's Mercy Hospital

Sherry M. Mabry Children's Mercy Hospital

Michael F. Nyp Children's Mercy Hospital

See next page for additional authors

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Authors

Sheng Xia, Heather Menden, Nicholas Townley MD, Sherry M. Mabry, Michael F. Nyp, Donald W. Thibeault, and Venkatesh Sampath

DELTA LIKE 4 (DLL4), AN ENDOTHELIAL SPECIFIC NOTCH LIGAND IS CRITICAL FOR LUNG VASCULAR ARBORIZATION AND ALVEOLARIZATION



Children's Mercy Kansas City, Department of Pediatrics, Division of Neonatology, Kansas City, MO

Introduction

- Human and mouse lung development progress temporally from the embryonic, pseudo-glandular, canalicular, saccular and alveolar phase.
- Vascular development parallels lung development, resulting in an arborized network with terminal capillaries in close proximity to alveoli.
- The relative contribution of vasculogenesis vs. angiogenesis to distal lung development remains unclear. Further, the contribution of endothelial cell (EC) angiogenesis to lung development is unknown.
- Herein, we investigated the role of Delta like 4 (DLL4), a regulator of embryonic angiogenesis in vascular arborization and alveolarization.

Hypothesis and Objectives

Hypothesis: DLL4-dependent angiogenesis is essential for distal vascular arborization and normal alveolarization.

Objectives: 1) To map DLL4 during mouse and human lung development using wild-type and *Dll4-LacZ* mice. 2) To investigate the effect of DLL4 haploinsufficiency and EC-specific Dll4 deletion on lung vascular development and alveolarization.

Methods

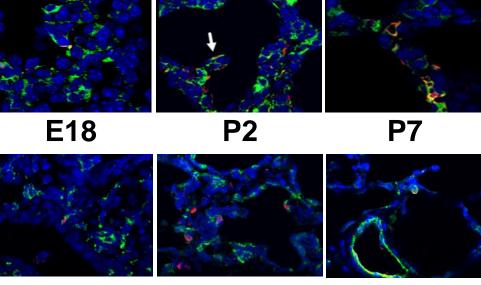
- 1. DLL4 mapping in mouse & human: DLL4 & PECAM IF/IHC in WT, $DLL4^{+/lacZ}$ mice, human autopsy lung samples, X-gal in $DLL4^{+/lacZ}$ mice.
- 2. Mouse gene expression: Whole lung RNA expression by qRT-PCR.
- 3. Mouse lung morphometry: Radial alveolar count (RAC) quantification was done to assess alveolarization in WT and DLL4+/lacZ mice.
- **4.** Tamoxifen-inducible DII4 deletion: DII4 +/Loxp (F) X DII4 +/Loxp; VE-cad-CreERT2 (M). Female injected with Tamoxifen on P2, P3, P4.

Results

DLL4 expression from Canalicular to Alveolar Phase





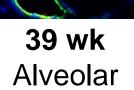


DLL4/ PECAM/DAPI (nucleus)

Arrow inset represents DLL4 expression in alveolar septum in PECAM+ cells. 63x; n=3.

22 wk Canalicular

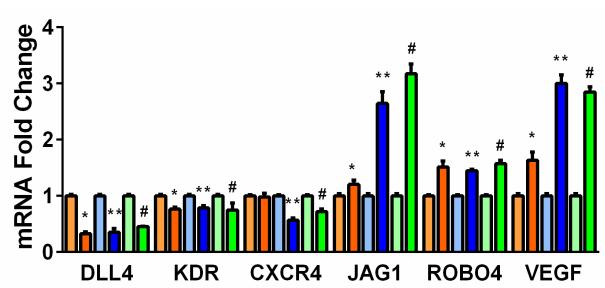
27 wk Saccular



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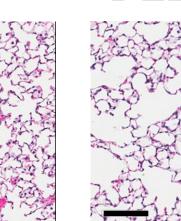
Results

DLL4 Lung Mapping Using DII4+/lacZ Reporter Mice



DLL4 Haploinsufficiency Causes Defective Alveolarization in Mice

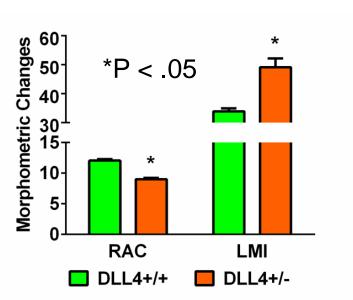




Inflation-fixed P28 mouse lungs Scale bar = $100 \,\mu m$

DLL4+/-

DLL4/ PECAM/ Fast Red (nucleus)



A. X-gal (blue) indicates

capillary EC.

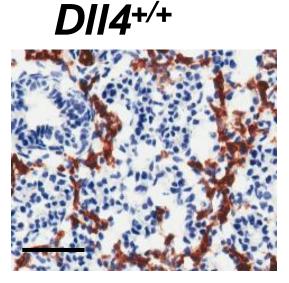
DLL4 in distal lung

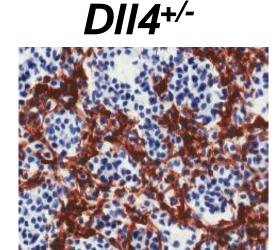
B. Arrow - DLL4 in alveolar

(lung EC). 40x; n=3.

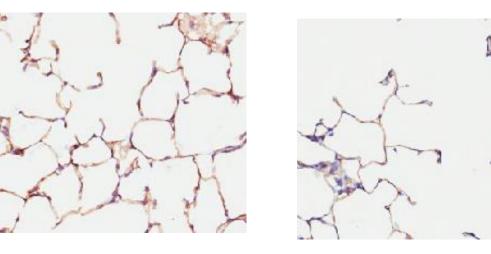
septum in PECAM+ cells

Aberrant Pulmonary Vascular Development in DII4+/- Mice

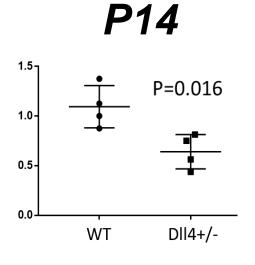




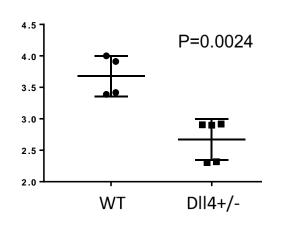
E17.5 Canalicular Stage





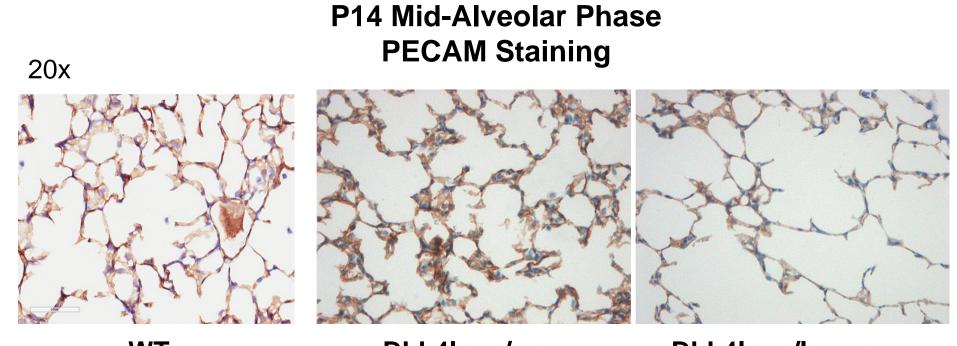


Blood vessels >10µM/40X field



Capillaries <10µM/40X field

DII4 Deletion in EC Impairs Lung alveolarization



WT

Results

DLL4 Haploinsufficiency Causes Defective Alveolarization in Mice

DOL28 WT

INC School of Medicine

🗖 DOL14 Het DOL28 Het

*p<0.01 (DOL4 WT vs. DOL4 Het) **p<0.01 (DOL14 WT vs. DOL14 Het) [#]p<0.01 (DOL28 WT vs. DOL28 Het)

DII4 +/Loxp (F) crossed with DII4 +/Loxp; VE-cad-CreERT2 (M) Dam treated with tamoxifen - P4 and P5

DLL4loxp/loxp;cre

Conclusion

DLL4 is expressed from the canalicular to the alveolar phase of lung development in mice & humans, indicating that DLL4dependent angiogenesis contributes to vascular development. DLL4+/- mice have decreased expression of tip cell EC fate markers, with increased expression of stalk cell EC markers and VEGF. DLL4+/- mice have dysmorphic microvasculature, impaired angiogenesis and disrupted alveolar development. Pilot data suggest that EC-DII4 deletion impairs alveolarization. **Future:** Investigate how EC-DLL4 directs alveolarization, and DII4's role in defective Bronchopulmonary Dysplasia vasculature.

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DLL4loxp/+;cre