

RESEARCH ARTICLE

Epithelial inactivation of *Yy1* abrogates lung branching morphogenesis

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ABSTRACT

Yin Yang 1 (YY1) is a multifunctional zinc-finger-containing transcription factor that plays crucial roles in numerous biological processes by selectively activating or repressing transcription, depending upon promoter contextual differences and specific protein interactions. In mice, *Yy1* null mutants die early in gestation whereas *Yy1* hypomorphs die at birth from lung defects. We studied how the epithelial-specific inactivation of *Yy1* impacts on lung development. The *Yy1* mutation in lung epithelium resulted in neonatal death due to respiratory failure. It impaired tracheal cartilage formation, altered cell differentiation, abrogated lung branching and caused airway dilation similar to that seen in human congenital cystic lung diseases. The cystic lung phenotype in *Yy1* mutants can be partly explained by the reduced expression of *Shh*, a transcriptional target of YY1, in lung endoderm, and the subsequent derepression of mesenchymal *Fgf10* expression. Accordingly, SHH supplementation partially rescued the lung phenotype *in vitro*. Analysis of human lung tissues revealed decreased YY1 expression in children with pleuropulmonary blastoma (PPB), a rare pediatric lung tumor arising during fetal development and associated with *DICER1* mutations. No evidence for a potential genetic interplay between murine *Dicer* and *Yy1* genes during lung morphogenesis was observed. However, the cystic lung phenotype resulting from the epithelial inactivation of *Dicer* function mimics the *Yy1* lung malformations with similar changes in *Shh* and *Fgf10* expression. Together, our data demonstrate the crucial requirement for YY1 in lung morphogenesis and identify *Yy1* mutant mice as a potential model for studying the genetic basis of PPB.

KEY WORDS: YY1, SHH, FGF10, DICER1, Lung branching, Pleuropulmonary blastoma

INTRODUCTION

YY1 is a ubiquitous zinc finger transcription factor that contains diverse domains, enabling a plethora of protein-protein interactions. YY1 can recruit co-activators or co-repressors, which determine whether YY1 will execute inhibitory or activating functions on targets (Deng et al., 2010). In regulating a multitude of genes, YY1 plays crucial functions in numerous biological processes, including cell proliferation and differentiation, X-chromosome inactivation,

and embryogenesis (Donohoe et al., 2007; Nicholson et al., 2011). The *Yy1* null mutation in mice results in peri-implantation lethality, a phenotype precluding investigation of YY1 requirement at later developmental stages (Donohoe et al., 1999). Using mouse lines carrying a *Yy1* conditional allele and tissue-specific *Cre*-expressing transgenes, *Yy1* was found to widely participate in developmental processes. However, despite the fact that mice expressing 25% of normal YY1 levels die at birth from respiratory failure due to collapsed lungs, little is known about the role of *Yy1* in lung morphogenesis (Affar et al., 2006).

Lung development is subdivided into five overlapping periods (Morrissey and Hogan, 2010). The embryonic phase is characterized by the formation of the ventral diverticulum that arises from the laryngotracheal groove of the foregut endoderm. Subsequently, this diverticulum, destined to become the trachea, divides into the left and right lung buds. At the pseudoglandular stage, the two primary buds elongate and undergo stereotypical branching via complex epithelium-mesenchyme interactions to form the respiratory tree. This is followed by the canalicular and saccular stages, during which the progressive differentiation of the pulmonary epithelium, the expansion of the vasculature with thinning of the mesenchyme, and the formation of functional air-blood barriers happen. Alveologenesis occurs after birth and is characterized by the formation of alveoli, the final gas-exchange units. Recent data indicate that the transition from branching morphogenesis to epithelial cell differentiation depends on control mechanisms involving key players of lung development such as *Fgf10* and *SOX9*. These latter act by regulating the delicate balance between distal and proximal endodermal progenitors, promoting branching and preventing precocious alveolar differentiation (Chang et al., 2013; Volckaert et al., 2013; Yang and Chen, 2014).

We showed that the specific ablation of *Yy1* function in lung mesenchyme causes neonatal death of mutant pups due to collapsed lungs, a phenotype similar to that of *Yy1* hypomorph mutants (Bérubé-Simard et al., 2014). In the present study, we investigated the impact of the epithelial-specific inactivation of *Yy1* in the developing lung. The loss of *Yy1* function in lung epithelium also resulted in death at birth. It affected tracheal cartilage formation, cell differentiation and lung branching, leading to the formation of large cysts. The reduced expression of *Shh* and the subsequent upregulation of *Fgf10* expression likely contributed to the branching defects in *Yy1* mutants. Accordingly, *in vitro* SHH supplementation partially rescued the lung phenotype of *Yy1* mutant explants. The cysts in *Yy1* mutants mimicked the lung phenotype of *Dicer* mutants with similar *Shh* and *Fgf10* expression variations (Harris et al., 2006). DICER is a ribonuclease III essential for the biogenesis of mature microRNAs (Bartel, 2004). Cystic lesions characterize several pulmonary diseases such as congenital cystic adenomatoid malformation (CCAM) and pleuropulmonary blastoma (PPB), the latter being a rare dysontogenetic neoplasm

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of childhood evolving from a cystic to a solid state over time. Mutations in the human *DICER1* gene are found in PPB patients (Hill et al., 2008; Messinger et al., 2015). Analysis of lung tissue from CCAM patients revealed no major changes in YY1 expression. Conversely, decreased YY1 expression was detected in PPB children. In summary, our results demonstrate the essential requirement for YY1 in lung morphogenesis and identify YY1 as a potential factor involved in the molecular pathogenesis of PPB.

RESULTS

Loss of epithelial *Yy1* function causes defective lung morphogenesis

To circumvent the early embryonic lethality of the *Yy1* null mutants and address *Yy1* function in lung morphogenesis, we specifically deleted the *Yy1* gene in lung epithelium using the *Shh*^{+Cre} deleter mouse line (supplementary material Fig. S1; Harfe et al., 2004). All *Yy1*^{flox/flox};*Shh*^{+Cre} newborns died at birth from respiratory failure, whereas *Yy1*^{+flox};*Shh*^{+Cre} mice were viable and normal (Table 1). Analysis of lungs from embryonic (E) day 18.5 *Yy1*^{flox/flox};*Shh*^{+Cre} embryos revealed a disorganized architecture with the presence of dilated fluid-filled sacs (Fig. 1). At E12.5, lungs from *Yy1*^{flox/flox};*Shh*^{+Cre} embryos presented two hypoplastic lobes in contrast to the expected asymmetric pattern of four right lobes and one left lobe. Moreover, lung epithelium of mutant specimens had an abnormal stratified structure.

To identify the causes of the lung phenotype, we looked at cell proliferation. At E12.5, reduced immunostaining for cyclin D1, a marker for the G1/S transition, for BrdU, which labels cells in S-phase, and for pHH3, a marker for cells in late G2 and mitosis, was specifically observed in lung epithelium of *Yy1*^{flox/flox};*Shh*^{+Cre} mutants (Fig. 2A,B; supplementary material Fig. S2). Apoptosis, as indicated by cleaved caspase-3 immunostaining, was also more important in the multilayered lung epithelium of *Yy1*^{flox/flox};*Shh*^{+Cre} specimens (Fig. 2C,D). At E14.5, apoptosis was increased in lung mesenchyme from mutants (supplementary material Fig. S2). Together, reduced epithelial proliferation and augmented apoptosis contribute to the abnormal lung structure in *Yy1*^{flox/flox};*Shh*^{+Cre} mutants.

Altered lung patterning and cell differentiation in *Yy1*^{flox/flox};*Shh*^{+Cre} mutants

To investigate whether the proximal-distal patterning of airway epithelium was disrupted in *Yy1*^{flox/flox};*Shh*^{+Cre} mutants, we examined the expression of the transcription factors sex-determining region Y (SRY)-box 2 and 9 (SOX2 and SOX9), two lineage commitment markers of proximal and distal endodermal lung progenitor cells, respectively (Tian et al., 2011). In controls, SOX2 was expressed in the epithelium of trachea and proximal airways, whereas SOX9 was present in the epithelium of distal tubular tips and in the mesenchyme lining proximal airways. In

mutants, SOX2 and SOX9 epithelial expression also displayed a reciprocal pattern, but SOX2 expression was downregulated and restricted to a more proximal region (Fig. 2E-H). We further characterized epithelial cell specification by examining the expression of NKX2-1 and FOXA2, two transcription factors essential for lung branching and epithelial cell differentiation (Kimura et al., 1996; Minoo et al., 1999; Wan et al., 2004). No major difference was observed (Fig. 2I-L).

In agreement with the lack of SOX2 expression along the cysts, no secretory club (Clara) cells or ciliated cells, normally found along the proximal airway epithelium, were detected in the cystic epithelium (Fig. 2M-P). In contrast, Types I and II pneumocytes were present along the cyst-lining epithelium (Fig. 2Q-X). Microvascular development, as detected by PECAM immunostaining, also occurred within the mesenchyme surrounding the cysts (Fig. 2Y,Z).

In E12.5 controls, expression of alpha smooth muscle actin (α SMA), a marker of airway smooth muscle differentiation, was detected in myofibroblasts surrounding blood vessels and conducting airways, but excluded from the tip of growing buds. In mutants, α SMA-positive cells were lacking around cysts, but still present around pulmonary vasculature (Fig. 2AA, BB). Thus, *Yy1* epithelial ablation prevents branching morphogenesis with no major consequences on the specification of the distal epithelium. However, it interferes with the differentiation or the survival of airway myofibroblasts.

Abnormal formation of cartilage rings and impaired cell differentiation in trachea from *Yy1*^{flox/flox};*Shh*^{+Cre} mutants

The trachea from *Yy1*^{flox/flox};*Shh*^{+Cre} mutants appeared thinner with disorganized cartilage rings (Fig. 1A-B). Alcian Blue staining verified the abnormal banding pattern (Fig. 3A,B). Tracheal stenosis was confirmed by the measurement of the external diameter (625 \pm 26 μ m versus 339 \pm 25 μ m; $P < 0.001$) and the luminal surface area (61 \pm 16 μ m² versus 4.6 \pm 2 μ m²; $P < 0.01$), both of which were significantly smaller in mutants. The trachea was also longer in mutants (2029 \pm 130 μ m versus 2487 \pm 47 μ m; $P < 0.01$) (Fig. 3A-E). We analyzed the expression of SOX9, a master regulator of chondrogenesis essential for tracheal cartilage patterning (Park et al., 2010). In E14.5 controls, the punctuated SOX9 expression pattern reflected future cartilage rings. Conversely, mutants displayed continuous SOX9 expression along the upper airways (Fig. 3F,G). Thus, epithelial YY1 controls the condensation of SOX9-positive mesenchymal cells into precartilage nodules.

We assessed the expression of p63, a marker of basal cells, which are known to generate ciliated, club and goblet cells (Rock et al., 2009). Basal cells were distributed irregularly along the proximal airways of *Yy1*^{flox/flox};*Shh*^{+Cre} embryos (Fig. 3H,I). Although the number of p63-positive basal cells was not noticeably altered, ciliated, club and goblet cells were scarce in mutants (Fig. 3H-M). SOX2 is crucial for tracheal cartilage

Table 1. Ratios of genotypes in litters from crosses between *Yy1*^{flox/+};*Shh*^{+Cre} and *Yy1*^{flox/flox} mice

Age	# litters	# pups	<i>Shh</i> ^{+/+}		<i>Shh</i> ^{+Cre}	
			<i>Yy1</i> ^{flox/+}	<i>Yy1</i> ^{flox/flox}	<i>Yy1</i> ^{flox/+}	<i>Yy1</i> ^{flox/flox}
Expected (%)			(25)	(25)	(25)	(25)
E12.5	21	144	31 (21.5)	41 (28.5)	37 (25.7)	35 (24.3)
E14.5	7	54	19 (35.2)	6 (11.1)	15 (27.8)	14 (25.9)
E18.5	9	59	13 (22.1)	16 (27.1)	12 (20.3)	18 (30.5)
P21	13	74	26 (35.1)	21 (28.4)	27 (36.5)	0 (0)

The percentage obtained is indicated in parentheses.

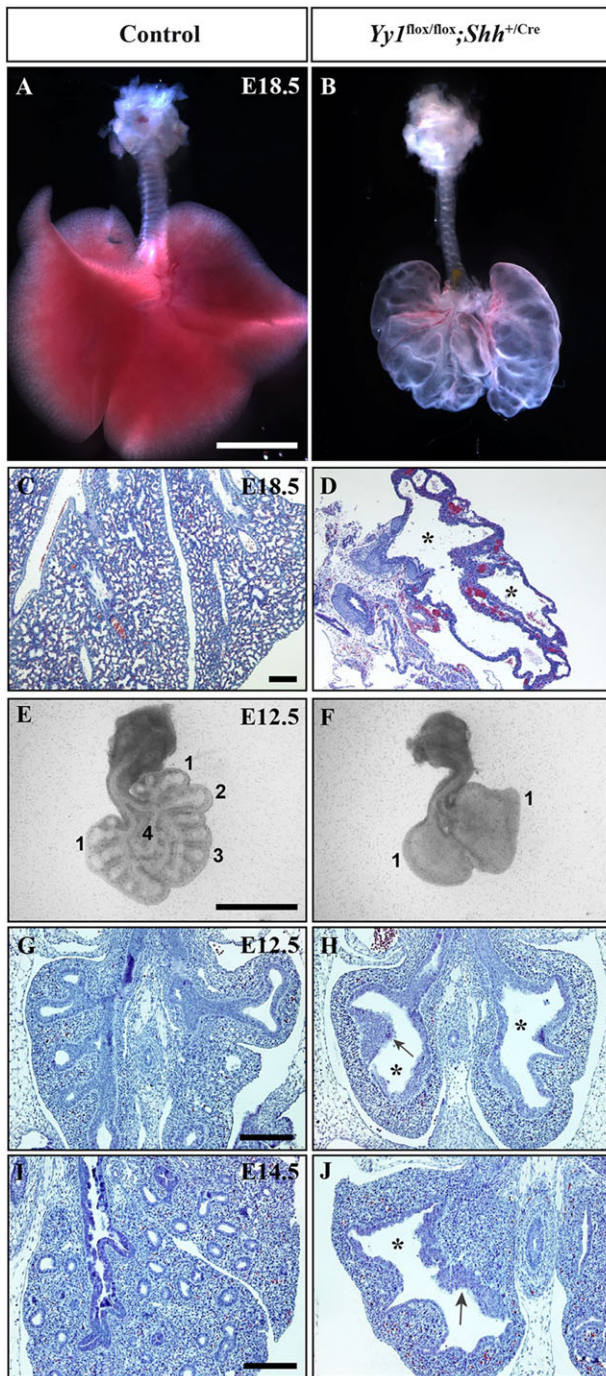


Fig. 1. *Yy1* is crucial for lung branching morphogenesis. (A,B) Lungs from E18.5 *Yy1^{flox/flox};Shh^{+/-Cre}* embryos (B) showed dilated sacs instead of the well-formed lobes seen in controls (A). (C,D,G-J) H&E-stained lung sections revealed that as early as E12.5, *Yy1^{flox/flox};Shh^{+/-Cre}* mutants displayed defective airway branching causing cyst formation (asterisks) with an abnormal stratified epithelium (arrows). (E,F) Whole-mount lungs from E12.5 control and *Yy1^{flox/flox};Shh^{+/-Cre}* lungs presented left isomerism. Labels 1-4 represent lobes. Scale bars: 2 mm in A,B; 200 μ m in C,D,G-J; 1 mm in E,F.

patterning and epithelial cell differentiation (Que et al., 2009). In controls, SOX2 was expressed in all airway epithelial cells. In mutants, SOX2 signal was unevenly spread along the tracheal epithelium, a result reminiscent of the abnormal SOX2 expression in lungs from E12.5 *Yy1^{flox/flox};Shh^{+/-Cre}* embryos (Figs 2E,F and 3N,O). These data underscore the non-cell-autonomous role of

YY1 in tracheal cartilage patterning and its cell-autonomous function in airway epithelial cell differentiation.

***Yy1* epithelial inactivation affects the expression of key players of lung development**

We assessed by qRT-PCR if the *Yy1* mutation caused the misregulation of molecules with established roles in lung morphogenesis. During branching, FGF10 is dynamically expressed in mesenchymal clustered cells, inducing the activation of the ERK/MAPK pathway in the adjacent epithelium to control the directional outgrowth of lung buds (Bellusci et al., 1997). At E14.5, *Fgf10* expression was 2.7-fold higher in *Yy1^{flox/flox};Shh^{+/-Cre}* lungs compared with *Yy1^{flox/+};Shh^{+/-Cre}* controls. Consequently, expression of *Bmp4*, *Spry2* and *Etv4* genes, known targets of lung FGF10 signaling, was upregulated in mutants (Fig. 4A; Weaver et al., 2000; Mailloux et al., 2001; Liu et al., 2003). Reduced expression of *Etv5*, another target of FGF10 signaling, was also observed. *In situ* hybridization assays revealed an expanded spatial distribution of *Fgf10* mRNA in lungs from E12.5 mutants (Fig. 4B,C). Accordingly, increased expression of phospho-ERK (pERK), the activated form of ERK, was observed in the cystic epithelium of *Yy1^{flox/flox};Shh^{+/-Cre}* embryos (Fig. 4D,E). Expression of *Fgf9*, *Hoxa5*, *Hoxb5*, *Foxp1*, *Foxp2*, *Foxp4*, *Hdac1*, and *Hdac2* genes, all involved in lung branching, was also monitored (Aubin et al., 1997; White et al., 2006; Shu et al., 2007; Wang et al., 2011, 2013). With the exception of a reduction in expression of *Fgf9* and *Foxp2*, no change in gene expression was detected (Fig. 4A).

YY1 positively regulates *Shh* expression

Shh null mutants present rudimentary lung sacs and tracheal-bronchial cartilage defects that mirror the *Yy1^{flox/flox};Shh^{+/-Cre}* phenotype. Moreover, increased *Fgf10* and *Bmp4* lung expression levels are common denominators in *Yy1* and *Shh* mutants. Finally, α SMA-positive cells are absent around the bronchi in *Shh^{-/-}* mutants (Litingtung et al., 1998; Picicelli et al., 1998; van Tuyl et al., 2007). We examined *Shh* expression in *Yy1* mutants. The *Shh^{Cre}* allele used to generate the *Yy1* epithelial deletion is a Cre knock-in producing a *Shh* null allele (Harfe et al., 2004). To eliminate any bias that might result from *Shh* haploinsufficiency, *Shh* expression levels were measured by qRT-PCR in lungs from E14.5 *Yy1^{flox/flox};Shh^{+/-Cre}* mutants and compared to *Yy1^{+/-flox};Shh^{+/-Cre}* specimens. *Shh* expression, as well as that of *Ptc1*, *Hip1* and *Foxf1*, targets of the SHH pathway, was significantly decreased in *Yy1^{flox/flox};Shh^{+/-Cre}* lungs (Fig. 4F; Picicelli et al., 1998; Chuang and McMahon, 1999; Mahlapuu et al., 2001).

Shh expression is confined to lung epithelium, suggesting that *Shh* may be a direct target of YY1. Sequence comparison with TFSEARCH and TESS databases revealed putative YY1 binding sites clustered into five domains spread along the 4.5 kb upstream sequences relative to the transcription start site (TSS) of the *Shh* murine gene. To establish whether these sites were effective *in vivo*, we performed ChIP assay on cross-linked chromatin from lungs of E14.5 wild-type embryos. We observed high occupancy by YY1 of *Shh* upstream sequences and the *Tra2b* (also known as *Sfrs10*) positive control, and no binding with the *Rcor3* and *Shh* locus negative controls (Fig. 4G; Kang et al., 2009). Thus, YY1 can physically interact with *Shh* regulatory sequences *in vivo* during lung development. Using a transfection reporter assay in HEK293 cells, we demonstrated that YY1 activated transcription of the luciferase reporter gene placed under the control of 5 kb upstream sequences of the *Shh* murine gene (supplementary

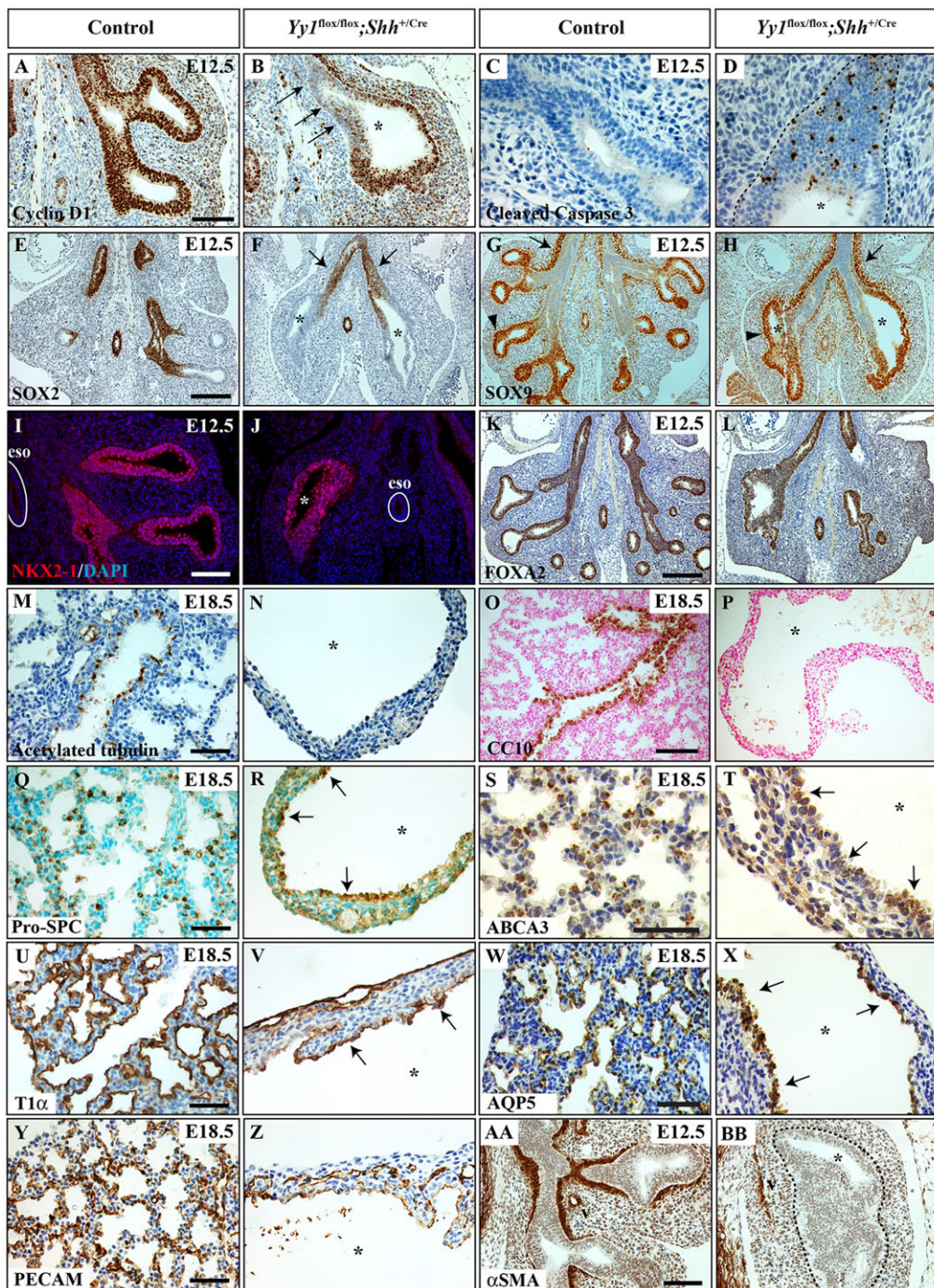


Fig. 2. YY1 controls lung epithelial proliferation and apoptosis, patterning and airway myofibroblast differentiation.

(A,B) Reduced cyclin D1 epithelial immunostaining (arrows) revealed decreased cell proliferation in lung epithelium of E12.5 $Yy1^{flox/flox}; Shh^{+/Cre}$ embryos. (C,D) Cleaved caspase-3 immunostaining showed massive apoptosis in lung epithelium of mutants. (E-H) SOX2 (E,F) and SOX9 (G,H) immunostaining assays revealed that SOX2 and SOX9 displayed reciprocal epithelial expression patterns in the developing airways from E12.5 control and $Yy1^{flox/flox}; Shh^{+/Cre}$ embryos. In mutants, SOX2 expression was downregulated (arrows) and restricted to a more anterior territory. Arrowheads indicate SOX9 expression. (I-L) Lungs from E12.5 controls and $Yy1^{flox/flox}; Shh^{+/Cre}$ mutants demonstrated comparable NKX2-1 (I,J) and FOXA2 (K,L) expression in airway epithelium. (M-P) Neither ciliated nor club cells, as detected by IHC with acetylated tubulin (M,N) and CC10 (O,P) specific markers, respectively, were observed in cyst epithelium of E18.5 $Yy1^{flox/flox}; Shh^{+/Cre}$ mutants. (Q-X) Cysts were lined by Type II and Type I pneumocytes as revealed by proSP-C (Q,R), ABCA3 (S,T), and T1 α (U,V) and AQP5 (W,X) specific markers, respectively (arrows). (Y,Z) A microvascular network detected by PECAM IHC was present in the parenchyma forming the cyst walls. (AA, BB) At E12.5, no α SMA expression was detected around the bronchi in $Yy1^{flox/flox}; Shh^{+/Cre}$ specimens. Asterisks indicate cysts. eso, esophagus; v, vasculature. Scale bars: 50 μ m in C,D,M,N,Q-Z; 100 μ m in A,B,I,J,O,P,AA, BB; 200 μ m in E-H, K,L.

material Fig. S3). Taken together, these data support the notion that YY1 is a transcriptional activator of *Shh* expression in lung epithelium.

$Yy1^{flox/flox}; Tg^{Nkx2-1Cre}$ and $Yy1^{flox/flox}; Shh^{+/Cre}$ mice share lung phenotype characteristics

To inactivate *Yy1* in the developing lung endoderm in a *Shh* haploinsufficiency-independent context, we used the BAC-*Nkx2-1-Cre* transgenic deleter line (Xu et al., 2008). $Yy1^{flox/flox}; Tg^{+/Nkx2-1Cre}$ embryos exhibited cysts in the proximal region of the lobes, a phenotype consistent with the gradient of activity of the recombinase (Fig. 5A-I). As observed in $Yy1^{flox/flox}; Shh^{+/Cre}$ mutants, Types I and II pneumocytes, but not club and ciliated cells, were present along the

cystic epithelium (supplementary material Fig. S4). Decreased *Shh* and *Hip1* and increased *Fgf10* expression levels were detected in lungs from $Yy1^{flox/flox}; Tg^{+/Nkx2-1Cre}$ embryos, providing additional evidence that epithelial YY1 regulates *Shh* expression (Fig. 5J).

SHH partially rescues the lung phenotype of $Yy1^{flox/flox}; Shh^{+/Cre}$ mutants

We tested whether addition of recombinant mouse SHH (rmSHH) could rescue the *Yy1* lung phenotype *in vitro*. Lungs from E12.5 $Yy1^{flox/flox}; Shh^{+/Cre}$ embryos cultured in control media failed with branch. They showed a loss of α SMA immunoreactivity around the cysts and an increased number of apoptotic cells when compared with controls (Fig. 6). Addition

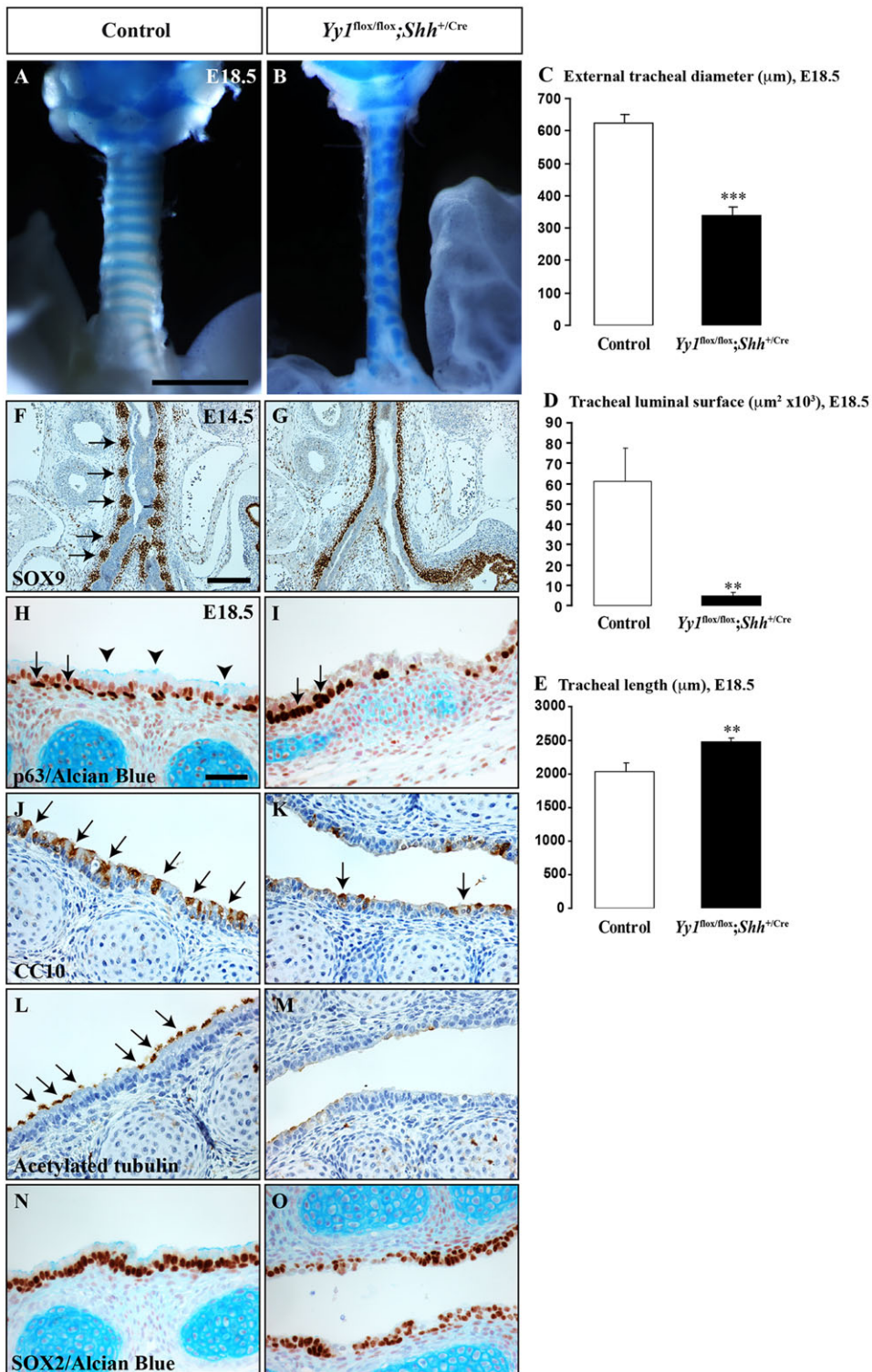


Fig. 3. Abnormal cartilage ring patterning and epithelial cell differentiation in the trachea of *Yy1^{flox/flox};Shh^{+/Cre}* embryos.

(A,B) As shown by Alcian Blue staining, the C-shaped cartilaginous rings, segmentally distributed along the trachea in controls, were replaced by irregular patches of cartilage in E18.5 *Yy1^{flox/flox};Shh^{+/Cre}* embryos. (C-E) Reduced external tracheal diameter and luminal surface, and increased tracheal length were observed in E18.5 *Yy1^{flox/flox};Shh^{+/Cre}* mutants. Values are expressed as mean \pm s.d.; ** P <0.01, *** P <0.001; n =3-4/genotype. (F,G) Unsegmented SOX9 expression was observed along the upper airways of E14.5 *Yy1^{flox/flox};Shh^{+/Cre}* embryos when compared with controls. Arrows indicate punctuated SOX9 expression. (H-M) p63 (H,I), CC10 (J,K) and acetylated tubulin (L,M) immunostaining assays detected basal, club and ciliated cells, respectively (arrows). Mucus-producing goblet cells were detected with Alcian Blue (arrowheads, H-I). In mutants, basal cells were irregularly distributed along the proximal airways and a near-complete loss of club, goblet and ciliated cells was observed. (N,O) SOX2-positive cells were less abundant and more dispersed along the tracheal epithelium of mutants. Scale bars: 1 mm in A,B; 200 μm in F,G; 50 μm in H-O.

of rmSHH to *Yy1^{flox/+};Shh^{+/Cre}* explants caused an increased overall size, probably due to the thickened mesenchymal layer (Fig. 6C,D). The rmSHH treatment of *Yy1^{flox/flox};Shh^{+/Cre}* explants did not rescue the branching defect (Fig. 6G,H). However, apoptosis in rmSHH-supplemented *Yy1^{flox/flox};Shh^{+/Cre}* lungs returned to control levels (Fig. 6M-P). Moreover, α SMA expression at the periphery of cysts was partially recovered, corroborating the importance of SHH in airway smooth muscle cell specification (Fig. 6I-L).

Decreased YY1 expression in PPB lung specimens

The presence of lung cysts in *Yy1^{flox/flox};Shh^{+/Cre}* and *Yy1^{flox/flox};Tg^{+/Nkx2-1Cre}* mutants mimics features of CCAM and PPB, two human pediatric cystic lung diseases (Stocker, 2009). To define the clinical relevance of our findings, YY1 immunostaining was performed on lung sections from children suffering from types I and II CCAM, and types I, II and III PPB and compared with lung specimens from age-matched patients who died from a non-pulmonary cause (supplementary material Table S1). PPB evolves

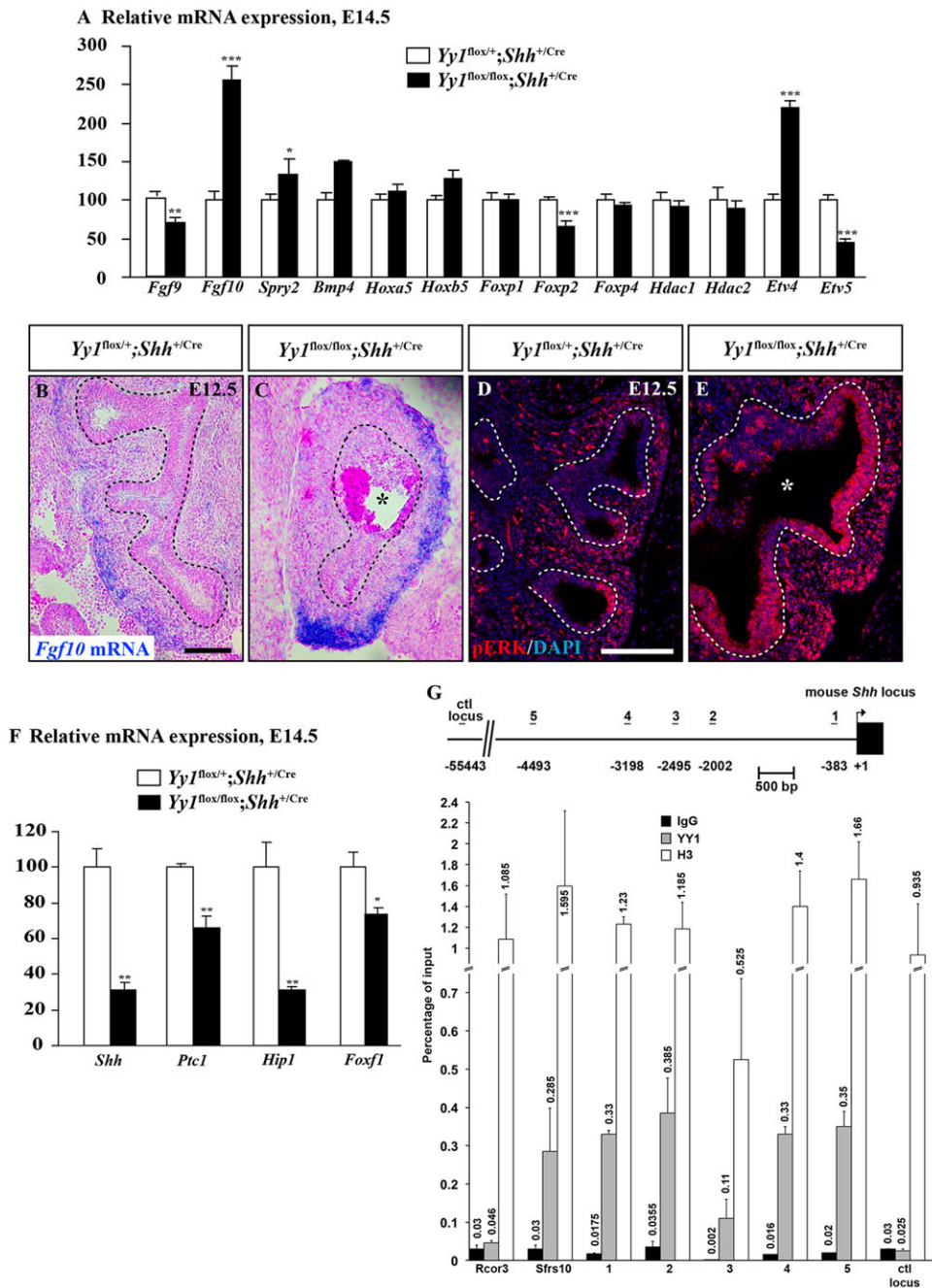


Fig. 4. Epithelial *Yy1* inactivation affects *Fgf10* and *Shh* expression. (A) qRT-PCR expression analysis revealed increased levels of *Fgf10*, *Spry2* and *Etv4* and decreased levels of *Fgf9*, *Foxp2* and *Etv5* in lungs from E14.5 *Yy1*^{flox/flox};*Shh*^{+Cre} embryos when compared with *Yy1*^{flox/+};*Shh*^{+Cre} specimens. Values are expressed as mean±s.e.m.; **P*<0.05, ***P*<0.01, ****P*<0.001. (B,C) Increased *Fgf10* expression was detected by *in situ* hybridization in lung mesenchyme surrounding the dilated airways of E12.5 *Yy1*^{flox/flox};*Shh*^{+Cre} embryos. (D,E) Phospho-ERK (pERK) expression was stronger and more abundant in the cyst epithelium in E12.5 mutants, reflecting the increased FGF10 signaling activity. (F) qRT-PCR analyses showed the reduced expression of *Shh* and its target genes *Ptc1*, *Hip1* and *Foxf1* in lungs from E14.5 *Yy1*^{flox/flox};*Shh*^{+Cre} mutants. Values are expressed as mean±s.e.m.; **P*<0.05, ***P*<0.01. (G) Upper panel: schematic representation and position relative to the TSS of the five putative YY1 binding sites in the *Shh* promoter. The position of the qPCR fragments is indicated. Lower panel: ChIP analysis of endogenous *Shh* regulatory sequences in E14.5 lung. Chromatin was immunoprecipitated with rabbit IgG, anti-YY1 and anti-histone H3 antibodies. Recruitment of YY1 and H3 on regulatory sequences of the *Shh* locus was evaluated by qPCR and indicated as the percentage of input. YY1 bound to the *Shh* genomic regions spanning the YY1 binding sites whereas no binding was observed with the *Shh* negative control. The data indicated correspond to mean±s.d. of two independent experiments. Asterisks indicate cysts. Dashed lines indicate the demarcation between the airway epithelium and the mesenchyme. Scale bars: 100 μm in B,C; 200 μm in D,E.

through sequential pathological changes; the early stage (Type I) is characterized by a pure cystic architecture that progresses over time into a mixed cystic and solid lesion (Type II), which further evolves into a purely solid aggressive tumor with anaplastic undifferentiated sarcomatous components (Type III; Messinger et al., 2015). In controls and CCAM specimens, YY1 protein was strongly expressed in lung epithelium and mesenchyme. In specimens from all three types of PPB, YY1 staining was reduced, as assessed by IHC semiquantitative scoring (Fig. 7). These findings suggest a pathogenic role for YY1 in PPB.

No genetic interplay between *Yy1* and *Dicer* genes in lung morphogenesis

PPB is an extremely rare lung sarcoma that arises during fetal lung development and occurs in young children. PPB is associated with

germ line and somatic mutations in the *DICER1* gene (Hill et al., 2009; Pugh et al., 2014). In mice, the conditional loss of *Dicer* function in lung epithelium causes neonatal death due to defective lung branching and airway dilation, similar to the phenotype of *Yy1*^{flox/flox};*Shh*^{+Cre} mutants (Fig. 8A-D; Harris et al., 2006). These resemblances raised questions about a potential interplay between *Yy1* and *Dicer* genes in lung morphogenesis. We assessed *Dicer* expression levels between lungs from E14.5 *Yy1*^{flox/+};*Shh*^{+Cre} and *Yy1*^{flox/flox};*Shh*^{+Cre} embryos. No significant difference was found, indicating that YY1 did not control *Dicer* expression during lung formation (Fig. 8E). Likewise, a *Dicer* epithelial mutation did not perturb *Yy1* lung expression in *Dicer*^{flox/flox};*Shh*^{+Cre} mutants at the RNA and protein levels (Fig. 8F; not shown).

As reported, the *Dicer* mutation caused increased *Fgf10* expression (*P*<0.001) (Fig. 8F; Harris et al., 2006). We also found

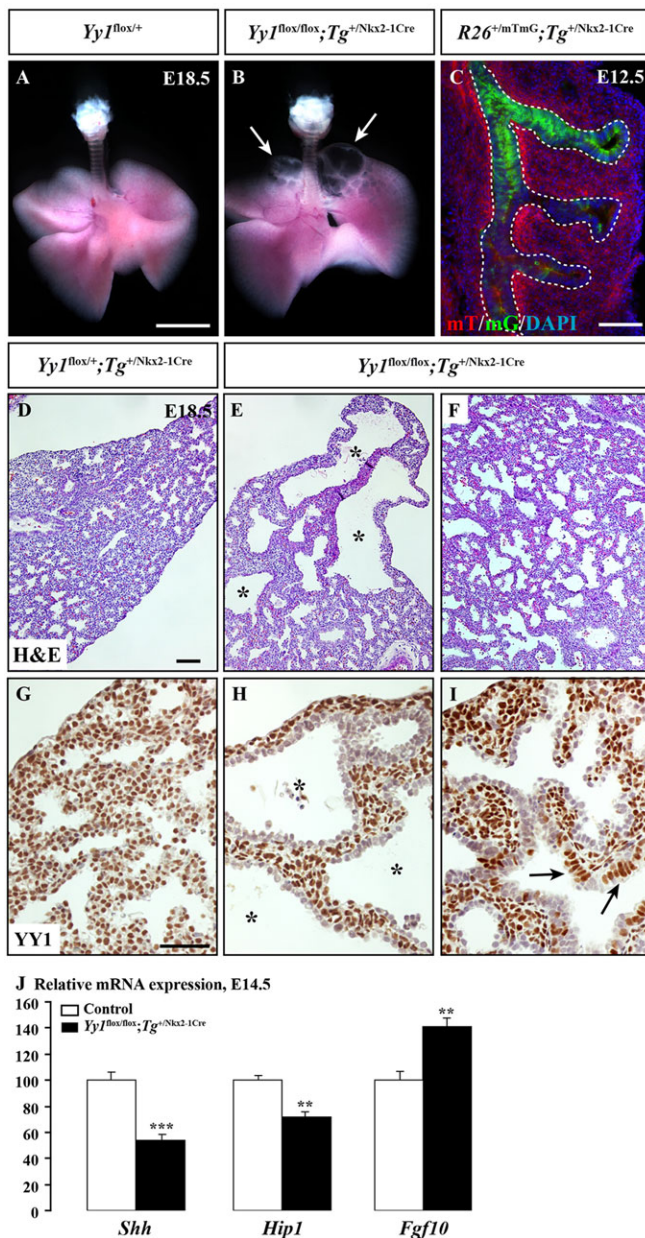


Fig. 5. *Yy1* inactivation in the developing lung endoderm with the *Nkx2-1Cre* transgenic mouse line causes cyst formation. (A,B) Lungs from E18.5 *Yy1^{lox/lox};Tg^{+/Nkx2-1Cre}* embryos exhibited cysts in the proximal region of the lobes (arrows). (C) A robust Cre activity was detected in the proximal respiratory epithelium from E12.5 *R26^{+/mTmG};Tg^{+/Nkx2-1Cre}* embryos. (D-F) H&E-stained lung sections revealed that E18.5 *Yy1^{lox/lox};Tg^{+/Nkx2-1Cre}* mutants presented defective airway branching as shown by the formation of cysts (asterisks). Panels E and F correspond to the proximal and distal parts of the lung, respectively. (G-I) YY1 immunostaining demonstrated a near-complete loss of YY1 expression in the epithelium lining cysts in the proximal portion of the lungs from E18.5 *Yy1^{lox/lox};Tg^{+/Nkx2-1Cre}* mutants (H). The deletion was incomplete in distal lung as shown by YY1-positive cells (arrows; I). (J) qRT-PCR analysis revealed decreased *Shh* and *Hip1* expression and increased *Fgf10* levels in lungs from E14.5 *Yy1^{lox/lox};Tg^{+/Nkx2-1Cre}* embryos. Values are expressed as mean±s.e.m.; ** $P<0.01$, *** $P<0.001$. Asterisks indicate cysts. Scale bars: 2 mm in A,B; 100 μ m in C-F; 50 μ m in G-I.

reduced expression of *Shh* ($P<0.01$) and its targets *Ptc1* ($P<0.01$) and *Hip1* ($P<0.05$) in *Dicer^{lox/lox};Shh^{+/Cre}* specimens compared with *Dicer^{lox/lox};Shh^{+/Cre}* controls (Fig. 8F). Therefore, both *Yy1* and

Dicer genes regulate *Shh* expression and subsequent lung branching morphogenesis.

To address if genetic interactions occur between the *Yy1* and *Dicer* genes *in vivo*, we generated *Yy1^{lox/+};Dicer^{lox/+};Shh^{+/Cre}* double heterozygous animals. Like single heterozygous mutants, *Yy1^{lox/+};Dicer^{lox/+};Shh^{+/Cre}* embryos did not present lung defects, suggesting no genetic interplay (not shown). Accordingly, *Shh* expression levels were similar in lungs from single and double heterozygous mutants and comparable to that detected in *Shh^{+/Cre}* specimens (Fig. 8G). Thus, the homozygous mutation of either *Yy1* or *Dicer* in lung epithelium is required to significantly reduce *Shh* expression.

Microarray analysis of RNA from lungs of E14.5 *Shh^{+/Cre}*, *Yy1^{lox/lox};Shh^{+/Cre}* and *Dicer^{lox/lox};Shh^{+/Cre}* embryos was performed to assess the molecular consequences of the *Yy1* and *Dicer* mutations in lung epithelium (supplementary material Fig. S5). An analysis of the total number of differentially expressed genes with a fold change ≥ 1.5 and $P<0.05$ revealed an overlap of 210 genes of which only seven were regulated in opposite direction. This suggested that a core pattern of altered gene expression is associated with airway dilation common to the two models. A heat map was established. In addition to confirming the modified expression of *Shh*, *Hip1* and *Fgf10* genes, it revealed the upregulation of *Cdkn1a*, known to be repressed by YY1 (Affar et al., 2006). *Cdkn1a* encodes the cyclin-dependent kinase inhibitor p21, and its upregulation concurred with the reduced proliferation observed in mutants. The detected decrease in expression of *Elf5*, *Irx2*, *Irx3* and *Irx5* genes, encoding transcriptional regulators of lung morphogenesis, also corroborates the observed lung phenotype. Inhibition of *Irx* expression has been shown to cause reduced lung branching (van Tuyl et al., 2006). *Elf5* misexpression in lung epithelium also disrupts branching, suggesting that precise levels of *Elf5* are required for lung morphogenesis (Metzger et al., 2008).

***Yy1^{lox/lox};Tg^{+/Nkx2-1Cre}* mice present a PPB-like phenotype**

The neonatal lethality of *Yy1^{lox/lox};Shh^{+/Cre}* mice precluded the study of lung tumor formation and progression. However, proliferation assays in lungs from E18.5 *Yy1^{lox/lox};Shh^{+/Cre}* and *Dicer^{lox/lox};Shh^{+/Cre}* embryos revealed a dramatic increase in cell proliferation of the cystic walls, suggesting tissue overgrowth (Fig. 9A-D).

The lung phenotype of *Yy1^{lox/lox};Tg^{+/Nkx2-1Cre}* embryos was less severe than the *Yy1^{lox/lox};Shh^{+/Cre}* phenotype. However, increased proliferation was also detected in lungs from E18.5 *Yy1^{lox/lox};Tg^{+/Nkx2-1Cre}* embryos (supplementary material Fig. S4K,L). Most *Yy1^{lox/lox};Tg^{+/Nkx2-1Cre}* mutants died at birth but some survived up to weaning age (Table 2). Out of 86 pups, seven *Yy1^{lox/lox};Tg^{+/Nkx2-1Cre}* mice born alive survived until weaning or were sacrificed due to health problems. The lungs of these seven mice exhibited cysts in the proximal region and histological analyses showed a multiloculated structure with septa of variable thicknesses around cysts (Fig. 9E-G). Elevated proliferation was observed in the disorganized mass (Fig. 9I,J). Vimentin immunostaining confirmed the mesenchymal nature of the tissue (Fig. 9K,L). Putative primitive small mesenchymal cells, a hallmark of type I PPB, were also detected (Fig. 9H). Finally, NKX2-1 and T1 α positive cells revealed the alveolar nature of the cyst epithelium (Fig. 9O,P). Altogether, these characteristics are reminiscent of the histologic features of an evolving type I PPB (Hill et al., 2008).

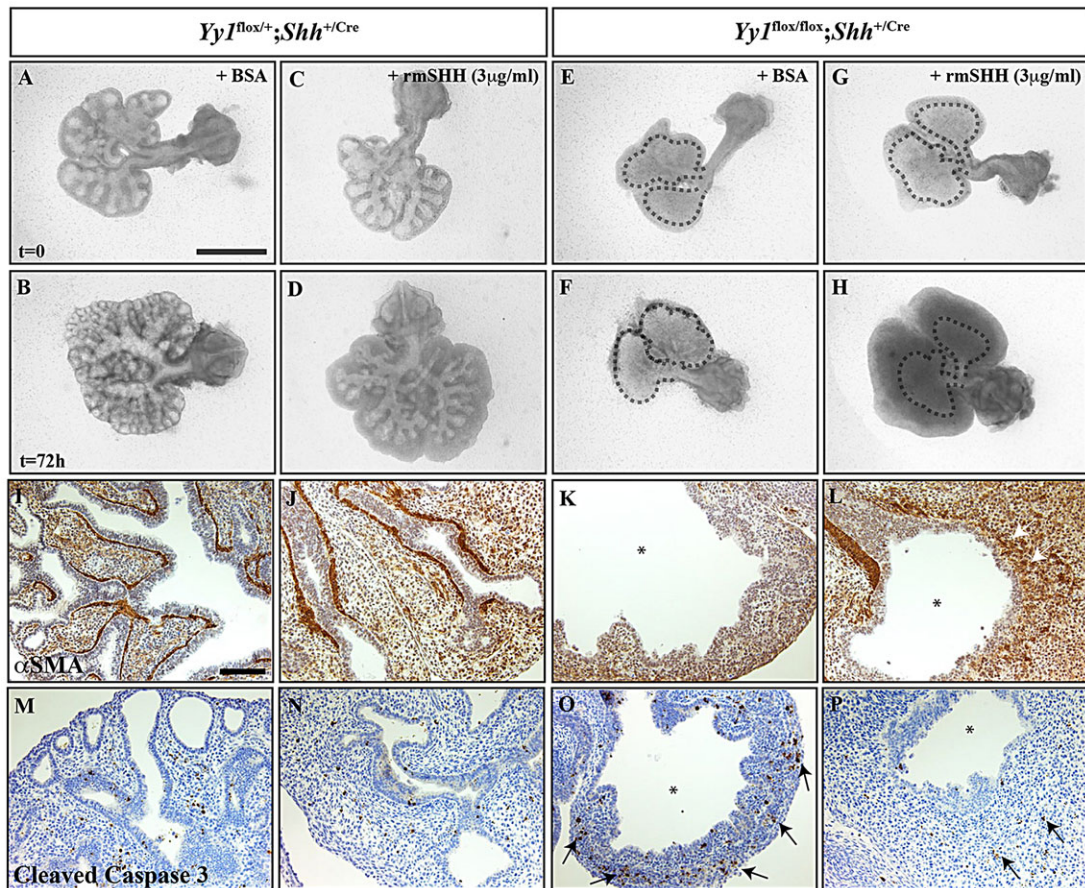


Fig. 6. Addition of rmSHH partially rescues the airway smooth muscle cells and the apoptotic defects in $Yy1^{flox/flox};Shh^{+/Cre}$ lung explants. Lung explants from E12.5 $Yy1^{flox/+};Shh^{+/Cre}$ (A-D,I,J,M,N) and $Yy1^{flox/flox};Shh^{+/Cre}$ (E-H,K,L,O,P) embryos were cultured for 3 days with (C,D,G,H,J,L,N,P) or without (A,B,E,F,I,K,M,O) rmSHH. BSA-treated $Yy1^{flox/flox};Shh^{+/Cre}$ explants exhibited cysts and failed to branch (E,F). They showed a loss of α SMA immunoreactivity around the cysts (I,K) and an increased number of apoptotic cells (M,O). Addition of rmSHH caused an increased size of the mesenchymal layer in both $Yy1^{flox/+};Shh^{+/Cre}$ and $Yy1^{flox/flox};Shh^{+/Cre}$ explants but did not improve lung branching in mutants (C,D,G,H). rmSHH partially restored α SMA expression in $Yy1^{flox/flox};Shh^{+/Cre}$ explants (I-L). The number of apoptotic cells in $Yy1^{flox/flox};Shh^{+/Cre}$ lungs supplemented with rmSHH returned to control levels (M-P). Arrows indicate cleaved caspase 3-positive cells. Asterisks indicate cysts, which are delineated by a dashed line. ($n=3-5$ per condition). Scale bars: 1 mm in A-H; 100 μ m in I-P.

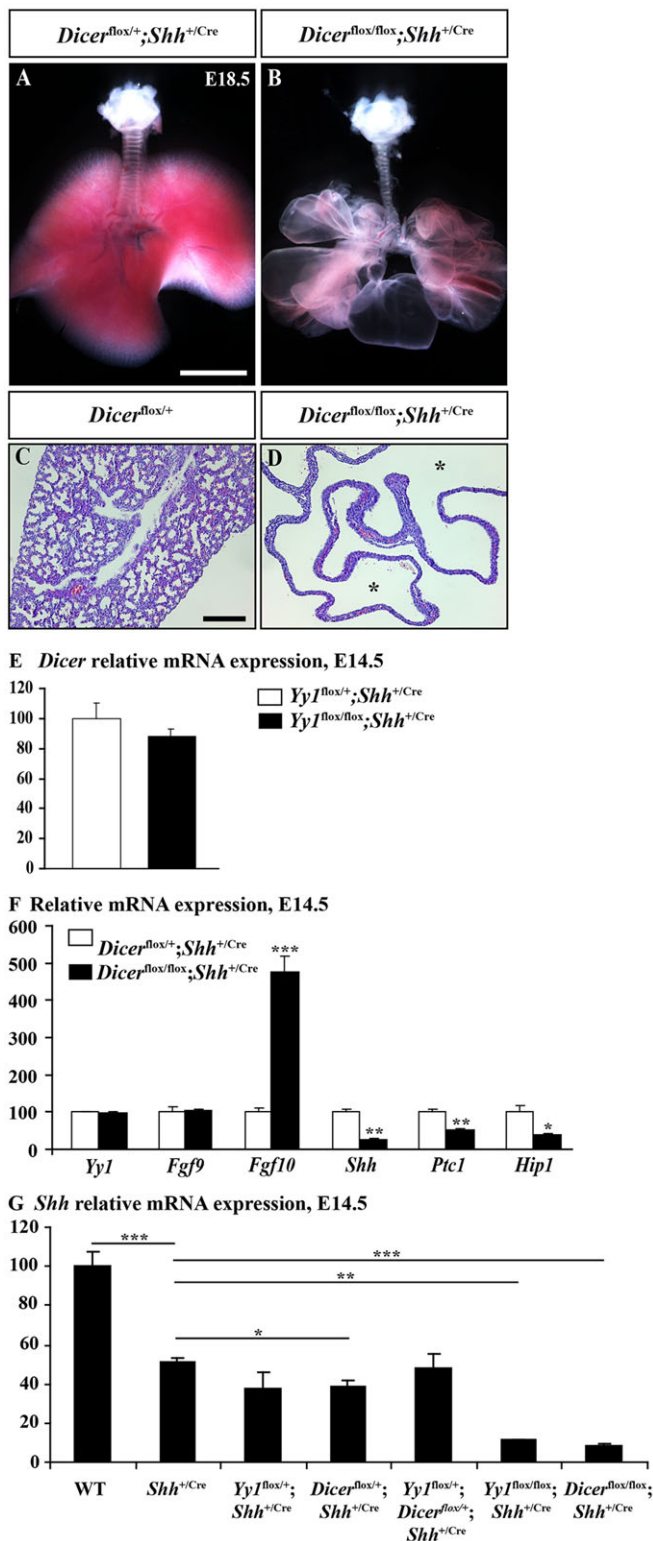
DISCUSSION

YY1 is a multifunctional transcription factor member of the Polycomb group protein family, and its actions are controlled by protein-protein interactions (Gordon et al., 2006). YY1 plays pivotal roles throughout development. In the lung, there is a dosage-dependent requirement for YY1 with a lower limit of expression for survival as hypomorph mutants die at birth from respiratory distress due to collapsed lungs (Affar et al., 2006). This phenotype was reproduced when we specifically ablated *Yy1* function in the mesenchyme. Moreover, *Yy1* lung mesenchymal deletion affects epithelial cell differentiation indicating an YY1 non-cell autonomous role (Bérubé-Simard et al., 2014). Here, we demonstrated that the *Yy1* mutation in lung epithelium also impacts on lung formation. *Yy1* function is thus required in both mesenchymal and epithelial cell layers for the correct development of the respiratory tract.

Epithelial inactivation of the *Yy1* gene inhibited lung branching but maintained distal epithelial cell differentiation, which agrees with the notion of negative correlation between branching and alveolar differentiation (Chang et al., 2013). The *Yy1* mutation also caused the formation of large cysts that mimicked the *Shh* lung phenotype (Litingtung et al., 1998; Picicelli et al., 1998). *Shh* expression was decreased in *Yy1* mutant lungs independently of the *Cre* allele used. SHH, a diffusible factor secreted by the lung

epithelium, inhibits *Fgf10* expression in lung mesoderm (Bellusci et al., 1997). Consequently, *Fgf10* expression was upregulated in *Yy1* mutants, resulting in the loss of *Fgf10*-expressing foci in lung mesenchyme that normally trigger branching events. Gain in *Fgf10* expression may generate a driving force favoring the cystic dilation of airways in *Yy1* mutants (Fig. 10). This mechanistic explanation is further supported by the observations that trans-uterine intraparenchymal microinjections of a vector carrying an *Fgf10* transgene induce cystic lung malformations in rat fetuses that recapitulate the phenotype of human congenital cystic lung diseases (Gonzaga et al., 2008). Therefore, our work uncovers a functional cascade sequentially implicating *Yy1*, *Shh* and *Fgf10* genes in lung branching coordination.

One *Shh* wild-type allele remains in $Yy1^{flox/flox};Shh^{+/Cre}$ specimens, raising concerns about the impact of *Shh* haploinsufficiency on the lung phenotype. We have shown that: (1) $Yy1^{+/flox};Shh^{+/Cre}$ specimens did not present lung anomalies; (2) *Shh* expression was reduced in $Yy1^{flox/flox};Shh^{+/Cre}$ specimens compared with $Yy1^{flox/+};Shh^{+/Cre}$ samples; and (3) $Yy1^{flox/flox};Tg^{+/Nkx2-1Cre}$ and $Yy1^{flox/flox};Shh^{+/Cre}$ mutants share a similar lung phenotype. Moreover, *Shh*^{+/-} mutants do not present lung anomalies and variations in *Fgf10* expression (van Tuyl et al., 2007). These data provide concordant evidence for a role of YY1 in the regulation of *Shh* expression in the lung



Despite the resemblances between *Yy1*^{fllox/fllox};*Shh*^{+Cre} and *Dicer*^{fllox/fllox};*Shh*^{+Cre} mutant lung phenotypes and the important overlap in differentially expressed lung genes, production of *Yy1*^{fllox/+};*Dicer*^{fllox/+};*Shh*^{+Cre} double heterozygous mice and expression analyses did not provide evidence for a genetic interaction between *Yy1* and *Dicer* genes during lung morphogenesis. Therefore, *Dicer* and *Yy1* may act via distinct pathways to control lung development.

Fig. 8. Loss of *Dicer* function in the developing lung epithelium phenocopies the lung defects of *Yy1* mutants. (A,B) Lung from E18.5 *Dicer*^{fllox/fllox};*Shh*^{+Cre} embryos showed dilated sacs. (C,D) H&E-stained lung sections revealed that lungs from E18.5 *Dicer*^{fllox/fllox};*Shh*^{+Cre} mutants exhibited defects in airway branching causing cyst formation (asterisks). (E) *Dicer* expression was unchanged in lungs from E14.5 *Yy1*^{fllox/fllox};*Shh*^{+Cre} embryos, as detected by qRT-PCR. (F) qRT-PCR analysis revealed increased levels of *Fgf10* expression and reduced expression levels for *Shh*, *Ptc1*, and *Hip1* in the lungs of E14.5 *Dicer*^{fllox/fllox};*Shh*^{+Cre} embryos when compared with *Dicer*^{fllox/+};*Shh*^{+Cre} specimens. (G) Comparative analysis of *Shh* expression levels in lungs from E14.5 *Yy1*;*Dicer*;*Shh*^{+Cre} mutant embryos. *Shh* expression levels in single and double heterozygous mutants were comparable to that of *Shh*^{+Cre} specimens corresponding to ~40-50% of the levels in wild-type (WT) controls. In *Yy1*^{fllox/fllox};*Shh*^{+Cre} embryos and *Dicer*^{fllox/fllox};*Shh*^{+Cre} embryos, *Shh* expression levels were ~10% of that of WT specimens. (E-G) Values are expressed as mean±s.e.m.; **P*<0.05, ***P*<0.01, ****P*<0.001. Scale bars: 2 mm in A,B; 200 μm in C,D.

Although our data pointed toward a direct transcriptional regulation of *Shh* expression by YY1, the mechanisms of DICER action in lung epithelium remain elusive (Harris et al., 2006).

YY1 acts on target genes via the recruitment of co-factors, and elucidating YY1 partners may unveil the molecular mechanisms underlying the lung phenotype. For instance, YY1 recruits protein modifiers, such as Ezh2 and histone deacetylases (HDAC), that mediate posttranslational modifications involved in chromatin remodeling (Deng et al., 2010). The epithelial deletion of *Ezh2* function does not cause cyst formation (Snitow et al., 2015). Conversely, the combined epithelial mutations of *Hdac1* and *Hdac2* genes led to airway dilation and to upregulated *Cdkn1a* expression, raising the possibility that YY1 and HDAC act together to some extent during lung morphogenesis (Wang et al., 2013).

Aberrations in lung developmental processes may give rise to structural abnormalities, such as congenital cystic diseases that encompass a spectrum of rare but clinically significant conditions like CCAM and PPB. PPB is characterized in its earliest manifestation by large cysts lined by an alveolar-type epithelium and is often mistaken for CCAM. However, cyst epithelium from CCAM presents a bronchial-like appearance (Hill et al., 2008). There is no known genetic cause for CCAM (Kotecha et al., 2012). Conversely, heterozygous germ line *DICER1* mutations were identified as one major predisposing factor for developing PPB or related conditions such as cystic nephroma and ovarian tumors (Boman et al., 2006; Hill et al., 2009; Slade et al., 2011; Messinger et al., 2015; Pugh et al., 2014). It is proposed that the loss of *DICER1* function occurs primarily in the non-neoplastic epithelial component of the tumor, leading to dysregulation of tumor-promoting factors that stimulate proliferation of the mesenchyme and predispose primitive mesenchymal cells to sarcomatous transformation (Hill et al., 2009). Heterozygous germ line *DICER1* loss-of-function mutations were found in patients with both sporadic and familial PPB. However, carriers with one *DICER1* mutated allele are phenotypically normal, suggesting that secondary genetic events are required for tumor formation. About 30% of PPB patients do not have mutations in the *DICER1* gene further supporting the implication of other genes (Messinger et al., 2015).

A whole exome sequencing analysis of PPB patients revealed somatic *DICER1* missense mutations, but no *YY1* mutations were found (Pugh et al., 2014). The somatic *DICER1* mutations were mainly localized in the RNaseIIIb domain responsible for the cleavage of the 3' end from the -5p strand of microRNA precursors. Mutations of the RNaseIIIb domain were shown to cause a -3p mature microRNA strand bias due to the loss of -5p strand cleavage of the pre-microRNAs (Anglesio et al., 2013).

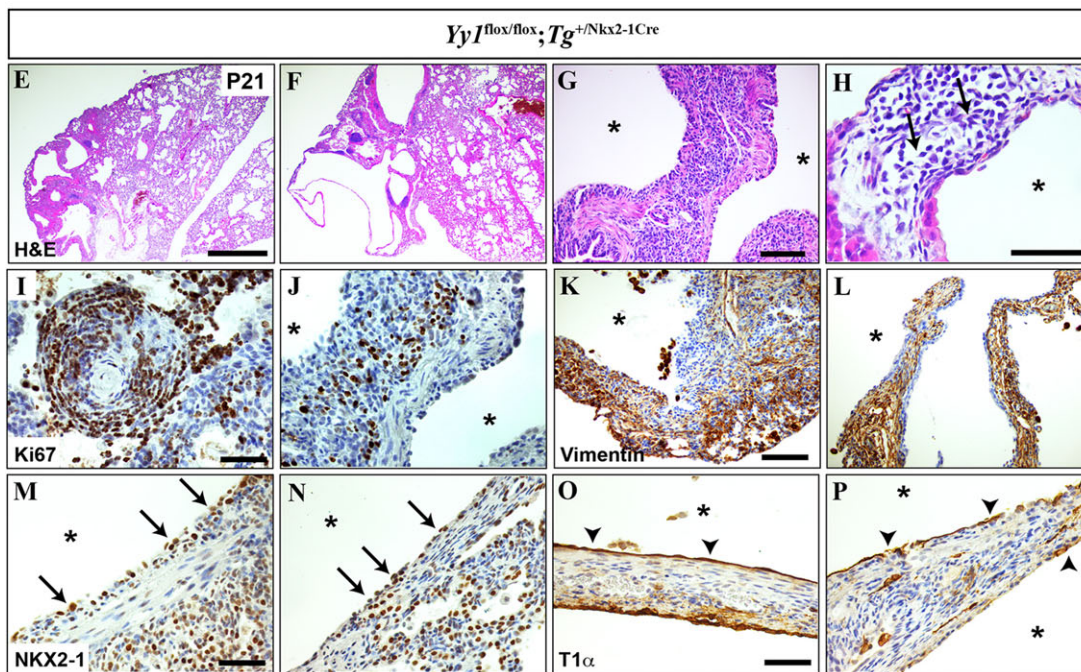
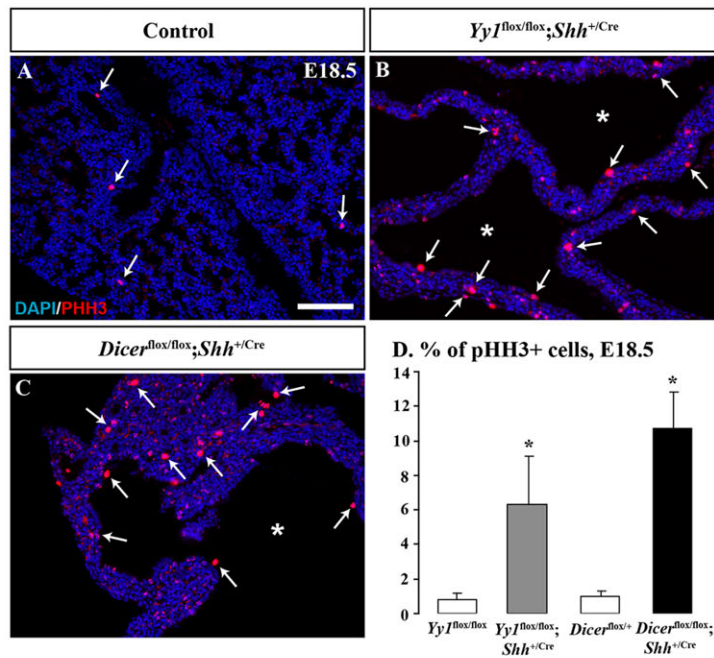


Fig. 9. *Yy1*^{flox/flox}; *Tg*^{+/Nkx2-1Cre} mice present characteristics of an evolving type I pleuropulmonary blastoma-like phenotype. (A-D) pHH3-positive cells (A-C, arrows) revealed increased proliferation (D) in lungs from E18.5 *Yy1*^{flox/flox}; *Shh*^{+/Cre} and *Dicer*^{flox/flox}; *Shh*^{+/Cre} embryos. Values are expressed as mean±s.d.; **P*<0.05. (E-H) H&E-stained lung sections from P21 *Yy1*^{flox/flox}; *Tg*^{+/Nkx2-1Cre} mice exhibited alterations in lung architecture with multilocular cysts and variable septal thickness. Arrows in H indicate primitive small mesenchymal cells within the cyst walls. (I,J) Ki67 IHC showed high proliferation levels of mesenchymal cells within the cystic wall. (K-N) Vimentin immunostaining confirmed the mesenchymal identity of the cystic walls (K,L) whereas NKX2-1 IHC established the epithelial nature of the cells lining the cyst (M,N; arrows). (O,P) T1α-positive cells (arrowheads) along the cystic epithelium revealed its alveolar nature. Asterisks indicate cysts. Scale bars: 100 μm in A-C,G,K,L; 1 mm in E,F; 50 μm in H-J,M-P.

Serum microRNA profiling from a PPB patient carrying a somatic mutation in the *DICER1* RNaseIIIb domain showed the overexpression of microRNAs largely derived from the -3p strand (Murray et al., 2014). Among the microRNAs presenting the higher fold change were *let-7a-3p* and *let-7b-3p*, which are predicted to target the human *YY1* gene (<http://mirdb.org>). It is tempting to speculate that abnormal levels of particular

miRNAs may alter YY1 expression providing a mechanistic explanation for the reduced YY1 levels seen in lungs from PPB patients (Fig. 10).

Because *YY1* expression and function are closely associated with cell-cycle progression and apoptosis, the potential role of YY1 in cancer was extensively explored. Both increased and decreased *YY1* expression levels were linked to a wide range of cancers (Wang et al.,

Table 2. Ratios of genotypes in litters from crosses between *Yy1^{flox/+};Tg^{+/Nkx2-1Cre}* and *Yy1^{flox/flox}* mice

Age	# litters	# pups	<i>Tg^{+/Nkx2-1Cre}</i>			
			<i>Yy1^{flox/+}</i>	<i>Yy1^{flox/flox}</i>	<i>Yy1^{flox/+}</i>	<i>Yy1^{flox/flox}</i>
Expected (%)			(25)	(25)	(25)	(25)
E14.5	3	23	9 (39.1)	7 (30.4)	2 (8.8)	5 (21.7)
E18.5	2	14	4 (28.6)	4 (28.6)	3 (21.4)	3 (21.4)
P21	14	86	17 (19.8)	37 (43)	25 (29.1)	7 (8.1)

The percentage obtained is indicated in parentheses.

2008; Nicholson et al., 2011). Analysis of *Yy1^{flox/flox};Shh^{+/Cre}* lungs at late gestation revealed a burst in cell proliferation in both epithelium and mesenchyme that may reflect a potential subsequent malignant transformation. The reduced expressivity of the lung phenotype of *Yy1^{flox/flox};Tg^{+/Nkx2-1Cre}* mutants allowed us to overcome the neonatal death of *Yy1^{flox/flox};Shh^{+/Cre}* pups. The cystic architectural and histological features observed in mutant mice were reminiscent of an evolving type I PPB, further supporting the notion that reduced expression of YY1 may participate in PPB pathogenesis. It must be underscored that the unrepressed *Fgf10* expression seen in *Yy1* mutants should also be considered to play a role in cancer development as *Fgf10* overexpression in the lung during the postnatal period can cause multifocal pulmonary tumors (Clark et al., 2001).

In conclusion, we present evidence of the crucial role played by the transcription factor YY1 in the developing epithelium of the respiratory tract for lung branching morphogenesis. Although further works are needed to fully elucidate the underlying mechanisms, SHH appears to be a key player in mediating YY1 function in the lung. Our *Yy1* mouse models also reproduced characteristics of the rare early childhood cancer PPB and indicated that a more precise gene dosage mouse model operating within the context of a developmental window should

provide a strong model for the exploration of PPB molecular pathogenesis.

MATERIALS AND METHODS

Mice, genotyping and tissue collection

Yy1^{flox/flox} mice were obtained from Dr Shi (Affar et al., 2006). The *Rosa26* reporter line [*Gt(ROSA)26Sor^{tm4}(ACTB-tdTomato,-EGFP)^{Luo}*], the *Dicer1^{flox/flox}* conditional line (*Dicer1^{tm1Bdh/J}*), and the *Shh^{Cre}* [*Shh^{tm1(EGFP/cre)Cjt}*] and *Tg^{+/Nkx2-1Cre}* [*Tg^{(Nkx2-1-cre)2Sand}*] deleter strains were purchased from The Jackson Laboratory (Harfe et al., 2004, 2005; Muzumdar et al., 2007; Xu et al., 2008). As only individuals carrying the *Yy1^{flox/flox};Shh^{+/Cre}*, *Yy1^{flox/flox};Nkx2-1-Cre* and *Dicer1^{flox/flox};Shh^{+/Cre}* genotypes presented defects, all the other genotypes were referred hereafter as controls except when specified. Age of the embryos was estimated by considering the morning of the day of the vaginal plug as E0.5. Experimental specimens were genotyped by PCR analyses. Lungs were collected at E12.5, E14.5, E18.5 or postnatal day (P) 21 as described (Boucherat et al., 2014). For RNA extraction, lungs were snap-frozen in N₂. Experiments were performed according to the guidelines of the Canadian Council on Animal Care and approved by the institutional animal care committee.

Human tissues

This study was conducted with anonymised specimens of CCAM and PPB patients from the Department of Pathology of Hôpital Necker-Enfants Malades, Paris, France and from McGill University, Montréal, Canada and

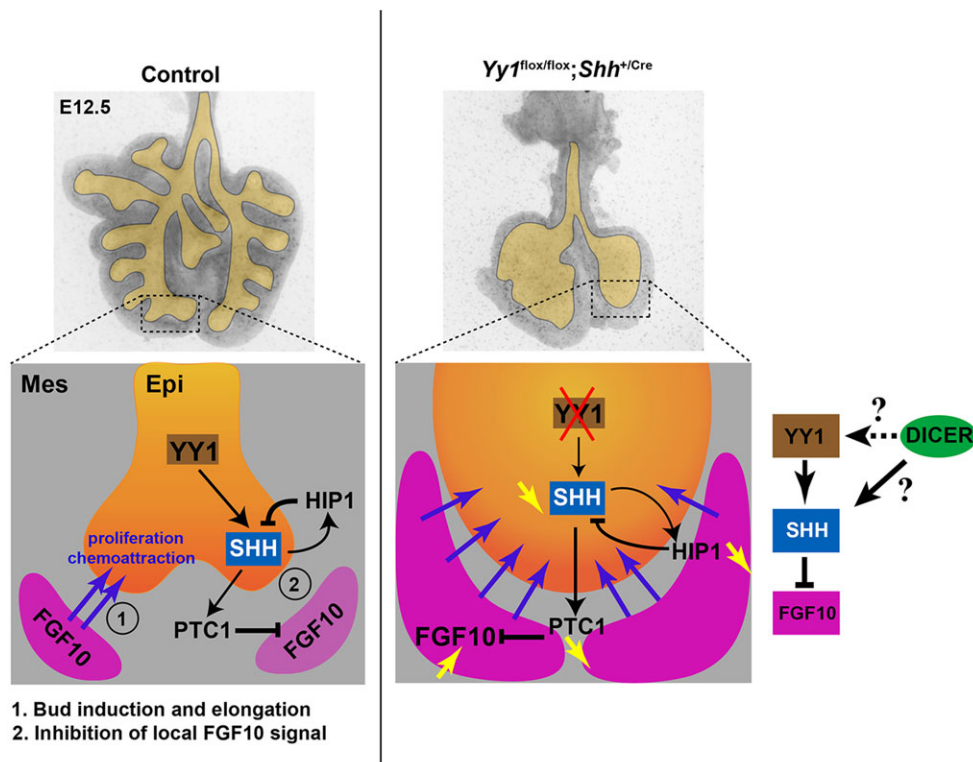


Fig. 10. Model for the action of YY1 during lung branching morphogenesis.

During lung branching morphogenesis, FGF10 localization and expression are dynamically regulated within the mesenchyme to induce epithelial proliferation and budding. Once the epithelial buds start to elongate, activation of the SHH signaling pathway occurs as a negative feedback mechanism. Consequently, the subepithelial mesenchyme proliferates, FGF10 expression levels near the bud tips decrease and FGF10-expressing foci shift laterally, promoting a next round of branching. In the absence of *Yy1* function in lung epithelium, *Shh* expression is reduced, causing an increased and diffuse expression of FGF10 throughout the mesenchyme. Lung branching is impaired, resulting in rudimentary lungs with large epithelial cysts. The unrestricted and elevated FGF10 expression represents a reasonable primary cause of airway dilation in the developing lung. Similar changes are observed in *Dicer1^{flox/flox}*, *Shh^{+/Cre}* mice. However, the mechanisms of DICER action in lung epithelium are still unknown.

approved by the respective Ethics Committee. Controls corresponded to normal lung specimens from the Department of Pathology of Hôpital Raymond Poincaré, Garches, France. They were collected at autopsy from age-matched children who died from a non-pulmonary cause (sudden infant death syndrome). Characteristics of patients are summarized in supplementary material Table S1.

Histology, immunohistochemistry (IHC) and immunofluorescence (IF) analyses

Experiments were performed as described (Boucherat et al., 2014). The Cyanine 3 Tyramide Signal Amplification Kit (PerkinElmer) was used for pERK detection. Antibodies are listed in supplementary material Table S2.

Proliferation and apoptosis

Experiments were performed as described (Boucherat et al., 2014). Three to four random fields were taken, for an average number of 650 cells per field, from four to five embryos per genotype.

In situ hybridization

RNA *in situ* hybridization was performed on 15 µm cryosections of E12.5 embryos (Schaeren-Wiemers and Gerfin-Moser, 1993). A 584-bp mouse *Fgf10* cDNA fragment was used for the digoxigenin-labeled riboprobe (provided by Dr B. Hogan, Duke University Medical Center, USA). Experiments were performed on four specimens per genotype.

Alcian Blue cartilage staining

Dissected respiratory tracts from E18.5 embryos were stained in a solution of 0.03% Alcian Blue and 20% acetic acid prepared in 95% ethanol. The external tracheal diameter was measured at five rostrocaudal locations along the most linear portion of the trachea and quantified using NIH ImageJ software. Tracheal luminal surface was measured using Leica SCN 400 F SlideScanner and SlidePath Gateway Software.

Quantitative RT-PCR (qRT-PCR)

Lung total RNA was isolated from individual E14.5 embryos. qRT-PCR experiments were performed as described (Boucherat et al., 2012). Three to eight specimens were used per genotype tested. Primer sequences are listed in supplementary material Table S3A.

Chromatin immunoprecipitation (ChIP) assays

Lungs from E14.5 wild-type embryos were collected and ChIP assay was performed with rabbit anti-YY1 antibody, rabbit anti-histone H3, or control rabbit IgG as described (Bérubé-Simard et al., 2014). qPCR-ChIP analyses were performed with primers specific to each domain containing YY1 binding sites (supplementary material Table S3B). The values for the samples immunoprecipitated by anti-YY1, anti-histone H3, or control IgG were recorded as the percentage relative to input. ChIP results were confirmed by two independent experiments. qPCR was performed in triplicate for each sample. ChIP efficiency was calculated by dividing the amount of PCR product obtained with the immunoprecipitated DNA by the amount obtained with the input DNA (Aparicio et al., 2005).

Transactivation assays

Human pCMV-YY1 and control pCMV-GFP-LpA expression vectors were obtained from Drs G. Blanck (University of South Florida, USA) and R. Aasland (University of Bergen, Norway), respectively. A 5-kb *Bg/II-XhoI* genomic fragment located between positions -5016 bp and +60 bp of the mouse *Shh* gene (relative to TSS) was cloned into the pGL3 basic luciferase reporter expression plasmid (Promega). HEK293 cells were transiently co-transfected in 24-well plates (40,000 cells/well) with 0.3 µg/well of luciferase reporter construct and 0.2 µg/well of YY1 or control expression vectors using the FuGENE 6 transfection reagent (Promega). The RL-SV40 *Renilla reniformis* luciferase expression vector (Promega) was used as an internal control for transfection efficiency (0.001 µg/well). Luciferase activity was measured 48 h after transfection with the Dual-Luciferase Reporter Assay System (Promega). Transfections were performed in triplicate in at least two independent experiments. Data

from a representative experiment are presented as the fold induction±s.d. of normalized relative luciferase activity.

Lung explant cultures

Experiments were performed as described (Boucherat et al., 2014). Lungs were kept for 72 h in serum-free DMEM/F12 medium (Gibco) in presence or not of rmSHH protein, N-Terminus (R&D Systems) at a concentration of 3 µg/ml. BSA was used as control.

Microarray analysis

Total RNA was isolated from lungs of E14.5 *Shh*^{+Cre}, *Yy1*^{flox/flox}; *Shh*^{+Cre} and *Dicer*^{flox/flox}; *Shh*^{+Cre} embryos (*n*=4/genotype). RNA quality and quantity assessment, cDNA probe preparation, hybridization to the Affymetrix Mouse Gene 2.0ST Array and image scan were performed at the Genome Quebec Innovation Centre at McGill University (Montréal, Canada). Data were pre-processed and normalized using Affymetrix Power Tools with the rma-sketch method. Raw and normalized data were uploaded to the NCBI Gene Expression Omnibus database (<http://www.ncbi.nlm.nih.gov/projects/geo>) with the accession number GSE66171 according to MIAME standards (Edgar et al., 2002). Significantly modulated probes were identified using the empirical Bayes statistics available in limma (Smyth, 2004). Probes were considered to be significantly modulated when the Benjamini-Hochberg-adjusted was significant to *P*<0.05.

Statistical analyses

Student's *t*-test was performed for comparative studies. A significance level inferior to 5% (*P*<0.05) was considered statistically significant.

Acknowledgements

We thank Drs Y. Shi for *Yy1* mice, R. Aasland, G. Blanck, B. Hogan and G. Singh for sharing reagents, P. Joubert and B. Têtu for advice in histopathology, M. Simard for judicious comments, V. Nadeau for technical help and E. Paquet for microarray data analysis.

Competing interests

The authors declare no competing or financial interests.

Author contributions

O.B. and L.J. designed the experiments; O.B., K.L.-T, F.-A.B.-S., N.H., L.B. performed the experiments; G.L., W.D.F. and C.D. provide human specimens, O.B., K.L.-T, F.-A.B.-S., N.H., L.B., J.C. and L.J. analyzed the data; O.B. and L.J. wrote the paper.

Funding

This work was supported by grants from the Canadian Institutes of Health Research [MOP-15139, L.J.; MOP-97801, J.C.], Natural Sciences and Engineering Research Council of Canada [194559, L.J.], The Cancer Research Society (L.J.) and Université Laval Foundation (L.J.).

Supplementary material

Supplementary material available online at <http://dev.biologists.org/lookup/suppl/doi:10.1242/dev.120469/-DC1>

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