

The Transcription Factors Grainyhead-like 2 and NK2-Homeobox 1 Form a Regulatory Loop That Coordinates Lung Epithelial Cell Morphogenesis and Differentiation^{*§}

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Background: *Grhl2* regulates cell-junction gene transcription in several epithelia but has not been fully characterized in lungs.

Results: In lung epithelial cells GRHL2 regulates cell-cell interaction genes, collective cell migration, and *Nkx2-1* transcription. Conversely, NKX2-1 regulates transcription of *Grhl2*.

Conclusion: A *Grhl2*- and *Nkx2-1*-positive transcriptional loop coordinates morphogenesis and differentiation of lung epithelial cells.

Significance: This regulatory loop reinforces normal lung epithelial cell identity.

The Grainyhead family of transcription factors controls morphogenesis and differentiation of epithelial cell layers in multicellular organisms by regulating cell junction- and proliferation-related genes. Grainyhead-like 2 (*Grhl2*) is expressed in developing mouse lung epithelium and is required for normal lung organogenesis. The specific epithelial cells expressing *Grhl2* and the genes regulated by *Grhl2* in normal lungs are mostly unknown. In these studies we identified the NK2-homeobox 1 transcription factor (*Nkx2-1*) as a direct transcriptional target of *Grhl2*. By binding and transcriptional assays and by confocal microscopy we showed that these two transcription factors form a positive feedback loop *in vivo* and in cell lines and are co-expressed in lung bronchiolar and alveolar type II cells. The morphological changes observed in flattening lung alveolar type II cells in culture are associated with down-regulation of *Grhl2* and *Nkx2-1*. Reduction of *Grhl2* in lung epithelial cell lines results in lower expression levels of *Nkx2-1* and of known *Grhl2* target genes. By microarray analysis we identified that in addition to *Cadherin1* and *Claudin4*, *Grhl2* regulates other cell interaction genes such as semaphorins and their receptors, which also play a functional role in developing lung epithelium. Impaired collective cell migration observed in *Grhl2* knock-

down cell monolayers is associated with reduced expression of these genes and may contribute to the altered epithelial phenotype reported in *Grhl2* mutant mice. Thus, *Grhl2* functions at the nexus of a novel regulatory network, connecting lung epithelial cell identity, migration, and cell-cell interactions.

During lung organogenesis, the progenitor cell layer lining the developing airways maintains its epithelial characteristics while proliferating and differentiating into multiple cell types to form a functional lung (1). Genes that establish and maintain cell polarity and cell junctions must be coordinately regulated with cell-specific differentiation programs to generate multiple lung epithelial cell types (2–4).

In *Drosophila*, the CP2 family transcription factor Grainyhead (*Grh*) contributes to tracheal organogenesis by regulating the size of the apical membrane and morphogenesis of tracheal epithelial cells (5). In vertebrates, three homologs of *Grh*, Grainyhead like-1, -2, and -3 (*Grhl1*, *Grhl2*, and *Grhl3*), share high similarity in their DNA binding sequence and in the carboxyl terminal dimerization region but display a different pattern of expression in the lung and other epithelial tissues (6). Previous *in situ* hybridization analyses indicate that *Grhl2* is the only family member that is highly expressed in distal lung epithelium throughout development, although the particular cells expressing *Grhl2* have not been identified nor has its functional role in lung epithelium. *Grhl1* and *Grhl3*, in contrast, are expressed in proximal lung epithelium until E15.5, but later their expression is reduced in bronchi/brechchioles and is undetectable in the alveolar lung epithelium (6).

Grhls seem to have conserved functions controlling cell shape, cell growth, cell proliferation, and cell fate (7–14). They maintain epithelial cell characteristics by regulating cell-cell junction genes including the desmosomal cadherin Desmoglein-1 (*Dsg1*) by *Grhl1* and *Claudin1* (*Cldn1*) and Occludin

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(*Ocdn*) by *Grhl3*. Recently, E-cadherin (*Cdh1*) and *Cldn4* were identified as direct transcriptional targets of *Grhl2*. *Grhl2* null mutant mice die by embryonic day E11.5 (15) due to defects in neural tube closure and defective apical junction complex composition in epithelial tissues. Expression patterns of *Cdh1* and *Cldn4* were drastically reduced in foregut endoderm and otic epithelium as well as in the surface ectoderm, indicating that apical junction genes are regulated *in vivo* by *Grhl2*. Due to the early lethality of *Grhl2* null mutants, studies at later stages of development were not possible. Similarly, *N*-ethyl-*N*-nitrosourea (ENU)⁴-generated *Grhl2* mutant mice die by E12.5 due to defects in neural tube closure and heart development (16). Apical junction gene expression in epithelial organs was also reduced. A few embryos that survived to E18.5 had smaller lungs, disorganized epithelial apical junctions, and collapsed alveolar sacs, suggesting a functional role for *Grhl2* in lung development and regulation of lung epithelial genes.

Herein we identify genes regulated by *Grhl2* in lung epithelial cells and provide evidence for a novel positive transcriptional feedback loop between *Grhl2* and the homeobox transcription factor *Nkx2-1* in embryonic lung. The critical role of *Nkx2-1* in regulating epithelial cell proliferation and differentiation and of *Grhl2* in regulating cell-cell interactions and epithelial structure suggest that the *Grhl2-Nkx2-1*-positive feedback loop may serve to differentiate and reinforce distal lung epithelial phenotypes.

EXPERIMENTAL PROCEDURES

Cell Lines and Tissues—E10 is a spontaneously immortalized adult mouse lung epithelial cell line (17) provided by Dr. A. Malkinson (University of Colorado) and Dr. R. J. Ruch (University of Toledo). MLE15 (18) is an adult mouse lung epithelial cell line provided by Dr. J. A. Whitsett (Cincinnati Children's Hospital Medical Center). Cells were cultured in conditions described previously (19). Lung tissues used in chromatin immunoprecipitation analysis and immunohistochemistry were dissected from CD1 mice (Charles River Laboratories). All experiments were performed according to protocols approved by the Institutional Animal Care and Use Committee at Boston University Medical Center in accordance with National Institute of Health guidelines.

Isolation and Culture of Murine Alveolar Type II Cells—Cells were isolated from SFTPC-Gfp mice provided by Dr. J. K. Heath (University of Birmingham, Birmingham, UK) (20) using a previously described method (21) with some modifications. SFTPC-Gfp mice were anesthetized, and tracheae were exposed and cannulated with a 20-gauge luer stub adapter. Lungs were perfused with 10 ml of 1× Phosphate Buffered Saline (PBS) via the pulmonary artery, and 1–2 ml of dispase (2.4 units/ml, Roche Diagnostics, #14435800) was rapidly instilled through the tracheal cannula followed by 0.5 ml of low-melting agarose solution warmed at 55 °C. Lungs were immediately covered with ice for 2 min to gel the agarose, removed from the animals, and then incubated in dispase II/collagenase A (Roche Diagnostics) (2.4 units/ml/0.1%) solu-

tion for 45 min at 37 °C. After this incubation, lung tissues were treated with DNase 1 (0.025 mg/ml, Qiagen) on ice for 5 min. Cells in suspension were then filtered through a 100-μm filter and centrifuged at 800 rpm for 6 min at 4 °C. Cells were resuspended with 1 ml of RBC lysis buffer (Invitrogen) for 90 s before adding DMEM (Invitrogen) without serum (6 ml for each lung). FBS (Invitrogen) (0.5 ml) was added to the bottom of the tube, and cells were centrifuged at 800 rpm for 5 min at 4 °C. The pellet was resuspended in PF10 (PBS supplemented with 10% FBS), and cells were filtered just before sorting in a MoFlow instrument (Dako Cytomation) at the Boston University School of Medicine flow cytometry core. Cells with high GFP and high side-scatter signals were either collected in RLT buffer (RNAeasy kit, Qiagen) for RNA isolation or in DMEM medium supplemented with 20% FBS for culturing on fibronectin-coated or plain 24-well culture plates (Corning).

RNA Purification and Real Time RT (qRT)-PCR—Total RNA was isolated from mouse lung tissue, sorted cells, and cell lines using the RNAeasy kit (Qiagen) and treated with DNase1 (Qiagen). Isolated RNA was reverse-transcribed (RT) using TaqMan Reverse Transcription Reagents (Applied Biosystems) following the manufacturer's protocols. The StepOnePlus system (Applied Biosystems) and commercially available TaqMan gene expression assays (Applied Biosystems) were used for qRT-PCR analyses (supplemental Table S1). Reactions were performed with TaqMan PCR master mix or Fast TaqMan PCR master mix (Applied Biosystems). Relative expression levels normalized to *Gapdh* were determined using the comparative $2^{-\Delta\Delta CT}$ method.

Plasmid Construction—Full-length *Grhl2* cDNA was subcloned from the pGADT7-HA-*Grhl2* vector provided by Dr. Bogi Andersen (University of California, Irvine, CA). Briefly, the *Grhl2* cDNA was amplified by PCR using primers 5'-CAA GCG GCC GCC ATG TCA CAA GAG TCG GAC-3' and 5'CGC TGA TGG AGA TCT GAG GAT CCA TTC-3' that contain Not1 and BamH1 adaptors, respectively. This fragment was inserted in place of the dsRed gene in the dual promoter-reporter lentiviral plasmid, pCMV-dsred-UBC-Gfp (22), to generate pCMV-*Grhl2*-UBC-Gfp. The construct was verified by restriction enzyme digestion and by sequencing. This *Grhl2* vector was used in overexpression experiments in MLE15 cells. The genomic DNA fragment containing the 5' end region of the mouse *Nkx2-1* gene (−339 to −2230 bp from the second ATG site) (supplemental Fig. S1) was generated by PCR using genomic DNA from mouse 129/Ola ES cells, cloned into the pCR-BluntII-TOPO shuttle plasmid, and then subcloned into KpnI and HindIII sites of pGL3 basic vector (Promega). The −350-bp fragment of the *Nkx2-1* proximal promoter (−3 to −352 bp from the second ATG site) was generated by PCR and cloned in the pGL3 basic vector (Promega) (supplemental Fig. S1). The constructs were verified by sequencing and were identified as −2kbNkx2-1Luc and −0.35kbNkx2-1Luc. Two fragments in the first intron of the *Grhl2* gene that bind NKX2-1 protein (region H (high binding) and region L (low binding) (Fig. 6A)) were amplified by PCR from genomic DNA, cloned in pGEM T vector (Promega), and subcloned in the KpnI and SacI sites 5' to the SV40 promoter in the pGL3 promoter vector (Promega) (supplemental Fig. S1). The constructs were verified

⁴The abbreviations used are: ENU, *N*-ethyl-*N*-nitrosourea; qRT-PCR, quantitative real-time PCR; FDR, false discovery rate.

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by sequencing and identified as H-*Grhl2* Sv40Luc and L-*Grhl2* SVv40Luc.

Lentivirus Production and Transduction—To knock down *Grhl2* gene expression, a replication incompetent, vesicular stomatitis virus G-pseudotyped lentiviral vector that contained a Gfp tracking reporter gene along with either a non-targeting control shRNA (RHS4346) or the shRNA targeting *Grhl2* (V2LMM_20130) was employed (Open Biosystems). Lentiviruses were packaged and concentrated as described previously (19). MLE15 cell infections with lentiviral vectors were performed for 16 h in the presence of Polybrene (5 µg/ml) at a multiplicity of infection of 50. Stable cell lines were generated by puromycin dihydrochloride (5 µg/ml, Sigma) selection for 6 days. Cells were harvested after selection and analyzed for transduction efficiency and viability by flow cytometry analysis of GFP expression and exclusion of propidium iodide. Knockdown of *Grhl2* in the infected population was quantified at the mRNA level by qRT-PCR and at the protein level by immunostaining and Western blot analyses. Similarly, *Grhl2* was overexpressed in E10 cells by transduction of packaged bicistronic construct pCMV-*Grhl2*-UBC-Gfp, and cell morphology was compared with the cells transfected with pCMV-dsred-UBC-Gfp. *Nkx2-1* gene expression knockdown was performed as described previously (19) using a mixture of three lentiviral clones (TRCN0000020449, TRCN0000020450, and TRCN0000086264 Open Biosystems) targeting *Nkx2-1* mRNA of mouse, rat, and human origin.

Chromatin Immunoprecipitation Assays (CHIP)-PCR—Mouse lung buds (5–7 per reaction) were dissected from E11.5 day embryos and fixed in 1% formaldehyde in 1× PBS at room temperature for 15 min as described previously (23). Samples were sonicated using a Branson 450 dismembrator to achieve a chromatin fragment size of 500–2000 bp. To immunoprecipitate chromatin fragments, we incubated the samples (equal amount of DNA) at 4 °C overnight with rabbit GRHL2 (HPA00482, Sigma) or rabbit NKX2-1 (07-601, Millipore-Upstate) antibodies. Control experiments were performed with the corresponding IgG isotype (Santa Cruz Biotechnology) to determine nonspecific binding. Pre-absorbed protein A/G beads (Santa Cruz Biotechnology) were used to immunoprecipitate chromatin-antibody complexes. Equal volumes of immunoprecipitated DNA solution, 10% of the input DNA fragments, and genomic DNA were amplified by PCR using oligonucleotides spanning 228 bp of the proximal promoter of *Nkx2-1* (sense primer, 5'-GCA CAC TCT TTT GGT GGT GA-3'; antisense primer, 5'-GCA ACC AAC TTG GGG AGT TA-3') and 224 bp of the *Grhl2* first intron (sense primer, 5'-GGG TTA CGT GGC TGC TTC A-3'; antisense primer, 5'-CGT CAG GTT GCT AAG GGC A-3'). Binding of GRHL2 to the *Cldn4* promoter and to the *Cdh1* enhancer region was also determined with samples immunoprecipitated using the rabbit GRHL2 antibody and quantified by qPCR on StepOne-Plus equipment using oligonucleotides (IDT) previously described (15). Data were normalized to the input and compared with IgG control.

Electrophoretic Mobility Shift Assays (EMSA)—The DNA probes (supplemental Fig. S1) were annealed as described previously (24) and labeled with the Biotin 3' End DNA labeling kit

(Thermo Scientific). The assay was carried out using the Thermo Light Shift Chemiluminescent EMSA kit (Thermo Scientific). Samples were run in 5% TBE acrylamide gels (Bio-Rad) and detected using the Chemiluminescent Nucleic Acid Detection Module (Thermo Scientific). MLE15 cells nuclear proteins were extracted as described previously (24). 10 µg of protein was used per reaction. All binding reactions were carried out at room temperature. For competition experiments, unlabeled oligonucleotides (100-fold) were incubated with proteins for 10 min before the addition of the labeled oligonucleotide. For super shift analysis, samples were preincubated with 5 µl of NKX2-1 antibody (ab76013, Abcam), GRHL2 antibody (HPA004820, Sigma), or nonspecific rabbit IgG (Santa Cruz, sc-2027).

Immunocytochemistry—MLE15 cells were grown in conditions described previously on glass coverslips to reach 70–80% confluence, fixed with 4% formaldehyde in 1× PBS, and processed for immunostaining. For GRHL2 nuclear staining, MLE15 cells were treated with 0.01% Triton X-100 (Fisher) in 1× PBS, blocked with 1× PBS containing 2% FBS for 30 min, and incubated overnight at 4 °C with rabbit GRHL2 antibody (HPA00482, Sigma; 1:500 dilution). After three washes with 1× PBS, cells were incubated with secondary anti-rabbit FITC-conjugated antibody (Invitrogen, 1:2000 dilution) for 1 h at room temperature. Images were taken on a Leica inverted microscope or on a Leitz Aristoplan microscope. For phalloidin staining, cells were processed in a similar fashion and stained with Alexa Fluor 594 Phalloidin (Invitrogen) according to the manufacturer's protocol. Confocal imaging was performed using a Zeiss 710 microscope. Images were processed using Metamorph microscopy image analysis software to generate maximum projection image and orthogonal images of MLE15 stained with phalloidin.

Immunohistochemistry—E9.5 whole embryos as well as E15.5, E18.5, and adult dissected lungs were fixed in freshly prepared 4% paraformaldehyde in 1× PBS, pH 7.4, at 4 °C for 16 h. These samples were embedded in paraffin following standard processing with ethanol dehydration as described previously (25). Sections (6 µm) of these tissues were deparaffinized and hydrated by standard methods. Antigen retrieval was performed using an Antigen Unmasking solution (Vector Laboratories, Inc.). Endogenous peroxidase was quenched with 3% H₂O₂ in methanol for 15 min. Blocking was performed with 2% normal goat serum in 1× PBS for 1 h at room temperature. The tissues were incubated with monoclonal rabbit NKX2-1 antibody (ab76013, Abcam, 1:500 dilution) for 16 h at 4 °C or rabbit GRHL2 antibody (HPA00482, Sigma, 1:250 dilution) in 1× PBS + 0.2% Triton X-100 (PBST) and washed with 1× PBS (5 min, twice). Antibody binding was detected with the Vectastain Elite ABC kit (Vector Laboratories) that includes incubation with biotinylated tyramide signal amplification (TSA) reagent (PerkinElmer Life Sciences) and diaminobenzidine as substrate. Images were taken using a Leitz Aristoplan microscope.

For GRHL2 immunofluorescence analyses and double staining for NKX2-1 and CDH1, 6-µm tissue sections were deparaffinized at 60 °C for 1 h and hydrated by incubation in Histo-Choice Clearing Agent (Sigma) for 6 min followed by

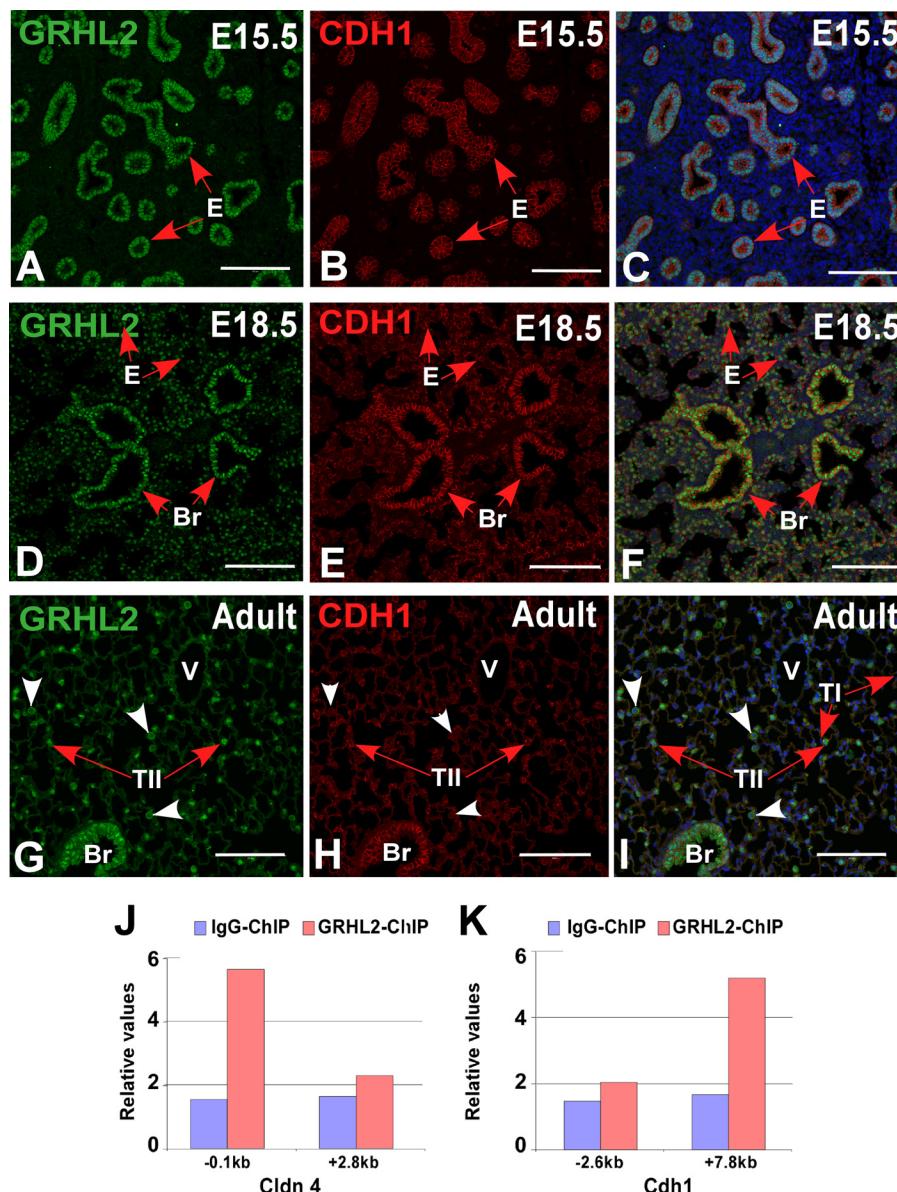


FIGURE 1. Expression and binding patterns of GRHL2 in mouse lung development. Immunofluorescence co-localization analyses of GRHL2 and one of its known targets CDH1 in days E15.5 (A–C) and E18.5 (D–F) and in adult lung (G–I) are shown. ChIP-qPCR analysis of GRHL2 binding to *Cldn4* regulatory regions (−0.1 and +2.8 kb) (J) and to *Cdh1* regulatory regions (−2.6 and +7.8 kb) (K) *in vivo* in E11.5 mouse lung. Bars, A–I, 100 μ m. E, epithelium; Br, bronchi/bronchioles; V, blood vessel; TII, alveolar type II cells; TI, alveolar type I cells.

incubations in progressive ethanol dilutions as previously described. Antigen retrieval was performed in antigen unmasking solution (Vector Laboratories) and heating in a microwave (8 min on high power and 17 min on low power). The slides were then cooled down to room temperature for 1 h. Permeabilization was performed by incubation in 1× TN buffer (20 mM Tris-HCl, pH 7.4, 150 mM NaCl) containing 0.5% Triton X-100 at room temperature for 1 h, blocked in 3% BSA in 1× TN containing 0.5% Tween 20 at room temperature for 1 h, incubated with GRHL2 antibody (HPA00482, Sigma, 1:350) or monoclonal NKX2-1 antibody (Seven Hills, 8G7G3-1, 1:350) or CDH1 antibody (BD Biosciences, 610182, 1:500) in 1× TNT (1× PBS containing 5% goat serum, 0.2% BSA, and 0.05% Tween 20) at 4 °C overnight, washed in 1× TNT for 5 min, and then exposed to Alexa Fluor-labeled antibodies. We used Alexa Fluor 488 goat anti-rabbit IgG (H+L) (A11008, Invitrogen,

1:200 dilution) in 1× TNT for 5 min and Alexa Fluor 647 goat anti-mouse IgG (H+L) (A21235, Invitrogen, 1:200). Sections were washed in 1× TNT for 5 min and mounted under coverslips with Prolong Gold with DAPI (Invitrogen). Confocal images were collected using a Zeiss 710 microscope. Autofluorescence from the red blood cells was removed by spectral imaging. The images were then processed with Zen 2009 LE software (Carl Zeiss).

Transient Co-transfection and Luciferase Assays—These assays were performed in MLE15 cells as described previously (19). Cells were co-transfected with empty pGL3 vector, −0.35kbNkx2-1Luc or −2kbNkx2-1Luc and pCMV-dsred-UBC-Gfp or pCMV-*Grhl2*-UBC-Gfp by the DEAE-dextran/chloroquine method (24). pGL3 promoter vector Sv40Luc, H-Sv40Luc, or L-Sv40Luc were co-transfected with pCMV empty vector or pCMV-*Nkx2-1* expression construct provided

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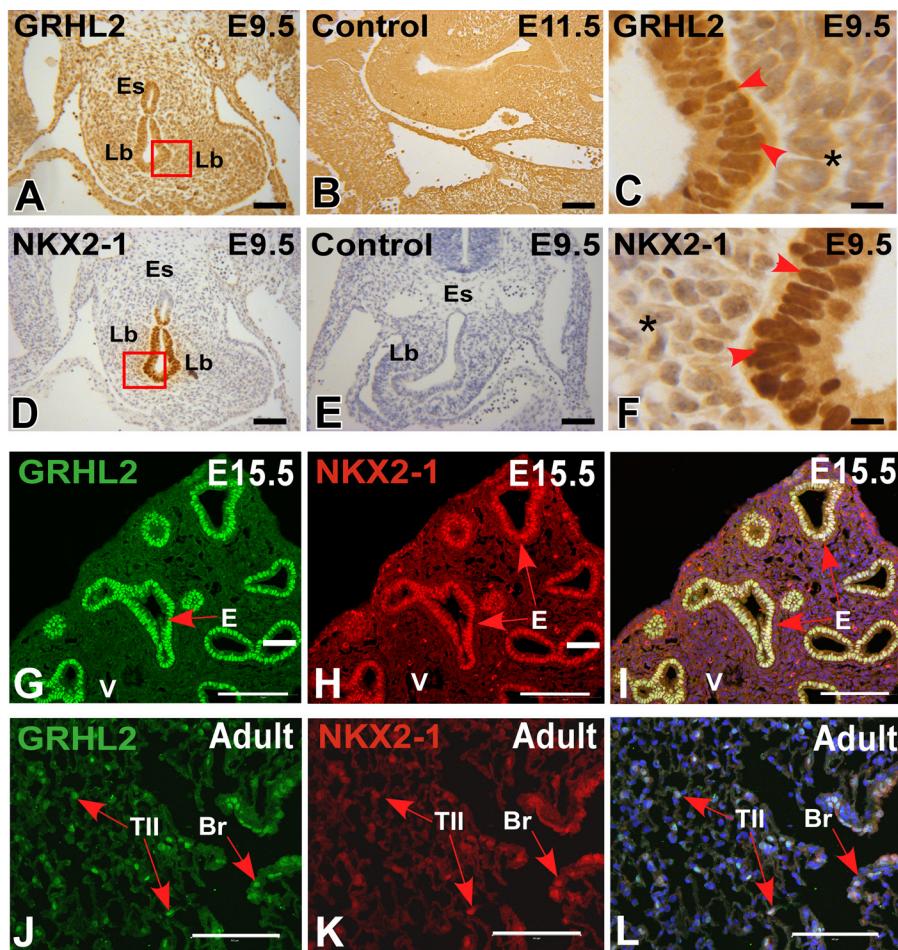


FIGURE 2. Co-localization of GRHL2 and NKX2-1 in mouse lung development. Shown are immunohistochemical analyses of GRHL2 at E9.5 (A) and its corresponding control (B) and GRHL2 expression in the nuclei of lung epithelial cells (C). Immunohistochemical analyses of NKX2-1 protein expression in E9.5 mouse embryos (D) and its corresponding control (E) are shown. F, NKX2-1 expression in the nuclei of lung epithelial cells (F). Immunofluorescence co-localization of NKX2-1 and GRHL2 proteins in day E15.5 lung (G–I) and in adult lung (J–L). Lb, lung buds; Es, esophagus; E, epithelium; V, blood vessel; Br, bronchi/bronchioles; TII, alveolar type II cells; *, negative mesenchymal cell nuclei. Bars A, B, D, and E = 50 μ m, C and F = 5 μ m, and G–L = 100 μ m.

by Dr. Roberto Di Lauro (Stazione Zoologica A. Dohrn, Napoli, Italy). In all experiments renilla luciferase was used to normalize for transfection efficiency. Firefly and renilla luciferase activities were measured after 48 h with the dual luciferase system (Promega).

Gene Array Experiments—All procedures were performed at Boston University Microarray Resource Facility following protocols described in the GeneChip® Whole Transcript (WT) Sense Target Labeling Assay Manual (Affymetrix). Samples were analyzed on the GeneChip Mouse Gene Array 1.0ST (Affymetrix). Data were analyzed using the Robust Multi-Array Analysis (RMA) algorithm (26, 27). Differential expression between experimental and control samples was examined using Welch's *t* test and a false discovery rate (FDR) correction (28). Genes with a *p* \leq 0.02 and an FDR adjusted *p* value $<$ 0.3 were considered to be differentially expressed. Data have been deposited in the GEO repository, accession number GSE 40729.

RESULTS

NKX2-1 and GRHL2 Protein Expression Patterns Overlap in Developing Lung Epithelium—We characterized the expression pattern of GRHL2 protein by immunohistochemistry and immunofluorescence at different stages of lung development.

GRHL2 protein was detected in the lung epithelium at E9.5 and E15.5 and in bronchiolar and alveolar epithelial cells at E18.5 and adult in a pattern similar to the expression pattern of the message RNA (6) and of its target *Cdh1* (Fig. 1, A–I). To evaluate whether GRHL2 protein expressed in the lung epithelium binds *in vivo* to the same targets identified in a kidney cell line (15), we performed ChIP-PCR analyses using E11.5 mouse lungs. GRHL2 binds to the proximal promoter region (-0.1 kb) of *Cldn4* but not to the intronic region ($+2.8$ kb) (Fig. 1J) and to the intronic region ($+7.8$ kb) of the *Cdh1* gene but not to the proximal promoter region (-2.6 kb) (Fig. 1K). We compared the patterns of expression of GRHL2 and the essential lung transcription factor NKX2-1. At E9.5, when mouse lung morphogenesis begins, GRHL2 has a wider field of expression than NKX2-1. Although the expression of both proteins overlaps in the forming lung buds, GRHL2 can be detected in the dorsal and ventral esophagus cells, whereas NKX2-1 is only detected in the ventral cells (Fig. 2, A–F). By immunofluorescence and confocal microscopy, we found considerable overlap in the expression of GRHL2 and NKX2-1 in E15.5 (Fig. 2, G–I) and adult lungs (Fig. 2, J–L). Similar expression patterns were also detected in primary lung epithelial cells. Surfactant protein

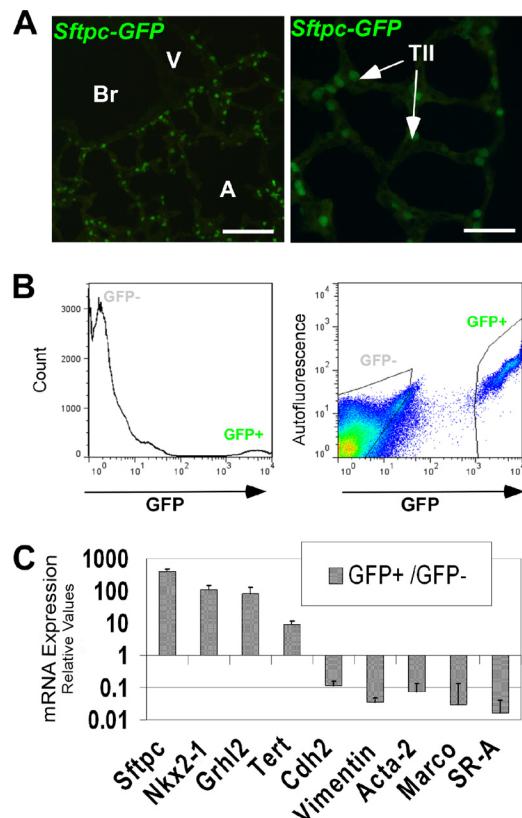


FIGURE 3. **Grhl2 expression in alveolar type II cells.** *A*, fluorescence microscopy analyses show the expression pattern of GFP in the distal lung epithelium of an adult 3.7 kb-SFTPC-Gfp transgenic mouse used for alveolar type II isolation. *Left panel* bar = 50 μ m. *Br*, bronchioles; *A*, alveoli. *Right panel* bar = 25 μ m. *TII*, alveolar type II cells. *B*, cell sorting of GFP-positive cells from 3.7KbSFTPC-Gfp lungs. A histogram shows that ~5% of the cells are GFP-positive (*left panel*); gates used to collect GFP-positive (*GFP+*) and GFP-negative (*GFP-*) cells (*right panel*) are shown. *C*, shown is a gene expression profile of GFP+ cells isolated from 3.7KbSFTPC-Gfp lungs. Relative quantities of different cell-type markers in GFP+ cells compared with GFP- cells are shown. The data sets were normalized against *Gapdh* ($n = 3$). Data represent the mean \pm S.D.

C-Gfp (SFTPC-Gfp) mouse lungs (20) (Fig. 3A) have been used previously to isolate by fluorescence-activated flow cytometry enriched populations of alveolar type II epithelial cells. Using this method, we obtained from total lung cell suspensions ~5% high fluorescence-intensity cells (Fig. 3B). GFP-positive cells have higher expression levels of *Nkx2-1* and *Grhl2* than GFP-negative cells, as determined by qRT-PCR (Fig. 3C). Other markers of epithelial type II cells, such as *Sftpc* and *Cdh1*, were detected at higher levels in GFP-positive cells, whereas levels of markers of alveolar type I cells (*Podoplanin* (*Pdpn*), also known as *T1 α*), mesenchymal cells (*Cadherin2* (*Cdh2*), *Vimentin* (*Vim*), α -smooth muscle actin-2 (*Acta2*)) and of macrophages (macrophage receptor with collagenous structure (*Marco*)) were low.

Grhl2 Regulates Nkx2-1 in Lung Epithelial Cells—To analyze the regulatory network of the *Grhl2* gene, we used two mouse lung epithelial cell lines. MLE15 are cuboidal cells expressing several alveolar epithelial type II cell-specific genes, like *Sftpb* and *Nkx2-1*, whereas E10 are squamous cells expressing alveolar epithelial type I genes like *Pdpn* and *Caveolin1* (*Cav1*). The mRNA expression levels of *Grhl2*, *Cdh1*, *Cldn4*, and *Ocdn* in MLE15 are 2–3 orders of magnitude higher than the levels in

E10 cells (Fig. 4A). Immunostaining results corroborated the high expression level of GRHL2 protein in MLE15 compared with E10 cells (Fig. 4, *B–E*). MLE15 cells express high levels of *Grhl2*, whereas *Grhl1* and *Grhl3* are low or undetected by qRT-PCR (data not shown), making this cell system attractive for studying genes regulated only by *Grhl2*. We infected MLE15 cells with *Grhl2*-shRNA lentivirus. Reduction of *Grhl2* mRNA by >80% (Fig. 4F) results in $69 \pm 12\%$ lower GRHL2 protein levels determined by Western blot and immunofluorescence (Fig. 4, *G–I*). *Grhl2* target genes are also down-regulated (Fig. 4J). *Nkx2-1* mRNA and its downstream target, surfactant protein B (*Sftpb*) (Fig. 5A), were significantly reduced. To determine whether *Nkx2-1* is a direct target of GRHL2 *in vivo*, we performed ChIP with E11.5 lungs (Fig. 5B). A GRHL2 antibody was used to immunoprecipitate chromatin fragments bound by GRHL2. Immunoprecipitated DNA fragments were amplified with specific primers for an *Nkx2-1* regulatory region. A higher signal of the PCR product in GRHL2 over isotype control indicates that GRHL2 binds specifically to the *Nkx2-1* promoter in lung epithelium at E11.5. We also performed EMSAs to further characterize binding of GRHL2 to three regions containing putative GRHL2 binding sites. These sites were selected based on the conserved sequence of Grhl family member binding sites in mammals and other organisms (supplemental Fig. S1). Three regions were tested (Fig. 5C), P1 located -2230 bp from the second ATG (+1) and upstream of the first ATG (at -1842 bp), P2 located -328 bp from the second ATG (+1), and P3. P1 forms a complex with MLE15 nuclear proteins under the conditions tested (Fig. 5D). This complex can be competed by a 100-fold excess of unlabeled oligonucleotide. The addition of GRHL2 antibody but not of IgG interferes with complex formation, indicating specific binding of GRHL2 protein. No DNA-protein complexes were detected with P2 and P3. To determine the transcriptional response of these two *Nkx2-1* promoter regions to GRHL2 overexpression, we co-transfected luciferase constructs containing -0.35 or -2 kb of the *Nkx2-1* 5'-flanking region with a *Grhl2* expression vector (CMV-*Grhl2*-UBC-Gfp) into MLE15 cells (Fig. 5E). A 14-fold increase in the level of luciferase activity with the 2-kb fragment containing the P1 region supports a direct role of GRHL2 in regulation of *Nkx2-1*.

NKX2-1 Regulates Grhl2 in Lung Epithelial Cells—In a previous study we analyzed direct transcriptional targets of NKX2-1 in developing lung by ChiP-on-chip (23). In that analysis we observed that NKX2-1 binds to a DNA region within the first intron of the *Grhl2* gene (Fig. 6A), suggesting a regulatory loop between *Nkx2-1* and *Grhl2*. To test this hypothesis, we investigated whether *Nkx2-1* directly regulates *Grhl2* in lung epithelial cells. *Nkx2-1* mRNA was reduced in MLE15 cells by transduction of *Nkx2-1* shRNA lentivirus (19) (Fig. 6B). Levels of *Grhl2* mRNA expression and of one of its regulated genes, the telomerase reverse transcriptase gene (*Tert*) were reduced by ~50 and ~70%, respectively. Reduction of *Nkx2-1* mRNA results in changes in cell phenotype as cells tend to move apart, reducing the appearance of cell clusters (supplemental Fig. S2). To confirm whether *Grhl2* is a direct target of NKX2-1 *in vivo*, we performed ChIP with E11.5 lungs (Fig. 6C). A NKX2-1 antibody was used to immunoprecipitate chromatin fragments bound by NKX2-1. Immunoprecipitated DNA fragments were

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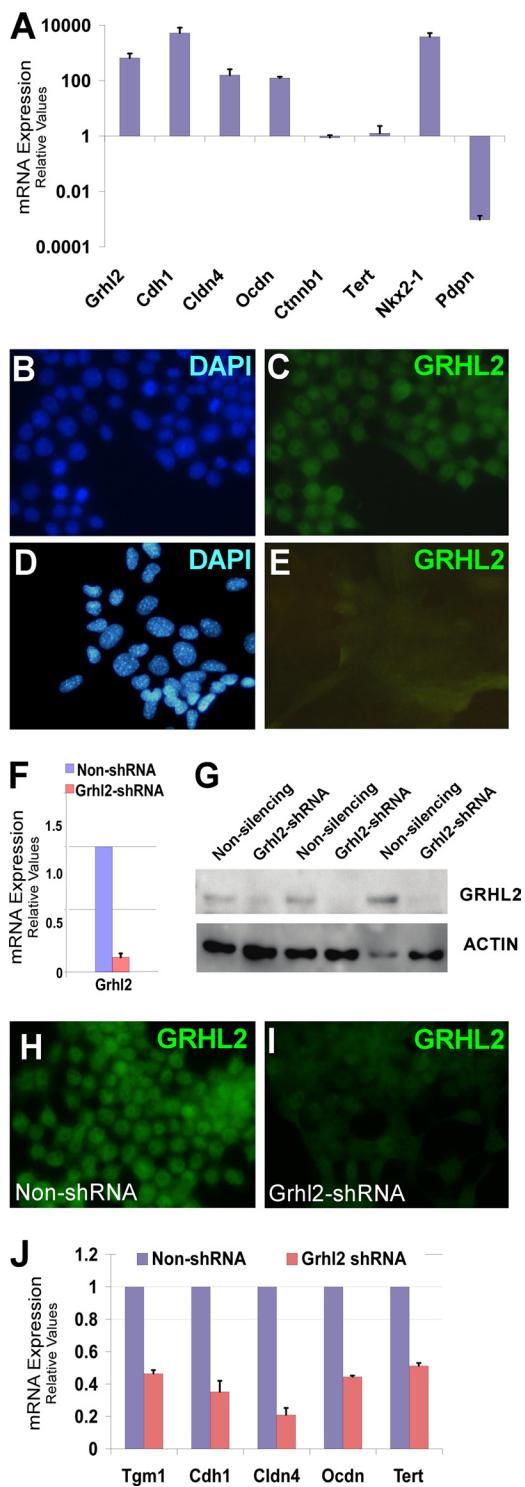


FIGURE 4. Down-regulation of Grhl2 affects expression of its target genes. **A**, qRT-PCR analysis of the expression levels of *Grhl2*, its potential targets, and type I epithelial cell marker *Pdpn* in lung epithelial MLE15 and E10 cells is shown. The data sets were normalized against *Gapdh* ($n = 3$). Data represent the mean \pm S.D. Immunofluorescence analysis of GRHL2 expression in MLE15 cells (**B** and **C**) and in E10 cells (**D** and **E**) is shown. **F**, qRT-PCR expression analysis of *Grhl2* in MLE15 cells infected with *Grhl2*-shRNA V2LMM_20130 (Open Biosystems) or non-silencing-shRNA. Data set was normalized against *Gapdh* ($n = 3$). Data represent the mean \pm S.D. **G**, the Western blot shows down-regulation of GRHL2 protein in MLE15 cells by shRNA in three independent transductions. Data were normalized to β -actin. **H–I**, immunofluorescence analysis of GRHL2 protein expression in MLE15 cells infected with *Grhl2* shRNA or non-silencing control is shown. **J**, by qRT-PCR analysis, we determined that knockdown of *Grhl2* in MLE15 cells lowers

amplified with specific primers spanning *Grhl2* regulatory regions. A higher signal of the PCR product in NKX2-1 over isotype control indicates that NKX2-1 binds specifically to the *Grhl2* intronic region in lung epithelium at E11.5. We also performed EMSAs to further characterize binding of NKX2-1 to three regions of the *Grhl2* intron containing putative NKX2-1 binding sites. These sites were selected based on the conserved core sequence of NKX2-1 (CTTG/CAAG) binding site (supplemental Fig. S1). Three regions were tested, H1 and H2, corresponding to peak H highly bound by NKX2-1, and L1, corresponding to peak L weakly bound by NKX2-1 (Fig. 6, *A* and *D*). Two probes (H1 and L1) form complexes with MLE15 nuclear proteins under the conditions tested (Fig. 6*E*). These complexes can be competed by a 100-fold excess of unlabeled oligonucleotide. The addition of NKX2-1 antibody but not of IgG reduces the mobility of the complexes, super-shifting the bands and indicating specific binding of NKX2-1 protein. No DNA-protein complexes were detected with H2. To determine the transcriptional response of these two *Grhl2* intronic regions to *Nkx2-1* overexpression, we co-transfected SV40 promoter-pGL3 luciferase constructs (SV40Luc) containing fragments H or L of the *Grhl2* intron 5' upstream of the SV40 minimal promoter with an *Nkx2-1* expression vector (pCMV-*Nkx2-1*) into MLE15 cells. A 4-fold increase in the level of luciferase activity with the H fragment supports a direct role of *Nkx2-1* in regulation of *Grhl2* (Fig. 6*F*). Fragment L, although bound by NKX2-1, does not increase the luciferase activity in response to *Nkx2-1* overexpression.

Grhl2 and Its Known Targets Are Down-regulated When Isolated Type II Cells Flatten in Culture—Isolated lung alveolar type II cells cultured on fibronectin-coated dishes are an *ex vivo* model that mimics many of the plasticity features associated with the transition from a type II alveolar epithelial cell phenotype into a type I-like phenotype. These features include changes in cell morphology, from cuboidal to flat, a decrease in expression of well characterized type II markers, and an increase in expression of the type I cell alveolar epithelial markers (19, 29, 30).

GFP-positive cells isolated by cell sorting from SFTPC-Gfp adult mouse lungs were cultured on fibronectin-coated plates for 7 days (Fig. 7*A*). We observed down-regulation of the type II markers *Sftpc* and *Nkx2-1* and up-regulation of the type I cell marker *Pdpn* (Fig. 7*B*). The expression level of *Grhl2* was substantially down-regulated, and its downstream target *Cdh1* was down-regulated 3.2-fold. This suggested either a functional role for *Grhl2* in maintaining the cuboidal cell shape of type II epithelial cells or conversely that a change in cell shape regulates the expression of *Grhl2*.

Grhl2 Regulates Lung Epithelial Cell Shape, Migration, and Expression Levels of Cell-Cell Interaction Genes—We further evaluated whether changes in *Grhl2* levels were a cause or an effect of the alteration in epithelial cell shape using MLE15 and E10 cell lines (19). To determine the effect of *Grhl2* on the regulation of genes involved in cell shape, we used MLE15 cells infected with lentivirus expressing *Grhl2*-shRNA or a non-si-

the expression of its target genes. The data sets were normalized against *Gapdh* ($n = 3$). Data represent the mean \pm S.D.

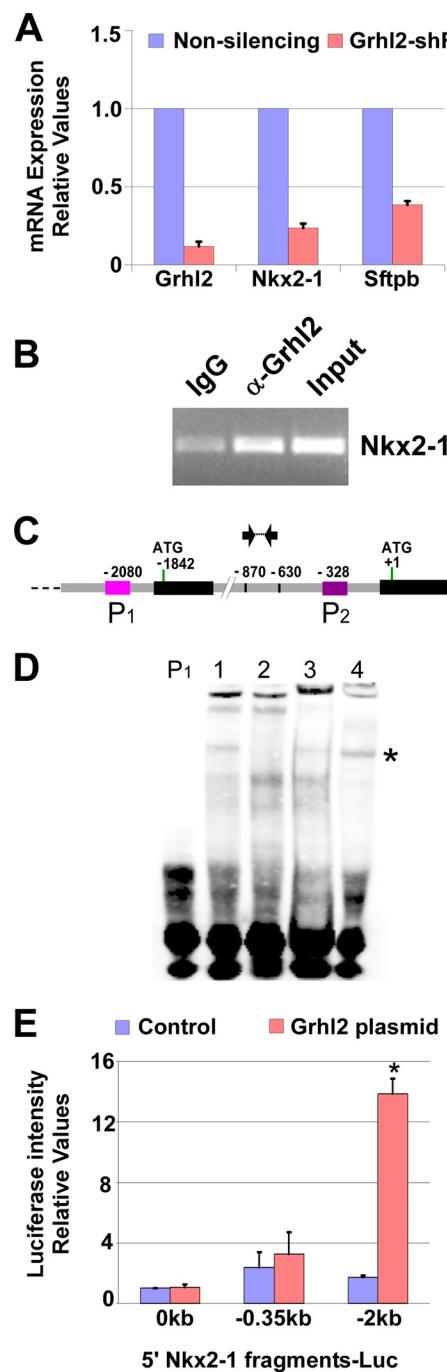


FIGURE 5. **GRHL2 binds to *Nkx2-1* promoter and activates its transcription.** *A*, qRT-PCR expression analysis of *Nkx2-1* and its target *Sftpb* in MLE15 cells infected with *Grhl2*-shRNA or non-silencing-shRNA. Data sets were normalized against *Gapdh* ($n = 3$). Data represent the mean \pm S.D. *B*, ChIP-PCR analysis shows GRHL2 binding to the *Nkx2-1* promoter in the embryonic lung. ChIP was performed with chromatin from E11.5 day lung buds immunoprecipitated with GRHL2-specific antibody or its isotype control. Immunoprecipitated and input DNA fragments were amplified using primers corresponding to *Nkx2-1*. *C*, a scheme of the *Nkx2-1* gene depicting 5' regulatory regions, first and second exons (black boxes), and first intron (gray boxes). Numbers are relative to the second ATG (+1). Pink (P1) and purple boxes (P2) represent probes used in EMSAs. Arrows indicate oligonucleotides used in ChIP-PCR analyses. *D*, EMSA analysis shows binding of MLE15 nuclear proteins to probe P1 (lane 1), competition with 100-fold unlabeled probe (lane 2), interference of complex formation using GRHL2 antibody (lane 3), and control with IgG. The asterisk marks the P1-protein complex. *E*, the *Nkx2-1* promoter is activated in the presence of exogenous GRHL2. MLE15 cells were transfected with either empty luciferase plasmid (pGL3; 0-Luc-plasmid), *Nkx2-1* promoter luciferase plasmids -0.35kbNkx2-1Luc or -2.1KbNkx2-1Luc , and co-

lencing shRNA and investigated the change in cell morphology and in expression patterns of selected cell junction genes. Reduced levels of *Grhl2* relaxed the cuboidal morphology of MLE15 cells into an expanded cell phenotype and reorganize F-actin in the apical and basal sides of the cells (Fig. 8, *A–D*), suggesting that GRHL2 may directly or indirectly regulate cell-cell interaction and cytoskeletal genes or their arrangement. Confocal x-z orthogonal images of phalloidin-stained cells (Fig. 8, *E* and *F*) showed a drastic reorganization of F-actin, leading to relaxation of the cuboidal shape. Conversely, mis-expression of CMV-*Grhl2*-UBC-Gfp in E10 cells (supplemental Fig. S3) changes the squamous and elongated phenotype of E10 cells into a cuboidal phenotype favoring formation of small cell clusters. We tested the effect of reduced *Grhl2* levels on the migratory behavior of MLE15 cell monolayers upon induced injury. Scratch assays of confluent and serum-starved MLE15 cells infected with non-silencing control or *Grhl2*-shRNA showed that lower levels of *Grhl2* alter migration patterns, particularly the collective migration of the epithelial layer, as cells detach from each other and move in smaller clusters (Fig. 9, *A–D*). Scratch healing analysis using Tscratch software indicated that cells with reduced levels of *Grhl2* closed $\sim 30\%$ more of the wound than control cells (Fig. 9*E*) (31). Reduction in the expression of cell junction genes in *Grhl2*-shRNA-infected cells could contribute to the different migration pattern of these cells.

Identification of Downstream Targets of GRHL2 in Lung Epithelial Cells—To identify additional downstream genes regulated by GRHL2 in lung epithelial cells, we performed cDNA microarray analyses in MLE15 cells infected with *Grhl2*-shRNA and compared with a non-silencing control. Among the probe sets represented in the gene expression microarray, we found 286 down-regulated and 218 up-regulated genes in *Grhl2*-shRNA cells compared with control cells ($p < 0.02$; FDR adjusted p value <0.3 , -fold change >1.25). *Grhl2* knockdown leads to down-regulation of genes that could be direct or indirect targets of GRHL2 in lung epithelial cells. To explore the potential biological functions of the down-regulated genes in *Grhl2* knockdown cells, annotations were derived from GO biological processes, and significant annotated functional groups were evaluated using GATHER (32). Among the top functional groups of genes containing more than 10 genes and a Bayes factor >1 were “morphogenesis” and “lipid metabolism,” both processes involved in organ development and in the regulation of cell shape (data not shown).

Among the genes involved in morphogenesis there were several semaphorins, a repulsion-attraction family of genes expressed in lung epithelium (33, 34). This prompted us to further evaluate changes in expression within known families of cell-cell interaction genes. We determined that several members of the semaphorin family and their corresponding receptors as well as Notch and Ephrin genes were reduced when *Grhl2* levels were lower (Table 1). Many of these genes are expressed in the lung epithelium in a pattern similar to *Grhl2* at E14.5 (35).

transfected with CMV-*Grhl2*-UBC-Gfp or CMV-dsred-UBC-Gfp plasmids. Firefly luciferase was normalized to renilla luciferase levels ($n = 2–3$). Data represent the mean \pm S.D. The asterisk indicates $p \leq 0.05$.

Grhl2 and Nkx2-1 Transcriptional Loop

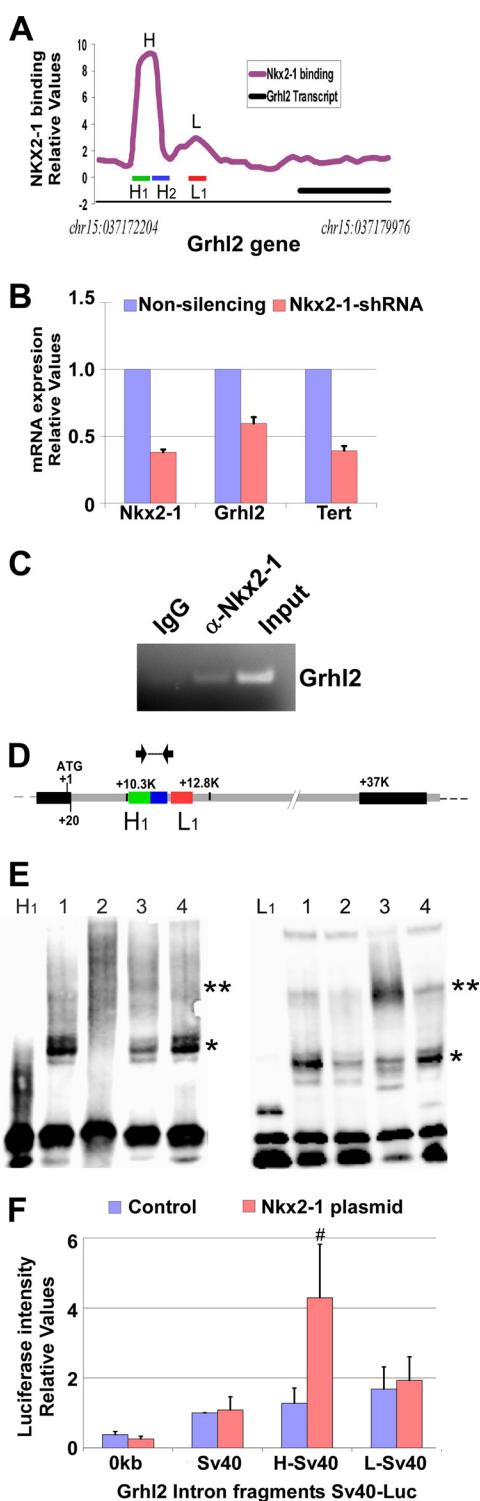


FIGURE 6. NKX2-1 binds to Grhl2 intronic regions and increases transcription. *A*, shown is the binding pattern of NKX2-1 protein to the Grhl2 intron identified in a global ChIP-on-chip analysis of NKX2-1 binding profiles in E11.5 mouse lung epithelium (23). *H* and *L* identify the binding regions; *H1*, *H2*, and *L1* identify the probes used in EMSAs. *B*, shown is qRT-PCR analysis of *Nkx2-1*, *Grhl2*, and *Tert* expression in MLE15 cells infected with *Nkx2-1*-shRNA or non-silencing-shRNA. The data set was normalized against *Gapdh* ($n = 3$). *C*, a representative ChIP-PCR analysis shows NKX2-1 binding to the *Grhl2* intron. ChIP was performed with chromatin from E11.5 day lung bud immunoprecipitated with NKX2-1-specific antibody or its isotype control. Immunoprecipitated and input DNA fragments were amplified using primers corresponding to the *Grhl2* intron. *D*, shown is a scheme of the *Grhl2* gene depicting, first, and second exons (black boxes) and first intron (gray box).

We validated by qRT-PCR arrays (Applied Biosystems) 13 of 20 genes selected from the microarray analysis (Fig. 10). These included the cell-cell interaction genes *Sema3b*, *Sema3c*, and their receptor *Nrp2*, the cell fate specification gene *Notch1*, and the Kruppel-like factor 12 (*Klf12*). *Sema3a*, *Plxnb1*, and *Plxnb2*, although validated by qRT-PCR, are expressed in mesenchymal or subepithelial cells during lung development (34).

DISCUSSION

We studied GRHL2-mediated gene regulation in the lung and explored the molecular mechanisms by which GRHL2 may regulate epithelial characteristics and differentiation of alveolar cells. We present evidence that GRHL2 directly regulates the key lung transcription factor *Nkx2-1*. Notably, *Grhl2* and *Nkx2-1* bind to each other's promoter *in vivo*, forming a positive feedback regulatory loop. Because *Nkx2-1* is critical for distal lung cell differentiation and *Grhl2* regulates cell morphogenesis, we propose that these two factors sustain each other's expression to maintain epithelial cell features during differentiation of distinct lung epithelial cell phenotypes.

Because of the larger expression domain of GRHL2 in the foregut endoderm at E9.5 and the overlapping patterns thereafter, we initially hypothesized that GRHL2 regulates *Nkx2-1* expression. *In vivo* and *in vitro* experiments proved that GRHL2 binds to a particular region of the *Nkx2-1* promoter and activates its transcriptional activity. Similarly, NKX2-1 protein binds to the first intron of the *Grhl2* gene and regulates its expression. Although the identification of the particular *cis*-element in these regulatory regions will require further deletion and mutation analyses, the results presented indicate the presence of a regulatory loop connecting these transcription factors during mouse lung epithelial cell differentiation.

Positive transcriptional feedback loops have been described during pattern formation in development of several organisms such as zebrafish and mouse. These loops restrict transcriptional activity to give rise to different cell types in specific spatial and temporal domains or to reinforce initial cell fate decisions, providing robustness in gene expression to resist variations of external signals (36). The factors involved in any positive feedback loop are not necessarily involved in initiation of their own expression (36). In early *Grhl2* ENU mutant lungs (16), the expression of *Nkx2-1* is not altered, suggesting that the initial transcription does not depend on *Grhl2* alone. Other members of the *Grhl* family present in the foregut endoderm, such as

Numbers are relative to the ATG (+1). Green (*H1*), blue (*H2*), and red boxes (*L1*) represent probes used in EMSAs. Arrows indicate oligonucleotides used in ChIP-PCR analyses. *E*, left panel, EMSA analysis shows binding of MLE15 nuclear proteins to probe *H1* (lane 1), competition with 100-fold unlabeled probe (lane 2), super-shift of the complex using NKX2-1 antibody (lane 3), and control with IgG. The single asterisk marks the *H1*-protein complex. The double asterisk marks the super-shifted complex. Right panel, EMSA analysis shows binding of MLE15 nuclear proteins to probe *L1* (lane 1), competition with 100-fold unlabeled probe (lane 2), super-shift of complex formation using *Nkx2-1* antibody (lane 3), and control with IgG. The single asterisk marks the *L1*-protein complex. The double asterisk marks the super-shifted complex. *F*, MLE15 cells were co-transfected with empty luciferase plasmid (pGL3; 0-Luc-plasmid), Sv40 promoter-luciferase (Sv40Luc), *Grhl2* intron fragment *H* inserted 5' to Sv40-luciferase plasmid (*H-Sv40Luc*), or *Grhl2* intron fragment *L* inserted 5' to Sv40-luciferase plasmid (*L-Sv40Luc*) with CMV-*Nkx2-1* or pCMV plasmids. Firefly luciferase was normalized to Renilla luciferase levels ($n = 5$). Data represent the mean \pm S.E. # indicates $p \leq 0.05$.

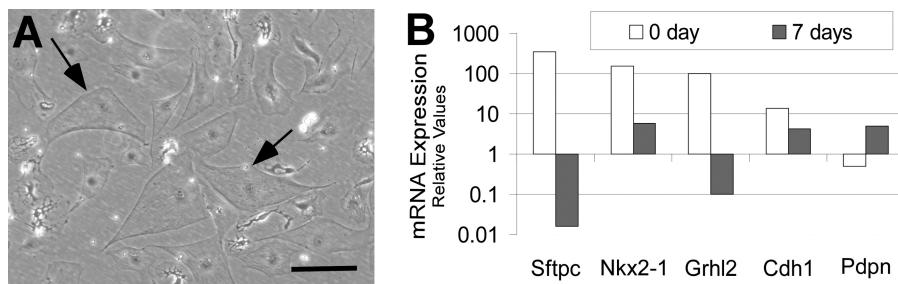


FIGURE 7. **Grhl2 is down-regulated as alveolar type II cells flatten in culture.** *A*, shown are changes in the cell morphology of sorted GFP-positive cells upon culturing on fibronectin. Bar = 10 μ m. *B*, gene expression analyses by qRT-PCR of Grhl2 type I and type II marker genes in freshly isolated versus cultured alveolar type II cells are shown.

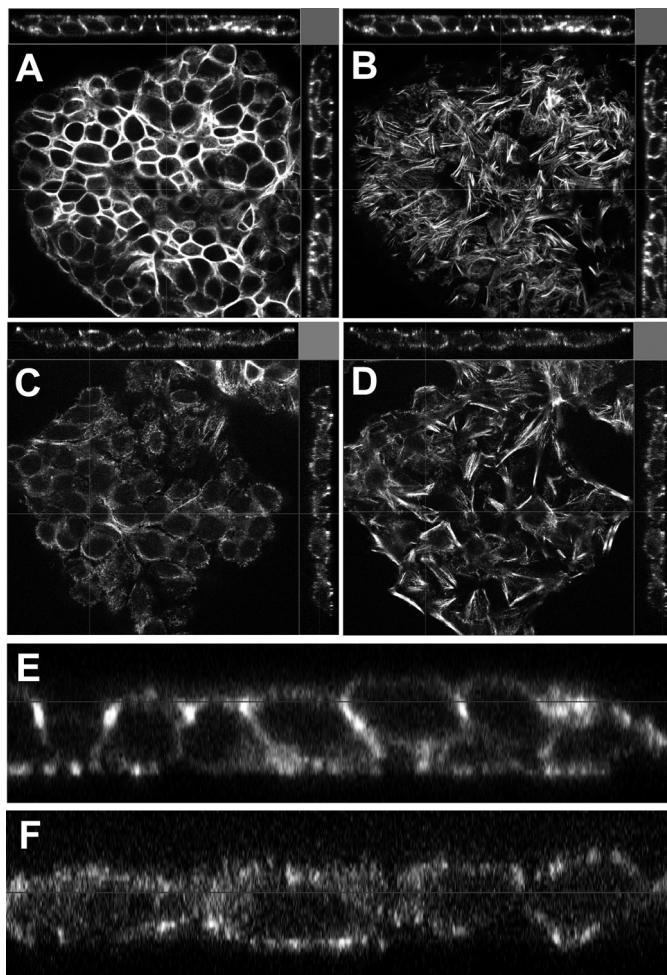


FIGURE 8. **Reduced Grhl2 expression affects morphology of lung epithelial cells.** Confocal orthogonal images of phalloidin stained MLE15 cells transduced with non-silencing shRNA (non-shRNA) apical XY image (*A*), basal XY image (*B*), or with Grhl2-shRNA lentivirus apical XY image (*C*) and basal XY image (*D*). Bar = 20 μ m. Shown is zoom-in of the orthogonal XZ image of MLE15 transduced with non-silencing shRNA (*E*) and of the orthogonal XZ image of MLE15 transduced with Grhl2 shRNA (*F*).

Grhl1 and/or *Grhl3*, or other transcription factors may initiate *Nkx2-1* expression. Later in gestation, when *Grhl2* is the only member of the *Grhl* family expressed at high levels in the distal lung epithelium, the effect of its specific loop may be more evident. The few viable E18.5 *Grhl2* ENU mutant fetuses were reported to have smaller lungs and disorganized cell junctions (16). It is possible that at E18.5, the absence of *Grhl2* may affect expression levels or patterns of *Nkx2-1* and result in altered epi-

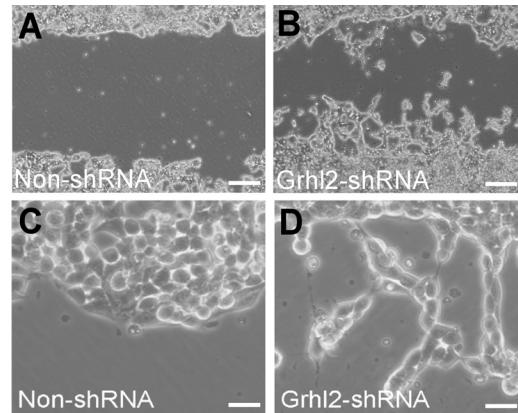


FIGURE 9. **Reduced Grhl2 expression affects migration pattern of lung epithelial cells.** Shown is representative wound healing image of MLE15 cells transduced with non-silencing shRNA (non-shRNA) (*A*) or with Grhl2-shRNA lentivirus 48 h after scratch (*B*). Bar = 50 μ m. Knockdown of Grhl2 alters migration of MLE15 cells in wound healing scratch assays. *C* and *D*, higher magnification images show the difference in migratory pattern with reduced levels of Grhl2. Bar = 10 μ m. *E*, quantification of the scratch healing using Tscratch software ($n = 3$) is shown. Data represent the mean \pm S.E. The asterisk indicates $p \leq 0.05$.

thelial cell fates, although this has not been evaluated. *Nkx2-1*^{-/-} mice, in contrast, have an abnormal lung with rudimentary main bronchi and an absence of peripheral lung structures (37, 38). Despite the importance of *Nkx2-1* in lung morphogenesis, there is limited information concerning the transcriptional regulation of *Nkx2-1* (39–41) or the restriction of its expression to lung bronchiolar or alveolar type II cells. These findings add *Grhl2* as a new factor in the *Nkx2-1* transcriptional network controlling lung development.

The alveolar epithelium includes cuboidal type II and flattened type I cells. These two types of epithelial cells with distinct morphological characteristics and functions lie adjoined to each other in the distal lung to form the alveoli. During lung development, any alteration in the differentiation of type I and type II epithelial cells may cause an aberrant alveolar morphol-

Grhl2 and Nkx2-1 Transcriptional Loop

TABLE 1

Selected cell-cell interaction and other lung genes regulated by Grhl2 in MLE15 cells

($p < 0.02$; FDR adjusted p value <0.3 , fold-change >1.25).

| Symbol | Description | -Fold change | <i>p</i> value | FDR | Lung expression E14.5 | Probeset |
|---------|--|--------------|----------------|------|-----------------------|----------|
| Fabp5 | Fatty acid-binding protein 5 | -3.1 | 0.0014 | 0.19 | | 10585699 |
| Fgb | Fibrinogen β chain | -2.6 | 0.0058 | 0.23 | | 10498981 |
| Hapln1 | Hyaluronan and proteoglycan link protein 1 | -2.3 | 0.0003 | 0.17 | | 10406519 |
| Klf12 | Kruppel-like factor 12 | -2.1 | 0.0083 | 0.25 | | 10422013 |
| Cldn8 | Claudin 8 | -1.9 | 0.0061 | 0.24 | E + | 10440647 |
| Sema3a | Semaphorin 3A | -1.5 | 0.0003 | 0.17 | | 10519717 |
| Sema3c | Semaphorin 3C | -1.4 | 0.0051 | 0.23 | E + + | 10519886 |
| Cldn4 | Claudin 4 | -1.3 | 0.0471 | 0.39 | E + + | 10534395 |
| Efna5 | Ephrin A5 | -1.3 | 0.0008 | 0.17 | | 10452419 |
| Sema6d | Semaphorin 6D | -1.3 | 0.0033 | 0.21 | | 10475544 |
| Notch1 | Notch gene homolog 1 | -1.3 | 0.0124 | 0.27 | E + + | 10481056 |
| Sftpd | Surfactant associated protein D | -1.3 | 0.0114 | 0.27 | E + | 10419096 |
| Nrp2 | Neuropilin 2 | -1.2 | 0.0064 | 0.24 | E + | 10346843 |
| Efna1 | Ephrin A1 | -1.2 | 0.0035 | 0.22 | E + + | 10499536 |
| Plxnb1 | Plexin B1 | -1.2 | 0.0011 | 0.19 | E + + | 10589368 |
| Plxna2 | Plexin A2 | -1.2 | 0.0002 | 0.15 | | 10352867 |
| Grhl2 | Grainyhead-like 2 | -1.2 | 0.0602 | 0.42 | E + + | 10423770 |
| Ephb3 | Eph receptor B3 | -1.2 | 0.0060 | 0.24 | | 10434559 |
| Notch2 | Notch gene homolog 2 | -1.2 | 0.0053 | 0.23 | | 10494595 |
| Plxna4 | Plexin A4 | -1.2 | 0.0047 | 0.23 | | 10543802 |
| Plxna1 | Plexin A1 | -1.2 | 0.0113 | 0.27 | | 10546184 |
| Enpp2 | Ectonucleotide pyrophosphatase/phosphodiesterase 2 | 2.2 | 0.0066 | 0.24 | | 10428619 |
| Scgb3a1 | Secretoglobin, family 3A, member 1 | 2.4 | 0.0018 | 0.19 | E + | 10375608 |
| Hddc2 | HD domain containing 2 | 2.5 | 0.0054 | 0.23 | | 10362394 |
| Bex1 | Brain expressed gene 1 | 2.9 | 0.0005 | 0.17 | | 10606868 |

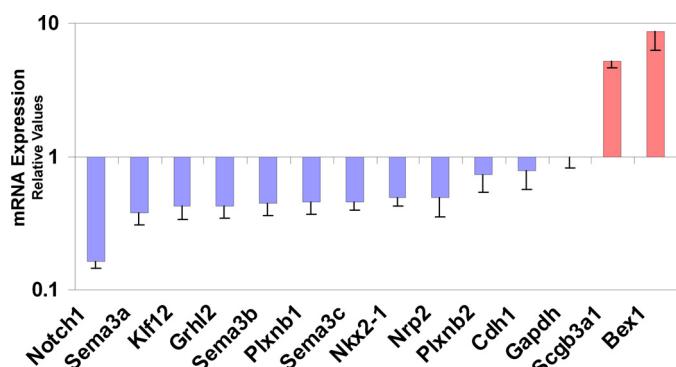


FIGURE 10. Down-regulation of Grhl2 affects the expression of other cell-cell interaction genes. Genes involved in cell-cell interaction identified in the microarray analysis were validated by qRT-PCR. 13 of 20 genes selected were confirmed ($n = 2$). Data represent the mean \pm S.D.

ogy, which in turn may lead to respiratory failure at birth. Similarly, during lung injury, alterations in alveolar structure compromise the respiratory capacity and may lead to disease (42). GRHL2 protein is co-expressed with NKX2-1 in alveolar type II cells. Conversely, flattened cells facing the alveolar lumen that have an elongated nucleus and are negative for NKX2-1-like type I cells- are also negative for GRHL2. Our analysis of GRHL2 function in primary alveolar type II cells and lung cell lines showed that there is a link between GRHL2 expression and cell morphogenesis, migration, and gene expression. Changes in type II alveolar cell morphology, as they flatten in culture, is accompanied by a reduction in *Grhl2* expression. This reduction could be a cause or a consequence of the changes in cell shape. To address this issue, we down-regulated the expression of *Grhl2* in the lung epithelial cell line MLE15. Changes in the levels of expression of cell-cell interaction genes and differences in the pattern of collective cell migration of the epithelial layer indicate that *Grhl2* down-regulation is a cause rather than a consequence of any of these changes. Importantly, changes in cell shape and migration in MLE15 cells reflects

mostly the effect of *Grhl2* in those processes, as these cells do not express detectable levels of *Grhl1* or *Grhl3*. In contrast, in the E18.5 *Grhl2* ENU or null mutant models, the presence of *Grhl1* or *Grhl3* in lung epithelium in early development may compensate for the lack of *Grhl2*, masking the effects of individual members of this family of transcription factors. Nevertheless, the lung phenotype of the few *Grhl2* ENU mutant mice that survived to E18.5 supports the hypothesis that *Grhl2* participates in lung epithelial cell differentiation. Due to the lethality of homozygous *Grhl2* mutations in mice, the ultimate role of *Grhl2* in alveolar sac morphogenesis will need to be evaluated *in vivo* by conditional ablation of *Grhl2* expression in type II cells. These experiments are beyond the scope of the present report but will allow the validation of the functional role of *Grhl2* *in vivo*.

GRHL2 directly regulates transcription of *Cdh1* in lung bud epithelium. NKX2-1 also regulates *Cdh1* expression in a human adenocarcinoma cell line (43), although this regulation could be direct or indirect. Our finding that *Nkx2-1* forms a feed-forward regulatory loop with *Grhl2* supports an indirect link between *Nkx2-1* and *Cdh1* in the lung epithelial layer as it differentiates into distinct cell types to form a functional respiratory tree. Indeed, *Grhl2* and *Nkx2-1* are among the top 25 candidate genes identified in a microarray analysis of genes involved in EMT through regulation of *Cdh1* (44). The maintenance of the regulatory loop between *Grhl2* and *Nkx2-1* would thus be expected to favor an epithelial phenotype by preventing the differentiation of epithelial cells into mesenchymal cells, such as myofibroblasts, in fibrotic lung diseases.

Grhl2 knockdown affects cell migration after scratch-wounding in culture as well as the expression of *Tert* in lung epithelial MLE15 cells. This evidence together with the reported regulation of *RhoGEF19* by GRHL2 (7), the impaired injury/repair process in the cochlea in a family segregating DFNA28, the progressive hearing loss associated with a *GRHL2*

mutation (45), and the down-regulation of *Grhl2* observed in microarray analyses of hyperoxia injured lung (GEO GDS248 hyperoxic lung injury) suggest that *Grhl2* plays an important role in the lung injury/repair process. *Grhls* also regulate proliferation of some human carcinoma cell lines (46). *Tert*, a gene regulated by GRHL2 (47, 48) in humans, is crucial in lung morphogenesis during development (49) as well as in injury/repair in mice and humans (50). These data and the co-localization reported for *Tert* in type II cells lead us to infer that GRHL2 may also regulate *Tert* in mouse lung alveolar cells. Genetic mutations in hTERT within human families predispose them to idiopathic pulmonary fibrosis (51–53), pointing to a potential role of GRHL2 in lung fibrosis through regulation of *Tert* and possibly other lung epithelial genes.

Microarray analyses revealed novel genes regulated by GRHL2 in lung epithelial cells. Expression of several members of the semaphorin family and their corresponding receptors were reduced when *Grhl2* levels were down-regulated. Although the regulation of these genes by *Grhl2* could be direct or indirect, the change in expression of attraction-repulsion genes further supports the role of *Grhl2* in balancing collective epithelial cell movements and differentiation. As *Grhl2*, semaphorins are also important for the development of different organs, neural cell migration (16, 54), angiogenesis, tumorigenesis, and the immune response (34, 55). Previous analysis of the role of semaphorins in lung development suggested that semaphorins may control branching morphogenesis and promote epithelial proliferation (33, 34). Therefore, we propose that through regulation of these genes, *Grhl2* contributes to lung organogenesis.

Notably, we also showed that *Grhl2* regulates expression of *Klf12*, a Kruppel-like transcription factor that directly represses expression of the helix-loop-helix transcription factor *AP-2 α* (56). *AP-2 α* , in turn, directly regulates expression of *Cdh1* (57). Although both GRHL2 and *AP-2 α* are positive regulators of *Cdh1*, experiments in *Grhl2* or *AP-2 α* mutant embryos indicated that these factors do not regulate each other's expression in neural tissues (16). Our findings suggest that *Grhl2* can directly activate *Cdh1* expression or inhibit *Cdh1* expression by activating *Klf12*; these are potential mechanisms for fine-tuning *Cdh1* expression levels to allow simultaneous epithelial layer migration and cell differentiation. Down-regulation of *Grhl2* in MLE15 cells also caused reduced expression of genes such as *Nkx2-1*, *Sftpb*, and surfactant protein D (*Sftpd*) and increased the expression of a tracheal/bronchial gene secretoglobin 3a1 (*Sctgb3a1*) (58), suggesting a role for GRHL2 in proximal-distal cell differentiation in lung development. The early lethality of *Grhl2* null mutant embryos limits further *in vivo* lung analyses.

The *Grhl2-Nkx2-1*-positive feedback loop, the cell-cell interaction genes regulated by GRHL2, and the *Nkx2-1* direct targets that we recently identified in lung development (23) expand the knowledge about the regulatory networks in distal lung epithelium. The *Grhl2-Nkx2-1* loop may be conserved in lung development and disease similarly to the conservation of *Nkx2-1* regulatory patterns in development and tumors (23). *Nkx2-1* haploinsufficiency in humans results in abnormal airway and alveolar morphogenesis (59). In lung tumors, *Nkx2-1* controls

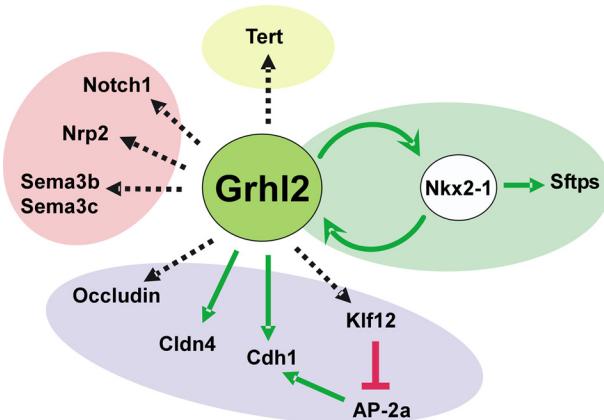


FIGURE 11. Summary of genes regulated by GRHL2 identified in these studies. GRHL2 regulates lung cell differentiation genes (*Nkx2-1* feed forward loop, green oval), cell junction genes (purple oval), and cell-cell interaction genes (pink oval) as well as *Tert*. Solid green or red lines represent direct regulation. Dotted black arrows represent direct or indirect regulation.

cell differentiation and limits metastatic potential (60, 61). It is possible that through the *Grhl2-Nkx2-1*-positive feedback loop, *Nkx2-1* regulates *Cdh1* and other cell-cell interaction genes controlling cell metastatic potential.

In summary, we identified novel cell-cell interaction genes regulated by *Grhl2* that are known to participate in lung morphogenesis and cell differentiation. Most importantly, we identified a positive transcriptional feedback loop between *Grhl2* and *Nkx2-1* that mutually and positively affects their expression levels (Fig. 11). We, therefore, propose that the *Grhl2-Nkx2-1*-positive feedback loop serves to reinforce distal lung epithelial gene expression, epithelial layer integrity, and maintenance of particular epithelial cell fates.

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The Transcription Factors Grainyhead-like 2 and NK2-Homeobox 1 Form a Regulatory Loop That Coordinates Lung Epithelial Cell Morphogenesis and Differentiation

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