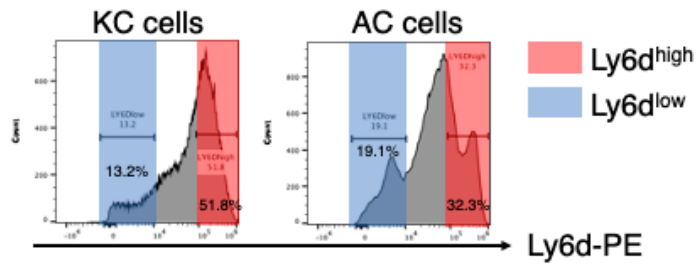
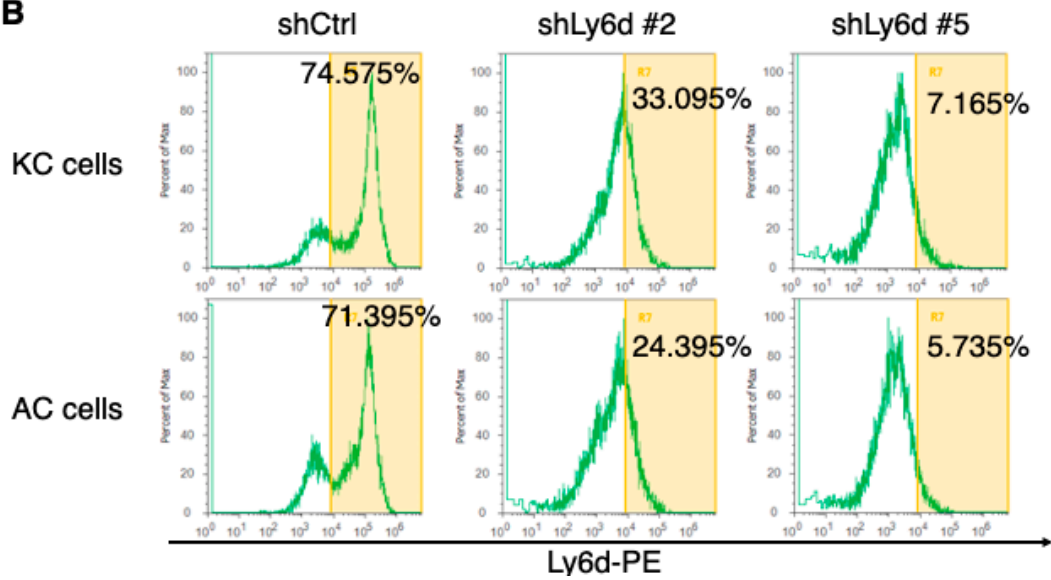


Supplementary Figure S1. The expressions of KRAS, ALK, and GFP. (A) Immunoblot analysis of KRAS, ALK, and GFP in cultured *Cdkn2a*^{-/-} lung cells, KC, and AC cells. β -Actin was examined as a loading control. (B - E) The whole immunoblotting image for KRAS (B), ALK (C), GFP (D), and β -Actin (E).

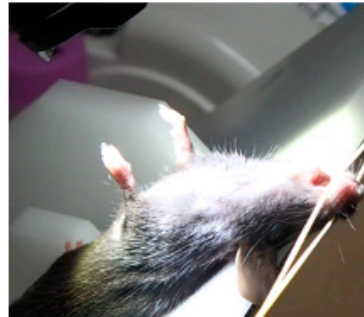
A**B**

Supplementary Figure S2. Ly6d expression in KC and AC cells. (A) Gating strategy for isolation of Ly6d^{high} and Ly6d^{low} cells from KC and AC cells by FACS. (B) Validation of shRNA-mediated knockdown of Ly6d expression in KC and AC cells by flow cytometry.

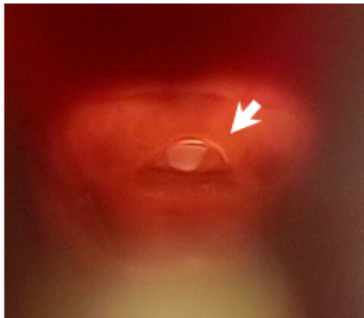
1. Place anesthetized mouse on a tilted platform



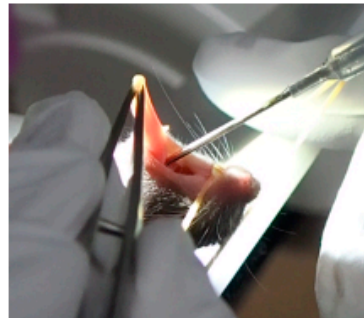
2. Direct a light toward the upper chest of the mouse



3. Visualize the glottis of the mouse (arrow)



4. Insert a cannula into the airway of the mouse



Supplementary Figure S3. Procedure for intratracheal injection.

Supplementary Table S1. List of PCR primer sequences. Fw, forward; Rv, reverse.

Gene symbol		Sequence (5' → 3')
<i>Psca</i>	Fw	TCATCTGTGCTGTGCATGAAT
	Rv	GCTCACTGCAACCATGAAGA
<i>Slurp1</i>	Fw	GTCACCATGGGACTGTGGTT
	Rv	TTGCTCAGTGCAAGATGGAA
<i>Lypd2</i>	Fw	GCATCCAACCTGTGTCACCAC
	Rv	GTCAGAATTGCAGCAGGACA
<i>Slurp2</i>	Fw	TCCACCCACTGTGTCATCAT
	Rv	GGAGTGTGGTCAGTCCCTGT
<i>Lynx1</i>	Fw	TGCCTGAGCTCTTGGTCTCT
	Rv	TACCAACACCGCACGAAGT
<i>Ly6d</i>	Fw	AGACAGCTCTGCTCGTCCTC
	Rv	CTGTAGTCGGAGGTGCATGA
<i>Ly6k</i>	Fw	CAGGCATAGCTGTTCTGCG
	Rv	CCTAGTGGCCTTGCTCGTAG
<i>Gml</i>	Fw	TCTTGTGCTGTGGGAGTGTC
	Rv	CTGGAGAGGATCAAGGCAAG
<i>Ly6e</i>	Fw	ACAGCACAGGCAGGAAGACT
	Rv	TCGGTATTATCTTCGGGGC
<i>Ly6a</i>	Fw	GGCAGATGGGTAAGCAAAGA
	Rv	CAATTACCTGCCCCTACCCT
<i>Ly6h</i>	Fw	AATTGGCCAGGGTGCACT
	Rv	ATGAAGAGCCTCGGTCTGG
<i>Gpihbp1</i>	Fw	CTGGAGCAGCTCTGTGTCTG
	Rv	ACCAACATGATCCCTGGAAG
<i>Gapdh</i>	Fw	GTGAAGGTCGGTGTGAACG
	Rv	GACCATGTAGTTGAGGTCAATG

Supplementary Materials and Methods

Immunoblot analysis

Cultured lung cells, KC and AC cells were collected by digesting Matrigel using Cell Dissociation Solution (Corning, Corning, NY, USA). Cells were lysed with SDS sample buffer (2% SDS, 10% glycerol, 50 mM Tris-HCl [pH 6.8]) containing 1 M DTT for fractionation by SDS-PAGE. The separated proteins were transferred to a polyvinylidene difluoride membrane and exposed to anti-KRAS antibody (C-19; Santa Cruz Biotechnology, Dallas, TX, USA), anti-ALK antibody (D5F3; Cell Signaling Technology, Danvers, MA, USA), anti-GFP antibody (FL, Santa Cruz Biotechnology), and anti- β -Actin antibody (sc-47778, Santa Cruz Biotechnology). Immune complexes were detected with horseradish peroxidase-conjugated anti-rabbit antibody (Agilent, Santa Clara, CA, USA) or anti-mouse antibody (Agilent) and enhanced chemiluminescence reagents (Chemi-Lumi One L, Nacalai Tesque, Kyoto, Japan).