Cancer is a multistep process that involves mutations and other alterations in oncogenes and tumour suppressor genes. Genome sequencing studies have identified a large collection of genetic alterations that occur in human cancers. However, the determination of which mutations are causally related to tumorigenesis remains a major challenge. Here we describe a novel CRISPR/Cas9-based approach for rapid functional investigation of candidate genes in well-established autochthonous mouse models of cancer. Using a Kras\(^{G12D}\)-driven lung cancer model, we performed functional characterization of a panel of tumour suppressor genes with known loss-of-function alterations in human lung cancer. Cre-dependent somatic activation of oncogenic Kras\(^{G12D}\) combined with CRISPR/Cas9-mediated genome editing of tumour suppressor genes resulted in lung adenocarcinomas with distinct histopathological and molecular features. This rapid somatic genome engineering approach enables functional characterization of putative cancer genes in the lung and other tissues using autochthonous mouse models. We anticipate that this approach can be used to systematically dissect the complex catalogue of mutations identified in cancer genome sequencing studies.

Lung cancer genome sequencing studies have revealed a multitude of recurrent mutations and copy number alterations. However, the determination of which mutations are causally related to tumorigenesis remains a major challenge. Genetically engineered mouse models of lung cancer have assisted in the functional characterization of putative driver events identified in human lung tumours, but these require modification of the germ line and cannot be performed in a highly parallel manner.

Recent work from our laboratory has demonstrated the feasibility of using the CRISPR clustered regularly interspaced short palindromic repeats/Cas9 system to directly mutate cancer genes in the liver following hydrodynamic delivery of plasmids carrying the CRISPR components, which relies on the efficient transfection of hepatocytes. To rapidly interrogate cancer genes in the lung and other tissues, we developed pSECC (Fig. 1a), a lentiviral-based system that delivers both the CRISPR system and Cre recombinase. In this setting, CRISPR-induced mutation of genes can be examined in the context of several of the well-studied conditional Cre/loxP-based mouse models of lung cancer and other cancer types. To test this system, we used genetically engineered mouse models of lung adenocarcinoma, in which tumours are induced in loxP-Stop-loxP Kras\(^{G12D/+}\) (hereafter referred to as Kras\(^{LSL-G12D/+}\)) or Kras\(^{LSL-G12D/+}\); p53\(^{-/-}\) mice upon intratracheal administration of lentiviral vectors expressing Cre recombinase.

To validate pSECC, we developed the Green-Go (GG) reporter cell line, which expresses GFP following exposure to Cre (Extended Data Fig. 1a–c). To assess the efficiency of Cas9 in tumours in vivo, we targeted a Cre-activatable tdTomato knock-in reporter allele with pSECC lentiviruses expressing a single guide RNA (sgRNA) against tdTomato (sgTom) or an empty vector control (Extended Data Fig. 1d, e). At 10 weeks post-infection, we assessed knockdown of tdTomato expression by immunohistochemistry. We observed that 28% of tumours lacked tdTomato expression, suggesting that the system was functional in vivo by editing an endogenous allele in the context of a lung tumour (Extended Data Fig. 2a–e). Importantly, animals infected with empty pSECC rarely contained non-tumour-Tomato-expressing cells (data not shown), indicating that there is minimal infection of non-epithelial cells when using a low lentiviral titre.

We then proceeded to functionally characterize tumour suppressor genes using this approach. Loss of Nkx2 homeobox 1 (Nkx2-1), a master regulator of lung development, or phosphatase and tensin homologue (Pten), a negative regulator of oncogenic PI(3)K/Akt signalling, accelerates lung tumorigenesis in Kras\(^{LSL-G12D/+}\); p53\(^{-/-}\) lung tumour models. We infected Kras\(^{LSL-G12D/+}\); and Kras\(^{LSL-G12D/+}\); p53\(^{-/-}\) animals with pSECC vectors expressing validated sgPten, sgNkx2-1 and controls (sgTom and empty vector) to induce lung tumours. Ten weeks post-infection, we euthanized animals to assess the effects of CRISPR/Cas9-mediated gene editing in tumours by histopathology, surrogate assays and deep sequencing of the targeted alleles (Fig. 1a). All animals expressing sgRNAs targeting Pten or Nkx2-1 contained tumours with marked histopathological differences compared to controls (Fig. 1b, d and Extended Data Fig. 3a–d).

Animals infected with sgNkx2-1-pSECC developed mucinous adenocarcinomas typified by the presence of elongated cells, mucin production and glandular rearrangements, in agreement with previous Cre/loxP-based (Nkx2-1\(^{fl/fl}\)) data (Fig. 1b). The majority of tumours (61%, 54/88 tumours) from sgNkx2-1-pSECC animals lacked Nkx2-1 expression (compared to 0/33 tumours from controls) (Fig. 1b, c). Importantly, 85% (46/54 tumours) of these Nkx2-1-negative tumours stained positively for mucin (Fig. 1c), a biomarker of mucinous adenocarcinomas. Thus, although a subset of tumours appeared to partially or fully escape CRISPR-mediated deletion of Nkx2-1, we were able to observe clear phenotypes by examining the full spectrum of tumours generated by sgNkx2-1-pSECC.

Animals infected with sgPten-pSECC demonstrated complete loss of Pten protein in 74% of tumours (40/54 tumours), which was accompanied by a concomitant increase in pAkt (S473), a downstream biomarker of increased PI(3)-kinase pathway activity (Fig. 1d, e). These results mimic previously published data using a Pten\(^{fl/fl}\) allele in Kras\(^{LSL-G12D/+}\); p53\(^{-/-}\) mice. Collectively, these data indicate that CRISPR/Cas9-based gene editing leads to loss-of-function mutations in this model and closely parallels what is seen with the use of traditional conditional alleles.

We next used this system to study adenosomatous polyposis coli (Apc), a tumour suppressor whose functional role in lung adenocarcinoma has not been characterized. Of note, Apc is found in a region that frequently undergoes copy number loss in human lung cancer. We infected animals with pSECC lentiviruses expressing a validated sgRNA targeting Apc.
Apc. At 10 weeks post-infection, we observed a striking difference in the histopathology of sgApc tumours compared to controls (Fig. 1f and Extended Data Fig. 3e). Importantly, tumours from Kras\(^{LSL-G12D^-}\);A\(^{fl/fl}\) mice, which express a conditional allele of Apc\(^{18}\), exhibited identical histopathology (Fig. 1f). Tumours with Cas9-mediated deletion of Apc were highly dedifferentiated, invasive and had a significant stromal component (Fig. 1f). The majority of these tumours (78%, 91/117) stained strongly for nuclear β-catenin, a marker of Apc mutation in colon cancer and other settings\(^{19}\) (Fig. 1f, g). Furthermore, 77% (70/91) of tumours with nuclear β-catenin stained positive for the transcription factor Sox9, which might reflect a distal embryonic differentiation state\(^{20,21}\). Of note, we observed a statistically significantly higher number of Sox9-positive tumours in Kras\(^{LSL-G12D^-}\);p53\(^{ΔΔ}\);sgApc (29/33, or 88%) than in Kras\(^{LSL-G12D^-}\)-sgApc mice (41/58 tumours, or 71%), suggesting a possible role for p53 in regulating this change in differentiation (Extended Data Fig. 6b, c).

To further characterize the differentiation state of sgApc tumours, we stained serial sections for lung differentiation markers, including Sox2, Clara cell secretory protein (CCSP), surfactant protein C (SP-C), p63, Nkx2-1 and Sox9 (Extended Data Fig. 6a)\(^{22}\). Tumours from Kras\(^{LSL-G12D^-}\);p53\(^{ΔΔ}\)-sgTom mice stained positively for CCSP, SP-C and Nkx2-1 and negatively for Sox2, p63 and Sox9. In contrast, tumours

**Figure 1** CRISPR/Cas9-mediated somatic gene editing in an autochthonous mouse model of lung cancer. a, pSECC lentiviruses are intratracheally delivered into mouse lungs to delete genes of interest. DNA extracted from tumour-bearing lungs is analysed by high-throughput sequencing and surveyor assays to identify gene-editing events. The remaining tissue is analysed by histopathology. b, Representative haematoxylin and eosin (H&E) and immunohistochemistry (IHC) staining of serial sections from lung tumours of mice 10 weeks after infection with sgTom-pSECC (left panel) or sgNkx2-1-pSECC (right panel). Alcian Blue/PAS (periodic acid-Schiff) stain for mucin. Note the accumulation of mucin only in tumours from sgNkx2-1-pSECC mice. c, Contingency tables demonstrating anti-correlation between Nkx2-1 expression and mucin production (PAS stain) (two-sided Fisher’s exact test, \(P < 0.0001\)). d, Representative H&E and IHC stainings of serial sections from lung tumours of mice 10 weeks after infection with sgTom-pSECC (left panel) or sgPten-pSECC (right panel). Dashed lines demarcate tumour boundaries on each consecutive histological section. e, Contingency tables demonstrating anti-correlation between Pten expression and Akt phosphorylation (two-sided Fisher’s exact test, \(P < 0.0001\)). f, Representative H&E and IHC stainings of serial sections from lung tumours of mice 10 weeks after infection with sgTom-pSECC (left panel) or sgApca-pSECC (middle panel). The far right panel corresponds to serial sections from lung tumours of Kras\(^{LSL-G12D^-}\)\(^{ΔΔ}\);A\(^{fl/fl}\) mice 18 weeks after infection with Adeno-Cre. g, Contingency tables demonstrating positive correlation between β-catenin expression and Sox9 expression (two-sided Fisher’s exact test, \(P < 0.0001\)). These data are representative of at least 3 independent Kras\(^{LSL-G12D^-}\)\(^{ΔΔ}\) or Kras\(^{LSL-G12D^-}\)\(^{ΔΔ}\);p53\(^{ΔΔ}\) mice infected with each pSECC sgRNA. All scale bars, 0.05 mm.
from sgApc mice frequently stained positively for SP-C, Nkx2-1 and Sox9 and negatively for CGSP, Sox2 and p63. A large number of tumours from sgApc mice had areas with low levels or complete absence of Nkx2-1, which correlated with the levels of the Nkx2-1 transcriptional target SP-C \( \text{P}^2 \) (16/52 tumours, or 31\%) (Extended Data Fig. 6d). These data indicate that these tumours are poorly differentiated and that hyperactivation of the canonical Wnt signalling pathway through loss of Apc in Kras-driven lung adenocarcinomas results in tumours with varying degrees of differentiation. These results also mimic what we observed in tumours from Apc conditional knockout mice (Fig. 1f and Extended Data Fig. 6e) and recapitulate recent observations in a Braf\( ^{V600E} \)-driven mouse model of lung adenocarcinoma upon Wnt pathway hyperactivation \( ^{23} \).

Our initial analysis demonstrated histological and pathway-specific differences upon deletion of these tumour suppressors in lung tumours. To assess the overall impact of these alterations on tumorigenesis, we measured tumour burden and grade in both Kras\(^{LSL-G12D/+}\) and Kras\(^{LSL-G12D/+}\); p53\(^{fl/fl}\) animals. Deletion of Pten and Apc significantly increased overall tumour burden, which correlated with higher tumour grades (Grade 3 and 4) (Fig. 2a–c and Extended Data Fig. 3f, g). Nkx2-1 deletion had a significant effect on overall tumour burden only in Kras\(^{LSL-G12D/+}\); p53\(^{fl/fl}\) animals; however, we observed a striking transition to highly dedifferentiated mucinous adenocarcinomas in both Kras\(^{LSL-G12D/+}\) and Kras\(^{LSL-G12D/+}\);p53\(^{fl/fl}\) mice (Fig. 2a–c, Extended Data Fig. 3f, g and Extended Data Fig. 4a–d). Conversely, Apc deletion had a significant effect on tumour burden only in Kras\(^{LSL-G12D/+}\);p53\(^{fl/fl}\) mice (Fig. 2a and Extended Data Fig. 3f, g). Deletion of all three genes led to increased BrdU incorporation, suggesting that the increased tumour burden is partly due to increased proliferation (Extended Data Fig. 3h). These data demonstrate the tumour suppressive role of Nkx2-1, Pten and Apc in the context of oncogenic Kras. Furthermore, the unique histopathology observed for each targeted tumour suppressor gene in this Kras-driven model illustrates the potential of this approach to rapidly model the effects of cooperative genetic events in lung tumorigenesis and progression.

Using this \textit{in vivo} somatic genome editing approach, we observed inter- and intra-tumoural heterogeneity in terms of CRISPR-based loss-of-function of \textit{Pten} in sgPten animals (Fig. 2d, e and Extended Data Fig. 5). Clones that acquired loss of \textit{Pten} had increased PI(3)K/Akt signalling and may, therefore, have had a selective advantage over tumours that retained wild-type \textit{Pten} within the same animal. We observed that tumours with complete or sub-clonal loss of \textit{Pten} were significantly larger than tumours that retained \textit{Pten} (Fig. 2d, e).

The histopathological and immunohistochemistry analyses indicate that the pSECC system is highly efficient \textit{in vivo}, leading to robust target-specific phenotypic differences in lung tumours. To confirm Cas9-mediated editing of the alleles and precisely characterize the events at single-nucleotide resolution, we performed deep sequencing of target samples, using the control samples as a background model to analyse the mutational rate revealed that sgTarget samples were enriched for mutations within 7 bp upstream of the PAM sequences in predicted cutting sites, strongly suggesting that they are not secondary consequences of tumour progression (Fig. 3d and Extended Data Fig. 7a–c). The maximum per-base mutation frequency observed in sgTarget samples was 71.7\% in Nkx2-1, 66.06\% in \textit{Pten} and 39.91\% in \textit{Apc} (in contrast to control samples: 0.11\%, 0.73\% and 0.14\%, respectively). On average, 27.48\% \( \pm 10.3 \) (Nkx2-1), 44.64\% \( \pm 5.3 \) (\textit{Pten}) and 13.54\% \( \pm 5.3 \) (\textit{Apc}) read fragments covering this 7 bp locus harboured indels in sgTarget samples. Across all sgTarget samples, > 94\% of observed indels constituted non-synonymous frame-altering events (Extended Data Fig. 7d, e and Supplementary Tables 3 and 4).

Several studies have reported that Cas9 can bind to sites in the genome other than the intended target site \(^{4–5}\), which could result in unintended editing at an off-target (OT) site. To assess off-target editing, we analysed the three predicted \(^{24}\) loci (Supplementary Tables 2) for each sgRNA by deep sequencing. We observed negligible off-target editing (Extended Data Fig. 8). On average, 0.048% \( \pm 0.031 \) (for sgNkx2-1), 0.26% \( \pm 0.096 \) (for sgPten) and 0.051% \( \pm 0.027 \) (for sgApc) of read fragments harboured indels in the off-target sites (Supplementary Tables 5–7). This data suggests that the observations reported for each of the sgRNAs arise from deletion of the intended target and not from editing of another gene.

The goal of cancer genomics is to identify genetic events that underlie cancer initiation and progression. The functional interrogation of
putative cancer genes in appropriate experimental models will elucidate which mutations identify bona fide cancer genes. This study presents a novel approach to rapidly evaluate human cancer genome candidates and assess cooperativity between genetic events in the context of well-established mouse models of lung cancer. Moreover, our ability to model different lung adenocarcinoma subtypes allows for the detailed study of subtype-specific molecular mechanisms controlling disease initiation and progression. We anticipate that this approach can be readily adapted to many existing Cre/loxP-based genetically engineered mouse models of several cancer types to facilitate the rapid functional assessment of new hypotheses generated by cancer genome studies.

Online Content Methods, along with any additional Extended Data display items and Source Data, are available in the online version of the paper; references unique to these sections appear only in the online paper.

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Author Contributions F.J.S.-R., T.P. and T.J. designed the study; F.J.S.-R., T.P., R.R., M.R.B. and L.S. performed experiments; T.T. generated K48-Δ1120/Δ1120; A.B. conducted bioinformatic analyses; N.S.J. generated GG cells; R.T.B. provided pathology and L.S. performed experiments; T.T. generated CRISPR-mediated direct mutation of cancer genes in the mouse liver. W.X. wrote the manuscript with comments from all authors.

Author Information Illumina MiSeq sequence datasets have been deposited into the NCBI repository under BioProjectID PRJNA256245. Reprints and permissions information is available at www.nature.com/reprints. The authors declare no competing financial interests. Readers are welcome to comment on the online version of the paper. Correspondence and requests for materials should be addressed to T.J. (tjacks@mit.edu).
LETTER

Deep sequencing and bioinformatic analysis of Cas9 target loci. For each target gene or potential off-target site, a genomic region containing the target sequence was amplified using Herculase II Fusion DNA polymerase and gel purified (primer sequences are shown in Supplementary Table 1). Sequencing libraries were prepared from 50 ng of PCR product using the Nextera DNA Sample Preparation Kit (Illumina) and sequenced on Illumina MiSeq machines. In order to retain high-quality sequence for mutation analysis, Illumina MiSeq reads (150 bp paired-end) were trimmed to 100mer paired end reads to drop lower quality 3’ ends of reads. Traces of Nextera adapters were clipped from PE1 and PE2 100mer reads using the FASTX toolkit (Hannon Lab, CSHL). Reads greater than 15 nucleotides in length were retained. Additionally, reads with 50% or more bases below a base quality threshold of Q30 were dropped from subsequent analysis. Reference sequences with 10 bp genomic flanks were indexed using the Burrows–Wheeler Aligner (BWA) IS linear time algorithm and reads were aligned using the BWA aligner. Reads with mapping quality greater than zero were retained. Overlapping alignments of paired end reads due to short inserts were resolved in order to avoid double counting of coverage and/or mutations observed in a single fragment. In order to minimize alignment ambiguity in the presence of mutations (including indels), the GATK Toolkit was used to realign pooled cohorts mapping to a given locus. Mutations (base substitutions, insertions and deletions) were assessed using a combination of Samtools and Annovar (indel quantification and annotation), NGSUtils/BAMutils software suite (total mutations per position), and custom scripts. Mutation frequencies were adjusted for sample purity (see next section) and per base substitution, insertion, and deletion frequencies were determined. Significance of overall mutation rates across 10 bp flanking the target locus was assessed using the Wilcoxon rank sum test comparing control and sgTarget sample events. Positional enrichment for mutation frequency compared to control samples was assessed using the custom simulation of minimum likelihood (Smallsim) algorithm and was mean centred and scaled (z-scores) across a 10 bp flank on either side of the PAM sequence in each sample. A number of other utilities/tools were used to enable various parts of the analysis, including: BEDTools, the Integrated Genome Viewer (IGV) and Picard (http://broadinstitute.github.io/picard). Statistical analyses and sequence enrichment plots were implemented in R (http://www.R-project.org). Illumina MiSeq sequence data sets have been deposited into the NCBI repository under BioProject ID PRJNA256245.

Tumour purity correction. Lung lobes and microdissected tumour genomic DNA was used to perform real-time PCR based analysis to detect the relative levels of the un-recombined 
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alleles (from non-tumour tissue) using forward primer: 5‘-CTCTGGGCTAGGGCACTGTC-3‘ and reverse primer: 5‘-AGCTA GCCACATGGCTGTAGTTACTGC-3‘. To correct for DNA loading of each sample, we amplified the ch5 10054507–10054621 region using forward primer: 5‘-GGAGAATTTAGGGCATGTCCTC-3‘ and reverse primer: 5‘-CTTCTCC CAGTGCATTATTTGA-3‘. Real-time PCR reactions were performed using KAPA Fast SYBR master mix in a Roche LightCycler Real-Time PCR instrument. To calculate percent purity we performed the following calculations for each sample: 

\[
\text{Relative Tumour Purity} = \frac{\Delta Ct_{\text{Target}}} {\Delta Ct_{\text{Internal Control}}} \\
\Delta Ct = Ct_{\text{Target}} - Ct_{\text{Internal Control}} \\
\Delta Ct_{\text{Target}} = Ct_{\text{Target}} - Ct_{\text{Non-Tumour}} \\
\Delta Ct_{\text{Internal Control}} = Ct_{\text{Internal Control}} - Ct_{\text{Non-Tumour}}
\]

To normalize for sample loading followed by analysis of 

\[
\frac{\Delta Ct_{\text{Target}}}{\Delta Ct_{\text{Internal Control}}}
\]

to tumour purity to lung tissue from 
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animals that were not infected with Cre. To validate the assay, we generated mouse embryonic fibroblasts from 
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mice treated with Cre recombinase (or control FlpO recombinase). Purity values are reported in Supplementary Table 3.

Statistics. P values were determined by Student’s t-test for all measurements of tumour burden and IHC quantifications except for contingency tables, in which Fisher’s exact test or Chi-square test were used. All error bars denote s.e.m.

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Extended Data Figure 1 | In vitro validation of pSECC. a, The Green-Go Cre-reporter cell line used to validate pSECC lentiviruses in vitro. Upon infection with a Cre-containing lentivirus, such as pSECC, cells become GFP\(^{1}\), allowing for purification of pSECC-containing cells by FACS. Red and blue triangles denote pairs of loxP sites, with red loxP sites being able to recombine only with other red loxP sites and blue loxP sites being able to recombine only with other blue loxP sites. b, Validation of sgPten-pSECC. Numbers below the bands denote quantitation of protein level relative to empty vector control. c, Validation of sgNkx2-1-pSECC in a cell line that expresses Nkx2-1. d, e, Validation of sgTom-pSECC by fluorescence activated cell sorting (FACS). Briefly, a cell line obtained from a \(Kras^{LSL-G12D}^{+/-}\), \(p53^{fl/fl}\), \(Rosa26^{LSL-tdTomato/LSL-tdTomato}\) mouse was infected with either empty-pSECC (d) or sgTom-pSECC (e) and cultured for 10 days post-infection, after which time the cells were collected and analysed by FACS.
Extended Data Figure 2 | In vivo validation of pSECC. a, Representative H&E and tdTomato IHC staining of serial sections from lung tumours of \( \text{Kras}^{\text{LSL-G12D}^{+/+}}; \text{p}53^{\text{fl/fl}}; \text{Rosa26LSL-tdTomato/LSL-tdTomato} \) mice infected with Empty-pSECC. b–d, Representative H&E and IHC staining of serial sections from negative (b), mixed (c) and positive (d) lung tumours of \( \text{Kras}^{\text{LSL-G12D}^{+/+}}; \text{p}53^{\text{fl/fl}}; \text{Rosa26LSL-tdTomato/LSL-tdTomato} \) mice infected with sgTom-pSECC \( (n = 6) \). e, Distribution of lung tumours from all mice infected with sgTom-pSECC \( (n = 6) \) that were scored as negative, mixed or positive based on tdTomato IHC.
Extended Data Figure 3 | Histological analysis of lung tumours obtained from mice infected with pSECC lentiviruses. a–e, Representative H&E images of lung tumours obtained from mice infected with Empty-pSECC (a), sgTom-pSECC (b), sgNkx2-1-pSECC (c), sgPten-pSECC (d), and sgApc-pSECC (e). f, g, Quantification of tumour burden (total tumour area/total lung area) in Kras<sup>LSL-G12D/+</sup> (f) or Kras<sup>LSL-G12D/+; p53<sup>fl/fl</sup></sup> (g) animals 10 weeks after infection with pSECC lentiviruses expressing: control (empty or sgTom, n = 4 tumours) and Kras<sup>LSL-G12D/+; p53<sup>fl/fl</sup></sup> (n = 7)), sgNkx2-1 (Kras<sup>LSL-G12D/+; p53<sup>fl/fl</sup></sup> (n = 2) and Kras<sup>LSL-G12D/+; p53<sup>fl/fl</sup></sup> (n = 3)), sgApc (Kras<sup>LSL-G12D/+; p53<sup>fl/fl</sup></sup> (n = 3) and Kras<sup>LSL-G12D/+; p53<sup>fl/fl</sup></sup> (n = 6)) and sgPten (Kras<sup>LSL-G12D/+; p53<sup>fl/fl</sup></sup> (n = 4) and Kras<sup>LSL-G12D/+; p53<sup>fl/fl</sup></sup> (n = 3)). h, Quantification of BrdU incorporation (BrdU<sup>+</sup> cells per mm<sup>2</sup>) to assess proliferation of tumour cells from lung tumours in Kras<sup>LSL-G12D/+; p53<sup>fl/fl</sup></sup> animals 10 weeks after infection with pSECC lentiviruses expressing: control (empty or sgTom, n = 4 tumours), sgNkx2-1 (n = 11 tumours), sgApc (n = 10 tumours) and sgPten (n = 15 tumours). Mice were given a pulse of BrdU for 4 h before being euthanized. n.s., not significant, *P < 0.05, **P < 0.01, ***P < 0.001 obtained from two-sided Student’s t-test. All error bars denote s.e.m.
Extended Data Figure 4 | IHC-based analysis of mice infected with sgNkx2-1-pSECC. a–c, Negative (a), mixed (b) and positive (c) lung tumours of mice infected with sgNkx2-1-pSECC. d, Distribution of Nkx2-1 IHC staining status in all sgNkx2-1-pSECC infected animals (n = 8) represented as percent of negative, mixed and positive tumours. Positive tumour, ~100% of the tumour cells stained positive for Nkx2-1. Mixed tumour, at least ~30% of tumour cells stained positive for Nkx2-1. Negative tumour, <25% of the tumour cells stained positive for Nkx2-1.
Extended Data Figure 5 | IHC-based analysis of mice infected with sgPten-pSECC. a–c, Negative (a), mixed (b) and positive (c) lung tumours of mice infected with sgPten-pSECC ($n = 9$). Positive tumour, $>100\%$ of the tumour cells stained positive for Pten. Mixed tumour, at least $30\%$ of tumour cells stained positive for Pten. Negative tumour, $<25\%$ of the tumour cells stained positive for Pten. Dashed line in b demarcates the positive/negative tumour area.
Extended Data Figure 6 | IHC-based analysis of Kras\textsuperscript{LSL-G12D/+} and Kras\textsuperscript{LSL-G12D/+}; p53\textsuperscript{fl/fl}-sgApc tumours. a, Representative H&E and IHC staining of serial sections from Kras\textsuperscript{LSL-G12D/+}; p53\textsuperscript{fl/fl}-sgTom (control, denoted as KP-sgTom here), Kras\textsuperscript{LSL-G12D/+}–sgApc (denoted as K-sgApc here) and Kras\textsuperscript{LSL-G12D/+}; p53\textsuperscript{fl/fl}-sgApc (denoted as KP-sgApc here) lung tumours. CCSP, Clara cell secretory protein; SP-C, surfactant protein C. b, Contingency table demonstrating a statistically significantly higher number of β-catenin/Sox9 double-positive tumours in Kras\textsuperscript{LSL-G12D/+}; p53\textsuperscript{fl/fl}-sgApc mice (29/33 tumours, 88%) vs K-sgApc mice (41/58 tumours, 71%) (one-sided chi-square test, \(P<0.05\)). c, Percentage of all tumours that stained positive for nuclear β-catenin that stained positive or negative for Sox9 in Kras\textsuperscript{LSL-G12D/+} and Kras\textsuperscript{LSL-G12D/+}; p53\textsuperscript{fl/fl}-sgApc mice. d, Contingency table demonstrating a statistically significantly higher number of tumours with Nkx2-1 low/negative areas (which are also SP-C low/negative) in sgApc-pSECC animals compared to sgTom-pSECC control animals (two-sided Fisher’s exact test, \(P<0.0001\)). e, Representative IHC staining of serial sections from an Nkx2-1 Low/Neg lung tumour obtained from a Kras\textsuperscript{LSL-G12D/+}; Apc\textsuperscript{fl/fl} mouse 18 weeks after infection with Adeno-Cre. Inset shows Sox9 staining. Low/neg = tumour that had areas with clear downregulation or complete loss of Nkx2-1 or SP-C as assessed by IHC staining.
Extended Data Figure 7 | Representative examples of indels observed in lungs and tumours from mice infected with pSECC lentiviruses.

a–c. Representative indels observed in the Nkx2-1 (a), Pten (b) and Apc (c) locus from sgNkx2-1T1, sgPtenL1 and sgApcT3 samples, respectively. Left panel, details of sequence alignments around the PAM sequence. Right panel, overview of sequence alignments around the PAM sequence. Deletions and insertions are highlighted in black and purple bars, respectively. Inset in a depicts a magnification of an insertion. d, Distribution of indels (in-frame insertions, frameshift insertions, in-frame deletions and frameshift deletions) observed in samples from mice infected with sgNkx2-1-pSECC, sgPten-pSECC and sgApc-pSECC. Amp, mutations across whole PCR amplicon; PAM, mutations across 7 base pair region upstream of the PAM sequence. e, Table summarizing percentages of indels from total mutant reads (left percentage indicates Amp (mutations across whole PCR amplicon) and right percentage indicates PAM (mutations across 7 base pair region upstream of the PAM sequence). All error bars denote s.e.m.
Extended Data Figure 8 | Off-target analysis. a–i, Analysis of off-target editing for sgNkx2-1 (a–c), sgPten (d–f) and sgApc (g–i). Briefly, potential off-target cutting at the top three predicted off-target sites (obtained from [http://crispr.mit.edu/](http://crispr.mit.edu/); see Supplementary Table 2) for each sgRNA was assayed by Illumina MiSeq. Each plot corresponds to the fraction of bases mutated per position in 10 bp flanks on either side of the PAM sequence (highlighted in red). Samples were obtained from entire lobes (L) from mice 10 weeks after infection with pSECC lentiviruses expressing sgNkx2-1, sgPten, sgApc or sgTom (control).