Here, we describe the multiple lentiviral expression (MuLE) system that allows multiple genetic alterations to be introduced simultaneously into mammalian cells. We created a toolbox of MuLE vectors that constitute a flexible, modular system for the rapid engineering of complex polycistronic lentiviruses, allowing combinatorial gene overexpression, gene knockdown, Cre-mediated gene deletion, or CRISPR/Cas9-mediated (where CRISPR indicates clustered regularly interspaced short palindromic repeats) gene mutation, together with expression of fluorescent or enzymatic reporters for cellular assays and animal imaging. Examples of tumor engineering were used to illustrate the speed and versatility of performing combinatorial genetics using the MuLE system. By transducing cultured primary mouse cells with single MuLE lentiviruses, we engineered tumors containing up to 5 different genetic alterations, identified genetic dependencies of molecularly defined tumors, conducted genetic interaction screens, and induced the simultaneous CRISPR/Cas9-mediated knockout of 3 tumor-suppressor genes. Intramuscular injection of MuLE viruses expressing oncogenic $H-Ras^{G12V}$ together with combinations of knockdowns of the tumor suppressors cyclin-dependent kinase inhibitor 2A ($Cdkn2a$), transformation-related protein 53 ($Trp53$), and phosphatase and tensin homolog ($Pten$) allowed the generation of 3 murine sarcoma models, demonstrating that genetically defined autochthonous tumors can be rapidly generated and quantitatively monitored via direct injection of polycistronic MuLE lentiviruses into mouse tissues. Together, our results demonstrate that the MuLE system provides genetic power for the systematic investigation of the molecular […]
A versatile modular vector system for rapid combinatorial mammalian genetics

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Here, we describe the multiple lentiviral expression (MuLE) system that allows multiple genetic alterations to be introduced simultaneously into mammalian cells. We created a toolbox of MuLE vectors that constitute a flexible, modular system for the rapid engineering of complex polycistronic lentiviruses, allowing combinatorial gene overexpression, gene knockdown, Cre-mediated gene deletion, or CRISPR/Cas9-mediated (where CRISPR indicates clustered regularly interspaced short palindromic repeats) gene mutation, together with expression of fluorescent or enzymatic reporters for cellular assays and animal imaging. Examples of tumor engineering were used to illustrate the speed and versatility of performing combinatorial genetics using the MuLE system. By transducing cultured primary mouse cells with single MuLE lentiviruses, we engineered tumors containing up to 5 different genetic alterations, identified genetic dependencies of molecularly defined tumors, conducted genetic interaction screens, and induced the simultaneous CRISPR/Cas9-mediated knockout of 3 tumor-suppressor genes. Intramuscular injection of MuLE viruses expressing oncogenic H-RasG12V together with combinations of knockdowns of the tumor suppressors cyclin-dependent kinase inhibitor 2A (Cdkn2a), transformation-related protein 53 (Trp53), and phosphatase and tensin homolog (Pten) allowed the generation of 3 murine sarcoma models, demonstrating that genetically defined autochthonous tumors can be rapidly generated and quantitatively monitored via direct injection of polycistronic MuLE lentiviruses into mouse tissues. Together, our results demonstrate that the MuLE system provides genetic power for the systematic investigation of the molecular mechanisms that underlie human diseases.

Introduction

In the study of many biological processes, it is often desirable to be able to genetically manipulate multiple genes simultaneously. A prime example is tumor modeling in mice. The application of genomic technologies to the study of cancers has revolutionized our understanding of the genetic landscapes of tumors (1, 2); however, functional studies are required to make sense of these cataloguing efforts to determine how different combinations of the many candidate genetic mutations dictate tumor phenotypes and to provide accurate models that can be used in preclinical studies to identify mutation-specific therapies. Germline genetic manipulation techniques have allowed the generation of lines of mice in which genes can be deleted, mutated, silenced, or overexpressed in temporal and cell-type-specific manners (3). These approaches have provided many insights into cancer development and progression; however, the generation of genetically modified mice and their interbreeding to generate compound mutants are time consuming and costly processes. In addition, tumor modeling ex vivo using primary cells is often complicated by the limited period for which these cells can be cultured and the lack of appropriate genetic tools that allow multiple genetic alterations to be introduced simultaneously into these cells. Current gene-delivery approaches aiming to introduce multiple genetic alterations using plasmid or viral vectors often require cumbersome cloning approaches to generate complicated polycistronic vector constructs and/or multiple rounds of transfection or transduction and necessitate the introduction of multiple selective markers to identify and isolate the appropriate cells.

To address these issues, we have developed the multiple lentiviral expression (MuLE) system, which is based on MultiSite Gateway cloning (4) and allows the easy and flexible generation of polycistronic, replication-incompetent ecotropic or amphotropic lentiviruses. This system allows complex combinatorial genetic alterations to be introduced into mammalian cells by infection ex vivo and in vivo with a single vector. The use of lentiviral gene delivery permits the transduction of a wide variety of dividing and nondividing cells, and the integration of the provirus into the genome provides heritability of the introduced genetic alterations. We present examples of cancer engineering ex vivo and in vivo to demonstrate that this system, when used alone or in combination with germline genetic approaches, provides new experimental genetic power in cultured mammalian cells and in mice.

Results

Generation of complex polycistronic lentiviruses using the MuLE vector toolbox. To be able to quickly and systematically conduct combinatorial genetic experiments in cultured mammalian cells and
in mouse tissues, we designed a genetic system that allows complex lentiviral gene-delivery vectors to be generated in a very flexible, rapid, and user-friendly manner. We generated a toolbox of building-block vectors that allows multiple genetic elements to be combinatorially recombined into a family of lentiviral gene-delivery vectors. A series of 104 MultiSite Gateway–compatible Entry vectors (MuLE Entry vectors, Supplemental Tables 1 and 2; supplemental material available online with this article; doi:10.1172/JCI79743DS1) that allow several different types of genetic manipulations has been generated. Conventional restriction enzyme cloning is not practical for the rapid, high-throughput, and user-friendly production of lentiviral vectors containing multiple distinct genetic elements.
ing is first used to clone genetic inserts downstream of a promoter of choice, with the entire promoter-insert genetic element being flanked by specific attl and attR sites (Figure 1A). MuLE Entry vectors can be recombined efficiently in an overnight reaction via an attl-attR recombination with a variety of Gateway destination lentiviral vectors that we have generated. The choice of different attl-attR sites provides directional specificity to the recombination reactions and allows either 1, 2, 3, or 4 MuLE Entry vectors to be simultaneously recombined into the Destination vector. Using this system, complex multicistronic mammalian expression vectors containing 2, 3, 4, or 5 independent genetic elements can be rapidly generated (Figure 1B). MuLE vectors can be packaged to generate amphotropic or ecotropic lentiviruses, which can then be used for infection of cultured cell lines, primary cells ex vivo or cells in mouse tissues in vivo (Figure 1C). This entire procedure can be completed within 7 to 8 days.

We generated Entry vectors containing 3 different RNA polymerase (pol) II promoters (CMV, SV40, SFFV) and a multiple cloning site (MCS) to generate genetic elements allowing constitutive cDNA overexpression (Figure 2A). An Entry vector with a CMV promoter and loxP sites flanking the MCS allows for Cre-mediated conditional removal of inserted genes (Figure 2A) and Entry vectors containing a doxycycline-inducible (DOX-inducible) CMV/TO promoter allow inducible gene expression (Figure 2B). Supplemental Table 3 shows the restriction enzyme sites of the MCS of each vector. Promoterless Entry vectors containing only a MCS flanked by different combinations of attl-attR sites (Figure 2C) represent flexible elements that facilitate the generation of diverse types of complex vector constructs. We generated Entry vectors to allow U6 promoter–driven expression of short guide RNAs (sgRNAs) for CRISPR/Cas9-mediated (where CRISPR indicates clustered regularly interspersed short palindromic repeats) gene mutation (Figure 2D). Entry vectors with 3 different constitutive pol III promoters (U6, 7SK, H1) allowed the expression of shRNAs (Figure 2E). For constitutive expression of shRNAs in microRNA-30 (miR-30) format, we generated Entry vectors based on the pSM2 plasmid (5), which includes the 5′ and 3′ flanking sequences from miR-30 driven by the U6, 7SK, or H1 pol III promoters (Figure 2F). A DOX-inducible gene knockdown vector was generated by using the CMV/TO promoter to drive shRNA–miR-30 expression (Figure 2G). In addition to these flexible cloning vectors, we created several ready-to-use Entry vectors containing cDNAs for hCas9, fluorescent proteins (EGFP, mCherry, near-infrared fluorescent protein [iRFP], tdTomato), enzymatic reporters (luciferase, lacZ), drug resistance (puromycin), and tamoxifen-inducible Cre recombinase (Cre-ER<sup>29</sup>) (Figure 2H). To allow a high degree of flexibility in combining different numbers and types of Entry vectors by recombination cloning, most of the described vectors are available with multiple combinations of different attl-attR sites (Supplemental Tables 1 and 2). We further extended our vector toolbox by generating a series of lentiviral Destination vectors based on modifications of the previously described pLenti XI-Puro-DEST vector (6), allowing cells to be marked with drug resistance (puromycin, neomycin) or a variety of fluorescent (EGFP, iRFP, IFP1.4) or luminescent (luciferase) proteins (Figure 2I) to facilitate cellular assays as well as live animal-imaging studies. We performed numerous recombination reactions with the MuLE system and found that approximately 90%, 65%, or 25% of all bacterial colonies isolated from recombinations involving 2, 3, or 4 Entry vectors, respectively, contained the desired product (Figure 2J). Thus, highly complicated, multicistronic MuLE expression vectors can be easily cloned. These complex vectors can be stably replicated in bacteria and show no evidence of unwanted genetic recombination (Supplemental Figure 1). Similarly to all lentiviral vectors, there is an inverse relationship between the size of the MuLE provirus and the titer of the generated virus, with proviruses up to 7.5 kb yielding titers of approximately 10<sup>6</sup> CFU/ml and very large (12.5 kb) proviruses yielding titers of approximately 10<sup>4</sup> CFU/ml (Supplemental Figure 2). All of these vectors have been deposited with Addgene and can be obtained individually or as a 96-vector set. Addgene reference numbers for each vector are provided in Supplemental Tables 1 and 2.

While the MuLE system is compatible with the VSV-G amphotropic envelope protein (Supplemental Figure 5 shows an example), for biosafety reasons, all lentiviruses used to transduce murine cells in this study were pseudotyped with an ecotropic envelope protein from the Moloney murine leukemia virus that has been described as allowing lentiviral infection of murine fibroblast and hematopoietic cells, but not of human cells (7, 8). Despite obvious biosafety advantages, this envelope has surprisingly found very little use in research laboratories. We further examined the cellular tropism of MuLE lentiviruses pseudotyped with this envelope and found that they infect a variety of cultured primary cells, including mouse embryonic fibroblasts (MEFs), embryonic stem cells, kidney epithelial cells, endometrial epithelial cells, hepatocytes, and aortic endothelial cells as well as immortalized myoblast, melanoma, lung carcinoma, and colorectal carcinoma cell lines, but were not able to infect several human cell lines, including 786-O, 293T (Figure 3A), HeLa, and MCF-7 (data not shown). Cell-type–specific gene expression is also possible using MuLE vectors. Cloning EGFP downstream of the renal epithelium–specific Ksp1.3 promoter in a MuLE vector (Figure 3B) allowed expression of EGFP in renal epithelial cells but not in MEFs (Figure 3C). Thus, ecotropic MuLE viruses can infect a broad spectrum of target cells and can be utilized to induce gene expression in a cell-type–specific manner.

In order to validate the functionality of the vector toolbox for the genetic manipulation of primary cells, we conducted a series of experiments involving transduction of primary MEFs. Unless otherwise stated, all experiments in this publication were conducted at an MOI of 1. These validation studies demonstrated that MuLE vectors are capable of the following: (a) inducing shRNA-mediated knockdown from different pol III promoters (Supplemental Figure 3, A–C); (b) inducing simultaneous double-gene knockdowns from a single vector in a manner in which the strengths of the knockdowns are independent of the order of the knockdown cassettes in the vector (Supplemental Figure 3, D and E) and are equivalent to the extent of knockdowns obtained from vectors expressing the single shRNAs (Supplemental Figure 3, F and G); (c) inducing shRNA–miR-30–mediated constitutive (Supplemental Figure 3, H–J) and inducible (Supplemental Figure 3, K–M) gene knockdown; (d) inducing tamoxifen-inducible Cre-ER<sup>29</sup>–mediated gene deletion (Supplemental Figure 4, A and B); and (e) inducing the
Engineering genetically complex tumors from primary cells using single MuLE viruses. To demonstrate the utility of the MuLE system in cancer modeling, we performed experiments in MEFs showing that a single virus can reproduce several phenotypes that are known to result from genetic cooperation between oncogenes and tumor-suppressor genes. Overexpression of an oncogenic form of simultaneous expression of 2 shRNAs, 2 cDNAs, and a drug-resistance gene from 5 different expression cassettes in a single vector (Supplemental Figure 4, C–E). Transduction of human A-375 melanoma cells showed that MuLE viruses expressing iRFP or luciferase proteins allow quantitative monitoring of tumor burden in xenograft experiments (Supplemental Figure 5).

Figure 2. The MuLE vector toolbox. (A and B) MuLE Entry vectors for pol II promoter–driven constitutive expression (A) or DOX-inducible expression (B). (C) Promoterless MuLE Entry vectors. (D) MuLE Entry vectors for U6–driven expression of sgRNAs. (E and F) MuLE Entry vectors for shRNA–based (E) and shRNA–miR-30–based (F) gene knockdown using pol III promoters and (G) a DOX-inducible miR-30–based shRNA expression vector. Restriction enzyme sites for cloning are shown. (H) Schematic representation of MuLE Entry vectors for expression of hCas9, fluorescent proteins (EGFP, mCherry, iRFP, tdTomato), firefly-luciferase, β-galactosidase (LacZ), puromycin resistance, or Cre-ERT2. P various promoters. In all panels, attA and attB denote that multiple combinations of MultiSite Gateway attL-attR sites are available for these vectors. (I) Schematic representation of Destination vectors modified from the pLenti X1 series to contain the different expression cassettes shown. (J) Quantification (mean ± SD) of recombination efficiencies of n independent MultiSite Gateway attL-attR recombinations using 2, 3, or 4 MuLE Entry vectors.
H-Ras (G12V) in combination with loss of function of p53 results in cellular transformation (9, 10). We generated a tricistronic lentiviral expression vector designed to simultaneously express shRNA–miR-30 against transformation-related protein 53 (Trp53) and express oncogenic H-RasG12V and puromycin resistance (Figure 4A) as well as control viruses expressing neither or only 1 of these elements. Western blot analysis of puromycin-selected primary MEFs demonstrated the expected knockdown of p53 and overexpression of H-RasG12V (Figure 4B), and transformation assays demonstrated that Trp53 knockdown alone allowed colony formation following plating at low density (Figure 4C), but that loss of contact inhibition (foci formation) and anchorage-independent growth occurred only in cells with simultaneous knockdown of Trp53 and overexpression of H-RasG12V, as expected (Figure 4, D–F). Similarly, the use of a single MuLE vector to simultaneously knock down Trp53 and overexpress Myc (Figure 4G and H) allowed growth of cells at low density (Figure 4D) and reproduced the known effect of loss of Trp53 function in rescuing Myc-induced apoptosis in MEFs (ref. 11 and Figure 4).

To further prove the utility of our system in tumor modeling, we generated a multicistronic vector designed to knock down Trp53, overexpress H-RasG12V from the inducible CMV/TO promoter, and express iRFP to allow monitoring of tumor development (Figure 5A). Blasticidin-resistant MEFs after infection with pLenti-CMV-TetR-Blast (6) were transduced with the MuLE vector and selected with puromycin. Western blot analysis indicated constitutive p53 knockdown, but H-RasG12V overexpression occurred only upon induction with DOX (Figure 5B). These cells grew efficiently as tumors when injected subcutaneously in immunocompromised SCID/beige mice only upon addition of DOX to the drinking water (Figure 5, C–E), demonstrating that MuLE vectors can be employed to generate regulatable and quantitatively monitorable models of cancer based on cooperative genetic interactions.

To further demonstrate the potential of MuLE vectors for engineering and monitoring genetically complex tumors, we generated a pentacistronic vector to simultaneously express shRNA–miR-30 against the Retinoblastoma (Rb1) tumor-suppressor gene and shRNA against the phosphatase and tensin homolog (Pten) tumor-suppressor gene as well as to express oncogenic H-RasG12V, Cre-ERT2, and the puromycin resistance gene (Figure 4A). We transduced (MOI = 2) primary MEFs harboring floxed alleles of the von Hippel–Lindau (Vhl) and Trp53 genes (Vhlfl/fl Trp53fl/fl) and treated puromycin-selected cells with 300 nM 4-hydroxytamoxifen (4-OHT) or vehicle for a period of 3 days. Cells transduced with the lentivirus and treated with 4-OHT showed reduction of VHL abundance (accompanied by stabilization of HIF1A), reduction of PTEN, pRb, and p53 protein levels, and overexpression of H-RasG12V, (Figure 6B) thus demonstrating the functionality of each expression cassette in the MuLE vector. To analyze the potential of these cells to form genetically defined tumors in vivo, we additionally transduced the cells with a MuLE virus expressing EGFP and iRFP, thereby generating within 1 week pools of cells that harbored 5 separate genetic changes and expression of 3 marker genes. Cells that were treated with 4-OHT grew marginally faster in xenograft...
assays than those that were not treated (Figure 6C and Supplemental Figure 6, A and B). Importantly, cells that were isolated from the tumors 3 weeks after injection retained EGFP expression (data not shown) and displayed the same changes in protein abundance that were present in the cells before injection (Figure 6D), demonstrating that MuLE viruses permit the introduction of stable genetic alterations.

We took advantage of this system to perform proof-of-principle therapeutic target identification experiments to show that MuLE viruses can be employed to assess which genes are necessary for the growth of tumors with a defined set of genetic mutations. Using the same workflow described above, we infected primary MEFs derived from WT, 

Hif1afl/fl, Hif2afl/fl, or Hif1afl/fl Hif2afl/fl mice with the virus shown in Figure 6A to assess the contribution of the hypoxia-inducible factor α (HIF1α and HIF2α) transcription factors to tumor development in the background of oncogenic 

H-RasG12V together with loss of function of Rb1 and Pten. mRNA analyses confirmed that 4-OHT treatment induced the anticipated downregulation of Hif1a and/or Hif2a and of several Hif1a target genes in these cells (Figure 7A). Xenograft experiments revealed that HIF1α, but not HIF2α, is necessary for efficient tumor growth in this model (Figure 7, B–E). All tumors described in Figures 6 and 7 displayed a similar histological appearance of sarcomatoid cells growing in a storiform pattern, with numerous tumor giant cells with bizarre nuclei (Supplemental Figure 6, C and D). We took advantage of this large set of imaging data to perform an analysis of the utility of iRFP to monitor tumor growth. iRFP is a new fluorescent protein that has been proposed to be

![Figure 4. Combinatorial genetics using MuLE vectors.](image-url)

(A) Lentiviral vector to simultaneously knock down Trp53 and overexpress oncogenic H-RasG12V. (B) Western blot analysis of primary MEFs transduced with the indicated lentiviruses. (C) Crystal violet staining of the same cells 14 days after seeding at low density and (D) 14 days after seeding in a focus formation assay. (E) Representative images of the same cells seeded in a soft agar colony assay after 3 weeks of growth. Scale bars: 200 μm. (F) Quantification of the foci and colonies growing in assays from D and E. (G) Lentiviral vector generated to simultaneously knock down Trp53 and overexpress Myc. (H) Western blot analysis of primary MEFs transduced with the indicated lentiviruses. (I) Crystal violet staining of the same cells 14 days after seeding at low density. (J) Quantification of viable cells 3 days after transduction with the indicated lentiviruses. All graphs depict mean ± SD. Student’s t test, n = 3. **P < 0.01; ***P < 0.001.
excellent for animal imaging due to the low absorbance by mouse tissues of the emitted light (12–14). In each of the 5 experiments described above, there was a strong linear correlation between the iRFP signal and the measured tumor volume over time, with Pearson correlation coefficient \( R^2 \) values ranging from 0.79 to 0.94 in the experiments (Supplemental Figure 6E), supporting the utility of iRFP as an excellent marker protein to track tumor formation and progression in mouse imaging studies.

**Combinatorial CRISPR/Cas9-mediated genetic engineering using single MuLE vectors.** The discovery of the applicability of the CRISPR/Cas9 system to mammalian cells has provided a powerful new tool to efficiently target genetic mutations to defined loci (15, 16), and very recent publications illustrate the power of this system for tumor engineering in mice (17–20). We reasoned that the ease and flexibility of cloning provided by the MuLE system would make it an ideal experimental platform to generate vectors that allow the introduction of multiple simultaneous genetic manipulations using CRISPR/Cas9. We first generated tricistronic MuLE vectors expressing single sgRNAs targeting exon 7 or exon 8 of the Trp53 locus or exon 2 of the cyclin-dependent kinase inhibitor 2A (Cdkn2a) locus together with expression of hCas9 and puromycin resistance (Supplemental Figure 7A) and infected primary MEFs. Two functionally validated Trp53 sgRNAs (21) and 3 newly designed Cdkn2a sgRNAs, but not a scrambled control sgRNA (21), induced efficient mutation of their respective target genes, as shown by Surveyor assays (Supplemental Figure 7B). Both Trp53 sgRNAs caused the generation of truncated p53 protein species, and 2 of 3 Cdkn2a sgRNAs caused loss of p16 and p19 protein expression (Supplemental Figure 7C), verifying that CRISPR/Cas9 genome engineering is compatible with the MuLE system.

To demonstrate that cooperative genetic tumor modeling can be achieved using CRISPR/Cas9-mediated gene knockouts together with oncogene overexpression from a single vector, we generated tetracistronic MuLE vectors designed to express either scrambled sgRNA or sgRNAs targeting Trp53 exon 7 or exon 8 as well as to express H-RasG12V, hCas9, and puromycin resistance (Figure 8A) and used these to infect primary MEFs. All viruses expressing sgRNA against Trp53, but not the control virus, allowed colony formation following plating of cells at low density (Figure 8B), but tumor formation in xenograft assays only resulted from the combination of sgRNA against Trp53 with H-RasG12V expression (Figure 8C). Western blotting of independent cell lines isolated from independent tumor xenografts of each genotype confirmed the overexpression of H-RasG12V and the presence of numerous truncated mutant p53 protein species (Figure 8D).

Previous approaches using the CRISPR/Cas9 system to generate cells harboring multiple mutations required the cotransfection of multiple plasmids, the simultaneous infection with multiple viruses, and the use of transgenic mice that express hCas9 or that carry germine floxed alleles (17, 18, 20, 22, 23). The multicistronic building-block MuLE viral system allows multiple sgRNAs to be expressed together with hCas9 from a single viral construct. To prove that this approach of targeting multiple loci simultaneously with a single vector is feasible, we generated pentacistronic MuLE viruses, each simultaneously expressing sgRNAs against the Trp53, Pten, and Vhl tumor-suppressor genes, hCas9, and puromycin resistance (Figure 8E). Two different sgRNAs targeting Trp53 exon 7 or exon 8, 3 different sgRNAs targeting Pten exon 1 or exon 2, and 3 different sgRNAs targeting Vhl exon 1 were combined to generate 3 independent MuLE viruses expressing different sets of sgRNAs. After infection and selection for drug resistance, cells were plated at low density to assess cellular transformation. All viruses allowed the formation of colonies (Figure 8F), indicative of loss of p53 function, and a total of 21 clonal immortalized cell lines were derived from isolated colonies. Western blotting revealed the complete loss or reduction of expression of PTEN and VHL and presence of truncated p53 protein species in 16, 14, and 20 clones, respectively (Figure 8G). All clones displayed levels
of the p53-inducible p21 protein equivalent to levels in Vhl/Trp53 null MEFs, verifying that functional p53 knockout occurred in all clones. Sixteen clones displayed stabilization of HIF1α protein, and 14 clones displayed elevated levels of P-S473-Akt, indicative of functional loss of VHL and PTEN, respectively (Figure 8G). Since some clones retained protein expression of p53, VHL, and/or PTEN, we further validated that mutations arose at all loci by using deep sequencing to genotype the regions of Pten, Trp53, and Vhl targeted by the sgRNAs in each of 10 clones (Table 1). Half of the clones showed insertions or deletions of all 6 alleles, and half showed mutations of 5 of 6 alleles, demonstrating that infection with a single MuLE CRISPR/Cas9 vector can simultaneously induce mutations in multiple target genes.

Combination of genetic screening using MuLE vectors. Another advantage of the building-block nature of the MuLE system is that it provides a platform to facilitate population-based genetic interaction screens. As proof of principle that this is feasible, we conducted a screen to identify negative regulators of the cell cycle whose loss of function can cooperate with H-RasG12V to allow transformation of MEFs. We assembled an shRNA library (designated shRNA X) in the commercially available pLKO.1 vector backbone, comprising 4 to 6 shRNAs against each member of the Cdkn1, Cdkn2, Trp53, and Rb gene families (Figure 9A). PCR was used to amplify all shRNAs, including the U6 promoter, from a pooled DNA preparation of this library, and the product was cloned en masse into a MuLE Entry vector to generate an Entry shRNA X library, which was recombined to generate a library of tricistronic MuLE vectors, each expressing a single shRNA from the library together with H-RasG12V and EGFP. These MuLE vectors and control vectors expressing either nonsilencing shRNA, the shRNA X library alone, or H-RasG12V alone were transduced into MEFs at an MOI of 0.1 to avoid multiple integrations into the same cell. Transduced cells were plated at low density in the presence of a 30-fold excess of WT MEFs for a focus formation assay. The combination of the shRNA X library plus H-RasG12V resulted in an increased number of foci (Figure 9B). Eight EGFP-positive foci were isolated and expanded as cell lines. PCR followed by DNA sequencing was used to isolate and identify the integrated shRNA, revealing that 7 of these clones contained shRNA against Cdkn2a (clone TRCN0000222731) and 1 contained shRNA against Trp53 (clone TRCN0000012360). Western blotting verified the knockdown of p19 and p16 protein expression in the Cdkn2a shRNA–expressing cell lines and of p53 in the Trp53 shRNA–expressing cell line (Figure 9C). Reengineering MuLE viruses to express the isolated Cdkn2a shRNA together with H-RasG12V and iRFP expression cassettes, as well as relevant control viruses, confirmed that this genetic cooperation causes transformation of MEFs and allows these cells to grow as xenografts in immunocompromised mice (Figure 9, D and E). Loss of function of Cdkn2a or of Trp53 has been shown to cooperate with activated H-Ras in causing transformation of MEFs (24), validating the effectiveness of screening for cooperative genetic events with the MuLE system.

Generation of genetically complex, quantitatively monitorable autochthonous sarcomas in mice using single MuLE viruses. Having established the power of the MuLE system for combinatorial genetics in primary cells ex vivo and based on experiments show-
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Figure 7. Hif1a but not Hif2a is necessary for efficient growth of Pten/Rb1-deficient, H-RasG12V-expressing tumors. (A) mRNA expression analysis of the indicated genes in WT, Hif1aflox/flox, Hif2aflox/flox, and Hif1aflox/flox Hif2aflox/flox MEFs that were transduced with lentivirus generated from the vector shown in Figure 6A 96 hours after induction of CreER2 with 300 nm 4-OHT. Shown are ratios of 4-OHT treated to EtOH treated. (B–E) In vivo fluorescence imaging at the day of euthanasia (top left panels), excised tumors (top right panels), and longitudinal tumor growth (bottom panels) of mice injected with cells described in A that had been treated with EtOH (left flank) or 4-OHT (right flank) prior to subcutaneous injection. Color intensity in B–E is the same as in Figure 6C. Scale bars: 1 cm. All graphs depict mean ± SD. Student’s t test, n = 3–6. *P < 0.05; **P < 0.01.

ing that combinatorial genetic manipulations via lentiviral injection can generate brain tumors (25), we tested whether autotrophic tumors could also be induced in mice via direct transduction of cells with MuLE viruses in vivo. Using R26-lox-STOP-lox-tdTomato mice (26) as a reporter system to monitor cellular infection, we found that injection of ecotropic viruses expressing Cre into the gastrocnemius muscle of mice induced tdTomato fluorescence in muscle fibers and small cells adjacent to muscle fibers (Figure 10A). Mouse models have demonstrated that oncogenic K-Ras expression combined with loss of Trp53 or Cdkn2a function in skeletal muscle cells of different stages of differentiation causes several types of soft tissue sarcomas, including embryonal and pleomorphic rhabdomyosarcomas (RMS) as well as undifferentiated pleomorphic sarcomas (UPS) with or without myogenic differentiation (27–30). The RAS-signaling pathway is frequently activated in childhood RMS tumors due to oncogenic mutations in the HRAS, KRAS, or NRAS genes or homozygous deletions of NF1 (31–35). Some RMS tumors also harbor loss-of-function mutations or gene deletions in the CDKN2A, TP53, and PTEN genes (32–34, 36, 37), and systematic analysis of the status of the TP53 and CDKN2A loci, as well as of their respective proteins, revealed that UPS tumors almost universally display loss of 1 or more components of the p53 pathway (38). To determine whether MuLE viruses could be utilized to model human sarcomas, we generated a series of MuLE vectors designed to systematically investigate the single and combinatorial effects of gain of H-Ras function and loss of Cdkn2a, Trp53, and Pten functions. All MuLE vectors expressed luciferase to mark infected cells, allowing quantitative monitoring of tumor development. Concentrated ecotropic lentiviruses (10^7 CFU/ml) were injected once into each gastrocnemius muscle of 18-day-old SCID/beige immunocompromised mice. Injection of empty control virus, viruses expressing only
Figure 8. Combinatorial genetics using the CRISPR/Cas9 system in MuLE vectors. (A) Schematic of MuLE vector expressing sgRNA against Trp53 and expressing H-RasG12V, hCas9, and puromycin resistance. (B) MEFs were infected with the indicated viruses expressing sgRNAs targeting Trp53 exon 7 (Ex7) or exon 8 with or without H-RasG12V expression, plated at low density 6 days after transduction, and stained with crystal violet 14 days after plating. (C) Growth of cells as tumor xenografts and images of tumors derived from the combination of Trp53 exon 7 or exon 8 sgRNAs with H-RasG12V overexpression. (D) Western blot analysis of tumor cells that were isolated from 3 separate tumors of each genotype 5 weeks after cell injection. (E) Schematic of MuLE vector simultaneously expressing sgRNAs against Trp53, Pten, and Vhl and expressing hCas9 and puromycin resistance. (F) MEFs infected with 3 independent combinations of different sgRNAs formed colonies when plated at low density 10 days after viral transduction. (G) Western blotting of cell lines (lanes 1-21) derived from colonies that formed after infection with viruses expressing the indicated combinations of sgRNAs.
tion. Histological analysis of the tumors revealed that they were undifferentiated sarcomas with pleomorphic and rhabdoid features. Tumors contained undifferentiated round to spindle cells and an admixture of polygonal cells with densely eosinophilic cytoplasm in spindle, tadpole, and racquet-like contours (Figure 10, L and M) demonstrating myogenic differentiation in some cells. Injections of shRNA-Cdkn2a plus H-RasG12V demonstrated myogenic differentiation in the malignant cells (Figure 10, G, I, and K) and the demonstration by electron microscopy of sarcomere formation in the malignant cells (Figure 10, A–H). Heterologous foci of tumors and tumor-derived cancer cell lines in mice.

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<tr>
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The sequences of the 2 alleles of each locus are listed; underlined dots represent a base deletion and underlined text a base insertion compared with the WT sequence.

Discussion

We show proof-of-principle examples that illustrate the user friendliness, speed, versatility, and genetic power of the MuLE system in primary mammalian cells. Complex genetic manipulations involving combinatorial gene overexpression, knockdown, and knockout can be achieved using single viral infections. The simultaneous marking of infected cells with a variety of reporter cassettes facilitates cellular studies and in vivo imaging studies. By expressing multiple sgRNAs together with hCas9, single MuLE vectors are able to simultaneously target genetic mutations to multiple loci, providing a platform that harnesses the genetic power of the CRISPR/Cas9 system for performing combinatorial genetic manipulations. Combinatorial genetic screening experiments are also possible by combining libraries of Entry vectors with Entry vectors that encode defined genetic alterations. The ease of combining different genetic elements using this system represents a tool that will greatly facilitate systematic combinatorial genetic studies in mammalian cells. Although this study focuses on tumor biology, the MuLE system will be widely applicable to numerous areas of biological investigation. It is hoped that other investigators will utilize the building-block nature of the system to generate new Entry vectors that will expand the range of possible genetic manipulations.

While there are several theoretical concerns that could potentially be envisaged as limiting the utility of complex MuLE lentiviruses, our data indicate that these do not represent significant problems. While some lentiviral vector plasmids are unstable in bacteria, MuLE vector plasmids are not prone to unwanted recombination during bacterial propagation. Another potential source of recombination in lentiviral vectors is at the level of...
own experiments using repeated U6 promoter elements showed efficient knockdown or CRISPR/Cas9-mediated knockout of all genes that were targeted by the shRNAs or sgRNAs, suggesting that high levels of recombined viruses are not being produced in a manner that negatively affects achieving the desired genetic modulations. Another theoretical drawback of lentiviral vectors is the potential for insertional mutagenesis, although new generation HIV-based self-inactivating (SIN) lentiviral vectors (such as the vector that the MuLE system is based on) appear to not suffer from the problems of deleterious insertional mutagenesis that were observed with initial γ-retroviral gene-delivery vectors.

Figure 9. Combinatorial genetic screening using MuLE vectors. (A) Schematic representation of the workflow to generate a MuLE Entry vector shRNA library targeting the listed genes (shRNA X) and the final tricistronic lentiviral expression vector library that was used to screen for shRNAs that cause cell transformation in cooperation with oncogenic H-RasG12V overexpression. (B) Quantification of foci that were formed when WT MEFs were transduced with the indicated lentiviruses (MOI = 0.1). (C) Western blot analysis of EGFP-expressing cell clones derived from foci harboring shRNA against the indicated gene. (D) In vivo fluorescence images of mice that were subcutaneously injected with WT MEFs that had been infected with a MuLE virus expressing the identified shRNA against Cdkn2a alone plus iRFP or in combination with H-RasG12V and iRFP. (E) Tumor growth in the same mice monitored by longitudinal in vivo fluorescence imaging. All graphs depict mean ± SD. Student’s t test, n = 3. **P < 0.01; ***P < 0.001.

reverse transcription of the viral genome. Some studies have described that repeated genetic elements can undergo recombination-mediated deletion at low frequency in viral infections (39, 40), while other studies show that recombination of repeated elements was not detectable (41, 42). For this reason, we sought to provide options that would allow users to avoid the use of repeated elements in MuLE vectors in the unlikely event that unwanted recombination would prove to be an experimental problem. We generated Entry vectors with multiple different pol II and pol III promoters to allow gene and shRNA expression without incorporating repeated elements into final MuLE expression vectors. Our
Lentiviral vectors have been shown to be safe in recent gene therapy trials (43, 44). Several lines of evidence demonstrate that random insertional mutagenesis of MuLE vectors does not contribute to the observed phenotypes in this study. Every experiment included all control vectors, including vectors with empty inserts, or expressing only single or double combinations of genes to control for potential effects of insertional mutagenesis. For example, as shown in Figure 4, E and F, cellular transformation and anchorage-independent growth only occurred when Trp53 was knocked down together with H-RasG12V overexpression. Cells that were infected with MuLE viruses expressing only shRNA-Trp53 or only H-RasG12V represent genetically sensitized cells that could be envisaged as being prone to oncogenic transformation by insertional mutagenesis. However, transformed cells that could grow in soft agar were never observed in these infections. Similarly, in all of the sarcoma tumor-modeling experiments, none of the relevant empty, single, and double control vectors induced tumors. Tumors arose rapidly in 100% of mice after injection of the shRNA-Cdkn2a plus H-RasG12V, shRNA-Trp53 plus H-RasG12V, and shRNA-Trp53 plus shRNA-Pten plus H-RasG12V combinatorial MuLE viruses with almost uniform kinetics, indicating that the tumors are driven by the genetic alterations introduced by the MuLE viruses and not as a result of rare random cooperating mutagenic events. One real practical limitation to the complexity of cloned MuLE vectors is that, as with all lentiviruses, viral titer decreases proportionally to the size of the provirus (Supplemental Figure 2 and ref. 45). In our hands, proviruses up to approximately 12 kb were able to generate experimentally usable viral titers; however, users are advised to generate the smallest possible proviruses if achieving a very high viral titer is experimentally important. The MuLE Entry vector toolbox is theoretically also compatible with other Gateway cloning-based plasmid and viral delivery systems, potentially allowing for larger and more complicated vectors and providing additional opportunities for targeting different cell types.

A great promise of the MuLE system is the potential to allow genetic manipulation of somatic cells directly in mice and potentially in other mammals. Bypassing extensive germline transgenic approaches has major benefits in terms of time and cost, and the lentiviral-mediated somatic genetics approach also mimics the fact that many cancer-associated genetic alterations are acquired in somatic cells. In this context, by engineering 3 new autochthonous mouse models of sarcoma, we show that MuLE viruses can be used to systematically assess the contribution of tumor suppressors and oncogenes to tumor formation in vivo. We show that the combinations of oncopgenic H-RasG12V expression plus knockdown of Cdkn2a, Trp53, or both Trp53 and Pten cause the formation of undifferentiated sarcomas with pleomorphic and rhabdoid features from skeletal muscles in mice. The histological similarity of these tumor models to existing transgenic mouse muscle–derived sarcoma models that are driven by oncogenic K-RasG12V in Trp53 null, Trp53 point mutant, or Cdkn2a null genetic backgrounds (27–30) demonstrates that MuLE viruses can be employed to recapitulate transgenic tumor models. Interestingly, additional Pten knockdown in shRNA-Trp53 plus H-RasG12V tumors caused phosphorylation of AKT, demonstrating hyperactivation of the PI3K pathway, but did not lead to any obvious differences in the phenotype of these tumors. It is noteworthy that Trp53/Pten double-knockdown or Pten knockdown plus H-RasG12V expression did not lead to tumor formation. These observations are potentially consistent with a recent report that activating PI3KCA and inactivating PTEN mutations in embryonal RMS mostly occur concurrently with mutations in MYOD1 that functionally block myogenic differentiation (46). Thus, in muscle-derived sarcomas, the Ras and PI3K pathways appear to cooperate with different sets of genes to cause tumor formation.

The fact that MuLE-derived tumors are marked with luciferase and arise within weeks with uniform kinetics in every mouse will be advantageous for preclinical therapeutic studies, as tumor burden is quantitatively monitorable using a simple live-animal-imaging approach. In comparison, the generation of similar models using conventional genetic approaches would require the interbreeding of at least 4 to 5 different germline-modified transgenic mice (for example, tissue-specific Cre expression; homozygous floxed Cdkn2a, Trp53, and/or Pten alleles; Cre-inducible alleles of oncogenic H-RasG12V; and luciferase). The flexibility and speed of cloning that the MuLE system allows also paves the way for a systematic approach to model sarcomas by introducing other combinations of genetic alterations that occur in these tumor types, without the need for extensive crossing of germline-modified mice. By generating autochthonous tumors via direct injection of MuLE viruses into existing genetically modified mouse strains, it should also be possible to rapidly test, for example, the role of potential genetic modifiers or the immune system in tumor progression.

Lentiviral vectors allow gene transfer into dividing and nondividing cells and are increasingly being used for somatic cell transgenics, including oncogene delivery (25, 47–49). The most common envelope protein used for pseudotyping lentiviral vectors is from vesicular stomatitis virus (VSV-G) and allows the infection of human cells (50, 51). Here, we have presented numerous examples of tumor engineering using the MuLE system, employing an ecotropic envelope protein from the Moloney murine leukemia virus (7, 52), which offers significant biosafety advantages (8). To the best of our knowledge, this envelope protein has not been previously employed for lentiviral delivery in vivo. In the context of tumor modeling in mice using lentiviral vectors that are potentially oncogenic in humans, the use of an ecotropic envelope prevents possible infection of researchers and also allows the use of conventional biosafety level 1 cell-culture and animal housing facilities, making this system available to almost all research groups. Our preliminary ex vivo and in vivo studies indicate that ecotropic MuLE viruses are able to transduce a wide variety of cell types, suggesting wide applicability of this system. Given the broad tropism of ecotropic MuLE viruses, achieving restricted infection of a particular cell type in vivo via a simple injection may not be possible in some settings. Our proof-of-principle demonstration that cell-type–specific expression can be achieved using a kidney-specific promoter element in the MuLE system may provide a solution to this issue. We also envisage that existing tissue-specific and/or inducible Cre, rtTA, and tTA mouse lines could provide the opportunity to spatially and temporally restrict gene expression from MuLE vectors containing loxP sites or tetracycline-responsive elements.
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TECHNICAL ADVANCE

A

R26-LSL-tdTomato

Ctrl Cre Ctrl

Cre Cre

B

C

D

E

F

shCdkn2a + H-Ras

G

shCdkn2a + H-Ras

H

shTrp53 + H-Ras

I

shTrp53 + H-Ras

J

shTrp53 + shPten + H-Ras

K

shTrp53 + shPten + H-Ras

L

M

N

O

shCdkn2a + H-Ras

shTrp53 + H-Ras

shPten + H-Ras

shTrp53 + shPten + H-Ras

Ras

p53

Pten

Akt

α-Actin
Figure 10. Generation of 3 autochthonous mouse models of undifferen-
tiated sarcoma using MuLE vectors. (A) Intramuscular injection of
ROSAS26- lox- STOP- lox- tdTomato mice with control or Cre-expressing
virus. Bottom left panel shows infected myocytes, and small cells adjac-
to to myocytes are seen at higher magnification (arrowheads, bottom
right panel). (B) Bioluminescence imaging 3 and 31 days after injection of
3 \times 10^6 functional viral particles into each gastrocnemius muscle of
18-day-old SCID/beige mice with MuLE-luciferase viruses expressing
combinations of shRNA against Cdkn2a, Trp53, and Pten with or without
expression of H-RasG12V. (C) Quantification (mean ± SD) of luminescent
signal intensities over time after injection. ‘Sacrifice of all mice in these
cohorts by this time point. (D) A tumor (arrow) in a mouse injected with
the shCdkn2a plus H-RasG12V MuLE virus only in the right gastrocnemius
muscle. (E) Histological image of the tumor (T) from D surrounded by
muscle tissue (M). (F–K) Representative histology of tumors derived from
injection of shCdkn2a plus H-RasG12V (F and G), shTrp53 plus H-RasG12V
(H and I), and shTrp53 plus shPTEN plus H-RasG12V (J and K) viruses.
Arrowheads in G, I, and K highlight pleomorphic rhobdoid cells. (L) EM
showing an example of a tumor cell with sarcomere formation; M and N
and show higher magnification of the regions in L marked with an arrowhead
and arrows, respectively, showing Z-bands or irregular masses of Z-band
material with converging filaments. (D) Western blot analysis of indepen-
dent cell lines (lanes 1–8) derived from independent tumors of the indi-
cated genotypes. MEFs, muscle tissue, and C2C12 myoblast cells served as
controls. Scale bars: 50 μm (A and F–K); 10 μm (L); 500 nm (M and N).

Methods
Cells. Primary MEFs were isolated from relevant WT or floxed strains
and were frozen in aliquots at passage 2. MEFs were cultured in
DMEM plus 10% FCS in cell-culture incubators supplemented with
5% CO₂ and maintained at 5% O₂. If not stated otherwise, cells were transduced at a MOI between 1 and 2. MEFs were incubated overnight
in virus-containing medium in the presence of 4 μg/ml polybrein (Sig-
ma-Aldrich, no. H9268). Drug selection was performed 48 hours after
transduction using the following concentrations: 3 μg/ml puromycin,
10 μg/ml blasticidin, 500 μg/ml G418. Human melanoma A-375 cells
(ATCC, no. CRL-1619) were cultured in DMEM plus 10% FCS. C2C12 cells were obtained from Sigma-Aldrich (no. 91031101) and cultured in
DMEM plus 15% FCS. HEK293T (ATCC, no. CRL-1619) were cultured in DMEM
plus 10% FCS. NIH3T3 cells (ATCC, no. CRL-1615) were cultured in DMEM plus 10% FCS. LLC-1, B16-F10, and MC-38 cell lines were a gift of Labor Borgis
(Institute of Physiology, University of Zurich) and cultured in DMEM
plus 10% FCS. Mouse embryonic stem cells were a gift of Kurt Bürki
(Institute of Physiology, University of Zurich) and cultured in N2B27+
2i medium. Murine primary kidney epithelial cells were isolated and cultured as described before (53). Endometrial epithelial cells were isolated and cultured as described (54). Murine aortic endothelial
cells and immortalized mouse hepatocytes were gifts of Rok Humar
(Research Unit, Division of Internal Medicine, University Hospital of
Zurich, Zurich, Switzerland) and Nora Rösch (Institute of Molecular
Health Sciences, ETH Zurich, Zurich, Switzerland), respectively, and
were cultured in DMEM plus 10% FCS. Cells were isolated from dis-
sected tumors by digestion for 1 hour at 37°C with 1 mg/ml collagenase
II ( Gibco; Life Technologies), washed twice with PBS, and cultured in DMEM plus 10% FCS. All cells were kept in cell-culture incubators supplemented with 5% CO₂ and maintained at 20% O₂.

Cloning of MuLE vectors. Details of the cloning of all parental
MuLE Entry and Destination vectors, the experimental Entry vec-
tors, and experimental MuLE expression vectors that were used in this
study are provided in Supplemental Methods.

Cloning of sgRNAs, surveyor assays, and next generation sequenc-
ing. Design and cloning of sgRNAs used in this study are described in
Supplemental Methods. For Surveyor assays, 350- to 450-bp fragments
surrounding the sgRNA target site were amplified using proof-reading
PCR (Expand High Fidelity, Roche). Primers and PCR conditions are
described in Supplemental Methods. For mixed cell populations, hetero-
duplexes were identified by reannealing the PCR product alone by heating
to 95°C followed by gradual cooling. Heteroduplexes were digested with
Surveyor nuclease (2 μl heteroduplex, 1 μl Surveyor Enhancer S, 1 μl Sur-
veyor Nuclease S, 1 μl 0.15 M MgCl₂) for 60 minutes at 42°C and subjected to
10% polyacrylamide gel electrophoresis using 1× TBE as running buff-
er. For sequencing of the Trp53, Vhl, and Pten loci in cell clones, gene seg-
ments surrounding the sgRNA-binding site were amplified from genomic
DNA using primers described in Supplemental Methods, followed by liga-
tion to barcoded adapters and pooled sequencing using Ion Torrent PGM.
The full protocol is provided in Supplemental Methods. Sequencing data
were deposited in the NCBI’s BioProject database (ID 272478).

Lentivirus production and titration. Lentivirus was prepared using calcium
phosphate-mediated transfection of subconfluent HEK293T cells cultured in DMEM plus 10% FCS. For a 10-cm dish, lentiviral vector (8 μg) was cotransfected with the lentiviral packaging vec-
tor pSAX2 (Addgene, no. 12260) and either the ecotropic envelope
(pCMV-Eco, Cellbiolabs, no. RV-112) or the amphotropic envelope
(pMD2.G, Addgene, no. 12259). Concentration of lentiviral particles
for in vivo injection was done via centrifugation for 5 hours at
11,000 g over a layer of 20% sucrose (55). In cases in which the viral
construct contained a drug-resistance gene, determination of func-
tional viral titer was performed by drug-resistance colony assay with
NIH3T3 cells as described (56). In cases in which fluorescent reporters
expressed from the constructs, titers were determined using FACS as
described (56). For determination of titers with luciferase as the only
reporter, we generated an EGFP-luciferase-expressing lentiviral vec-
tor and used this to generate a reference curve, which allowed us to
determine the titer of viruses by measuring luciferase activity in cells
that had been transduced with different dilutions of the virus.

Antibodies, Western blotting, and immunohistochemistry. Western
blotting and immunohistochemistry were conducted as described (57)
using antibodies against the following epitopes: H-Ras (Santa Cruz Bio-
technology Inc., no. sc-520), β-actin (Sigma-Aldrich, A2228), p16 (Santa Cruz
Biotechnology Inc., no. sc-1207), p19 (Santa Cruz Biotechnology Inc.,
no. sc-32748), Myc (Sigma-Aldrich, no. M5546), VHL (Santa Cruz
Biotechnology Inc., sc-5575), p53 (Novocastra, NCL-p53-CM5p), PTEN
(Santa Cruz Biotechnology Inc., no. sc-7974), GFP (Life Technologies,
no. G10362), β1 (Cell Signaling, no. 9313), HIF1α (Novus, no. NB100-
479), p21 (FS) (no. sc-6246), phospho-Akt (Ser473) (193H12) (Cell Sig-
pling, no. 4058), and Akt (pan) (C67E7) (Cell Signaling no. 4691).
The full scans of all Western blots are available in the Supplemental Material.

Real-time PCR. Real-time PCR was performed as described (57)
using the following primer pairs: SI2 (5′-GAAGCCTGCAAAAGCTT-
TAGA-3′, 5′-AACGTGAAACACACCTC-3′), Hi5a (5′-TGCT-
CATCGTGGCAGGACCTTC-3′, 5′-CCCTCATGTTCAACATGGATAGA-3′),
Hi5a (5′-GAGGAGGAGGAAATCCCGTG-3′, 5′-CTGATGGCGCACG
CCATGATG-3′), Vgl5a (5′-CTTTGTGCAAGGGAGGGCAAGACG-3′,
5′-ACAATCGAAGATCGTTGAGG-3′, 5′-GGAGCTCCTGAGGCTG
ATAGTC-3′, 5′-CTCCTGAGAAGTCTGGGGG-3′), and
Pgk1 (5′-TCCTGCTGAATCCATTCATCTG-3′, 5′-CAGGCACTTTCTC-
GACTTCTGGG-3′).

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In vivo imaging. Noninvasive in vivo multispectral fluorescence and bioluminescence imaging were performed using the IVIS Spectrum (PerkinElmer) with Living Image software (version 4.4). Mice were anesthetized using a vapor isoflurane inhalation narcosis system. For necrosis initiation, 3% isoflurane (Attane, MINRAD Inc.) in O2, at a flow rate of 3 to 5 μl/min was given via an inhalation mask with the mouse placed on a warming plate. During imaging, the isoflurane levels were reduced to 1% to 2%. For fluorescence time-course measurements of tumor xenografts, mice were shaved and imaged at the indicated time points after cell injection. The filter channels used for calculation of tumor growth with iRFP were Ex.675/Em.725 and for mCherry Ex.570/Em.620. All measurements were performed in epifluorescence mode if not otherwise stated. For the quantification of the total radiant efficiency, a region of interest was drawn around the tumor and total radiant efficiency was measured. For bioluminescence imaging, mice were injected subcutaneously with 150 mg/kg D-luciferin (Caliper, no. 122796) and imaged when peak signal intensities were reached, usually 13 to 17 minutes after injection.

Mice. SCID/beige mutant mice (C.B-17/ScidHsd-Prkdc<sup>scid</sup>Ly5m<sup>h<sup>2</sup></sup>) and C57Bl/6 and BALB/c mice were obtained from Harlan Laboratories and housed under standard conditions. Vh<sup>fl/fl</sup>Trp53fl/fl, Pten<sup>fl/fl</sup>, Hif1a<sup>fl/fl</sup>, and Hif2α<sup>fl/fl</sup> mice have been previously described (53, 58–60). B6.Cg-Gt(ROSA)26Sortm14(CAG-tdTomato)Hze<sup>+/−</sup> mice were obtained from Jackson Laboratory (stock no. 007914). For xenograft experiments, mice were anesthetized and injected subcutaneously with 5 × 10<sup>6</sup> cells suspended in 50% Matrigel (BD, no. 354230). In parallel to in vivo imaging studies, tumor size was measured using a caliper. For intramuscular injections of lentiviruses, 18- to 20-day-old juvenile mice were anesthetized and injected with concentrated ectropic lentivirus into the gastrocnemius muscle.

Statistics. Statistical significance was determined by 2-tailed Student’s t test, with a P value of less than 0.05 considered to be statistically significant.

Study approval. All mouse experiments were approved by the Veterinary Office of the Canton of Zurich under the licences 06/2013, 43/2015 and 137/2013.

Acknowledgments

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