



Emerging Technologies to Create Inducible and Genetically Defined Porcine Cancer Models

Lawrence B. Schook^{1,2*}, Laurie Rund², Karine R. Begnini³, Mariana H. Remião³, Fabiana K. Seixas³ and Tiago Collares^{3*}

¹ University of Illinois Cancer Center, University of Illinois at Chicago, Chicago, IL, USA, ² Department of Animal Sciences, University of Illinois at Urbana-Champaign, Champaign, IL, USA, ³ Postgraduate Program in Biotechnology, Biotechnology Unit, Technology Development Center, Federal University of Pelotas, Pelotas, Brazil

There is an emerging need for new animal models that address unmet translational cancer research requirements. Transgenic porcine models provide an exceptional opportunity due to their genetic, anatomic, and physiological similarities with humans. Due to recent advances in the sequencing of domestic animal genomes and the development of new organism cloning technologies, it is now very feasible to utilize pigs as a malleable species, with similar anatomic and physiological features with humans, in which to develop cancer models. In this review, we discuss genetic modification technologies successfully used to produce porcine biomedical models, in particular the Cre-*loxP* System as well as major advances and perspectives the CRISPR/Cas9 System. Recent advancements in porcine tumor modeling and genome editing will bring porcine models to the forefront of translational cancer research.

Keywords: oncopigs, cancer, biotechnology, transgenesis, genome editing

INTRODUCTION

Animal models have played a central role over the centuries in scientific investigations of human disease and treatment strategies. Genetic strategies for the development of cancer models using human mutations in targeted oncogenic pathways demonstrated that porcine fibroblasts could be transformed *in vitro* and could be tumorigenic with four to six gene alterations (Adam et al., 2007). The authors used retroviral vectors carrying pairs of human and murine oncogenic cDNAs (hTERT and p53^{DD}, cyclin D1 and CDK4^{R24C}, and c-Myc^{T58A}, and H-Ras^{G12V}) to transform porcine fibroblasts. These altered cells showed a transformed phenotype in culture and formed tumors following autologous transfer. These induced changes demonstrated that the pig/tumorigenic pathway recapitulated those observed in human much more closely than murine cells (Adam et al., 2007). Although this approach was limited because the animals needed to be immuno-suppressed for tumors to grow *in vivo*, this work was the first to demonstrate that genetically defined tumors could be induced in a large animal (Schook et al., 2015a).

Recent innovations in reproductive, cloning and transgene technologies have enhanced efficacy and efficiency or producing targeted porcine genome modifications. With the successful cloning of animals by somatic cell nuclear transfer (SCNT), it is now possible to produce genetically modified pigs from genetically engineered somatic donor cells using a wide variety of techniques from random genomic insertion of plasmid DNA (Hyun et al., 2003; Watanabe et al., 2005), to genomic integration of transduced retroviral or lentiviral vectors (Lai et al., 2002; Park et al., 2002), and to modern genome editing with molecular methods using endonucleases such as transposases,

OPEN ACCESS

Edited by:

Ji Luo,
National Institute of Health, USA

Reviewed by:

Christina Stuelten,
National Cancer Institute, USA
Bryan Raymond George Williams,
Monash Institute of Medical
Research, Australia

*Correspondence:

Lawrence B. Schook
schook@illinois.edu;
Tiago Collares
tiago.collares@pq.cnpq.br

Specialty section:

This article was submitted to
Cancer Genetics,
a section of the journal
Frontiers in Genetics

Received: 03 November 2015

Accepted: 13 February 2016

Published: 29 February 2016

Citation:

Schook LB, Rund L, Begnini KR,
Remião MH, Seixas FK and Collares T
(2016) Emerging Technologies
to Create Inducible and Genetically
Defined Porcine Cancer Models.
Front. Genet. 7:28.
doi: 10.3389/fgene.2016.00028

recombinases, and programmable nucleases (Zhou et al., 2015). Genetic modification technologies successfully used to produce porcine biomedical models, in particular the *Cre-loxP* System as well as major advances and perspectives the CRISPR/Cas9 System will be presented in this mini review.

Cre-loxP SYSTEM

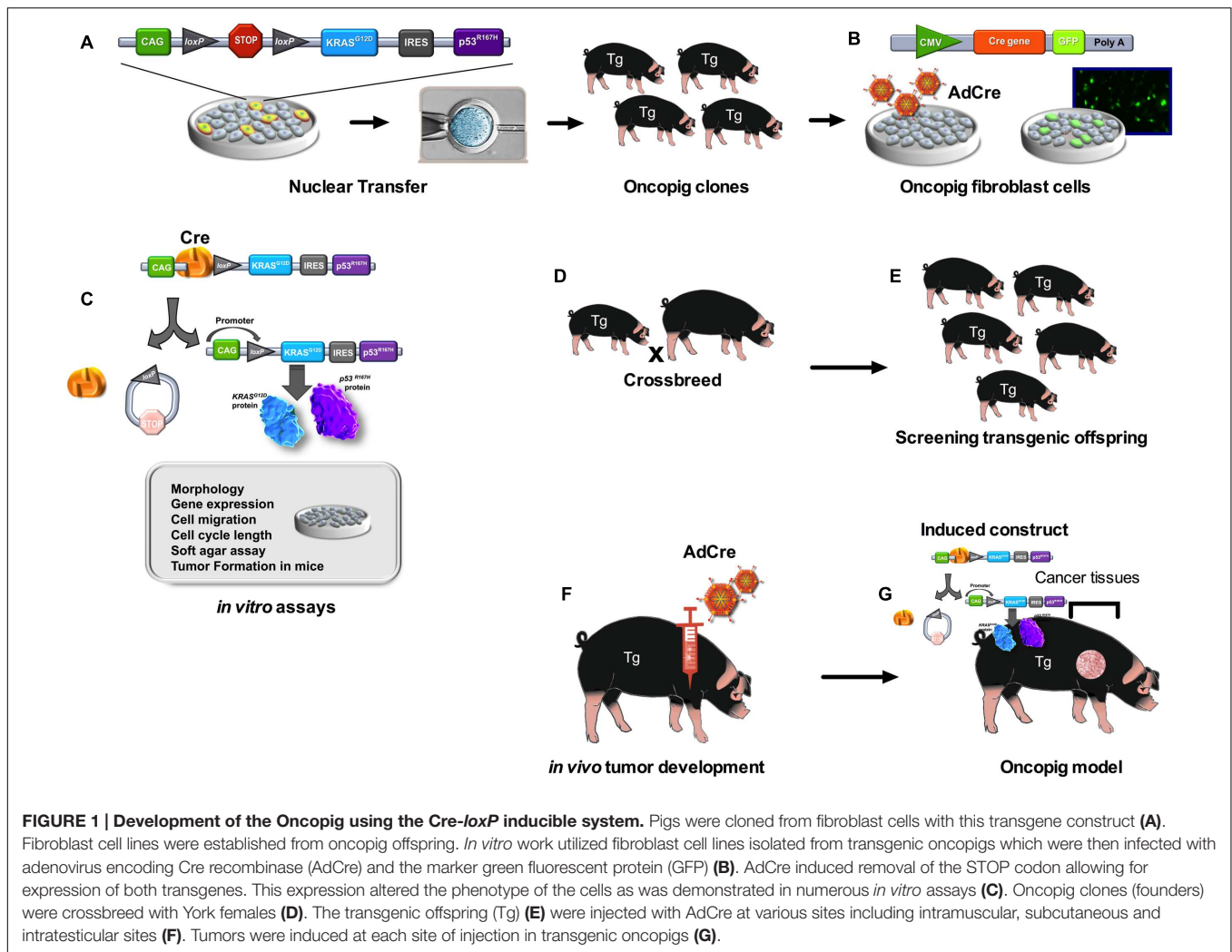
The ability to activate latent genes in defined tissues and at defined times is a key factor to enable the development of inducible temporally and spatially regulated cancer models. The activation of an oncogenic mutation(s) in a chosen tissue could mimic the spontaneous somatic events that initiate many human cancers and enable replication of diverse cancer types using the same mutant gene(s) (Flisikowska et al., 2013). Those conditional gene expressions are now well established in mice using site-specific recombinase (SSR) systems that allow the precise recombination between genomic sites, resulting in deletion or inversion of the intervening sequences (Frese and Tuveson, 2007; Oh-McGinnis et al., 2010). The use of SSR technology in genome manipulation has been demonstrated to effectively resolve complex transgene insertions to single copy, remove unwanted DNA, and precisely insert DNA into known genomic target sites (Wang et al., 2011). Site-specific recombination occurs at a specific sequence or recognition site and involves cleavage and reunion leading to integration, deletion or inversion of a DNA fragment without the gain or loss of nucleotides (Wang et al., 2011). Because of the efficiency of the SSR systems, it can be applied to conditional deletions of relatively short coding sequences or regulatory elements but also to more extensive chromosomal rearrangement strategies (Oh-McGinnis et al., 2010).

Cre-recombinase system is one of the best-studied and most commonly used SSR in mammalian cell cultures. Since its first use for mammalian genome editing in 1988 (Sauer and Henderson, 1988) many adaptations have expanded the utility of the Cre system from flies to mammalian cells beyond mouse to include porcine and humans cell lines (Lanza et al., 2012). Cre-recombinase is derived from the bacteriophage P1 and recognizes a distinct sequence-specific motif termed as recombination target sites (*loxP*) catalyzing efficient conservative DNA rearrangements (Wirth et al., 2007). The *loxP* site is a 34 bp palindromic sequence with an 8-bp asymmetric spacer region (Feng et al., 1999; Siegel et al., 2001; Araki et al., 2002; Sauer, 2002; Schnutgen et al., 2003; Garcia-Otin and Guillou, 2006) and acts upon the neighboring DNA sequences. The *Cre-loxP* system is a bidirectional tyrosine recombinase that enables the recombinase-mediated genetic cross-over between two identical *loxP* recognition sites promoting intermolecular or intramolecular recombination. Intermolecular recombination is a translocation between two DNA fragments with corresponding *loxP* sites, while the intramolecular recombination involves removal of genetic material between two *loxP* sites, with the last one been the preferred function of Cre-recombinase (Feng et al., 1999). Because of the identical nature of the recognition sites, the recombination reaction is fully reversible, although

intramolecular recombination (excision) is highly favored over intermolecular reactions (integration) (Wang et al., 2011).

One of the most powerful and widely used applications of the *Cre-loxP* system is in conditional gene expression (Gu et al., 1994). This strategy allows for tissue and time-specific gene expression when recombination is triggered by Cre-recombinase, and is even more important in cancer models where oncogenic activation in a chosen tissue could mimic the spontaneous somatic events that initiate many human cancers (Schook et al., 2015a). Endogenous engineered mice are usually conditional alleles constructed by the insertion of a transcriptional and translational *LoxStopLox* 'stop' cassette between the promoter and first coding exon of the oncogenic allele. Providing the expression of an active Cre-recombinase, the stop cassette is excised and the mutant oncogene is subsequently expressed (de Alboran et al., 2001; Jackson et al., 2001). In pigs, this conditional gene expression strategy has been used to promote oncogenic expression in three cancer models (Leuchs et al., 2012; Li et al., 2015; Schook et al., 2015b). Leuchs et al. (2012) have generated gene-targeted pigs with a conditionally activated oncogenic mutant form of p53, which in latent form is a gene knockout. The construction used a porcine BAC vector with CAGGS promoter-mCherry cassette (in reverse orientation) as a fluorescent counter-selectable marker; a short arm of homology corresponding to a region of TP53 intron 1 from a point of exon 2 to a *PmlI* restriction enzyme site of exon 2; a floxed transcriptional termination cassette (LSL); and a region extending from the *PmlI* site in intron 1 to a point of exon 11 that includes a G to A substitution in exon 5 changing arginine to histidine in codon 167 (R167H) (Leuchs et al., 2012). In this same model, viable gene-targeted pigs carrying a latent *Kras*^{G12D} mutant allele that could be activated by Cre-recombinase was constructed (Li et al., 2015). The KRAS-neo vector comprised: a short homology arm in KRAS intron 1; a transcriptional stop cassette comprising: a *loxP* site; adenoviral splice acceptor; promoterless neomycin phosphotransferase resistance gene (*neo*); three poly-adenylation signals derived from SV40, bovine growth hormone and cytomegalovirus; and a second *loxP* site inserted into a *Clal* site in KRAS intron 1; and a region of porcine KRAS extending from the *Clal* site in intron 1 to a *SaclI* site in intron 2, which also included an engineered G to A point mutation within exon 2 that results in a glycine to aspartic acid substitution at codon 12 (G12D) (Li et al., 2015). Both KRAS and TP53 transgenic pigs cells were transduced with 5 μ M of Cre protein produced *in vitro* with the vector pTriEx-HTNC (Addgene plasmid 13763; Leuchs et al., 2012; Li et al., 2015).

Transgenic oncopigs (Figure 1) have also been engineered to contain oncogenic *Kras*^{G12D} and dominant-negative p53^{R167H} downstream of a *LoxP*-polyA(STOP)-*LoxP* sequence (LSL) and CAG promoter (Schook et al., 2015b). Site-directed mutagenesis was then used to introduce the oncogenic G12D mutation into the porcine KRAS cDNA and the R167H mutation was chosen for TP53 as its human equivalent (R175H) is commonly found in human cancers as well as the cancer predisposition Li-Fraumeni Syndrome. These two cDNAs were then introduced into a Cre-inducible vector, followed by the aforementioned LSL sequence,



$KRAS^{G12D}$, an IRES sequence to allow for bicistronic expression, $TP53^{R167H}$ and a poly A sequence. This design allows for co-expression of both $KRAS^{G12D}$ and $TP53^{R167H}$ in ostensibly any cells of the pig by transient expression of AdCre (Ad5CMVCre-GFP, AdGFP, Gene Transfer Vector Core; Schook et al., 2015b). These pig models have resulted in tumorigenic profiles *in vitro* (Leuchs et al., 2012; Li et al., 2015) and *in vivo* (Schook et al., 2015b) and the results obtained with these three cancer pig models are shown in Table 1.

CRISPR/Cas9 SYSTEM

The discovering of molecules that recognize specific sequences of DNA was one of the most important advances in gene editing technology allowing site specific genetic modifications to be made. These DNA binding proteins include the zinc fingers and transcriptional activator-like effector (TALE; Wood et al., 2011; Gaj et al., 2013). When they are fused to nucleases, they generate a double-strand break (DSB) in the DNA at the desired genomic loci, triggering the endogenous DNA repair machinery (Gaj

et al., 2013; Zhu et al., 2014); if fused to transcription factors or inhibitor molecules, they can bind to promoter regions of target genes, modulating gene expression (Gilbert et al., 2014; Kearns et al., 2014). However, there is a disadvantage of utilizing these proteins that interact with DNA: production of these proteins involves a complicated and more expensive assembly process (Pan et al., 2014).

An easier, cheaper, and yet highly efficient tool for directed genome editing appeared to be more worthwhile and profitable than proteins: the CRISPR (clustered regularly interspaced short palindromic repeat)/Cas (CRISPR associated proteins) system. This system is simpler than zinc fingers and TALEs because the CRISPR/Cas system uses the RNA-DNA interaction for genome loci recognition, which is more specific than protein-DNA (Gasiunas and Siksnys, 2013; Pan et al., 2014).

CRISPR/Cas system has been recently discovered as an adaptive immune system of some bacteria and archaea and protects them against invading viruses and plasmids (Barrangou et al., 2007). The transcription of the repeat-spacer elements from CRISPR locus generates a precursor non-coding CRISPR RNA (pre-crRNA) that later will be cleaved to have short CRISPR

TABLE 1 | Porcine models for cancer developed by different strategies.

Author	Gene	Technique	Inducible strategy	Survival	Location	Phenotype	Tumor progression
Yamakawa et al., 1999	<i>v-Ha-ras</i>	Pronuclear microinjection	No	-	-	No	No
McCalla-Martin et al., 2010	<i>Gli2</i>	DNA vector + SCNT	No	Euthanized due to bacterial infection	-	Not informed	Not informed
Luo et al., 2011	<i>BRCA-1 (KO)</i>	Adenovirus + SNCT and HMC	No	18 days	Breast	Not informed	Not informed
Flisikowska et al., 2013	<i>APC</i>	DNA vector + SCNT	No	At least 1 year	Intestinal polyps	Yes	Yes
Leuchs et al., 2012	<i>TP53</i>	DNA vector + SNCT	Cre-recombination	Not informed	-	Yes	Not yet
Sieren et al., 2014	<i>TP53</i>	rAAV + SCNT	No	Not informed	Several locations	Yes	Yes
Li et al., 2015	<i>Kras</i>	DNA vector + SNCT	Cre-recombination	Yes	-	Not informed	Not informed
Schook et al., 2015b	<i>TP53 + Kras</i>	DNA vector + SNCT	Cre-recombination	Yes	Several locations	Yes	Yes

SCNT, Somatic Cell Nuclear Transfer; HMC, Handmade Cloning.

RNAs (crRNA) (Garneau et al., 2010; Jinek et al., 2012). The crRNA will be homologous to the DNA or RNA from foreign sequences, and when the invasion occurs, the crRNA will be directed just by Watson-Crick base pairing (Jinek et al., 2012; Wade, 2015). There are different types of CRISPR systems in different organisms (I–III), and the one that have been most developed as a new tool for genome editing, the CRISPR/Cas9 system, is the type II CRISPR originating from *Streptococcus pyogenes* SF370 (Jinek et al., 2012; Qi et al., 2013). The type II is different from types I and III, that crRNA hybridize with another RNA molecule, the *trans*-activating crRNA (tracrRNAs), to direct Cas9 protein to specific DNA sequences (Jinek et al., 2012; Mali et al., 2013; Doudna and Charpentier, 2014). For genome editing, the researchers created a single chimeric guide RNAs (sgRNA), which is a fusion of a precursor crRNA and a transactivating crRNA (tracrRNA) (Jinek et al., 2012; Pan et al., 2014). Beyond the polymerization, the genome sequence from invader has a complementary genome sequence containing a tri-nucleotide protospacer adjacent motif (PAM) that will be required for initial binding of Cas9 protein (Guilinger et al., 2014). Cas9 protein has an endonuclease activity that cleaves on both strands a few nucleotides away from the PAM generating DSB, preventing the invader genome translation (Jinek et al., 2012). This has been used to generate knockin and knockout transgenic animals, as the DSB activates the endogenous DNA repair machinery by non-homologous joining (Ma et al., 2014; Flemr and Buhler, 2015; Yang, 2015; Zhu et al., 2015).

However, the study of CRISPR/Cas9 identified a new application for Cas9: without its nuclease activity, Cas9 protein, attached to a molecule that modulates gene expression, could bind to the promoter region of some gene of interest, changing the genic expression pattern (Qi et al., 2013). The catalytically dead Cas9 (dCas9), lacking endonuclease activity, contains two mutations in the nuclease domains (D10A and H840A) (Choudhary et al., 2015). Since dCas9 was reported, new studies have been described using it for genome regulation creating different segments to use this tool: CRISPRi, for gene interference, and CRISPRa, for activation of gene translation. When these strategies uses an effector domain attached to dCas9, it can be called CRISPRe. For gene interference (CRISPRi), dCas9 recognizes sgRNA attached to the promoter region of target gene, impairing transcription (Qi et al., 2013). However, this strategy is not efficient for gene repression in eukaryotic cells, so dCas9 can be fused to a transcription repression domain to enhance gene knockdown (Gilbert et al., 2013). The most described strategy for CRISPRi is dCas9 fused to a KRAB (Krüppel-associated box domain of Kox1), a repressive chromatin modifier domain, which have been demonstrating increased gene expression repression in relation to dCas9 alone (Gilbert et al., 2013, 2014). Some authors mention that CRISPRi can be an alternative strategy to RNAi for repressing gene expression in mammalian cells (Gilbert et al., 2013).

Another approach for using dCas9 is fused to transcriptional activator domains, which can be called CRISPRa (Gilbert et al., 2014) or CRISPR-on system (Cheng et al., 2013a), to induce expression of target genes. To achieve that, dCas9 fused to the transcriptional activator is guided by the sgRNA complementary

to the promoter region of the gene. The well-characterized tetramer of herpes simplex virus protein, VP16 (VP64) is one of the most reported transcription activator attached to dCas9 and it has been shown to induce gene expression in eukaryotic cells, including human cells (Gilbert et al., 2013, 2014; Maeder et al., 2013; Perez-Pinera et al., 2013; Kearns et al., 2014). Some studies also report that target genes can be simultaneously artificially activated by just adding complementary sgRNAs of promoters of each one of the interest genes (Cheng et al., 2013b; Maeder et al., 2013). This strategy has been tested in human and mouse transformed cells, as well as in ES cells, in one-cell embryo (Cheng et al., 2013b).

The use of CRISPR/Cas9 strategy to build an animal for model of cancer disease is a recently developed approach. For lung adenocarcinoma, Maddalo et al. (2014) describe a methodology of *in vivo* chromosomal rearrangement using CRISPR/Cas9 delivered by virus infection. Rearranging chromosomes by fusing EML4 and ALK genes generated a new murine model for lung adenocarcinoma. An *in vivo* somatic cancer mutation in adult animals was described by Xue et al. (2014), which they developed a different strategy using a hydrodynamic delivery of plasmids with CRISPR components that occasioned to efficient hepatocyte transfection to edit oncogenes and suppressor-tumor genes.

Most frequently, rodents are used to test new strategies for genome editing with CRISPR/Cas9 system to develop cancer and other biomedical models of human disease. However, a new strategy for enrichment of cells with chromosomal deletions made by CRISPR/Cas9 to generate cancer genotype was developed in porcine embryonic fibroblasts (He et al., 2015). For employment in xenotransplants, CRISPR/Cas9 technology has already been applied to inactivate porcine endogenous retroviruses in porcine kidney epithelial cell line (Yang et al., 2015).

Not only modifications in genome sequence can induce cancer phenotype, epigenetic modifications can also be a target to develop animal models for cancer. Falahi et al. (2015) supposes that dCas9 can contribute for epigenome engineering to develop animals for cancer study. Effector domains attached to dCas9 could generate epigenetic mutations known to evolve to different cancer types. Also using dCas9, attached or not to KRAB domain, initial studies in human cells HEK293 and HEK293T, showed repression of TP53 (Lawhorn et al., 2014).

The recent advances generated by CRISPR/Cas9 system in genome editing are extremely important for development of new strategies to generate animal models of cancer. The simplicity, low cost, and low off-target effects put this strategy as one alternative not only for ZFN and TALEN, but also for RNAi technology and *Cre-loxP* systems.

REFERENCES

- Adam, S. J., Rund, L. A., Kuzmuk, K. N., Zachary, J. F., Schook, L. B., and Counter, C. M. (2007). Genetic induction of tumorigenesis in swine. *Oncogene* 26, 1038–1045. doi: 10.1038/sj.onc.1209892
- Araki, K., Araki, M., and Yamamura, K. (2002). Site-directed integration of the cre gene mediated by Cre recombinase using a combination of mutant lox sites. *Nucleic Acids Res.* 30:e103. doi: 10.1093/nar/gnf102

PERSPECTIVES

To unite *Cre-loxP* and CRISPR/Cas9 system has been a promising approach to develop animal models for cancer. *Cre-loxP* affords to conditional gene expression, while CRISPR/Cas9 can be used for target gene insertion and also for gene expression regulation. Some promising works already showed how these technologies can be used together. Using *Cre-loxP* system for induced expression, Sánchez-Rivera et al. (2014) used a system with CRISPR/Cas9 and Cre recombinase to evaluate new candidates for cancer genome, developing adenocarcinoma by editing tumor-suppressor genes sequences in mice models. A different association of both techniques is a study that a mouse model had Cas9 expressed by Cre dependence, and when expressed in conjunction with sgRNAs for Kras, p53, and LKB1 genes, it generated a change of function of those proteins, taking to macroscopic tumors of adenocarcinoma pathology (Platt et al., 2014). Probably, the next step is to standardize those techniques and employ them for a next-generation models for human cancer (Sanchez-Rivera and Jacks, 2015), and pigs fits for those purpose.

AUTHOR CONTRIBUTIONS

TC: acquisition of data, data analysis/interpretation, drafting of the manuscript and figure; FS: acquisition of data, data analysis/interpretation, drafting of the manuscript; KB: acquisition of data, data analysis/interpretation, drafting of the manuscript and table; MR: acquisition of data, data analysis/interpretation, drafting of the manuscript; LR: critical revision of the manuscript, drafting of the manuscript and figure; LS: critical revision of the manuscript. All authors approved the manuscript.

FUNDING

TC, FS, and KB were supported by the Brazilian Scholarship Program “Science Without Borders” promoted by the National Counsel of Technological and Scientific Development. LS was supported by USDA (AG 58-5438-2-307), the U.S. National Institutes of Health (was supported by USDA (AG 58-5438-2-307), the U.S. National Institutes of Health (CA 153132), the Edward William and Jane Marr Gutgsell Foundation, and the Cooperative Research Program for Agriculture Science and Technology Development (No. PJ009103), Rural Development Administration, Republic of Korea.

- Barrangou, R., Fremaux, C., Deveau, H., Richards, M., Boyaval, P., Moineau, S., et al. (2007). CRISPR provides acquired resistance against viruses in prokaryotes. *Science* 315, 1709–1712. doi: 10.1126/science.1138140
- Cheng, A. W., Wang, H., Yang, H., Shi, L., Katz, Y., Theunissen, T. W., et al. (2013a). Multiplexed activation of endogenous genes by CRISPR-on, an RNA-guided transcriptional activator system. *Cell Res.* 23, 1163–1171. doi: 10.1038/cr.2013.122

- Cheng, E. H., Chen, S. U., Lee, T. H., Pai, Y. P., Huang, L. S., Huang, C. C., et al. (2013b). Evaluation of telomere length in cumulus cells as a potential biomarker of oocyte and embryo quality. *Hum. Reprod.* 28, 929–936. doi: 10.1093/humrep/det004
- Choudhary, E., Thakur, P., Pareek, M., and Agarwal, N. (2015). Gene silencing by CRISPR interference in mycobacteria. *Nat. Commun.* 6, 62–67. doi: 10.1038/ncomms7267
- de Alboran, I. M., O'hagan, R. C., Gartner, F., Malynn, B., Davidson, L., Rickert, R., et al. (2001). Analysis of C-MYC function in normal cells via conditional gene-targeted mutation. *Immunity* 14, 45–55. doi: 10.1016/S1074-7613(01)00088-7
- Doudna, J. A., and Charpentier, E. (2014). Genome editing. The new frontier of genome engineering with CRISPR-Cas9. *Science* 346:1258096. doi: 10.1126/science.1258096
- Falahi, F., Sgro, A., and Blancafort, P. (2015). Epigenome engineering in cancer: fairytale or a realistic path to the clinic? *Front. Oncol.* 5:22. doi: 10.3389/fonc.2015.00022
- Feng, Y. Q., Seibler, J., Alami, R., Eisen, A., Westerman, K. A., Leboulch, P., et al. (1999). Site-specific chromosomal integration in mammalian cells: highly efficient CRE recombinase-mediated cassette exchange. *J. Mol. Biol.* 292, 779–785. doi: 10.1006/jmbi.1999.3113
- Flemer, M., and Buhler, M. (2015). Single-step generation of conditional knockout mouse embryonic stem cells. *Cell Rep.* 12, 709–716. doi: 10.1016/j.celrep.2015.06.051
- Flisikowska, T., Kind, A., and Schnieke, A. (2013). The new pig on the block: modelling cancer in pigs. *Transgenic Res.* 22, 673–680. doi: 10.1007/s11248-013-9720-9
- Frese, K. K., and Tuveson, D. A. (2007). Maximizing mouse cancer models. *Nat. Rev. Cancer* 7, 645–658. doi: 10.1038/nrc2192
- Gaj, T., Gersbach, C. A., and Barbas, C. F. 3rd. (2013). ZFN, TALEN, and CRISPR/Cas-based methods for genome engineering. *Trends Biotechnol.* 31, 397–405. doi: 10.1016/j.tibtech.2013.04.004
- Garcia-Otin, A. L., and Guillouf, F. (2006). Mammalian genome targeting using site-specific recombinases. *Front. Biosci.* 11:1108–1136. doi: 10.2741/1867
- Garneau, J. E., Dupuis, M. E., Villion, M., Romero, D. A., Barrangou, R., Boyaval, P., et al. (2010). The CRISPR/Cas bacterial immune system cleaves bacteriophage and plasmid DNA. *Nature* 468, 67–71. doi: 10.1038/nature09523
- Gasiunas, G., and Siksnys, V. (2013). RNA-dependent DNA endonuclease Cas9 of the CRISPR system: holy Grail of genome editing? *Trends Microbiol.* 21, 562–567. doi: 10.1016/j.tim.2013.09.001
- Gilbert, L. A., Horlbeck, M. A., Adamson, B., Villalta, J. E., Chen, Y., Whitehead, E. H., et al. (2014). Genome-scale CRISPR-mediated control of gene repression and activation. *Cell* 159, 647–661. doi: 10.1016/j.cell.2014.09.029
- Gilbert, L. A., Larson, M. H., Morsut, L., Liu, Z., Brar, G. A., Torres, S. E., et al. (2013). CRISPR-mediated modular RNA-guided regulation of transcription in eukaryotes. *Cell* 154, 442–451. doi: 10.1016/j.cell.2013.06.044
- Gu, H., Marth, J. D., Orban, P. C., Mossmann, H., and Rajewsky, K. (1994). Deletion of a DNA polymerase beta gene segment in T cells using cell type-specific gene targeting. *Science* 265, 103–106. doi: 10.1126/science.8016642
- Guilinger, J. P., Thompson, D. B., and Liu, D. R. (2014). Fusion of catalytically inactive Cas9 to FokI nuclease improves the specificity of genome modification. *Nat. Biotechnol.* 32, 577–582. doi: 10.1038/nbt.2909
- He, Z., Shi, X., Du, B., Qin, Y., Cong, P., and Chen, Y. (2015). Highly efficient enrichment of porcine cells with deletions induced by CRISPR/Cas9 using dual fluorescence selection. *J. Biotechnol.* 214, 69–74. doi: 10.1016/j.jbiotec.2015.07.011
- Hyun, S., Lee, G., Kim, D., Kim, H., Lee, S., Nam, D., et al. (2003). Production of nuclear transfer-derived piglets using porcine fetal fibroblasts transfected with the enhanced green fluorescent protein. *Biol. Reprod.* 69, 1060–1068. doi: 10.1095/biolreprod.102.014886
- Jackson, E. L., Willis, N., Mercer, K., Bronson, R. T., Crowley, D., Montoya, R., et al. (2001). Analysis of lung tumor initiation and progression using conditional expression of oncogenic K-ras. *Genes Dev.* 15, 3243–3248. doi: 10.1101/gad.943001
- Jinek, M., Chylinski, K., Fonfara, L., Hauer, M., Doudna, J. A., and Charpentier, E. (2012). A programmable dual-RNA-guided DNA endonuclease in adaptive bacterial immunity. *Science* 337, 816–821. doi: 10.1126/science.1225829
- Kearns, N. A., Genga, R. M., Enuameh, M. S., Garber, M., Wolfe, S. A., and Maehr, R. (2014). Cas9 effector-mediated regulation of transcription and differentiation in human pluripotent stem cells. *Development* 141, 219–223. doi: 10.1242/dev.103341
- Lai, L., Park, K. W., Cheong, H. T., Kuhhholzer, B., Samuel, M., Bonk, A., et al. (2002). Transgenic pig expressing the enhanced green fluorescent protein produced by nuclear transfer using colchicine-treated fibroblasts as donor cells. *Mol. Reprod. Dev.* 62, 300–306. doi: 10.1002/mrd.10146
- Lanza, A. M., Dyess, T. J., and Alper, H. S. (2012). Using the Cre/lox system for targeted integration into the human genome: loxFAS-loxP pairing and delayed introduction of Cre DNA improve gene swapping efficiency. *Biotechnol. J.* 7, 898–908. doi: 10.1002/biot.201200034
- Lawhorn, I. E., Ferreira, J. P., and Wang, C. L. (2014). Evaluation of sgRNA target sites for CRISPR-mediated repression of TP53. *PLoS ONE* 9:e113232. doi: 10.1371/journal.pone.0113232
- Leuchs, S., Saalfrank, A., Merkl, C., Flisikowska, T., Edlinger, M., Durkovic, M., et al. (2012). Inactivation and inducible oncogenic mutation of p53 in gene targeted pigs. *PLoS ONE* 7:e43323. doi: 10.1371/journal.pone.0043323
- Li, S., Edlinger, M., Saalfrank, A., Flisikowski, K., Tschukes, A., Kurome, M., et al. (2015). Viable pigs with a conditionally-activated oncogenic KRAS mutation. *Transgenic Res.* 24, 509–517. doi: 10.1007/s11248-015-9866-8
- Luo, Y., Li, J., Liu, Y., Lin, L., Du, Y., Li, S., et al. (2011). High efficiency of BRCA1 knockout using rAAV-mediated gene targeting: developing a pig model for breast cancer. *Transgenic Res.* 20, 975–988. doi: 10.1007/s11248-010-9472-8
- Ma, Y., Ma, J., Zhang, X., Chen, W., Yu, L., Lu, Y., et al. (2014). Generation of eGFP and Cre knockin rats by CRISPR/Cas9. *FEBS J.* 281, 3779–3790. doi: 10.1111/febs.12935
- Maddalo, D., Manchado, E., Concepcion, C. P., Bonetti, C., Vidigal, J. A., Han, Y. C., et al. (2014). In vivo engineering of oncogenic chromosomal rearrangements with the CRISPR/Cas9 system. *Nature* 516, 423–427. doi: 10.1038/nature13902
- Maeder, M. L., Linder, S. J., Cascio, V. M., Fu, Y., Ho, Q. H., and Joung, J. K. (2013). CRISPR RNA-guided activation of endogenous human genes. *Nat. Methods* 10, 977–979. doi: 10.1038/nmeth.2598
- Mali, P., Yang, L., Esvelt, K. M., Aach, J., Guell, M., Dicarlo, J. E., et al. (2013). RNA-guided human genome engineering via Cas9. *Science* 339, 823–826. doi: 10.1126/science.1232033
- McCalla-Martin, A. C., Chen, X., Linder, K. E., Estrada, J. L., and Piedrahita, J. A. (2010). Varying phenotypes in swine versus murine transgenic models constitutively expressing the same human Sonic hedgehog transcriptional activator, *K5-HGLI2ΔN*. *Transgenic Res.* 19, 869–887. doi: 10.1007/s11248-010-9362-0
- Oh-McGinnis, R., Jones, M. J., and Lefebvre, L. (2010). Applications of the site-specific recombinase Cre to the study of genomic imprinting. *Brief Funct. Genomics* 9, 281–293. doi: 10.1093/bfpg/elq017
- Pan, A., Weintraub, N. L., and Tang, Y. (2014). Enhancing stem cell survival in an ischemic heart by CRISPR-dCas9-based gene regulation. *Med. Hypoth.* 83, 702–705. doi: 10.1016/j.mehy.2014.09.022
- Park, K. W., Lai, L., Cheong, H. T., Cabot, R., Sun, Q. Y., Wu, G., et al. (2002). Mosaic gene expression in nuclear transfer-derived embryos and the production of cloned transgenic pigs from ear-derived fibroblasts. *Biol. Reprod.* 66, 1001–1005. doi: 10.1095/biolreprod66.4.1001
- Perez-Pinera, P., Kocak, D. D., Vockley, C. M., Adler, A. F., Kabadi, A. M., Polstein, L. R., et al. (2013). RNA-guided gene activation by CRISPR-Cas9-based transcription factors. *Nat. Methods* 10, 973–976. doi: 10.1038/nmeth.2600
- Platt, R. J., Chen, S., Zhou, Y., Yim, M. J., Swiech, L., Kempton, H. R., et al. (2014). CRISPR-Cas9 knockin mice for genome editing and cancer modeling. *Cell* 159, 440–455. doi: 10.1016/j.cell.2014.09.014
- Qi, L. S., Larson, M. H., Gilbert, L. A., Doudna, J. A., Weissman, J. S., Arkin, A. P., et al. (2013). Repurposing CRISPR as an RNA-guided platform for sequence-specific control of gene expression. *Cell* 152, 1173–1183. doi: 10.1016/j.cell.2013.02.022
- Sanchez-Rivera, F. J., and Jacks, T. (2015). Applications of the CRISPR-Cas9 system in cancer biology. *Nat. Rev. Cancer* 15, 387–395. doi: 10.1038/nrc3950
- Sánchez-Rivera, F. J., Papagiannakopoulos, T., Romero, R., Tammela, T., Bauer, M. R., Bhutkar, A., et al. (2014). Rapid modelling of cooperating genetic events in cancer through somatic genome editing. *Nature* 516, 428–431. doi: 10.1038/nature13906

- Sauer, B. (2002). Cre/lox: one more step in the taming of the genome. *Endocrine* 19, 221–228. doi: 10.1385/ENDO:19:3:221
- Sauer, B., and Henderson, N. (1988). Site-specific DNA recombination in mammalian cells by the Cre recombinase of bacteriophage P1. *Proc. Natl. Acad. Sci. U.S.A.* 85, 5166–5170. doi: 10.1073/pnas.85.14.5166
- Schnutgen, F., Doerflinger, N., Calleja, C., Wendling, O., Chambon, P., and Ghyselinck, N. B. (2003). A directional strategy for monitoring Cre-mediated recombination at the cellular level in the mouse. *Nat. Biotechnol.* 21, 562–565. doi: 10.1038/nbt811
- Schook, L. B., Collares, T. V., Darfour-Oduro, K. A., De, A. K., Rund, L. A., Schachtschneider, K. M., et al. (2015a). Unraveling the swine genome: implications for human health. *Annu. Rev. Anim. Biosci.* 3, 219–244. doi: 10.1146/annurev-animal-022114-110815
- Schook, L. B., Collares, T. V., Hu, W., Liang, Y., Rodrigues, F. M., Rund, L. A., et al. (2015b). A genetic porcine model of cancer. *PLoS ONE* 10:e0128864. doi: 10.1371/journal.pone.0128864
- Siegel, R. W., Jain, R., and Bradbury, A. (2001). Using an in vivo phagemid system to identify non-compatible loxP sequences. *FEBS Lett.* 499, 147–153. doi: 10.1016/S0014-5793(01)02541-8
- Sieren, J. C., Meyerholz, D. K., Wang, X. J., Davis, B. T., Newell, J. D. Jr., Hammond, E., et al. (2014). Development and translational imaging of a TP53 porcine tumorigenesis model. *J. Clin. Invest.* 124, 4052–4066. doi: 10.1172/JCI75447
- Wade, M. (2015). High-throughput silencing using the CRISPR-Cas9 system: a review of the benefits and challenges. *J. Biomol. Screen.* 20, 1027–1039. doi: 10.1177/1087057115587916
- Wang, Y., Yau, Y. Y., Perkins-Balding, D., and Thomson, J. G. (2011). Recombinase technology: applications and possibilities. *Plant Cell Rep.* 30, 267–285. doi: 10.1007/s00299-010-0938-1
- Watanabe, S., Iwamoto, M., Suzuki, S., Fuchimoto, D., Honma, D., Nagai, T., et al. (2005). A novel method for the production of transgenic cloned pigs: electroporation-mediated gene transfer to non-cultured cells and subsequent selection with puromycin. *Biol. Reprod.* 72, 309–315. doi: 10.1095/biolreprod.104.031591
- Wirth, D., Gama-Norton, L., Riemer, P., Sandhu, U., Schucht, R., and Hauser, H. (2007). Road to precision: recombinase-based targeting technologies for genome engineering. *Curr. Opin. Biotechnol.* 18, 411–419. doi: 10.1016/j.copbio.2007.07.013
- Wood, A. J., Lo, T. W., Zeitler, B., Pickle, C. S., Ralston, E. J., Lee, A. H., et al. (2011). Targeted genome editing across species using ZFNs and TALENs. *Science* 333:307. doi: 10.1126/science.1207773
- Xue, W., Chen, S., Yin, H., Tammela, T., Papagiannakopoulos, T., Joshi, N. S., et al. (2014). CRISPR-mediated direct mutation of cancer genes in the mouse liver. *Nature* 514, 380–384. doi: 10.1038/nature13589
- Yamakawa, H., Nagai, T., Harasawa, R., Yamagami, T., Takahashi, J., Ishikawa, K., et al. (1999). Production of Transgenic Pig Carrying MMTV/v-Ha-ras. *J. Reprod. Dev.* 45, 111–118. doi: 10.1262/jrd.45.111
- Yang, L., Guell, M., Niu, D., George, H., Lesha, E., Grishin, D., et al. (2015). Genome-wide inactivation of porcine endogenous retroviruses (PERVs). *Science* 350, 1101–1104. doi: 10.1126/science.aad1191
- Yang, X. (2015). Applications of CRISPR-Cas9 mediated genome engineering. *Mil Med. Res.* 2:11. doi: 10.1186/s40779-015-0038-1
- Zhou, X., Xin, J., Fan, N., Zou, Q., Huang, J., Ouyang, Z., et al. (2015). Generation of CRISPR/Cas9-mediated gene-targeted pigs via somatic cell nuclear transfer. *Cell Mol. Life Sci.* 72, 1175–1184. doi: 10.1007/s00018-014-1744-7
- Zhu, Z., Gonzalez, F., and Huangfu, D. (2014). The iCRISPR platform for rapid genome editing in human pluripotent stem cells. *Methods Enzymol.* 546, 215–250. doi: 10.1016/B978-0-12-801185-0.00011-8
- Zhu, Z., Verma, N., Gonzalez, F., Shi, Z. D., and Huangfu, D. (2015). A CRISPR/Cas-mediated selection-free knockin strategy in human embryonic stem cells. *Stem Cell Rep.* 4, 1103–1111. doi: 10.1016/j.stemcr.2015.04.016

Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

The reviewer CS and handling Editor declared their shared affiliation, and the handling Editor states that the process nevertheless met the standards of a fair and objective review.

Copyright © 2016 Schook, Rund, Begnini, Remião, Seixas and Collares. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) or licensor are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.