The robust, high-throughput, and temporally regulated roxCre and loxCre reporting systems for genetic modifications *in vivo*

Mengyang Shi[1,5], Jie Li[1,5], Xiuxiu Liu[1], Kuo Liu[2], Wenjuan Pu[1], Wendong Weng[1], Shaohua Zhang[1], Huan Zhao[1], Kathy O. Lui[3], Bin Zhou[1,2,4,6,*]

[1]New Cornerstone Science Laboratory, State Key Laboratory of Cell Biology, CAS Center for Excellence in Molecular Cell Science, Shanghai Institute of Biochemistry and Cell Biology, Chinese Academy of Sciences, University of Chinese Academy of Sciences, Shanghai, China

[2]Key Laboratory of Systems Health Science of Zhejiang Province, School of Life Science, Hangzhou Institute for Advanced Study, University of Chinese Academy of Sciences, Hangzhou, Chinace

[3]Department of Chemical Pathology; and Li Ka Shing Institute of Health Sciences, The Chinese University of Hong Kong, Prince of Wales Hospital, Hong Kong, China

[4]School of Life Science and Technology, ShanghaiTech University, Shanghai, China

[5]These authors contributed equally

[6]Lead contact

[*]Correspondence: zhoubin@sibs.ac.cn

Abstract

Cre-loxP technology, a cornerstone in fate mapping and *in vivo* gene function studies, faces challenges in achieving precise and efficient conditional mutagenesis through inducible systems. In this study, we introduce two innovative genetic tools designed to overcome these limitations. The first, loxCre, facilitates conditional gene targeting by allowing any CreER line to induce Cre expression with significantly enhanced efficiency. The second, roxCre, enables DreER-mediated Cre release, paving the way for intersectional genetic manipulation that permits both increased precision and efficiency. Both tools incorporate a fluorescent reporter for genetic lineage tracing, revealing efficient gene knockout in cells marked by the reporter simultaneously. These strategies hold great potential for precise and efficient exploration of lineage-specific gene functions, marking a significant advancement in genetic research methodologies.

Introduction

Genetic modification such as gene knockout in a precise spatiotemporal manner forms the basis for understanding cellular and molecular mechanisms in multiple biological processes[1, 2]. The powerful Cre-loxP recombination system is the most widely used approach employed to knockout or over-express specific genes *in vivo*[3]. Cre recombinase, derived from bacteriophage P1 gene, targets short palindrome DNA sequence loxP (34bp) and recombines two loxP-flanked sequences for deletion[4, 5]. Application of the Cre-loxP system in mouse genetics has revolutionized gene functional analysis and significantly advanced our understanding of many developmental and pathophysiological processes at the molecular level[6, 7]. To date, most of the mouse genes have been successfully targeted by two loxP sites (floxed allele), which could be recombined by cell-type specific Cre lines. For many *in vivo* genetic studies, a tissue-specific gene knockout approach has been widely used as a gold standard to dissect gene function in a particular cell type during a biological process.

While constitutively active Cre is robust and efficient in recombination for gene deletion, Cre lacks temporal control when the promoter driving Cre is active from the embryonic to the adult stage. Therefore, a temporally active Cre is needed to avoid early mortality as a result of the ablation of an indispensable gene at a key developmental stage. To overcome the temporal limitation of Cre, CreER, the fusion of Cre with a mutated estrogen receptor, is generated to

enable its activation only after tamoxifen (Tam) treatment, and interaction of ER with Tam induces nuclear translocation of CreER from the cytoplasm[8]. As a result, the activity of inducible CreER could be controlled by Tam treatment, thus restricting CreER activity at a specific time window and permitting gene functional analysis at a higher spatiotemporal resolution than Cre[9]. As the loxP-flanked sequences and their locations in genome or chromatin influence its accessibility for recombination, the recombination efficiency of CreER varies significantly among different floxed alleles, ranging from easy- to inert-to-recombine alleles[10]. For example, CreER has high efficiency in targeting the easy-to-recombine alleles such as generic reporter *R26-tdT*[11], but exhibits low efficiency on some other reporters such as *R26-Confetti*[12], or may not excise some floxed gene alleles that are inert or resistant for targeting, leading to failure of gene deletion in some tdT^+ cells and causing false-positive tracing fate (Figure 1A). In other words, tdT expression may not always necessarily denote gene deletion in the labeled cells. The notion that the pattern of Cre-expression based on a reporter (e.g. *R26-tdT*) represents the pattern of Cre-mediated recombination is less rigorous[13]. It is therefore not correct to assume the successful deletion of the gene of interest based on the deletion of another gene (or reporter) by the same CreER mouse line.

One way to increase the recombination efficiency of CreER is to treat mice with many times of Tam, as the high dosage of Tam in theory could increase the chances of CreER-mediated recombination. However, the increased dosage of Tam treatment is toxic and could lead to many side effects on the phenotype, creating confounding effects on the study[14, 15]. To increase the inducible recombination efficiency of floxed alleles, Lao et al. reported a mosaic mutant analysis with spatial and temporal control of recombination (MASTR), in which initial FlpoER-mediated recombination induces GFPCre expression, that subsequently recombines floxed alleles effectively in GFP^+ cells[16]. Similarly, a Flp-induced mosaic analysis system with Cre or Tomato (MASCOT) has been reported to express Cre and reporter simultaneously in a cell[17]. While MASTR and MASCOT enable spatiotemporal labeling of mosaic mutant cells utilizing the current resources of floxed mouse lines, it may not be suitable for effective gene knockout at a tissue/population level due to low recombination efficiency initiated by Flp recombinase. In order to improve the inducible recombination efficiency, Tian et al. have also reported a self-cleaved inducible CreER (sCreER) that switches inducible CreER into a constitutively active Cre,

effectively recombining floxed allele for gene manipulation[10]. However, self-cleaved inducible Cre mice have to be newly generated each time for targeting different promoters, which is time-consuming and the system is not compatible with current resources largely based on CreER mouse lines. A recent study has reported an iSuRe-Cre strategy to induce and report Cre-dependent genetic modifications utilizing conventional CreER tools[18]. The reporter expression reflects gene knockout in cells by *iSuRe-Cre*. However, two limitations hinder its widespread applications. First, without any induction, the *iSuRe-Cre* transgene is leaky in some tissues such as heart and skeletal muscle. Second, the initial recombination for releasing the *iSuRe-Cre* allele induced by CreER is far from efficient in some tissues compared with easy-to-recombine alleles such as *R26-tdT*[18], limiting its usage for efficient gene knockout at a tissue level. Thus, a new method is needed to ensure gene knockout in cells as effectively and efficiently as recombination on easy-to-recombine alleles, thus allowing efficient gene deletion at a population level.

Conventional Cre-loxP system uses tissue-specific promoter to drive Cre, thus the precision of genetic targeting solely depends on promoter activity. Now it is known that many promoters are not as specific as previously reported[19–21]. The ectopic or unwanted Cre expression in other cell types may result in unspecific cell labeling or gene deletion, leading to many confounding issues or controversies in multiple fields of research[22–24]. In addition, some cell types cannot be clearly distinguished by one marker from another, and thus could not be specifically and genetically targeted by only one gene promoter-driven recombinase[25]. To circumvent this limit, dual recombinases-mediated genetic targeting utilizes two promoters to independently drive Cre and Dre[26]. Dre is another bacteriophage recombinase that targets rox sites and is orthogonal to the Cre-loxP system[27]. While cells of interest could be labeled by specific reporters that are responsive to dual recombinases, gene deletion requires a final readout on singular Cre recombinase for targeting floxed gene alleles.

In this study, we developed two genetic strategies: roxCre and loxCre, in which the *rox-Stop-rox* (RSR) and *loxP-Stop-loxP* cassettes are respectively inserted into the *Cre* coding region, such that Cre transcription and translation are not properly carried out before the removal of the *Stop* cassette. We did not detect any spontaneous leakiness by roxCre or loxCre. We found that CreER-induced loxCre effectively deletes genes in reporter-labeled cells, ensuring efficient gene

knockout for the evaluation of lineage-specific gene function. In addition, DreER-induced roxCre efficiently deletes genes in cells for both specific and efficient intersectional genetic studies. We expect that these tools would overcome the issues of inefficient and non-specific genetic modifications, enhancing our ability to more precisely manipulate genes in a particular cell lineage for a better understanding of gene functions in multiple biomedical fields.

Results

Design of roxCre for DreER-induced mCre expression

A sequential genetic approach has been recently reported to delete genes[28], which uses Dre-induced CreER expression by removing the *loxP-Stop-loxP* sequence ahead. As aforementioned, the recombination carried out by the released CreER for excising floxed gene alleles may not be as efficient as reporters (e.g. *R26-tdT*). A possible strategy to control Cre activity is to engineer a rox-flanked *Stop* sequence ahead of its DNA (*R26-RSR-Cre*) that will be removed after Dre-rox recombination (Figure 1B). However, such a strategy is not ideal, as the upstream transcriptional *Stop* sequence might not completely prevent *Cre* transcription, leading to the leakiness issue associated with the *iSuRe-Cre* strategy[18], for instance (Figure 1C). We then inserted a longer *Stop* sequence or reversed the direction of the *Cre* DNA by generating *R26-RSR-Cre2* and *R26-R-reverseCre-R* lines, respectively, that still displayed heavy leakiness as Cre was independently activated for recombination without being crossed with a Dre mouse line (Figure 1D-G). To enable efficient dual recombinases-mediated gene knockout, we need to design a strategy for DreER-induced expression of a constitutively active Cre, and the system should not exhibit leakiness.

We first generated roxCre, in which an RSR cassette was inserted into the coding region of *Cre*, splitting *Cre* into N- and C-terminal segments (Figure 1H). After removal of RSR by Dre-rox recombination, Cre is recombined containing one remaining rox site within its coding sequence, hereafter termed as modified Cre (mCre, Figure 1H). To ensure that mCre has the same recombination efficiency as a conventional Cre, we inserted a rox sequence at 12 different sites of *Cre* individually, most of which were located between helix domains (Figure 1I). As insertions of additional amino acids of rox would change the original sequence of Cre, potentially leading to reduced activity, we screened for their recombination efficiency by

transfecting cells with 12 versions of mCre (mCre1 to mCre12) and the responding GFP reporter (Figure 1J). We found that mCre1, 2, 3, 4, 6, 7, 10, and 11 displayed a comparable efficiency as the conventional Cre; whereas mCre5, 8, 9, and 12 exhibited impaired recombination efficiency (Figure 1K, L). These data demonstrate that at least some of the mCre generated in our study were as efficient as the conventional Cre *in vitro*.

Next, we sought to screen for the most robust version of mCre *in vivo*. We generated four roxCre knock-in mice based on mCre1, 4, 7, and 10 as determined from the above *in vitro* screening (Figure S1-S2). In these roxCre lines, RSR inserted into *Cre* was removed by Dre-rox recombination resulting in the generation of mCre (Figure 1J). We then used hepatocyte- and endothelial cell-specific promoters albumin (Alb) and VE-cadherin (Cdh5) to drive roxCre, and generate *Alb-roxCre1-tdT*, *Cdh5-roxCre4-tdT*, *Alb-roxCre7-GFP*, and *Cdh5-roxCre10-GFP* knock-in mice, respectively (Figure S1-S2). No fluorescence signal was detected in *Alb-roxCre1-tdT*, *Cdh5-roxCre4-tdT*, *Alb-roxCre7-GFP*, and *Cdh5-roxCre10-GFP* knock-in mice (Figure S1-S2), demonstrating no reporter leakiness without Dre-rox recombination. These mice were crossed with *R26-GFP*[29] or *R26-tdT*[11] reporters. Nor did we detect leakiness as a result of any Cre activity when they were crossed with *R26-tdT* or *R26-GFP* reporter (Figure S1-S2). These data demonstrate that roxCre was functionally efficient yet non-leaky.

DreER-induced mCre robustly recombines inert alleles

The mCre system was originally designed for inducible gene knockout to study biological functions. To more rigorously evaluate the recombination effectiveness among different versions of mCre, we used an inducible DreER to temporally and specifically release mCre in hepatocytes or endothelial cells. Although mCre efficiently labeled virtually all targeted cells (Figure S3A-E), a strong Cre is not needed to drive easy-to-recombine alleles such as *R26-tdT* reporter in this case. To identify a strong mCre, we attempted to target one of the most inert alleles for recombination, *R26-Confetti*[12], which is often used as reporters for clonal analysis due to its sparse labeling with rare recombination events[10, 12]. In the Cre-loxp recombination principle, Cre can remove the sequence between two loxp sites that have the same transcription direction and turn over the sequence in the middle of two loxp sites that have the opposite transcription

direction. As the recombination efficiency of CreER is limited, *R26-Confetti* only can be recombined into YFP, or nGFP (nuclear GFP), or mCFP (membrane CFP), or RFP (Figure S3F).

We first crossed *R26-DreER;R26-Confetti* mice with *Alb-roxCre1-tdT* (group 1) and *Alb-roxCre7-GFP* (group 2) mice, and Tam-induced Dre-rox recombination yielded *Alb-mCre1-tdT* and *Alb-mCre7-GFP* mice, respectively (Figure 2A). In this system, the constitutively active mCre would completely remove the sequence between two loxp sites that have the same transcription direction, and recombine *R26-Confetti* into two sets of reporter pairs: YFP-nGFP pair and mCFP-RFP pair, which could be detected in a single hepatocyte due to the poly-nuclear or polyploidy feature of hepatocytes (Figure 2A). In each pair, reporters can equally express, as mCre is strong enough to turn over the sequence in the middle of two loxp sites that have the opposite transcription direction (Figure S3G). Therefore, we used YFP/mCFP to detect mCre recombination efficiency and tdT/RFP or GFP to trace its endogenous reporter activity that was driven by Alb promoter in *Alb-mCre1-tdT* and *Alb-mCre7-GFP* mice, respectively (Figure 2A). Besides, *Alb-CreER*^{26}*;R26-Confetti* was used as a control under the same Tam treatment (group 3, Figure 2A).

In fact, DreER-rox recombination in hepatocytes is not 100%. Therefore, we quantified YFP and mCFP expression in hepatocytes with DreER-rox recombination to evaluate the strength of mCre activity. We found sparse YFP$^+$ or mCFP$^+$ hepatocytes (0.39 ± 0.04%) in *Alb-CreER;R26-Confetti* mice. On the other hand, 99.94 ± 0.02% of tdT$^+$ hepatocytes (DreER recombined) were YFP$^+$ or mCFP$^+$ in *R26-DreER;Alb-roxCre1-tdT;R26-Confetti* mice; and 76.83 ± 3.20% of GFP$^+$ hepatocytes (DreER recombined) were YFP$^+$ or mCFP$^+$ in *R26-DreER;Alb-roxCre7-GFP;R26-Confetti* mice (Figure 2B, C). Similarly, we found that 39.81 ± 2.61% of tdT$^+$ endothelial cells and 74.15 ± 3.64% of GFP$^+$ endothelial cells in small intestine were YFP$^+$ or mCFP$^+$ in *R26-DreER;Cdh5-roxCre4-tdT;R26-Confetti* and *R26-DreER;Cdh5-roxCre10-GFP;R26-Confetti* mice, respectively, compared with sparse labeling in *Cdh5-CreER;R26-Confetti* mice (Figure 2D-F). Similar results were also observed in other tissues and organs (Figure S4). These data demonstrate that inducible DreER controlled roxCre activation, and mCre1 was the most efficient recombinase assisting the recombination of inert alleles *in vivo*.

Dre-induced mCre efficiently deletes floxed genes

Next, we examined the efficiency of gene knockout by Dre-induced mCre. We generated the *Alb-roxCre1-tdT;Ctnnb1*$^{flox/flox}$ mouse line to examine if Dre-induced mCre could efficiently delete *Ctnnb1* gene that encodes β-catenin (Ctnnb1^{flox})[30]. To enhance the precision and specificity of targeted cells, we employed intersectional genetic targeting using dual recombinases, and generated the *Cyp2e1-DreER* mouse line that labeled peri-central hepatocytes and also renal epithelial cells (Figure S5). Crossing *Cyp2e1-DreER* with hepatocyte-specific *Alb-roxCre1-tdT* mice achieved genetic targeting of peri-central hepatocytes exclusively, circumventing issues associated with ectopic targeting of renal cells by *Cyp2e1-DreER* line (Figure S5) or unwanted targeting of peri-portal hepatocytes by *Alb-CreER* line[26].

We generated *Cyp2e1-DreER;Alb-roxCre1-tdT;Ctnnb1*$^{flox/flox}$ triple knock-in mice (mutant), in which Tam-induced DreER-rox recombination yielded mCre that subsequently targeted floxed Ctnnb1 allele (Figure 3A). We treated mutant mice and their littermate control *Cyp2e1-DreER;Alb-roxCre1-tdT;Ctnnb1*$^{flox/+}$ mice with Tam at 8 weeks of age, and purified tdT^{+} hepatocytes for analysis at 3 days after Tam treatment (Figure 3B). qRT-PCR analysis showed significantly reduced *Ctnnb1* and *Glul* in tdT^{+} hepatocytes of the mutant group, compared to that of the control group (Figure 3C, D). We then collected liver samples at 3 days (D) and 4 weeks (W) post-Tam for further analysis (Figure 3E). Immunostaining for tdT, β-catenin, GS, and E-CAD on liver sections revealed similar levels of β-catenin and GS expression in the 3D mutant liver than the control liver (Figure 3F). However, expression of β-catenin and GS almost disappeared in the 4W mutant sample compared with the control (Figure 3G), suggesting efficient deletion of *Ctnnb1* and down-regulation of Wnt signaling downstream gene target GS. Western blotting against β-catenin and qRT-PCR against *Ctnnb1* of purified tdT^{+} hepatocytes at 4W post-Tam revealed a near complete loss of Ctnnb1 at both the protein and mRNA levels (Figure 3H-K). Additionally, Wnt downstream target genes including *Glul*, *Axin2*, *Cyp1a2*, *Cyp2e1*, *Oat*, *Tcf7*, *Lect2*, *Tbx3*, *Slc1a2*, and *Rhbg* were significantly reduced in the mutant than the control group (Figure 3L). Taken together, these data demonstrate that DreER-induced mCre enabled efficient intersectional genetic manipulation in the cells of interest.

***R26-loxCre-tdT* is efficiently recombined by CreER**

Having successfully established roxCre1 for Dre-induced mCre expression, we iterated the system to enable mCre induction by CreER, a more widely available tool for biomedical research. We replaced the rox sites of roxCre1 with loxP sites by inserting a *loxP-Stop-loxP* cassette into *Cre* cDNA at the same locus as roxCre1(Figure 4A). We then generated a new *R26-loxCre-tdT* mouse line, in which mCre and tdT would be expressed simultaneously after *Stop* removal, allowing tdT as a surrogate marker for mCre detection in the same cell (Figure 4C). We then designed two experiments to verify the baseline leakiness of Cre expression (Figure 4B) and to examine if mCre/tdT could be expressed after *Stop* removal (Figure 4C), respectively, by *R26-loxCre-tdT;R26-tdT* (Strategy 1) and by injecting *AAV8-Cre* virus into *R26-loxCre-tdT* mice (Strategy 2). Immunostaining for tdT on tissue sections revealed no tdT expression in Strategy 1 demonstrating minimal to negligible leakiness at the baseline. The robust tdT expression in the liver in Strategy 2 showed mCre/tdT was allowed to express after *Stop* removal by AAV2/8-hTBG-Cre(Figure 4D).

We then confirm if the new design can improve the labeling efficiency compared to the conventional approach by crossing endothelial cell-specific *Cdh5-CreER* with the conventional reporter *R26-tdT* (Strategy 3) and the new reporter *R26-roxCre-tdT* mice (strategy 4) under the same protocol of Tam treatment (Figure 4E). Immunostaining for tdT and VE-Cad on tissue sections revealed no difference in the percentage of VE-Cad$^+$ endothelial cells expressing tdT between strategies 3 and 4 (Figure 4F- G), suggesting that recombination of the *R26-roxCre-tdT* allele by CreER was as efficient and easy as *R26-tdT* which is an easy-to-recombine reporter mouse line. Therefore, *R26-roxCre-tdT* can be served as an easy-to-recombine reporter, enabling efficient recombination to generate mCre/tdT in CreER-expressing cells, without leakiness.

***R26-loxCre-tdT* adaptor ensures efficient recombination**

We next examined whether mCre released from loxCre could also display high recombination efficiency for the inert-to-recombine allele *R26-Confetti*. We generated the triple knock-in *Cdh5-CreER;R26-loxCre-tdT;R26-Confetti* mouse line in which mCre was released for confetti recombination upon Tam treatment, and *Cdh5-CreER;R26-Confetti* was used as a control (Figure 5A). Both mouse strains were treated with Tam at 8 weeks of age and their tissue samples were collected for analysis at one week post-Tam (Figure 5B). As tdT expression is also detected in

R26-loxCre-tdT after the first recombination by CreER, we only compared the YFP and mCFP fluorescence signals from *R26-Confetti* (Figure 5A). Wholemount-fluorescence imaging showed significantly more YFP and mCFP signals in the retina of the *Cdh5-CreER;R26-loxCre-tdT;R26-Confetti* mice, compared with that of *Cdh5-CreER;R26-Confetti* mice (Figure 5C). Immunostaining on frozen sections revealed significantly higher percentage of endothelial cells expressing YFP/mCFP in various vascularized organs of the *Cdh5-2a-CreER;R26-loxCre-tdT;R26-Confetti* mice, compared with that of the *Cdh5-2a-CreER;R26-Confetti* mice (Figure 5D). Of note, virtually all tdT^+ endothelial cells simultaneously expressed YFP/mCFP (Figure 5D). These data demonstrate that the *R26-loxCre-tdT* line could be used as an adaptor to enhance CreER-mediated recombination with a simultaneous expression of tdT as an accurate indicator.

***R26-loxCre-tdT* ensures gene deletion in tdT^+ cells**

To examine if *R26-roxCre-tdT* enables *Alb-CreER* to more efficiently delete genes, we crossed *Alb-CreER;R26-roxCre-tdT* with $Ctnnb1^{flox/flox}$ to generate *Alb-CreER;R26-roxCre-tdT*;$Ctnnb1^{flox/flox}$ mice for evaluating the efficiency of releasing mCre and also mCre-mediated *Ctnnb1* deletion in tdT^+ hepatocytes (Figure 6A). We also crossed *Alb-CreER;R26-tdT2* with $Ctnnb1^{flox/flox}$ to generate *Alb-CreER;R26-tdT2;*$Ctnnb1^{flox/flox}$ mice to evaluate recombination efficiency of both *R26-tdT2* and $Ctnnb1^{flox/flox}$ alleles (Figure 6A). Additionally, *Alb-CreER;R26-roxCre-tdT*;$Ctnnb1^{flox/+}$ mice were used as a control for heterozygous gene deletion (Figure 6B). We treated *Alb-CreER;R26-tdT2;*$Ctnnb1^{flox/flox}$ and *Alb-CreER;R26-roxCre-tdT*;$Ctnnb1^{flox/+}$ mice with five dosages of Tam, and treated *Alb-CreER;R26-roxCre-tdT;*$Ctnnb1^{flox/flox}$ mice only with one dosage of Tam, and collected tissues for analysis to determine *Ctnnb1* gene deletion at 4 weeks after Tam treatment (Figure 6B). Immunostaining data showed that β-catenin, GS, E-CAD, and Cyp2e1 were significantly reduced in *Alb-CreER;R26-roxCre-tdT;*$Ctnnb1^{flox/flox}$ mice, but were readily detectable in both *Alb-CreER;R26-tdT2;*$Ctnnb1^{flox/flox}$ and *Alb-CreER;R26-roxCre-tdT*;$Ctnnb1^{flox/+}$ mice (Figure 6C, D). Of note, a small number of hepatocytes continued to express Cyp2e1 in *Alb-CreER;R26-roxCre-tdT;*$Ctnnb1^{flox/flox}$ mice (Figure 6D). These Cyp2e1-expressing hepatocytes were unanimously tdT^-, and not a single tdT^+ hepatocyte expressed Cyp2e1 (Figure 6D), suggesting that these $Cype2e1^+tdT^-$ hepatocytes were likely not targeted by *Alb-CreER* for the recombination to release mCre. Western blotting of β-catenin and qRT-PCR of *Ctnnb1* of isolated hepatocytes revealed almost complete deletion of Ctnnb1 in the

Alb-CreER;R26-tdT2;Ctnnb1$^{flox/flox}$ mice compared with those in control groups (Figure 6E-G). Additionally, those Wnt downstream or regulated genes including *Glul*, *Axin2*, *Cyp1a2*, *Cyp2e1*, *Oat*, *Tcf7*, *Lect2*, *Tbx3*, *Slc1a2*, and *Rhbg* were significantly reduced in hepatocytes derived from the *Alb-CreER;R26-tdT2;Ctnnb1*$^{flox/flox}$ mice, compared with those from the *Alb-CreER;R26-tdT2;Ctnnb1*$^{flox/flox}$ and *Alb-CreER;R26-roxCre-tdT;Ctnnb1*$^{flox/+}$ mice, respectively (Figure 6H).

Discussion

In this study, we respectively generated two genetic tools based on roxCre and loxCre to enhance the effectiveness of Cre-mediated gene deletion even under the control of a less efficient DreER or CreER driver. The *R26-loxCre-tdT* line permitted efficient recombination to release mCre, which greatly enhanced the effectiveness of subsequent gene deletion in tdT$^+$ cells. The inclusion of a fluorescent reporter with roxCre or loxCre allele allowed us to readily detect cells with gene deletion, thus facilitating further detailed characterization of these cells both *in situ* and *ex vivo*. Furthermore, the strategy of roxCre further enhances the precision for cell fate mapping with genetic deletion simultaneously in Dre$^+$Cre$^+$ cells through intersectional genetic targeting. For example, we have showcased the power of roxCre for genetic manipulation in peri-central hepatocytes, resulting in efficient gene knockout in tdT$^+$ hepatocytes specifically. Additionally, we have demonstrated the efficient deletion of *Wls* and *Ctnnb1* genes in tdT$^+$ endothelial cells and hepatocytes, respectively, by combining *R26-loxCre-tdT* adaptors with CreER lines. Altogether, we have demonstrated that the *R26-loxCre-tdT* mouse line could be widely applicable to any CreER-bearing mouse for enhancing its effectiveness in genetic modification with simultaneous cell lineage tracing in multiple biomedical research fields.

Coupling gene deletion with reporter activity can greatly facilitate the accuracy of studying gene function in targeted cells by *in situ* investigation with or without specific cell isolation for *ex vivo* experiments. Assuming successful gene deletion in the cell type of interest because of efficient cell labeling by a lineage-specific reporter mediated by the same Cre is incorrect, as recombination efficiency varies significantly among different mouse lines. In fact, many factors can potentially impact Cre-loxP recombination efficiency. Concerning the recombinase enzyme, the type of recombinase, the strength of the promoter that drives the recombinase, the expression level of the recombinase, and the translocation efficiency of an inducible recombinase such as

CreER from the cytoplasm to the nucleus after Tam induction could all influence the recombination efficiency. For the targeted DNA sites, the genomic loci where targeted sites are placed, loxP-flanked sequence length, and even the inserted DNA sequence *per se* could also influence the recombination efficiency. We clearly observed the differences in recombination efficiency of different alleles (e.g. *R26-tdT*, *R26-tdT2*, and *R26-Confetti*), even located in the same genomic locus (e.g. Rosa26) mediated by the same CreER under the same Tam treatment protocol, as evidenced by high efficiency in *R26-tdT*, medium efficiency in *R26-tdT2*, and low efficiency in *R26-Confetti*. Importantly, *R26-loxCre-tdT* was found to be efficient for recombination mediated by CreER, which was comparable to that of the *R26-tdT* allele, thus enabling efficient labeling of cells of interest. Furthermore, *R26-loxCre-tdT* also enabled CreER to efficiently ablate genes in any cells that had been genetically traced with tdT, synchronizing genetic modification and lineage tracing in the labeled cells.

Despite the high demand in the quest for efficient genetic modification in genetically labeled cells is significant, no efficient tool has been available to date. A previous study attempted to generate an adaptor for CreER to efficiently delete genes in reporter-labeled cells by a transgenic *iSuRe-Cre*[18]. However, the leakiness of *iSuRe-Cre* on the Rosa26 locus was too high in the male germline. Therefore, a transgenic mouse line was reconstructed subjected to *iSuRe-Cre* mediated recombination, which still displays consistently leaky recombination in some tissues such as cardiac and skeletal muscle[18]. The leakiness was likely due to the inability of the *Stop* cassette to completely block the transcription of *Cre* in *iSuRe-Cre*. Consistently, we also found leakiness as a result of *Cre* transcription even in the presence of *Stop* ahead in the *R26-Stop-Cre* mouse line (Figure 1C). These leaky events, despite being at different levels, demonstrate that transcription of recombinases such as Cre could not be easily inhibited. Instead of blocking its transcription by *Stop*, we inserted *Stop* in between the N- and C-terminus of *Cre*, thus preventing Cre expression completely at the translational level. We found that the removal of *Stop* rendered mCre, which efficiently recombined different types of alleles examined in our study. Additionally, we also observed inefficient recombination in hepatocyte-specific *Alb-CreER;iSuRe -Cre* mice, rendering this line less effective in conditional gene deletion in hepatocytes at a population level. In contrast, *R26-loxCre-tdT* was an easy-to-recombine allele for *Alb-CreER* mediated recombination, allowing tdT labeling and gene deletion simultaneously in the majority of

hepatocytes. Therefore, our *R26-loxCre-tdT* line has circumvented the leakiness issue associated with *iSuRe-Cre*, exhibited as an easy-to-recombine allele for CreER targeting, and synchronized fate mapping with efficient genetic modification robustly, efficiently and temporally in the same cells.

The efficiency of CreER-loxP recombination depends not only on the location of the loxP-containing alleles but also on the strength of CreER driven by an active promoter. For instance, *R26-loxCre-tdT* was an easy-to-recombine allele compared with other tested R26 alleles, but the recombination efficiency to release mCre for subsequent recombination depends on the strength of the selected CreER lines for genetic targeting. It is known that different CreER mouse lines vary significantly for recombination, largely due to the differences in expression levels of CreER and Tam induction efficacy. Therefore, we should select a relatively stronger CreER line for efficient recombination on *R26-loxCre-tdT* to release mCre for gene deletion at a population level. In addition to bulk gene deletion, the *R26-loxCre-tdT* also offers an alternative way for mosaic analysis of gene function by sparse labeling. Previous studies have used Cre-loxP to over-express gene and fluorescence reporter simultaneously in a cell for functional genetic mosaic (ifgMosaic) analysis[31]. While ifgMosaic could over-express some dominantly negative genes for functional study, this system could not take advantage of rich resources from many conventionally available floxed mouse lines for loss-of-function study. Mosaic mutant analysis with spatial and temporal control of recombination (MASTR) utilized FlpoER for tissue-specific expression of GFPCre to delete floxed genes[16]. Compared with FlpoER lines, CreER is a more broadly used recombinase that most laboratories use in gene deletion experiments. A recent study using dual recombinase-mediated cassette exchange (MADR) also permits stable labeling of mutant cells expressing transgenic elements from defined chromosomal loci[32]. However, the use of viruses and electroporation, albeit convenient and rapid, are inefficient to specifically target any type of cells for *in vivo* studies. The powerful mosaic analysis with double markers (MADM) enables simultaneous lineage tracing of a pair of mutant and control sibling cells with distinct fluorescence reporters, allowing precise mosaic analysis of gene function in any cell[33, 34]. Nevertheless, the MADM cassette has to be combined with the mutant null allele, which is not readily available for synchronizing reporter expression and genetic modification in cells of interest. The ensured gene deletion with tdT reporter by *R26-roxCre-tdT* could be coupled with

any present CreER and loxP alleles for potential mosaic analysis. This could be achieved by adjusting Tam at a lower dosage so that the recombination efficiency to release mCre could be low in cells of interest for sparse tdT labeling. We believe that *R26-loxCre-tdT* could be useful for robust, efficient, and temporal gene deletion at a population level and mosaic analysis at a single-cell level with improvement in the future.

In this study, we deleted the Wnt signaling gene *Ctnnb1* in hepatocytes to present how roxCre and loxCre respectively enable efficient recombination. The impact of deleting *Ctnnb1* in a specific hepatocyte lineage on homeostasis, development, and injury recovery requires further investigation. Besides, roxCre and loxCre should be involved in more gene functional studies, in which knocking out genes continuously leads to embryonic death due to their crucial role in development.Both loxCre and roxCre incorporate a fluorescent reporter tdT for genetic lineage tracing, revealing efficient gene knockout in cells marked by the reporter simultaneously. In other words, one reporter tdT can only show cells with gene knockout. One direction to improve the current *R26-loxCre-tdT* for mosaic analysis in the future is to generate a second distinct surrogate reporter for wild-type cells as an internal control during mosaic analysis, similar to that in the MADM system.

Materials and Methods

Mice

Experiments using mice (Mus musculus) were carried out with the study protocols (SIBCB-S374-1702-001-C1) approved by the Institutional Animal Care and Use Committee of Center for Excellence in Molecular Cell Science (CEMCS), Shanghai Institute of Biochemistry and Cell Biology, Chinese Academy of Sciences. The *R26-DreER*[35], *Alb-CreER*[26], *R26-GFP*[29], *R26-tdT*[11], *R26-tdT2*[36], *R26-RSR-tdT*[37], *R26-Confetti*[12],and *Ctnnb1-flox*[30] mouse lines were used as previously described. New knock-in mouse lines *R26-RSR-Cre*, *R26-RSR-Cre2*, *R26-R-reverseCre-R*, *Cdh5-CreER*, *Alb-roxCre1-tdT*, *Alb-roxCre7-GFP*, *Cdh5-roxCre4-tdT*, *Cdh5-roxCre10-GFP*, *Cyp2e1-DreER*, and *R26-loxCre-tdT* were generated by homologous recombination using CRISPR/Cas9 technology. These new mouse lines were generated by the Shanghai Model Organisms Center, Inc. (SMOC). These mice were bred in a C57BL6/ICR mixed background. All mice were housed at the laboratory Animal center of the Center for Excellence in Molecular Cell Science in a Specific Pathogen Free facility with individually ventilated cages. The room has controlled temperature (20–25 °C), humidity (30–70%), and light (12 hours light-dark cycle). For the determination of the embryonic period of sampling, the day on which the vaginal plug was examined in female mice was recorded as E0.5. As the sex is not relevant to the topic of this study, male and female mice ranging in age from E13.5 to 12W (week) were allocated and mixed into the experimental groups in this study. No data in mouse experiments were excluded.

Genomic PCR

Genomic DNA was prepared from the mouse toes or embryonic tails. Tissues were precipitated by centrifugation at maximum speed for 1 min at room temperature. After that, tissues were lysed by lysis buffer (100 mM Tris-HCl, 5 mM EDTA, 0.2% SDS, 200 mM NaCl, and 100 μg/mL Proteinase K) at 55 °C overnight. About 750μL pure ethanol was added to the lysis mixture and mixed thoroughly, followed by centrifugation at maximum speed for 5 min at room temperature to collect the DNA precipitation. Then the supernatant was discarded and the mixture was dried at 55 °C for 1 hour. About 100–200μL double-distilled H_2O was added to dissolve the DNA. The genomic PCR primer pairs were designed for the mutant alleles spanning

both endogenous genomic fragments and insert fragments. All the new genomic PCR primer sequences would be provided upon request.

In vitro screening of mCre for efficient recombination

To test if the remaining rox sequence after Dre-rox recombination would impact Cre activity, the pCDNA3.1[38] (Invitrogen, V79020) was used to express 12 types of modified Cre (mCre). For testing the Cre activity, 500ng pCDNA3.1-mCre plasmids, or pCDNA3.1 (negative control), or pHR-CMV-nlsCRE (positive control, Invitrogen, 12265) were mixed with 500ng pCAG-loxp-stop-loxp-ZsGreen (Addgene, 51269) plasmid. *In-vitro* transfection experiment protocol is performed according to that described previously[39]. For cell culture medium, DMEM (ThermoFisher, 11965092) was supplemented with 10% fetal bovine serum (fetal bovine serum, 10099141c) for preparing fresh complete culture medium. Poly-D-Lysine (ThermoFisher, A3890401) precoated coverslips (biosharp, BS-14-RC) or 10 cm dish (ThermoFisher, 150466) overnight to dry, and sterile water was used to flush it for three times. A cryogenic vial of QBI293 was put in a 37°C water bath, then thawed liquid contents into a 50 mL conical tube (Corning,430829) prefilled with 5 mL prewarmed fresh complete culture medium. After centrifuging at 125 × g for 5 min, the supernatant was discarded and cells were resuspended in 10 mL complete medium. One-third of the cells were plated in a 10 cm coated dish and incubated in cultures at 37°C. On the second day, the culture medium was discarded, and the cell layer was briefly rinsed with prewarmed PBS (Gibco, C10010500BT). After that, PBS was removed and 1 mL 0.25% Trypsin solution (with EDTA, Gibco, 25200072) was added. The cells were incubated at 37°C for 3 min., and a 6mL complete growth medium was added to aspirate cells by gently pipetting. The medium-containing cells were transferred to a 50 mL conical tube and centrifuged at 125 × g for 5 min. The supernatant was discarded, and cells were resuspended in 5.2 mL complete culture medium. 24 well plates were placed into coated coverslips and added 1mL complete culture medium. 200μL cells were added to every experimental well and incubated cultures at 37°C for 9 hours to grow up to ~80%. The new prewarmed fresh complete culture medium replaced the old for 1mL per well. Lipofectamine™ 3000 Transfection Reagent (Thermo Fisher, L3000015) was used for plasmid transfection. Samples were incubated in cultures at 37°C for 24 hours. Samples were washed by PBS once and mounted on slides by

VECTASHIELD®. Antifade Mounting Medium with DAPI (Vector, H-1200). Images were acquired using an Olympus microscope (Olympus, BX53).

Tamoxifen injection and sample collection

For induction of CreER or DreER, tamoxifen (Sigma, T5648) was dissolved in corn oil and was administered to mice by oral gavage. *Cyp2e1-DreER;Alb-roxCre1-tdT;Ctnnb1*$^{flox/+}$ and *Cyp2e1-DreER;Alb-roxCre1-tdT;Ctnnb1*$^{flox/flox}$ mice were treated with 0.05 mg Tam per gram mouse body weight for one dose. For comparing tdT^{+} cells, *Cyp2e1-DreER;R26-RSR-tdT* and *Cyp2e1-DreER;Alb-roxCre1-tdT* mice were treated with 0.05 mg Tam per gram mouse body weight. *R26-DreER;Alb-roxCre1-tdT;R26-Confetti*, *R26-DreER;Cdh5-roxCre4-tdT;R26-Confetti*, *R26-DreER;Alb-roxCre7-GFP;R26-Confetti*, *R26-DreER;Cdh5-roxCre10-GFP;R26-Confetti*, *R26-DreER; R26-DreER;Alb-roxCre7-GFP;R26-GFP*, *Alb-CreER;R26-Confetti*, *Cdh5-CreER;R26-Confetti*, *Alb-CreER;R26-tdT2;R26-Confetti*, *Alb-CreER;iSuRe-Cre;R26-Confetti*, *Alb-CreER;R26-loxCre-tdT;R26-Confetti*, *Alb-CreER;R26-loxCre-tdT;Ctnnb1*$^{flox/flox}$, and *Cyp2e1-DreER;R26-RSR-tdT* mice were treated with 0.2 mg Tam per gram mouse body weight for one dose. *Cdh5-CreER;R26-tdT*, *Cdh5-CreER;R26-loxCre-tdT*, *Cdh5-CreER;R26-Confetti*, *Cdh5-CreER;R26-loxCre-tdT;R26-Confetti*, *Alb-CreER;R26-tdT2;Ctnnb1*$^{flox/flox}$, and *Alb-CreER;R26-loxCre-tdT;Ctnnb1*$^{flox/+}$ mice were treated with 0.2 mg Tam per gram mouse body weight one dose per day for 5 days. *R26-RSR-Cre2;R26-GFP* and *R26-R-reverseCre-R;R26-tdT* mice were sacrificed at 8 weeks old for analysis. Mice, both males and females, at the age of 8–12 weeks were used for experiments with similar-aged mice for both control and experimental groups.

Whole-mount imaging and sectioning

The tissue samples were fixed with 4% paraformaldehyde (PFA) (Sigma, P6148-500g, wt/vol in PBS) for 1 or 2 hours (stomach, intestine, colon) at 4°C, followed by washing with PBS three times. The fixed tissues were placed in an agarose-filled petri dish for bright-field and fluorescence imaging by a Zeiss stereoscopic microscope (AxioZoom V16). For cryo-sections, tissues were sectioned to slides of 10-μm thickness after dehydration by 30% sucrose (Sinopharm, H-10021463, wt/vol in PBS) overnight and pre-embedding with OCT (Sakura) at 4 °C for 1 hour.

Immunostaining

Immunostaining was performed as previously described[40]. 0.2% PBST was prepared with 0.2% (vol/vol) Triton X-100(Sigma, T9284) dissolved in PBS. Tissue sections were blocked with 2.5% normal donkey serum and 0.1% 4'6-diamidino-2-phenylindole (DAPI, Vector lab, D21490) dissolved in 0.2% PBST for 30 mins after washing with PBS three times. The tissue sections were incubated with primary antibody diluted in 0.2% PBST at 4°C overnight. On the next day, sections were incubated with secondary antibodies diluted in 0.2% PBST at room temperature for 30 min, followed by PBS washing for three times. The slides were washed with PBS for three times. The slides were mounted with a mounting medium (Vector Lab). For weak signals, the endogenous peroxidase activity was quenched before blocking. Horseradish peroxidase or biotin-conjugated secondary antibodies and a Tyramide signal amplification kit (PerkinElmer) were used after incubating the primary antibodies. For primary antibodies of murine origin, mouse immunoglobulins were blocked with an anti-mouse Fab antibody (Jackson, 715-007-003, 1:100). The included primary antibodies are listed as follows: tdT (Rockland, 600-401-379, 1:500; or Rockland, 200-101-379, 1:500), GFP (Invitrogen, A11122, 1:500), GFP (Rockland, 600-101-215M, 1:500), GFP (Nacalai, 04404-84; 1:500), GS (Abcam, ab49873, 1:1000), β-catenin (BD Pharmingen, 610153, 1:200), E-CAD (R&D, AF748, 1:500), Ep-CAM (Abcam, ab92382, 1:500), VE-Cad (R&D, AF1002, 1:100), and CYP2E1(Abcam, ab28146, 1:100). The corresponding secondary antibodies (JIR or Abcam) were diluted according to the instructions. Images were captured by using a Nikon confocal (Nikon A1 FLIM) or an Olympus confocal (FV3000), and captured images were analyzed by Image J2 (version 2.9.0/1.54f) and Photoline (version 23.02) software. We collected five random fields from each liver section for quantification. Mutant and control sections were processed at the same time to avoid potential batch differences during staining. Imaging of all immunostained slides was performed under the same exposure and contrast conditions using the same confocal microscope.

Imaging of Confetti fluorescence reporters

All the fluorescence images of *R26-confetti* were taken by a Leica confocal (Leica SP8 will). All the sections were collected freshly and blocked with 0.2% PBST containing DAPI for 30 mins after washing with PBS three times. The parameters of excitement light and emission light for all channels are set as follows: DAPI (ex.405nm, em.415-450nm), CFP (ex.458nm, em.463-481nm),

GFP (ex.488nm, em.495-511nm), YFP (ex.520nm, em.527-545nm), tdT (ex.546nm, em.555-590nm), markers for antibody staining (ex.647nm, em.657-700nm). Captured images were analyzed by Image J2 (version 2.9.0/1.54f) and Photoline (version 23.02) software.

Hepatocyte dissociation

Mouse primary hepatocytes were isolated by the two-step collagenase perfusion method, which was modified from a previous protocol[41]. Briefly, mice were anesthetized and the liver was exposed through an incision in the lower abdomen. The inferior vena cava and portal vein were also exposed. A needle was inserted into the inferior vena cava and secured with a hemostatic clamp around the needle. The portal vein was cut immediately when the mouse liver was perfused with a perfusion medium buffer (containing 0.5 mM EGTA) for 5 minutes using a peristaltic pump. Then, the liver was perfused with medium containing collagenase type I (150 U/ml; Gibco, 17100-017) for 2-5 min to adequately digest the liver. After the gallbladder was removed, the liver was dissected with cold DMEM to free the hepatic cells. Then the cell suspension was passed through a 70-μm cell strainer (BD Biosciences, 352350) and centrifuged at 50 x g for 3 min at 4 °C. The supernatant was removed, and cells were resuspended in Percoll (GE Healthcare, 17-0891-01)/DMEM/10 x PBS (Sangon, B548117, diluted in 1:1) (1:1:0.1) mixture and centrifuged at 300 x g for 5 min at 4 °C. After the supernatant was removed, cells were dissected with cold DMEM and centrifuged at 50 x g for 3 min at 4 °C. Purified hepatocytes were collected for FACS analysis, qRT-PCR or Western Blot analysis.

Flow cytometric analysis and isolation of tdT$^+$ hepatocytes

Cells were stained with the following antibodies: anti-mouse CD45 APC (eBioscience, 17-0451-82, 1:400), anti-mouse CD140a APC (eBioscience, 17-1401-81, 1:100), anti-mouse CD31 APC (eBioscience, 17-0311-82, 1:40). All antibodies were diluted in 1~2mL DMEM (according to cells number) and cells were incubated with antibodies for 30 min in the dark at 4 °C. After staining, 10mL DMEM was added to stop staining, and the cell suspension was passed through a 70-μm cell strainer. Cells were centrifuged at 50 x g for 3 min at 4 °C and dissected with relevant suspended solution. For no DAPI control group, their suspended solution is 0.1% Dnase I (Worthington, LS002139, diluted by DMEM), otherwise, the suspended solution is 0.1% Dnase I mixed with 0.1% DAPI. Hepatocytes were sorted by FACS Aria SORP machine. Cell viability

was assessed with DAPI staining. $APC^{-}tdT^{+}$ hepatocytes were collected for subsequent experiments.

Total RNA extraction and qRT-PCR

Total RNA was extracted from the liver of indicated mice or hepatocytes isolated from indicated mice as previously described[42]. Cells or homogenized tissues were lysed with Trizol (Invitrogen, 15596018), and total RNA was extracted according to the manufacturer's instructions. For each sample, 1 μg of total RNA was reversely transcribed into cDNA using the Prime Script RT kit (Takara, RR047A). The SYBR Green qPCR master mix (Thermo Fisher Scientific, 4367659) was used and quantitative RT-PCR was performed on QuantStudio 6 Real-Time PCR System (Thermo Fisher Scientific). *Gapdh* was used as the internal control. Sequences of all primers would be provided upon request.

Western blot

Liver tissues were collected at the indicated stages. All samples were lysed in RIPA lysis buffer (Beyotime, P0013B) containing protease inhibitors (Roche, 11836153001) for 30 min on ice, and then centrifuged at 15,000 x g for 15 min to collect the supernatant. All samples were separated with 30μL for testing the protein concentration by Pierce™ BCA Protein Assay kits (Thermo Scientific™,23227). The remaining samples were mixed with 5 x loading buffer (Beyotime, p0015L) and boiled at 100 °C for 5 min. 40μg samples were added to every well in precast gradient gels (Beyotime, P0469M) with 1x running buffer (Epizyme, PS105S, should be diluted into 1x). After running, samples were transferred onto Immobilon® PVDF membranes (Millipore, IPVH00010). After blocking in blocking buffer (Epizyme, PS108P), the membranes were incubated with primary antibodies (diluted by primary antibody dilution buffer (Epzyme, PS114)) overnight at 4 °C, then washed for three times and incubated with HRP-conjugated secondary antibodies (diluted by 1xTBST (Epzyme, PS103S, should be diluted into 1x)). Samples were incubated with chemiluminescent HRP substrate (Thermo Fisher Scientific, WBKLS0500), and related signals were detected by MiniChemi 610 Plus (Biogp). The following antibodies were used: β-catenin (BD Pharmingen, 610153, 1: 5000), GAPDH (Proteintech, 60004-1-IG, 1:2000), β-actin (Epzyme, LF202,1:1000), HRP-donkey-a-mouse (JIR, 715-035-150, 1:5000), and HRP-goat a rabbit IgG (JIR, 111-035-047, 1:5000).

AAV injection

The AAVs used in this study were purchased from Taitool Biotechnologies company (Shanghai, China). Briefly, AAVs were produced with cis-plasmids containing the full TBG promoter, which is specifically active in hepatocytes, and Cre expression is under the control of TBG promoter, replication incompetent AAV2/8-TBG-Cre virus (AAV8-TBG-Cre, S0657-8-H5) was packaged and purified before application to mice. Mice were injected intraperitoneally at 1x 10^{11} genome copies of the virus per mouse. The above newly generated virus as well as targeting plasmids will be provided upon request.

Statistical analysis

Statistical analyses were performed using GraphPad Prism (Version 9.5.1). All the continuous variables were expressed as means ± standard error of the mean (SEM). One-way ANOVA was used to detect statistical significance between three experimental groups. The statistical difference between the two experimental groups was determined using the unpaired student's *t*-test. The probability (*P*) values of less than 0.05 were considered statistically significant.

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Author contributions

M.S., J.L., and B.Z. designed the study, analyzed the data, and wrote the manuscript; X.L., K.L., W.P., W.W., S.Z., and H.Z. bred the mice and performed experiments; K.O.L. edited the manuscript and provided intellectual input; B.Z. conceived, supervised, and organized the study.

Declaration of interests

The authors declare no competing interests.

Data availability

All data included in this study are included in the manuscript. The raw data are available upon request.

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Figure legends

Figure 1. Design of roxCre for DreER-induced mCre expression

(A) A schematic showing genetic labeling and/or gene knockout in CreER-expressing cells.

(B, D, F) Strategies for DreER-induced Cre expression.

(C) Crossing with R26-tdT mice, R26-RSR-Cre exhibits leakiness in E13.5 embryos. Scale bars, yellow 1mm. Each image is representative of 5 individual biological samples.

(E) Examination of Cre leakiness by whole-mount fluorescence images of organs collected from R26-RSR-Cre2;R26-26-GFP adult mice. Scale bars, yellow 1mm. Each image is representative of 5 individual biological samples.

(G) Examination of Cre leakiness by whole-mount fluorescence images of organs collected from R26-R-reverseCre-R;R26-tdT adult mice. Scale bars, yellow 1mm. Each image is representative of 5 individual biological samples.

(H) A schematic showing the design for roxCre and mCre.

(I) A schematic showing mCre1 to mCre12 by insertion of rox into Cre cDNA at 12 different loci. Two dual-recombinase designs for inducible Cre expression once tried before.

(J) Examination of mCre for recombination efficiency by the *loxP-Stop-loxP-GFP* reporter.

(K) Immunofluorescence images of cells stained with GFP on cells. Scale bars, white, 100 µm. Each image is representative of 5 individual biological samples.

(L) Quantification of the percentage of cells expressing GFP in each group. Data are the means ± SEM; n = 5.

Figure 2. DreER-induced mCre robustly recombines inert alleles

(A) A schematic showing the experimental design to test the recombination efficiency of mCre1 and mCre7 on the *R26-Confetti* allele. Tam-induced DreER-rox recombination leads to mCre1/tdT or mCre7/GFP expression in hepatocytes in strategy 1 or 2, respectively. Strategy 3 uses conventional *Alb-CreER* as control. YFP and mCFP signals are used for the examination of recombination on the *R26-Confetti* allele.

(B) Fluorescence images of YFP and mCFP on liver sections collected from mice in strategy 1-3. Strategy 1 and 3 exhibit tdT or GFP in hepatocytes after DreER-rox recombination, respectively. Scale bars, 100 µm. Each image is representative of 5 individual biological samples.

(C) Quantification of the percentage of hepatocytes (Heps) expressing either YFP and/or mCFP in strategy 4-6. Data are the means ± SEM; n = 11 mice for each strategy; ****$P < 0.0001$ by one-way ANOVA.

(D) A schematic showing the experimental strategy to test the recombination efficiency of mCre4 and mCre10 on *R26-Confetti* allele. Tam-induced recombination leads to mCre4/tdT or mCre10/GFP expression in endothelial cells (ECs) in strategy 4 or 5, respectively. Strategy 6 uses conventional *Cdh5-CreER* as control. YFP and mCFP signals are used for the examination of recombination on *R26-Confetti* allele.

(E) Fluorescence images of YFP and mCFP on intestinal sections collected from mice in strategy 4-6. Strategy 4 and 5 exhibit tdT or GFP in ECs after DreER-rox recombination, respectively. Scale bars, 100 µm. Each image is representative of 5 individual biological samples.

(F) Quantification of the percentage of ECs expressing either YFP and/or mCFP in strategy 4-6. Data are the means ± SEM; n = 11 mice for each strategy; ****$P < 0.0001$ by one-way ANOVA.

Figure 3. Dre-induced mCre efficiently deletes genes

(A) A schematic showing the experimental design for Dre-induced mCre expression and the subsequent gene deletion.

(B) A schematic showing the experimental strategy.

(C and D) qRT-PCR analysis of the relative expression of *Ctnnb1* (C) and Glul (D) in the tdT^+ hepatocytes. Data are the means ± SEM; n = 5; ****$P < 0.0001$ by student's *t*-test.

(E) A schematic showing the experimental strategy.

(F) Immunostaining for tdT, β-Catenin, GS, and E-CAD on liver sections collected on the day 3 post-Tam. PV, portal vein; CV, central vein. Scale bars, yellow 1mm; white 100 µm. Each image is representative of 5 individual biological samples.

(G) Immunostaining for tdT, β-Catenin, GS, and E-CAD on liver sections collected at week 4 post-Tam. Scale bars, yellow 1mm; white 100 µm. Each image is representative of 5 individual biological samples.

(H) A schematic showing the experimental strategy.

(I) Western blotting of β-Catenin and GAPDH in tdT^+ cells.

(J) Quantification of the relative expression of β-Catenin protein. Data are the means ± SEM; n = 3. **$P < 0.01$ by student's *t* test.

(K) qRT-PCR analysis of the relative expression of *Ctnnb1* in the tdT^+ cells. Data are the means ± SEM; n = 6. ****$P < 0.0001$ by student's *t* test.

(L) qRT-PCR analysis of the relative expression of *Glul*, *Axin2*, *Cyp1a2*, *Cyp2e1*, *Oat*, *Tcf7*, *Lect2*, *Tbx3*, *Slc1a2*, and *Rhbg* in the tdT^+ cells. Data are the means ± SEM; n = 6. ***$P < 0.001$, ****$P < 0.0001$ by student's *t* test.

Figure 4. *R26-loxCre-tdT* is efficiently recombined by CreER

(A) A schematic showing the knock-in strategy for the generation of the *R26-loxCre-tdT* allele. In this line, tdT is used to denote the mCre expression after the removal of *Stop*.

(B) A schematic showing the experimental strategy 1 to examine the leakiness of *R26-loxCre-tdT* mice.

(C) A schematic showing the experimental strategy 2 to test mCre/tdT expression by AAV8-TBG-Cre.

(D) Immunostaining for tdT on tissue sections shows tdT expression in the liver and pancreas when recombination was intiated by AAV8-TBG-Cre. Scale bars, white 100 µm. Each image is representative of 5 individual biological samples.

(E) A schematic showing the experimental strategies 3 and 4 for comparing the CreER-mediated first recombination efficiency between *Cdh5-CreER;R26-tdT* and *Cdh5-CreER;R26-loxCre-tdT* mice.

(F) Immunostaining for tdT and VE-Cad on liver sections collected on day 7 post-Tam. Scale bars, white 100 µm. Each image is representative of 5 individual biological samples.

(G) Quantification of the percentage of $VE\text{-}Cad^+$ ECs expressing tdT. Data are the means ± SEM; n= 6. ****$P < 0.0001$ by student's *t*-test.

Figure 5. *R26-loxCre-tdT* enables CreER to recombine *R26-Confetti* efficiently

(A) Schematics showing the experimental design. In *Cdh5-CreER;R26-loxCre-tdT;R26-Confetti* mice, Tam-induced CreER-loxp recombination switches *R26-loxCre-tdT* into *R26-mCre-tdT* allele with simultaneous tdT labeling and expression of mCre, which subsequently targets *R26-Confetti* (right panel). The conventional *Cdh5-CreER;R26-Confetti* mice are used as control.

(B) A schematic showing the experimental strategy.

(C) Whole-mount YFP and mCFP fluorescence images of retina collected from two mice groups. Scale bars, white 100 µm. Each image is representative of 5 individual biological samples.

(D) Immunofluorescence images of YFP, mCFP, and VE-Cad on tissue sections show significantly more YFP^+ and/or $mCFP^+$ ECs in the *Cdh5-CreER;R26-loxCre-tdT;R26-Confetti* mice compared with that of *Cdh5-CreER;R26-Confetti* mice (left panel). The right panel shows the quantification of ECs expressing YFP and/or mCFP. Data are the means ± SEM; n = 5. ****$P < 0.0001$ by student's *t*-test. Scale bars, white 100 µm. Each image is representative of 5 individual biological samples.

Figure 6. *R26-loxCre-tdT* enables *Alb-CreER* to efficiently knockout genes in hepatocytes

(A) Schematics showing the experimental designs for Ctnnb1 gene knockout using either *Alb-CreER* or *Alb-CreER;R26-loxCre-tdT* mice.

(B) A schematic showing the experimental strategy. *Alb-CreER;R26-tdT2;Ctnnb1*$^{flox/flox}$ and *Alb-CreER;R26-loxCre-tdT;Ctnnb1*$^{flox/+}$ mice were injected with Tam for 5 times, while *Alb-CreER;R26-loxCre-tdT;Ctnnb1*$^{flox/flox}$ mice were injected with Tam for one time.

(C) Immunostaining for tdT, β-Catenin, GS, and E-CAD on liver sections. Scale bars, yellow 1mm; white 100 µm. Each image is representative of 5 individual biological samples.

(D) Immunostaining for tdT and CYP2E1 on liver sections. Scale bars, yellow 1mm; white 100 µm. Each image is representative of 5 individual biological samples.

(E) Western blotting of β-Catenin and β-Actin expression in hepatocytes, n = 3.

(F) Quantification of β-Catenin expression. Data are the means ± SEM; n = 5.

(G) qRT-PCR analysis of the relative expression of *Ctnnb1* in the hepatocytes.

(H) qRT-PCR analysis of the relative expression of *Glul*, *Axin2*, *Cyp2e1*, *Oat*, *Tcf7*, *Lect2*, *Tbx3*, and *Slc1a2* in hepatocytes. Data are the means ± SEM; n = 7; ns, non-significant; *$P < 0.05$, **$P < 0.01$, ***$P < 0.001$, ****$P < 0.0001$ by one-way ANOVA.

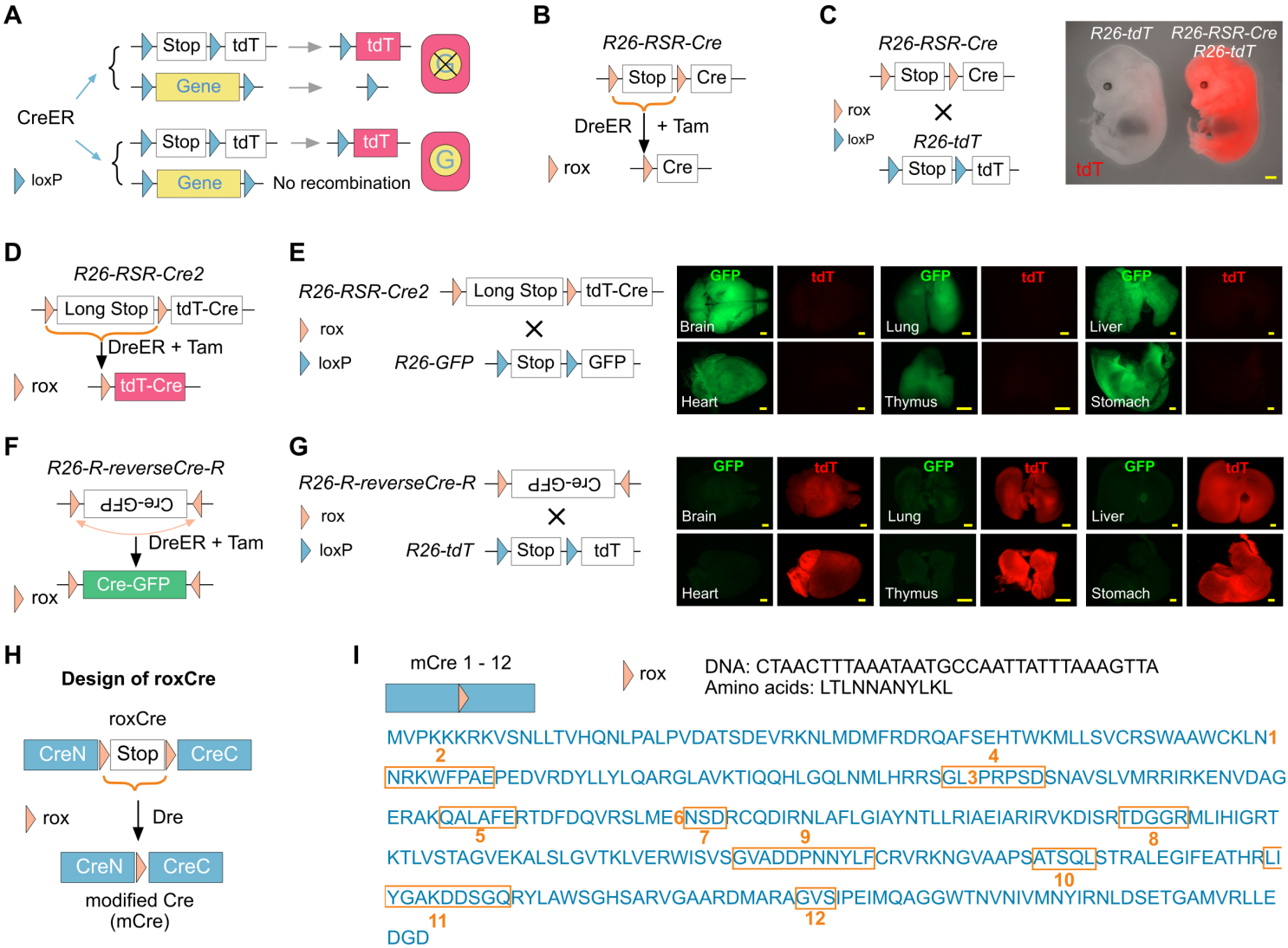
A
Stop
tdT
Gene
CreER
loxP
No recombination
B
R26-RSR-Cre
Stop
Cre
DreER
+ Tam
rox
C
R26-RSR-Cre
R26-tdT
R26-RSR-Cre
R26-tdT
tdT
D
R26-RSR-Cre2
Long Stop
tdT-Cre
DreER + Tam
E
R26-RSR-Cre2
R26-GFP
GFP
tdT
Brain
Lung
Liver
Heart
Thymus
Stomach
F
R26-R-reverseCre-R
Cre-GFP
G
H
Design of roxCre
roxCre
CreN
CreC
Dre
modified Cre
(mCre)
I
mCre 1 - 12
DNA: CTAACTTTAAATAATGCCAATTATTTAAAGTTA
Amino acids: LTLNNANYLKL
MVPKKKRKVSNLLTVHQNLPALPVDATSDEVRKNLMDMFRDRQAFSEHTWKMLLSVCRSWAAWCKLN
NRKWFPAEPEDVRDYLLYLQARGLAVKTIQQHLGQLNMLHRRSGLPRPSDSNAVSLVMRRIRKENVDAG
ERAKQALAFERTDFDQVRSLMENSDRCQDIRNLAFLGIAYNTLLRIAEIARIRVKDISRTDGGRMLIHIGRT
KTLVSTAGVEKALSLGVTKLVERWISVSGVADDPNNYLFCRVRKNGVAAPSATSQLSTRALEGIFEATHRLI
YGAKDDSGQRYLAWSGHSARVGAARDMARAGVSIPEIMQAGGWTNVNIVMNYIRNLDSETGAMVRLLE
DGD
J
Stop
GFP
Cre
loxP
L
GFP %
mCre
No Cre
Cre
K
Mutant Cre (mCre)
Control
GFP DAPI
No Cre
Cre

Figure1

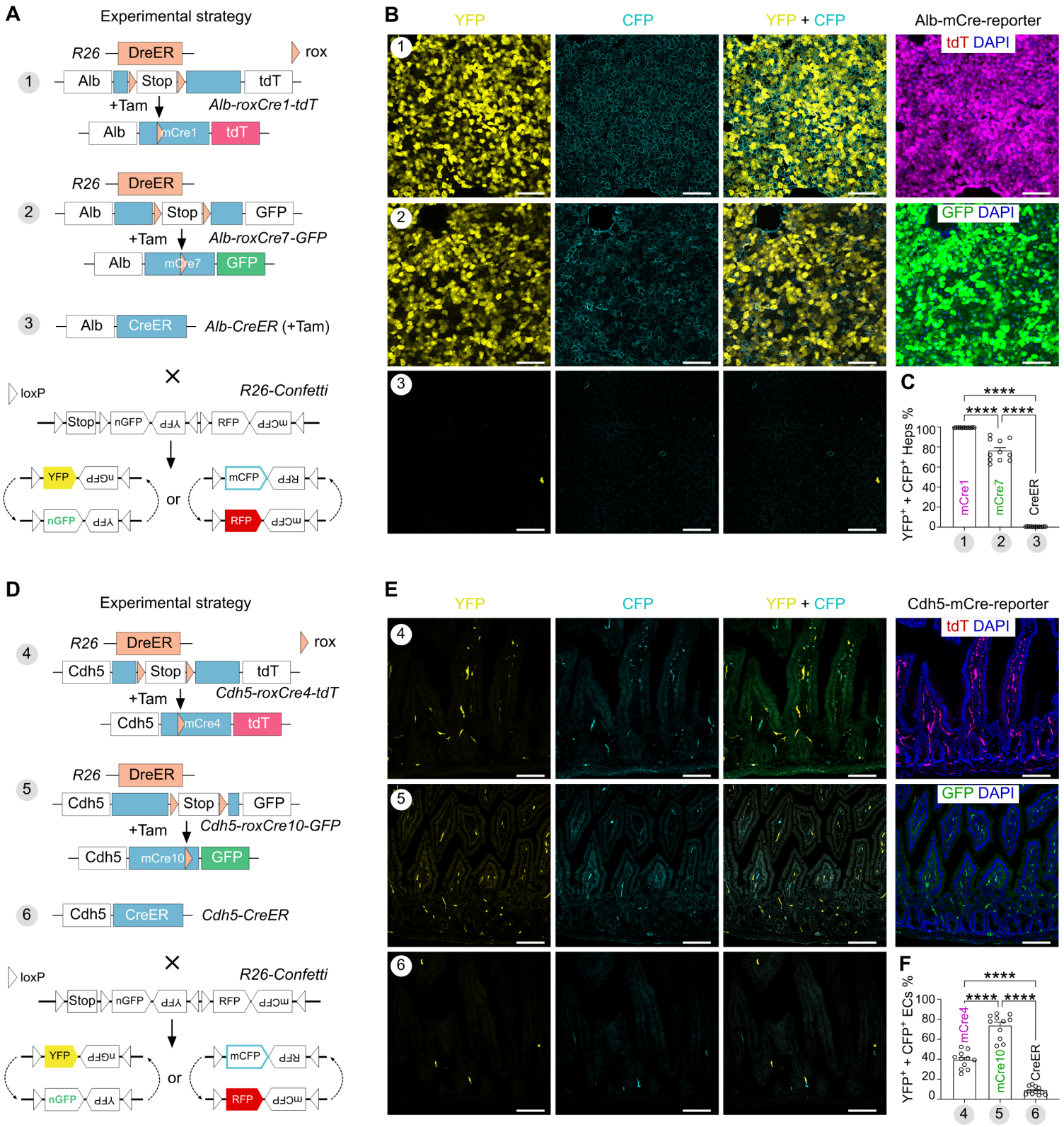
A
Experimental strategy
R26
DreER
rox
1
Alb
Stop
tdT
+Tam
Alb-roxCre1-tdT
mCre1
2
GFP
Alb-roxCre7-GFP
mCre7
3
CreER
Alb-CreER (+Tam)
loxP
R26-Confetti
nGFP
YFP
RFP
mCFP
or
B
YFP
CFP
YFP + CFP
Alb-mCre-reporter
tdT DAPI
GFP DAPI
C
YFP+ + CFP+ Heps %
****
mCre1
mCre7
CreER
D
Experimental strategy
4
Cdh5
Cdh5-roxCre4-tdT
mCre4
5
Cdh5-roxCre10-GFP
mCre10
6
Cdh5-CreER
E
Cdh5-mCre-reporter
F
YFP+ + CFP+ ECs %

Figure2

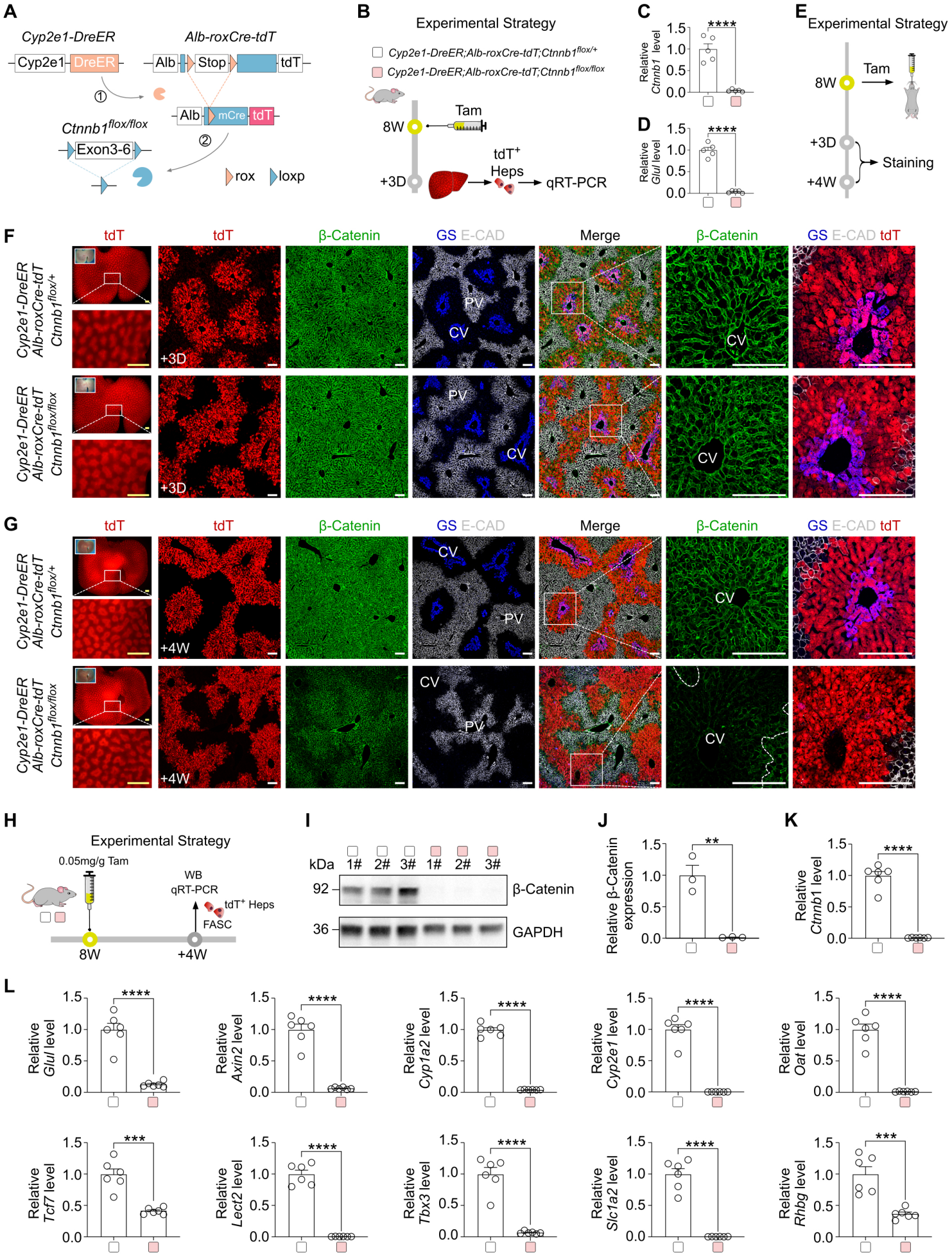
A
Cyp2e1-DreER
Alb-roxCre-tdT
Cyp2e1
DreER
Alb
Stop
tdT
Alb
mCre
tdT
Ctnnb1flox/flox
Exon3-6
rox
loxp
B
Experimental Strategy
Cyp2e1-DreER;Alb-roxCre-tdT;Ctnnb1flox/+
Cyp2e1-DreER;Alb-roxCre-tdT;Ctnnb1flox/flox
Tam
8W
+3D
tdT+ Heps
qRT-PCR
C
Relative Ctnnb1 level
D
Relative Glul level
E
Experimental Strategy
Tam
8W
+3D
+4W
Staining
F
tdT
β-Catenin
GS E-CAD
Merge
GS E-CAD tdT
Cyp2e1-DreER Alb-roxCre-tdT Ctnnb1flox/+
Cyp2e1-DreER Alb-roxCre-tdT Ctnnb1flox/flox
+3D
PV
CV
G
+4W
H
Experimental Strategy
0.05mg/g Tam
WB
qRT-PCR
tdT+ Heps
FASC
8W
+4W
I
kDa
1# 2# 3# 1# 2# 3#
92
β-Catenin
36
GAPDH
J
Relative β-Catenin expression
K
Relative Ctnnb1 level
L
Relative Glul level
Relative Axin2 level
Relative Cyp1a2 level
Relative Cyp2e1 level
Relative Oat level
Relative Tcf7 level
Relative Lect2 level
Relative Tbx3 level
Relative Slc1a2 level
Relative Rhbg level

Figure3

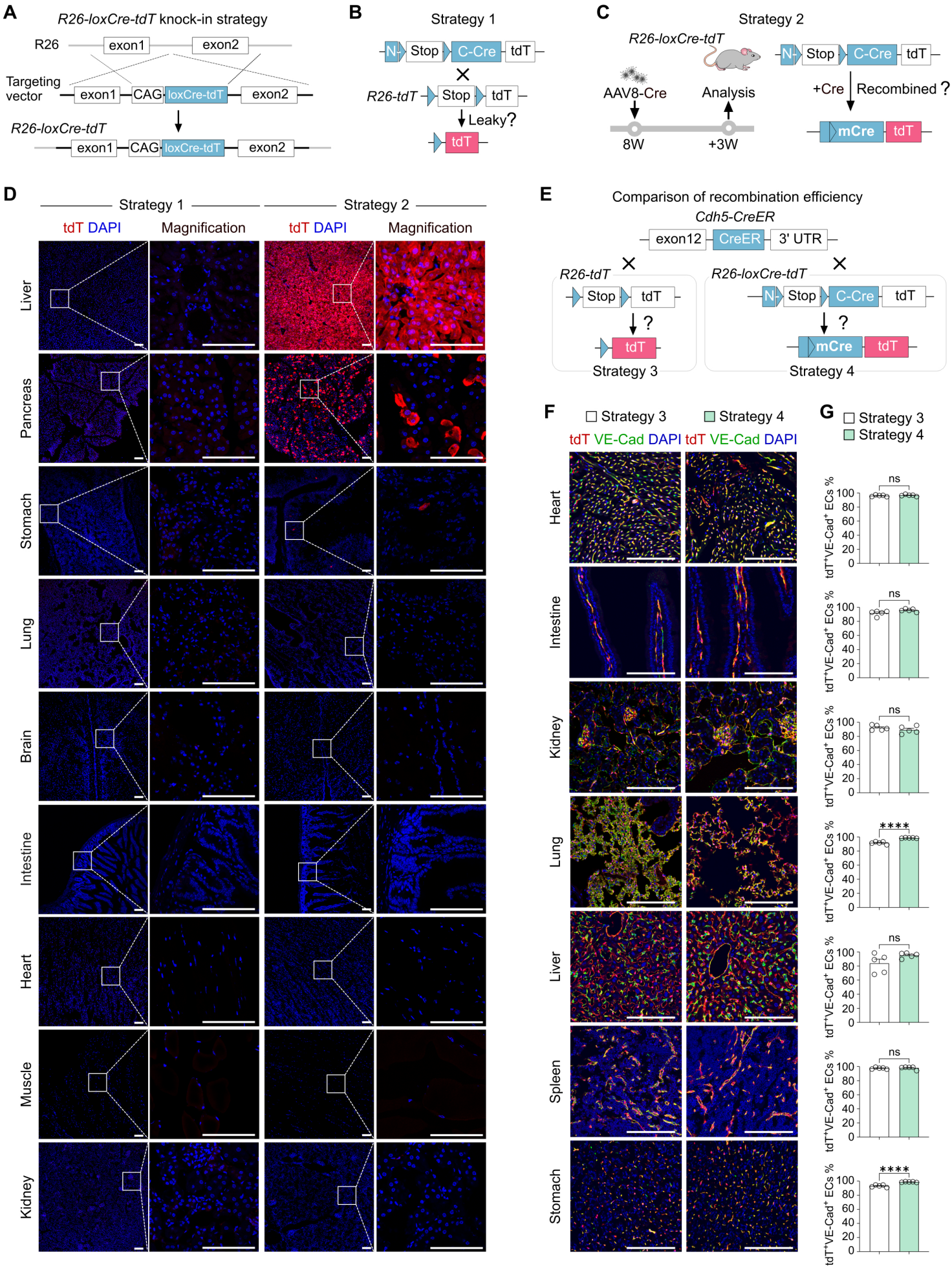
A
R26-loxCre-tdT knock-in strategy
R26
exon1
exon2
Targeting vector
exon1
CAG
loxCre-tdT
exon2
R26-loxCre-tdT
exon1
CAG
loxCre-tdT
exon2
B
Strategy 1
N
Stop
C-Cre
tdT
R26-tdT
Stop
tdT
Leaky?
tdT
C
Strategy 2
R26-loxCre-tdT
AAV8-Cre
Analysis
8W
+3W
N
Stop
C-Cre
tdT
+Cre
Recombined ?
mCre
tdT
D
Strategy 1
Strategy 2
tdT DAPI
Magnification
tdT DAPI
Magnification
Liver
Pancreas
Stomach
Lung
Brain
Intestine
Heart
Muscle
Kidney
E
Comparison of recombination efficiency
Cdh5-CreER
exon12
CreER
3' UTR
R26-tdT
Stop
tdT
?
tdT
Strategy 3
R26-loxCre-tdT
N
Stop
C-Cre
tdT
?
mCre
tdT
Strategy 4
F
Strategy 3
Strategy 4
tdT VE-Cad DAPI
tdT VE-Cad DAPI
Heart
Intestine
Kidney
Lung
Liver
Spleen
Stomach
G
Strategy 3
Strategy 4
tdT+VE-Cad+ ECs %
100
80
60
40
20
0
ns
****

Figure4

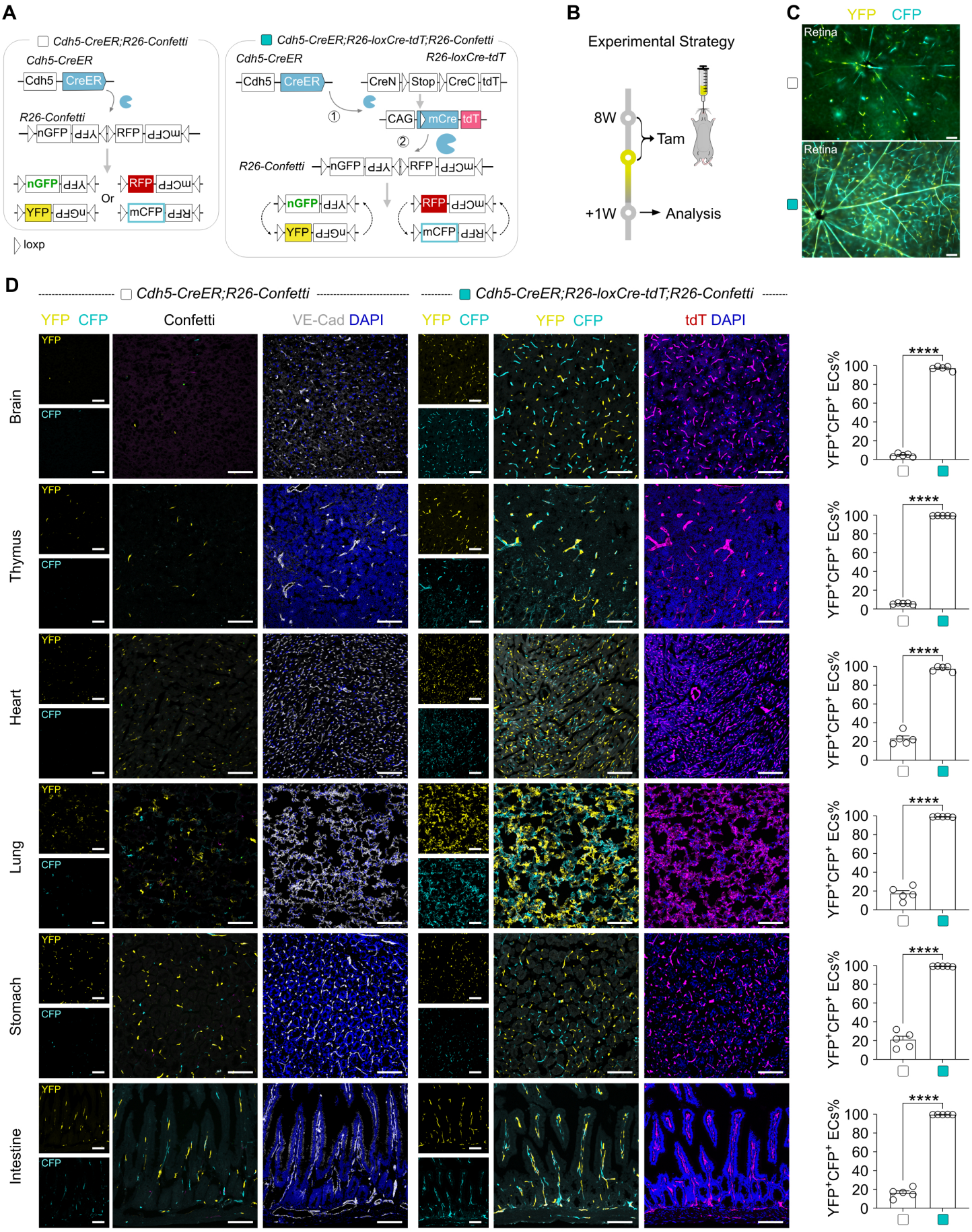
A
Cdh5-CreER;R26-Confetti
Cdh5-CreER
Cdh5
CreER
R26-Confetti
nGFP
YFP
RFP
mCFP
Or
loxp
Cdh5-CreER;R26-loxCre-tdT;R26-Confetti
R26-loxCre-tdT
CreN
Stop
CreC
tdT
CAG
mCre
①
②
B
Experimental Strategy
8W
Tam
+1W
Analysis
C
YFP CFP
Retina
D
Confetti
VE-Cad DAPI
tdT DAPI
Brain
Thymus
Heart
Lung
Stomach
Intestine
YFP+CFP+ ECs%
****

Figure5

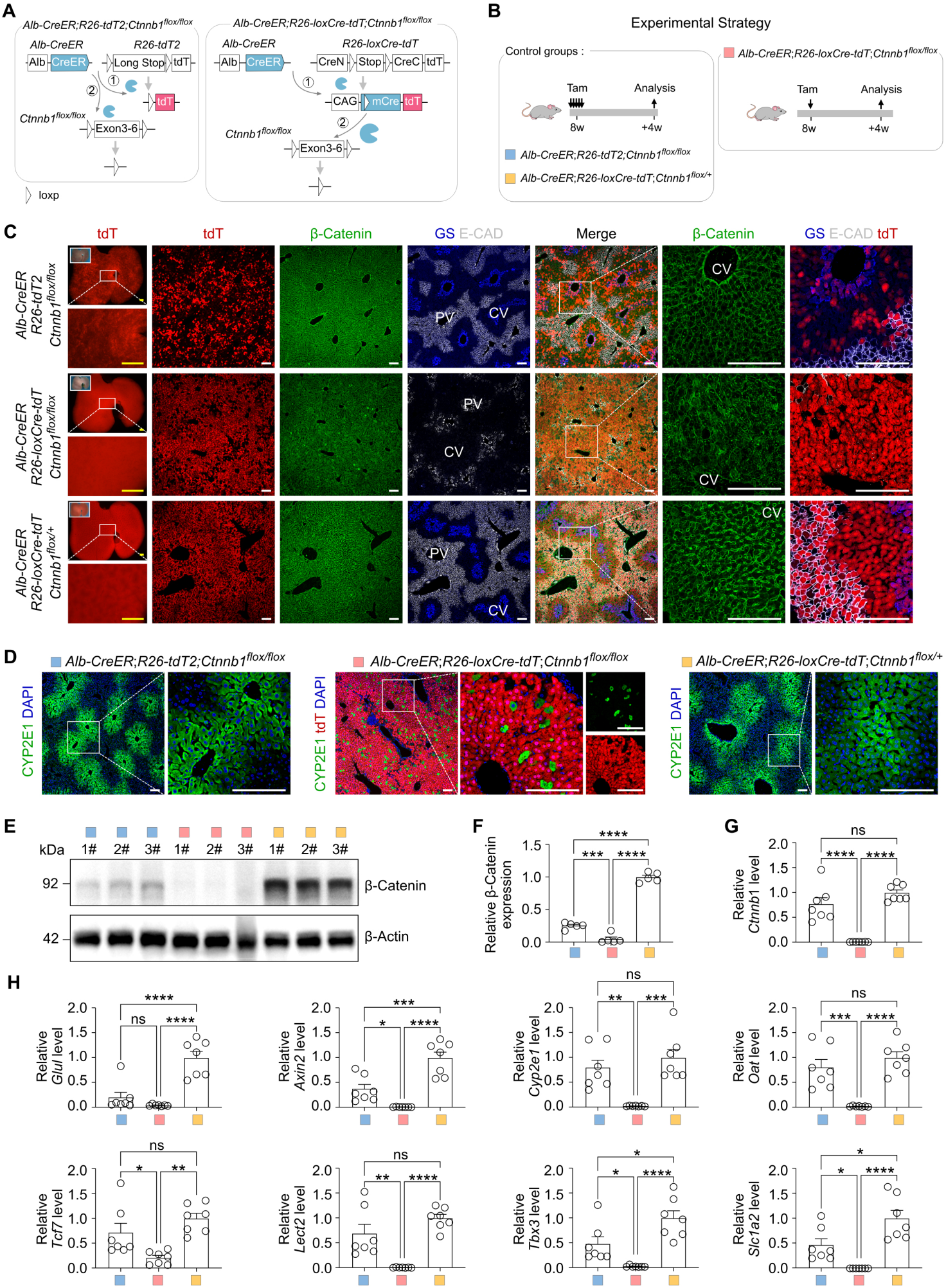
A
Alb-CreER;R26-tdT2;Ctnnb1flox/flox
Alb-CreER
R26-tdT2
Alb
CreER
Long Stop
tdT
Ctnnb1flox/flox
Exon3-6
Alb-CreER;R26-loxCre-tdT;Ctnnb1flox/flox
R26-loxCre-tdT
CreN
Stop
CreC
CAG
mCre
loxp
B
Experimental Strategy
Control groups :
Tam
Analysis
8w
+4w
Alb-CreER;R26-tdT2;Ctnnb1flox/flox
Alb-CreER;R26-loxCre-tdT;Ctnnb1flox/+
C
tdT
β-Catenin
GS E-CAD
Merge
PV
CV
D
CYP2E1 DAPI
CYP2E1 tdT DAPI
E
kDa
1#
2#
3#
92
42
β-Catenin
β-Actin
F
Relative β-Catenin expression
G
Relative Ctnnb1 level
H
Relative Glul level
Relative Axin2 level
Relative Cyp2e1 level
Relative Oat level
Relative Tcf7 level
Relative Lect2 level
Relative Tbx3 level
Relative Slc1a2 level
ns
*
**
***
****

Figure6