



Published in final edited form as:

Transgenic Res. 2017 June ; 26(3): 411–417. doi:10.1007/s11248-017-0018-1.

Spontaneous Recombinase Activity of Cre-ERT2 *In Vivo*

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Abstract

Inducible Cre-ERT recombinase technology is widely used for gene targeting studies. The second generation of inducible Cre-ERT recombinase, hemizygous B6.129S-Tg(UBC-cre/ERT2)1Ejb/J (hereafter abbreviated as *Cre-ERT2*), a fusion of a mutated estrogen receptor and Cre recombinase, was engineered to be more efficient and specific than the original Cre-ERT. The putative mechanism of selective Cre-mediated recombination is Cre sequestration in the cytoplasm in the basal state with translocation to the nucleus only in the presence of tamoxifen. We utilized both a reporter mouse (B6.129 (Cg)-*Gt(ROSA)26Sor^{tm4}(ACTB-tdTomato,-EGFP)Luo/J*) and endothelin converting enzyme-1 (*Ece1*) floxed transgenic mouse line to evaluate Cre-ERT2 activity. We observed spontaneous Cre activity in both settings. Unintended Cre activity is a confounding factor that has a potentially large impact on data interpretation. Thus, it is important to consider background Cre activity in experimental design.

Keywords

Tamoxifen; Estrogen Receptor; Recombination; Cre-activity

Introduction

The development of conditional knockout alleles has enabled the study of gene function in a variety of settings that were previously beyond the scope of simple gene-targeting strategies. Tissue-specific Cre drivers permit the assessment of anatomically restricted gene ablation and examination of embryonically lethal mutations in in post-natal animals. Furthermore, drug regulated Cre strains (Feil et al., 1997; Gossen and Bujard, 1992; Johansen et al., 2002; Pluta et al., 2005) can also be used to circumvent developmental phenotypes by delaying gene ablation until a desired time point.

Inducible Cre-ERT recombinase is one of several systems that has allowed researchers to manipulate gene expression in a time dependent manner. The second generation of inducible Cre-ERT recombinase was developed to be more sensitive and specific than the original Cre-ERT (Indra et al., 1999; Jaisser, 2000). Hemizygous B6.129S-Tg (UBC-cre/ERT2)1Ejb/J (hereafter abbreviated as *Cre-ERT2*), a second generation inducible Cre-ERT recombinase, mouse line was generated through lentiviral transgenesis (Lois et al., 2002; Ruzankina et al., 2007). The construct has the common feature of a Cre recombinase-mutated estrogen receptor (Cre-ERT2) protein fusion that is expressed under the control of the Ubiquitin C (UBC) promoter (Ruzankina et al., 2007). Previous *in vitro* experiments showed that the Cre-ERT2 was limited to the cytoplasm in the absence of tamoxifen (Feil et al., 1997). Past reports have also claimed that Cre-ERT2 is activated exclusively by tamoxifen or 4-hydroxytamoxifen (Feil et al., 1997; Indra et al., 1999; Ruzankina et al., 2007). In the presence of tamoxifen, Cre-ERT2 translocates to the nucleus and mediates recombination (Feil et al., 1997; Indra et al., 1999), but in its absence Cre-ERT2 is limited to the cytoplasm. Background Cre activity can preclude the accurate interpretation of gene function. Therefore, it is important to assess Cre-recombinase activity in the absence of tamoxifen.

Reporter mice are often used to evaluate the spatial and temporal extent of Cre-mediated recombination. In a double fluorescent Cre recombinase reporter mouse B6.129(Cg)-*Gt(ROSA)26Sor^{tm4}(ACTB-tdTomato,-EGFP)Luo/J* (hereafter abbreviated as *mT/mG*), cells express membrane targeted tdTomato (*mT*) prior to Cre excision and membrane targeted enhanced green fluorescent protein (EGFP) (*mG*) following Cre excision (Muzumdar et al., 2007). The *mT/mG* cassette is expressed under the control of the CMV β -actin enhanced promoter (pCA). The double fluorescent marker system allows visualization of both recombined and non-recombined cells.

We used the *mT/mG* mouse line to evaluate Cre-ERT2 activity *in vivo*. In addition, we also utilized *Ece1^{flox/flox}* mice to confirm Cre activity in another targeted gene system. *Ece1* encodes for endothelin converting enzyme-1, a key enzyme in the proteolytic processing of big endothelins (big ETs) to mature, active endothelins (ETs). Here, we report that Cre-ERT2 exhibits spontaneous activity *in vivo*, and that its extent varies among individual animals and among tissues within the same animal.

Materials and Methods

Targeted Mouse Line Production

The generation of *mT/mG* and *Cre-ERT2* mice has been described (Lois et al., 2002; Muzumdar et al., 2007; Ruzankina et al., 2007). Both *mT/mG* and *Cre-ERT2* mice were purchased from Jackson Laboratory. *mT/mG* and *Cre-ERT2* mice were intercrossed to generate *mT/mG* and *Cre-ERT2 mT/mG* mice (Fig. 1a). By 8 ± 1 weeks, the mice were either left untreated, or treated with vehicle (98% corn oil, 2% ethanol) or tamoxifen (MP Biomedicals) at a concentration of 0.5 μ M/g of body weight dissolved in 98% corn oil and 2% ethanol once daily for 5 days via intraperitoneal (IP) injection. Both *mT/mG* and *Cre-ERT2 mT/mG* were maintained to an age of 10 ± 1 weeks.

The conditional *Ece1* knockout mouse construct was purchased from the knockout mouse project (KOMP). The construct was made on a C57BL/6N background. Four generation backcrossing with C57BL/6J were performed to generate *Ece1* KOMP mice with C57BL/6J background. The *Ece1* KOMP mice were then initially bred to transgenic mice harboring FLP recombinase (B6.Cg-Tg (ACTFLPe)9205Dym/J, purchased from Jackson laboratory) and subsequently bred to EIIA Cre recombinase (B6.FVB-Tg(EIIA-cre)C5379Lmgd/J, purchased from Jackson laboratory) mice to generate the *Ece1^{fllox}* and *Ece1* alleles, respectively. *Ece1^{+/fllox}* mice were intercrossed to generate *Ece1^{fllox/fllox}* mice. *Ece1^{-/-}* mice were then intercrossed with *Cre-ERT2* mice to generate *Cre-ERT2 Ece1^{-/-}* mice. *Ece1^{fllox/fllox}* and *Cre-ERT2 Ece1^{-/-}* mice were intercrossed to generate *Ece1^{+fllox}*, *Ece1^{-fllox}*, *Cre-ERT2 Ece1^{+fllox}* and *Cre-ERT2 Ece1^{-fllox}* mice (Fig. 1b). All experimental animals were left untreated or injected with either vehicle or tamoxifen at 8 ± 1 weeks of age.

Mice were housed 2–5 mice/500 cm² cage with a 12 h light-dark cycle, given acidified water, and fed laboratory rodent chow 5001 (PMI Nutrition International, Richmond, IN, USA) *ad libitum* according to the guidelines set forth by “The Guide for the Care and Use of Laboratory Animals”. All animal procedures were approved by the William S. Middleton VA Hospital Institutional Animal Care and Use Committee (IACUC).

Genotyping

DNA was isolated from tail snips and tissue samples using the IBI genomic DNA mini kit (Peosta, IA USA) following manufacturer’s protocol. For the PCR reaction, we used Failsafe PCR system (Epicentre Technologies, Madison WI) with PreMix J for the (*mT/mG* transgene) and Premix H for *Ece1^{fllox}* allele genotyping.

The *mT/mG* transgene and *Cre-ERT2* allele were identified by PCR from tail DNA preparations. *Ece1* KOMP, *Ece1^{fllox}* and *Ece1* alleles were genotyped by PCR using tail and tissue DNA preparations. The list of primers and PCR setting is summarized in Table 1.

The transgene containing the *Cre* allele was identified by quantitative Real-time PCR reactions, performed in a StepOne RT-PCR instrument (Life Technologies, Carlsbad, CA, USA) using TaqMan assay (Life Technologies, Carlsbad, CA, USA) following manufacturer’s recommended protocol. The TaqMan copy number assay is Mr00635245_cn (Life Technologies, Carlsbad, CA, USA).

Tissue Preparation and Histology

Mice were euthanized using isoflurane followed by cervical dislocation. For cryosection preparation, kidney, liver, heart and lung were perfused with 4% cold paraformaldehyde in 0.1 M phosphate buffered saline (PBS), fixed for 6-24 hours at 4°C, cryoprotected in 30% sucrose overnight at 4°C and embedded in OCT, following a previously described method (Muzumdar et al., 2007). Samples were stored at –80°C until use. Ten micron sections were obtained using a Leica CM1800 cryostat, washed three times with PBS, counterstained with DAPI, washed an additional three times and mounted.

Data and Image Analysis

Two-dimensional and z-stack confocal images were obtained with a Nikon C1 confocal microscope and EZ-C1 software (v. 3.91) in 1.0 μm slices with a 400x objective. Images were processed with Image J (FIJI).

Results

UBC Cre-ERT2 Activity in *mT/mG* Mice

Reporter mice that only carried the *mT/mG* allele (n=6) did not emit mG fluorescence in the green channel in any of the tissues examined (Fig. 2), indicating that mG labeling was Cre dependent as previously shown (Muzumdar et al., 2007). In the tamoxifen treated *Cre-ERT2 mT/mG* mice (n=6), we observed approximately ~50-90% of mG positive cells in all tissues examined (Fig. 2). Both vehicle (n=4) and untreated *Cre-ERT2 mT/mG* (n=4) mice also unexpectedly showed >50% and 10-80% mG positive cells, respectively, in the kidney and liver (Fig. 2, 3). In the kidney, mG positive cells were observed primarily in the mesangium and tubules. In the liver, mG labeling was observed in the hepatocytes. In heart and lung, there were ~5-30% of mG positive cells observed in untreated groups for both tissues (Fig. 2, 3). Moreover, mG labeling was mainly observed in the cardiomyocytes and in the alveoli and bronchi of the lung. There was also a small degree of overlap in the localization of mT and mG fluorescence, suggesting co-localization of recombined and non-recombined cells. The degree of Cre efficiency in the double transgenic progeny of *mT/mG* and *Cre-ERT2* mice varied among animals. Spontaneous Cre activity was detected in all tissues examined in untreated *Cre-ERT2 mT/mG* mice (Fig. 2, 3). mG positive cells were detected in the untreated and vehicle treated *Cre-ERT2 mT/mG* mice indicating that Cre-mediated recombination occurred in the absence of tamoxifen. Furthermore, *mG* expression in the untreated group excludes unrecognized estrogenic activity of the corn oil/ethanol vehicle as the basis for Cre-mediated recombination in the absence of tamoxifen.

Cre-ERT2 Activity in *Ece1^{flox/flox}* Mice

To verify our findings on the Cre activity of *Cre-ERT2* mice, we used an *Ece1^{flox/flox}* targeted mouse line. We monitored for the presence of the various *Ece1* alleles under various conditions. We compared tail DNA samples of untreated *Cre-ERT2 Ece1^{+/flox}* (n=3) mice with *Ece1^{+/flox}* (n=3) and *Cre-ERT2 Ece1^{-/flox}* (n=3) at 3 weeks of age. Using 5 μl PCR reaction samples, we did not detect the *Ece1* allele band (~300 bp) in *Cre-ERT2 Ece1^{+/flox}* and *Ece1^{+/flox}* DNA (Fig. 4a). However, when 20 μl PCR reaction samples were electrophoresed, the *Ece1* allele band was detected in some of *Cre-ERT2 Ece1^{+/flox}* DNA samples (Fig. 4b). We also observed a diminishing *Ece1^{flox}* allele band (~1100 bp) and a more prominent *Ece1* allele band in the 17-week old vehicle treated *Cre-ERT2 Ece1^{-/flox}* mice (n=2) in both the liver and kidney using 5 μl PCR reaction volumes (Fig. 4c). Collectively, these results indicated spontaneous Cre activity in the absence of tamoxifen and suggest continued Cre activity over time.

Discussion

Using the *mT/mG* reporter mice, our data show that the UBC *Cre-ERT2* mouse line has substantial tamoxifen-independent Cre activity. The Cre-recombinase activity also varies among individuals, tissues and cell types. There are several possible explanations for these findings. Some organs may have higher expression of the reporter gene, higher expression of Cre-recombinase or a combination of the two. The Cre may have higher efficiency in specific cell types because the levels of circulating tamoxifen or endogenous estrogen are higher in certain tissues. There are also differences in cell turnover rates and blood flow in different organ systems that need to be considered. Regardless of the factors that may contribute to the variability of the Cre activity, our study demonstrates that *Cre-ERT2* is active in the absence of tamoxifen. It also further confirms previous finding on tamoxifen independent UBC-driven Cre-ERT2 activity *in vivo* (Seime et al., 2015). Sanger sequencing of the UBC-driven *Cre-ERT2* gene did not show any mutations that could potentially render the Cre-ERT2 perpetually active (data not shown). Many of the Cre-ERT2 lines have been reported to have spontaneous Cre recombination regardless of its specific promoter (Leone et al., 2003; Mishra et al., 2015; Papoutsi et al., 2015; Seime et al., 2015), which demonstrate that it is an inherent property of the fusion protein. The ubiquitin C promoter is widely and highly expressed, which enabled us to detect ligand independent Cre-recombinase activity in many tissues at surprisingly high levels. Our findings differ from those reported previously in showing more generalized and more extensive Cre-recombinase activity. Thus, investigators should consider the impact of spontaneous Cre activity on their studies.

To address the potential difference in Cre-recombination efficiency between the reporter construct and a specific floxed gene, we performed genomic DNA genotyping for the *Ece1* allele using our *Ece1* floxed mouse constructs. In the untreated *Cre-ERT2 Ece1^{+/-}flox* mice, the *Ece1* allele was observed at 3 weeks of age, before tamoxifen treatment. Collectively, these findings demonstrate the need for caution of unintended recombination activity in many inducible Cre strains. Although the use of tamoxifen increases the Cre efficiency, the results stress the importance on the use of appropriate controls and the inclusion of both sexes when studying conditional knockout alleles.

Spontaneous Cre activity can be a confounding factor in analyzing gene function and must be considered during experimental design. Without the use of appropriate controls, meaningful results can be overlooked. In many cases, the use of only a vehicle treated control group may not be sufficient to characterize the phenotypic outcomes of homozygous gene deletion. Due to the unintended intrinsic Cre activity, vehicle treated controls may have a comparable degree of gene deletion to tamoxifen treated animals, masking significant genotype-dependent effects. As a result, some changes in phenotypes may be understated. Littermates that do not harbor the Cre transgene are valuable controls that should be included in tamoxifen inducible gene ablation studies.

These experiments have both strengths and limitations. The strengths include the use of both reporter mice and inducible *Ece1* knockout mice to evaluate Cre activity in two targeted gene systems. We also examined the Cre-recombinase activity in multiple tissues

allowing for comparison of the Cre efficiency in different organ systems. We excluded vehicle as a possible inducer of Cre activity. A study limitation is that we did not perform quantitative genotyping for *Ece1* alleles; thus, we were unable to confirm the effect of sexual dimorphism on the Cre-ERT2 activity. Further experiments are required to quantitatively identify possible sexual dimorphism of spontaneous Cre-recombinase activity. Additionally, we only examined Cre activity over a small range of times. Therefore, we were unable to address the accumulation of recombined cells beyond young adulthood. Recent findings show that cross contamination of exogenous tamoxifen can occur in animals that are housed together through exposure to the general environment and/or coprophagous behavior (Brake et al., 2004). In our study, there were untreated and tamoxifen treated mice that were housed in the same cage. Thus, we were unable to exclude the possibility of tamoxifen cross contamination may contribute to the Cre activity observed in the untreated animals. However, the detection of the *Ece1* allele in 3 week old mice, prior to drug treatment, demonstrates that coprophagia is not a sufficient explanation for spontaneous Cre recombinase activity.

Despite the shortcomings of some Cre constructs, the Cre-recombinase system has been proven to be a very useful tool in biological studies. The Cre-ERT2 system has been used to deduce many gene functions over the years. Although new approaches may be required to refine this powerful tool, fully understanding the extent of Cre activity will help researchers better characterize the outcomes of their studies.

Acknowledgements

We would like to thank Baozhi Yuan and Harmut Weiler for helpful discussions, and Aron Geurts and Michael Grzybowski for Sanger Sequencing and analysis. This material is based on work supported in part by award I21 RX1440, by the Office of Research and Development, Rehabilitation Research and Development Service, and a Merit Review Award from the Office of Research and Development, Biomedical and Laboratory Research and Development Service, the Department of Veterans Affairs, and performed in the Geriatrics Research, Education, and Clinical Center at the William S. Middleton Memorial Veterans Hospital. The views expressed here are those of the authors, and the contents do not represent the views of the U.S. Department of Veterans Affairs or the United States Government.

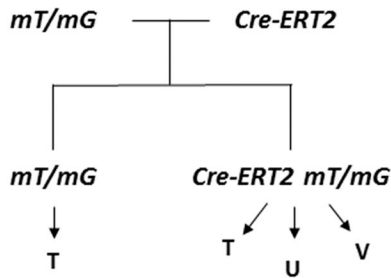
Research reported in this publication was supported in part by National Institute of Arthritis and Musculoskeletal and Skin Diseases of the National Institute of Health under Award Number NIH AR54753. The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institute of Health.

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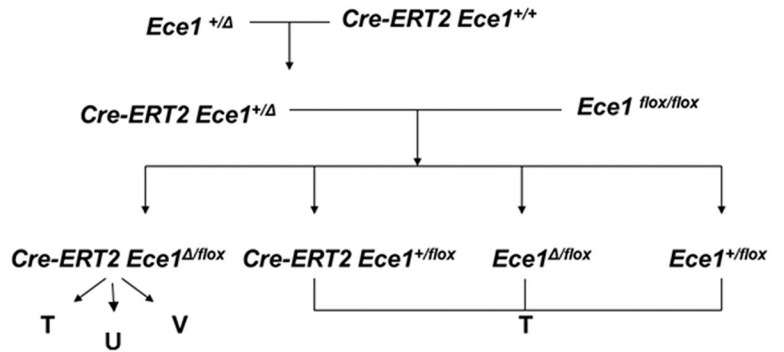
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a Generation of inducible *Cre-ERT2 mT/mG* Mice



b Generation of inducible *Ece1* KO Mice



Untreated (U)

Vehicle (V)

Tamoxifen (T)= 0.5 μ M/g of body weight

Fig. 1. Generation of *Cre-ERT2 mT/mG* mice and inducible *Ece1* Knockout (KO) mice.

a A schematic showing the generation of *Cre-ERT2 mT/mG* mice. Mice harboring *mT/mG* gene were treated with tamoxifen. Mice harboring *Cre-ERT2 mT/mG* gene were left untreated or treated with tamoxifen or vehicle at 8 ± 1 weeks of age. **b** A schematic showing the generation of inducible *Ece1* KO mice. T Mice harboring *Ece1^{flox/flox}* allele were intercrossed with *Cre-ERT2 Ece1^{+/-}* mice to generate the experimental animals. These mice were then treated with either tamoxifen or vehicle at 8 ± 1 weeks of age.

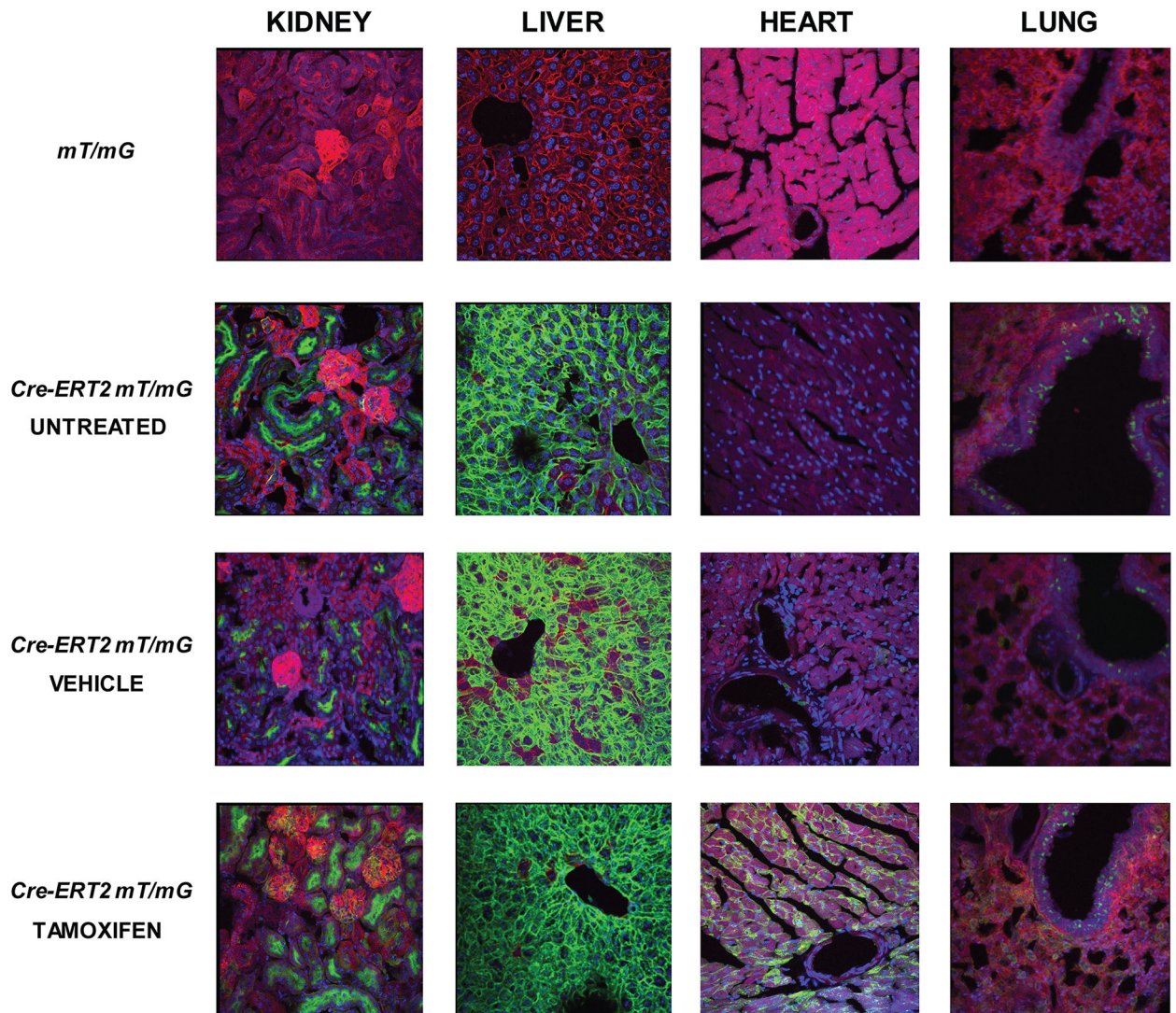


Fig 2. mT/mG labeling of *mT/mG* and *Cre-ERT2 mT/mG* mice.

Fixed tissue sections of *mT/mG* mice (n=6) did not show mG labeling for all tissues examined. Fixed tissue sections of *Cre-ERT2 mT/mG* mice showed mosaic pattern of mT and mG labeling in the kidney, liver, heart and lung tissue sections. Tamoxifen treated *Cre-ERT2 mT/mG* mice (n=6) showed > 50% Cre efficiency as represented by the percent of mG labeling in all tissues examined. mG labeling was observed in tissue sections of *Cre-ERT2 mT/mG* untreated (n=4) and vehicle (n=4) treated groups, indicating spontaneous Cre activity in the absence of tamoxifen.

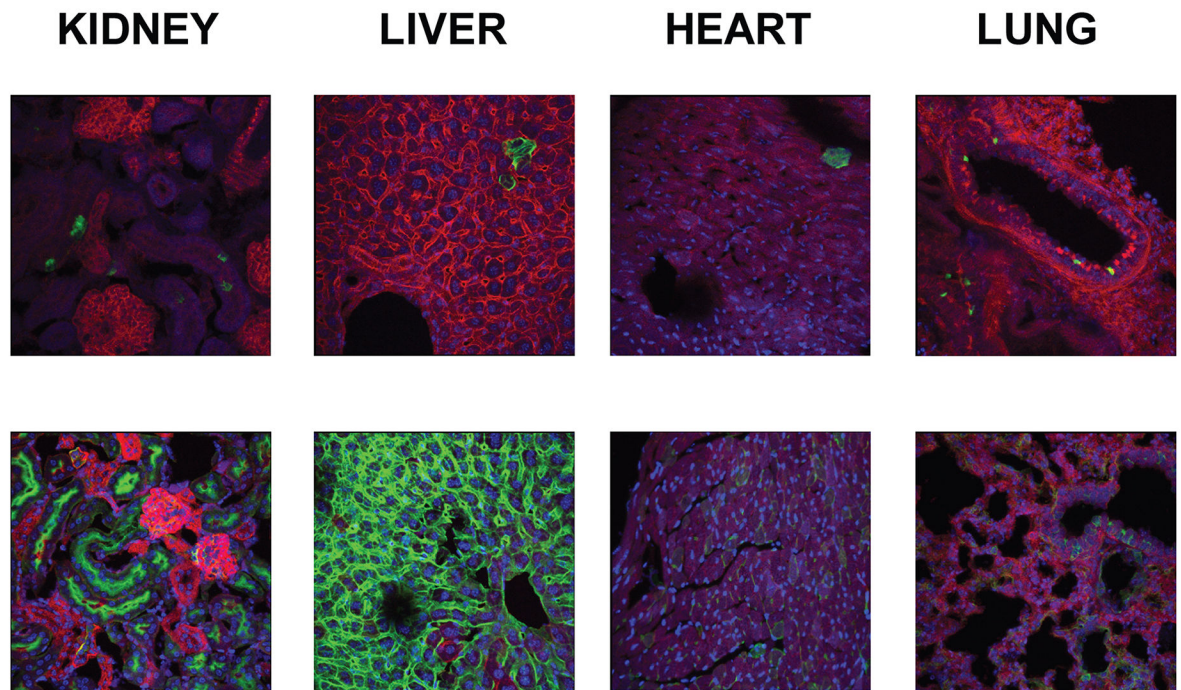


Fig 3. Cre activity in untreated 10 ± 1 weeks old *Cre-ERT2 mT/mG* mice.

These are representative images of tissue sections from two different animal samples (I, II) to show the range of the spontaneous Cre activity. Spontaneous Cre activity was observed in kidney, liver, heart, and lung tissue sections. Additionally, the Cre activity appears to be animal dependent.

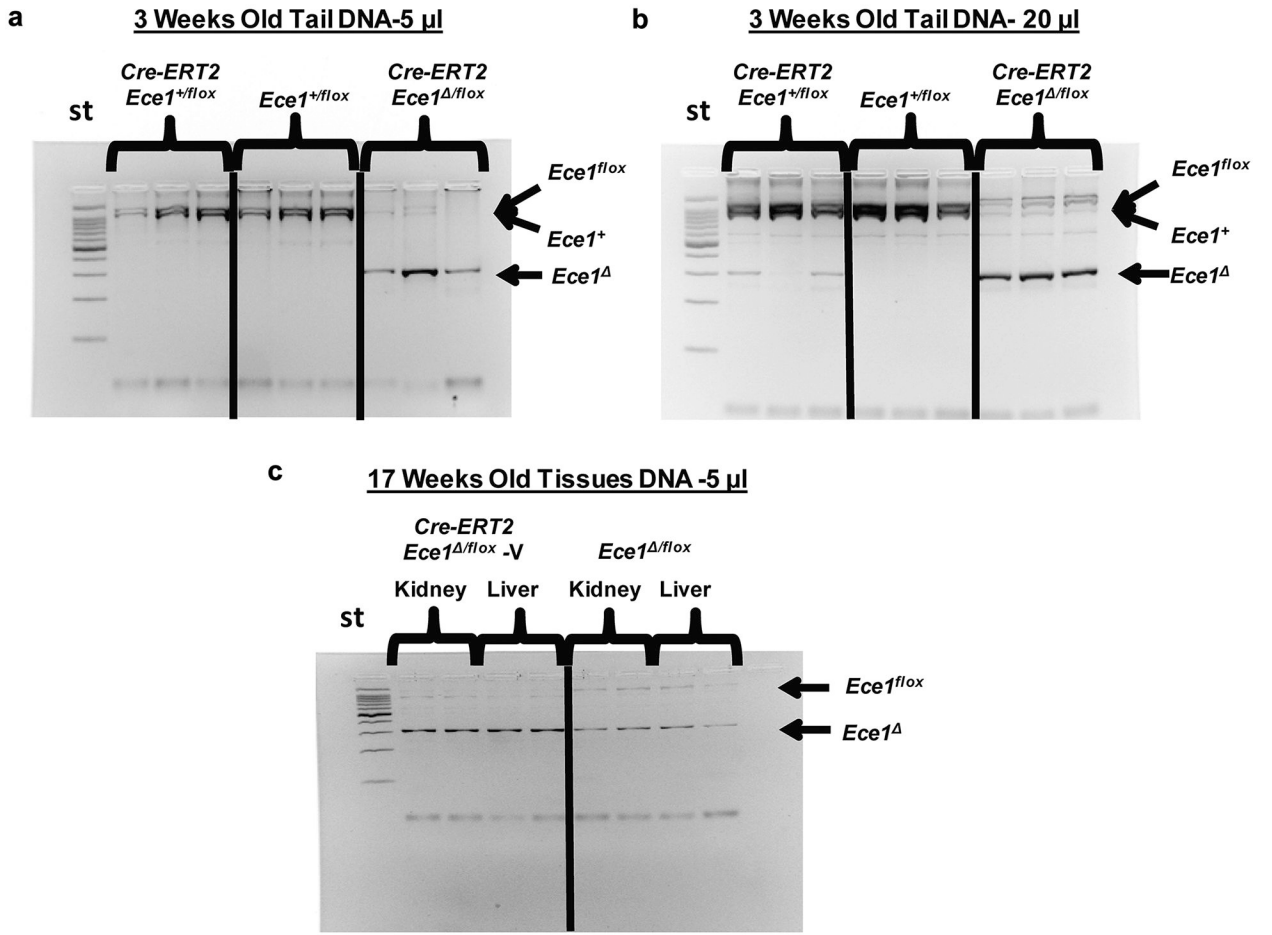


Fig 4. Cre activity analysis in *Ece1* floxed mice.

St= standard, *Ece1^{flox}* = ~1100 bp, *Ece1⁺* = ~1000 bp, *Ece1^Δ* = ~300 bp (a) Using a 5 µl PCR reaction volume, genomic DNA from 3 weeks old tail preparations showed an undetectable *Ece1* band in the untreated *Cre-ERT2 Ece1^{+/flox}* (n=3) and *Ece1^{+/flox}* (n=3) samples, while a weaker *Ece1^{flox}* band and a more intense *Ece1* band were observed in the untreated *Cre-ERT2 Ece1^{Δ/flox}* samples (n=3). (b) Using a 20 µl PCR reaction volume, genomic DNA from untreated 3 weeks old tail preparation showed *Ece1* band detected in *Cre-ERT2 Ece1^{+/flox}* mice (n=3), while none was observed in the *Ece1^{+/flox}* samples (n=3). (c) Genomic DNA from 17 week old mice kidney and liver showed a less intense *Ece1^{flox}* band and a more intense *Ece1* band for vehicle treated *Cre-ERT2 Ece1^{Δ/flox}*.

Table 1.

Primers and PCR conditions

Target	Primers	Annealing Temperature
<i>Gt(Rosa)26Sor (mT/mG)</i>	5' CCA TAT ATG GAG TTC CGC GT 3' 5' TAA CCT GGT GTG TGG GCG TTG T 3' 5' CGG GCC ATT TAC CGT AAG TTA T 3'	61 °C /45 sec For 35 cycles
<i>EcoI Exon 6</i>	5' TTA GTC AGG GTT ACT ATT GCT G 3' 5' TAG AGA AGG ATT AGT CAG TAC AG 3'	60°C/ 30 sec –0.5°C/Cycle For 10 cycles Then 50°C/ 30 sec For 40 cycles

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