

C57BL/6JGpt-Pf4-P2A-iCre

Strain Name: C57BL/6JGpt-Pf4em1Cin(P2A-iCre)/Gpt

Strain Type: Knock-in Strain Number: T005328 Background: C57BL/6JGpt

Description

This mouse strain expresses codon optimized iCre recombinase [1] under the control of the mouse endogenous *Pf4* promoter, P2A-iCre was inserted upstream of the stop codon TAG of mouse *Pf4* gene by CRISPR/Cas9 technology. When crossed with a strain with loxP site flanked sequence in its genome, Cre-mediated recombination will result in excision of the DNA fragment between the two loxPs in megakaryocytes and platelets. Recombinase activity was also detected in a proportion of cells in lung, spleen, kidney, liver, intestine and heart.

Strategy

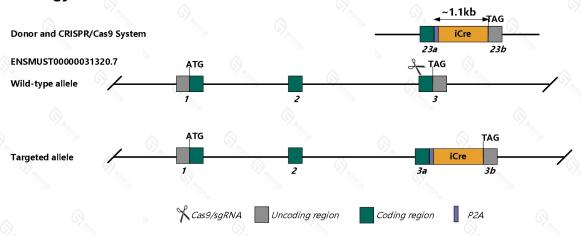


Fig.1 Schematic diagram of C57BL/6JGpt-Pf4-P2A-iCre model strategy.

Applications

1. Cre tool mice for specific induction of loxP recombination in megakaryocytes and platelets [2].

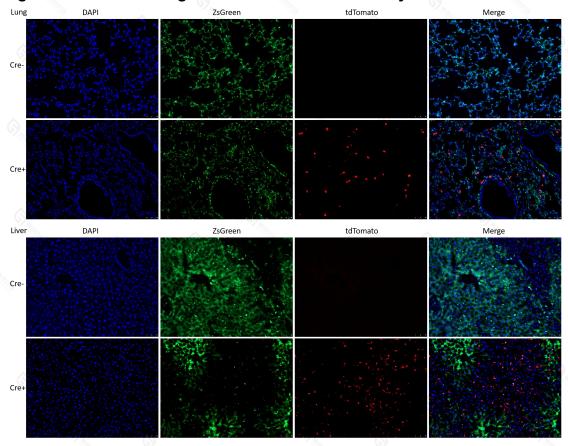


Data support

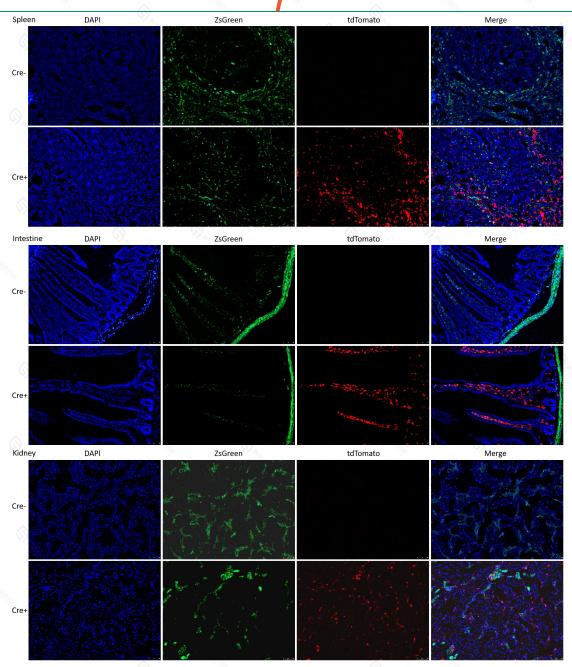
1. Validation methods & notes

Pf4-P2A-iCre mice was crossed with CAG-loxp-ZsGreen-Stop-loxp-tdTomato mice with ubiquitous reporter expression (hereafter referred as CAG-G/R mice), Cre-mediated recombination will lead to excision of ZsGreen and the stop cassette and expression of tdTomato, thus loss of green fluorescence and gain of red fluorescence will indicate Cre activity. Fluorescence imaging of frozen sections were performed to exhibit Cre activity in various tissues and organs. Imaging of sections were performed under a 200x microscopy. Note: these results may only represent the activity of Cre in this strain at the identical stage. Recombinase activity may be different at other stages in your application.

2. Images of tissues and organs with obvious Cre activity









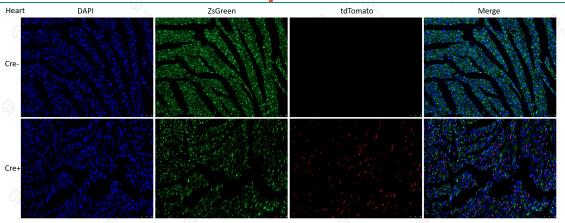


Fig 2. Fluorescence imaging of tissues and organs with obvious Cre activity.

Organ name was indicated in the left top of each subfigure group. Cre-: CAG-G/R single positive individuals; Cre+: Pf4-P2A-iCre, CAG-G/R double positive individuals.

3. Images of tissues and organs with little or no Cre activity

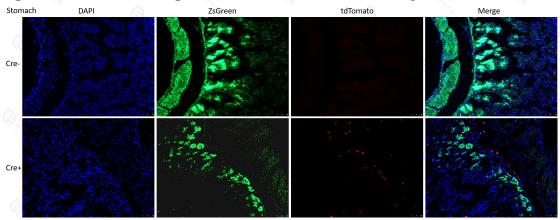


Fig 3. Fluorescence imaging of tissues and organs with little or no Cre activity.

Organ name was indicated in the left top of each subfigure group. Cre-: CAG-G/R single positive individuals; Cre+: Pf4-P2A-iCre, CAG-G/R double positive individuals.

4. Flow cytometry analysis of cells with Cre activity



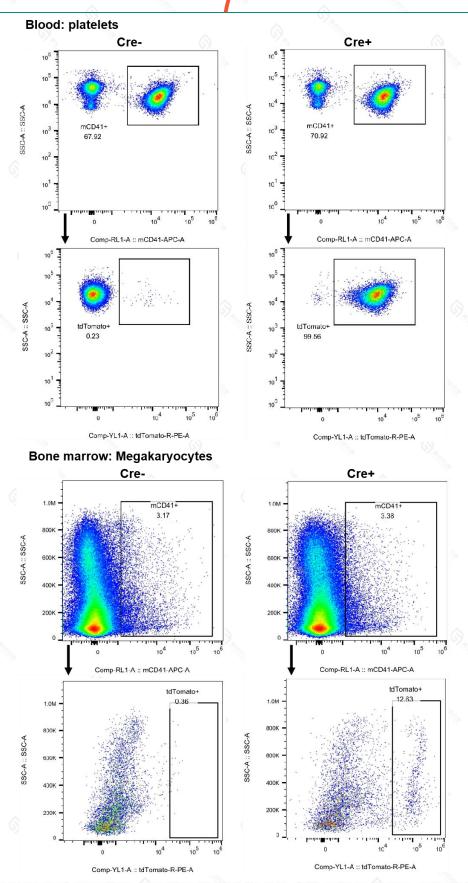




Fig 4. Flow cytometry analysis of cells with Cre activity

Organ name was indicated in the left top of each subfigure group. Cre-: CAG-G/R single positive individuals; Cre+: Pf4-P2A-iCre, CAG-G/R double positive individuals. Bone marrow-derived cells and whole blood cells were harvested and analyzed for tdTomato expression with flow cytometry.

Reference

- 1.Shimshek D R, Kim J, Hübner M R, et al. "Codon-improved Cre recombinase (iCre) expression in the mouse." genesis, 2002, 32(1): 19-26.
- 2.Tiedt R, Schomber T, Hao-Shen H, et al. Pf4-Cre transgenic mice allow the generation of lineage-restricted gene knockouts for studying megakaryocyte and platelet function in vivo. Blood, 2007, 109(4): 1503-6.