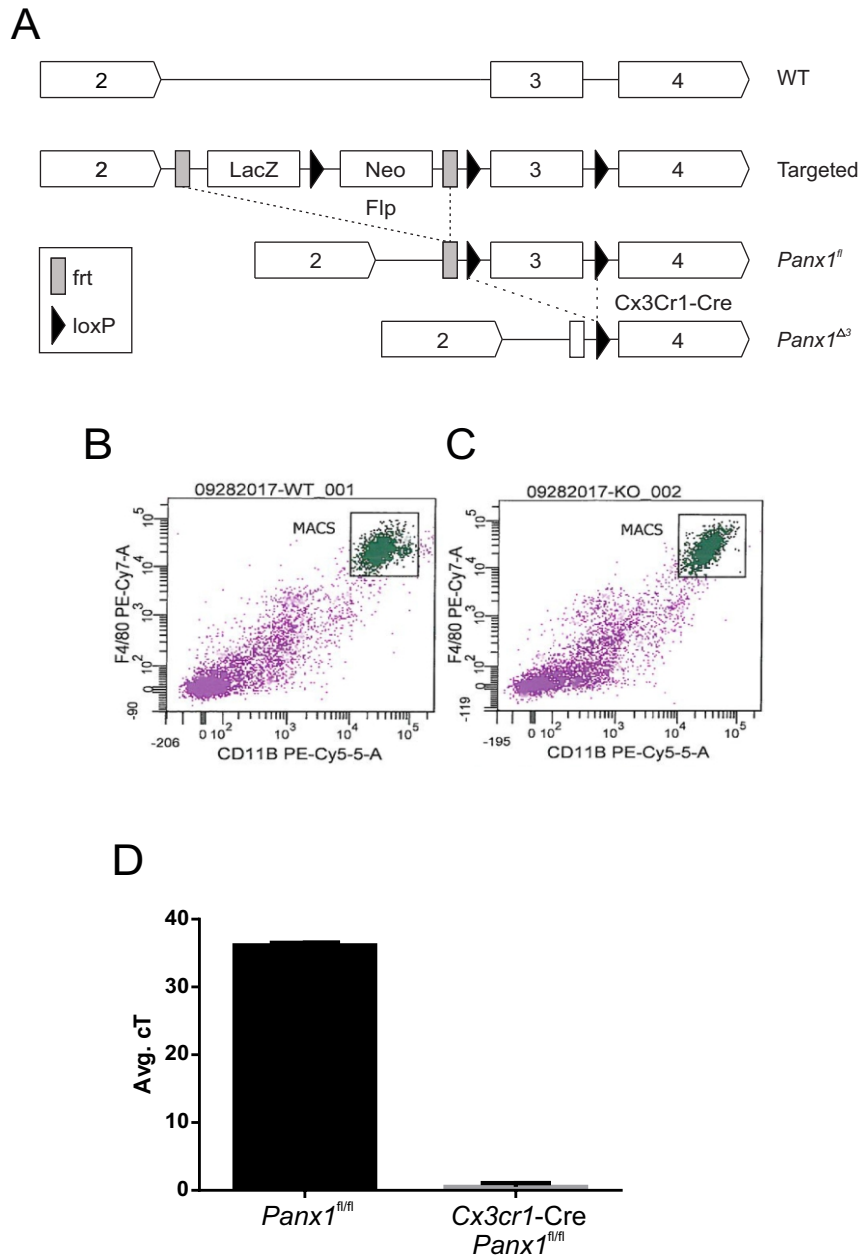


Supplementary Figure 1



Supplementary Fig. 1 Generation of *Cx3cr1-Cre::Panx1*^{fl/fl} transgenic mice

Mice harboring *Panx1* deletion was generated using the cre-loxP system (34). *Panx1*^{tm1a(KOMP)Wtsi} mice containing 2 loxP sequences flanking exon 3 of the *Panx1* gene were obtained from KOMP repository, and then bred to mice harboring FLP1 recombinase gene under the control of the human ACTB promoter (Jax mice: ACTB:FLPe B6N, stock number 019100) to generate homozygous *Panx1*^{fl/fl} mice. These mice were crossed with C57BL/6J mice that express Cre recombinase under the direction of the *Cx3cr1* promoter (Jax mice: B6N(Cg)-*Cx3cr1*^{tm1.1(cre)Jung/J}, stock number 025524) and then backcrossed 8 generations to yield the myeloid conditional knockout *Cx3cr1-Cre::Panx1*^{fl/fl} mice. To generate myeloid reporter mice with *Panx1* deletion mutation, we crossed *Panx1*^{fl/fl} and *Cx3cr1-Cre::Panx1*^{fl/fl} mice with *Cx3Cr1*^{EGFP/EGFP} mice.

(A) Targeting strategy for conditional deletion of *Panx1*. Peripheral macrophages were isolated using FACS from **(B)** Floxed and **(C)** conditional- knockout mice. **(D)** mRNA expression of wild-type *panx1* in peritoneal macrophages. Statistical significance was evaluated using one way ; $n \geq 3$ per group. N.D. Non-detected