Genes | Genomes | Genetics

A Multifunctional Mutagenesis System for Analysis of Gene Function in Zebrafish

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Figure S1 Number of inserts per line. Southern analysis was performed on 42 representative trapped lines to estimate the number of inserts per line. A) Representative Southern blot analysis. Lane 1: Positive control containing vector DNA and AB genomic DNA. Lane 2: Negative control containing AB genomic DNA. Lanes 3-11: Outcrosses lines *Tg(DsDELGT4)ws0188* and *Tg(DsDELGT4)ws0196* showing multiple integrations per line. B) Estimated number of inserts per line.

GFP Enhancer Trap



Figure S2 The spatial-temporal distribution of enhancer trap reporter expression. Graphs show the number of lines expressing GFP in various tissues at 1 dpf, 2 dpf, 3-4 dpf and 5-7 dpf. CNS: central nervous system, MHB: mid-hindbrain boundary, FMB: forebrain-midbrain boundary.



Figure S3 Dynamic expression of the enhancer trap reporter in various tissues. The pie charts show the relative number of lines expressing GFP in various tissues at 1 dpf, 2 dpf, 3-4 dpf and 5-7 dpf. CNS: central nervous system, MHB: mid-hindbrain boundary, FMB: forebrain-midbrain boundary.

mCherry Protein Trap



Figure S4 The spatial-temporal distribution of protein trap reporter expression. Graphs show the number of lines expressing mCherry in various tissues at 1 dpf, 2 dpf, 3-4 dpf and 5-7 dpf. CNS: central nervous system, MHB: mid-hindbrain boundary, FMB: forebrain-midbrain boundary.



Figure S5 Dynamic expression of the protein trap reporter in various tissues. The pie charts show the relative number of lines expressing mCherry in various tissues at 1 dpf, 2 dpf, 3-4 dpf and 5-7 dpf. CNS: central nervous system, MHB: mid-hindbrain boundary, FMB: forebrain-midbrain boundary.

Injection of Cre Capped mRNA into embryos with two insertions on the same chromosome PCR for primer pairs 1,2,3,4 to check chromosome rearrangement in embryo with both *Tg(DsDELGT4)ws0310* and *Tg(DsDELGT4)ws2629* insertions on chr21 after Cre injection Cre/lox induced lox2272 lox2272 deletion 1 I 1 Primer 1 Primer Primer Primer 1 I 1 1 1 1 Pair 4 1 Pair 1 Pair 2 Pair 3

В

А



Figure S6 Cre-lox mediated targeted large deletion. (A) Schematic of a Cre-induced recombination experiment. Cre recombinase capped mRNA was injected into a batch of one-cell stage embryos harboring two insertions on the same chromosome. Individual injected embryos were collected for PCR with six pair of primers to detect deletion events in embryos. In this experiment, the injected embryos carry two insertions *Tg(DsDELGT4)ws0310* and *Tg(DsDELGT4)ws2629* on chr21, 25Mb away from each other. (B) PCR results with six pairs of primers shows the expected deletion in embryo 4 (lane 7). Lane 1 shows PCR results of a wild-type embryo for each primer pair, Lane 2 shows PCR positive control for each primer pair. Lane 3 shows ladders. Lanes 4-9 show PCR products with each primer pair for injected embryos 1-6. Images showing PCR products from embryo 5 and embryo 6 (lanes 8,9) were cropped from a different part of the same gel.

Tables S1-S20

Available for download as Excel files at www.g3journal.org/lookup/suppl/doi:10.1534/g3.114.015842/-/DC1 Table S1 DsDELGT4 lines with reporter expression in liver from day0 to day7 Table S2 DsDELGT4 lines with reporter expression in pancreas from day0 to day7 Table S3 DsDELGT4 lines with reporter expression in intestine from day0 to day7 Table S4 DsDELGT4 lines with reporter expression in swim bladder from day0 to day7 Table S5 DsDELGT4 lines with reporter expression notochord from day0 to day7
 Table S6
 DsDELGT4 lines with reporter expression in pronephric duct from day0 to day7
Table S7 DsDELGT4 lines with reporter expression in skeletal muscle cells from day0 to day7
 Table S8
 DsDELGT4 lines with reporter expression in heart tube from day0 to day7
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