

Supplementary Data

Untargeted Metabolomics of Slc13a5 Deficiency Reveal Critical Liver-Brain Axis for Lipid Homeostasis

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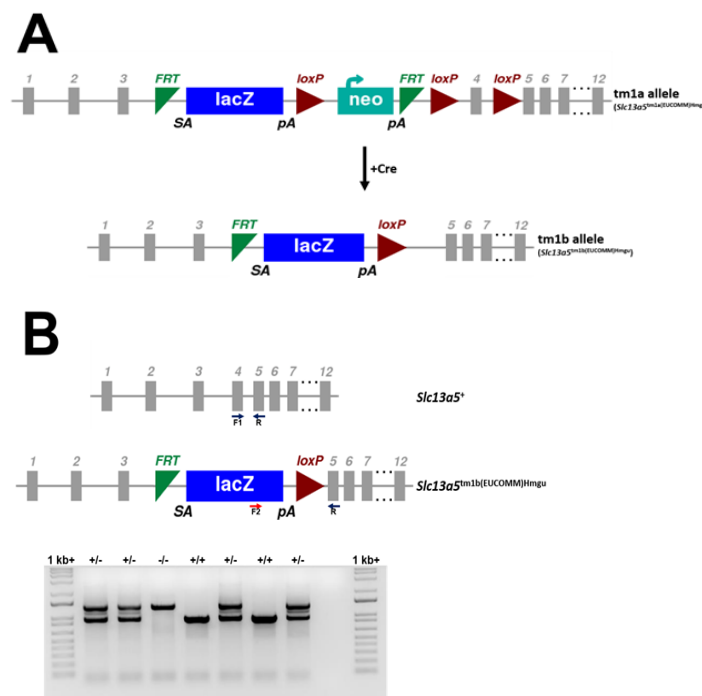


Figure S1. Generation and Genotyping of Slc13a5 Deficiency Mice. **A.** EUCOMM production of mouse model of Slc13a5 deficiency (C57BL/6N-Slc13a5^{tm1b}(EUCOMM)Hmgu/Ieg). The *tm1a* allele has a *lacZ* gene trap cassette and a neomycin resistant transgene flanked by indicated FRT and loxP sites inserted in the intron between exons 3 and 4 of *Slc13a5*, as well as an additional loxP site inserted in the intron between exons 4 and 5. The *Slc13a5*^{tm1b} mutant allele is generated by Cre-mediated excision of *Slc13a5* exon 4 and the *neo* cassette from the *Slc13a5*^{tm1a} allele, which results in a frameshift mutation, leaving the *lacZ* gene trap intact. Partial genomic maps of *Slc13a5*^{tm1a} and *Slc13a5*^{tm1b} alleles are depicted. **B.** Genotyping of Slc13a5-deficient mice (C57BL/6N-Slc13a5^{tm1b}(EUCOMM)Hmgu/Ieg). The wild type allele PCR amplicon is 919 bp, and the mutant allele PCR amplicon is 1301 bp. Partial genomic maps of *Slc13a5*⁺ and *Slc13a5*^{tm1b} alleles with indicated PCR primer binding sites, as well as a representative picture of agarose gel electrophoretic analysis of PCR genotyping results, are shown. “1 kb+” = 1 kb Plus DNA ladder. “+/+” = *Slc13a5*^{+/+}; “+/-” = *Slc13a5*^{+/tm1b}; and “-/-” = *Slc13a5*^{tm1b/tm1b} [1].

1. Skarnes WC, Rosen B, West AP, Koutsourakis M, Bushell W, Iyer V, Mujica AO, Thomas M, Harrow J, Cox T, et al. A conditional knockout resource for the genome-wide study of mouse gene function. *Nature*, 2011. 474(7351): p. 337-42.

Online eTable S1. Slc13a5 Deficiency Mouse Tissue Metabolite Flux. Interactive table: [https://app.powerbi.com/view?](https://app.powerbi.com/view?r=eyJrIjojN2I4YTM3ZjMtZGE1MS00ODk0LThlNmItYWZlY2IzZjY2ZGE1IiwidCI6IjZiYmUyYmZmLWNjMzQtNDBIYi05OTZmLWZiNGY2ZjI5MTdiZiIsImMiOiJ9)

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