

# Aldh7A1-KO

|                     |                                                |
|---------------------|------------------------------------------------|
| <b>Nomenclature</b> | C57BL/6Smoc- <i>Aldh7A1</i> <sup>em1Smoc</sup> |
| <b>Cat. NO.</b>     | TBD                                            |
| <b>Strain State</b> | Developing                                     |

## Gene Summary

|                                      |                       |                                    |
|--------------------------------------|-----------------------|------------------------------------|
| <b>Gene Symbol</b><br><b>Aldh7a1</b> | <b>Synonyms</b>       | Atq1, D18Wsu181e                   |
|                                      | <b>NCBI ID</b>        | <a href="#">110695</a>             |
|                                      | <b>MGI ID</b>         | <a href="#">108186</a>             |
|                                      | <b>Ensembl ID</b>     | <a href="#">ENSMUSG00000053644</a> |
|                                      | <b>Human Ortholog</b> | ALDH7A1                            |

## Model Description

Exon 2-5 of Aldh7A1 gene was deleted to generate Aldh7A1 knockout mice.

\*Literature published using this strain should indicate: Aldh7A1-KO mice (Cat. NO. TBD) were purchased from Shanghai Model Organisms Center, Inc..

## Disease Connection

|                                      |                     |                                                                                                                                                                                                                                                                                                                          |
|--------------------------------------|---------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>Pyridoxine-Dependent Epilepsy</b> | <b>Phenotype(s)</b> | <a href="#">MGI:6491217</a>                                                                                                                                                                                                                                                                                              |
|                                      | <b>Reference(s)</b> | Al-Shekaili HH, Petkau TL, Pena I, Lengyell TC, Verhoeven-Duif NM, Ciapaite J, Bosma M, van Faassen M, Kema IP, Horvath G, Ross C, Simpson EM, Friedman JM, van Karnebeek C, Leavitt BR, A novel mouse model for pyridoxine-dependent epilepsy due to antiquitin deficiency. Hum Mol Genet. 2020 Nov 25;29(19):3266-3284 |

## Validation Data

No data