

STOCK *Atp13a2*^{tm1.1Wtd} /J

Stock No: 028387 | *Atp13a2* floxed

 Targeted Mutation

Typically mice are recovered in 10-14 weeks. Contact Customer Service to place an order or for more information.

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endolysosomal dysfunction, α -synuclein accumulation, and neurodegeneration associated with the onset of Parkinson's disease.

Donating Investigator

William Dauer, University of Michigan Medical School

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GENETIC OVERVIEW

Genetic Background

Generation

Atp13a2^{tm1.1Wtd}

Alele Type

Targeted (Conditional ready (e.g. floxed), No functional change)

Gene Symbol

Atp13a2

Gene Name

ATPase type 13A2

VIEW GENETICS

RESEARCH APPLICATIONS

Neurobiology Research

Research Tools

VIEW ALL RESEARCH APPLICATIONS

BASE PRICE

Starting at:

\$2,854.50 Domestic price Cryo Recovery

V I E W P R I C E L I S T

Details

Detailed Description

Atp13a2 floxed mice have exons 2-3 of the ATPase type 13A2 (*Atp13a2*) gene flanked by *loxP* site, ATP13A2 encodes a transmembrane lysosomal ATPase that is involved in intracellular cation homeostasis. Mutations in *Atp13a2* have been associated with the onset of Kufor–Rakeb syndrome (KRS), characterized by early-onset Parkinsonism with diffuse cerebral and cerebellar atrophy and neurodegeneration. Homozygotes are viable and fertile. When crossed with a strain expressing Cre Recombinase, resulting offspring will contain a premature stop codon, abolishing gene expression.

Atp13a2 null mice exhibit age-related neuropathological changes, including reactive astrocytosis, lipofuscinosis, protein aggregation, and lysosomal accumulation in multiple brain regions, including the cortex, cerebellum, hippocampus, and striatum by 1 month of age. This was followed by onset of lipofuscinosis and autofluorescence by 3 months of age, and the accumulation of lysosomal proteins LAMP1, LAMP2, and lysosomal lipid BMP by 6 months. The aggregation of ubiquitinated proteins and p62 is evident by 12 months. No α -synuclein related abnormalities are observed in mice up to 18 months of age. Modulating α -synuclein levels by intercrossing these mice with α -synuclein null mice or α -synuclein overexpressing mice (B6;C3-Tg(Prnp-SNCA*A53T)83Vle/J Stock No. [004479](#)) did not change the onset or extent of pathological change.

Development

Control Suggestions

Selected References

Genetics

Atp13a2^{tm1.1Wtd}

⊖ Disease/Phenotype

[+ Disease Terms](#)

[+ Research Areas By Phenotype](#)

[+ Mammalian Phenotype Terms by Genotype](#)

[+ References](#)

⊖ Technical Support

C O N T A C T T E C H N I C A L S U P P O R T

Genotyping Protocols

Standard PCR:[Atp13a2](#)

[Genotyping resources and troubleshooting](#)

Breeding Considerations

When maintaining a live colony, homozygous mice may be bred together.

[Additional Breeding and Husbandry Support](#)

Citation

When using the Atp13a2 floxed mouse strain in a publication, please [cite the originating article\(s\)](#) and include JAX stock #028387 in your Materials and Methods section.

Animal Health Reports

[Facility Barrier Level Descriptions](#)

Production of mice from cryopreserved embryos or sperm occurs in a maximum barrier room, [G200](#)

⊖ Pricing & Availability



Cryo

Typically mice are recovered in 10-14 weeks. Contact Customer Service to place an order or for more information.

Domestic International

Pricing effective for USA, Canada and Mexico shipping destinations

CRYORECOVERY - DOMESTIC PRICING

SERVICE/PRODUCT	DESCRIPTION	PRICE
Cryo Recovery	Heterozygous or wildtype for Atp13a2<tm1.1Wtd>	\$2,854.50

RELATED PRODUCTS AND SERVICES

Frozen Mouse Embryo	STOCK Atp13a2<tm1.1Wtd>/J Frozen Embryo	\$2595.00
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TERMS OF USE

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Q U E S T I O N S A B O U T T E R M S O F U S E

ADDITIONAL USE RESTRICTIONS APPLY

[Use of MICE by companies or for-profit entities requires a no-fee JAX Leap License prior to shipping.](#)

LICENSING INFORMATION

Phone: 207-288-6470

Email: TechTran@jax.org

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