

B6.129S7-B4galt1^{tm2Shur} /J

Stock No: **006943**

 Congenic, Targeted Mutation

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glycosidic molecular interactions and function, and cell-to-extracellular matrix (ECM) interactions.

Donating Investigator

Barry D Shur, Emory University

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GENETIC OVERVIEW

Genetic Background

Generation

B4galt1^{tm2Shur}

Alele Type

Targeted (Modified isoform(s))

Gene Symbol

B4galt1

Gene Name

UDP-Gal:betaGlcNAc beta 1,4- galactosyltransferase, polypeptide 1

VIEW GENETICS

RESEARCH APPLICATIONS

Cell Biology Research

Endocrine Deficiency Research

Reproductive Biology Research

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

These mice carry a mutant allele that has a point mutation in the first translation initiation codon in exon 1, which initiates translation of the long isoform of beta 1,4-galactosyltransferase. The second translation initiation codon in exon 1 is not affected. These mice express only the shorter isoform of beta 1,4-galactosyltransferase. No long isoform protein is detected in mammary tissue by Western blot analysis. Mice that are homozygous for the targeted mutation are viable, fertile, normal in size and do not display any gross physical or behavioral abnormalities. Total Beta 1,4-galactosyltransferase activity is reduced to 72% of wildtype levels in mammary gland epithelial cells while activity on mammary epithelial cell surfaces is diminished by over 60%. Sperm and testis exhibit near wildtype levels of enzyme activity and glycoprotein galactosylation. The short isoform is expressed ectopically in sperm. Although able to undergo normal acrosomal exocytosis induced by calcium ionophore, homozygous sperm do not exhibit acrosome reaction to zona pellucida glycoproteins or anti-galactosyltransferase antibodies. Homozygous females exhibit abnormal mammary gland morphology including excessive mammary epithelial duct branching. Approximately 20% of homozygous females do not support their young. Mutants have elevated levels of expression of metalloproteinases (Mmp14 and Mmp7). Laminin expression in basal lamina is abnormal. This mutant mouse strain may be useful in studies of glycosidic molecular interactions and function, and cell-to-extracellular matrix (ECM) interactions.

This strain was transferred from the collection of the Consortium for Functional Glycomics.

Development

Control Suggestions

Selected References

Genetics

B4galt1^{tm2Shur}

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

Pyrosequencing:[B4galt1](#)

[Genotyping resources and troubleshooting](#)

Breeding Considerations

When maintaining a live colony, these mice can be bred as homozygotes.

[Additional Breeding and Husbandry Support](#)

Citation

When using the B6.129S7-*B4galt1*^{tm2Shur}/J mouse strain in a publication, please [cite the originating article\(s\)](#) and include JAX stock #006943 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
Recovery

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CRYORECOVERY - DOMESTIC NOT-FOR-PROFIT & ACADEMIC PRICING

SERVICE/PRODUCT	DESCRIPTION	PRICE
Cryo Recovery	Heterozygous for B4galt1<tm2Shur>	\$2,854.50

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LICENSING INFORMATION

Phone: 207-288-6470

Email: TechTran@jax.org

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