

C3(Cg)-*Dscam*^{3J}/GrsrJ

Stock No: 006046

 Spontaneous Mutation

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defects associated with some cases of Down syndrome.

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GENETIC OVERVIEW

Genetic Background

Generation

Dscam^{3J}

Alele Type

Spontaneous

Gene Symbol

Dscam

Gene Name

DS cell adhesion molecule

VIEW GENETICS

RESEARCH APPLICATIONS

Developmental Biology Research

Neurobiology Research

Cell Biology Research

VIEW ALL RESEARCH APPLICATIONS

BASE PRICE

Starting at:

Details

Detailed Description

Dscam^{3J}, a point mutation causing an R1018P amino acid substitution in the second fibronectin domain, causes a less severe phenotype than *Dscam*^{2J}, *Dscam*^{del17}, or *Dscam* targeted null mutants. The phenotype of *Dscam* null mutants varies with genetic background and includes respiratory distress, associated perinatal lethality, changes in C4 ventral root and pre-inspiratory neuron signaling, and an abnormal response to hypercapnia. *Dscam*^{3J} homozygotes display kyphosis, domed skull, muscle stiffness, and less difficulty in the righting response than is found in *Dscam*^{2J} homozygotes, which have a truncation in the extracellular domain of this protein. *Dscam*^{3J} homozygotes have enlarged central and lateral ventricles of the brain.

Characterization of the retina shows that, similar to *Dscam*^{2J} homozygotes, *Dscam*^{3J} homozygotes have expanded inner plexiform, inner nuclear and retinal ganglion cell layers, but, distinct from *Dscam*^{2J} homozygotes, the inner nuclear layer is evenly laminated. The retinal ganglion cells have defects in arborization and soma spacing, but the dopaminergic amacrine cells have reduced defects in arborization and soma spacing compared with those of the retinas of *Dscam*^{2J} homozygotes. Retinas of *Dscam*^{3J} homozygotes also have increased incidence of juxtaposed dopaminergic amacrine cells, occasional loose fasciculation of dopaminergic amacrine cell neurites, and increased numbers of bNOS amacrine cells, which also have abnormal spacing and appear hypertrophied, but the severity of these defects is less severe than the defects caused by other *Dscam* mutant alleles. The defects in neurite lamination, laminar specificity of type 2 and type 6 cone bipolar cells, and the disrupted targeting of retinal ganglion cell axons to the dorsal lateral geniculate nucleus that are found in *Dscam*^{2J} and other *Dscam* mutant homozygotes are not distinct phenotypes in *Dscam*^{3J} homozygotes. Subcellular localization of DSCAM protein from *Dscam*^{3J} homozygotes shows increased retention in the cell bodies in the retinal ganglion cell layer consistent with a model of mis-localization of this mutant protein. (See Schramm et al., 2013 for more detail.)

Development

Control Suggestions

Selected References

Genetics

Dscam^{3J}

⊖ 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

End Point Analysis: [Dscam<3J>](#)

[Genotyping resources and troubleshooting](#)

Mating System

Heterozygote x Heterozygote

Citation

When using the C3(Cg)-*Dscam*^{3J}/GrsrJ mouse strain in a publication, please [cite the originating article\(s\)](#) and include JAX stock #006046 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



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CRYORECOVERY - DOMESTIC PRICING

SERVICE/PRODUCT	DESCRIPTION	PRICE
Cryo Recovery	Heterozygous for Dscam<3J>	\$2,854.50

RELATED PRODUCTS AND SERVICES

Frozen Mouse Embryo	C3(CB17)-Dscam<3J>/GrsrJ	\$2595.00
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