

## STOCK Rb(11.13)4Bnr/J

Stock No: 000729

 Chromosome Aberration, Robertsonian

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reproduction/meiosis, birth defects, aneuploidy generation, other genomic studies or genetic mapping.

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## GENETIC OVERVIEW

### Genetic Background

### Generation

#### *Rd3<sup>rd3</sup>*

#### Alele Type

Spontaneous

#### Gene Symbol

*Rd3*

#### Gene Name

retinal degeneration 3

#### *Mc1r<sup>E-tob</sup>*

#### Alele Type

Spontaneous

#### Gene Symbol

*Mc1r*

#### Gene Name

melanocortin 1 receptor

### Marker(s)

#### Marker Symbol

Rb(11.13)4Bnr

#### Marker Name

Robertsonian translocation, Chr 11 and 13, Universitat Bonn/Rhein 4

Rb(11.13)4Bnr

Robertsonian translocation, Chr 11 and 13, Universitat Bonn/Rhein 4

VIEW GENETICS

# RESEARCH APPLICATIONS

Endocrine Deficiency Research  
Sensorineural Research

VIEW ALL RESEARCH APPLICATIONS

## Details

### Important Note

This strain is homozygous for retinal degeneration 3, *rd3*.

### Detailed Description

This inbred strain bears a Robertsonian translocation, Rb(11.13)4Bnr, and a spontaneous eye mutation, *Rd3<sup>rd3</sup>*, and was derived from the sibling inbreeding of an F1 cross between a female from the outbred NMRI stock and a wild trapped *Mus poschiavinus* male from Val Poschiavo, Switzerland. It may be useful for studies of retinal degeneration, cancer, reproduction/meiosis, birth defects, aneuploidy generation, other genomic studies or genetic mapping.

### Development

## Genetics

### *Rd3<sup>rd3</sup>*

### *Mc1r<sup>E-tob</sup>*

### Rb(11.13)4Bnr

### Rb(11.13)4Bnr

## Disease/Phenotype

### Disease Terms

### Research Areas By Phenotype

### Mammalian Phenotype Terms by Genotype

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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  
[Genotyping resources and troubleshooting](#)

Mating System  
Homozygote x Homozygote  
Appearance  
albino  
Related Genotype:  $Tyr^c/Tyr^c$   $Mc1r^{E-tob}/Mc1r^{E-tob}$

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