



B 6 1 2 9 S - T c (H S A 2 1) 1 T y b E m c f / J

Stock No: 010801 | Tc1



AVAILABLE FOR REGISTERING INTEREST

R E G I S T E R I N T E R E S T

Please contact **Technical Support** for more information

Overview

Also Known As: Tc1, Tc(HSA21)91-1Emcf

This strain is currently unavailable due to replenishing of cryopreserved stocks.

These Tc1 mice contain 42Mb (approximately 90%) of a freely segregating human fragment of Chromosome 21 *Hsa21* containing 269 genes, including most of the gene orthologs located on mouse Chromosome 10 (*Mmu10*), *Mmu16*, and *Mmu17*, which have been found to contribute to human Down Syndrome (DS). These mice may be useful for studying the genes involved in human chromosome aneuploidy and its role in DS.

Donating Investigator

Elizabeth MC Fisher, UCL Institute of Neurology

R E A D M O R E +

GENETIC OVERVIEW

Genetic Background

Generation

Tc(HSA21)1TybEmcf

Allele Type

Not Applicable (Inserted expressed sequence, Humanized sequence)

Gene Symbol

Tc(HSA21)1TybEmcf

Gene Name

transchromosomal, human 21, line 1, Victor Tybulewicz and Elizabeth MC Fisher

[VIEW GENETICS](#)

RESEARCH APPLICATIONS

Neurobiology Research

Developmental Biology Research

Mouse/Human Gene Homologs

[VIEW ALL RESEARCH APPLICATIONS](#)

Details

Detailed Description

Mice carrying a human fragment of Chromosome 21 (*Hsa21*) are viable, fertile, and normal in size, only the female carriers consistently transmits the mutation to the germline. When maintained on a background other than (C57BL/6 X 129S8/SvEv) germline transmission is completely abolished. These Tc1 mice contain 42Mb (approximately 83%) of a freely segregating *Hsa21* containing 269 genes, including most of the gene orthologs located on mouse Chromosome 10 (Mmu10), Mmu16, and Mmu17, which have been found to contribute to human Down Syndrome (DS). This mouse strain represents the most complete model of DS, exhibiting alterations in behavior, learning, memory, synaptic plasticity, cerebellar neuronal number, heart development, mandible size, defects in motor coordination, perturbed hematopoiesis, and reduced tumor angiogenesis. These mice may be useful for studying the genes involved in human chromosome aneuploidy and its role in DS.

Development

Control Suggestions

Selected References

Genetics

Tc(HSA21)1TybEmcf

Disease/Phenotype

Disease Terms

[+ Research Areas By Genotype](#)

[+ Mammalian Phenotype Terms by Genotype](#)

[+ References](#)

[- Technical Support](#)

C H A T O  F L I N E

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: [Tc\(Hsa21\)1TybEmcf](#)

[Genotyping resources and troubleshooting](#)

Breeding Considerations

When maintaining a live colony, Tc1 females are bred to B6129S8F1/J males (Stock No. [012868](#)) to maintain the live colony. Tc1 males do not consistently transmit mutation to germline, and transmission is completely lost on some congenic backgrounds.

[Additional Breeding and Husbandry Support](#)

Mating System

Heterozygote x F1

Citation

When using the Tc1 mouse strain in a publication, please [cite the originating article\(s\)](#) and include JAX stock #010801 in your Materials and Methods section.

STRAIN INTEREST REGISTRATION

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This information helps us manage the colony build and better meet the broad needs of the research community.

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Frequency

Product

Comment

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