

## STOCK Tg(tetO-CHRM3\*)1Blr/J

Stock No: **014093** | TRE-hM3Dq

 Transgenic

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

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(inducible/reversible) expression of hM3Dq; a mutant G protein-coupled receptor (GPCR) that activates the canonical G<sub>q</sub> pathway specifically following administration of the pharmacologically inert molecule clozapine-*N*-oxide (CNO). TRE-hM3Dq transgenic mice may be useful for chemogenetic/pharmacogenetic applications to study receptor-specific functions or general downstream cellular signaling emanating from the activated GPCR.

### Donating Investigator

Bryan L Roth, University of North Carolina at Chapel Hill

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## GENETIC OVERVIEW

Genetic Background

Generation

### Tg(tetO-CHRM3\*)1Blr

#### Alele Type

Transgenic (Inducible, Inserted expressed sequence)

VIEW GENETICS

## RESEARCH APPLICATIONS

Research Tools

Neurobiology Research

Diabetes and Obesity Research

Immunology, Inflammation and Autoimmunity Research

Cell Biology Research

Cancer Research

## BASE PRICE

Starting at:

\$2,854.50 Domestic price Cryo Recovery

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### Details

#### Detailed Description

Hemizygous TRE-hM3Dq transgenic mice are viable and fertile, with no reported phenotypic abnormalities. The TRE-hM3Dq transgene has a modified Tet response element (TRE or *tetO*) upstream of the mutant G protein-coupled receptor, hM3Dq (a human muscarinic 3 receptor with two amino acid substitutions (Y149C<sup>3,35</sup>/A239G<sup>3,46</sup>) that abolish receptor affinity for the native ligand, acetylcholine (ACh), but allow receptor binding and subsequent activation by the small pharmacologically inert molecule clozapine-*N*-oxide (CNO)). When bred with another mouse expressing tetracycline-controlled transactivator protein (tTA) or reverse tetracycline-controlled transactivator protein (rtTA), hM3Dq expression in the resulting double mutant offspring can be regulated with tetracycline or its analog doxycycline (dox). As designed, TRE-hM3Dq transgenic mice have no reported levels of hM3Dq activity in tTA(+dox) or rtTA(-dox) cell types. In TRE-hM3Dq transgenic tTA(-dox) or TRE-hM3Dq transgenic rtTA(+dox) cells types, hM3Dq expression results in activation of the canonical G<sub>q</sub> pathway only following administration of CNO.

These TRE-hM3Dq transgenic mice may be useful to study receptor-specific functions or general downstream cellular signaling emanating from the activated G protein-coupled receptor (GPCR). For example, when bred to a strain expressing tTA in forebrain neurons (Camk2a-tTA; see Stock No. [003010](#)), these transgenic mice are a model allowing *in vivo* chemical control of neuronal activity, neuronal firing, and non-neuronal signaling.

In addition, breeding TRE-hM3Dq mice with cfos-htTA mice (Stock No. [018306](#)) generates double transgenic offspring with hM3Dq expression in excitatory neurons that are sufficiently active to drive the c-fos promoter. Those hM<sub>3</sub>D<sub>q</sub><sup>fos</sup> double transgenic mice allow CNO ligand-induced, artificial reactivation of neurons associated with a specific memory. Temporal modulation may also be employed using dox. Such hM<sub>3</sub>D<sub>q</sub><sup>fos</sup> double transgenic mice allow one to generate and artificially-activate a synthetic memory trace, and may be useful in studying the spatial pattern of activity at the time of learning and retrieval.

*Of note, several chemogenetic/pharmacogenetic tool strains are available from The Jackson Laboratory Repository; including these Tet-responsive TRE-hM3Dq transgenic mice (Stock No. [014093](#)), the adora2A-rM3Ds transgenic mice (Stock No. [017863](#)), the Tet-responsive TRE-hM4Di transgenic mice (Stock No. [024114](#)), the Cre-inducible R26-LSL-Gi-DREADD mice (R26-hM4Di/mCitrine; Stock No. [026219](#)) and the Cre-inducible CAG-LSL-Gq-DREADD mice (Tg-hM3Dq/mCitrine [formerly R26-LSL-Gq-DREADD]; Stock No. [026220](#)).*

For CNO protocol, see the [detailed protocol for dissolving CNO](#) obtained from the attachments section of the [Designer Receptors Exclusively Activated by Designer Drugs DREADD wiki webpage](#).

[+ Development](#)

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[+ Expression Data](#)

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[+ Control Suggestions](#)

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[+ Selected References](#)

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## [- Genetics](#)

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[+ Tg\(tetO-CHRM3\\*\)1Blr](#)

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## [- Disease/Phenotype](#)

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[+ Disease Terms](#)

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[+ Research Areas By Phenotype](#)

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[+ Mammalian Phenotype Terms by Genotype](#)

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[+ References](#)

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## [- 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:[\\*\\* human CHRM3\\*Y149C](#)

Pyrosequencing:[\\*\\* human CHRM3\\*A239G](#)

Standard PCR:[Tg\(tetO-CHRM3\\*\)1Blr](#)

Separated PCR:[Tg\(tetO-CHRM3\\*\)1Blr](#)

Standard PCR:[Tg\(tetO-CHRM3\\*\)1Blr](#)

[Genotyping resources and troubleshooting](#)

## Breeding Considerations

When maintaining a live colony, transgene carrier mice may be bred together or to wildtype (noncarrier) mice from the colony. The donating investigator reports they have not tried to generate homozygous mice (November 2011).

### Additional Breeding and Husbandry Support

## Citation

When using the TRE-hM3Dq mouse strain in a publication, please [cite the originating article\(s\)](#) and include JAX stock #014093 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

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
<a href="#">Cryo Recovery</a>	Hemizygous or Non carrier for Tg(tetO-CHRM3*)1Blr	\$2,854.50

### RELATED PRODUCTS AND SERVICES

<a href="#">Frozen Mouse Embryo</a>	STOCK Tg(tetO-CHRM3*)1Blr/J Frozen Embryo	\$2595.00
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## PAYMENT TERMS AND CONDITIONS

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### TERMS OF USE

[General Terms and Conditions](#)

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### LICENSING INFORMATION

Phone: 207-288-6470

Email: [TechTran@jax.org](mailto:TechTran@jax.org)

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