

Resource Summary Report

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B6.129S4(C)-Mecp2^{tm1Jae}/Mmucd

RRID:MMRRC_011918-UCD

Type: Organism

Proper Citation

RRID:MMRRC_011918-UCD

Organism Information

URL: https://www.mmrrc.org/catalog/sds.php?mmrrc_id=11918

Proper Citation: RRID:MMRRC_011918-UCD

Description: Mus musculus with name B6.129S4(C)-Mecp2^{tm1Jae}/Mmucd from MMRRC.

Species: Mus musculus

Notes: Research areas: Models for Human Disease, Neurobiology; Mutation Type: other ; Collection:

Phenotype: tremors [MP:0000745]| decreased brain size [MP:0000774]| obese [MP:0001261]| decreased body weight [MP:0001262]| ataxia [MP:0001393]| hypoactivity [MP:0001402]| impaired coordination [MP:0001405]| abnormal pilomotor reflex [MP:0001492]| abnormal respiration [MP:0001943]| premature death [MP:0002083]| decreased brain weight [MP:0002175]| abnormal neuron morphology [MP:0002882]| increased susceptibility to age related obesity [MP:0003212]| prolonged QT interval [MP:0003233]| increased heart rate variability [MP:0003928]| abnormal heartbeat [MP:0004085]| increased response of heart to induced stress [MP:0004485]| abnormal behavior [MP:0004924]| cachexia [MP:0005150]| decreased heart rate [MP:0005333]| decreased body temperature [MP:0005534]| abnormal hippocampus CA2 region morphology [MP:0008265]| ventricular tachycardia [MP:0008950]| ventricular premature beat [MP:0009732]

Affected Gene: Mecp2

Catalog Number: 011918-UCD

Background: other

Database: Mutant Mouse Resource and Research Center (MMRRC)

Database Abbreviation: MMRRC

Source References: [PMID:11242118](#), [PMID:12432090](#), [PMID:14593183](#)

Alternate IDs: MMRRC_11918-UCD, MMRRC_011918, MMRRC_11918

Organism Name: B6.129S4(C)-*Mecp2*^{tm1Jae}/Mmucd

Record Creation Time: 20230308T054913+0000

Record Last Update: 20250419T223010+0000

Ratings and Alerts

No rating or validation information has been found for B6.129S4(C)-*Mecp2*^{tm1Jae}/Mmucd.

No alerts have been found for B6.129S4(C)-*Mecp2*^{tm1Jae}/Mmucd.

Data and Source Information

Source: [Integrated Animals](#)

Source Database: Mutant Mouse Resource and Research Center (MMRRC)

Usage and Citation Metrics

We found 10 mentions in open access literature.

Listed below are recent publications. The full list is available at [FDI Lab - SciCrunch.org](#).

Rupert DD, et al. (2023) Selective Deletion of Methyl CpG Binding Protein 2 from Parvalbumin Interneurons in the Auditory Cortex Delays the Onset of Maternal Retrieval in Mice. *The Journal of neuroscience : the official journal of the Society for Neuroscience*, 43(40), 6745.

Jiang Y, et al. (2021) Rett syndrome linked to defects in forming the MeCP2/Rbfox/LASR complex in mouse models. *Nature communications*, 12(1), 5767.

Trovato F, et al. (2020) Modelling genetic mosaicism of neurodevelopmental disorders in vivo by a Cre-amplifying fluorescent reporter. *Nature communications*, 11(1), 6194.

Lavery LA, et al. (2020) Losing Dnmt3a dependent methylation in inhibitory neurons impairs neural function by a mechanism impacting Rett syndrome. *eLife*, 9.

Du F, et al. (2016) Acute and crucial requirement for MeCP2 function upon transition from early to late adult stages of brain maturation. *Human molecular genetics*, 25(9), 1690.

Sztainberg Y, et al. (2015) Reversal of phenotypes in MECP2 duplication mice using genetic rescue or antisense oligonucleotides. *Nature*, 528(7580), 123.

Zhang Y, et al. (2014) Deep-brain magnetic stimulation promotes adult hippocampal neurogenesis and alleviates stress-related behaviors in mouse models for neuropsychiatric disorders. *Molecular brain*, 7, 11.

Zhang W, et al. (2014) Loss of MeCP2 from forebrain excitatory neurons leads to cortical hyperexcitation and seizures. *The Journal of neuroscience : the official journal of the Society for Neuroscience*, 34(7), 2754.

Nguyen MV, et al. (2013) Oligodendrocyte lineage cells contribute unique features to Rett syndrome neuropathology. *The Journal of neuroscience : the official journal of the Society for Neuroscience*, 33(48), 18764.

Nguyen MV, et al. (2012) MeCP2 is critical for maintaining mature neuronal networks and global brain anatomy during late stages of postnatal brain development and in the mature adult brain. *The Journal of neuroscience : the official journal of the Society for Neuroscience*, 32(29), 10021.