

# Resource Summary Report

Generated by [Kravitz](#) on Sep 17, 2026

[Apoe<sup>tm1\(APOE\\*2\)</sup>Mae/Apoe<sup>tm1\(APOE\\*2\)</sup>Mae](#)

RRID:MGI:3695702

Type: Organism

## Proper Citation

RRID:MGI:3695702

## Organism Information

**URL:**

**Proper Citation:** RRID:MGI:3695702

**Description:** Allele Detail: Targeted This is a legacy resource.

**Species:** Mus musculus

**Notes:** Allele Detail: Targeted This is a legacy resource.

**Phenotype:** liver/biliary system phenotype, increased circulating cholesterol level, increased susceptibility to atherosclerosis, abnormal lipid level, increased circulating triglyceride level, abnormal lipid homeostasis, abnormal retinal pigment epithelium morphology, increased circulating cholesterol level, retinal degeneration, hyperlipidemia, abnormal Bruch membrane morphology, atherosclerotic lesions, abnormal lipid level, abnormal lipid homeostasis, xanthoma, increased circulating triglyceride level

**Affected Gene:** Apoe

**Genomic Alteration:** tm1(APOE\*2)Mae

**Catalog Number:** 3695702

**Background:** involves: 129P2/OlaHsd \* C57BL/6

**Database:** MGI, Mouse Genome Informatics MGI

**Database Abbreviation:** MGI

**Availability:** Availability unknown check source stock center

**Source References:** [PMID:11076954](#), [PMID:9649566](#), [PMID:16079201](#)

**Organism Name:** Apoe<sup>tm1(APOE\*2)</sup>Mae/Apoe<sup>tm1(APOE\*2)</sup>Mae

**Record Creation Time:** 20240120T190535+0000

**Record Last Update:** 20240130T201950+0000

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## Ratings and Alerts

No rating or validation information has been found for Apoe<sup>tm1(APOE\*2)Mae</sup>/Apoe<sup>tm1(APOE\*2)Mae</sup>.

No alerts have been found for Apoe<sup>tm1(APOE\*2)Mae</sup>/Apoe<sup>tm1(APOE\*2)Mae</sup>.

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## Data and Source Information

**Source:** [Integrated Animals](#)

**Source Database:** MGI, Mouse Genome Informatics MGI

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## Usage and Citation Metrics

We found 3 mentions in open access literature.

**Listed below are recent publications.** The full list is available at [Kravitz](#).

Labuza A, et al. (2025) Aging, Rather than Genotype, Is the Principal Contributor to Differential Gene Expression Within Targeted Replacement APOE2, APOE3, and APOE4 Mouse Brain. Brain sciences, 15(10).

Peng KY, et al. (2024) Apolipoprotein E2 Expression Alters Endosomal Pathways in a Mouse Model With Increased Brain Exosome Levels During Aging. Traffic (Copenhagen, Denmark), 25(5), e12937.

Shinohara M, et al. (2020) APOE2 is associated with longevity independent of Alzheimer's disease. eLife, 9.