

# Resource Summary Report

Generated by [FDI Lab - SciCrunch.org](#) on May 17, 2025

## C57BL/6J-Lyst<sup>bg-Btlr</sup>/Mmucd

RRID:MMRRC\_010470-UCD

Type: Organism

### Proper Citation

RRID:MMRRC\_010470-UCD

### Organism Information

**URL:** [https://www.mmrrc.org/catalog/sds.php?mmrrc\\_id=10470](https://www.mmrrc.org/catalog/sds.php?mmrrc_id=10470)

**Proper Citation:** RRID:MMRRC\_010470-UCD

**Description:** Mus musculus with name C57BL/6J-Lyst<sup>bg-Btlr</sup>/Mmucd from MMRRC.

**Species:** Mus musculus

**Notes:** Research areas: Cell Biology, Immunology and Inflammation, Models for Human Disease, Virology; Mutation Type: chemically induced mutation ; Collection: Beutler Mutagenetix

**Phenotype:** diluted coat color [MP:0000371]| increased susceptibility to bacterial infection [MP:0002412]| increased susceptibility to viral infection [MP:0002418]| abnormal foot pigmentation [MP:0009379]| abnormal NK cell physiology [MP:0010766]| decreased tail pigmentation [MP:0011277]| decreased ear pigmentation [MP:0011279]

**Affected Gene:** Lyst

**Catalog Number:** 010470-UCD

**Background:** chemically induced mutation

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

**Database Abbreviation:** MMRRC

**Alternate IDs:** MMRRC\_10470-UCD, MMRRC\_010470, MMRRC\_147

**Organism Name:** C57BL/6J-Lyst<sup>bg-Btlr</sup>/Mmucd

**Record Creation Time:** 20230308T054905+0000

**Record Last Update:** 20250510T103400+0000

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## Ratings and Alerts

No rating or validation information has been found for C57BL/6J-Lyst<sup>bg-Btlr</sup>/Mmucd.

No alerts have been found for C57BL/6J-Lyst<sup>bg-Btlr</sup>/Mmucd.

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## Data and Source Information

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

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

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## Usage and Citation Metrics

We found 1 mentions in open access literature.

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

Kögl T, et al. (2024) Patients and mice with deficiency in the SNARE protein SYNTAXIN-11 have a secondary B cell defect. The Journal of experimental medicine, 221(7).