# **Resource Summary Report**

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# Mouse HapMap Imputation Genotype Resource

RRID:SCR 002576

Type: Tool

## **Proper Citation**

Mouse HapMap Imputation Genotype Resource (RRID:SCR\_002576)

#### **Resource Information**

URL: http://mouse.cs.ucla.edu/mousehapmap/

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**Description:** Genetic maps for 94 inbred strains of mouse and imputed genotypes using the NIEHS / Perlegen resequencing resource. Combining with the 13,094 Wellcome Trust SNPs (Single-nucleotide polymorphisms), a set of 132,285 SNPs was compiled and is available for download. Using the mouse HapMap resource, it is possible to accurately impute the genotypes of the 94 strains at the 8 million SNPs discovered by the NIEHS/Perlegen mouse resequencing project. They imputed the genotypes at the NIEHS/Perlegen SNPs from the mouse HapMap SNPs and an additional set of 7,570 gap-filling SNPs provided by NIEHS/Perlegen. Since each NIEHS/Perlegen SNP probe has different quality, they classified roughly half of the SNPs as "high-quality" SNPs, which do not have missing genotype at any of the 15 resequenced strains. The imputed genotypes are available for the high-quality SNPs, which has estimated error rate of 0.27% for high-confidence imputed genotypes. In addition, the imputed genotypes for all 8 million SNPs are also available for download. Their estimated error rate is 0.37% for high-confidence imputed genotypes.

Abbreviations: Mouse HapMap

Resource Type: data or information resource, data set

**Keywords:** genetic variation, genotype, haplotype, imputed genotype, inbred mouse strain, imputed, phenotype, single-nucleotide polymorphism

Funding: NIEHS

Resource Name: Mouse HapMap Imputation Genotype Resource

Resource ID: SCR\_002576

**Alternate IDs:** nif-0000-21752

**Record Creation Time:** 20220129T080214+0000

**Record Last Update:** 20250407T215329+0000

### Ratings and Alerts

No rating or validation information has been found for Mouse HapMap Imputation Genotype Resource.

No alerts have been found for Mouse HapMap Imputation Genotype Resource.

#### Data and Source Information

Source: SciCrunch Registry

### **Usage and Citation Metrics**

We found 5 mentions in open access literature.

Listed below are recent publications. The full list is available at FDI Lab - SciCrunch.org.

Rudy RF, et al. (2019) A Genome-Wide Analysis of the Penumbral Volume in Inbred Mice following Middle Cerebral Artery Occlusion. Scientific reports, 9(1), 5070.

Zhou Y, et al. (2019) Integrative system genetic analysis reveals mRNA-lncRNA network associated with mouse spontaneous lung cancer susceptibility. Oncotarget, 10(3), 339.

Rau CD, et al. (2015) High-Density Genotypes of Inbred Mouse Strains: Improved Power and Precision of Association Mapping. G3 (Bethesda, Md.), 5(10), 2021.

Adams A, et al. (2013) Presphenoidal synchondrosis fusion in DBA/2J mice. Mammalian genome: official journal of the International Mammalian Genome Society, 24(1-2), 54.

Park CC, et al. (2011) Gene networks associated with conditional fear in mice identified using a systems genetics approach. BMC systems biology, 5, 43.