

Resource Summary Report

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B6.FVB-Tg(Myh6-cre)2182Mds/J

RRID:IMSR_JAX:011038

Type: Organism

Proper Citation

RRID:IMSR_JAX:011038

Organism Information

URL: <https://www.jax.org/strain/011038>

Proper Citation: RRID:IMSR_JAX:011038

Description: Mus musculus with name B6.FVB-Tg(Myh6-cre)2182Mds/J from IMSR.

Species: Mus musculus

Notes: gene symbol note: myosin; heavy polypeptide 6; cardiac muscle; alpha||transgene insertion 2182; Michael D Schneider|myosin; heavy polypeptide 6; cardiac muscle; alpha||transgene insertion 2182; Michael D Schneider; mutant strain: Myh6||Tg(Myh6-cre)2182Mds|Myh6||Tg(Myh6-cre)2182Mds

Affected Gene: myosin; heavy polypeptide 6; cardiac muscle; alpha||transgene insertion 2182; Michael D Schneider|myosin; heavy polypeptide 6; cardiac muscle; alpha||transgene insertion 2182; Michael D Schneider

Genomic Alteration: transgene insertion 2182; Michael D Schneider

Catalog Number: JAX:011038

Database: International Mouse Resource Center IMSR, JAX

Database Abbreviation: IMSR

Availability: live

Alternate IDs: IMSR_JAX:11038

Organism Name: B6.FVB-Tg(Myh6-cre)2182Mds/J

Record Creation Time: 20230509T193303+0000

Record Last Update: 20250412T090536+0000

Ratings and Alerts

No rating or validation information has been found for B6.FVB-Tg(Myh6-cre)2182Mds/J.

Warning: Warning. Researchers have noted that this genotype does not sufficiently model human hypoplastic left heart syndrome.

Data and Source Information

Source: [Integrated Animals](#)

Source Database: International Mouse Resource Center IMSR, JAX

Usage and Citation Metrics

We found 77 mentions in open access literature.

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

Ge W, et al. (2025) Rnd3 protects against doxorubicin-induced cardiotoxicity through inhibition of PANoptosis in a Rock1/Drp1/mitochondrial fission-dependent manner. *Cell death & disease*, 16(1), 2.

Shu J, et al. (2024) PARP1 Promotes Heart Regeneration and Cardiomyocyte Proliferation. *International journal of biological sciences*, 20(5), 1602.

Niu X, et al. (2024) lncRNA Oip5-as1 inhibits excessive mitochondrial fission in myocardial ischemia/reperfusion injury by modulating DRP1 phosphorylation. *Cellular & molecular biology letters*, 29(1), 72.

Akins KA, et al. (2024) Runx1 is sufficient but not required for cardiomyocyte cell-cycle activation. *American journal of physiology. Heart and circulatory physiology*, 327(2), H377.

Gural B, et al. (2024) Novel Insights into Post-Myocardial Infarction Cardiac Remodeling through Algorithmic Detection of Cell-Type Composition Shifts. *bioRxiv : the preprint server for biology*.

Wei X, et al. (2024) Tert promotes cardiac regenerative repair after MI through alleviating ROS-induced DNA damage response in cardiomyocyte. *Cell death discovery*, 10(1), 381.

Qu Z, et al. (2024) The positive feedback loop of the NAT10/Mybbp1a/p53 axis promotes cardiomyocyte ferroptosis to exacerbate cardiac I/R injury. *Redox biology*, 72, 103145.

Tang Y, et al. (2024) Nucleolin myocardial-specific knockout exacerbates glucose metabolism disorder in endotoxemia-induced myocardial injury. *PeerJ*, 12, e17414.

Davenport A, et al. (2024) Comparative analysis of two independent Myh6-Cre transgenic mouse lines. *Journal of molecular and cellular cardiology plus*, 9.

Bi X, et al. (2024) Characterization of ferroptosis-triggered pyroptotic signaling in heart failure. *Signal transduction and targeted therapy*, 9(1), 257.

Sun SJ, et al. (2024) Gasdermin-E-mediated pyroptosis drives immune checkpoint inhibitor-associated myocarditis via cGAS-STING activation. *Nature communications*, 15(1), 6640.

Li W, et al. (2024) Cardiac corin and atrial natriuretic peptide regulate liver glycogen metabolism and glucose homeostasis. *Cardiovascular diabetology*, 23(1), 383.

Yang H, et al. (2024) Dysregulated RBM24 phosphorylation impairs APOE translation underlying psychological stress-induced cardiovascular disease. *Nature communications*, 15(1), 10181.

Duan Q, et al. (2024) Deptor protects against myocardial ischemia-reperfusion injury by regulating the mTOR signaling and autophagy. *Cell death discovery*, 10(1), 508.

Grimes KM, et al. (2024) MEK1-ERK1/2 signaling regulates the cardiomyocyte non-sarcomeric actin cytoskeletal network. *American journal of physiology. Heart and circulatory physiology*, 326(1), H180.

Sweeney M, et al. (2024) Interleukin 11 therapy causes acute left ventricular dysfunction. *Cardiovascular research*, 120(17), 2220.

Sun JT, et al. (2024) PEX3 promotes regenerative repair after myocardial injury in mice through facilitating plasma membrane localization of ITGB3. *Communications biology*, 7(1), 795.

Han J, et al. (2024) Cardiomyocyte-derived USP28 negatively regulates antioxidant response and promotes cardiac hypertrophy via deubiquitinating TRIM21. *Theranostics*, 14(16), 6236.

Su C, et al. (2024) Tudor-SN promotes cardiomyocyte proliferation and neonatal heart regeneration through regulating the phosphorylation of YAP. *Cell communication and signaling : CCS*, 22(1), 345.

Liu L, et al. (2024) 14-3-3 binding motif phosphorylation disrupts Hdac4-organized condensates to stimulate cardiac reprogramming. *Cell reports*, 43(4), 114054.