

Resource Summary Report

Generated by [NIF](#) on Jul 30, 2026

pcDNA3.1-MEF2C-HA

RRID:Addgene_32515

Type: Plasmid

Proper Citation

RRID:Addgene_32515

Plasmid Information

URL: <http://www.addgene.org/32515>

Proper Citation: RRID:Addgene_32515

Insert Name: Mef2c

Organism: Mus musculus

Bacterial Resistance: Ampicillin

Defining Citation: [PMID:19704004](#)

Vector Backbone Description: Backbone Marker:Invitrogen; Backbone Size:5400; Vector Backbone:pcDNA3.1; Vector Types:Mammalian Expression; Bacterial Resistance:Ampicillin

Plasmid Name: pcDNA3.1-MEF2C-HA

Record Creation Time: 20220422T222141+0000

Record Last Update: 20260725T074551+0000

Ratings and Alerts

No rating or validation information has been found for pcDNA3.1-MEF2C-HA.

No alerts have been found for pcDNA3.1-MEF2C-HA.

Data and Source Information

Source: [Addgene](#)

Usage and Citation Metrics

We found 7 mentions in open access literature.

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

Górska AA, et al. (2021) Muscle-specific Cand2 is translationally upregulated by mTORC1 and promotes adverse cardiac remodeling. *EMBO reports*, 22(12), e52170.

Nath SR, et al. (2020) MEF2 impairment underlies skeletal muscle atrophy in polyglutamine disease. *Acta neuropathologica*, 140(1), 63.

Mukai J, et al. (2019) Recapitulation and Reversal of Schizophrenia-Related Phenotypes in Setd1a-Deficient Mice. *Neuron*, 104(3), 471.

Wang C, et al. (2019) Methyltransferase-like 21c methylates and stabilizes the heat shock protein Hspa8 in type I myofibers in mice. *The Journal of biological chemistry*, 294(37), 13718.

Pröschel C, et al. (2017) Epilepsy-causing sequence variations in SIK1 disrupt synaptic activity response gene expression and affect neuronal morphology. *European journal of human genetics : EJHG*, 25(2), 216.

Xia JB, et al. (2016) Hypoxia/ischemia promotes CXCL10 expression in cardiac microvascular endothelial cells by NFkB activation. *Cytokine*, 81, 63.

Sakai Y, et al. (2013) Neuroendocrine phenotypes in a boy with 5q14 deletion syndrome implicate the regulatory roles of myocyte-specific enhancer factor 2C in the postnatal hypothalamus. *European journal of medical genetics*, 56(9), 475.