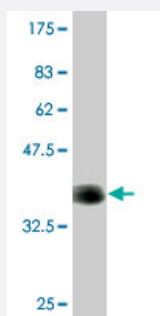


# CACNA1F monoclonal antibody (M05), clone 1H6

Catalog # H00000778-M05

Size 100 ug

## Applications



Western Blot detection against Immunogen (36.74 KDa) .

## Specification

<b>Product Description</b>	Mouse monoclonal antibody raised against a partial recombinant CACNA1F.
<b>Immunogen</b>	CACNA1F (NP_005174, 1878 a.a. ~ 1977 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.
<b>Sequence</b>	LHVPGTHSDPSHGKRGSAADSLVEAVLISEGLGLFARDPRFVALAKQEIADACRLTDEMDNAAS DLLAQGTSSLYSDEESILSRFDEEDLGDEMACVHAL
<b>Host</b>	Mouse
<b>Reactivity</b>	Human
<b>Interspecies Antigen Sequence</b>	Mouse (90)
<b>Isotype</b>	IgG3 Kappa
<b>Quality Control Testing</b>	Antibody Reactive Against Recombinant Protein. Western Blot detection against Immunogen (36.74 KDa) .
<b>Storage Buffer</b>	In 1x PBS, pH 7.4
<b>Storage Instruction</b>	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

## Applications

- Western Blot (Recombinant protein)

[Protocol Download](#)

- ELISA

## Gene Info — CACNA1F

Entrez GeneID	<a href="#">778</a>
GeneBank Accession#	<a href="#">NM_005183</a>
Protein Accession#	<a href="#">NP_005174</a>
Gene Name	CACNA1F
Gene Alias	AIED, COD3, CORDX, CORDX3, CSNB2, CSNB2A, CSNBX2, Cav1.4, JM8, JMC8, OA2
Gene Description	calcium channel, voltage-dependent, L type, alpha 1F subunit
Omim ID	<a href="#">300071</a> <a href="#">300110</a> <a href="#">300476</a>
Gene Ontology	<a href="#">Hyperlink</a>
Gene Summary	<p>This gene encodes a member of the alpha-1 subunit family; a protein in the voltage-dependent calcium channel complex. Calcium channels mediate the influx of calcium ions into the cell upon membrane polarization and consist of a complex of alpha-1, alpha-2/delta, beta, and gamma subunits in a 1:1:1:1 ratio. The alpha-1 subunit has 24 transmembrane segments and forms the pore through which ions pass into the cell. There are multiple isoforms of each of the proteins in the complex, either encoded by different genes or the result of alternative splicing of transcripts. Alternate transcriptional splice variants of the gene described here have been observed but have not been thoroughly characterized. Mutations in this gene have been shown to cause incomplete X-linked congenital stationary night blindness type 2 (CSNB2). [provided by RefSeq]</p>
Other Designations	Cav1.4alpha1

## Pathway

- [Arrhythmogenic right ventricular cardiomyopathy \(ARVC\)](#)
- [Calcium signaling pathway](#)

- [Cardiac muscle contraction](#)
- [GnRH signaling pathway](#)
- [Hypertrophic cardiomyopathy \(HCM\)](#)
- [MAPK signaling pathway](#)
- [Vascular smooth muscle contraction](#)

## Disease

- [Retinal Diseases](#)
- [Rhinitis](#)