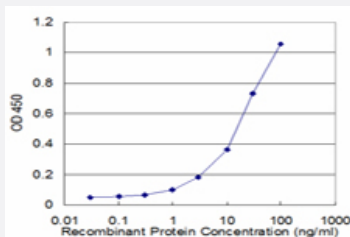


# GJB1 monoclonal antibody (M08), clone 1F5

Catalog # H00002705-M08

Size 100 ug

## Applications



### Sandwich ELISA (Recombinant protein)

Detection limit for recombinant GST tagged GJB1 is approximately 1ng/ml as a capture antibody.

## Specification

<b>Product Description</b>	Mouse monoclonal antibody raised against a partial recombinant GJB1.
<b>Immunogen</b>	GJB1 (AAH22426, 75 a.a. ~ 174 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.
<b>Sequence</b>	RLWSLQLILVSTPALLVAMHVAHQHIEKKMLRLEGHGDPLHLEEVKRHKVHISGTLWWTYISVV FRLLFEAVFMVVFYLLYPGYAMVRLVKCDVYPCP
<b>Host</b>	Mouse
<b>Reactivity</b>	Human
<b>Interspecies Antigen Sequence</b>	Mouse (97); Rat (97)
<b>Isotype</b>	IgG1 Kappa
<b>Quality Control Testing</b>	Antibody Reactive Against Recombinant Protein.
<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

- Sandwich ELISA (Recombinant protein)

Detection limit for recombinant GST tagged GJB1 is approximately 1ng/ml as a capture antibody.

[Protocol Download](#)

- ELISA

## Gene Info — GJB1

Entrez GeneID [2705](#)

GeneBank Accession# [BC022426](#)

Protein Accession# [AAH22426](#)

Gene Name GJB1

Gene Alias CMTX, CMTX1, CX32

Gene Description gap junction protein, beta 1, 32kDa

Omim ID [302800 304040](#)

Gene Ontology [Hyperlink](#)

**Gene Summary** This gene encodes a member of the gap junction protein family. The gap junction proteins are membrane-spanning proteins that assemble to form gap junction channels that facilitate the transfer of ions and small molecules between cells. According to sequence similarities at the nucleotide and amino acid levels, the gap junction proteins are divided into two categories, alpha and beta. Mutations in this gene cause X-linked Charcot-Marie-Tooth disease, an inherited peripheral neuropathy. Alternatively spliced transcript variants encoding the same protein have been found for this gene. [provided by RefSeq]

**Other Designations** OTTHUMP00000023502|OTTHUMP00000023503|OTTHUMP00000023504|connexin 32

## Disease

- [Charcot-Marie-Tooth Disease](#)
- [Deafness](#)
- [Genetic Diseases](#)

- [Genetic Predisposition to Disease](#)
- [Hearing Loss](#)
- [Ovarian Neoplasms](#)