

## FGF13 rabbit monoclonal antibody

Catalog # H00002258-K Size 100 ug x up to 3

Specification	
Product Description	Rabbit monoclonal antibody raised against a human FGF13 peptide using ARM Technology.
Immunogen	A synthetic peptide of human FGF13 is used for rabbit immunization.  Customer or Abnova will decide on the preferred peptide sequence.
Host	Rabbit
Library Construction	Non-fusion antibody library from rabbit spleen ( <u>ARM Technology</u> ).
Expression	Overexpression vector and transfection into 293H cell line.
Reactivity	Human
Purification	Protein A
Isotype	lgG
Quality Control Testing	Antibody reactive against human FGF13 peptide by ELISA and mammalian transfected lysate by W estern Blot.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.
Deliverable	Up to three rabbit lgG clones of 100 ug each will be delivered to customer.
Note	<ol> <li>Customer may provide cell or tissue lysate for antibody screening.</li> <li>Rabbit monoclonal antibody generated by ARM technology is amenable to antibody engineering in cluding F(ab)<sub>2</sub>, lgG, scFv and different Fc and non-Fc conjugates per customer request.</li> </ol>

## **Applications**

Western Blot (Transfected lysate)

Protocol Download



ELISA

Gene Info — FGF13	
Entrez GenelD	2258
GeneBank Accession#	FGF13
Gene Name	FGF13
Gene Alias	FGF2, FHF2
Gene Description	fibroblast growth factor 13
Omim ID	300070
Gene Ontology	<u>Hyperlink</u>
Gene Summary	The protein encoded by this gene is a member of the fibroblast growth factor (FGF) family. FGF f amily members possess broad mitogenic and cell survival activities, and are involved in a variety of biological processes, including embryonic development, cell growth, morphogenesis, tissue re pair, tumor growth, and invasion. This gene is located in a region on chromosome X, which is ass ociated with Borjeson-Forssman-Lehmann syndrome (BFLS), making it a possible candidate gen e for familial cases of the BFLS, and for other syndromal and nonspecific forms of X-linked menta I retardation mapping to this region. Alternative splicing of this gene at the 5' end results in several transcript variants encoding different isoforms with different N-termini. [provided by RefSeq
Other Designations	OTTHUMP00000024143 OTTHUMP00000024144 fibroblast growth factor homologous factor 2

## Pathway

- MAPK signaling pathway
- Melanoma
- Pathways in cancer
- Regulation of actin cytoskeleton

## Disease

- Alzheimer disease
- Cardiovascular Diseases



- Diabetes Complications
- Metabolic Syndrome X
- Neoplasms
- Osteoporosis