

FGF10 polyclonal antibody

Catalog # PAB16154 Size 100 ug

Specification	
Product Description	Rabbit polyclonal antibody raised against FGF10.
Immunogen	Human FGF10.
Host	Rabbit
Reactivity	Human, Mouse, Rat
Form	Lyophilized
Isotype	lgG
Recommend Usage	ELISA (1-2 ug/mL) Western Blot (2-10 ug/mL) Immunohistochemistry (2-10 ug/mL) The optimal working dilution should be determined by the end user.
Storage Buffer	Lyophilized from PBS.
Storage Instruction	Store at -20°C on dry atmosphere. After reconstitution with sterile 10mM PBS pH 7.4, store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot
- Immunohistochemistry
- Enzyme-linked Immunoabsorbent Assay

Gene Info — FGF10



Entrez GenelD	<u>2255</u>
Protein Accession#	<u>O15520</u>
Gene Name	FGF10
Gene Alias	-
Gene Description	fibroblast growth factor 10
Omim ID	<u>149730</u> <u>180920</u> <u>602115</u>
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 protein exhibits mitogenic activity for keratinizing epidermal cells, but essentially no activity for fibroblasts, which is similar to the biological activity of FGF7. St udies of the mouse homolog of suggested that this gene is required for embryonic epidermal mor phogenesis including brain development, lung morphogenesis, and initiation of lim bud formation. This gene is also implicated to be a primary factor in the process of wound healing. [provided by RefSeq
Other Designations	keratinocyte growth factor 2 produced by fibroblasts of urinary bladder lamina propria

Pathway

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

Disease

- Abnormalities
- Attention Deficit Disorder with Hyperactivity
- Cleft Lip



- Cleft Palate
- Genetic Predisposition to Disease
- <u>Hyperparathyroidism</u>
- Hypospadias
- Tourette Syndrome