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Datasheet for ABIN7317752

FGFR2 Protein (His tag,Fc Tag)



Overview

Quantity:	100 μg
Target:	FGFR2
Origin:	Human
Source:	HEK-293 Cells
Protein Type:	Recombinant
Biological Activity:	Active
Purification tag / Conjugate:	This FGFR2 protein is labelled with His tag,Fc Tag.

Product Details

Purpose:	Recombinant Human FGFR2/CD332 Protein (His & Fc Tag)(Active)
Sequence:	Met 1-Glu 377
Characteristics:	A DNA sequence encoding the human FGFR2 (NP_000132.3) extracellular domain (Met 1-Glu 377) was fused with the C-terminal polyhistidine-tagged Fc region of human IgG1 at the C-terminus.
Purity:	> 90 % as determined by reducing SDS-PAGE.
Endotoxin Level:	< 1.0 EU per µg as determined by the LAL method.
Biological Activity Comment:	Measured by its ability to inhibit FGF-acidic (aFGF/FGF1) dependent proliferation of Balb/C 3T3 mouse fibroblasts. The ED50 for this effect is typically 0.5-2.5 ng/ml.

Target Details

Target:	FGFR2	

Target Details

Alternative Name:	FGFR2/CD332 (FGFR2 Products)	
Background:	Background: FGFR2, also known as CD332, belongs to the fibroblast growth factor receptor	
	subfamily where amino acid sequence is highly conserved between members and throughout	
	evolution. FGFR2 acts as cell-surface receptor for fibroblast growth factors and plays an	
	essential role in the regulation of cell proliferation, differentiation, migration and apoptosis, and	
	in the regulation of embryonic development. It is required for normal embryonic patterning,	
	trophoblast function, limb bud development, lung morphogenesis, osteogenesis and skin	
	development. FGFR2 plays an essential role in the regulation of osteoblast differentiation,	
	proliferation and apoptosis, and is required for normal skeleton development. It also promotes	
	cell proliferation in keratinocytes and imature osteoblasts, but promotes apoptosis in	
	differentiated osteoblasts. FGFR2 signaling is down-regulated by ubiquitination, internalization	
	and degradation. Mutations that lead to constitutive kinase activation or impair normal CD332	
	maturation, internalization and degradation lead to aberrant signaling. Over-expressed FGFR2	
	promotes activation of STAT1. Defects in CD3322 are the cause of Crouzon syndrome,	
	Jackson-Weiss syndrome, Apert syndrome, Pfeiffer syndrome, Beare-Stevenson cutis gyrata	
	syndrome, familial scaphocephaly syndrome, lacrimo-auriculo-dento-digital syndrome and	
	Antley-Bixler syndrome without genital anomalies or disordered steroidogenesis.Immune	
	Checkpoint Immunotherapy Cancer Immunotherapy Targeted Therapy	
	Synonym: BBDS;BEK;BFR-1;CD332;CEK3;CFD1;ECT1;JWS;K-SAM;KGFR;TK14;TK25	
Molecular Weight:	67.6 kDa	
NCBI Accession:	NP_000132	
Pathways:	RTK Signaling, Fc-epsilon Receptor Signaling Pathway, EGFR Signaling Pathway, Neurotrophin	
	Signaling Pathway, Regulation of Muscle Cell Differentiation, Skeletal Muscle Fiber	
	Development, Growth Factor Binding	
Application Details		
Restrictions:	For Research Use only	
Handling		
Format:	Lyophilized	
Reconstitution:		
	Please refer to the printed manual for detailed information.	
Buffer:	Lyophilized from sterile PBS, pH 7.4	

Handling

Storage:	4 °C,-20 °C,-80 °C
Storage Comment:	Generally, lyophilized proteins are stable for up to 12 months when stored at -20 to -80°C. Reconstituted protein solution can be stored at 4-8°C for 2-7 days. Aliquots of reconstituted
	samples are stable at < -20°C for 3 months.