

OriGene Technologies, Inc.

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Product datasheet for TP723097

FGF 23 (FGF23) (NM_020638) Human Recombinant Protein

Product data:

Product Type:	Recombinant Proteins	
Description:	Purified recombinant protein of Human fibroblast growth factor 23 (FGF23).	
Species:	Human	
Expression Host:	E. coli	
Expression cDNA Clone or AA Sequence:	MYPNASPLLG SSWGGLIHLY TATARNSYHL QIHKNGHVDG APHQTIYSAL MIRSEDAGFV VITGVMSRRY LCMDFRGNIF GSHYFDPENC RFQHQTLENG YDVYHSPQYH FLVSLGRAKR AFLPGMNPPP YSQFLSRRNE IPLIHFNTPI PRRHTRSAED DSERDPLNVL KPRARMTPAP ASCSQELPSA EDNSPMASDP LGVVRGGRVN THAGGTGPEG CRPFAKFI	
Tag:	Tag Free	
Predicted MW:	22.5 kDa	
Concentration:	lot specific	
Purity:	>95% as determined by SDS-PAGE and Coomassie blue staining	
Buffer:	Lyophilized from a 0.2 μ M filtered solution of 20mM phosphate buffer,100mM NaCl, pH 7.2	
Endotoxin:	Endotoxin level is < 0.1 ng/μg of protein (< 1 EU/μg)	
Storage:	Store at -80°C.	
Stability:	Stable for at least 6 months from date of receipt under proper storage and handling conditions.	
RefSeq:	<u>NP 065689</u>	
Locus ID:	8074	
UniProt ID:	<u>Q9GZV9</u>	
RefSeq Size:	3018	
Cytogenetics:	12p13.32	
RefSeq ORF:	753	
Synonyms:	ADHR; FGFN; HFTC2; HPDR2; HYPF; PHPTC	



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Summary:	This gene encodes a member of the fibroblast growth factor family of proteins, which possess broad mitogenic and cell survival activities and are involved in a variety of biological processes. The product of this gene regulates phosphate homeostasis and transport in the kidney. The full-length, functional protein may be deactivated via cleavage into N-terminal and C-terminal chains. Mutation of this cleavage site causes autosomal dominant hypophosphatemic rickets (ADHR). Mutations in this gene are also associated with hyperphosphatemic familial tumoral calcinosis (HFTC). [provided by RefSeq, Feb 2013]
Protein Families Protein Pathway	 Druggable Genome, Secreted Protein MAPK signaling pathway, Melanoma, Pathways in cancer, Regulation of actin cytoskeleton

Product images:

200-	
116—	
97	
55-	
37—	
22-	
14 <u>-</u> 6-	

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