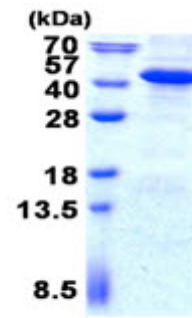


AR50713PU-S**Human Inositol monophosphatase 3 / IMPA3 (34-359, His-tag) - Purified**

Alternate names:	IMPAD1, IMPase 3, Inositol monophosphatase domain-containing protein 1, Inositol-1(or 4)-monophosphatase 3, Myo-inositol monophosphatase A3
Quantity:	0.1 mg
Concentration:	1 mg/ml (determined by Bradford assay)
Background:	Inositol monophosphatase 3, also known as IMPAD1, is a member of the inositol monophosphatase family. IMPAD1 is localized to the Golgi apparatus and catalyzes the hydrolysis of phosphoadenosine phosphate (PAP) to adenosine monophosphate (AMP). Mutations in this gene are a cause of GRAPP type chondrodysplasia with joint dislocations, and a pseudogene of this gene is located on the long arm of chromosome 1.
Uniprot ID:	Q9NX62
NCBI:	NP_060283
GeneID:	54928
Species:	Human
Source:	E. coli
Format:	State: Liquid purified protein Purity: >90% by SDS - PAGE Buffer System: 20 mM Tris-HCl buffer (pH 8.0) containing 2M Urea, 20% glycerol
Description:	Recombinant human IMPAD1 protein, fused to His-tag at N-terminus, was expressed in E.coli. AA Sequence: MGSSHHHHHH SSGLVPRGSH MSGRFSFLFG LGGEPGGGAA GPAAAADGGT VDLREMLAVS VLA AVRGGDE VRRVRESNVL HEKSKGKTRE GAEDKMTSGD VLSNRKMFYL LKTAFPSVQI NTEEHVDAAD QEVILWDHKI PEDILKEVTT PKEVPAESVT VWIDPLDATQ EYTEDLRKYV TTMVCVAVNG KPMLGVIHKP FSEYTAWAMV DGGSNVKARS SYNEKTPRIV VSRSHSGMVK QVALQTFGNQ TTIIPAGGAG YKVLALLDVP DKSQEKADLY IHVTYIKKWD ICAGNAILKA LGGHMTTLSG EEISYTGSDG IEGLLASIR MNHQALVRKL PDLEKTGHK Molecular weight: 37.6 kDa (349aa)
Storage:	Store undiluted at 2-8°C for one week or (in aliquots) at -20°C to -80°C for longer. Avoid repeated freezing and thawing. Shelf life: one year from despatch.
General Readings:	Vissers L E., et al. (2011) Am J Hum Genet. 88(5):608-15. Kalujnaia S., et al. (2010) FASEB J. 24(10):3981-91.

Pictures:



15% SDS-PAGE (3ug)