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IMPAD1, human recombinant

CATALOG #: 7812-10 10 μg

ALTERNATE NAMES: Inositol monophosphatase 3, GPAPP, IMP 3, IMP-3,

IMPA3

SOURCE: E. Coli

PURITY: > 90% by SDS - PAGE

MOL. WEIGHT: 37.6 kDa (349 aa, 34-359 aa + His Tag)

FORM: Liquid

FORMULATION: 0.25 mg/ml solution in PBS (pH 7.4).

STORAGE CONDITIONS:

Can be stored at 4°C short term (1-2 weeks). For long term storage, aliquot and store at -20°C or -70°C. Avoid repeated freezing and thawing cycles.

DESCRIPTION:

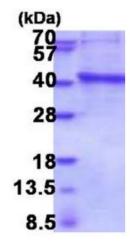
IMPAD1, also known as Inositol monophosphatase 3, is a member of the inositol monophosphatase family. IMPAD1 is localized to the Golgi apparatus and catalyzes the hydrolysis of phosphoadenosine phosphate (PAP) to adenosine monophosphate (AMP). Mutations in this gene are a cause of GRAPP type chondrodysplasia with joint dislocations, and a pseudogene of this gene is located on the long arm of chromosome 1. Recombinant human IMPAD1 protein, fused to His-tag at N-terminus, was expressed in E.coli and purified by conventional chromatography, after refolding of the isolated inclusion bodies in a renaturation buffer.

AMINO ACID SEQUENCE:

MGSSHHHHHH SSGLVPRGSH MGSGRFSLFG LGGEPGGGAA **GPAAAADGGT VDLREMLAVS** VLAAVRGGDE 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 IEGGLLASIR MNHQALVRKL PDLEKTGHK

BIOLOGICAL ACTIVITY:

Specific activity > 3300 pmole/min/ μ g, its ability to dephosphorylate adenosine 3'5'-diphosphate sodium salt at pH 7.5, 25°C.



15% SDS-PAGE (3ug)

Human Recombinant IMPAD1

RELATED PRODUCTS:

Proteins and Enzymes

FOR RESEARCH USE ONLY! Not to be used in humans.

