

Product datasheet

Recombinant Mouse BAPX1 protein (Tagged) ab226436

1 Image

Description

Product name	Recombinant Mouse BAPX1 protein (Tagged)	
Purity	> 90 % Immunogen affinity purified.	
Expression system	Escherichia coli	
Accession	P97503	
Protein length	Full length protein	
Animal free	No	
Nature	Recombinant	
Species	Mouse	
Sequence	MAVRGSGLTPFSIQAILNKKEERGGLATPEGRPAPGGTE VAVTAAPAVC CWRIFGETEAGALGGAEDSLLASPARTRTAVGQSAESPG GWSDSALSSEE NEGRRRCADVPGASGTGRARVTLGLDQPGCELHAAKDL EEEAPVRSDSEM SASVSGDHSPRGEDDSVSPGGARVPLRGAAGSGASG GQAGGVEEEEPEA APKPRKKRSRAAFSHAQVFELERRFNHQRYLSGPERADL AASLKLTTETQV KWFQNRRTYKTKRRQMAADLLASAPAAKKVAVKVLVRDD QRQYLPGEVLR PPSLLPLQPSYYYPYCLPGWALSTCAAAAAGTQ	
Predicted molecular weight	40 kDa including tags	
Amino acids	1 to 333	
Tags	His tag N-Terminus	
Additional sequence information	N-terminal 10xHis tag and C-terminal Myc tag.	

Specifications

Our [Abpromise guarantee](#) covers the use of **ab226436** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications SDS-PAGE

Form Liquid

Preparation and Storage

Stability and Storage Shipped at 4°C. Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle.
Constituents: 50% Glycerol (glycerin, glycerine), Tris buffer

General Info

Function Transcriptional repressor that acts as a negative regulator of chondrocyte maturation. PLays a role in distal stomach development; required for proper antral-pyloric morphogenesis and development of antral-type epithelium. In concert with GSC, defines the structural components of the middle ear; required for tympanic ring and gonium development and in the regulation of the width of the malleus.

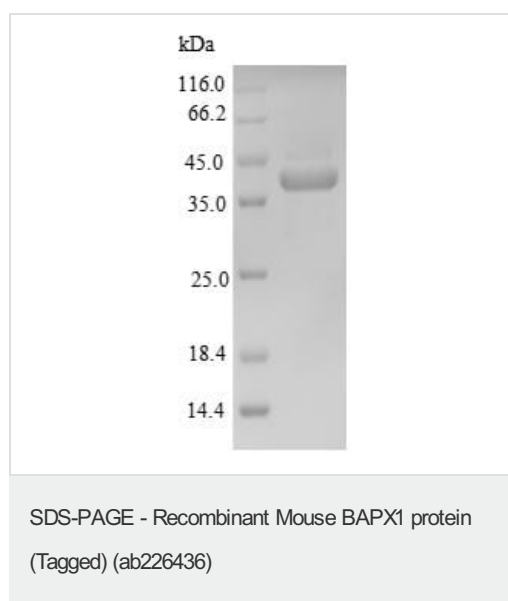
Tissue specificity Expressed at highest levels in cartilage, bone (osteosarcoma) and gut (small intestine and colon), whereas moderate expression is seen in trachea and brain. Expressed in visceral mesoderm and embryonic skeleton.

Involvement in disease Defects in NKX3-2 are the cause of spondylo-megaepiphyseal-metaphyseal dysplasia (SMMD) [MIM:613330]. It is a skeletal dysplasia characterized by disproportionate short stature with a short and stiff neck and trunk, relatively long limbs that may show flexion contractures of the distal joints, delayed and impaired ossification of the vertebral bodies, the presence of large epiphysea ossification centers and wide growth plates in the long tubular bones and numerous pseudoepiphyses of the short tubular bones in hands and feet.

Sequence similarities Belongs to the NK-3 homeobox family.
Contains 1 homeobox DNA-binding domain.

Cellular localization Nucleus.

Images



(Tris-Glycine gel) Discontinuous SDS-PAGE (reduced) analysis of ab226436 with 5% enrichment gel and 15% separation gel.

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