

Product datasheet

Recombinant Human beta Actin protein ab235730

Description

Product name	Recombinant Human beta Actin protein	
Purity	> 85 % SDS-PAGE.	
Expression system	Mammalian	
Accession	P60709	
Protein length	Full length protein	
Animal free	No	
Nature	Recombinant	
Species	Human	
Sequence	<p>MDDDIAALVVDNGSGMCKAGFAGDDAPRAVFPSPVGRPR HQGVMVGMGQK DSYVGDEAQSKRGILTLKYPIEHGIVTNWDDMEKIWHHTFY NELRVAPEE HPVLLTEAPLNPKANREKMTQIMFETFNTPAMYVAIQAVLS LYASGRITG VMDSGDGVTHTVPIYEGYALPHAILRLDLAGRDLTDYLMKI LTERGYSF TTAEREMRDIKEKLCYVALDFEQEMATAASSSSLEKSYE LPDGQVITI GNERFRCPEALFQPSFLGMESCGIHETTFNSIMKCDVDIR KDLYANTVLS GGTMYPGIADRMQKEITALAPSTMKIKIAPPKYSVWIG GSILASLS TFQQMWISKQEYDESGPSVHRKCF</p>	
Predicted molecular weight	42 kDa	
Amino acids	1 to 375	

Specifications

Our [Abpromise guarantee](#) covers the use of **ab235730** 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. Store at -20°C or -80°C. Avoid freeze / thaw cycle.

Constituents: Tris buffer, 50% Glycerol (glycerin, glycerine)

General Info

Function

Actins are highly conserved proteins that are involved in various types of cell motility and are ubiquitously expressed in all eukaryotic cells.

Involvement in disease

Defects in ACTB are a cause of dystonia juvenile-onset (DYTJ) [MIM:607371]. DYTJ is a form of dystonia with juvenile onset. Dystonia is defined by the presence of sustained involuntary muscle contraction, often leading to abnormal postures. DYTJ patients manifest progressive, generalized, dopa-unresponsive dystonia, developmental malformations and sensory hearing loss.

Sequence similarities

Belongs to the actin family.

Post-translational modifications

ISGylated.

Cellular localization

Cytoplasm > cytoskeleton. Localized in cytoplasmic mRNP granules containing untranslated mRNAs.

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