



KBTBa mouse mAb

Catalog No	BYmab-11089
Isotype	IgG
Reactivity	Human;Rat
Applications	WB
Gene Name	KBTD10 KRP1
Protein Name	KBTBa
Immunogen	Synthesized peptide derived from human KBTBa AA range: 268-318
Specificity	This antibody detects endogenous levels of KBTBa at Human/Rat
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	
Observed Band	
Cell Pathway	Cytoplasm . Cytoplasm, cytoskeleton . Cell projection, pseudopodium . Cell projection, ruffle . Cytoplasm, myofibril, sarcomere, M line . Sarcoplasmic reticulum membrane . Endoplasmic reticulum membrane . Predominantly cytoplasmic but can colocalize with F-actin at the membrane ruffle-like structures at the tips of transformation-specific pseudopodia. .
Tissue Specificity	Sarcomeric muscle.
Function	function:Required for pseudopod elongation in transformed cells. Substrate-specific adapter of an E3 ubiquitin-protein ligase complex which mediates the ubiquitination and subsequent proteasomal degradation of target proteins.,pathway:Protein modification; protein ubiquitination.,PTM:Ubiquitinated and probably targeted for proteasome-independent degradation.,similarity:Contains 1 BTB (POZ) domain.,similarity:Contains 5 Kelch repeats.,subcellular location:Predominantly cytoplasmic but can co-localize with F-actin at the membrane ruffle-like structures at the tips of transformation-specific pseudopodia.,subunit:Interacts with NRAP (By similarity). Part of a complex that

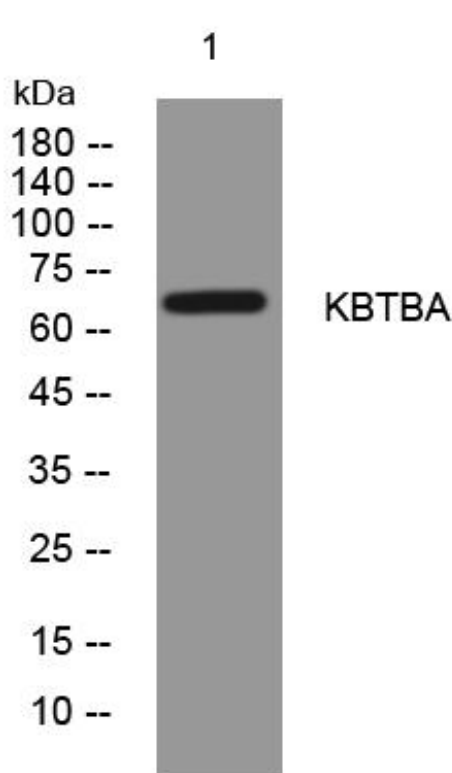
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contains CUL3, RBX1 and KBTBD10.,tissue specificity:Sarcomeric muscle.,

Background	This gene is a member of the kelch-like family. The encoded protein contains a BACK domain, a BTB/POZ domain, and 5 Kelch repeats. This protein is thought to function in skeletal muscle development and maintenance. Mutations in this gene have been associated with nemaline myopathy (NM), a rare congenital muscle disorder. [provided by RefSeq, Mar 2015],
matters needing attention	Avoid repeated freezing and thawing!
Usage suggestions	This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



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