



KBTBAb rabbit pAb

Catalog No	YP-Ab-11089
Isotype	IgG
Reactivity	Human;Rat
Applications	WB;IHC
Gene Name	KBTD10 KRP1
Immunogen	Synthesized peptide derived from human KBTBAA range: 268-318
Specificity	This antibody detects endogenous levels of KBTBAA at Human/Rat
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Polyclonal, Rabbit,IgG
Purification	The antibody was affinity-purified from rabbit serum by affinity-chromatography using specific immunogen.
Dilution	WB 1:500-2000;IHC-p 1:50-300
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	KBTBAA;KBTD10 KRP1
Observed Band	
Calculated Molecular Weight	
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



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],

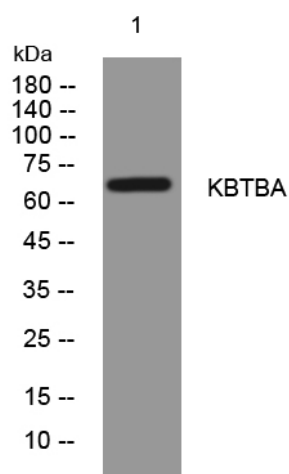
matters needing attention

Avoid repeated freezing and thawing!

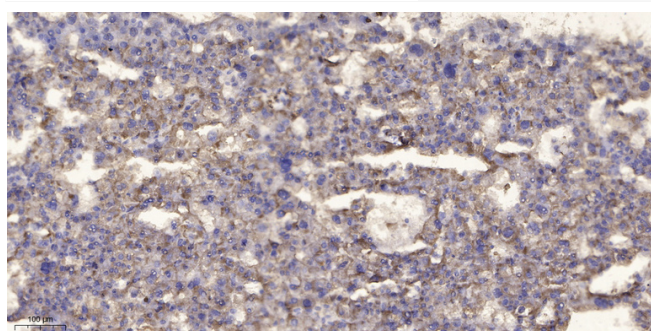
Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western blot analysis of lysates from HCT116 cells, primary antibody was diluted at 1:1000, 4° over night



Immunohistochemical analysis of paraffin-embedded human liver cancer. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 45min).