



# SCYL1BP1 Polyclonal Antibody

<b>Catalog No</b>	BYab-15046
<b>Isotype</b>	IgG
<b>Reactivity</b>	Human;Mouse;Rat
<b>Applications</b>	WB;ELISA;IHC
<b>Gene Name</b>	GORAB
<b>Protein Name</b>	RAB6-interacting golgin
<b>Immunogen</b>	The antiserum was produced against synthesized peptide derived from the N-terminal region of human GORAB. AA range:1-50
<b>Specificity</b>	SCYL1BP1 Polyclonal Antibody detects endogenous levels of SCYL1BP1 protein.
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source</b>	Polyclonal, Rabbit,IgG
<b>Purification</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Dilution</b>	WB 1:500-2000;IHC-p 1:50-300; ELISA 2000-20000
<b>Concentration</b>	1 mg/ml
<b>Purity</b>	≥90%
<b>Storage Stability</b>	-20°C/1 year
<b>Synonyms</b>	GORAB; NTKLBP1; SCYL1BP1; RAB6-interacting golgin; N-terminal kinase-like-binding protein 1; NTKL-BP1; NTKL-binding protein 1; hNTKL-BP1; SCY1-like 1-binding protein 1; SCYL1-BP1; SCYL1-binding protein 1
<b>Observed Band</b>	45kD
<b>Cell Pathway</b>	Cytoplasm . Golgi apparatus .
<b>Tissue Specificity</b>	Embryo,Pancreas,Testis,Trachea,
<b>Function</b>	caution:It is uncertain whether Met-1 or Met-26 is the initiator.,disease:Defects in GORAB are the cause of geroderma osteodysplasticum (GO) [MIM:231070]; also known as geroderma osteodysplastica or Walt Disney dwarfism. GO is a rare autosomal recessive disorder characterized by lax, wrinkled skin, joint laxity and a typical face with a prematurely aged appearance. Skeletal signs include severe osteoporosis leading to frequent fractures, malar and mandibular hypoplasia and a variable degree of growth retardation.,similarity:Belongs to the GORAB family.,subunit:Interacts with SCYL1 (By similarity). Interacts with RCHY1 and RAB6A/RAB6.,

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**Background**

golgin, RAB6 interacting(GORAB) Homo sapiens This gene encodes a member of the golgin family, a group of coiled-coil proteins localized to the Golgi. The encoded protein may function in the secretory pathway. The encoded protein, which also localizes to the cytoplasm, was identified by interactions with the N-terminal kinase-like protein, and thus it may function in mitosis. Mutations in this gene have been associated with geroderma osteodysplastica. Alternatively spliced transcript variants have been described. [provided by RefSeq, Mar 2009],

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**matters needing attention**

Avoid repeated freezing and thawing!

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**Usage suggestions**

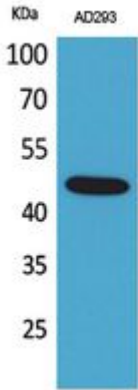
This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

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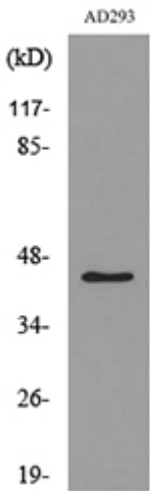
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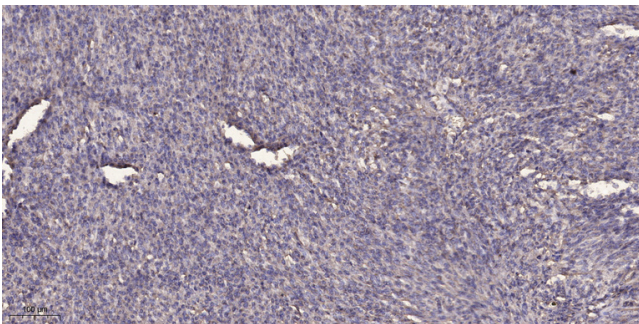
## Products Images



Western Blot analysis of AD293 cells using SCYL1BP1 Polyclonal Antibody. Secondary antibody(catalog#:RS0002) was diluted at 1:20000



Western blot analysis of lysate from AD293 cells, using GORAB Antibody.



Immunohistochemical analysis of paraffin-embedded human Colon 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).