

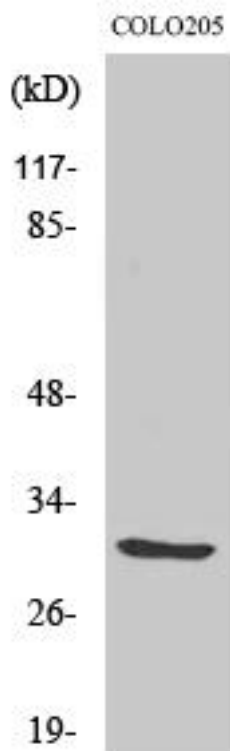


# ABHD11 Monoclonal Antibody

<b>Catalog No</b>	YP-mAb-02450
<b>Isotype</b>	IgG
<b>Reactivity</b>	Human;Rat;Mouse;
<b>Applications</b>	WB
<b>Gene Name</b>	ABHD11
<b>Protein Name</b>	Abhydrolase domain-containing protein 11
<b>Immunogen</b>	The antiserum was produced against synthesized peptide derived from human ABHD11. AA range:161-210
<b>Specificity</b>	ABHD11 Monoclonal Antibody detects endogenous levels of ABHD11 protein.
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source</b>	Monoclonal, Mouse,IgG
<b>Purification</b>	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Dilution</b>	WB 1:500-1:2000
<b>Concentration</b>	1 mg/ml
<b>Purity</b>	≥90%
<b>Storage Stability</b>	-20°C/1 year
<b>Synonyms</b>	ABHD11; WBSCR21; PP1226; Abhydrolase domain-containing protein 11; Williams-Beuren syndrome chromosomal region 21 protein
<b>Observed Band</b>	32kD
<b>Cell Pathway</b>	mitochondrion,
<b>Tissue Specificity</b>	Ubiquitously expressed.
<b>Function</b>	caution:It is uncertain whether Met-1 or Met-10 is the initiator.,sequence caution:Wrong choice of frame.,similarity:Belongs to the AB hydrolase superfamily.,tissue specificity:Ubiquitously expressed.,
<b>Background</b>	This gene encodes a protein containing an alpha/beta hydrolase fold domain. This gene is deleted in Williams syndrome, a multisystem developmental disorder caused by the deletion of contiguous genes at 7q11.23. [provided by RefSeq, Mar 2016],
<b>matters needing attention</b>	Avoid repeated freezing and thawing!
<b>Usage suggestions</b>	This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.



## Products Images



Western Blot analysis of various cells using ABHD11 Monoclonal Antibody