

Product Data Sheet

Anti-Beta-catenin Antibody

Catalog #	Source	Popotivity	Applications			
Catalog #	Source	Reactivity	Applications			
CPA9841	Mouse	Н	WB, IH			
Description	on Mouse monoclonal antibody to Beta-catenin					
Immunogen	KLH-	conjugated synthetic	peptide encompassing a sequence within human			
	Beta	Beta-catenin. The exact sequence is proprietary.				
Purification	The	antibody was purified	by immunogen affinity chromatography.			
Specificity	Reco	gnizes endogenous le	vels of Beta-catenin protein.			
Clonality	Mon	oclonal				
Conjugation						
Form	Mou	Mouse IgG2b. Liquid in PBS containing 50% glycerol, 0.2% BSA and 0.01% sodium				
	azide	2.				
Dilution	WB (1/500 - 1/1000), IH (1/1	1/300)			
Gene Symbol	CTNI	NB1				
Alternative Na	ames CTNI	NB; Catenin beta-1; Be	ta-catenin			
Entrez Gene	1499	1499 (Human)				
SwissProt	P352	22 (Human)				
Storage/Stabi	lity Ship	ped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid			
	freez	e/thaw cycles.				

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

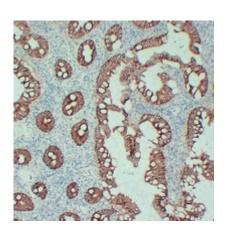
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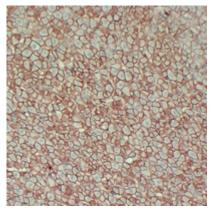
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Immunohistochemical analysis of Beta-catenin staining in human colon carcinoma formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then inc



Immunohistochemical analysis of Beta-catenin staining in human liver formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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