

Product Data Sheet

Anti-Beta-catenin Antibody

Catalog #	Source	Reactivity	Applications			
CPA9840	Mouse	Н	IH			
Description	N	Mouse monoclonal antibody to Beta-catenin				
Immunogen	К	(LH-conjugated synthetic p	eptide encompassing a sequence within human			
	В	eta-catenin. The exact sec	uence is proprietary.			
Purification	Т	he antibody was purified b	by immunogen affinity chromatography.			
Specificity	R	ecognizes endogenous lev	els of Beta-catenin protein.			
Clonality	Ν	Aonoclonal				
Conjugation						
Form	Ν	/louse IgG2b. Liquid in PBS	containing 50% glycerol, 0.2% BSA and 0.01% sodium			
	a	zide.				
Dilution	IF	H (1/100 - 1/300)				
Gene Symbol	C	TNNB1				
Alternative Na	ames C	TNNB; Catenin beta-1; Bet	a-catenin			
Entrez Gene	1	.499 (Human)				
SwissProt	P	935222 (Human)				
Storage/Stabi	lity S	hipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid			
	fr	reeze/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

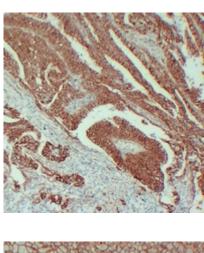
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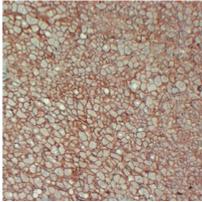
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Immunohistochemical analysis of Beta-catenin staining in human colon carcinoma formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then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