

## **Product Data Sheet**

### **Anti-SPTBN5** Antibody

Catalog #	Source	Reactivity	Applications		
CPA5987	Rabbit	Н	WB, IH, IC		
Description		Rabbit polyclonal antibody	to SPTBN5		
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
		region of human SPTBN5. T	he exact sequence is proprietary.		
Purification		The antibody was purified I	oy immunogen affinity chromatography.		
Specificity		Recognizes endogenous lev	els of SPTBN5 protein.		
Clonality	nality Polyclonal				
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
		and 0.01% sodium azide.			
Dilution WB (1/500		WB (1/500 - 1/1000), IH (1/5	0 - 1/1000), IH (1/50 - 1/200), IC (1/100 - 1/500)		
Gene Symbol		SPTBN5			
Alternative Names		BSPECV; HUBSPECV; HUSPECV; Spectrin beta chain non-erythrocytic 5; Beta-V			
		spectrin			
Entrez Gene		51332 (Human)			
SwissProt		Q9NRC6 (Human)			
<b>Storage/Stability</b> Shipped at 4°C. Upon delivery aliquot and stor		Shipped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid		
		freeze/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

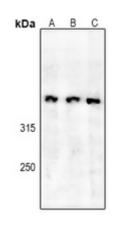
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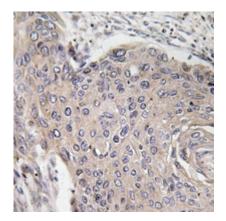
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Western blot analysis of SPTBN5 expression in HEK293T (A), SGC7901 (B), Panc1 (C) whole cell lysates. (Predicted band size: 416 kD; Observed band size: 416 kD)



Immunohistochemical analysis of SPTBN5 staining in human lung cancer 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.



Immunocytochemistry analysis of SPTBN5 staining in HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a HRP-conjugated secondary antibody in PBS at room temperature. DAB was used as the chromogen.

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