

Product Data Sheet

Anti-B3GALTL Antibody

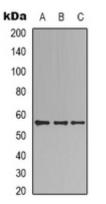
Catalog #	Source	Reactivity	Applications		
CPA7208	Rabbit	Н	WB, IH		
Description		Rabbit polyclonal antibody	to B3GALTL		
Immunogen		KLH-conjugated synthetic p	eptide encompassing a sequence within the C-term		
	I	region of human B3GALTL.	The exact sequence is proprietary.		
Purification		The antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous lev	els of B3GALTL protein.		
Clonality		Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	i	and 0.01% sodium azide.			
Dilution	,	WB (1/500 - 1/1000), IH (1/1	00 - 1/200)		
Gene Symbol		B3GALTL			
Alternative Na	ames	B3GTL; Beta-1,3-glucosyltra	nsferase; Beta3Glc-T; Beta-3-glycosyltransferase-like		
Entrez Gene		145173 (Human)			
SwissProt		Q6Y288 (Human)			
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	1	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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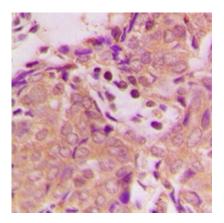




For research purposes only, not for human use

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Western blot analysis of B3GALTL expression in HEK293T (A), MCF7 (B), HepG2 (C) whole cell lysates. (Predicted band size: 56 kD; Observed band size: 57 kD)



Immunohistochemical analysis of B3GALTL staining in human breast 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.

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