

Product Data Sheet

Anti-PEX11G Antibody

Catalog #	Source	Reactivity	Applications		
CPA4149	Rabbit	H, M, R, Mk	WB, IH		
Description	ļ	Rabbit polyclonal antibody to PEX11G			
Immunogen	I	KLH-conjugated synthetic pep	tide encompassing a sequence within the N-term		
	I	region of human PEX11G. The	exact sequence is proprietary.		
Purification	-	The antibody was purified by	mmunogen affinity chromatography.		
Specificity	I	Recognizes endogenous levels	of PEX11G protein.		
Clonality Polyclonal					
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ä	and 0.01% sodium azide.			
Dilution	v	WB (1/500 - 1/1000), IH (1/100	- 1/200)		
Gene Symbol	I	PEX11G			
Alternative Na	ames	PEX11C; Peroxisomal membra	ne protein 11C; Peroxin-11C; Peroxisomal biogenesis		
	1	factor 11C; Protein PEX11 hor	nolog gamma; PEX11-gamma		
Entrez Gene		92960 (Human); 69129 (Mouse)			
SwissProt	(Q96HA9 (Human); Q6P6M5 (I	/louse)		
Storage/Stabi	lity S	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid			
	f	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

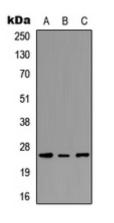
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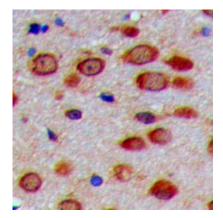


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Western blot analysis of PEX11G expression in MCF7 (A), NS-1 (B), PC12 (C) whole cell lysates. (Predicted band size: 26 kD; Observed band size: 26 kD)



Immunohistochemical analysis of PEX11G staining in human brain 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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