

Product Data Sheet

Anti-MCT12 Antibody

Catalog #	Source	Reactivity	Applications		
CPA5215	Rabbit	Η, Μ	WB, IH		
Description	R	Rabbit polyclonal antibody	o MCT12		
Immunogen	K	(LH-conjugated synthetic pe	eptide encompassing a sequence within the center		
	r	egion of human MCT12. Th	e exact sequence is proprietary.		
Purification	Т	he antibody was purified b	y immunogen affinity chromatography.		
Specificity	R	Recognizes endogenous leve	els of MCT12 protein.		
Clonality	Р	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	and 0.01% sodium azide.			
Dilution	V	NB (1/500 - 1/1000), IH (1/10	0 - 1/200)		
Gene Symbol	S	SLC16A12			
Alternative Na	ames N	MCT12; Monocarboxylate t	ansporter 12; MCT 12; Solute carrier family 16 member		
	1	.2			
Entrez Gene		387700 (Human); 240638 (Mouse)			
SwissProt	C	Q6ZSM3 (Human); Q8BGC3	(Mouse)		
Storage/Stabi	lity S	hipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fi	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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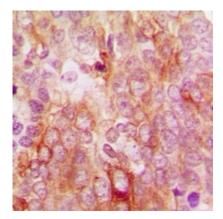
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kDa A B C 130 95 72 55 Western blot analysis of MCT12 expression in HEK293T (A), A549 (B), HepG2 (C) whole cell lysates. (Predicted band size: 56 kD; Observed band size: 60 kD)



Immunohistochemical analysis of MCT12 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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