

Product Data Sheet

Anti-COX6C Antibody

Catalog #	Source	Reactivity	Applications		
CPA5938	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Rabb	it polyclonal antibody	to COX6C		
Immunogen	KLH-o	conjugated synthetic p	eptide encompassing a sequence within the center		
	regio	region of human COX6C. The exact sequence is proprietary.			
Purification	The a	The antibody was purified by immunogen affinity chromatography.			
Specificity	Reco	gnizes endogenous lev	els of COX6C protein.		
Clonality	Polyc	lonal			
Conjugation					
Form	Liqui	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and ().01% sodium azide.			
Dilution	WB (1/500 - 1/1000), IH (1/5	0 - 1/200), IF/IC (1/100 - 1/500)		
Gene Symbol	COX6	C			
Alternative Na	ames Cytoo	chrome c oxidase subu	nit 6C; Cytochrome c oxidase polypeptide VIc		
Entrez Gene	1345	(Human); 12864 (Mo	use)		
SwissProt	P096	69 (Human); Q9CPQ1	(Mouse)		
Storage/Stabi	lity Shipp	oed at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid		
	freez	e/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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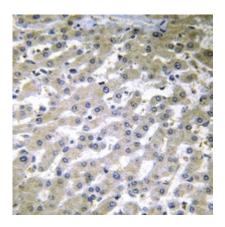
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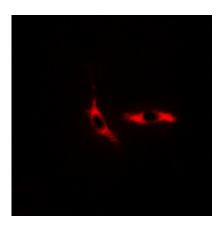
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Western blot analysis of COX6C expression in rat brain (A), mouse brain (B) whole cell lysates. (Predicted band size: 8 kD; Observed band size: 13 kD)



Immunohistochemical analysis of COX6C staining in human liver 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.



Immunofluorescent analysis of COX6C staining in HepG2 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 Alexa Fluor 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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