

Product Data Sheet

Anti-COX5B Antibody

Catalog #	Source	Reactivity	Applications		
CPA3867	Rabbit	H, M, R, B, P	WB, IH		
Description	Ral	bbit polyclonal antibody to	O COX5B		
Immunogen	KLł	H-conjugated synthetic per	otide encompassing a sequence within the N-term		
	reg	ion of human COX5B. The	exact sequence is proprietary.		
Purification	The	e antibody was purified by	immunogen affinity chromatography.		
Specificity	Re	cognizes endogenous level	ls of COX5B protein.		
Clonality	Pol	Polyclonal			
Conjugation					
Form	Liq	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	d 0.01% sodium azide.			
Dilution	WE	3 (1/500 - 1/1000) <i>,</i> IH (1/100) - 1/200)		
Gene Symbol	CO	X5B			
Alternative Na	ames Cyt	ochrome c oxidase subuni	it 5B mitochondrial; Cytochrome c oxidase polypeptide		
	Vb				
Entrez Gene	132	29 (Human); 94194 (Rat)			
SwissProt	P10	0606 (Human); P19536 (M	ouse); P12075 (Rat)		
Storage/Stabi	l <mark>ity</mark> Shi	pped at 4°C. Upon delivery	y aliquot and store at -20°C for one year. Avoid		
	fre	eze/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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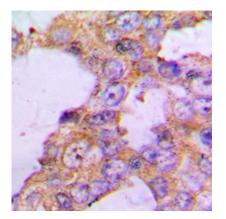


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kDa A B C Western blot analysis of COX5B expression in HEK293T (A), Raw264.7 (B), H9C2 (C) whole cell lysates. (Predicted band size: 13 kD; Observed band size: 14 kD)



Immunohistochemical analysis of COX5B 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.

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