

## **Product Data Sheet**

### Anti-COX4-1 Antibody

Catalog #	Source	Reactivity	Applications	
CPA3866	Rabbit	Н	WB, IH, IF/IC	
Description		Rabbit polyclonal antibody	v to COX4-1	
Immunogen		KLH-conjugated synthetic	peptide encompassing a sequence within the N-term	
		region of human COX4-1.	The exact sequence is proprietary.	
Purification		The antibody was purified	by immunogen affinity chromatography.	
Specificity		Recognizes endogenous le	vels of COX4-1 protein.	
Clonality		Polyclonal		
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
		and 0.01% sodium azide.		
Dilution	,	WB (1/500 - 1/1000), IH (1/:	100 - 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol	1	COX4I1		
Alternative N	ames	COX4; Cytochrome c oxida	se subunit 4 isoform 1, mitochondrial; Cytochrome c	
		oxidase polypeptide IV; Cy	tochrome c oxidase subunit IV isoform 1; COX IV-1	
Entrez Gene		1327 (Human)		
SwissProt		P13073 (Human)		
Storage/Stabi	ility	Shipped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid	
		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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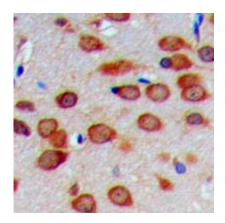
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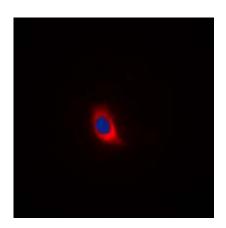
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Western blot analysis of COX4-1 expression in HepG2 (A), Jurkat (B) whole cell lysates.



Immunohistochemical analysis of COX4-1 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX. w



Immunofluorescent analysis of COX4-1 staining in HeLa 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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