

Product Data Sheet

Anti-COX4-1 Antibody

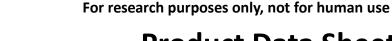
Catalog #	Source	Reactivity	Applications
CQA8238	Mouse	H, M, R, Ha, G, Mk	WB, IH, IF/IC, IP, FC
Description	Мс	ouse monoclonal antibody	to COX4-1
Immunogen	A s	synthetic peptide correspo	nding to carboxyl terminal residues of human COX IV
Purification	The	e antibody was purified by	immunogen affinity chromatography.
Specificity	Re	cognizes endogenous leve	ls of COX4-1 protein.
Clonality	Мс	onoclonal	
Conjugation			
Form	Liq	uid in PBS containing 50%	glycerol, 0.5% BSA and 0.02% sodium azide, pH 7.3.
Dilution	WE	3 (1/500 - 1/1000), IH (1/50	- 1/100), IF/IC (1/50 - 1/100), IP (1/10 - 1/50), FC (1/50 -
	1/1	.00)	
Gene Symbol	CO	X4I1	
Alternative N	ames CO	X4; Cytochrome c oxidase	subunit 4 isoform 1 mitochondrial; Cytochrome c
	oxi	dase polypeptide IV; Cyto	chrome c oxidase subunit IV isoform 1; COX IV-1
Entrez Gene	13	27 (Human); 12857 (Mous	e); 29445 (Rat)
SwissProt	P1	3073 (Human); P19783 (M	ouse); P10888 (Rat)
Storage/Stabi	lity Shi	ipped at 4°C. Upon deliver	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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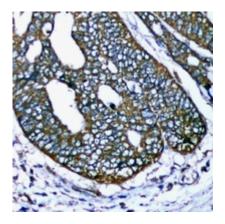


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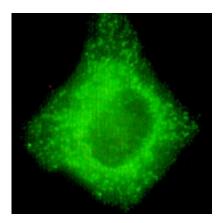
kDa
A
B
C
D
E
F

130
95
 -<

Western blot analysis of COX4-1 expression in CHOk1 (A), COS7 (B), NIH3T3 (C), Hela (D), C6 (E), mouse skeletal muscel (F) whole cell lysates. (Predicted band size: 20 kD; Observed band size: 17 kD)



Immunohistochemical analysis of COX4-1 staining in human colon 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 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 AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark.

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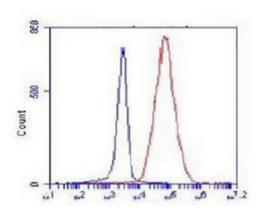
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Flow cytometric analysis of COX4-1 in K562 (red). Blue line histogram represents the isotype control.

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