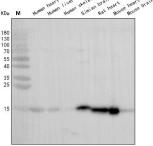


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## Anti-COX4I1 Antibody, Mouse Monoclonal, C8T

COX4, Cytochrome c oxidase polypeptide IV Catalog: MA00294HuM10-C8T Size:100µL

Basic Info	Host	Species Reactivity
	Mouse	Human/Monkey/Mouse/Rat
	Conjugate	Clonality, Isotype
	None	/
	Concentration	Immunogen
	1mg/mL	Recombinant human COX4I1 protein
	Physical State	Purification
	Liquid	Immunogen affinity purified
	Clone No	Applications
	C8T	WB/IHC
Property	Form & Buffer: Supplied in PBS, 50% glycerol, PH7.4.	
	Specificity / Sensitivity: MA00294HuM10-C8T recognizes endogenous levels	
	of human COX4I1.	
Applications	The application notes include recommended starting dilutions;	
	optimal dilutions/concentrations should be determined by the end user.	
	WB: 0.5~5µg/mL IF	F: 5~20µg/mL
	IHC: 5~20µg/mL IC	CC: 5~20µg/mL
Usage and	Store at $4^{\circ}$ C for frequent use.	
Storage	Aliquot and store at $-20^\circ\!\mathrm{C}$ for 12 months.	
	Avoid repeated freezing/thawing and violent shaking.	
	Please centrifuge it, before using.	
QC Data	mart liver unland mark mort rain	Figure 1. Application in WB
	KDa M Human Human Simian Rat heave thouse by	Wastern blot analysis of extracts of various call lines ar



Western blot analysis of extracts of various cell lines and tissues, using COX4I1 antibody MA00294HuM10-C8T at  $3\mu$ g/mL Secondary antibody: HRP Mouse Anti-Rabbit IgG (H+L) at 1: 4000 dilution. Lysates/proteins: 25ug per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit. Exposure time: 0.5s.



## QC Data



## Figure 2. Application in IHC

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of monkey brain, monkey cerebellum, monkey left ventricle, monkey kidney sections labelling COX4I1 with purified MA00294HuM10-C8T at  $10\mu$ g/mL dilution. Heat mediated antigen retrieval was performed using Heat mediated antigen retrieval using citrate buffer (PH6.0). Tissue was counterstained with Hematoxylin. Mouse specific IHC polymer detection kit HRP/DAB secondary antibody was used at 1: 2000 dilution. PBS instead of the primary antibody was used as the negative control.

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