

Product Data Sheet

Anti-NDUFA1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA4114	Rabbit	Н, М	WB, IH		
Description		Rabbit polyclonal antibody t	o NDUFA1		
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the N-term		
	I	region of human NDUFA1. T	ne exact sequence is proprietary.		
Purification		The antibody was purified b	<i>immunogen affinity chromatography</i> .		
Specificity		Recognizes endogenous leve	ls of NDUFA1 protein.		
Clonality		Polyclonal			
Conjugation					
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 (1/50	- 1/100)		
Gene Symbol		NDUFA1			
Alternative Na	ames	NADH dehydrogenase [ubiq	uinone] 1 alpha subcomplex subunit 1; Complex		
		I-MWFE; CI-MWFE; NADH-u	piquinone oxidoreductase MWFE subunit		
Entrez Gene		4694 (Human); 54405 (Mou	se)		
SwissProt		O15239 (Human); O35683 (I	Mouse)		
Storage/Stabi	lity	Shipped at 4°C. Upon delive	y 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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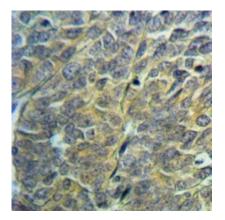
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A B

For research purposes only, not for human use

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Western blot analysis of NDUFA1 expression in HeLa (A), Raw264.7 (B) whole cell lysates. (Predicted band size: 8 kD; Observed band size: 9 kD)



Immunohistochemical analysis of NDUFA1 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.

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