

## **Product Data Sheet**

### **Anti-NDUFB5** Antibody

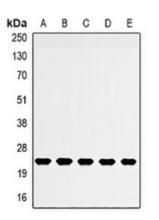
Catalog #	Source	Reactivity	Applications		
CQA2103	Rabbit	H, M, R	WB, IH		
Description	Rab	bit polyclonal antibody	to NDUFB5		
Immunogen	Rec	ombinant full length pro	tein of human NDUFB5		
Purification	The	antibody was purified b	y immunogen affinity chromatography.		
Specificity	Rec	ognizes endogenous lev	els of NDUFB5 protein.		
Clonality	Poly	clonal			
Conjugation					
Form	Liqu	uid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB	(1/500 - 1/2000), IH (1/50	) - 1/100)		
Gene Symbol	NDU	JFB5			
Alternative Na	ames NAE	OH dehydrogenase [ubio	uinone] 1 beta subcomplex subunit 5 mitochondrial;		
	Con	nplex I-SGDH; CI-SGDH;	NADH-ubiquinone oxidoreductase SGDH subunit		
Entrez Gene 47		4711 (Human); 66046 (Mouse)			
SwissProt	O43	8674 (Human); Q9CQH3	(Mouse)		
Storage/Stabi	<b>lity</b> Ship	oped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	free	ze/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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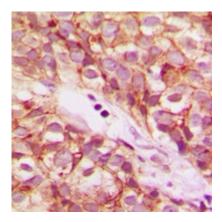




For research purposes only, not for human use

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Western blot analysis of NDUFB5 expression in HepG2 (A), SW480 (B), mouse liver (C), mouse brain (D), rat kidney (E) whole cell lysates. (Predicted band size: 15; 21 kD; Observed band size: 22 kD)



Immunohistochemical analysis of NDUFB5 staining in human breast 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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