

Product Data Sheet

Anti-UCP2 Antibody

Catalog #	Source	Reactivity	Applications		
CPA5439	Rabbit	H, M, R, B, D, S	WB, IH		
Description	Rab	bit polyclonal antibody to	UCP2		
Immunogen	KLH	-conjugated synthetic per	tide encompassing a sequence within the center		
	regi	on of human UCP2. The e	xact sequence is proprietary.		
Purification	The	antibody was purified by	immunogen affinity chromatography.		
Specificity	Rec	ognizes endogenous level	s of UCP2 protein.		
Clonality	Polyclonal				
Conjugation					
Form	Liqu	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	UCF	2			
Alternative Na	ames SLC	25A8; Mitochondrial unco	upling protein 2; UCP 2; Solute carrier family 25		
	mer	mber 8; UCPH			
Entrez Gene	735	1 (Human); 22228 (Mouse	e); 54315 (Rat)		
SwissProt	P55	851 (Human); P70406 (M	ouse); P56500 (Rat)		
Storage/Stabi	lity Ship	oped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid		
	free	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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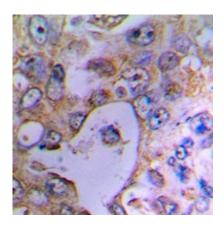
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Western blot analysis of UCP2 expression in HEK293T (A), Raw264.7 (B), PC12 (C) whole cell lysates. (Predicted band size: 33 kD; Observed band size: 35 kD)



Immunohistochemical analysis of UCP2 staining in human lung 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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