

Product Data Sheet

Anti-Pyruvate Carboxylase Antibody

Catalog #	Source	Reactivity	Applications		
CPA1857	Rabbit	H, M, R, B, C, Mk, P, S, Z	WB, IH		
Description	Rat	Rabbit polyclonal antibody to Pyruvate Carboxylase			
Immunogen	KLF	I-conjugated synthetic peptid	e encompassing a sequence within the center		
	reg	region of human Pyruvate Carboxylase. The exact sequence is proprietary.			
Purification	The	e antibody was purified by im	munogen affinity chromatography.		
Specificity	Rec	cognizes endogenous levels of	f Pyruvate Carboxylase protein.		
Clonality	Pol	yclonal			
Conjugation					
Form	Liqu	uid in 0.42% Potassium phosp	ohate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	d 0.01% sodium azide.			
Dilution	WB	8 (1/500 - 1/1000), IH (1/50 - 1/1	100)		
Gene Symbol	PC				
Alternative Na	ames Pyr	uvate carboxylase mitochond	lrial; Pyruvic carboxylase; PCB		
Entrez Gene	509	91 (Human); 25104 (Rat)			
SwissProt	P11	L498 (Human); Q05920 (Mous	se); P52873 (Rat)		
Storage/Stabi	lity Shi	pped at 4°C. Upon delivery ali	iquot 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

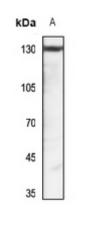
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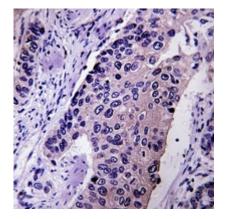
Cohesion

For research purposes only, not for human use

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Western blot analysis of Pyruvate Carboxylase expression in rat liver (A) whole cell lysates. (Predicted band size: 129 kD; Observed band size: 130 kD)



Immunohistochemical analysis of Pyruvate Carboxylase 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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