

Product Data Sheet

Anti-Catalase Antibody

Catalog #	Source	Reactivity	Applications		
CPA5904	Rabbit	H, M, R, B, D, Z	WB, IH		
Description	Rabb	Rabbit polyclonal antibody to Catalase			
Immunogen	KLH-c	onjugated synthetic peptide e	ncompassing a sequence within the center		
	regio	n of human Catalase. The exac	t sequence is proprietary.		
Purification	The a	ntibody was purified by immu	nogen affinity chromatography.		
Specificity	Reco	gnizes endogenous levels of Ca	talase protein.		
Clonality	Polyc	lonal			
Conjugation					
Form	Liquid	l in 0.42% Potassium phospha	e, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and O	.01% sodium azide.			
Dilution	WB (2	l/500 - 1/1000), IH (1/50 - 1/200)		
Gene Symbol	CAT				
Alternative Na	ames Catala	ase			
Entrez Gene	847 (Human); 12359 (Mouse); 2424	8 (Rat)		
SwissProt	P0404	40 (Human); P24270 (Mouse);	P04762 (Rat)		
Storage/Stabi	lity Shipp	ed at 4°C. Upon delivery aliqu	ot and store at -20°C for one year. Avoid		
	freeze	e/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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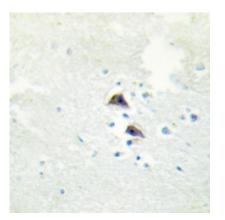
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kDa A B C D

For research purposes only, not for human use

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Western blot analysis of Catalase expression in rat prostate (A), moues kidney (B), A549 (C), LO2 (D) whole cell lysates. (Predicted band size: 59 kD; Observed band size: 60 kD)



Immunohistochemical analysis of Catalase staining in human brain 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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