

Product Data Sheet

Anti-Cytochrome P450 4Z1/2 Antibody

Catalog #	Source	Reactivity	Applications			
CPA2889	Rabbit	H, R	WB, IH			
Description	Rabl	Rabbit polyclonal antibody to Cytochrome P450 4Z1/2				
Immunogen	KLH	-conjugated synthetic p	eptide encompassing a sequence within the center			
	regio	on of human Cytochroi	ne P450 4Z1/2. The exact sequence is proprietary.			
Purification	The	antibody was purified	by immunogen affinity chromatography.			
Specificity	Reco	ognizes endogenous lev	els of Cytochrome P450 4Z1/2 protein.			
Clonality	Poly	clonal				
Conjugation						
Form	Liqu	id 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 (1/1	00 - 1/200)			
Gene Symbol	CYP4	4Z1; CYP4Z2P				
Alternative Na	ames					
Entrez Gene	1999	974 (Human)				
SwissProt	Q5V	VE4, Q8N1L4 (Human)				
Storage/Stabi	lity Ship	ped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid			
	free	ze/thaw cycles.				

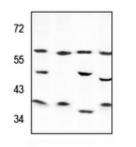
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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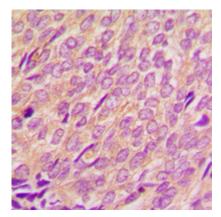
kDa A B C D



For research purposes only, not for human use

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Western blot analysis of Cytochrome P450 4Z1/2 expression in PC12 (A), MEF (B), Hela (C), MCF7 (D) whole cell lysates. (Predicted band size: 40 kD; Observed band size: 59; 40 kD)



Immunohistochemical analysis of Cytochrome P450 4Z1/2 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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