

Product Data Sheet

Anti-ZP2 Antibody

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Catalog #	Source	Reactivity	Applications	
CPA5354	Rabbit	H, M, R	WB, IH	
Description		Rabbit polyclonal antibody t	o ZP2	
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the center	
		region of human ZP2. The ex	act sequence is proprietary.	
Purification		The antibody was purified by	<i>immunogen affinity chromatography.</i>	
Specificity		Recognizes endogenous leve	ls of ZP2 protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/2000), IH (1/50	- 1/200)	
Gene Symbol		ZP2		
Alternative Na	ames	ZPA; Zona pellucida sperm-b	inding protein 2; Zona pellucida glycoprotein 2; Zp-2;	
		Zona pellucida protein A		
Entrez Gene		7783 (Human); 22787 (Mous	e); 81828 (Rat)	
SwissProt		Q05996 (Human); P20239 (N	1ouse); O54767 (Rat)	
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid	
		freeze/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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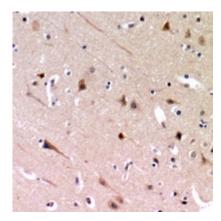
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Western blot analysis of ZP2 expression in H1792 (A), A549 (B), mouse muscle (C), rat muscle (D) whole cell lysates. (Predicted band size: 82 kD; Observed band size: 67 kD)



Immunohistochemical analysis of ZP2 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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