

Product Data Sheet

Anti-ZIP7 Antibody

Catalog #	Source	e Reactivity	Applications		
CPA2256	Rabbit	н, М, R, D, P	WB, IF/IC		
Description		Rabbit polyclonal antibody to	ZIP7		
Immunogen		KLH-conjugated synthetic pep	tide encompassing a sequence within the center		
		region of human ZIP7. The exact sequence is proprietary.			
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous level	s of ZIP7 protein.		
Clonality Polyclonal					
Conjugation					
Form Liquid in 0.42% Pota		Liquid in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IF/IC (1/1	00 - 1/500)		
Gene Symbol		SLC39A7			
Alternative N	ames	HKE4; RING5; Zinc transporte	r SLC39A7; Histidine-rich membrane protein Ke4;		
		Really interesting new gene 5	protein; Solute carrier family 39 member 7; Zrt-,		
		Irt-like protein 7; ZIP7			
Entrez Gene		7922 (Human); 14977 (Mouse	•)		
SwissProt		Q92504 (Human); Q31125 (N	ouse)		
Storage/Stabi	lity	Shipped at 4°C. Upon delivery	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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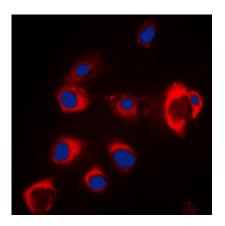
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Western blot analysis of ZIP7 expression in mouse kidney (A), rat kidney (B) whole cell lysates. (Predicted band size: 50 kD; Observed band size: 50 kD)



Immunofluorescent analysis of ZIP7 staining in MCF7 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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