

Product Data Sheet

Anti-Copine 8 Antibody

Catalog #	Source	Reactivity	Applications
CPA5204	Rabbit	H, M, R, B, Z	WB, IH
Description	Rab	bit polyclonal antibody t	o Copine 8
Immunogen	KLH	-conjugated synthetic pe	ptide encompassing a sequence within the N-term
	regi	on of human Copine 8. T	he exact sequence is proprietary.
Purification	The	antibody was purified by	<i>immunogen affinity chromatography.</i>
Specificity	Reco	ognizes endogenous leve	ls of Copine 8 protein.
Clonality	Poly	rclonal	
Conjugation			
Form	Liqu	id in 0.42% Potassium pl	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB	(1⁄500 - 1⁄1000), IH (1⁄10	0 - 1/200)
Gene Symbol	CPN	E8	
Alternative Na	ames Cop	ine-8; Copine VIII	
Entrez Gene	144	402 (Human); 66871 (Mo	ouse)
SwissProt	Q86	YQ8 (Human); Q9DC53 (Mouse)
Storage/Stabi	lity Ship	ped at 4°C. Upon deliver	y 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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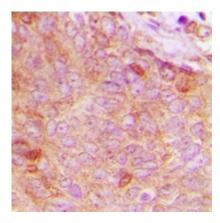
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kDa A B C D E 130
95
72
55 For research purposes only, not for human use

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Western blot analysis of Copine 8 expression in A549 (A), HEK293T (B), COS7 (C), CT26 (D), PC12 (E) whole cell lysates. (Predicted band size: 63 kD; Observed band size: 65 kD)



Immunohistochemical analysis of Copine 8 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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