

Product Data Sheet

Anti-ERGIC3 Antibody

Catalog #	Source	Reactivity	Applications		
CPA5162	Rabbit	H, M, R, B, Mk	WB, IH, IF/IC		
Description	Rabb	it polyclonal antibody to	ERGIC3		
Immunogen	KLH-	conjugated synthetic per	otide encompassing a sequence within the C-term		
	regio	on of human ERGIC3. The	e exact sequence is proprietary.		
Purification	The	antibody was purified by	immunogen affinity chromatography.		
Specificity	Reco	gnizes endogenous level	s of ERGIC3 protein.		
Clonality	Poly	clonal			
Conjugation					
Form	Liqui	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	0.01% sodium azide.			
Dilution	WB (1/500 - 1/1000), IH (1/100) - 1/200), IF/IC (1/100 - 1/500)		
Gene Symbol	ERGI	C3			
Alternative Na	ames C20c	orf47; ERV46; SDBCAG84	; Endoplasmic reticulum-Golgi intermediate		
	com	partment protein 3; Sero	logically defined breast cancer antigen NY-BR-84		
Entrez Gene	5161	.4 (Human); 66366 (Mou	se)		
SwissProt	Q9Y2	282 (Human); Q9CQE7 (N	Aouse)		
Storage/Stabi	lity Ship	oed at 4°C. Upon delivery	y aliquot and store at -20°C for one year. Avoid		
	freez	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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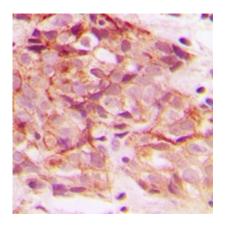
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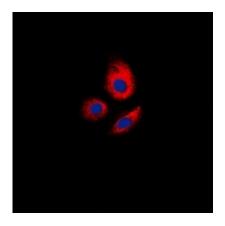
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Western blot analysis of ERGIC3 expression in rat brain (A) whole cell lysates. (Predicted band size: 43 kD; Observed band size: 43 kD)



Immunohistochemical analysis of ERGIC3 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.



Immunofluorescent analysis of ERGIC3 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 hidified 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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