

Product Data Sheet

Anti-HPC2 Antibody

Catalog #	Source	Reactivity	Applications		
CQA4355	Rabbit	H, M, R	WB, IH		
Description	Ra	abbit polyclonal antibody to	0 HPC2		
Immunogen	Re	ecombinant fusion protein	of human HPC2. The exact sequence is proprietary.		
Purification	Th	he antibody was purified by	immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous leve	s of HPC2 protein		
Clonality	Рс	olyclonal			
Conjugation					
Form	Lie	quid in 0.42% Potassium pl	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	ar	nd 0.01% sodium azide.			
Dilution	W	/B (1⁄500 - 1⁄1000), IH (1⁄50	- 1/200)		
Gene Symbol	EL	LAC2			
Alternative Names		HPC2; Zinc phosphodiesterase ELAC protein 2; ElaC homolog protein 2; Heredity			
	pr	rostate cancer protein 2; Ri	oonuclease Z 2; RNase Z 2; tRNA 3 endonuclease 2;		
	tR	Nase Z 2			
Entrez Gene	60	0528 (Human)			
SwissProt	Q	9BQ52 (Human)			
Storage/Stabi	lity Sh	nipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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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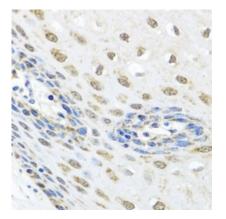
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For research purposes only, not for human use

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Western blot analysis of HPC2 expression in MCF7 (A) whole cell lysates. (Predicted band size: 50; 56; 87; 92 kD; Observed band size: 105 kD)



Immunohistochemical analysis of HPC2 staining in human esophagus 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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