

Product Data Sheet

Anti-FHIT Antibody

	Catalog #	Source	e Reactivity	Applications		
	CPA1425	Rabbit	: Н	WB, IH, IF/IC		
	Description		Rabbit polyclonal antibody to FHIT			
Immunogen			KLH-conjugated synthetic peptide encompassing a sequence within the center			
			region of human FHIT. The exact sequence is proprietary.			
Purification			The antibody was purified by immunogen affinity chromatography.			
Specificity			Recognizes endogenous levels of FHIT protein.			
	Clonality	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 - 1/1000), IH (1/100 - 1/200), IF/IC (1/100 - 1/500)			
	Gene Symbol		FHIT			
Alternative Names		mes	Bis(5'-adenosyl)-triphosphatase; AP3A hydrolase; AP3Aase; Diadenosine			
			5',5'''-P1,P3-triphosphate hydrolase; Dinucleosidetriphosphatase; Fragile histidine			
			triad protein			
	Entrez Gene		2272 (Human)			
	SwissProt		P49789 (Human)			
	Storage/Stabil	ge/Stability 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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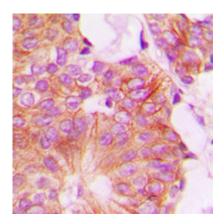
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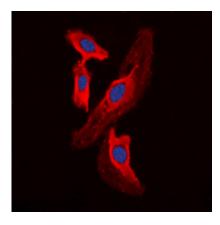
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Western blot analysis of FHIT expression in MCF7 (A), SGC7901 (B) whole cell lysates. (Predicted band size: 16 kD; Observed band size: 20 kD)



Immunohistochemical analysis of FHIT 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 FHIT staining in HEK293T 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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