

Product Data Sheet

Anti-DFF45 Antibody

Catalog #	Source	e Reactivity	Applications	
CPA1338	Rabbit	Н	WB, IH, IF/IC	
Description		Rabbit polyclonal antibody	to DFF45	
Immunogen		KLH-conjugated synthetic p	eptide encompassing a sequence within the center	
		region of human DFF45. Th	e exact sequence is proprietary.	
Purification		The antibody was purified	by immunogen affinity chromatography.	
Specificity		Recognizes endogenous lev	els of DFF45 protein.	
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 (1/5	0 - 1/100), IF/IC (1/50 - 1/200)	
Gene Symbol		DFFA		
Alternative Na	ames	DFF1; DFF45; DNA fragmer	tation factor subunit alpha; DNA fragmentation factor 45	
		kDa subunit; DFF-45; Inhib	tor of CAD; ICAD	
Entrez Gene		1676 (Human)		
SwissProt		O00273 (Human)		
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ery 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

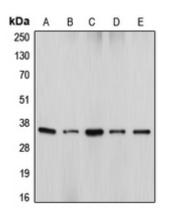
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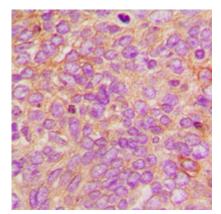


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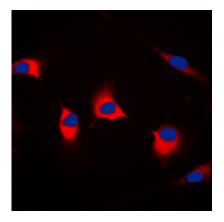
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Western blot analysis of DFF45 expression in HeLa (A), U937 (B), WR19L (C), HEK293T (D), Jurkat (E) whole cell lysates. (Predicted band size: 36 kD; Observed band size: 35 kD)



Immunohistochemical analysis of DFF45 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 DFF45 staining in HeLa 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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