

Product Data Sheet

Anti-FADD Antibody

Catalog #	Source	Reactivity	Applications	
CPA4639	Rabbit	H, Mk	WB, IH	
Description		Rabbit polyclonal antibody t	o FADD	
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the C-term	
		region of human FADD. The	exact sequence is proprietary.	
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	els of FADD 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 (1/10	0 - 1/200)	
Gene Symbol		FADD		
Alternative Na	ames	MORT1; FAS-associated dea	h domain protein; FAS-associating death	
		domain-containing protein;	Growth-inhibiting gene 3 protein; Mediator of receptor	
		induced toxicity; Protein FAI	D	
Entrez Gene		8772 (Human)		
SwissProt		Q13158 (Human)		
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry 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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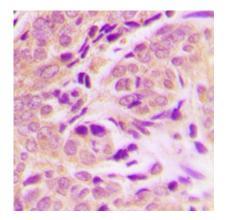
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Western blot analysis of FADD expression in A549 (A), HeLa (B) whole cell lysates. (Predicted band size: 23 kD; Observed band size: 26 kD)



Immunohistochemical analysis of FADD 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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