

Product Data Sheet

Anti-AATF Antibody

Catalog #	Source	Reactivity	Applications	
CPA2614	Rabbit	H, Mk	WB, IH, IF/IC	
Description	I	Rabbit polyclonal antibody t	o AATF	
Immunogen	I	KLH-conjugated synthetic pe	ptide encompassing a sequence within the N-term	
	ı	region of human AATF. The	exact sequence is proprietary.	
Purification	-	The antibody was purified b	y immunogen affinity chromatography.	
Specificity	I	Recognizes endogenous leve	els of AATF protein.	
Clonality	I	Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	ä	and 0.01% sodium azide.		
Dilution	N	WB (1/500 - 1/1000), IH (1/50	- 1/100), IF/IC (1/50 - 1/200)	
Gene Symbol	/	AATF		
Alternative Na	ames (CHE1; DED; Protein AATF; A	ooptosis-antagonizing transcription factor; Rb-binding	
	I	protein Che-1		
Entrez Gene		26574 (Human)		
SwissProt	(Q9NY61 (Human)		
Storage/Stabi	lity S	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid	
	f	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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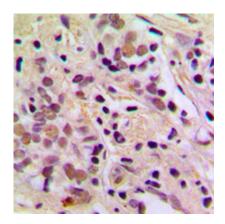
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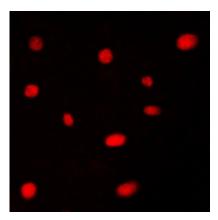
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Western blot analysis of AATF expression in Hela (A) whole cell lysates. (Predicted band size: 63 kD; Observed band size: 63 kD)



Immunohistochemical analysis of AATF 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 AATF 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.

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