

Product Data Sheet

Anti-NAT5 Antibody

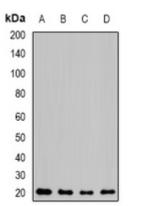
Catalog #	Source	e Reactivity	Applications	
CQA1540	Rabbit	H, M, R	WB, IH, IP	
Description		Rabbit polyclonal antibody to	DINAT5	
Immunogen		Recombinant full length prot	ein of human NAT5	
Purification		The antibody was purified by	immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	ls of NAT5 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 (1/50	- 1/200), IP (1/10 - 1/50)	
Gene Symbol		NAA20		
Alternative Names		NAT5; N-alpha-acetyltransferase 20; Methionine N-acetyltransferase;		
		N-acetyltransferase 5; N-terr	ninal acetyltransferase B complex catalytic subunit	
		NAA20; N-terminal acetyltra	nsferase B complex catalytic subunit NAT5; NatB	
		complex subunit NAT5; NatB	catalytic subunit	
Entrez Gene		51126 (Human); 67877 (Mou	ise)	
SwissProt		P61599 (Human); P61600 (N	louse)	
Storage/Stabi	ility	Shipped at 4°C. Upon deliver	y 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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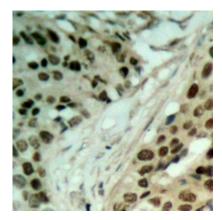




For research purposes only, not for human use

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Western blot analysis of NAT5 expression in MCF7 (A), mouse kidney (B), mouse heart (C), rat lung (D) whole cell lysates. (Predicted band size: 12; 20 kD; Observed band size: 20 kD)



Immunohistochemical analysis of NAT5 staining in human ovarian 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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