

## **Product Data Sheet**

## **Anti-NAT-15 Antibody**

Catalog #	Source	Reactivity	Applications		
CQA3327	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Ra	bbit polyclonal antibody	to NAT-15		
Immunogen	Re	combinant full length pro	otein of human NAT-15		
Purification	Th	e antibody was purified l	oy immunogen affinity chromatography.		
Specificity	Re	cognizes endogenous lev	rels of NAT-15 protein.		
Clonality	Ро	lyclonal			
Conjugation					
Form	Lic	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	d 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/2000), IH (1/5	0 - 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	NA	A60			
Alternative Na	ames HA	T4; NAT15; N-alpha-acet	yltransferase 60; Histone acetyltransferase type B protein		
	4;	HAT4; N-acetyltransferas	e 15; NatF catalytic subunit		
Entrez Gene 79903 (Human); 74		903 (Human); 74763 (Mo	ouse); 363545 (Rat)		
SwissProt	Q	H7X0 (Human); Q9DBU2	! (Mouse); Q3MHC1 (Rat)		
Storage/Stabil	l <mark>ity</mark> Sh	ipped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid		
	fre	eze/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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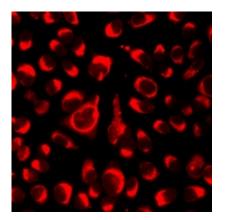
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# Cohesion

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Western blot analysis of NAT-15 expression in mouse brain (A), rat brain (B) whole cell lysates. (Predicted band size: 19; 20; 27; 28 kD; Observed band size: 27 kD)



Immunohistochemical analysis of NAT-15 staining in human colon 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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