

Product Data Sheet

Anti-CNOT1 Antibody

Catalog #	Source	Reactivity	Applications		
CQA4123	Rabbit	Н, М	WB, IF/IC		
Description	R	abbit polyclonal antibody	to CNOT1		
Immunogen	R	ecombinant fusion protein	of human CNOT1. The exact sequence is proprietary.		
Purification	Т	he antibody was purified b	y immunogen affinity chromatography.		
Specificity	R	ecognizes endogenous leve	els of CNOT1 protein		
Clonality	Р	olyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	nd 0.01% sodium azide.			
Dilution	V	VB (1/500 - 1/2000), IF/IC (1	/50 - 1/200)		
Gene Symbol	С	CNOT1			
Alternative Names		CDC39; KIAA1007; NOT1; CCR4-NOT transcription complex subunit 1;			
	С	CR4-associated factor 1; N	egative regulator of transcription subunit 1 homolog;		
	N	IOT1H; hNOT1			
Entrez Gene	2	3019 (Human); 234594 (M	ouse)		
SwissProt	А	5YKK6 (Human); Q6ZQ08 (Mouse)		
Storage/Stabil	l ity S	hipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fr	reeze/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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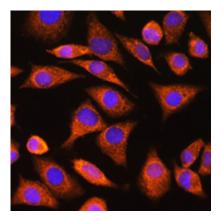
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Western blot analysis of CNOT1 expression in A549 (A), HeLa (B), mouse brain (C) whole cell lysates. (Predicted band size: 173; 241; 266 kD; Observed band size: 267 kD)



Immunofluorescent analysis of CNOT1 staining in L929 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 AF594-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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