

## **Product Data Sheet**

## **Anti-TIA1 Antibody**

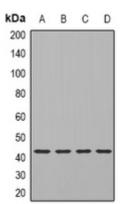
Catalog #	Source	Reactivity	Applications	
CQA2271	Rabbit	H, M, R	WB, IH	
Description		Rabbit polyclonal antibody t	D TIA1	
Immunogen		Recombinant full length prot	ein of human TIA1	
Purification		The antibody was purified by immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	ls of TIA1 protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium pl	nosphate, 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)	
Gene Symbol		TIA1		
Alternative Na	ames	Nucleolysin TIA-1 isoform p4	0; RNA-binding protein TIA-1; T-cell-restricted	
		intracellular antigen-1; TIA-1	; p40-TIA-1	
Entrez Gene		7072 (Human); 21841 (Mous	e)	
SwissProt		P31483 (Human); P52912 (N	louse)	
Storage/Stabi	lity	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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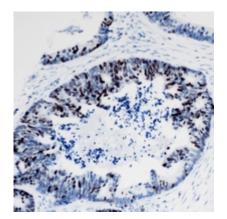




For research purposes only, not for human use

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Western blot analysis of TIA1 expression in SKOV3 (A), Hela (B), mouse brain (C), rat brain (D) whole cell lysates. (Predicted band size: 24; 41; 42 kD; Observed band size: 44 kD)



Immunohistochemical analysis of TIA1 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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