

## **Product Data Sheet**

## Anti-RING1b Antibody

Catalog #	Source	Reactivity	Applications	
CQA1238	Rabbit	H, M, R	WB, IH, IF/IC, IP, ChIP	
Description		Rabbit polyclonal antibody to	RING1b	
Immunogen		Recombinant full length prote	in of human RING1b	
Purification		The antibody was purified by	immunogen affinity chromatography.	
Specificity		Recognizes endogenous level	s of RING1b protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
		and 0.01% sodium azide.		
Dilution	,	WB (1/500 - 1/2000), IH (1/50 -	1/200), IF/IC (1/50 - 1/200), IP (1/50 - 1/200), ChIP (1/20 -	
		1/100)		
Gene Symbol		RNF2		
Alternative Names		BAP1; DING; HIPI3; RING1B; E3 ubiquitin-protein ligase RING2;		
		Huntingtin-interacting proteir	2-interacting protein 3; HIP2-interacting protein 3;	
		Protein DinG; RING finger pro	tein 1B; RING1b; RING finger protein 2; RING finger	
		protein BAP-1		
Entrez Gene		6045 (Human); 19821 (Mouse	e); 304850 (Rat)	
SwissProt		Q99496 (Human); Q9CQJ4 (N	ouse); Q4KLY4 (Rat)	
Storage/Stabi	lity	Shipped at 4°C. Upon delivery	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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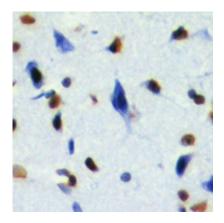
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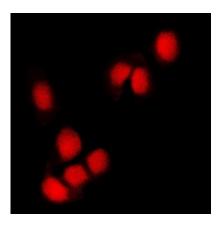
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Western blot analysis of RING1b expression in Jurkat (A), SHSY5Y (B), mouse testis (C) whole cell lysates. (Predicted band size: 37 kD; Observed band size: 30 kD)



Immunohistochemical analysis of RING1b staining in mouse brain 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 RING1b staining in U2OS 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 hidified 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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