

# **Product Data Sheet**

### **Anti-BRCA1** Antibody

Catalog #	Source	Reactivity	Applications		
CPA1107	Rabbit	H, M, R	WB, IH		
Description	Rabb	Rabbit polyclonal antibody to BRCA1			
Immunogen	KLH-	conjugated synthetic p	eptide encompassing a sequence within the center		
	regio	on of human BRCA1. Th	e exact sequence is proprietary.		
Purification	The a	antibody was purified b	y immunogen affinity chromatography.		
Specificity	Reco	gnizes endogenous lev	els of BRCA1 protein.		
Clonality	Polyc	clonal			
Conjugation					
Form	Liqui	d in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB (	1/500 - 1/1000), IH (1/1	00 - 1/200)		
Gene Symbol	BRCA	1			
Alternative Na	ames RNF5	3; Breast cancer type	L susceptibility protein; RING finger protein 53		
Entrez Gene 672 (Hu		Human); 12189 (Mous	ıman); 12189 (Mouse); 497672 (Rat)		
SwissProt	P383	98 (Human); P48754 (	Mouse); O54952 (Rat)		
Storage/Stabi	lity Ship	oed at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	freez	e/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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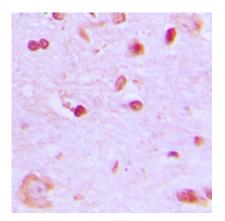
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Western blot analysis of BRCA1 expression in HeLa (A), mouse liver (B), rat liver (C) whole cell lysates. (Predicted band size: 7; 78-85; 202-210 kD; Observed band size: 210 kD)



Immunohistochemical analysis of BRCA1 staining in human 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.

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