

Product Data Sheet

Anti-MRE11 Antibody

Catalog #	Source	Reactivity	Applications		
CPA7174	Rabbit	H, Mk	WB, IH		
Description		Rabbit polyclonal antibody to MRE11			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
		region of human MRE11. Th	e exact sequence is proprietary.		
Purification		The antibody was purified by	v immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	ls of MRE11 protein.		
Clonality		Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/10	0 - 1/200)		
Gene Symbol		MRE11A			
Alternative Na	ames	HNGS1; MRE11; Double-stra	nd break repair protein MRE11A; Meiotic		
		recombination 11 homolog	l; MRE11 homolog 1; Meiotic recombination 11		
		homolog A; MRE11 homolog	A		
Entrez Gene		4361 (Human)			
SwissProt		P49959 (Human)			
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

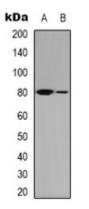
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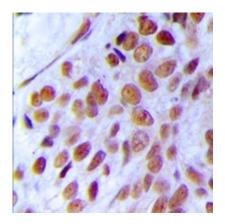


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Western blot analysis of MRE11 expression in K562 (A), Jurkat (B) whole cell lysates. (Predicted band size: 80 kD; Observed band size: 80 kD)



Immunohistochemical analysis of MRE11 staining in human breast 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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