

Product Data Sheet

Anti-PRMT2 Antibody

Catalog #	Source	Reactivity	Applications
CQA1281	Rabbit	H, M, R	WB, IH, IF/IC
Description	F	Rabbit polyclonal antibody to	D PRMT2
Immunogen	F	Recombinant full length prot	ein of human PRMT2
Purification	Т	The antibody was purified by	v immunogen affinity chromatography.
Specificity	F	Recognizes endogenous leve	ls of PRMT2 protein.
Clonality	P	Polyclonal	
Conjugation			
Form	L	iquid in 0.42% Potassium pl	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	а	and 0.01% sodium azide.	
Dilution	V	NB (1/500 - 1/2000), IH (1/5	0 - 1/200), IF/IC (1/50 - 1/100)
Gene Symbol	P	PRMT2	
Alternative Na	ames H	HMT1; HRMT1L1; Protein ar	ginine N-methyltransferase 2; Histone-arginine
	Ν	N-methyltransferase PRMT2	
Entrez Gene	3	3275 (Human)	
SwissProt	P	255345 (Human)	
Storage/Stabi	lity S	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid
	f	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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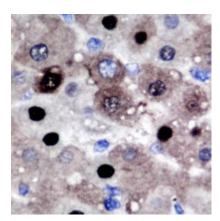
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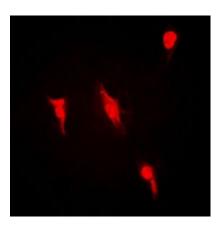
KDa <u>A</u> <u>B</u> <u>C</u> <u>D</u> <u>E</u> <u>F</u> <u>G</u> 130 95 72 55 43 34 26 17 For research purposes only, not for human use

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Western blot analysis of PRMT2 expression in MCF7 (A), Jurkat (B), HCT116 (C), mouse brain (D), mouse kidney (E), rat brain (F), rat kidney (G) whole cell lysates. (Predicted band size: 49 kD; Observed band size: 49 kD)



Immunohistochemical analysis of PRMT2 staining in human liver 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.



Immunofluorescent analysis of PRMT2 staining in Hela 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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