

## **Product Data Sheet**

### **Anti-DNMT3A Antibody**

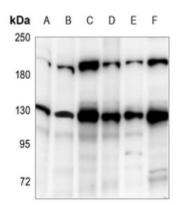
Catalog #	Source	Reactivity	Applications		
CPA3582	Rabbit	H, M, R	WB, IH		
Description		Rabbit polyclonal antibody to DNMT3A			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the N-term			
	I	region of human DNMT3A. T	he exact sequence is proprietary.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	ls of DNMT3A 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 (1/50	- 1/100)		
Gene Symbol		DNMT3A			
Alternative Na	ames	DNA (cytosine-5)-methyltrar	sferase 3A; Dnmt3a; DNA methyltransferase HsallIA;		
		DNA MTase HsallIA; M.Hsall	A		
Entrez Gene		1788 (Human); 13435 (Mouse); 444984 (Rat)			
SwissProt		Q9Y6K1 (Human); O88508 (Mouse); Q1LZ53 (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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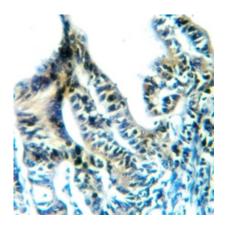




For research purposes only, not for human use

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Western blot analysis of DNMT3A expression in MEF (A), H9C2 (B), HCT116 (C), Jurkat (D), PC3 (E), HEK293T (F) whole cell lysates. (Predicted band size: 16; 81; 101 kD; Observed band size: 130 kD)



Immunohistochemical analysis of DNMT3A staining in human colorectal 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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