

## **Product Data Sheet**

## Anti-JMJD6 Antibody

Catalog #	Source	Reactivity	Applications		
-		-			
CQA1283	Rabbit	H, M, R	WB, IH, IF/IC, IP, ChIP		
Description	Ra	abbit polyclonal antibody t	D IMID6		
Immunogen	Re	ecombinant full length prot	ein of human JMJD6		
Purification	Tł	he antibody was purified by	immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous leve	ls of JMJD6 protein.		
Clonality	Pc	olyclonal			
Conjugation					
Form	Lie	quid in 0.42% Potassium pl	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	ar	nd 0.01% sodium azide.			
Dilution	W	/B (1/500 - 1/2000), IH (1/50	- 1/200), IF/IC (1/50 - 1/200), IP (1/20 - 1/50), ChIP (1/20 -		
	1/	(50)			
Gene Symbol	JN	MJD6			
Alternative Names		KIAA0585; PTDSR; Bifunctional arginine demethylase and lysyl-hydroxylase JMJD6;			
	Hi	istone arginine demethylas	e JMJD6; JmjC domain-containing protein 6; Jumonji		
	do	omain-containing protein 6	; Lysyl-hydroxylase JMJD6; Peptide-lysine 5-dioxygenase		
	JI	MJD6; Phosphatidylserine r	eceptor; Protein PTDSR		
Entrez Gene	23	3210 (Human); 107817 (Mo	ouse); 360665 (Rat)		
SwissProt	Q	6NYC1 (Human); Q9ERI5 (N	/louse); Q6AYK2 (Rat)		
Storage/Stabi	<b>lity</b> Sh	hipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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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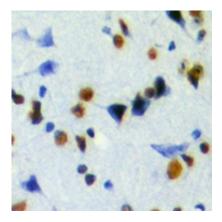
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# Cohesion

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Western blot analysis of JMJD6 expression in mouse heart (A), mouse liver (B) whole cell lysates. (Predicted band size: 43; 46; 47 kD; Observed band size: 46; 49 kD)



Immunohistochemical analysis of JMJD6 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.

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