

# **Product Data Sheet**

### Anti-Mms2 Antibody

Catalog #	Source	e Reactivity	Applications	
CQA1507	Rabbit	t H, M, R	WB, IH	
Description		Rabbit polyclonal antibody t	o Mms2	
Immunogen		Recombinant full length prot	ein of human Mms2	
Purification		The antibody was purified by	immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	ls of Mms2 protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium pl	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/2000), IH (1/50	- 1/200)	
Gene Symbol		UBE2V2		
Alternative Names		MMS2; UEV2; Ubiquitin-conjugating enzyme E2 variant 2; DDVit 1; Enterocyte		
		differentiation-associated fa	ctor 1; EDAF-1; Enterocyte differentiation-promoting	
		factor 1; EDPF-1; MMS2 hon	olog; Vitamin D3-inducible protein	
Entrez Gene		7336 (Human); 70620 (Mous	e); 287927 (Rat)	
SwissProt		Q15819 (Human); Q9D2M8	Mouse); Q7M767 (Rat)	
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y 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

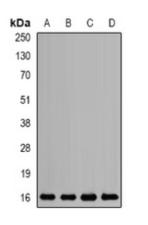
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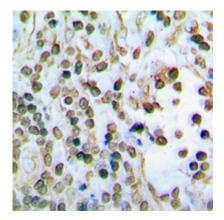


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Western blot analysis of Mms2 expression in SKOV3 (A), HL60 (B), mouse liver (C), rat brain (D) whole cell lysates. (Predicted band size: 16 kD; Observed band size: 16 kD)



Immunohistochemical analysis of Mms2 staining in rat spleen 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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