

## Anti-USP17L24 Antibody

Catalog #	Source	Reactivity	Applications
CQA9899	Rabbit	H	WB, FC
<b>Description</b>	Rabbit polyclonal antibody to USP17L24		
<b>Immunogen</b>	KLH-conjugated synthetic peptide encompassing a sequence within the C-terminal region of human USP17L24. The exact sequence is proprietary.		
<b>Purification</b>	The antibody was purified by immunogen affinity chromatography.		
<b>Specificity</b>	Recognizes endogenous levels of USP17L24 protein.		
<b>Clonality</b>	Polyclonal		
<b>Conjugation</b>			
<b>Form</b>	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.		
<b>Dilution</b>	WB (1/500 - 1/1000), FC (1/10 - 1/50)		
<b>Gene Symbol</b>	USP17L24; USP17L25; USP17L26; USP17L27; USP17L28; USP17L29; USP17L30		
<b>Alternative Names</b>	USP17; USP17H; USP17I; USP17J; USP17K; USP17L; USP17M; Ubiquitin carboxyl-terminal hydrolase 17-like protein 24; Deubiquitinating enzyme 17; Ubiquitin thioesterase 17; Ubiquitin-specific-processing protease 17		
<b>Entrez Gene</b>	7.28369728373728E+41 (Human)		
<b>SwissProt</b>	Q0WX57 (Human)		
<b>Storage/Stability</b>	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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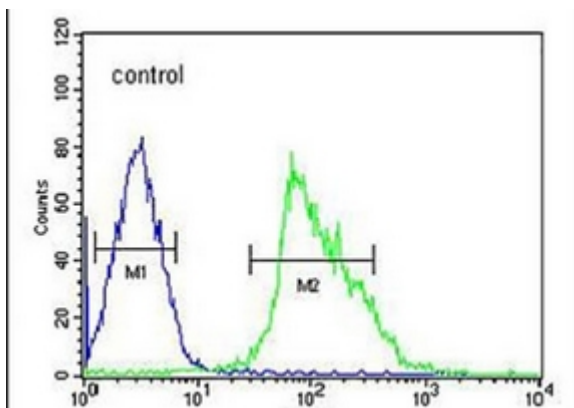
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# Product Data Sheet



Western blot analysis of USP17L24 expression in MADMB231 (A) whole cell lysates. (Predicted band size: 59 kD; Observed band size: 60 kD)



Flow cytometric analysis of HepG2 cells using Anti-USP17L24 Antibody. The cells were fixed with 2% paraformaldehyde (10 min) and then permeabilized with 90% methanol for 10 min. The cells were incubated in 2% bovine serum albumin to block non-specific protein-protein interactions followed by the antibody at 37 °C for 60 min. The secondary antibody Goat Anti-Rabbit IgG (H&L) - AF488 was incubated at 37 °C for 40 min. Isotype control antibody (blue line) was used under the same condition.

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