

Product Data Sheet

Recombinant Anti-MSH6 Rabbit mAb [KO/KD Validated]

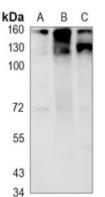
Catalog #	Source	Reactivity	Applications		
CMA1009	Rabbit	Н	WB, IH		
Description		Recombinant rabbit mono	clonal antibody to MSH6		
Immunogen		Recombinant fusion protei	n of human MSH6. The exact sequence is proprietary.		
Purification The antibody was purified by immunoge			by immunogen affinity chromatography.		
Specificity		Recognizes endogenous levels of MSH6 protein.			
Clonality		Monoclonal			
Conjugation					
Form	Form Liquid in PBS, pH 7.3, 50% glycerol, and 0.05% Proclin300.				
Dilution WB (1/500 - 1/1000), IH (1/100 - 1		WB (1/500 - 1/1000), IH (1/1	.00 - 1/200)		
Gene Symbol		MSH6			
Alternative Na	ames	GTBP; DNA mismatch repa	ir protein Msh6; hMSH6; G/T mismatch-binding protein;		
		GTBP; GTMBP; MutS-alpha	160 kDa subunit; p160		
Entrez Gene		2956 (Human)			
SwissProt P52701 (Human)					
Storage/Stabi	lity	Shipped at 4°C. Upon deliv	ery 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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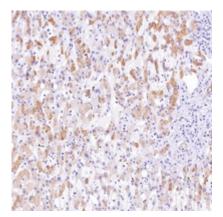
Coherion



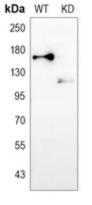
For research purposes only, not for human use

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Western blot analysis of MSH6 expression in HepG2 (A), PC3 (B), MCF7 (C) whole cell lysates. (Predicted band size: 152 kD; Observed band size: 160 kD)



Immunohistochemical analysis of MSH6 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.



Western blot analysis of MSH6 expression in wild type (WT) and knockdown (KD) HT1080 cell lysates.

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