## MSH6 (D60G2) XP® Rabbit mAb



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## For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W, IF-IC	<b>Reactivity:</b> H Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 160	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P52701	Entrez-Gene Id: 2956
Product Usage Information		<b>Application</b> Western Blotting Immunofluorescence	(Immunocytochem	nistry)		<b>Dilution</b> 1:1000 1:50
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		MSH6 (D60G2) ${\rm XP}^{\rm @}$ Rabbit mAb detects endogenous levels of total MSH6 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human MSH6.				
Background		The DNA mismatch repair system (MMR) repairs post-replication DNA, inhibits recombination between nonidentical DNA sequences, and induces both checkpoint and apoptotic responses following certain types of DNA damage (1). MSH2 (MutS homologue 2) forms the hMutS-α dimer with MSH6 and is an essential component of the mismatch repair process. hMutS-α is part of the BRCA1-associated surveillance complex (BASC), a complex that also contains BRCA1, MLH1, ATM, BLM, PMS2 proteins, and the Rad50-Mre11-NBS1 complex (2). Mutations in MSH6 and other MMR proteins have been found in a large proportion of hereditary nonpolyposis colorectal cancer (Lynch Syndrome), the most common form of inherited colorectal cancer in the Western world (3). Mutations in MSH6 have been shown to occur in glioblastoma in response to temozolomide therapy and to promote temozolomide resistance (4).				
Background References		1. O'Brien, V. and Brown, R. (2006) <i>Carcinogenesis</i> 27, 682-92. 2. Wang, Y. et al. (2000) <i>Genes Dev</i> 14, 927-39. 3. Plotz, G. et al. (2006) <i>J Mol Histol</i> 37, 271-83. 4. Yip, S. et al. (2009) <i>Clin Cancer Res</i> 15, 4622-9.				
Species Reactivity		Species reactivity is determined by testing in at least one approved application (e.g., western blot).				
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				
Applications Key		W: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)				
Cross-Reactivity Key		H: Human Mk: Monkey				
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