

MSH6 Antibody

Purified Mouse Monoclonal Antibody Catalog # AO1649a

Product Information

Application WB, IHC, FC, ICC, E

Primary Accession P52701
Reactivity Human
Host Mouse
Clonality Monoclonal

Clone Names3E1IsotypeIgG2aCalculated MW152786

Description This gene encodes a protein similar to the MutS protein. In E. coli, the MutS

protein helps in the recognition of mismatched nucleotides, prior to their repair. A highly conserved region of approximately 150 aa, called the Walker-A adenine nucleotide binding motif, exists in MutS homologs. The encoded protein of this gene combines with MSH2 to form a mismatch recognition complex that functions as a bidirectional molecular switch that exchanges ADP and ATP as DNA mismatches are bound and dissociated. Mutations in this gene have been identified in individuals with hereditary nonpolyposis

colon cancer (HNPCC) and endometrial cancer.

Immunogen Purified recombinant fragment of human MSH6 expressed in E. Coli.

Formulation Ascitic fluid containing 0.03% sodium azide.

Additional Information

Gene ID 2956

Other Names DNA mismatch repair protein Msh6, hMSH6, G/T mismatch-binding protein,

GTBP, GTMBP, MutS-alpha 160 kDa subunit, p160, MSH6, GTBP

Dilution WB~~1/500 - 1/2000 IHC~~1/200 - 1/1000 FC~~1/200 - 1/400 ICC~~N/A

E~~1/10000

Storage Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions MSH6 Antibody is for research use only and not for use in diagnostic or

therapeutic procedures.

Protein Information

Name MSH6 (HGNC:7329)

Synonyms GTBP

Function Component of the post-replicative DNA mismatch repair system (MMR).

Heterodimerizes with MSH2 to form MutS alpha, which binds to DNA mismatches thereby initiating DNA repair. When bound, MutS alpha bends the DNA helix and shields approximately 20 base pairs, and recognizes single base mismatches and dinucleotide insertion-deletion loops (IDL) in the DNA. After mismatch binding, forms a ternary complex with the MutL alpha heterodimer, which is thought to be responsible for directing the downstream MMR events, including strand discrimination, excision, and resynthesis. ATP binding and hydrolysis play a pivotal role in mismatch repair functions. The ATPase activity associated with MutS alpha regulates binding similar to a molecular switch: mismatched DNA provokes ADP-->ATP exchange, resulting in a discernible conformational transition that converts MutS alpha into a sliding clamp capable of hydrolysis-independent diffusion along the DNA backbone. This transition is crucial for mismatch repair. MutS alpha may also play a role in DNA homologous recombination repair. Recruited on chromatin in G1 and early S phase via its PWWP domain that specifically binds trimethylated 'Lys-36' of histone H3 (H3K36me3): early recruitment to

chromatin to be replicated allowing a quick identification of mismatch repair

to initiate the DNA mismatch repair reaction.

Cellular Location Nucleus, Chromosome, Note=Associates with H3K36me3 via its PWWP

domain

References

1. Hered Cancer Clin Pract. 2009 Dec 23;7(1):17. 2. Cancer Epidemiol Biomarkers Prev. 2009 Sep;18(9):2460-7.

Images

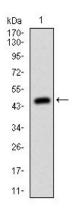
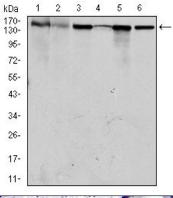


Figure 1: Western blot analysis using MSH6 mAb against human MSH6 (AA: 217-395) recombinant protein. (Expected MW is 45.5 kDa)

Figure 2: Western blot analysis using MSH6 mouse mAb against HEK293 (1), HCT116 (2), A549 (3), A431 (4), MCF-7 (5) and HepG2 (6) cell lysate.



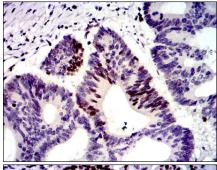


Figure 3: Immunohistochemical analysis of paraffin-embedded colon cancer tissues using MSH6 mouse mAb with DAB staining.

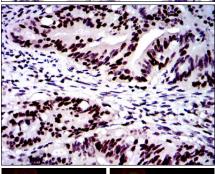


Figure 4: Immunohistochemical analysis of paraffin-embedded rectum cancer tissues using MSH6 mouse mAb with DAB staining.

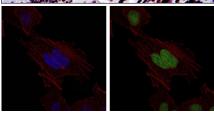


Figure 5: Immunofluorescence analysis of Hela cells using MSH6 mouse mAb (green). Blue: DRAQ5 fluorescent DNA dye. Red: Actin filaments have been labeled with Alexa Fluor-555 phalloidin.

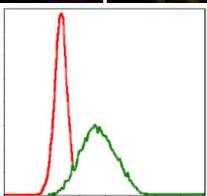


Figure 6: Flow cytometric analysis of MCF-7 cells using MSH6 mouse mAb (green) and negative control (red).

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