



MSH6 Polyclonal Antibody

E90983

Catalog Number: E90983**Amount:** 100ul

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).

Species: Rabbit**Isotype:** IgG

Storage/Stability: Store at -20oC or -80oC. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

Synonyms: MSH6;GTBP;HNPPC5;HSAP ;**Immunogen:** Recombinant protein of human MSH6**Purification:** Affinity purification**Reactivity:** H M R**Applications:** WB IHC**Molecular Weight:** 153kDa**Swiss-Prot No. :** P52701**Gene ID:** 2956

References: 1. O'Brien, V. and Brown, R. (2006) Carcinogenesis 27, 682-92. 2. Wang, Y. et al. (2000) Genes Dev 14, 927-39. 3. Plotz, G. et al. (2006) J Mol Histol 37, 271-83. 4. Yip, S. et al. (2009) Clin Cancer Res 15, 4622-9.

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