For Research Use Only

MLH1 Monoclonal antibody, PBS Only

Catalog Number: 60822-2-PBS



Purification Method:

Protein G purification

CloneNo.:

2D5A2

Basic Information

Catalog Number: 60822-2-PBS

Nanodrop:

Isotype:

lgG1

GenBank Accession Number:

BC006850

GeneID (NCBI):

UNIPROT ID:

P40692

Mouse Full Name:

100ug, Concentration: 1 mg/ml by

mutL homolog 1, colon cancer, nonpolyposis type 2 (E. coli)

Immunogen Catalog Number: Calculated MW: AG27723 756 aa. 85 kDa

Applications

Tested Applications:

IHC, Indirect ELISA

Species Specificity:

human

Background Information

MLH1, also named as COCA2, belongs to the DNA mismatch repair mutL/hexB family. It heterodimerizes with PMS2 to form MutL alpha which is a component of the post-replicative DNA mismatch repair system (MMR). MutL alpha (MLH1-PMS2) interacts physically with the clamp loader subunits of DNA polymerase III, suggesting that it may play a role to recruit the DNA polymerase III to the site of the MMR. MLH1 also implicated in DNA damage signaling, a process which induces cell cycle arrest and can lead to apoptosis in case of major DNA damages. MLH1 heterodimerizes with MLH3 to form MutL gamma which plays a role in meiosis. (PMID: 16873062, PMID: 18206974) Defects in MLH1 are the cause of hereditary non-polyposis colorectal cancer type 2 (HNPCC2). Defects in MLH1 are a cause of mismatch repair cancer syndrome (MMRCS). Defects in MLH1 are a cause of Muir-Torre syndrome (MTS). Defects in MLH1 are a cause of susceptibility to endometrial cancer.

Storage

Storage:

Store at -80°C.

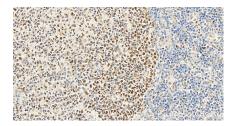
Storage Buffer:

PBS Only

in USA), or 1(312) 455-8498 (outside USA)

E: proteintech@ptglab.com W: ptglab.com

Selected Validation Data



Immunohistochemical analysis of paraffinembedded Human tonsillitis tissue slide using 60822-2-1g (MLH1 antibody) at dilution of 1:2000 (under 40x lens). This data was developed using the same antibody clone with 60822-2-PBS in a different storage buffer formulation.