For Research Use Only

MLH1 Polyclonal antibody

Catalog Number: 11697-1-AP

Featured Product

23 Publications

mutL homolog 1, colon cancer,

nonpolyposis type 2 (E. coli)

Calculated MW: 756 aa, 85 kDa



Basic Information

Catalog Number: 11697-1-AP

r: GenBank Accession Number: BC006850

GeneID (NCBI):

150ul , Concentration: 700 μ g/ml by 4292

Nanodrop and 453 $\mu g/ml$ by Bradford Full Name:

method using BSA as the standard;

Source: Rabbit Isotype:

IgG Observed MW:

Immunogen Catalog Number: 85 kDa

AG2319

Size

Purification Method:

Antigen affinity purification Recommended Dilutions:

WB 1:500-1:1000

IP 0.5-4.0 ug for IP and 1:500-1:1000

for WB IF 1:50-1:500

Applications

Tested Applications:

IF, IP, WB,ELISA

Cited Applications: ColP. IF. IHC. WB

Species Specificity:

human, mouse, rat

Cited Species:

human, mouse, zebrafish

Positive Controls:

WB: A431 cells, HEK-293 cells, HeLa cells, Jurkat cells,

human testis tissue

IP: HeLa cells,

IF: HeLa cells,

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. Western blot analysis with an MLH1 antibody detected a 85-100 kDa band. Full-length human MLH1 is specifically cleaved into degradation products of 40-45 kDa by caspase-3 (PMID: 15087450, PMID: 19603033). This antibody is specific to MLH1.

Notable Publications

Author	Pubmed ID	Journal	Application
Xuting Ran	35664732	Front Oncol	IHC
Jun Zhu	32396667	J Surg Oncol	IHC
Dazhang Bai	33949657	Hum Mol Genet	WB

Storage

Storage:

Store at -20°C. Stable for one year after shipment.

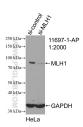
Storage Buffe

PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

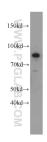
Aliquoting is unnecessary for -20°C storage

*** 20ul sizes contain 0.1% BSA

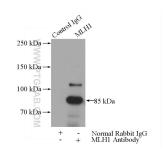
Selected Validation Data



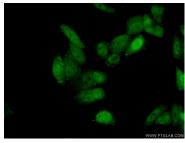
WB result of MLH1 antibody (11697-1-AP; 1:2000; incubated at room temperature for 1.5 hours) with sh-Control and sh-MLH1 transfected HeLa cells.



A431 cells were subjected to SDS PAGE followed by western blot with 11697-1-AP (MLH1 antibody) at dilution of 1:600 incubated at room temperature for 1.5 hours.



IP Result of anti-MLH1 (IP:11697-1-AP, 3ug; Detection:11697-1-AP 1:500) with HeLa cells lysate 3200ug.



Immunofluorescent analysis of (10% Formaldehyde) fixed HeLa cells using 11697-1-AP (MLH1 antibody) at dilution of 1:50 and Alexa Fluor 488-conjugated AffiniPure Goat Anti-Rabbit IgG(H+L).