

Anti-MUTYH Antibody

Rabbit polyclonal antibody to MUTYH

Catalog # AP60033

Product Information

Application	WB, IF/IC, IHC
Primary Accession	Q9UIF7
Other Accession	Q99P21
Reactivity	Human, Mouse, Rat, Monkey
Host	Rabbit
Clonality	Polyclonal
Calculated MW	60069

Additional Information

Gene ID	4595
Other Names	MYH; A/G-specific adenine DNA glycosylase; MutY homolog; hMYH
Target/Specificity	KLH-conjugated synthetic peptide encompassing a sequence within the center region of human MUTYH. The exact sequence is proprietary.
Dilution	WB~~WB (1/500 - 1/1000), IHC (1/100 - 1/200), IF/IC (1/100 - 1/500) IF/IC~~N/A IHC~~WB (1/500 - 1/1000), IHC (1/100 - 1/200), IF/IC (1/100 - 1/500)
Format	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.09% (W/V) sodium azide.
Storage	Store at -20 °C.Stable for 12 months from date of receipt

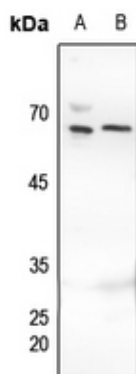
Protein Information

Name	MUTYH
Synonyms	MYH
Function	Involved in oxidative DNA damage repair. Initiates repair of A*oxoG to C*G by removing the inappropriately paired adenine base from the DNA backbone. Possesses both adenine and 2-OH-A DNA glycosylase activities.
Cellular Location	Nucleus. Mitochondrion {ECO:0000250 UniProtKB:Q99P21}

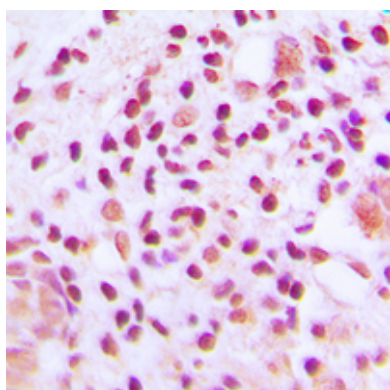
Background

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human MUTYH. The exact sequence is proprietary.

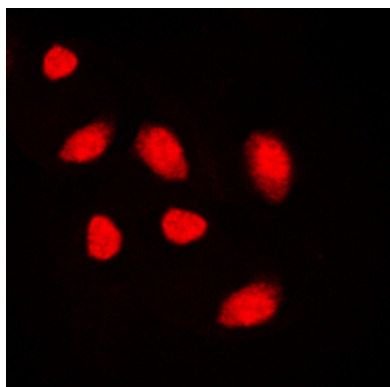
Images



Western blot analysis of MUTYH expression in HeLa (A), LOVO (B) whole cell lysates.



Immunohistochemical analysis of MUTYH staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of MUTYH staining in K562 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.