

Product datasheet for AP52779PU-N

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OriGene Technologies, Inc.

MYH (MUTYH) (Center) Rabbit Polyclonal Antibody

Product data:

Product Type: Primary Antibodies

Applications: FC, IF, WB

Recommended Dilution: Flow cytometry: 1/10-1/50.

Immunofluorescence: 1/10-1/50.

Western blot: 1/50-1/100. Enzyme immunoassay: 1/1000.

Reactivity: Human, Mouse

Host: Rabbit

Isotype: lg

Clonality: Polyclonal

Immunogen: Synthetic peptide - KLH conjugated - corresponding to the central region (between 352-

381aa) of human MUTYH.

Specificity: This antibody recognizes MUTYH.

Formulation: PBS with 0.09% (W/V) Sodium azide

State: Purified

State: Liquid purified Ig fraction

Concentration: lot specific

Purification: Prepared by Saturated Ammonium Sulfate (SAS) precipitation followed by dialysis against PBS

Conjugation: Unconjugated

Storage: Store undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.

Gene Name: mutY DNA glycosylase

Database Link: Entrez Gene 70603 MouseEntrez Gene 4595 Human

Q9UIF7

MYH (MUTYH) (Center) Rabbit Polyclonal Antibody - AP52779PU-N

Background: MYH is the eukaryotic homolog to the E. coli protein MutY. It was first identified in HeLa cells

and nicks DNA replication errors, specifically A/G mismatches. MYH is involved in oxidative DNA damage repair and initiates repair of A*oxoG to C*G by removing the inappropriately paired adenine base from the DNA backbone. It possesses both adenine and 2-OH-A DNA

glycosylase activities.

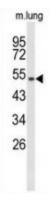
Synonyms: MYH, MutY homolog

Note: Molecular Weight: 60069 Da

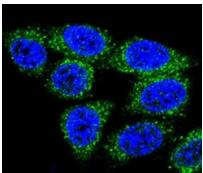
Protein Families: Druggable Genome, Stem cell - Pluripotency

Protein Pathways: Base excision repair

Product images:

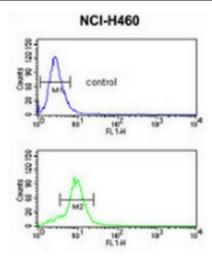


Western blot analysis of MUTYH (arrow) in mouse lung tissue lysates (35ug/lane) using MUTYH antibody.



Confocal immunofluorescent analysis of Hela cells using MUTYH antibody., followed by Alexa Fluor® 488-conjugated goat anti-rabbit IgG (green). DAPI was used to stain the cell nuclear (blue).





Flow cytometric analysis of NCI-H460 cells (bottom histogram) compared to a negative control cell (top histogram) using MUTYH antibody., followed by FITC-conjugated goat-antirabbit secondary antibodies.