GTBP (F-1): sc-271979



The Power to Question

BACKGROUND

The finding that mutations in DNA mismatch repair genes are associated with hereditary nonpolyposis colorectal cancer (HNPCC) has resulted in considerable interest in the understanding of the mechanism of DNA mismatch repair. Initially, inherited mutations in the MSH2 and MLH1 homologs of the bacterial DNA mismatch repair genes MutS and MutL were demonstrated at high frequency in HNPCC and were shown to be associated with microsatellite instability. A member of the mismatch repair family, GTBP (also designated MSH6), is a MSH2-related protein that binds to DNA containing G/T mismatches. Findings suggest that the mismatch-binding factor in human cells is composed of a heterodimer of GTBP and MSH2.

REFERENCES

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- 2. Palombo, F., et al. 1994. Mismatch repair and cancer. Nature 367: 417-418.
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- Papadopoulos, N., et al. 1994. Mutation of a MutL homolog in hereditary colon cancer. Science 263: 1625-1629.
- 5. Nicolaides, N.C., et al. 1994. Mutations of two PMS homologues in hereditary nonpolyposis colon cancer. Nature 371: 75-80.
- Prolla, T.A., et al. 1994. MLH1, Pms1, and Msh2 interactions during the initiation of DNA mismatch repair in yeast. Science 265: 1091-1092.
- 7. Palombo, F., et al. 1995. GTBP, a 160 kDa protein essential for mismatch-binding activity in human cells. Science 268: 1912-1914.
- 8. Shiwaku, H.O., et al. 1997. Alternative splicing of GTBP in normal human tissues. DNA Res. 4: 359-362.
- Ercoli, A., et al. 1999. hMSH2 and GTBP expression in advanced stage epithelial ovarian cancer. Br. J. Cancer 80: 1665-1671.

CHROMOSOMAL LOCATION

Genetic locus: MSH6 (human) mapping to 2p16.3.

SOURCE

GTBP (F-1) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 63-97 near the N-terminus of GTBP of human origin.

PRODUCT

Each vial contains 200 $\mu g \ lgG_1$ lambda light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-271979 P, (100 μg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

APPLICATIONS

GTBP (F-1) is recommended for detection of GTBP of human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000)

Suitable for use as control antibody for GTBP siRNA (h): sc-35528, GTBP shRNA Plasmid (h): sc-35528-SH and GTBP shRNA (h) Lentiviral Particles: sc-35528-V.

Molecular Weight of GTBP: 160 kDa.

Positive Controls: HeLa nuclear extract: sc-2120, K-562 whole cell lysate: sc-2203 or SW480 cell lysate: sc-2219.

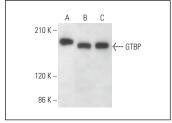
RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended:

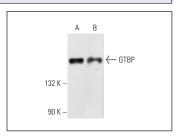
1) Western Blotting: use m-lgGλ BP-HRP: sc-516132 or m-lgGλ BP-HRP (Cruz Marker): sc-516132-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml).

3) Immunofluorescence: use m-lgGλ BP-FITC: sc-516185 or m-lgGλ BP-PE: sc-516186 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

DATA







GTBP (F-1): sc-271979. Western blot analysis of GTBP expression in HeLa (**A**) and A-431 (**B**) nuclear extracts.

SELECT PRODUCT CITATIONS

 Yang, M., et al. 2022. KDM6B promotes PARthanatos via suppression of O⁶-methylguanine DNA methyltransferase repair and sustained checkpoint response. Nucleic Acids Res. 50: 6313-6331.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

For research use only, not for use in diagnostic procedures.