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10-7010: Monoclonal Antibody to DNMT2 (Clone: ABM1H70)

Clonality: Monoclonal **Clone Name:** ABM1H70 Application: FACS.WB Reactivity: Mouse, Human Gene: TRDMT1 Gene ID: 1787 **Uniprot ID:** 014717 Format:

DNA (cytosine-5)-methyltransferase-like protein 2, DNA methyltransferase homolog HsallP, **Alternative Name:**

DNMT2

Purified

Isotype: Mouse IgG1, Kappa

A partial length recombinant protein from DNMT2 was used as the immunogen for this Immunogen Information:

antibody.

Description

DNMT2, also known as tRNA-aspartic acid methyltransferase 1 (TRDMT1), is highly conserved cytosine-C5 methyltransferase that introduces the C38 methylation of tRNAAsp in many species, including lower eukaryotes, plants, insects, and humans. DNMT2 contains all 10 sequence motifs that are conserved among m5C MTases, including the consensus S-adenosyl-Lmethionine-binding motifs and the active site ProCys dipeptide. It is a relatively small protein of 391 amino acids and lacks the large N-terminal domains present in the DNMT1 and DNMT3 families. DNMT2 is ubiquitously expressed with multiple mRNA species in most human and mouse adult tissues. It is variably expressed in human cancer cell lines.

Product Info

Amount: 25 μg / 100 μg

Purification: Protein G Chromatography

25 μg in 50 μl/100 μg in 200 μl PBS containing 0.05% BSA and 0.05% sodium azide. Sodium Content:

azide is highly toxic.

Store the antibody at 4°C, stable for 6 months. For long-term storage, store at -20°C. Avoid Storage condition:

repeated freeze and thaw cycles.

Application Note

WB: 4-6 μg/ml, FACS: 0.5-1 μg/10⁶

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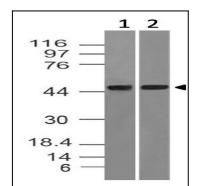


Fig-1: Western blot analysis of DNMT2. Anti-DNMT2 antibody (Clone: ABM1H70) was used at 4 μ g/ml on Jurkat and mouse embryonic liver lysates.

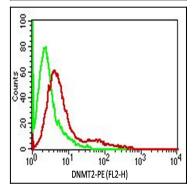


Fig-2: Intracellular Flow analysis of DNMT2 antibody in Hela cells using 0.5 μ g/ 10^6 cells of anti-DNMT2 antibody (ABM1H70). Green represents isotype control; red represents anti-DNMT 2 antibody. Goat anti-mouse PE conjugate was used as secondary antibody. (Cells were fixed with 4% paraformaldehyde for 10 min and washed with PBS by centrifuging at 1100 for 5 min followed by permeabilization for 20 min and washed again as mentioned above. Then cell were incubated with primary antibody for 45 min. and after washing the cells twice in PBS, incubated with conjugated secondary antibody for 30 min. Data acquisition was done after washing twice with PBS as mentioned above).