

A Reliable Research Partner in Life Science and Medicine

Elabscience®

DNMT1 Polyclonal Antibody

catalog number: E-AB-66957

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description

Reactivity Human; Mouse; Rat

Immunogen A synthetic peptide of human DNMT1.

Host Rabbit Isotype IgG

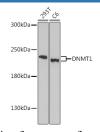
Purification Affinity purification

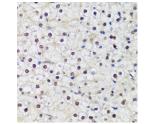
Buffer Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer and 50% glycerol.

Recommended Dilution Applications

WB 1:500-1:2000 IHC 1:50-1:200

Data

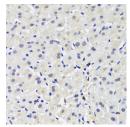




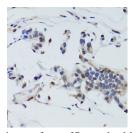
Western blot analysis of extracts of various cell lines using DNMT1 Polyclonal Antibody at dilution of 1:1000.

Observed-MV:230 kDa

Calculated-MV:144 kDa/183 kDa/184 kDa



Immunohistochemistry of paraffin-embedded Human liver using DNMT1 Polyclonal Antibody at dilution of 1:200 (40x lens).



Immunohistochemistry of paraffin-embedded Human liver Immunohistochemistry of paraffin-embedded Human gastric cancer using DNMT1 Polyclonal Antibody at dilution of 1:200 (40x lens).

cancer using DNMT1 Polyclonal Antibody at dilution of 1:200 (40x lens).

Preparation & Storage

Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles. Storage

Shipping The product is shipped with ice pack, upon receipt, store it immediately at the

temperature recommended.

Background

For Research Use Only

Toll-free: 1-888-852-8623 Web:www.elabscience.com

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Elabscience[®]

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This gene encodes an enzyme that transfers methyl groups to cytosine nucleotides of genomic DNA. This protein is the major enzyme responsible for maintaining methylation patterns following DNA replication and shows a preference for hemi-methylated DNA. Methylation of DNA is an important component of mammalian epigenetic gene regulation. Aberrant methylation patterns are found in human tumors and associated with developmental abnormalities. Variation in this gene has been associated with cerebellar ataxia, deafness, and narcolepsy, and neuropathy, hereditary sensory, type IE. Alternative splicing results in multiple transcript variants.

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