

## DNMT1 Polyclonal Antibody

catalog number: E-AB-93288

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

### Description

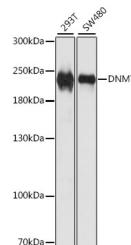
<b>Reactivity</b>	Human
<b>Immunogen</b>	Recombinant fusion protein of human DNMT1
<b>Host</b>	Rabbit
<b>Isotype</b>	IgG
<b>Purification</b>	Affinity purification
<b>Buffer</b>	Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer and 50% glycerol.

### Applications

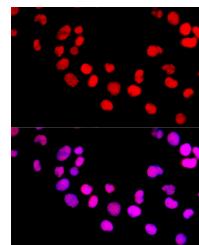
### Recommended Dilution

<b>WB</b>	1:500-1:2000
<b>IF</b>	1:50-1:200
<b>IP</b>	1:50-1:200

### Data



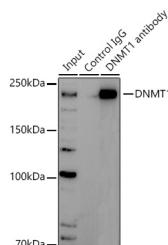
Western blot analysis of extracts of various cell lines using DNMT1 Polyclonal Antibody at 1:1000 dilution.



Immunofluorescence analysis of 293T cells using DNMT1 Polyclonal Antibody at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.

#### Observed-MW:Refer to figures

**Calculated-MW:144 kDa/183 kDa/184 kDa**



Immunoprecipitation analysis of 300ug extracts of Jurkat cells using 3ug DNMT1 Polyclonal Antibody. Western blot was performed from the immunoprecipitate using DNMT1 at a dilution of 1:1000.

#### Observed-MW:Refer to figures

**Calculated-MW:144 kDa/183 kDa/184 kDa**

### Preparation & Storage

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

**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](http://www.elabscience.com)

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Rev. V1.7

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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