

DNMT3A Antibody (C-term V897) Blocking Peptide Synthetic peptide Catalog # BP16264b

### **Specification**

## DNMT3A Antibody (C-term V897) Blocking Peptide - Product Information

Primary Accession

### <u>Q9Y6K1</u>

## DNMT3A Antibody (C-term V897) Blocking Peptide - Additional Information

Gene ID 1788

**Other Names** 

DNA (cytosine-5)-methyltransferase 3A, Dnmt3a, DNA methyltransferase HsallIA, DNA MTase HsallIA, MHsallIA, DNMT3A

Format

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

**Precautions** This product is for research use only. Not for use in diagnostic or therapeutic procedures.

# DNMT3A Antibody (C-term V897) Blocking Peptide - Protein Information

Name DNMT3A

Function

Required for genome-wide de novo methylation and is essential for the establishment of DNA methylation patterns during development (PubMed: <a href="http://www.uniprot.org/citations/12138111" target=" blank">12138111</a>, PubMed:<a href="http://www.uniprot.org/citations/16357870" target=" blank">16357870</a>, PubMed:<a href="http://www.uniprot.org/citations/30478443" target="blank">30478443</a>). DNA methylation is coordinated with methylation of histones (PubMed: <a href="http://www.uniprot.org/citations/12138111" target="\_blank">12138111</a>, PubMed:<a href="http://www.uniprot.org/citations/16357870" target="\_blank">16357870</a>, PubMed:<a href="http://www.uniprot.org/citations/30478443" target=" blank">30478443</a>). It modifies DNA in a non-processive manner and also methylates non-CpG sites (PubMed:<a href="http://www.uniprot.org/citations/12138111" target=" blank">12138111</a>, PubMed:<a href="http://www.uniprot.org/citations/16357870" target=" blank">16357870</a>, PubMed:<a href="http://www.uniprot.org/citations/30478443" target=" blank">30478443</a>). May preferentially methylate DNA linker between 2 nucleosomal cores and is inhibited by histone H1 (By similarity). Plays a role in paternal and maternal imprinting (By similarity). Required for methylation of most imprinted loci in germ cells (By similarity). Acts as a transcriptional corepressor for ZBTB18 (By similarity). Recruited to trimethylated 'Lys-36' of histone H3



(H3K36me3) sites (By similarity). Can actively repress transcription through the recruitment of HDAC activity (By similarity). Also has weak auto-methylation activity on Cys-710 in absence of DNA (By similarity).

#### **Cellular Location**

Nucleus. Chromosome Cytoplasm. Note=Accumulates in the major satellite repeats at pericentric heterochromatin {ECO:0000250|UniProtKB:088508}

#### **Tissue Location**

Highly expressed in fetal tissues, skeletal muscle, heart, peripheral blood mononuclear cells, kidney, and at lower levels in placenta, brain, liver, colon, spleen, small intestine and lung

## DNMT3A Antibody (C-term V897) Blocking Peptide - Protocols

Provided below are standard protocols that you may find useful for product applications.

#### <u>Blocking Peptides</u>

DNMT3A Antibody (C-term V897) Blocking Peptide - Images

## DNMT3A Antibody (C-term V897) Blocking Peptide - Background

CpG methylation is an epigenetic modification that isimportant for embryonic development, imprinting, and X-chromosomeinactivation. Studies in mice have demonstrated that DNAmethylation is required for mammalian development. This geneencodes a DNA methyltransferase that is thought to function in denovo methylation, rather than maintenance methylation. The proteinlocalizes to the cytoplasm and nucleus and its expression isdevelopmentally regulated. Alternative splicing results in multipletranscript variants encoding different isoforms. [provided byRefSeq].

### DNMT3A Antibody (C-term V897) Blocking Peptide - References

Holz-Schietinger, C., et al. J. Biol. Chem. 285(38):29091-29100(2010)Kelemen, L.E., et al. Cancer Epidemiol. Biomarkers Prev. 19(7):1822-1830(2010)Park, C.W., et al. J Cardiovasc Transl Res 3(3):290-295(2010)Haggarty, P., et al. PLoS ONE 5 (6), E11329 (2010) :Zhao, Z., et al. J. Biomed. Biotechnol. 2010, 737535 (2010) :