

DNMT1 Rabbit pAb

货号: B20856

产品信息

反应	Human,Mouse,Rat
宿主	Rabbit
克隆性	Polyclonal
预测反应	
应用	WB IHC IF/ICC IP ChIP
推荐浓度	WB: 1:500 - 1:1000 IHC: 1:50 - 1:200 IF/ICC: 1:50 - 1:200 IP: 1:500 - 1:1000 ChIP: 1:500 - 1:1000
理论分子量	144kDa/183kDa/184kDa
实测分子量	200KDa
形式	Liquid
保存条件	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.01% thiomersal,50% glycerol,pH7.3.
偶联物	Unconjugated
阳性对照	Jurkat
细胞定位	Nucleus
纯化	Affinity purification

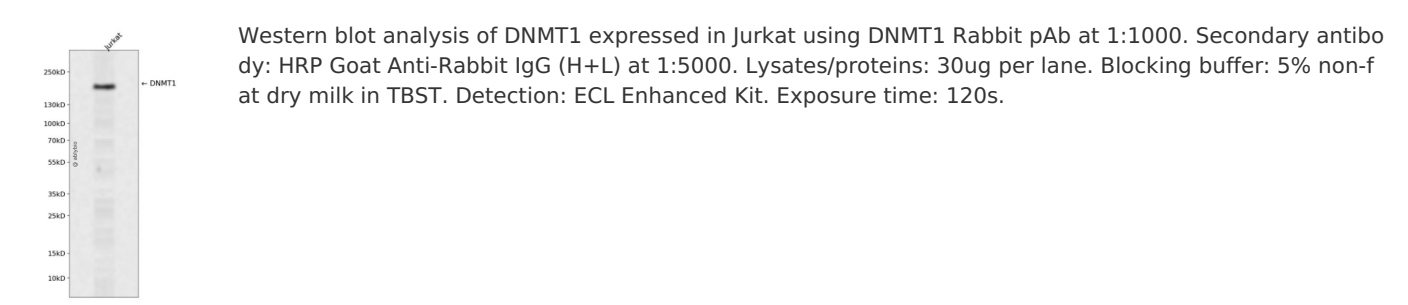
抗原信息

抗原信息	Recombinant fusion protein containing a sequence corresponding to amino acids 760-1109 of DNMT1 (NP_001370.1).
序列	GDCVSVIPDDSSKPLYLARVTALWEDSSNGQMFHAHWFCAGTDTVLGATSDPLELFLVDECEDMQLSYIHSKVKVIYKAP SENWAMEGGMDPESLLEGDDGKTYFYQLWYDQDYARFESPCKTQPTEDNKFKFCVSCARLAEMRQKEIPRVLEQLEDL DSRVLYYSATKNGILYRVGDGVYLPPEAFTFNIKSSPVKRPRKEPVDEDLYPEHYRKYSYIYKGSNLDAPEPYRIGRIKEIFC PKKSNRPNETDIKIRVNFYRPENTHKSTPASYHADINLLYWSDEEAVVDFKAVQGRCTVEYGEDLPECVQVYSMGGPN RFYFLEAYNAKSKSFEDPPNHARSPGNK

靶点信息

研究背景	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. [provided by RefSeq, Jan 2016]
基因ID	1786
基因名	DNMT1
Swiss	P26358
别名	ADCADN;AIM;CXXC9;DNMT;HSN1E;MCMT;m.Hsa1;DNMT1

产品验证



实验步骤

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