

# Product datasheet for KN217851BN

## DNMT1 Human Gene Knockout Kit (CRISPR)

### **Product data:**

#### OriGene Technologies, Inc.

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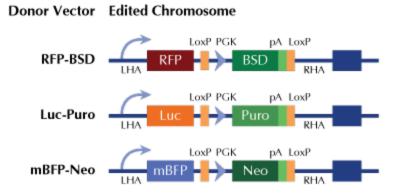
Product Type:	Knockout Kits (CRISPR)
Format:	2 gRNA vectors, 1 mBFP-Neo donor, 1 scramble control
Donor DNA:	mBFP-Neo
Symbol:	DNMT1
Locus ID:	1786
Components:	<ul> <li>KN217851G1, DNMT1 gRNA vector 1 in pCas-Guide CRISPR vector (GE100002)</li> <li>KN217851G2, DNMT1 gRNA vector 2 in pCas-Guide CRISPR vector (GE100002)</li> <li>KN217851BND, donor DNA containing left and right homologous arms and mBFP-Neo functional cassette.</li> <li>GE100003, scramble sequence in pCas-Guide vector</li> </ul>
Disclaimer:	These products are manufactured and supplied by OriGene under license from ERS. The kit is designed based on the best knowledge of CRISPR technology. The system has been functionally validated for knocking-in the cassette downstream the native promoter. The efficiency of the knock-out varies due to the nature of the biology and the complexity of the experimental process.
RefSeq:	<u>NM 001130823, NM 001318730, NM 001318731, NM 001379</u>
UniProt ID:	<u>P26358</u>
Synonyms:	ADCADN; AIM; CXXC9; DNMT; HSN1E; m.Hsal; MCMT
Summary:	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]



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#### **Product images:**



RFP, Luc, and mBFP will be under native gene promoter

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