

Product datasheet for **KN214878BN**

JMJD1C Human Gene Knockout Kit (CRISPR)

Product data:

Product Type:	Knockout Kits (CRISPR)
Format:	2 gRNA vectors, 1 mBFP-Neo donor, 1 scramble control
Donor DNA:	mBFP-Neo
Symbol:	JMJD1C
Locus ID:	221037
Components:	<p>KN214878G1, JMJD1C gRNA vector 1 in pCas-Guide CRISPR vector (GE100002)</p> <p>KN214878G2, JMJD1C gRNA vector 2 in pCas-Guide CRISPR vector (GE100002)</p> <p>KN214878BND, donor DNA containing left and right homologous arms and mBFP-Neo functional cassette.</p> <p>GE100003, scramble sequence in pCas-Guide vector</p>
Disclaimer:	<p>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.</p>
RefSeq:	<p>NM_001282948, NM_001318153, NM_001318154, NM_004241, NM_032776, NR_134512, NM_001322252, NM_001322254, NM_001322258</p>
UniProt ID:	Q15652
Synonyms:	TRIP8
Summary:	<p>The protein encoded by this gene interacts with thyroid hormone receptors and contains a jumonji domain. It is a candidate histone demethylase and is thought to be a coactivator for key transcription factors. It plays a role in the DNA-damage response pathway by demethylating the mediator of DNA damage checkpoint 1 (MDC1) protein, and is required for the survival of acute myeloid leukemia. Mutations in this gene are associated with Rett syndrome and intellectual disability. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Dec 2015]</p>



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Product images:

