

Product datasheet for KN211013BN

B Raf (BRAF) 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: B Raf Locus ID: 673

Components: KN211013G1, B Raf gRNA vector 1 in pCas-Guide CRISPR vector (GE100002)

KN211013G2, B Raf gRNA vector 2 in pCas-Guide CRISPR vector (GE100002)

KN211013BND, donor DNA containing left and right homologous arms and mBFP-Neo

functional cassette.

GE100003, scramble sequence in pCas-Guide vector

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 004333</u>, <u>NM 001354609</u>, <u>NR 148928</u>

UniProt ID: P15056

Synonyms: B-raf; B-RAF1; BRAF1; NS7; RAFB1

Summary: This gene encodes a protein belonging to the RAF family of serine/threonine protein kinases.

This protein plays a role in regulating the MAP kinase/ERK signaling pathway, which affects cell division, differentiation, and secretion. Mutations in this gene, most commonly the V600E mutation, are the most frequently identified cancer-causing mutations in melanoma, and have been identified in various other cancers as well, including non-Hodgkin lymphoma, colorectal cancer, thyroid carcinoma, non-small cell lung carcinoma, hairy cell leukemia and

adenocarcinoma of lung. Mutations in this gene are also associated with

cardiofaciocutaneous, Noonan, and Costello syndromes, which exhibit overlapping

phenotypes. A pseudogene of this gene has been identified on the X chromosome. [provided

by RefSeq, Aug 2017]



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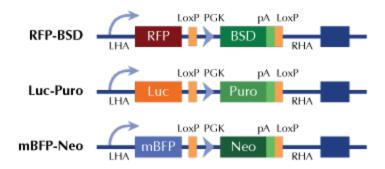
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Product images:

Donor Vector Edited Chromosome



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