

### Product datasheet for KN211292BN

## **GLDC 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

**GLDC** Symbol: Locus ID: 2731

**KN211292G1**, GLDC gRNA vector 1 in pCas-Guide CRISPR vector (GE100002) Components:

**KN211292G2**, GLDC gRNA vector 2 in pCas-Guide CRISPR vector (GE100002)

KN211292BND, 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.

NM 000170 RefSeq:

UniProt ID: P23378

Synonyms: GCE; GCSP; HYGN1

Summary: Degradation of glycine is brought about by the glycine cleavage system, which is composed of

> four mitochondrial protein components: P protein (a pyridoxal phosphate-dependent glycine decarboxylase), H protein (a lipoic acid-containing protein), T protein (a tetrahydrofolaterequiring enzyme), and L protein (a lipoamide dehydrogenase). The protein encoded by this gene is the P protein, which binds to glycine and enables the methylamine group from glycine to be transferred to the T protein. Defects in this gene are a cause of nonketotic

hyperglycinemia (NKH).[provided by RefSeq, Jan 2010]



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# **Product images:**

#### Donor Vector Edited Chromosome



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