

# Product datasheet for KN302893BN

# PD-L1 (Cd274) Mouse 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: PD-L1 Locus ID: 60533 KN302893G1, PD-L1 gRNA vector 1 in pCas-Guide CRISPR vector (GE100002) **Components:** KN302893G2, PD-L1 gRNA vector 2 in pCas-Guide CRISPR vector (GE100002) KN302893BND, 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 021893 RefSeq: UniProt ID: Q9EP73 Synonyms: A530045L16Rik; B7h1; Pdcd1l1; Pdcd1lg1; Pdl1 Summary: The protein encoded by this gene is an immune inhibitory receptor ligand that is expressed by hematopoietic and non-hematopoietic cells, such as T cells and B cells and various types of tumor cells. The encoded protein is a type I transmembrane protein that has immunoglobulin V-like and C-like domains. Interaction of this ligand with its receptor inhibits T-cell activation and cytokine production. During infection or inflammation of normal tissue, this interaction is important for preventing autoimmunity by maintaining homeostasis of the immune response. In tumor microenvironments, this interaction provides an immune escape for tumor cells through cytotoxic T-cell inactivation. Mice deficient for this gene display a

variety of phenotypes including decreased allogeneic fetal survival rates and severe experimental autoimmune encephalomyelitis. [provided by RefSeq, Sep 2015]



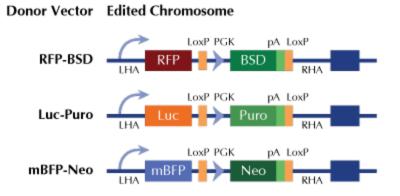
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### **Product images:**



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

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