

Product datasheet for **KN208745**

DDR2 Human Gene Knockout Kit (CRISPR)

Product data:

Product Type: Knockout Kits (CRISPR)
Format: 2 gRNA vectors, 1 GFP-puro donor, 1 scramble control
Donor DNA: GFP-puro
Symbol: DDR2
Locus ID: 4921
Components: **KN208745G1**, DDR2 gRNA vector 1 in pCas-Guide CRISPR vector (GE100002), Target Sequence: TGAGTAATAGCCATGTTACC
KN208745G2, DDR2 gRNA vector 2 in pCas-Guide CRISPR vector (GE100002), Target Sequence: GCAAAAGCTCAGGTTAATCC
KN208745D, donor DNA containing left and right homologous arms and GFP-puro functional cassette.

Homologous arm and GFP-puro sequences:

pUC vector backbone in gray; **Left arm sequence in blue**; **GFP-puro in green**; **Right arm in violet**

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AGAAGTAAGT TGGCCGAGT GTTATCACTC ATGGTTATGG CAGCACTGCA TAATTCTCTT ACTGTCATGC
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 TATTAATTGT TGCCGGGAAG CTAGAGTAAG TAGTTCGCCA GTTAATAGTT TGCGCAACGT TGTGGCCATT
 GCTACAGGCA TCGTGGTGTG ACGCTCGTCG TTTGGTATGG CTTCAATCAG CTCCGGTTCC CAACGATC

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:

[NM_001014796](#), [NM_006182](#), [NM_001354982](#), [NM_001354983](#)

UniProt ID:

[Q16832](#)

Synonyms:

MIG20a; NTRKR3; TKT; TYRO10

Summary:

This gene encodes a member of the discoidin domain receptor subclass of the receptor tyrosine kinase (RTKs) protein family. RTKs play a key role in the communication of cells with their microenvironment. The encoded protein is a collagen-induced receptor that activates signal transduction pathways involved in cell adhesion, proliferation, and extracellular matrix remodeling. This protein is expressed in numerous cell types and may also be involved in wound repair and regulate tumor growth and invasiveness. Mutations in this gene are the cause of short limb-hand type spondylometaphyseal dysplasia. [provided by RefSeq, Aug 2017]

Product images:

