

## Product datasheet for **KN413013**

### **RIM1 (RIMS1) Human Gene Knockout Kit (CRISPR)**

#### **Product data:**

Product Type:	Knockout Kits (CRISPR)
Format:	2 gRNA vectors, 1 linear donor
Donor DNA:	EF1a-GFP-P2A-Puro
Symbol:	RIM1
Locus ID:	22999



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**Components:**
**KN413013G1**, RIM1 gRNA vector 1 in pCas-Guide CRISPR vector (GE100002)

**KN413013G2**, RIM1 gRNA vector 2 in pCas-Guide CRISPR vector (GE100002)

**KN413013D**, Linear donor DNA containing LoxP-EF1a-tGFP-P2A-Puro-LoxP:

The sequence below is cassette sequence only. The linear donor DNA also contains proprietary target sequence.

LoxP-EF1a-tGFP-P2A-Puro-LoxP (2739 bp)

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ATAACTTCGT ATAATGTATG CTATACGAAG TTATCGTGAG GCTCCGGTGC CCGTCAGTGG GCAGAGCGCA
CATCGCCAC AGTCCCCGAG AAGTTGGGG GAGGGTTCGG CAATTGAACC GGTGCCTAGA GAAGGTGGCG
CGGGGTAAC TGGGAAAGTG ATGTCGTGTA CTGGCTCCGC CTTTTCCCG AGGGTGGGG AGAACCGTAT
ATAAGTCAG TAGTCGCCG GAACGTTCTT TTTCCGAACG GGTTCGCCG CAGAACACAG GTAAGTGCCG
TGTGTGGTTC CCGCGGGCCT GGCCTCTTTA CGGGTTATGG CCCTTGCGTG CCTTGAATTA CTTCCACCTG
GCTGCAGTAC GTGATTCTTG ATCCCGAGCT TCGGGTTGGA AGTGGGTGGG AGAGTTCGAG GCCTTGCGCT
TAAGGAGCCC CTTCGCCTCG TGCTTGAGTT GAGGCCTGGC CTGGGCGCTG GGGCCCGCG GTGCGAATCT
GGTGGCACCT TCGCGCCTGT CTCGCTGCTT TCGATAAGTC TCTAGCCATT TAAAATTTT GATGACCTGC
TGCAGCGCTT TTTTCTGGC AAGATAGTCT TGTAAATGCG GGCCAAGATC TGCACACTGG TATTTTCGTT
TTTGGGGCCG CGGGCGGCGA CGGGGCCCGT GCGTCCCAGC GCACATGTTC GGCAGGCGG GGCCTGCGAG
CGCGGCCACC GAGAATCGGA CGGGGGTAGT CTCAAGCTGG CCGGCCTGCT CTGGTGCCTG GCCTCGCGCC
GCCGTGTATC GCCCGCCCT GGGCGGCAAG GCTGGCCCGG TCGGCACCAG TTGCGTGAGC GGAAAGATGG
CCGCTTCCCG GCCCTGTGC AGGGAGCTCA AAATGGAGGA CGCGGCGCTC GGGAGAGCGG GCGGGTGAAGT
CACCCACACA AAGGAAAAGG GCCTTCCCGT CCTCAGCCGT CGCTTCATGT GACTCCACGG AGTACCGGGC
GCCGCTCAGG CACCTCGATT AGTTCTCGAG CTTTTGGAGT ACGTCGTCTT TAGTTGGGG GGAGGGGTTT
TATGCGATGG AGTTTCCCA CACTGAGTGG GTGGAGACTG AAGTTAGGCC AGCTTGGCAG TTGATGTAAT
TCTCCTTGGG ATTTGCCCTT TTTGAGTTTG GATCTTGGTT CATTCTCAAG CCTCAGACAG TGGTTCAAAG
TTTTTTCTT CCATTTCAAG TGTCGTGAAT GGAGAGCGAC GAGAGCGGCC TGCCCGCCAT GGAGATCGAG
TGCCGCATCA CCGGCACCCT GAACGGCGTG GAGTTCGAGC TGGTGGGCGG CGGAGAGGGC ACCCCCGAGC
AGGGCCGCAT GACCAACAAG ATGAAGAGCA CCAAAGGCGC CCTGACCTTC AGCCCTACC TGCTGAGCCA
CGTGATGGC TACGGCTTCT ACCACTTCGG CACCTACCCC AGCGGCTACG AGAACCCCTT CCTGCACGCC
ATCAACAACG GCGGCTACAC CAACACCCG ATCGAGAAGT ACGAGGACGG CGGCGTGCTG CACGTGAGCT
TCAGCTACCG CTACGAGGCC GGCCGCGTGA TCGGCGACTT CAAGGTGATG GGCACC GGCT TCCCCGAGGA
CAGCGTGATC TTCACCGACA AGATCATCCG CAGCAACGCC ACCGTGGAGC ACCTGCACCC CATGGCGGAT
AACGATCTGG ATGGCAGCTT CACCCGACC TTCAGCCTGC GCGACGCGG CTA CTACAGC TCCGTGGTGG
ACAGCCACAT GCACTTCAAG AGCGCCATCC ACCCCAGCAT CCTGCAGAAC GGGGGCCCA TGTTCCCTT
CCGCCCGTG GAGGAGGATC ACAGCAACAC CGAGTGGGG ATCGTGGAGT ACCAGCACG CTTCAAGACC
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CCCCAGGGCC GTACGCACCC TCGCCGCCG GTTCGCCGAC TACCCGCCA CGGCCACAC CGTCGATCCG
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AGGTGTGGT CGCGGACGAC GGCGCCCGG TGGCGGTCTG GACCACGCC GAGAGCGTCG AAGCGGGGGC
GGTGTTCGCC GAGATCGGCC CGCGCATGGC CGAGTTGAGC GGTTCGGG TGGCCCGCA GCAACAGATG
GAAGGCCTCC TGCGCCGCA CCGGCCAAG GAGCCCGCT GTTCTCTGGC CACCCTCGG GTCTCGCCG
ACCACCAGG CAAGGTCTG GGCAGCGCG TCGTGTCCC CGGAGTGGAG GCGGCCGAGC GCGCCGGGT
GCCCGCTTC CTGGAGACT CCGCGCCCG CAACCTCCC TTCTACGAGC GGCTCGGCT CACCGTCACC
GCCGACGTC AGGTGCCGA AGGACCGCG ACCTGGTGA TGACCCGCA GCCCGGTGCC TGAAACTTGT
TTATTGCAGC TTATAATGGT TACAAATAA GCAATAGCAT CACAAATTC ACAAATAAG CATTTTTTTC
ACTGCATTCT AGTTGTGGT TGTCCAACT CATCAATGTA TCTTAATAAC TTCGTATAAT GTATGCTATA CGAAGTTAT
    
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<b>Disclaimer:</b>	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.
<b>RefSeq:</b>	<a href="#">NM_001168407</a> , <a href="#">NM_001168408</a> , <a href="#">NM_001168409</a> , <a href="#">NM_001168410</a> , <a href="#">NM_001168411</a> , <a href="#">NM_014989</a> , <a href="#">NM_001350411</a> , <a href="#">NM_001350412</a> , <a href="#">NM_001350413</a> , <a href="#">NM_001350414</a> , <a href="#">NM_001350415</a> , <a href="#">NM_001350416</a> , <a href="#">NM_001350417</a> , <a href="#">NM_001350418</a> , <a href="#">NM_001350419</a> , <a href="#">NM_001350420</a> , <a href="#">NM_001350421</a> , <a href="#">NM_001350422</a> , <a href="#">NM_001350423</a> , <a href="#">NM_001350424</a> , <a href="#">NM_001350425</a> , <a href="#">NM_001350426</a> , <a href="#">NM_001350427</a> , <a href="#">NM_001350428</a> , <a href="#">NM_001350429</a> , <a href="#">NM_001350430</a> , <a href="#">NM_001350431</a> , <a href="#">NM_001350432</a> , <a href="#">NM_001350433</a> , <a href="#">NM_001350434</a> , <a href="#">NM_001350435</a> , <a href="#">NM_001350436</a> , <a href="#">NM_001350437</a> , <a href="#">NM_001350438</a> , <a href="#">NM_001350439</a> , <a href="#">NM_001350440</a> , <a href="#">NM_001350441</a> , <a href="#">NM_001350442</a> , <a href="#">NM_001350443</a> , <a href="#">NM_001350444</a> , <a href="#">NM_001350445</a> , <a href="#">NM_001350446</a> , <a href="#">NM_001350447</a> , <a href="#">NM_001350448</a> , <a href="#">NM_001350449</a> , <a href="#">NM_001350450</a> , <a href="#">NM_001350452</a> , <a href="#">NM_001350454</a> , <a href="#">NM_001350455</a> , <a href="#">NM_001350456</a> , <a href="#">NM_001350457</a> , <a href="#">NM_001350458</a> , <a href="#">NM_001350459</a> , <a href="#">NM_001350460</a> , <a href="#">NM_001350461</a> , <a href="#">NM_001350462</a> , <a href="#">NM_001350463</a> , <a href="#">NM_001350464</a> , <a href="#">NM_001350465</a> , <a href="#">NM_001350466</a> , <a href="#">NM_001350467</a> , <a href="#">NM_001350468</a> , <a href="#">NM_001350469</a> , <a href="#">NM_001350470</a> , <a href="#">NM_001350471</a> , <a href="#">NM_001350472</a> , <a href="#">NM_001350473</a> , <a href="#">NM_001350474</a>
<b>UniProt ID:</b>	<a href="#">Q86UR5</a>
<b>Synonyms:</b>	CORD7; RAB3IP2; RIM; RIM1
<b>Summary:</b>	The protein encoded by this gene is a RAS gene superfamily member that regulates synaptic vesicle exocytosis. This gene also plays a role in the regulation of voltage-gated calcium channels during neurotransmitter and insulin release. Mutations have suggested a role cognition and have been identified as the cause of cone-rod dystrophy type 7. Multiple transcript variants encoding different isoforms have been described for this gene. [provided by RefSeq, Mar 2012]

Product images:

