

## Product datasheet for **KN204456**

### DNA Polymerase gamma (POLG) 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:	DNA Polymerase gamma
Locus ID:	5428
Components:	<p><b>KN204456G1</b>, DNA Polymerase gamma gRNA vector 1 in pCas-Guide CRISPR vector (GE100002), Target Sequence: CCACCTTCCTCCAGAGCAGG</p> <p><b>KN204456G2</b>, DNA Polymerase gamma gRNA vector 2 in pCas-Guide CRISPR vector (GE100002), Target Sequence: CAGGCGCCGGTCCAGCTCCG</p> <p><b>KN204456D</b>, donor DNA containing left and right homologous arms and GFP-puro functional cassette.</p>

#### 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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CGCCAACACC CGCTGACGCG CCCTGACGGG CTTGTCTGCT CCCGGCATCC GCTTACAGAC AAGCTGTGAC
CGTCTCCGGG AGCTGCATGT GTCAGAGGTT TTCACCGTCA TCACCGAAAC GCGCGAGGCA GCTGCGGTAA
AGTCATCAG CGTGGTCGTG AAGCGATTCA CAGATGTCTG CCTGTTTCATC CGCGTCCAGC TCGTTGAGTT
TCTCCAGAAG CGTAAATGTC TGGCTTCTGA TAAAGCGGGC CATGTTAAGG GCGGTTTTTTT CCTGTTTGGT
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 ATACACTCCG CTATCGCTAC GTGACTGGGT CATGGCTGCG CCCCAGACCC

**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\\_001126131](#), [NM\\_002693](#)

UniProt ID: [P54098](#)

Synonyms: MDP1; MIRAS; MTDPS4A; MTDPS4B; PEO; POLG1; POLGA; SANDO; SCAE

**Summary:** Mitochondrial DNA polymerase is heterotrimeric, consisting of a homodimer of accessory subunits plus a catalytic subunit. The protein encoded by this gene is the catalytic subunit of mitochondrial DNA polymerase. The encoded protein contains a polyglutamine tract near its N-terminus that may be polymorphic. Defects in this gene are a cause of progressive external ophthalmoplegia with mitochondrial DNA deletions 1 (PEOA1), sensory ataxic neuropathy dysarthria and ophthalmoparesis (SANDO), Alpers-Huttenlocher syndrome (AHS), and mitochondrial neurogastrointestinal encephalopathy syndrome (MNGIE). Two transcript variants encoding the same protein have been found for this gene. [provided by RefSeq, Jul 2008]

### Product images:

