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Product datasheet for KN202135

ASAH1 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: ASAH1

Locus ID: 427

Components: **KN202135G1**, ASAH1 gRNA vector 1 in pCas-Guide CRISPR vector (GE100002), Target Sequence: GGGCTGGGAGAGAAAGCTCG
KN202135G2, ASAH1 gRNA vector 2 in pCas-Guide CRISPR vector (GE100002), Target Sequence: AAGCGCGCTGAGACTTGGGT
KN202135D, 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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TACAGGCATC GTGGTGTCAC GCTCGTCGTT TGGTATGGCT TCATTCAGCT CCGGTTCCCA ACGATC

GE100003, scramble sequence in pCas-Guide vector

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RefSeq: NM_001127505, NM_004315, NM_177924, NM_001363743

UniProt ID: Q13510

Synonyms: AC; ACDase; ASAH; PHP; PHP32; SMAPME

Summary: This gene encodes a member of the acid ceramidase family of proteins. Alternative splicing results in multiple transcript variants, at least one of which encodes a preproprotein that is proteolytically processed. Processing of this preproprotein generates alpha and beta subunits that heterodimerize to form the mature lysosomal enzyme, which catalyzes the degradation of ceramide into sphingosine and free fatty acid. This enzyme is overexpressed in multiple human cancers and may play a role in cancer progression. Mutations in this gene are associated with the lysosomal storage disorder, Farber lipogranulomatosis, and a neuromuscular disorder, spinal muscular atrophy with progressive myoclonic epilepsy. [provided by RefSeq, Oct 2015]

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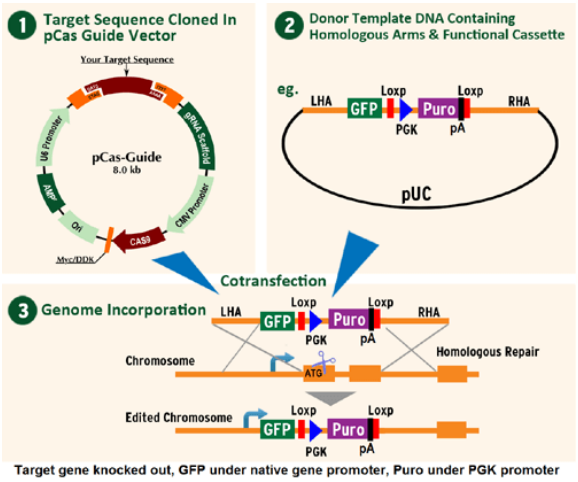

Target gene knocked out, GFP under native gene promoter, Puro under PGK promoter

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