

## Product datasheet for **KN202135LP**

### ASAH1 Human Gene Knockout Kit (CRISPR)

#### Product data:

Product Type:	Knockout Kits (CRISPR)
Format:	2 gRNA vectors, 1 Luciferase-Puro donor, 1 scramble control
Donor DNA:	Luciferase-Puro
Symbol:	ASAH1
Locus ID:	427
Components:	<p><b>KN202135G1</b>, ASAH1 gRNA vector 1 in pCas-Guide CRISPR vector (GE100002)</p> <p><b>KN202135G2</b>, ASAH1 gRNA vector 2 in pCas-Guide CRISPR vector (GE100002)</p> <p><b>KN202135LPD</b>, donor DNA containing left and right homologous arms and Luciferase-Puro functional cassette.</p> <p><b>GE100003</b>, scramble sequence in pCas-Guide vector</p>
RefSeq:	<a href="#">NM_001127505</a> , <a href="#">NM_004315</a> , <a href="#">NM_177924</a> , <a href="#">NM_001363743</a>
UniProt ID:	<a href="#">Q13510</a>
Synonyms:	AC; ACDase; ASAH; PHP; PHP32; SMAPME
Summary:	<p>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]</p>



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## Product images:

