

## Product datasheet for **KN214709BN**

### CLN5 Human Gene Knockout Kit (CRISPR)

#### Product data:

Product Type:	Knockout Kits (CRISPR)
Format:	2 gRNA vectors, 1 mBFP-Neo donor, 1 scramble control
Donor DNA:	mBFP-Neo
Symbol:	CLN5
Locus ID:	1203
Components:	<b>KN214709G1</b> , CLN5 gRNA vector 1 in pCas-Guide CRISPR vector (GE100002) <b>KN214709G2</b> , CLN5 gRNA vector 2 in pCas-Guide CRISPR vector (GE100002) <b>KN214709BND</b> , donor DNA containing left and right homologous arms and mBFP-Neo functional cassette. <b>GE100003</b> , 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:	<a href="#">NM_006493</a> , <a href="#">NM_001366624</a>
UniProt ID:	<a href="#">O75503</a>
Synonyms:	NCL
Summary:	This gene is one of eight which have been associated with neuronal ceroid lipofuscinoses (NCL). Also referred to as Batten disease, NCL comprises a class of autosomal recessive, neurodegenerative disorders affecting children. The genes responsible likely encode proteins involved in the degradation of post-translationally modified proteins in lysosomes. The primary defect in NCL disorders is thought to be associated with lysosomal storage function. [provided by RefSeq, Oct 2008]



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## Product images:

