

## Product datasheet for **KN202671LP**

### **TMEM138 Human Gene Knockout Kit (CRISPR)**

#### **Product data:**

<b>Product Type:</b>	Knockout Kits (CRISPR)
<b>Format:</b>	2 gRNA vectors, 1 Luciferase-Puro donor, 1 scramble control
<b>Donor DNA:</b>	Luciferase-Puro
<b>Symbol:</b>	TMEM138
<b>Locus ID:</b>	51524
<b>Components:</b>	<b>KN202671G1</b> , TMEM138 gRNA vector 1 in pCas-Guide CRISPR vector (GE100002) <b>KN202671G2</b> , TMEM138 gRNA vector 2 in pCas-Guide CRISPR vector (GE100002) <b>KN202671LPD</b> , donor DNA containing left and right homologous arms and Luciferase-Puro functional cassette. <b>GE100003</b> , scramble sequence in pCas-Guide vector
<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_001330281</a> , <a href="#">NM_016464</a> , <a href="#">NR_028473</a>
<b>UniProt ID:</b>	<a href="#">Q9NPI0</a>
<b>Synonyms:</b>	HSPC196
<b>Summary:</b>	This gene encodes a multi-pass transmembrane protein. Reduced expression of this gene in mouse fibroblasts causes short cilia and failure of ciliogenesis. Expression of this gene is tightly coordinated with expression of the neighboring gene TMEM216. Mutations in this gene are associated with the autosomal recessive neurodevelopmental disorder Joubert Syndrome. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Mar 2012]



[View online »](#)

## Product images:

