

## Product datasheet for **RC228123L3V**

### **MNX1 (NM\_001165255) Human Tagged ORF Clone Lentiviral Particle**

#### **Product data:**

<b>Product Type:</b>	Lentiviral Particles
<b>Product Name:</b>	MNX1 (NM_001165255) Human Tagged ORF Clone Lentiviral Particle
<b>Symbol:</b>	MNX1
<b>Synonyms:</b>	HB9; HLXB9; HOXHB9; SCRA1
<b>Mammalian Cell Selection:</b>	Puromycin
<b>Vector:</b>	pLenti-C-Myc-DDK-P2A-Puro (PS100092)
<b>Tag:</b>	Myc-DDK
<b>ACCN:</b>	NM_001165255
<b>ORF Size:</b>	567 bp
<b>ORF Nucleotide Sequence:</b>	The ORF insert of this clone is exactly the same as(RC228123).
<b>OTI Disclaimer:</b>	The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. <a href="#">More info</a>
<b>OTI Annotation:</b>	This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.
<b>RefSeq:</b>	<a href="#">NM_001165255.1</a> , <a href="#">NP_001158727.1</a>
<b>RefSeq ORF:</b>	570 bp
<b>Locus ID:</b>	3110
<b>UniProt ID:</b>	<a href="#">P50219</a>
<b>Cytogenetics:</b>	7q36.3
<b>Protein Families:</b>	Druggable Genome, ES Cell Differentiation/IPS
<b>Protein Pathways:</b>	Maturity onset diabetes of the young
<b>MW:</b>	20.4 kDa



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**Gene Summary:**

This gene encodes a nuclear protein, which contains a homeobox domain and is a transcription factor. Mutations in this gene result in Currarino syndrome, an autosomic dominant congenital malformation. Alternatively spliced transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Sep 2009]