

## Product datasheet for **KN202920RB**

### Twist (TWIST1) Human Gene Knockout Kit (CRISPR)

#### Product data:

Product Type:	Knockout Kits (CRISPR)
Format:	2 gRNA vectors, 1 RFP-BSD donor, 1 scramble control
Donor DNA:	RFP-BSD
Symbol:	Twist
Locus ID:	7291
Components:	<p><b>KN202920G1</b>, Twist gRNA vector 1 in pCas-Guide CRISPR vector (GE100002)</p> <p><b>KN202920G2</b>, Twist gRNA vector 2 in pCas-Guide CRISPR vector (GE100002)</p> <p><b>KN202920RBD</b>, donor DNA containing left and right homologous arms and RFP-BSD functional cassette.</p> <p><b>GE100003</b>, scramble sequence in pCas-Guide vector</p>
RefSeq:	<u><a href="#">NM_000474</a></u> , <u><a href="#">NR_149001</a></u>
UniProt ID:	<u><a href="#">Q15672</a></u>
Synonyms:	ACS3; bHLHa38; BPES2; BPES3; CRS; CRS1; CSO; SCS; TWIST
Summary:	<p>This gene encodes a basic helix-loop-helix (bHLH) transcription factor that plays an important role in embryonic development. The encoded protein forms both homodimers and heterodimers that bind to DNA E box sequences and regulate the transcription of genes involved in cranial suture closure during skull development. This protein may also regulate neural tube closure, limb development and brown fat metabolism. This gene is hypermethylated and overexpressed in multiple human cancers, and the encoded protein promotes tumor cell invasion and metastasis, as well as metastatic recurrence. Mutations in this gene cause Saethre-Chotzen syndrome in human patients, which is characterized by craniosynostosis, ptosis and hypertelorism. [provided by RefSeq, Jul 2020]</p>



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## Product images:

