

## Product datasheet for RC201618L1V

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## **GATA3 (NM\_002051) Human Tagged ORF Clone Lentiviral Particle**

**Product data:** 

Product Type: Lentiviral Particles

**Product Name:** GATA3 (NM\_002051) Human Tagged ORF Clone Lentiviral Particle

Symbol: GATA3

Synonyms: HDR; HDRS

Mammalian Cell

Selection:

None

**Vector:** pLenti-C-Myc-DDK (PS100064)

 Tag:
 Myc-DDK

 ACCN:
 NM\_002051

 ORF Size:
 1332 bp

**ORF Nucleotide** 

Sequence:

The ORF insert of this clone is exactly the same as(RC201618).

OTI Disclaimer:

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. More info

**OTI Annotation:** This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

RefSeq: <u>NM 002051.2</u>

 RefSeq Size:
 3067 bp

 RefSeq ORF:
 1332 bp

 Locus ID:
 2625

 UniProt ID:
 P23771

 Cytogenetics:
 10p14

 Domains:
 GATA





## GATA3 (NM\_002051) Human Tagged ORF Clone Lentiviral Particle - RC201618L1V

Protein Families: Adult stem cells, ES Cell Differentiation/IPS, Stem cell relevant signaling - JAK/STAT signaling

pathway, Transcription Factors

MW: 48 kDa

**Gene Summary:** This gene encodes a protein which belongs to the GATA family of transcription factors. The

protein contains two GATA-type zinc fingers and is an important regulator of T-cell

development and plays an important role in endothelial cell biology. Defects in this gene are the cause of hypoparathyroidism with sensorineural deafness and renal dysplasia. [provided

by RefSeq, Nov 2009]