

## Product datasheet for RC203132L4V

## OriGene Technologies, Inc.

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## RHAG (NM\_000324) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** RHAG (NM\_000324) Human Tagged ORF Clone Lentiviral Particle

Symbol: RHAG

Synonyms: CD241; OHS; OHST; RH2; Rh50; RH50A; Rh50GP; RHNR; SLC42A1

**Mammalian Cell** 

Selection:

Puromycin

**Vector:** pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

**ACCN:** NM\_000324 **ORF Size:** 1227 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC203132).

Sequence:

Cytogenetics:

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.

**RefSeg:** NM 000324.1

 RefSeq Size:
 1953 bp

 RefSeq ORF:
 1230 bp

 Locus ID:
 6005

 UniProt ID:
 Q02094

Domains: Ammonium\_transp

**Protein Families:** Druggable Genome, Transmembrane

6p12.3





## RHAG (NM\_000324) Human Tagged ORF Clone Lentiviral Particle - RC203132L4V

**MW:** 44.2 kDa

**Gene Summary:** The protein encoded by this gene is erythrocyte-specific and is thought to be part of a

membrane channel that transports ammonium and carbon dioxide across the blood cell membrane. The encoded protein appears to interact with Rh blood group antigens and Rh30 polypeptides. Defects in this gene are a cause of regulator type Rh-null hemolytic anemia

(RHN), or Rh-deficiency syndrome.[provided by RefSeq, Mar 2009]