

Product datasheet for RC202533L4V

OriGene Technologies, Inc.

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CXCL14 (NM_004887) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: CXCL14 (NM 004887) Human Tagged ORF Clone Lentiviral Particle

Symbol: CXCL14

Synonyms: BMAC; BRAK; KEC; KS1; MIP-2g; MIP2G; NJAC; SCYB14

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

ACCN: NM_004887

ORF Size: 333 bp

ORF Nucleotide

TI 005

OTI Disclaimer:

Sequence:

The ORF insert of this clone is exactly the same as(RC202533).

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 004887.3

 RefSeq Size:
 1989 bp

 RefSeq ORF:
 300 bp

 Locus ID:
 9547

 UniProt ID:
 095715

 Cytogenetics:
 5q31.1

Protein Families: Druggable Genome, Secreted Protein, Transmembrane

Protein Pathways: Chemokine signaling pathway, Cytokine-cytokine receptor interaction





ORIGENE

MW: 13.1 kDa

Gene Summary:

This antimicrobial gene belongs to the cytokine gene family which encode secreted proteins involved in immunoregulatory and inflammatory processes. The protein encoded by this gene is structurally related to the CXC (Cys-X-Cys) subfamily of cytokines. Members of this subfamily are characterized by two cysteines separated by a single amino acid. This cytokine displays chemotactic activity for monocytes but not for lymphocytes, dendritic cells, neutrophils or macrophages. It has been implicated that this cytokine is involved in the homeostasis of monocyte-derived macrophages rather than in inflammation. [provided by RefSeq, Sep 2014]