

Product datasheet for RC222687L1V

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

ZNF148 (NM_021964) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: ZNF148 (NM 021964) Human Tagged ORF Clone Lentiviral Particle

Symbol: ZNF148

Synonyms: BERF-1; BFCOL1; GDACCF; HT-BETA; pHZ-52; ZBP-89; ZFP148

Mammalian Cell

Selection:

None

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

 Tag:
 Myc-DDK

 ACCN:
 NM_021964

ORF Size: 2382 bp

ORF Nucleotide Sequence:

The ORF insert of this clone is exactly the same as(RC222687).

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 021964.1

 RefSeq Size:
 3032 bp

 RefSeq ORF:
 2385 bp

 Locus ID:
 7707

 UniProt ID:
 Q9UQR1

 Cytogenetics:
 3q21.2

Domains: zf-C2H2

Protein Families: Transcription Factors





ORIGENE

MW: 88.8 kDa

Gene Summary: The protein encoded by this gene is a member of the Kruppel family of zinc finger DNA

binding proteins. The encoded protein activates transcription of the T-cell receptor and intestinal alkaline phosphatase genes but represses transcription of the ornithine

decarboxylase, vimentin, gastrin, stomelysin, and enolase genes. Increased expression of this gene results in decreased patient survival rates from colorectal cancer, while mutations in this gene have been associated with global developmental delay, hypoplastic corpus

callosum, and dysmorphic facies. [provided by RefSeq, Feb 2017]