

## Product datasheet for RC205569L3V

## OriGene Technologies, Inc.

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## RFXAP (NM 000538) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

**Product Name:** RFXAP (NM\_000538) Human Tagged ORF Clone Lentiviral Particle

Symbol: RFXAP

Mammalian Cell Puromycin

Selection:

Vector:

pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK

**ACCN:** NM\_000538

ORF Size: 816 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC205569).

Sequence:
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 000538.2</u>

 RefSeq Size:
 2825 bp

 RefSeq ORF:
 819 bp

 Locus ID:
 5994

 UniProt ID:
 000287

 Cytogenetics:
 13q13.3

**Protein Families:** Transcription Factors

**Protein Pathways:** Antigen processing and presentation, Primary immunodeficiency

MW: 28.1 kDa







## **Gene Summary:**

Major histocompatibility (MHC) class II molecules are transmembrane proteins that have a central role in development and control of the immune system. The protein encoded by this gene, along with regulatory factor X-associated ankyrin-containing protein and regulatory factor-5, forms a complex that binds to the X box motif of certain MHC class II gene promoters and activates their transcription. Once bound to the promoter, this complex associates with the non-DNA-binding factor MHC class II transactivator, which controls the cell type specificity and inducibility of MHC class II gene expression. Mutations in this gene have been linked to bare lymphocyte syndrome type II, complementation group D. Transcript variants utilizing different polyA signals have been found for this gene. [provided by RefSeq, Jul 2008]