

## Product datasheet for RC215220L3V

## OriGene Technologies, Inc.

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## NBPF3 (NM\_032264) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

**Product Name:** NBPF3 (NM\_032264) Human Tagged ORF Clone Lentiviral Particle

Symbol: NBPF3
Synonyms: AE2

Mammalian Cell Puromycin

Selection:

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**Vector:** pLenti-C-Myc-DDK-P2A-Puro (PS100092)

 Tag:
 Myc-DDK

 ACCN:
 NM\_032264

 ORF Size:
 1899 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC215220).

OTI Disclaimer:

Sequence:

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 032264.3

 RefSeq Size:
 3783 bp

 RefSeq ORF:
 1902 bp

 Locus ID:
 84224

 UniProt ID:
 Q9H094

 Cytogenetics:
 1p36.12

 MW:
 72.8 kDa







## **Gene Summary:**

This gene is a member of the neuroblastoma breakpoint family (NBPF) which consists of dozens of recently duplicated genes primarily located in segmental duplications on human chromosome 1. This gene family has experienced its greatest expansion within the human lineage and has expanded, to a lesser extent, among primates in general. Members of this gene family are characterized by tandemly repeated copies of DUF1220 protein domains. DUF1220 copy number variations in human chromosomal region 1q21.1, where most DUF1220 domains are located, have been implicated in a number of developmental and neurogenetic diseases such as microcephaly, macrocephaly, autism, schizophrenia, cognitive disability, congenital heart disease, neuroblastoma, and congenital kidney and urinary tract anomalies. Altered expression of some gene family members is associated with several types of cancer. This gene family contains numerous pseudogenes. [provided by RefSeq, Feb 2013]