

## Product datasheet for RC220251L3V

## OriGene Technologies, Inc.

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## Huntingtin Associated Protein 1 (HAP1) (NM\_177977) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** Huntingtin Associated Protein 1 (HAP1) (NM\_177977) Human Tagged ORF Clone Lentiviral

Particle

Symbol: Huntingtin Associated Protein 1

Synonyms: HAP2; hHLP1; HIP5; HLP

Mammalian Cell

Selection:

Puromycin

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

 Tag:
 Myc-DDK

 ACCN:
 NM\_177977

**ORF Size:** 1857 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC220251).

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.

**RefSeq:** <u>NM 177977.1</u>

 RefSeq Size:
 3981 bp

 RefSeq ORF:
 1860 bp

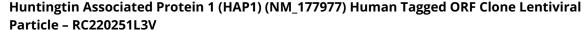
 Locus ID:
 9001

 UniProt ID:
 P54257

 Cytogenetics:
 17q21.2

**Protein Pathways:** Huntington's disease







**MW:** 69.5 kDa

**Gene Summary:** Huntington's disease (HD), a neurodegenerative disorder characterized by loss of striatal

neurons, is caused by an expansion of a polyglutamine tract in the HD protein huntingtin. This gene encodes a protein that interacts with huntingtin, with two cytoskeletal proteins (dynactin and pericentriolar autoantigen protein 1), and with a hepatocyte growth factor-regulated tyrosine kinase substrate. The interactions with cytoskeletal proteins and a kinase substrate suggest a role for this protein in vesicular trafficking or organelle transport. Several alternatively spliced transcript variants encoding different isoforms have been described for

this gene. [provided by RefSeq, Jul 2008]