

Product datasheet for RC220305L3

OriGene Technologies, Inc.

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Huntingtin Associated Protein 1 (HAP1) (NM 001079870) Human Tagged Lenti ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: Huntingtin Associated Protein 1 (HAP1) (NM_001079870) Human Tagged Lenti ORF Clone

Tag: Myc-DDK

Symbol: Huntingtin Associated Protein 1

Synonyms: HAP2; hHLP1; HIP5; HLP

Mammalian Cell

Selection:

Puromycin

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

E. coli Selection: Chloramphenicol (34 ug/mL)

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC220305).

Sequence:

Restriction Sites: Sgfl-Mlul

Cloning Scheme:





st The last codon before the Stop codon of the ORF.

ACCN: NM_001079870

ORF Size: 1806 bp





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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.

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube

containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

Reconstitution Method: 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of

shipping when stored at -20°C.

RefSeq: <u>NM 001079870.1</u>, <u>NP 001073339.1</u>

17q21.2

 RefSeq Size:
 3893 bp

 RefSeq ORF:
 1809 bp

 Locus ID:
 9001

 UniProt ID:
 P54257

Protein Pathways: Huntington's disease

MW: 67.4 kDa

Cytogenetics:

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]