



# Recombinant Human Huntingtin-associated protein 1 (HAP1)

<b>Product Code</b>	CSB-EP010129HU
<b>Storage</b>	Store at -20°C, for extended storage, conserve at -20°C or -80°C.
<b>Uniprot No.</b>	P54257
<b>Product Type</b>	Recombinant Protein
<b>Immunogen Species</b>	Homo sapiens (Human)
<b>Purity</b>	>85% (SDS-PAGE)
<b>Sequence</b>	MRPKRLGRCC AGSRLGPGDP AALTCAPSPS ASPAPEPSAQ PQARGTGQRV GSRATSGSQF LSEARTGARP ASEAGAKAGA RRPSAFSAIQ GDVRSMPDNS DAPWTRFVFQ GPFGRATGR GTGKAAGIWK TPAAYVGRRP GVSGPERAAF IRELEEALCP NLPPPVKKIT QEDVKVMLYL LEELLPPVWE SVTYGMVLQR ERDLNTAARI GQSLVKQNSV LMEENSKLEA LLGSAKEEIL YLRHQVNLRD ELLQLYSDSD EEDEDEEEEEE EEKEAEEEEQE EEEAEDLQC AHPCDAPKLI SQEALLHQHH CPQLEALQEK LRLLEENHQ LREEASQLDT LEDEEQMLIL ECVEQFSEAS QQMAELSEVL VLRLNEYERQ QQEVARLQAQ VLKLQQRCRM YGAETEKLQK QLASEKEIQM QLQEESVWVG SQLQLDLREKY MDCGGMLIEM QEEVKTLRQQ PPVSTGSATH YPYSVPLETL PGFQETLAE LRTSLRRMIS DPVYFMERNY EMPRGDTSSL RYDFRYSEDR EQVRGFEEAE GLMLAADIMR GEDFTPAEEF VPQEELGAAK KVPAAEEGVM EAEVSEETE GWEEVELELD EATRMNVVTS ALEASGLGPS HLDMNYVLQQ LANWQDAHYR RQLRWKMLQK GECPHGALPA ASRTSCRSSC R
<b>Source</b>	E.coli
<b>Target Names</b>	HAP1
<b>Protein Names</b>	Recommended name: Huntingtin-associated protein 1 Short name= HAP-1 Alternative name(s): Neuroan 1
<b>Expression Region</b>	1-671
<b>Notes</b>	Repeated freezing and thawing is not recommended. Store working aliquots at 4°C for up to one week.
<b>Tag Info</b>	Tag type will be determined during the manufacturing process.
<b>Protein Length</b>	Full length protein
<b>Target Details</b>	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.

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**Reconstitution**

We recommend that this vial be briefly centrifuged prior to opening to bring the contents to the bottom. Please reconstitute protein in deionized sterile water to a concentration of 0.1-1.0 mg/mL. We recommend to add 5-50% of glycerol (final concentration) and aliquot for long-term storage at -20°C/-80°C. Our default final concentration of glycerol is 50%. Customers could use it as reference.

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**Shelf Life**

The shelf life is related to many factors, storage state, buffer ingredients, storage temperature and the stability of the protein itself.

Generally, the shelf life of liquid form is 6 months at -20°C/-80°C. The shelf life of lyophilized form is 12 months at -20°C/-80°C.