



# Human HAP1 blocking peptide (CDBP1456)

This product is for research use only and is not intended for diagnostic use.

## PRODUCT INFORMATION

<b>Product Overview</b>	Blocking/Immunizing peptide for anti-HAP1 antibody
<b>Antigen Description</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.
<b>Species</b>	Human
<b>Conjugate</b>	Unconjugated
<b>Applications</b>	Apuri, BL, ELISA
<b>Format</b>	Lyophilized powder
<b>Size</b>	100 µg
<b>Preservative</b>	None
<b>Storage</b>	Shipped at ambient temperature, store at -20°C.

## GENE INFORMATION

<b>Gene Name</b>	<a href="#">Hap1 huntingtin-associated protein 1 [ Mus musculus ]</a>
<b>Official Symbol</b>	Hap1
<b>Synonyms</b>	HAP1; huntingtin-associated protein 1; HAP-1; MGC31449;

<b>Entrez Gene ID</b>	<a href="#">15114</a>
<b>mRNA Refseq</b>	<a href="#">NM_010404</a>
<b>Protein Refseq</b>	<a href="#">NP_034534</a>
<b>Pathway</b>	GABAergic synapse, organism-specific biosystem; GABAergic synapse, conserved biosystem; Huntingtons disease, organism-specific biosystem; Huntingtons disease, conserved biosystem;
<b>Function</b>	protein binding;