

Anti-HTT antibody



Description

Huntingtin is a protein linked to Huntington's disease, a neurodegenerative

disorder characterized by loss of striatal neurons.

Model STJ140045

Host Goat

Reactivity Avian, Bovine, Canine, Donkey, Feline, Goat, Guinea Pig, Hamster, Horse,

Human, Mouse, Other, Porcine, Rabbit, Rat, Sheep, Simian

Applications IF, IHC, WB

Immunogen Purified recombinant peptide derived from within residues 85 to 200 aa of

human HTT produced in E. coli.

Immunogen Region 85-200 aa

Gene ID 3064

Gene Symbol HTT

Dilution range Western blot 1:500-1:2,000 Immunofluorescence 1:500-1:2,000

Immunohistochemistry (paraffin) 1:500-1:2,000 Immunohistochemistry

(frozen) 1:500-1:2,000

Tissue Specificity Expressed in the brain cortex (at protein level). Widely expressed with the

highest level of expression in the brain (nerve fibers, varicosities, and nerve endings). In the brain, the regions where it can be mainly found are the cerebellar cortex, the neocortex, the striatum, and the hippocampal formation.

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Purification This antibody is epitope-affinity purified from goat antiserum.

Note For research use only (RUO).

Protein Name Huntingtin Huntington disease protein HD protein

Molecular Weight 348 kDa

Clonality Polyclonal

Conjugation Unconjugated

Isotype IgG

Formulation PBS, 20% glycerol and 0.05% sodium azide.

Concentration 3 mg/mL

Storage Instruction Store at -20°, and avoid repeated freeze-thaw cycles.

Database Links HGNC:48510MIM:143100

Alternative Names Huntingtin Huntington disease protein HD protein

Function May play a role in microtubule-mediated transport or vesicle function.

Sequence and Domain Family The N-terminal Gln-rich and Pro-rich domain has great conformational

flexibility and is likely to exist in a fluctuating equilibrium of alpha-helical,

random coil, and extended conformations.

Cellular Localization Cytoplasm. Nucleus. The mutant Huntingtin protein colocalizes with

AKAP8L in the nuclear matrix of Huntington disease neurons. Shuttles between cytoplasm and nucleus in a Ran GTPase-independent manner.

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