

YTHD2 rabbit pAb

Cat No.: ES11970

For research use only

Overview

Product Name YTHD2 rabbit pAb

Host species Rabbit
Applications WB;ELISA

Species Cross-Reactivity Human; Rat; Mouse;

Recommended dilutions WB 1:500-2000 ELISA 1:5000-20000

Immunogen Synthesized peptide derived from part region of

human protein

Specificity YTHD2 Polyclonal Antibody detects endogenous

levels of protein.

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and

0.02% sodium azide.

StorageStore at -20° C. Avoid repeated freeze-thaw cycles.Protein NameYTH domain family protein 2 (CLL-associated antigen)

KW-14) (High-glucose-regulated protein 8) (Renal

carcinoma antigen NY-REN-2)

Gene Name YTHDF2 HGRG8

Cellular localization Cytoplasm, cytosol . Cytoplasm, P-body . Cytoplasm,

Stress granule. Nucleus. Localizes to the cytosol and relocates to the nucleus following heat shock stress (PubMed:26458103). Can partition into different structures: into P-bodies in unstressed cells, and into stress granules during stress (PubMed:31292544).

Purification The antibody was affinity-purified from rabbit

antiserum by affinity-chromatography using

epitope-specific immunogen.

ClonalityPolyclonalConcentration1 mg/mlObserved band63kDHuman Gene ID51441Human Swiss-Prot NumberQ9Y5A9

Alternative Names

Background This gene encodes a member of the YTH (YT521-B

homology) superfamily containing YTH domain. The



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YTH domain is typical for the eukaryotes and is particularly abundant in plants. The YTH domain is usually located in the middle of the protein sequence and may function in binding to RNA. In addition to a YTH domain, this protein has a proline rich region which may be involved in signal transduction. An Alu-rich domain has been identified in one of the introns of this gene, which is thought to be associated with human longevity. In addition, reciprocal translocations between this gene and the Runx1 (AML1) gene on chromosome 21 has been observed in patients with acute myeloid leukemia. This gene was initially mapped to chromosome 14, which was later turned out to be a pseudogene. Alternatively spliced transcript variants encoding different isoforms have been identified in this gene

