

# RENT1 Antibody (N-term E22)

Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP1905a

### **Product Information**

**Application** WB, E **Primary Accession** Q92900 **Other Accession** Q9EPU0 Reactivity Human **Predicted** Mouse Host Rabbit Clonality Polyclonal Isotype Rabbit IgG **Calculated MW** 124345 7-37 **Antigen Region** 

### **Additional Information**

**Gene ID** 5976

Other Names Regulator of nonsense transcripts 1, 364-, ATP-dependent helicase RENT1,

Nonsense mRNA reducing factor 1, NORF1, Up-frameshift suppressor 1

homolog, hUpf1, UPF1, KIAA0221, RENT1

Target/Specificity This RENT1 antibody is generated from rabbits immunized with a KLH

conjugated synthetic peptide between 7-37 amino acids from the N-terminal

region of human RENT1.

**Dilution** WB~~1:1000 E~~Use at an assay dependent concentration.

**Format** Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide.

This antibody is prepared by Saturated Ammonium Sulfate (SAS) precipitation

followed by dialysis against PBS.

**Storage** Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions** RENT1 Antibody (N-term E22) is for research use only and not for use in

diagnostic or therapeutic procedures.

### **Protein Information**

Name UPF1 (HGNC:9962)

**Function** RNA-dependent helicase required for nonsense-mediated decay (NMD) of

aberrant mRNAs containing premature stop codons and modulates the

expression level of normal mRNAs (PubMed: 11163187, PubMed: 16086026, PubMed: 18172165, PubMed: 21145460, PubMed: 21419344, PubMed: 24726324). Is recruited to mRNAs upon translation termination and undergoes a cycle of phosphorylation and dephosphorylation; its phosphorylation appears to be a key step in NMD (PubMed: 11544179, PubMed:25220460). Recruited by release factors to stalled ribosomes together with the SMG1C protein kinase complex to form the transient SURF (SMG1-UPF1-eRF1-eRF3) complex (PubMed: 19417104). In EJC-dependent NMD, the SURF complex associates with the exon junction complex (EJC) (located 50-55 or more nucleotides downstream from the termination codon) through UPF2 and allows the formation of an UPF1-UPF2-UPF3 surveillance complex which is believed to activate NMD (PubMed:21419344). Phosphorylated UPF1 is recognized by EST1B/SMG5, SMG6 and SMG7 which are thought to provide a link to the mRNA degradation machinery involving exonucleolytic and endonucleolytic pathways, and to serve as adapters to protein phosphatase 2A (PP2A), thereby triggering UPF1 dephosphorylation and allowing the recycling of NMD factors (PubMed:12554878). UPF1 can also activate NMD without UPF2 or UPF3, and in the absence of the NMD-enhancing downstream EIC indicative for alternative NMD pathways (PubMed: 18447585). Plays a role in replication-dependent histone mRNA degradation at the end of phase S; the function is independent of UPF2 (PubMed:16086026, PubMed:18172165). For the recognition of premature termination codons (PTC) and initiation of NMD a competitive interaction between UPF1 and PABPC1 with the ribosome-bound release factors is proposed (PubMed: 18447585, PubMed: 25220460). The ATPase activity of UPF1 is required for disassembly of mRNPs undergoing NMD (PubMed:21145460). Together with UPF2 and dependent on TDRD6, mediates the degradation of mRNA harboring long 3'UTR by inducing the NMD machinery (By similarity). Also capable of unwinding double-stranded DNA and translocating on single-stranded DNA (PubMed:30218034).

**Cellular Location** 

Cytoplasm. Cytoplasm, P-body. Nucleus. Cytoplasm, perinuclear region {ECO:0000250|UniProtKB:Q9EPU0}. Note=Hyperphosphorylated form is targeted to the P-body, while unphosphorylated protein is distributed throughout the cytoplasm. Localized in the chromatoid bodies of round spermatids (By similarity). {ECO:0000250|UniProtKB:Q9EPU0}

**Tissue Location** 

Ubiquitous.

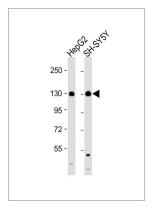
# **Background**

RENT1 is part of a post-splicing multiprotein complex involved in both mRNA nuclear export and mRNA surveillance. mRNA surveillance detects exported mRNAs with truncated open reading frames and initiates nonsense-mediated mRNA decay (NMD). When translation ends upstream from the last exon-exon junction, this triggers NMD to degrade mRNAs containing premature stop codons. This protein is located only in the cytoplasm. When translation ends, it interacts with the protein that is a functional homolog of yeast Upf2p to trigger mRNA decapping.

### References

Ohnishi, T., et al., Mol. Cell 12(5):1187-1200 (2003). Lykke-Andersen, J., Mol. Cell. Biol. 22(23):8114-8121 (2002). Carastro, L.M., et al., Nucleic Acids Res. 30(10):2232-2243 (2002). Mendell, J.T., et al., Science 298(5592):419-422 (2002). Serin, G., et al., Mol. Cell. Biol. 21(1):209-223 (2001).

## **Images**



All lanes: Anti-RENT1 Antibody (N-term E22) at 1:1000 dilution Lane 1: HepG2 whole cell lysate Lane 2: SH-SY5Y whole cell lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size: 124 kDa Blocking/Dilution buffer: 5% NFDM/TBST.

### **Citations**

- Amifostine Protects Mouse Liver Against Radiation-induced Autophagy Blockage.
- Two molecular pathways (NMD and ERAD) contribute to a genetic epilepsy associated with the GABA(A) receptor GABRA1 PTC mutation, 975delC, S326fs328X.

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