

## **THOC1 Antibody (N-term)**

Mouse Monoclonal Antibody (Mab)
Catalog # AM1950b

### **Specification**

## THOC1 Antibody (N-term) - Product Information

Application WB,E
Primary Accession O96FV9

Other Accession Q8R3N6, NP 005122.2

Reactivity
Predicted
Host
Clonality
Isotype
Calculated MW
Antigen Region

Human
Mouse
Mouse
Mouse
Monoclonal
IgG3,k
75666
257-285

# THOC1 Antibody (N-term) - Additional Information

### **Gene ID 9984**

### **Other Names**

THO complex subunit 1, Tho1, Nuclear matrix protein p84, p84N5, hTREX84, THOC1, HPR1

## Target/Specificity

This THOC1 antibody is generated from mice immunized with a KLH conjugated synthetic peptide between 257-285 amino acids from the N-terminal region of human THOC1.

### **Dilution**

WB~~1:500~1000

 $E\sim\sim$ Use at an assay dependent concentration.

#### **Format**

Purified monoclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein G column, 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**

THOC1 Antibody (N-term) is for research use only and not for use in diagnostic or therapeutic procedures.

# THOC1 Antibody (N-term) - Protein Information

### Name THOC1



## Synonyms HPR1

Function Component of the THO subcomplex of the TREX complex which is thought to couple mRNA transcription, processing and nuclear export, and which specifically associates with spliced mRNA and not with unspliced pre-mRNA (PubMed:15833825, PubMed:15998806, PubMed:17190602). Required for efficient export of polyadenylated RNA (PubMed:23222130). The THOC1-THOC2-THOC3 core complex alone is sufficient to bind export factor NXF1-NXT1 and promote ATPase activity of DDX39B/UAP56 (PubMed:33191911). TREX is recruited to spliced mRNAs by a transcription-independent mechanism, binds to mRNA upstream of the exon-junction complex (EJC) and is recruited in a splicing- and cap- dependent manner to a region near the 5' end of the mRNA where it functions in mRNA export to the cytoplasm via the TAP/NXF1 pathway (PubMed:15833825, PubMed:15998806, PubMed:17190602). Regulates transcriptional elongation of a subset of genes (PubMed:22144908). Involved in genome stability by preventing co-transcriptional R-loop formation (By similarity). May play a role in hair cell formation, hence may be involved in hearing (By similarity).

#### **Cellular Location**

[Isoform 1]: Nucleus speckle. Nucleus, nucleoplasm. Nucleus matrix. Cytoplasm. Note=Can shuttle between the nucleus and cytoplasm. Nuclear localization is required for induction of apoptotic cell death. Translocates to the cytoplasm during the early phase of apoptosis execution

#### **Tissue Location**

Ubiquitous. Expressed in various cancer cell lines. Expressed at very low levels in normal breast epithelial cells and highly expressed in breast tumors. Expression is strongly associated with an aggressive phenotype of breast tumors and expression correlates with tumor size and the metastatic state of the tumor progression

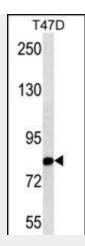
## **THOC1 Antibody (N-term) - Protocols**

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

### THOC1 Antibody (N-term) - Images





THOC1 Antibody (N-term) (Cat. #AM1950b) western blot analysis in T47D cell line lysates (35µg/lane). This demonstrates the THOC1 antibody detected the THOC1 protein (arrow).

# THOC1 Antibody (N-term) - Background

HPR1 is part of the TREX (transcription/export) complex, which includes TEX1 (MIM 606929), THO2 (MIM 300395), ALY (MIM 604171), and UAP56 (MIM 142560).

# **THOC1 Antibody (N-term) - References**

Davila, S., et al. Genes Immun. 11(3):232-238(2010) Liu, Y., et al. Mol. Psychiatry (2010) In press: Boyne, J.R., et al. PLoS Pathog. 4 (10), E1000194 (2008): Ferreira, M.A., et al. Nat. Genet. 40(9):1056-1058(2008) Yang, J., et al. Ann. Clin. Lab. Sci. 38(2):105-112(2008)