

Anti-TREX1 Rabbit Monoclonal Antibody

Catalog # ABO15326

Product Information

Application	WB, IHC, IF, ICC
Primary Accession	Q9NSU2
Host	Rabbit
Isotype	IgG
Reactivity	Human
Clonality	Monoclonal
Format	Liquid
Description	Anti-TREX1 Rabbit Monoclonal Antibody . Tested in WB, IHC, ICC/IF applications. This antibody reacts with Human.

Additional Information

Gene ID	11277
Other Names	Three-prime repair exonuclease 1, 3.1.11.2, 3'-5' exonuclease TREX1, Deoxyribonuclease III, DNase III, TREX1 {ECO:0000303 PubMed:10391904, ECO:0000312 HGNC:HGNC:12269}
Calculated MW	33212
Application Details	WB 1:500-1:2000 IHC 1:50-1:200 ICC/IF 1:50-1:200
Source	Eukaryota
Contents	Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol, 0.4-0.5mg/ml BSA.
Clone Names	Clone: 19T02
Immunogen	A synthesized peptide derived from human TREX1
Purification	Affinity-chromatography
Storage	Store at -20°C for one year. For short term storage and frequent use, store at 4°C for up to one month. Avoid repeated freeze-thaw cycles.

Protein Information

Name	TREX1 {ECO:0000303 PubMed:10391904, ECO:0000312 HGNC:HGNC:12269}
Function	Major cellular 3'-to-5' DNA exonuclease which digests single- stranded DNA (ssDNA) and double-stranded DNA (dsDNA) with mismatched 3' termini

(PubMed:[10391904](#), PubMed:[10393201](#), PubMed:[17293595](#)). Prevents cell-intrinsic initiation of autoimmunity (PubMed:[10391904](#), PubMed:[10393201](#), PubMed:[17293595](#)). Acts by metabolizing DNA fragments from endogenous retroelements, including L1, LTR and SINE elements (PubMed:[10391904](#), PubMed:[10393201](#), PubMed:[17293595](#)). Plays a key role in degradation of DNA fragments at cytosolic micronuclei arising from genome instability: its association with the endoplasmic reticulum membrane directs TREX1 to ruptured micronuclei, leading to micronuclear DNA degradation (PubMed:[33476576](#)). Micronuclear DNA degradation is required to limit CGAS activation and subsequent inflammation (PubMed:[33476576](#)). Unless degraded, these DNA fragments accumulate in the cytosol and activate the cGAS-STING innate immune signaling, leading to the production of type I interferon (PubMed:[33476576](#)). Prevents chronic ATM-dependent checkpoint activation, by processing ssDNA polynucleotide species arising from the processing of aberrant DNA replication intermediates (PubMed:[18045533](#)). Inefficiently degrades oxidized DNA, such as that generated upon antimicrobial reactive oxygen production or upon absorption of UV light (PubMed:[23993650](#)). During GZMA-mediated cell death, contributes to DNA damage in concert with NME1 (PubMed:[16818237](#)). NME1 nicks one strand of DNA and TREX1 removes bases from the free 3' end to enhance DNA damage and prevent DNA end reannealing and rapid repair (PubMed:[16818237](#)).

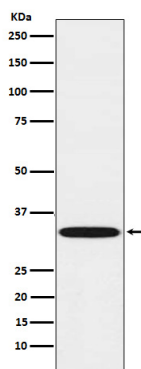
Cellular Location

Nucleus. Cytoplasm, cytosol. Endoplasmic reticulum membrane; Peripheral membrane protein. Note=Retained in the cytoplasm through the C-terminal region (By similarity). Localization to the endoplasmic reticulum membrane is required to direct TREX1 to ruptured micronuclei (PubMed:[33476576](#)). In response to DNA damage, translocates to the nucleus where it is specifically recruited to replication foci (PubMed:[16818237](#)). Translocation to the nucleus also occurs during GZMA-mediated cell death (PubMed:[16818237](#)) {ECO:0000250|UniProtKB:Q91XB0, ECO:0000269|PubMed:[16818237](#), ECO:0000269|PubMed:[33476576](#)}

Tissue Location

Detected in thymus, spleen, liver, brain, heart, small intestine and colon.

Images



Western blot analysis of TREX1 expression in Daudi cell lysate.

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