

TRIM7 Antibody (N-term) Blocking peptide
Synthetic peptide
Catalog # BP11979a**Specification**

TRIM7 Antibody (N-term) Blocking peptide - Product InformationPrimary Accession [O9C029](#)**TRIM7 Antibody (N-term) Blocking peptide - Additional Information****Gene ID** 81786**Other Names**

Tripartite motif-containing protein 7, Glycogenin-interacting protein, RING finger protein 90, TRIM7, GNIP, RNF90

Format

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

Precautions

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

TRIM7 Antibody (N-term) Blocking peptide - Protein Information**Name** TRIM7**Synonyms** GNIP, RNF90**Function**

E3 ubiquitin-protein ligase that have both tumor-promoting and tumor-suppressing activities and functions in several biological processes including innate immunity, regulation of ferroptosis as well as cell proliferation and migration (PubMed: [25851810](http://www.uniprot.org/citations/25851810), PubMed: [32853985](http://www.uniprot.org/citations/32853985), PubMed: [34062120](http://www.uniprot.org/citations/34062120)). Acts as an antiviral effector against multiple viruses by targeting specific viral proteins for ubiquitination and degradation including norovirus NTPase protein or SARS-CoV-2 NSP5 and NSP8 proteins (PubMed: [34062120](http://www.uniprot.org/citations/34062120), PubMed: [35982226](http://www.uniprot.org/citations/35982226)). Mechanistically, recognizes the C-terminal glutamine-containing motif usually generated by viral proteases that process the polyproteins and trigger their ubiquitination and subsequent degradation (PubMed: [35982226](http://www.uniprot.org/citations/35982226), PubMed: [35867826](http://www.uniprot.org/citations/35867826), PubMed: [35893676](http://www.uniprot.org/citations/35893676)).

target="_blank">35893676). Mediates 'Lys-63'-linked polyubiquitination and stabilization of the JUN coactivator RNF187 in response to growth factor signaling via the MEK/ERK pathway, thereby regulating JUN transactivation and cellular proliferation (PubMed:25851810). Promotes the TLR4-mediated signaling activation through its E3 ligase domain leading to production of pro-inflammatory cytokines and type I interferon (By similarity). Also plays a negative role in the regulation of exogenous cytosolic DNA virus-triggered immune response. Mechanistically, enhances the 'Lys-48'-linked ubiquitination of STING1 leading to its proteasome-dependent degradation (PubMed:32126128). Mediates the ubiquitination of the SIN3- HDAC chromatin remodeling complex component BRMS1 (PubMed:32853985). Modulates NCOA4-mediated ferritinophagy and ferroptosis in glioblastoma cells by ubiquitinating NCOA4, leading to its degradation (PubMed:36067704).

Cellular Location

Nucleus. Cytoplasm. Golgi apparatus

Tissue Location

Skeletal muscle and placenta, at lower levels in heart, brain and pancreas. Isoform 1 is widely expressed with high level in testis, kidney and heart.

TRIM7 Antibody (N-term) Blocking peptide - Protocols

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

- [Blocking Peptides](#)

TRIM7 Antibody (N-term) Blocking peptide - Images

TRIM7 Antibody (N-term) Blocking peptide - Background

The protein encoded by this gene is a member of the tripartite motif (TRIM) family. The TRIM motif includes three zinc-binding domains, a RING, a B-box type 1, a B-box type 2, and a coiled-coil region. The protein localizes to both the nucleus and the cytoplasm, and may represent a participant in the initiation of glycogen synthesis. Multiple transcript variants have been found for this gene, and some of them encode the same isoform. [provided by RefSeq].

TRIM7 Antibody (N-term) Blocking peptide - References

Zhai, L., et al. Arch. Biochem. Biophys. 421(2):236-242(2004) Skurat, A.V., et al. J. Biol. Chem. 277(22):19331-19338(2002) Reymond, A., et al. EMBO J. 20(9):2140-2151(2001)