

## Anti-TMEM173 Monoclonal Antibody Catalog # ABO14597

### Specification

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#### Anti-TMEM173 Monoclonal Antibody - Product Information

Application	WB, IF, ICC, FC
Primary Accession	<a href="#">Q86WV6</a>
Host	Rabbit
Isotype	Rabbit IgG
Reactivity	Human
Clonality	Monoclonal
Format	Liquid

#### Description

Anti-TMEM173 Monoclonal Antibody . Tested in WB, ICC/IF, Flow Cytometry applications. This antibody reacts with Human.

#### Anti-TMEM173 Monoclonal Antibody - Additional Information

**Gene ID** 340061

#### Other Names

Stimulator of interferon genes protein, hSTING, Endoplasmic reticulum interferon stimulator, ERIS, Mediator of IRF3 activation, hMITA, Transmembrane protein 173, STING1 (<a href="[http://www.genenames.org/cgi-bin/gene\\_symbol\\_report?hgnc\\_id=27962](http://www.genenames.org/cgi-bin/gene_symbol_report?hgnc_id=27962)" target="\_blank">HGNC:27962</a>)

#### Application Details

WB 1:500-1:2000<br>ICC/IF 1:100-1:500<br>FC 1:20

#### 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.

#### Immunogen

A synthesized peptide derived from human TMEM173 Facilitator of innate immune signaling that promotes the production of type I interferon (IFN-alpha and IFN-beta). Innate immune response is triggered in response to non-CpG double-stranded DNA from viruses and bacteria delivered to the cytoplasm.

#### 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.**

#### Anti-TMEM173 Monoclonal Antibody - Protein Information

Name STING1 ([HGNC:27962](#))

## Function

Facilitator of innate immune signaling that acts as a sensor of cytosolic DNA from bacteria and viruses and promotes the production of type I interferon (IFN-alpha and IFN-beta) (PubMed:[18724357](http://www.uniprot.org/citations/18724357)</a>, PubMed:[18818105](http://www.uniprot.org/citations/18818105)</a>, PubMed:[19433799](http://www.uniprot.org/citations/19433799)</a>, PubMed:[19776740](http://www.uniprot.org/citations/19776740)</a>, PubMed:[23027953](http://www.uniprot.org/citations/23027953)</a>, PubMed:[23747010](http://www.uniprot.org/citations/23747010)</a>, PubMed:[23910378](http://www.uniprot.org/citations/23910378)</a>, PubMed:[27801882](http://www.uniprot.org/citations/27801882)</a>, PubMed:[29973723](http://www.uniprot.org/citations/29973723)</a>, PubMed:[30842659](http://www.uniprot.org/citations/30842659)</a>, PubMed:[35045565](http://www.uniprot.org/citations/35045565)</a>, PubMed:[35388221](http://www.uniprot.org/citations/35388221)</a>, PubMed:[36808561](http://www.uniprot.org/citations/36808561)</a>, PubMed:[37832545](http://www.uniprot.org/citations/37832545)</a>, PubMed:[25704810](http://www.uniprot.org/citations/25704810)</a>, PubMed:[39255680](http://www.uniprot.org/citations/39255680)</a>). Innate immune response is triggered in response to non-CpG double-stranded DNA from viruses and bacteria delivered to the cytoplasm (PubMed:[26300263](http://www.uniprot.org/citations/26300263)</a>). Acts by binding cyclic dinucleotides: recognizes and binds cyclic di-GMP (c-di-GMP), a second messenger produced by bacteria, cyclic UMP-AMP (2',3'-cUAMP), and cyclic GMP-AMP (cGAMP), a messenger produced by CGAS in response to DNA virus in the cytosol (PubMed:[21947006](http://www.uniprot.org/citations/21947006)</a>, PubMed:[23258412](http://www.uniprot.org/citations/23258412)</a>, PubMed:[23707065](http://www.uniprot.org/citations/23707065)</a>, PubMed:[23722158](http://www.uniprot.org/citations/23722158)</a>, PubMed:[23747010](http://www.uniprot.org/citations/23747010)</a>, PubMed:[23910378](http://www.uniprot.org/citations/23910378)</a>, PubMed:[26229117](http://www.uniprot.org/citations/26229117)</a>, PubMed:[30842659](http://www.uniprot.org/citations/30842659)</a>, PubMed:[35388221](http://www.uniprot.org/citations/35388221)</a>, PubMed:[37379839](http://www.uniprot.org/citations/37379839)</a>). Upon binding to c-di-GMP, cUAMP or cGAMP, STING1 oligomerizes, translocates from the endoplasmic reticulum and is phosphorylated by TBK1 on the pLxIS motif, leading to recruitment and subsequent activation of the transcription factor IRF3 to induce expression of type I interferon and exert a potent anti-viral state (PubMed:[22394562](http://www.uniprot.org/citations/22394562)</a>, PubMed:[25636800](http://www.uniprot.org/citations/25636800)</a>, PubMed:[29973723](http://www.uniprot.org/citations/29973723)</a>, PubMed:[30842653](http://www.uniprot.org/citations/30842653)</a>, PubMed:[35045565](http://www.uniprot.org/citations/35045565)</a>, PubMed:[35388221](http://www.uniprot.org/citations/35388221)</a>). Exhibits 2',3' phosphodiester linkage-specific ligand recognition: can bind both 2'-3' linked cGAMP (2'-3'-cGAMP) and 3'-3' linked cGAMP but is preferentially activated by 2'-3' linked cGAMP (PubMed:[23747010](http://www.uniprot.org/citations/23747010)</a>, PubMed:[23910378](http://www.uniprot.org/citations/23910378)</a>, PubMed:[26300263](http://www.uniprot.org/citations/26300263)</a>). The preference for 2'-3'-cGAMP, compared to other linkage isomers is probably due to the ligand itself, whichs adopts an organized free- ligand conformation that resembles the STING1-bound conformation and pays low energy costs in changing into the active conformation (PubMed:[26150511](http://www.uniprot.org/citations/26150511)</a>). In addition to promote the production of type I interferons, plays a direct role in autophagy (PubMed:[30568238](http://www.uniprot.org/citations/30568238)</a>, PubMed:[30568238](http://www.uniprot.org/citations/30568238)</a>).

Following cGAMP-binding, STING1 buds from the endoplasmic reticulum into COPII vesicles, which then form the endoplasmic reticulum-Golgi intermediate compartment (ERGIC) (PubMed:<a href="http://www.uniprot.org/citations/30842662" target="\_blank">30842662</a>). The ERGIC serves as the membrane source for WIPI2 recruitment and LC3 lipidation, leading to formation of autophagosomes that target cytosolic DNA or DNA viruses for degradation by the lysosome (PubMed:<a href="http://www.uniprot.org/citations/30842662" target="\_blank">30842662</a>). Promotes autophagy by acting as a proton channel that directs proton efflux from the Golgi to facilitate MAP1LC3B/LC3B lipidation (PubMed:<a href="http://www.uniprot.org/citations/37535724" target="\_blank">37535724</a>). The autophagy- and interferon-inducing activities can be uncoupled and autophagy induction is independent of TBK1 phosphorylation (PubMed:<a href="http://www.uniprot.org/citations/30568238" target="\_blank">30568238</a>, PubMed:<a href="http://www.uniprot.org/citations/30842662" target="\_blank">30842662</a>). Autophagy is also triggered upon infection by bacteria: following c-di-GMP-binding, which is produced by live Gram- positive bacteria, promotes reticulophagy (By similarity). May be involved in translocon function, the translocon possibly being able to influence the induction of type I interferons (PubMed:<a href="http://www.uniprot.org/citations/18724357" target="\_blank">18724357</a>). May be involved in transduction of apoptotic signals via its association with the major histocompatibility complex class II (MHC-II) (By similarity).

### Cellular Location

Endoplasmic reticulum membrane; Multi-pass membrane protein {ECO:0000255, ECO:0000269|PubMed:30842659, ECO:0000269|PubMed:32690950}. Cytoplasm, perinuclear region. Endoplasmic reticulum-Golgi intermediate compartment membrane; Multi-pass membrane protein {ECO:0000255, ECO:0000269|PubMed:32690950}. Golgi apparatus membrane; Multi-pass membrane protein. Cytoplasmic vesicle, autophagosome membrane; Multi-pass membrane protein. Mitochondrion outer membrane; Multi-pass membrane protein. Cell membrane {ECO:0000250|UniProtKB:Q3TBT3}; Multi-pass membrane protein. Note=In response to double-stranded DNA stimulation, translocates from the endoplasmic reticulum through the endoplasmic reticulum-Golgi intermediate compartment and Golgi to post-Golgi vesicles, where the kinase TBK1 is recruited (PubMed:19433799, PubMed:29694889, PubMed:30842653, PubMed:30842659). Upon cGAMP-binding, translocates to the endoplasmic reticulum-Golgi intermediate compartment (ERGIC) in a process that is dependent on COPII vesicles; STING1-containing ERGIC serves as a membrane source for LC3 lipidation, which is a key step in autophagosome biogenesis (PubMed:30842662, PubMed:37832545). Localizes in the lysosome membrane in a TMEM203- dependent manner (By similarity). {ECO:0000250|UniProtKB:Q3TBT3, ECO:0000269|PubMed:19433799, ECO:0000269|PubMed:29694889, ECO:0000269|PubMed:30842653, ECO:0000269|PubMed:30842659, ECO:0000269|PubMed:30842662, ECO:0000269|PubMed:32690950, ECO:0000269|PubMed:37832545}

### Tissue Location

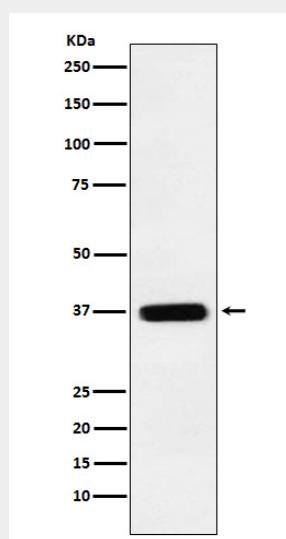
Ubiquitously expressed (PubMed:18724357, PubMed:18818105). Expressed in skin endothelial cells, alveolar type 2 pneumocytes, bronchial epithelium and alveolar macrophages (PubMed:25029335).

### Anti-TMEM173 Monoclonal Antibody - Protocols

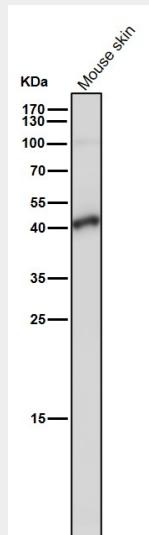
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)

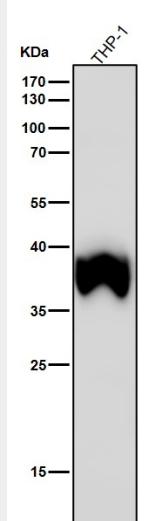
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

**Anti-TMEM173 Monoclonal Antibody - Images**

Western blot analysis of TMEM173 expression in HeLa cell lysate.



All lanes use the Antibody at 1:4K dilution for 1 hour at room temperature.



All lanes use the Antibody at 1:4K dilution for 1 hour at room temperature.