

## **TTC35 Antibody (Center)**

Affinity Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP18405c

## **Specification**

Reactivity

## TTC35 Antibody (Center) - Product Information

Application WB,E
Primary Accession Q15006

Other Accession <u>BOBNGO</u>, <u>Q9CRD2</u>, <u>Q5E993</u>, <u>Q8AVU9</u>, <u>Q6INS3</u>,

NP\_055488.1 Human, Mouse

65-91

Predicted Xenopus, Bovine, Rat

Host Rabbit
Clonality Polyclonal
Isotype Rabbit IgG
Calculated MW 34834

## TTC35 Antibody (Center) - Additional Information

#### **Gene ID 9694**

Antigen Region

#### **Other Names**

ER membrane protein complex subunit 2, Tetratricopeptide repeat protein 35, TPR repeat protein 35, EMC2, KIAA0103, TTC35

## Target/Specificity

This TTC35 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 65-91 amino acids from the Central region of human TTC35.

# **Dilution**

WB~~1:1000

## **Format**

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

## **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**

TTC35 Antibody (Center) is for research use only and not for use in diagnostic or therapeutic procedures.

#### TTC35 Antibody (Center) - Protein Information

Name EMC2 (<u>HGNC:28963</u>)



**Function** Part of the endoplasmic reticulum membrane protein complex (EMC) that enables the energy-independent insertion into endoplasmic reticulum membranes of newly synthesized membrane proteins (PubMed:29242231, PubMed:29809151, PubMed:30415835, PubMed:32439656, PubMed:32459176, PubMed:33964204). Preferentially accommodates proteins with transmembrane domains that are weakly hydrophobic or contain destabilizing features such as charged and aromatic residues (PubMed:20242231, PubMed:20800151, PubMed:30415835)

with transmembrane domains that are weakly hydrophobic or contain destabilizing features such as charged and aromatic residues (PubMed:29242231, PubMed:29809151, PubMed:30415835). Involved in the cotranslational insertion of multi-pass membrane proteins in which stop-transfer membrane-anchor sequences become ER membrane spanning helices (PubMed:29809151, PubMed:30415835). It is also required for the post-translational insertion of tail-anchored/TA proteins in endoplasmic reticulum membranes (PubMed:29242231, PubMed:29809151). By mediating the proper cotranslational insertion of N-terminal transmembrane domains in an N-exo topology, with translocated N- terminus in the lumen of the ER, controls the topology of multi-pass membrane proteins like the G protein-coupled receptors (PubMed:30415835). By regulating the insertion of various proteins in membranes, it is indirectly involved in many cellular processes (Probable).

#### **Cellular Location**

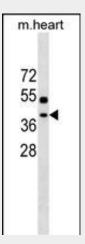
Endoplasmic reticulum membrane; Peripheral membrane protein; Cytoplasmic side Note=May also localize to the nuclear envelope {ECO:0000250|UniProtKB:Q9CRD2}

#### TTC35 Antibody (Center) - Protocols

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

- Western Blot
- Blocking Peptides
- Dot Blot
- <u>Immunohistochemistry</u>
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

## TTC35 Antibody (Center) - Images



TTC35 Antibody (Center) (Cat. #AP18405c) western blot analysis in mouse heart tissue lysates (35ug/lane). This demonstrates the TTC35 Antibody detected the TTC35 protein (arrow).

## TTC35 Antibody (Center) - Background





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TTC35 is also known as TPR repeat protein 35. TPR domains consist of a variable number of degenerate tandem 34 amino acid repeats. TPR domains have been suggested to have a variety of functions in proteins in various subcellular compartments and appear to function as targeting domains, mediating specific protein-protein interactions.

# TTC35 Antibody (Center) - References

Lamesch, P., et al. Genomics 89(3):307-315(2007) Dreger, M., et al. Proc. Natl. Acad. Sci. U.S.A. 98(21):11943-11948(2001) Hoja, M.R., et al. Exp. Cell Res. 259(1):239-246(2000)