# TOMM40 Rabbit mAb

Catalog No.: A24644 Recombinant



### **Basic Information**

### **Observed MW**

38kDa/39,41kDa

### **Calculated MW**

38kDa

### Category

Primary antibody

### **Applications**

WB,IP,IHC-P,IF/ICC,ELISA

#### **Cross-Reactivity**

Human, Mouse, Rat

#### CloneNo number

ARC63318

# **Background**

The protein encoded by this gene is localized in the outer membrane of the mitochondria. It is the channel-forming subunit of the translocase of the mitochondrial outer membrane (TOM) complex that is essential for import of protein precursors into mitochondria. Alternatively spliced transcript variants have been found for this gene.

### **Recommended Dilutions**

**WB** 1:2000 - 1:10000

IP 0.5μg-4μg antibody for 400μg-600μg extracts of

whole cells

IHC-P 1:100 - 1:500

**IF/ICC** 1:100 - 1:500

**ELISA** Recommended starting

concentration is 1 µg/mL.

Please optimize the
concentration based on
your specific assay
requirements.

### **Contact**

<u>a</u>	400-999-6126
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•	www.abclonal.com.cn

# **Immunogen Information**

**Gene ID Swiss Prot** 10452 096008

#### **Immunogen**

Recombinant protein (or fragment). This information is considered to be commercially sensitive.

### Synonyms

TOM40; PEREC1; C19orf1; PER-EC1; D19S1177E; TOMM40

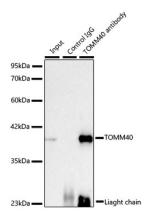
### **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

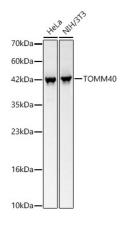
#### Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS containing 50% glycerol and 0.05% BSA, preserved with proclin300 or sodium azide (as specified on the Certificate of Analysis), pH 7.3.



Immunoprecipitation of TOMM40 from 500  $\mu g$  extracts of Hep G2 cells was performed using 2  $\mu g$  of TOMM40 Rabbit mAb (A24644). Rabbit Control IgG (AC005) was used to precipitate the Control IgG sample. IP samples were eluted with 1X Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using TOMM40 Rabbit mAb (A24644) at a dilution of 1:5000.

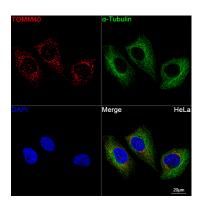


Western blot analysis of various lysates, using TOMM40 Rabbit mAb (A24644) at 1:4800 dilution. Secondary antibody: HRP-conjugated Goat anti-Rabbit lgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins:  $25\mu g$  per lane.

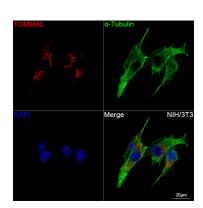
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

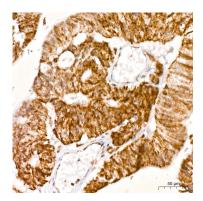
Exposure time: 5s.



Confocal imaging of HeLa cells using TOMM40 Rabbit mAb (A24644,at dilution of 1:300) (red). The cells were counterstained with  $\alpha$ -Tubulin Mouse mAb (AC012,dilution 1:400) (Green). DAPI was used for nuclear staining (blue). Objective: 100x.

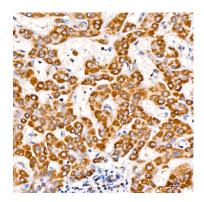


Confocal imaging of NIH/3T3 cells using TOMM40 Rabbit mAb (A24644,at dilution of 1:300) (Red). The cells were counterstained with  $\alpha$ -Tubulin Mouse mAb (AC012,dilution 1:400) (Green). DAPI was used for nuclear staining (blue). Objective: 100x.

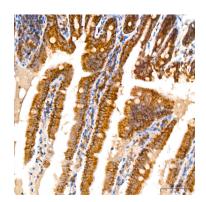


Immunohistochemistry analysis of paraffinembedded Human colon carcinoma using TOMM40 Rabbit mAb (A24644) at dilution of 1:400 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.

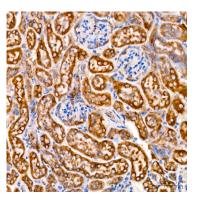
## **Validation Data**



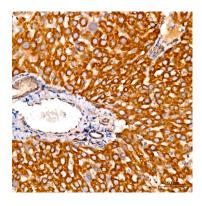
Immunohistochemistry analysis of paraffinembedded Human liver using TOMM40 Rabbit mAb (A24644) at dilution of 1:400 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Mouse colon using TOMM40 Rabbit mAb (A24644) at dilution of 1:400 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Mouse kidney using TOMM40 Rabbit mAb (A24644) at dilution of 1:400 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Rat liver using TOMM40 Rabbit mAb (A24644) at dilution of 1:400 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.