

# [KO Validated] TTC11/FIS1 Rabbit mAb

Catalog No.: A19666 KO Validated Recombinant 20 Publications

## **Basic Information**

## **Observed MW**

15kDa

#### **Calculated MW**

17kDa

### Category

Primary antibody

## **Applications**

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

### **Cross-Reactivity**

Human, Mouse, Rat

#### CloneNo number

ARC5010-03

## **Background**

Enables identical protein binding activity. Involved in several processes, including calcium-mediated signaling using intracellular calcium source; cellular calcium ion homeostasis; and mitochondrion organization. Acts upstream of or within mitochondrion morphogenesis. Located in mitochondrion and peroxisome. Is integral component of mitochondrial outer membrane and integral component of peroxisomal membrane. Part of protein-containing complex. Biomarker of Alzheimer's disease.

## **Recommended Dilutions**

**WB** 1:1000 - 1:6000

IHC-P 1:200 - 1:800

**IF/ICC** 1:100 - 1:400

**IP** 0.5μg-4μg antibody for

200µg-400µg extracts of

whole cells

**ELISA** Recommended starting

concentration is 1 µg/mL.

Please optimize the
concentration based on
your specific assay
requirements.

## **Contact**

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•		www.abclonal.com.cn

## Immunogen Information

 Gene ID
 Swiss Prot

 51024
 Q9Y3D6

### **Immunogen**

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

## **Synonyms**

TTC11; CGI-135; S1

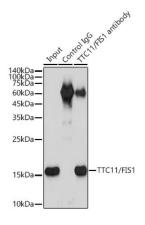
## **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

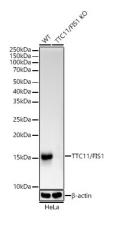
### Storage

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 TTC11/FIS1 from 300  $\mu$ g extracts of HeLa cells was performed using 3  $\mu$ g of TTC11/FIS1 antibody (A19666). Rabbit IgG isotype control (AC042) was used to precipitate the Control IgG sample. IP samples were eluted with 1X reducing Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using TTC11/FIS1 (A19666) at a dilution of 1:2000.

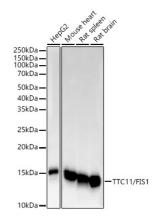


Western blot analysis of lysates from wild type (WT) and TTC11/FIS1 knockout (KO) HeLa(KO) cells, using [KO Validated] TTC11/FIS1 Rabbit mAb (A19666) at 1:1000 dilution incubated overnight at 4°C. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25 $\mu$ g per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 10s.



Western blot analysis of various lysates, using [KO Validated] TTC11/FIS1 Rabbit mAb (A19666) at 1:1000 dilution incubated overnight at  $4^{\circ}$ C.

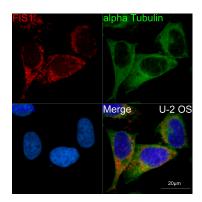
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

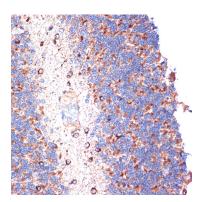
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

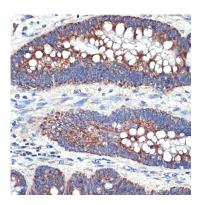
Exposure time: 10s.



Confocal imaging of U-2 OS cells using [KO Validated] TTC11/FIS1 Rabbit mAb (A19666,dilution 1:100)(Red). The cells were counterstained with  $\alpha$ -Tubulin Mouse mAb (AC012,dilution 1:400) (Green). DAPI was used for nuclear staining (blue). Objective:



Immunohistochemistry analysis of paraffinembedded Rat brain tissue using [KO Validated] TTC11/FIS1 Rabbit mAb (A19666) at a dilution of 1:100 (40x lens). Microwave antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Human colon carcinoma tissue using [KO Validated] TTC11/FIS1 Rabbit mAb (A19666) at a dilution of 1:100 (40x lens). Microwave antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.