# **TREX1 Rabbit mAb**

Catalog No.: A3819 Recombinant



### **Basic Information**

### **Observed MW**

33kDa/34kd

### **Calculated MW**

33kDa

### Category

SMab Recombinant Monoclonal Antibody

### **Applications**

WB,IP,ELISA

### **Cross-Reactivity**

Human

### CloneNo number

ARC0841

# **Background**

This gene encodes a nuclear protein with 3' exonuclease activity. The encoded protein may play a role in DNA repair and serve as a proofreading function for DNA polymerase. Mutations in this gene result in Aicardi-Goutieres syndrome, chilblain lupus, Cree encephalitis, and other diseases of the immune system. Alternative splicing results in multiple transcript variants.

# **Recommended Dilutions**

**WB** 1:500 - 1:2000

**IP** 0.5μg-4μg antibody for

200µg-400µg extracts of whole cells

Recommended starting concentration is 1

μg/mL. Please optimize the concentration based on your specific

assay requirements.

#### Contact

**ELISA** 

www.abclonal.com

## **Immunogen Information**

**Gene ID**Swiss Prot
11277
Q9NSU2

### **Immunogen**

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

### **Synonyms**

CRV; AGS1; DRN3; HERNS; RVCLS; TREX1

### **Product Information**

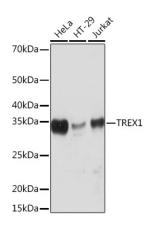
SourceIsotypePurificationRabbitIgGAffinity purification

#### Storage

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

Buffer: PBS with 0.02% sodium azide, 0.05% BSA, 50% glycerol, pH7.3.

### **Validation Data**



Western blot analysis of various lysates using TREX1 Rabbit mAb (A3819) at 1:1000 dilution.

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

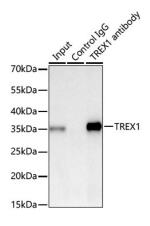
dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 10s.



Immunoprecipitation of TREX1 from 300  $\mu$ g extracts of HeLa cells was performed using 0.5  $\mu$ g of TREX1 Rabbit mAb (A3819). Rabbit IgG isotype control (AC005) 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 TREX1 Rabbit mAb (A3819) at a dilution of 1:1000.