# **CHM Rabbit pAb**

Catalog No.: A8345



# **Basic Information**

#### **Observed MW**

101kDa

### **Calculated MW**

73kDa

### Category

Primary antibody

### **Applications**

ELISA,WB

#### **Cross-Reactivity**

Human

# **Background**

This gene encodes component A of the RAB geranylgeranyl transferase holoenzyme. In the dimeric holoenzyme, this subunit binds unprenylated Rab GTPases and then presents them to the catalytic Rab GGTase subunit for the geranylgeranyl transfer reaction. Rab GTPases need to be geranylgeranyled on either one or two cysteine residues in their C-terminus to localize to the correct intracellular membrane. Mutations in this gene are a cause of choroideremia; also known as tapetochoroidal dystrophy (TCD). This X-linked disease is characterized by progressive dystrophy of the choroid, retinal pigment epithelium and retina. Alternatively spliced transcript variants have been found for this gene.

# **Recommended Dilutions**

**WB** 

1:500 - 1:2000

# Immunogen Information

Gene ID 1121 **Swiss Prot** 

P24386

#### **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 120-210 of human CHM (NP\_000381.1).

### **Synonyms**

TCD; GGTA; REP-1; DXS540; HSD-32; CHM

### **Contact**

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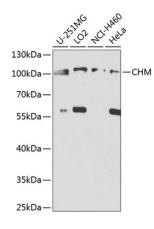
### **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

#### Storage

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

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



Western blot analysis of extracts of various cell lines, using CHM antibody (A8345) at 1:1000 dilution.\_Secondary antibody: HRP 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 Enhanced Kit (RM00021).\_Exposure time: 5s.