# **PHYH Rabbit pAb**

Catalog No.: A6304



# **Basic Information**

#### **Observed MW**

38kDa

### **Calculated MW**

39kDa

### Category

Primary antibody

## **Applications**

ELISA,WB

#### **Cross-Reactivity**

Human, Mouse

# **Background**

This gene is a member of the PhyH family and encodes a peroxisomal protein that is involved in the alpha-oxidation of 3-methyl branched fatty acids. Specifically, this protein converts phytanoyl-CoA to 2-hydroxyphytanoyl-CoA. Mutations in this gene have been associated with Refsum disease (RD) and deficient protein activity has been associated with Zellweger syndrome and rhizomelic chondrodysplasia punctata. Alternate transcriptional splice variants, encoding different isoforms, have been characterized.

# **Recommended Dilutions**

**WB** 

1:500 - 1:2000

# **Immunogen Information**

Gene ID 5264 **Swiss Prot** 

014832

#### **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 1-338 of human PHYH (NP\_006205.1).

## **Synonyms**

RD; LN1; PAHX; LNAP1; PHYH1; PHYH

## **Contact**

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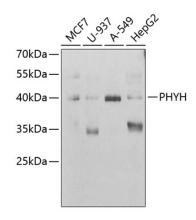
## **Product Information**

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 PHYH antibody (A6304) at 1:1000 dilution. Secondary antibody: HRP 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 Enhanced Kit (RM00021).

Exposure time: 90s.