

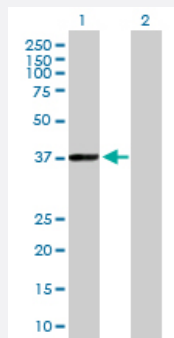
MaxPab®

PHYH purified MaxPab mouse polyclonal antibody (B01P)

Catalog # H00005264-B01P

Size 50 ug

Applications



Western Blot (Transfected lysate)

Western Blot analysis of PHYH expression in transfected 293T cell line ([H00005264-T01](#)) by PHYH MaxPab polyclonal antibody.

Lane 1: PHYH transfected lysate(37.18 kDa).

Lane 2: Non-transfected lysate.

Specification

Product Description

Mouse polyclonal antibody raised against a full-length human PHYH protein.

Immunogen

PHYH (NP_006205.1, 1 a.a. ~ 338 a.a) full-length human protein.

Sequence

MEQLRAAARLQIVLGHILGRPSAGAVVAHPTSGTISSASFHPQQFQYTLDDNNVLTLEQRKFYEENG
FLVIKNLVPDADIQRFRNEFEKICRKEVKPLGLTVMRDVTISKSEYAPSEKMITKVQDFQEDKELFR
YCTLPEILKYVECFTGPNIMAMHTMLINKPPDSGKKTSRHPLHQDLHYFPFRPSDLVCAWTAMEHI
SRNNGCLVVLPGTHKGSCLKPHDYKWEKGVNKMFGIQQDYEEKARVHLVMEKGDVFFHPLLI
HSGSQNKTKGFRKAISCHFASADCHYDVKGTSQENIEKEVVGIAHKFFGAENSVNLKDWMFRA
RLVKGERTNL

Host

Mouse

Reactivity

Human

Interspecies Antigen Sequence

Mouse (78); Rat (78)

Quality Control Testing

Antibody reactive against mammalian transfected lysate.

Storage Buffer

In 1x PBS, pH 7.4

Storage Instruction

Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot (Transfected lysate)

Western Blot analysis of PHYH expression in transfected 293T cell line ([H00005264-T01](#)) by PHYH MaxPab polyclonal antibody.

Lane 1: PHYH transfected lysate(37.18 KDa).

Lane 2: Non-transfected lysate.

[Protocol Download](#)

Gene Info — PHYH

Entrez GeneID [5264](#)

GeneBank Accession# [NM_006214](#)

Protein Accession# [NP_006205.1](#)

Gene Name PHYH

Gene Alias LN1, LNAP1, PAHX, PHYH1, RD

Gene Description phytanoyl-CoA 2-hydroxylase

Omim ID [266500 602026](#)

Gene Ontology [Hyperlink](#)

Gene Summary 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. [provided by RefSeq]

Other Designations phytanic acid oxidase|phytanoyl-CoA alpha hydroxylase|phytanoyl-CoA 2 oxoglutarate dioxygenase|phytanoyl-CoA alpha-hydroxylase|phytanoyl-CoA dioxygenase, peroxisomal|phytanoyl-CoA hydroxylase (Refsum disease)

Disease

- [Alzheimer Disease](#)

- [Genetic Predisposition to Disease](#)