

DNAXPAb

Hard-to-Find
Antibody

PLOD2 DNAXPab

Catalog # H00005352-W01P

Size 200 ug

Specification

Product Description Rabbit polyclonal antibody raised against a full-length human PLOD2 DNA using DNAX™ Immune technology.

Technology [DNAX™ Immune](#)

Immunogen Full-length human DNA

Sequence

MGGCTVKPQLLLLALVLHPWNPCLGADSEKPSSIPTDKLLVITVATKESDGFHRFMQSAKYFNYT
VKVLGQGEEWRGGDGINSIGGGQKVRMLKEVMEHYADQDDLVMFTECFDVIFAGGPPEVLKK
FQKANHKVVFAADGILWPDKRLADKYPVVHIGKRYLNSGGFIGYAPYVNRVQQWNLQDNDQDQ
FYTKVYIDPLKREAINITLDHKCKIFQTLNGAVDEVVLKFENGKARAKNTFYETLPVAINGNGPTKILL
NYFGNYVPNSWTQDNGCTLCEFDTVDL SAVDVHPNVSIGVFIEQPTPFLPRFLDILLTLDYPKEAL
KLFIHNKEVYHEKDIKVFFDKAKHEIKTIKVGPEENLSQAEARNMGMDFCRQDEKCDYFVSVDAD
VVLTNPRTLKILIEQNRKIAPLVTRHGKLWSNFWGALSPDGYARSEDYVDVQGNRVGVWNPY
MANVYLIKGKTLRSEMNERNYFVRDKLDPDMALCRNAREMTLQREKDSPTPETFQMLSPPKGVF
MYISNRHEFGRLSTANYNTSHYNNDLWQIFENPVDWKEKYINRDYSKIFTENVEQPCPDVFWFPI
FSEKACDELVEEMEHYGKWSGGKHHSRISGGYENVPTDDIHKQVDLENVWLHFIREFIAPVTL
KVFAGYYTKGFALLNFVVKYSPERQSLRPHHDASTFTINIALNNVGEDFQGGGCKFLRYNCSIES
PRKGWSFMHPGRLTHLHEGLPVKNGTRYIAVSFIDP

Host Rabbit

Reactivity Human

Purification Protein A

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)

[Protocol Download](#)

- Immunofluorescence (Transfected cell)

- Flow Cytometry (Transfected cell)

Gene Info — PLOD2

Entrez GeneID [5352](#)

GeneBank Accession# [NM_182943.2](#)

Protein Accession# [NP_891988.1](#)

Gene Name PLOD2

Gene Alias LH2, TLH

Gene Description procollagen-lysine, 2-oxoglutarate 5-dioxygenase 2

Omim ID [601865 609220](#)

Gene Ontology [Hyperlink](#)

Gene Summary

The protein encoded by this gene is a membrane-bound homodimeric enzyme that is localized to the cisternae of the rough endoplasmic reticulum. The enzyme (cofactors iron and ascorbate) catalyzes the hydroxylation of lysyl residues in collagen-like peptides. The resultant hydroxylysyl groups are attachment sites for carbohydrates in collagen and thus are critical for the stability of intermolecular crosslinks. Some patients with Ehlers-Danlos syndrome type VIB have deficiencies in lysyl hydroxylase activity. Mutations in the coding region of this gene are associated with Bruck syndrome. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq]

Other Designations lysine hydroxylase 2|lysyl hydroxylase 2|telopeptide lysyl hydroxylase

Pathway

- [Lysine degradation](#)

Disease

- [Cardiovascular Diseases](#)
- [Diabetes Mellitus](#)
- [Edema](#)
- [Genetic Predisposition to Disease](#)
- [Ovarian Neoplasms](#)