



# Recombinant Mouse Procollagen-lysine,2-oxoglutarate 5-dioxygenase 2 (Plod2)

<b>Product Code</b>	CSB-MP886417MO
<b>Abbreviation</b>	Plod2
<b>Storage</b>	The shelf life is related to many factors, storage state, buffer ingredients, storage temperature and the stability of the protein itself. Generally, the shelf life of liquid form is 6 months at -20°C/-80°C. The shelf life of lyophilized form is 12 months at -20°C/-80°C.
<b>Uniprot No.</b>	Q9R0B9
<b>Product Type</b>	Recombinant Protein
<b>Immunogen Species</b>	Mus musculus (Mouse)
<b>Purity</b>	≥85% (SDS-PAGE)
<b>Sequence</b>	VAEET PGRIPADKLL VITVATKEND GFHRFMNSAK YFNYYTVKVLG QGQEWRRGGDG MNSIGGGQKV RLLKEAMEHY ASQEDLVILF TECFDVVFAG GPEEVLLKFKQ KTNHKIVFAA DGLLWPKRL ADKYPVVHIG KRYLNSGGFI GYAPYISRLV QQWNLQDNDD DQLFYTKVYI DPLKREAFNI TLDHKCKIFQ ALNGATDEVV LKFENGKSRV KNTFYETLPV AINGNGPTKI LLNYFGNYVP NSWTQENGCA LCDVDTIDLS TVDVPPKVTL GVFIQPTPF LPRFLNLLLT LDYPKEALQL FIHNKEVYHE KDIKVFVDKA KHDISSIKIV GPEENLSQAE ARNMGMDFCR QDEKCDYYFS VDADVVLTPN RTLKFLIEQN RKIAPLVTR HGKLWSNFWG ALSPDGYAR SEDYVDIVQG NRVGWVNPY MANVYLIQGG TLRSEMNERN YFVRDKLDPD MALCRNARDM GVFMYSNRH EFGRLISTAN YNTSHLNDF WQIFENPVDW KEKYINRDYS KIFTENIVEQ PCPDVFWFPI FSERACDELV EEMEHYKWS GGKHHDSRIS GGYENVPTDD IHMKQIGLEN VWLHFIREFI APVTLKVFAG YYTKGFALLN FVVKYSPERQ RSLRPHHDAS TFTINIALNN VGEDFQGGGC KFLRYNCSIE SPRKGWSFMH PGRLTHLHEG LPVKNTRYI AVSFIDP
<b>Source</b>	Mammalian cell
<b>Target Names</b>	Plod2
<b>Protein Names</b>	Recommended name: Procollagen-lysine,2-oxoglutarate 5-dioxygenase 2 EC=1.14.11.4 Alternative name(s): Lysyl hydroxylase 2 Short name= LH2
<b>Expression Region</b>	26-737
<b>Notes</b>	Repeated freezing and thawing is not recommended. Store working aliquots at 4°C for up to one week.
<b>Tag Info</b>	Tag type will be determined during the manufacturing process.
<b>Protein Length</b>	Full Length of Mature Protein
<b>Target Details</b>	This protein 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.

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**Reconstitution**

We recommend that this vial be briefly centrifuged prior to opening to bring the contents to the bottom. Please reconstitute protein in deionized sterile water to a concentration of 0.1-1.0 mg/mL. We recommend to add 5-50% of glycerol (final concentration) and aliquot for long-term storage at -20°C/-80°C. Our default final concentration of glycerol is 50%. Customers could use it as reference.

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**Shelf Life**

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