



# Recombinant Human Peroxisomal bifunctional enzyme (EHHADH)

<b>Product Code</b>	CSB-YP600104HU
<b>Abbreviation</b>	EHHADH
<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>	Q08426
<b>Product Type</b>	Recombinant Protein
<b>Immunogen Species</b>	Homo sapiens (Human)
<b>Purity</b>	>85% (SDS-PAGE)
<b>Sequence</b>	MAEYTRLHNA LALIRLRNPP VNAISTLLR DIKEGLQKAV IDHTIKAIVI CGAEGKFSAG ADIRGFSAPR TFGLTLGHVV DEIQRNEKPV VAAIQGMAFG GGLELALGCH YRIAHAEAQV GLPEVTLGLL PGARGTQLLP RLTGVPAALD LITSGRRILA DEALKLGILD KVVNSDPVEE AIRFAQRVSD QPLESRRLCN KPIQSLPNMD SIFSEALLKM RRQHPGCLAQ EACVRAVQAA VQYPYEVGIK KEEELFLYLL QSGQARALQY AFFAERKANK WSTPSGASWK TASARPVSSV GVVGLGTMGR GIVISFARAR IPVIAVDSDK NQLATANKMI TSVLEKEASK MQQSGHPWSG PKPRLTSSVK ELGGVDLVIE AVFEEMSLKK QVFAELSAVC KPEAFLCTNT SALDVDEIAS STDRPHLVIG THFFSPAHEVM KLLLEVIPSQY SSPTTIATVM NLSKKIKKIG VVVGNCFGFV GNRMLNPYYN QAYFLLEEGS KPEEVDQVLE EFGFKMGPFV VSDLAGLDVG WKSARKGQGLT GPTLLPGTPA RKRGNRRYCP IPDVLCELGR FGQKTGKGWY QYDKPLGRIH KPDPWLSKFL SRYRKTHIE PRSISQDEIL ERCLYSLINE AFRILGEGIA ASPEHIDVVY LHGYGWPRHK GGPMFYASTV GLPTVLEKLQ KYRQNPDIQ QLEPSDYLLK LASQGNPPLK EWQSLAGSPS SKL
<b>Source</b>	Yeast
<b>Target Names</b>	EHHADH
<b>Protein Names</b>	Recommended name: Peroxisomal bifunctional enzyme Short name= PBE Short name= PBEF Including the following 2 domains: Enoyl-CoA hydratase/3,2-trans-enoyl-CoA isomerase EC= 4.2.1.17 EC= 5.3.3.8 3- hydroxyacyl-CoA
<b>Expression Region</b>	1-723
<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 protein
<b>Target Details</b>	This protein is a bifunctional enzyme and is one of the four enzymes of the



peroxisomal beta-oxidation pathway. The N-terminal region of the encoded protein contains enoyl-CoA hydratase activity while the C-terminal region contains 3-hydroxyacyl-CoA dehydrogenase activity. Defects in this gene are a cause of peroxisomal disorders such as Zellweger syndrome. Two transcript variants encoding different isoforms have been found for this gene.

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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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