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# Recombinant Human GCDH Protein (His Tag)

Catalog Number: PKSH032495

Note: Centrifuge before opening to ensure complete recovery of vial contents.

#### Description

Species Human

Source E.coli-derived Human GCDH protein Arg45-Lys438, with an N-terminal His

 Calculated MW
 45.0 kDa

 Observed MW
 41 kDa

 Accession
 Q92947

**Bio-activity** Not validated for activity

#### **Properties**

**Purity** > 95 % as determined by reducing SDS-PAGE.

**Concentration** Subject to label value.

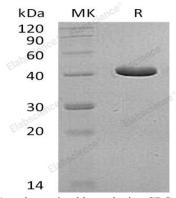
Endotoxin  $< 1.0 \text{ EU} \text{ per } \mu\text{g}$  of the protein as determined by the LAL method. Storage Storage Storage Storage winimize freeze-thaw cycles.

Shipping This product is provided as liquid. It is shipped at frozen temperature with blue ice/gel

packs. Upon receipt, store it immediately at < - 20°C.

**Formulation** Supplied as a 0.2 μm filtered solution of 20mM HEPES, 150mM NaCl, pH 7.4.

## **Data**



> 95 % as determined by reducing SDS-PAGE.

### Background

Glutaryl-CoA Dehydrogenase Mitochondrial (GCDH) is an enzyme that acts upon glutaryl-coenzyme A, creating crotonyl-coenzyme A. It plays a role in the metabolism of lysine, hydroxylysine and tryptophan. It uses electron transfer flavoprotein as its electron acceptor. Isoform Short is inactive Glutaryl-CoA and electron-transfer flavoprotein to (E)-but-2-enoyl-CoA, CO2 and reduced electron-transfer flavoprotein. A defect in this enzyme is associated with neurological condition glutaric acidemia type 1 and cause a progressive form of early-onset generalized dystonia.