

Human ENO1, His Tag, E. coli

Catalog Number LDG174PHE

Package 5 μg / 20 μg / 100 μg / Customized package

For full product information, images and publications, please visit our website.



Publications (2

Specifications

Species of Origin

Human

Affinity Tag

His Tag (N-term)

Purity

>98% as determined by SDS-PAGE analysis.

Endotoxin level

<0.1 EU per 1 μ g of the protein by the LAL method.

Expression system

Escherichia coli

Buffer

Lyophilized from a 0.2 μm filtered solution of PBS with 7 mM MgSO₄, pH 7.2

Molecular weight

The protein has a calculated MW of 47.85 kDa. The protein migrates as 55 kDa under reducing condition (SDS-PAGE analysis).

Form

Lyophilized

Background



Background

Enolase 1 (ENO1) also named alpha-enolase which is a glycolytic enzyme. It could convert the NAD+ to NADH by producing two ATP. ENO1 is a homodimer which is composed by two isozymes of enolase (2 α , 2 v, or 2 β). It also plays an important role in cancer model which can promote tumor cell proliferation and migration by the PI3K signaling pathway. ENO1 is a biomarker of prognostic and diagnostic cancer.

Uniprot ID

P06733 2

Synonyms

Alpha-enolase, 2-phospho-D-glycerate hydro-lyase, Cmyc promoter-binding protein, Phosphopyruvate hydratase, Plasminogen-binding protein

Sequence Note

Ser2-Lys434

Instruction

Reconstitution

It is recommended to reconstitute the lyophilized protein in sterile H_2O to a concentration of 200 $\mu g/mL$ and incubate the stock solution for at least 20 min to ensure sufficient re-dissolved.

Stability & Storage

This product is stable after storage at:

- -20°C for 12 months in lyophilized state from date of receipt.
- -20°C or -80°C for 1 month under sterile conditions after reconstitution.

Avoid repeated freeze/thaw cycles.

Shipping

The product is shipped with polar packs. Upon receipt, store it immediately at -20°C or lower for long term storage.

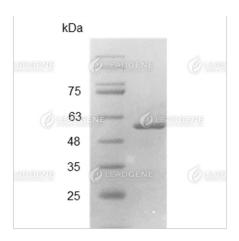
Image

Tainan Headquarter

Innovation & Research Center

CLD Center





SDS-PAGE analysis of recombinant human ENO1.

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