Product Name: Recombinant Human PIK3IP1 (C-6His) Catalog #: PHH1324



Summary

Name Phosphoinositide-3-kinase-interacting protein 1/PIK3IP1/HGFL

Purity Greater than 95% as determined by reducing SDS-PAGE

Endotoxin level <1 EU/µg as determined by LAL test.

Construction Recombinant Human Phosphoinositide-3-kinase-interacting Protein 1 is

produced by our Mammalian expression system and the target gene

encoding Ser22-Thr168 is expressed with a 6His tag at the C-terminus.

Accession # Q96FE7

Host **Human Cells**

Species Human

Predicted Molecular Mass 16.7 KDa

Formulation Lyophilized from a 0.2 µm filtered solution of 50mM Tris-HCl, 10mM reduced

Glutathione, pH 8.0.

Shipping The product is shipped at ambient temperature. Upon receipt, store it

immediately at the temperature listed below.

Store at ≤-70°C, stable for 6 months after receipt. Store at ≤-70°C, stable for 3 Stability&Storage

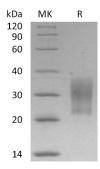
months under sterile conditions after opening. Please minimize freeze-thaw

cycles.

Reconstitution Always centrifuge tubes before opening. Do not mix by vortex or pipetting. It is

not recommended to reconstitute to a concentration less than 100µg/ml. Dissolve the lyophilized protein in distilled water. Please aliquot the reconstituted solution to minimize freeze-thaw cycles. Always centrifuge tubes before opening. Do not mix by vortex or pipetting. It is not recommended to reconstitute to a concentration less than 100µg/ml. Dissolve the lyophilized protein in distilled water. Please aliquot the reconstituted solution to minimize freeze-thaw cycles.

SDS-PAGE image



Background

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Alternative Names Kringle domain-containing protein HGFL; PIK3IP1; HGFL

Background Phosphoinositide-3-kinase-interacting protein 1(PIK3IP1) is an enzyme that in

humans is encoded by the PIK3IP1 gene.lt is a negative regulator of phosphatidylinositol-3-kinase (PI3K), suppresses the development of hepatocellular carcinoma. The gene encoding PIK3IP1 maps to human chromosome 22, which houses over 500 genes and is the second smallest human chromosome. Mutations in several of the genes that map to chromosome 22 are involved in the development of Phelan-McDermid syndrome, Neurofibromatosis

type 2, autism and schizophrenia.

Note

For Research Use Only, Not for Diagnostic Use.

Web: https://www.enkilife.com E-mail: order@enkilife.com techsupport@enkilife.com Tel: 0086-27-87002838