# Product Name: Recombinant Human FBPase1 (C-6His, E. Coli) Enkilife Catalog #: PEH0625

## **Summary**

Name FBPase 1/FBP1/Fructose-1,6-bisphosphatase 1

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

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

**Construction** Recombinant Human Fructose-1,6-Bisphosphatase 1 is produced by our E.coli

expression system and the target gene encoding Ala2-Gln338 is expressed

with a 6His tag at the C-terminus.

Accession # P09467

Host E.coli

Species Human

Predicted Molecular Mass 37.89 KDa

Formulation Supplied as a 0.2 µm filtered solution of 20mM Tris-HCl, 200mM NaCl, 1mM DTT,

1mM EDTA, 20% Glycerol, pH 8.0.

**Shipping** The product is shipped on dry ice/polar packs. Upon receipt, store it immediately

at the temperature listed below.

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

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

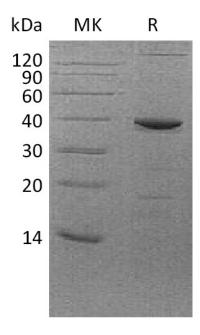
cycles.

Reconstitution

**SDS-PAGE** image

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### **Alternative Names**

Fructose-1; 6-Bisphosphatase 1; FBPase 1; D-Fructose-1; 6-Bisphosphate 1-Phosphohydrolase 1; FBP1; FBP

# **Background**

Fructose-1,6-Bisphosphatase 1 (FBPase 1) is a member of the FBPase class 1 family. FBPase 1 is a gluconeogenesis regulatory protein, which catalyzes the hydrolysis of fructose 1,6-bisphosphate to fructose 6-phosphate and inorganic phosphate. FBPase 1 can assume an active R-state, or an inactive T-state. FBPase 1 deficiency is inherited as an autosomal recessive disorder mainly in the liver and causes life-threatening episodes of hypoglycemia and metabolic acidosis in newborn infants or young children. FBPase 1 coupled with phosphofructokinase (PFK) is involved in the metabolism of pancreatic islet cells.

### Note

For Research Use Only, Not for Diagnostic Use.