# **Product Name: Recombinant Human HO-1**

Catalog #: PEH0803



### **Summary**

HO-1/HMOX1/HSP32 Name

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

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

Construction Recombinant Human Heme Oxygenase 1 is produced by our E.coli expression

system and the target gene encoding Met1-Thr261 is expressed.

Accession # P09601

Host E.coli

**Species** Human

**Predicted Molecular Mass** 29.86 KDa

**Formulation** Supplied as a 0.2 µm filtered solution of 20mM PB, 150mM NaCl, 1mM EDTA, pH

7.4.

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

at the temperature listed below.

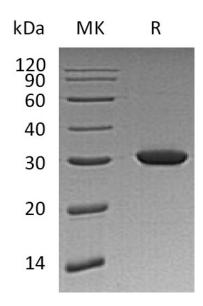
Store at ≤-70°C, stable for 6 months after receipt. Store at ≤-70°C, stable for 3 Stability&Storage

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

cycles.

Reconstitution

### **SDS-PAGE** image



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### **Alternative Names**

Heme Oxygenase 1; HO-1; HMOX1; HO; HO1

## **Background**

Heme Oxygenase 1 (HO-1) is an enzyme in endoplasmic reticulum that belongs to the heme oxygenase family. HO-1 cleaves the heme ring at the alpha methene bridge to form Biliverdin. Biliverdin is subsequently converted to Bilirubin by Biliverdin reductase. In physiological state, the highest activity of HO-1 is found in the spleen, where senescent erythrocytes are sequestrated and destroyed. HO-1 activity is highly inducible by its substrate heme and by various non-heme substances such as heavy metals, bromobenzene, endotoxin, oxidizing agents and UVA. HO-1 is involved in the regulation of cardiovascular function and response to a variety of stressors. Defects in HO-1 are the cause of Heme Oxygenase 1 deficiency, resulting in marked erythrocyte fragmentation and intravascular hemolysis, coagulation abnormalities, endothelial damage, and iron deposition in renal and hepatic tissues.

#### Note

For Research Use Only, Not for Diagnostic Use.

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