Product Name: Recombinant Human HO-1

Catalog #: PEH0803



Summary

Name HO-1/HMOX1/HSP32

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

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

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

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

at the temperature listed below.

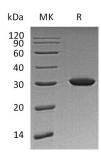
Stability&Storage Store at \leq -70°C, stable for 6 months after receipt. Store at \leq -70°C, stable for 3

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

cycles.

Reconstitution

SDS-PAGE image



Background

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

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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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