## **Product Name: Recombinant Human ARG1 (C-6His)**

Catalog #: PHH0097



#### **Summary**

Name Arginase-1/ARG1

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

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

Construction Recombinant Human Arginase-1 is produced by our Mammalian expression

system and the target gene encoding Met1-Lys322 is expressed with a 6His

tag at the C-terminus.

Accession # P05089

**Host** Human Cells

**Species** Human

Predicted Molecular Mass 35.6 KDa

Formulation Supplied as a 0.2 µm filtered solution of 20mM Tris-HCl, 150mM NaCl, 20%

Glycerol, 1mM DTT, pH 7.4.

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

at the temperature listed below.

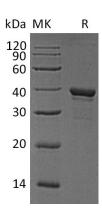
**Stability&Storage** Store at  $\leq$ -70°C, stable for 6 months after receipt. Store at  $\leq$ -70°C, stable for 3

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

cycles.

Reconstitution

### **SDS-PAGE** image



## **Background**

Alternative Names Arginase-1; Liver-type arginase; Type I arginase; ARG1

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#### **Background**

ARG1 is a member of the ureohydrolase family of enzymes. ARG1 can catalyze the hydrolysis of arginine to ornithine and urea. In the urea cycle, ARG1 catalyzes the fifth and final step, a series of biochemical reactions in mammals during which the body disposes of harmful ammonia. ARG1 is a cytosolic enzyme and expressed widely in the liver as part of the urea cycle, while it is also expressed in cells and tissues that lack a complete urea cycle, including lung. Inherited deficiency of this ARG1 causes argininemia, which is an autosomal recessive disorder characterized by hyperammonemia.

#### Note

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