

Arginase-1

PLD-537

Package : 可询价

Product Name : Arginase-1**Cat.No.:** PLD-537**Synonyms** : Liver-type arginase; Type I arginase**Application** : IHC-P**Reactivity** : Human**Host species** : Rabbit

Background

Arginase catalyzes the hydrolysis of arginine to ornithine and urea. At least two isoforms of mammalian arginase exist (types I and II) which differ in their tissue distribution, subcellular localization, immunologic crossreactivity and physiologic function. The type I isoform encoded by this gene, is a cytosolic enzyme and expressed predominantly in the liver as a component of the urea cycle. Inherited deficiency of this enzyme results in argininemia, an autosomal recessive disorder characterized by hyperammonemia. Two transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Sep 2011]

Gene ID

383

Swiss Prot

P05089

Synonyms

Liver-type arginase; Type I arginase

Reactivity

Human

Application

IHC-P

Recommended dilution

1:200-1:400

Host species

Rabbit

Clonality

Monoclonal

Isotype

IgG

Purity

Affinity Purification

Conjugation

Un-conjugated

Storage Stability

Store at -20°C. Supplied in 50mM Tris-Glycine(pH 7.4), 0.15M NaCl, 40% Glycerol, 0.01% sodium azide and 0.05% BSA. Stable for 12 months from date of receipt.