

## ARG1

**Reactivity:** Mouse Rat

**Tested applications:** WB

**Recommended Dilution:** WB 1:500 - 1:2000

**Calculated MW:** 35kDa

**Observed MW:** Refer to Figures

**Immunogen:**

Recombinant protein of human ARG1

**Storage Buffer:**

Store at -20. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

**Synonym:**

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

**Catalog #:** A1847

**Antibody Type:**

Polyclonal Antibody

**Species:** Rabbit

**Gene ID:** 383

**Isotype:** IgG

**Swiss Prot:** P05089

**Purity:** Affinity purification

For research use only.

**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.

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