

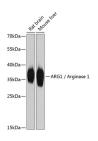
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# ARG1 / Arginase 1 Polyclonal Antibody

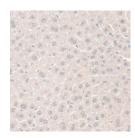
Catalog No.E-AB-60474ReactivityH,M,RStorageStore at -20°C. Avoid freeze / thaw cycles.HostRabbitApplicationsWB,IHC,IFIsotypeIgG

**Note:** Centrifuge before opening to ensure complete recovery of vial contents.

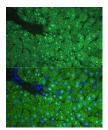
## **Images**



Western blot analysis of extracts of various cell lines using ARG1 / Arginase 1 Polyclonal Antibody at dilution of 1:2000.



Immunohistochemistry of paraffinembedded Mouse liver using ARG1 / Arginase 1 Polyclonal Antibody at dilution of 1:100 (40x lens).



Immunofluorescence analysis of Rat liver using ARG1 / Arginase 1 Polyclonal Antibody at dilution of 1:100. Blue: DAPI for nuclear staining.

## **Immunogen Information**

**Immunogen** Recombinant fusion protein of human ARG1 /

Arginase 1 (NP\_000036.2).

 GeneID
 383

 Swissprot
 P05089

**Synonyms** ARG1,arginase-1

#### **Product Information**

Calculated MW 25kDa/34kDa/35kDa

Observed MW 36kDa

**Buffer** PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

**Purify** Affinity purification

**Dilution** WB 1:500-1:2000 IHC 1:50-1:200 IF 1:50-1:200

#### **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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