

# **Product datasheet**

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# ARG64816 anti-Arginase 1 antibody

Package: 100 μg Store at: -20°C

# **Summary**

Product Description Goat Polyclonal antibody recognizes Arginase 1

Tested Reactivity Hu, Ms, Rat, Pig

Predict Reactivity Cow, Dog

Tested Application IHC-P, WB

Host Goat

Clonality Polyclonal

Isotype IgG

Target Name Arginase 1
Species Human

ImmunogenCFGLAREGNHKPIDConjugationUn-conjugated

Alternate Names EC 3.5.3.1; Type I arginase; Arginase-1; Liver-type arginase

# **Application Instructions**

Application table	Application	Dilution
	IHC-P	2 - 4 μg/ml
	WB	0.01 - 0.03 μg/ml
Application Note	IHC-P: Antigen Retrieval: Steam tissue section in Citrate buffer (pH 6.0).  WB: Recommend incubate at RT for 1h.  * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	
Observed Size	37 kDa	

### **Properties**

Form	Liquid	
Purification	Purified from goat serum by antigen affinity chromatography.	
Buffer	Tris saline (pH 7.3), 0.02% Sodium azide and 0.5% BSA.	
Preservative	0.02% Sodium azide	
Stabilizer	0.5% BSA	
Concentration	0.5 mg/ml	
Storage instruction	For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed	

before use.

Note

For laboratory research only, not for drug, diagnostic or other use.

#### Bioinformation

Background Arginase catalyzes the hydrolysis of arginine to ornithine and urea. At least two isoforms of mammalian

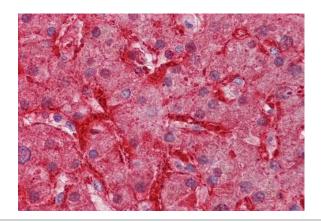
arginase exist (types I and II) which differ in their tissue distribution, subcellular localization, immunologic crossreactivity and physiologic function. The type I isoform encoded by this gene, is a cytosolic enzyme and expressed predominantly in the liver as a component of the urea cycle. Inherited 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]

Research Area Metabolism antibody; Signaling Transduction antibody

Calculated Mw 35 kDa

#### **Images**



#### ARG64816 anti-Arginase 1 antibody IHC-P image

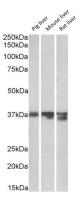
Immunohistochemistry: Paraffin-embedded Human liver stained with ARG64816 anti-Arginase 1 antibody at 3.75  $\mu$ g/ml dilution. Antigen Retrieval: Steam tissue section in Citrate buffer (pH 6.0).

#### Human liver

250kDa 150kDa 100kDa 75kDa 50kDa 37kDa 25kDa 20kDa

#### ARG64816 anti-Arginase 1 antibody WB image

Western blot: 35  $\mu g$  of Human liver lysate (in RIPA buffer) stained with ARG64816 anti-Arginase 1 antibody at 0.01  $\mu g/ml$  dilution and incubated at RT for 1 hour.



# ARG64816 anti-Arginase 1 antibody WB image

Western blot: 35  $\mu g$  of Pig liver, Mouse liver and Rat liver lysates (in RIPA buffer) stained with ARG64816 anti-Arginase 1 antibody at 0.03  $\mu g/ml$  dilution and incubated at RT for 1 hour.