abcam

Product datasheet

Recombinant human Liver Arginase protein ab168047

Description

Product name Recombinant human Liver Arginase protein

Biological activity Specific Activity: ~2.0U/µg protein. One unit is defined as the amount of enzyme that converts

1µmol of L-arginine to L-ornithine and urea per min. at 37°C, pH 9.5

Purity > 90 % SDS-PAGE.

Expression system Escherichia coli

Accession P05089

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Sequence MSAKSRTIGIIGAPFSKGQPRGGVEEGPTVLRKAGLLEKLK

EQECDVKDY

 ${\tt GDLPFADIPNDSPFQIVKNPRSVGKASEQLAGKVAEVKK}$

NGRISLVLGGD

HSLAIGSISGHARVHPDLGVIWVDAHTDINTPLTTTSGNLHG

QPVSFLLK

ELKGKIPDVPGFSWVTPCISAKDIVYIGLRDVDPGEHYILKT

LGIKYFSM

TEVDRLGIGKVMEETLSYLLGRKKRPIHLSFDVDGLDPSFT

PATGTPVVG

GLTYREGLYITEEIYKTGLLSGLDIMEVNPSLGKTPEEVTRT

VNTAVAIT LACFGLAREGNHKPIDYLNPPK

Predicted molecular weight 35 kDa

Amino acids 1 to 322

Specifications

Our <u>Abpromise guarantee</u> covers the use of ab168047 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications Functional Studies

SDS-PAGE

Form Liquid

Preparation and Storage

Stability and Storage Shipped at 4°C. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.

Constituents: 0.008% 2-Mercaptoethanol, 0.16% Tris HCI, 50% Glycerol (glycerin, glycerine),

0.013% Manganese chloride

This product is an active protein and may elicit a biological response in vivo, handle with caution.

General Info

Pathway Nitrogen metabolism; urea cycle; L-ornithine and urea from L-arginine: step 1/1.

Involvement in disease Defects in ARG1 are the cause of argininemia (ARGIN) [MIM:207800]; also known as

hyperargininemia. Argininemia is a rare autosomal recessive disorder of the urea cycle. Arginine is elevated in the blood and cerebrospinal fluid, and periodic hyperammonemia occurs. Clinical manifestations include developmental delay, seizures, mental retardation, hypotonia, ataxia,

progressive spastic quadriplegia.

Sequence similarities Belongs to the arginase family.

Cellular localization Cytoplasm.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

Our Abpromise to you: Quality guaranteed and expert technical support

- Replacement or refund for products not performing as stated on the datasheet
- Valid for 12 months from date of delivery
- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit https://www.abcam.com/abpromise or contact our technical team.

Terms and conditions

• Guarantee only valid for products bought direct from Abcam or one of our authorized distributors