# abcam

# Product datasheet

# Anti-ALAS2/ASB antibody [EPR15112(B)] - C-terminal ab184964



# ★☆☆☆ 1 Abreviews 8 References 6 Images

#### Overview

**Product name** Anti-ALAS2/ASB antibody [EPR15112(B)] - C-terminal

**Description** Rabbit monoclonal [EPR15112(B)] to ALAS2/ASB - C-terminal

Host species Rabbit

Tested applications Suitable for: WB, IP

**Species reactivity** Reacts with: Mouse, Rat, Human

**Immunogen** Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: NIH/3T3, PC-12, Rat brain, Jurkat, 293 and Human fetal liver. IP: K-562 cells.

**General notes**This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**<sup>®</sup> **patents**.

#### **Properties**

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

**Storage buffer** pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

Purity Protein A purified

Clonality Monoclonal
Clone number EPR15112(B)

**Isotype** IgG

1

#### **Applications**

#### The Abpromise quarantee

Our Abpromise guarantee covers the use of ab184964 in the following tested applications.

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

Application	Abreviews	Notes
WB		1/1000 - 1/2000. Detects a band of approximately 65 kDa (predicted molecular weight: 65 kDa).
IP		1/30 - 1/50.

#### **Target**

#### **Tissue specificity**

#### **Pathway**

#### Involvement in disease

Erythroid specific.

Porphyrin metabolism; protoporphyrin-IX biosynthesis; 5-aminolevulinate from glycine: step 1/1.

Defects in ALAS2 are a cause of anemia sideroblastic X-linked (XLSA) [MIM:300751]. Sideroblastic anemia is characterized by anemia of varying severity, hypochromic peripheral erythrocytes, systemic iron overload secondary to chronic ineffective erythropoiesis, and the presence of bone marrow ringed sideroblasts. Sideroblasts are characterized by iron-loaded mitochondria clustered around the nucleus. XLSA shows a variable hematologic response to pharmacologic doses of pyridoxine.

Defects in ALAS2 are the cause of erythropoietic protoporphyria X-linked dominant (XLDPT) [MIM:300752]. Porphyrias are inherited defects in the biosynthesis of heme, resulting in the accumulation and increased excretion of porphyrins or porphyrin precursors. They are classified as erythropoietic or hepatic, depending on whether the enzyme deficiency occurs in red blood cells or in the liver. XLDPT is a form of porphyria characterized biochemically by a high proportion of zinc-protoporphyrin in erythrocytes, in which a mismatch between protoporphyrin production and the heme requirement of differentiating erythroid cells leads to overproduction of protoporphyrin in amounts sufficient to cause photosensitivity and liver disease. Note=Gain of function mutations in ALS2 are responsible for XLDPT, but they can also be a possible aggravating factor in congenital erythropoietic porphyria and other erythropoietic disorders caused by mutations in other genes (PubMed:21309041).

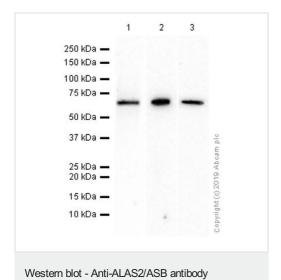
#### Sequence similarities

#### **Cellular localization**

Belongs to the class-II pyridoxal-phosphate-dependent aminotransferase family.

Mitochondrion matrix.

#### **Images**



[EPR15112(B)] - C-terminal (ab184964)

**All lanes :** Anti-ALAS2/ASB antibody [EPR15112(B)] - C-terminal (ab184964) at 1/1000 dilution (Purified)

Lane 1 : NIH/3T3 (Mouse embryonic fibroblast) whole cell lysates

Lane 2 : PC-12 (Rat adrenal gland pheochromocytoma) whole cell lysates

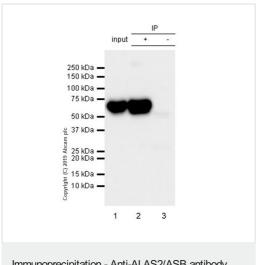
Lane 3: Rat brain lysates

Lysates/proteins at 15 µg per lane.

## **Secondary**

**All lanes :** Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

**Predicted band size:** 65 kDa **Observed band size:** 65 kDa



Immunoprecipitation - Anti-ALAS2/ASB antibody [EPR15112(B)] - C-terminal (ab184964) ab184964 (Purified) at 1:30 dilution (2  $\mu$ g) immunoprecipitating ALAS2/ASB in K-562 whole cell lysate.

Lane 1 (input): K-562 (Human chronic myelogenous leukemia lymphoblast) whole cell lysate 10 µg

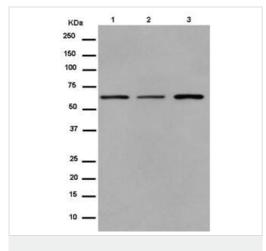
Lane 2 (+): ab184964 & K-562 whole cell lysate

Lane 3 (-): Rabbit monoclonal IgG (<u>ab172730</u>) instead of ab184964 in K-562 whole cell lysate

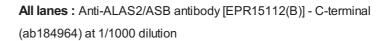
For western blotting, VeriBlot for IP Detection Reagent (HRP)

(ab131366) was used at 1:1000 dilution.

Blocking and diluting buffer: 5% NFDM/TBST.



Western blot - Anti-ALAS2/ASB antibody [EPR15112(B)] - C-terminal (ab184964)



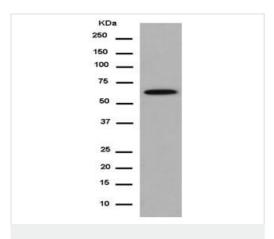
Lane 1 : K562 lysate Lane 2 : Jurkat lysate Lane 3 : 293 lysate

Lysates/proteins at 20 µg per lane.

# **Secondary**

**All lanes :** Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 65 kDa



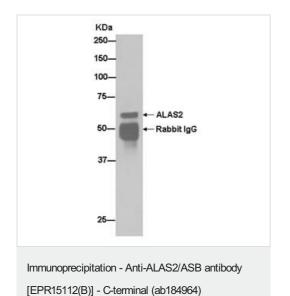
Western blot - Anti-ALAS2/ASB antibody [EPR15112(B)] - C-terminal (ab184964)

Anti-ALAS2/ASB antibody [EPR15112(B)] - C-terminal (ab184964) at 1/1000 dilution + Human fetal liver at 10 µg

# Secondary

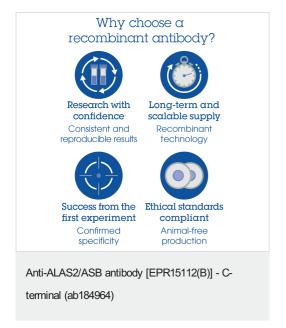
Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at 1/1000 dilution

Predicted band size: 65 kDa



Western blot analysis of immunoprecipitation pellet from 293 cell lysate immunoprecipitated using ab184964 at 1/50 dilution.

Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated secondary used at a 1/1000 dilution.



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