abcam

Product datasheet

Anti-LDB3 antibody [EPR10126] ab171936

Recombinant

RabMAb

2 Images

Overview

Product name Anti-LDB3 antibody [EPR10126]

Description Rabbit monoclonal [EPR10126] to LDB3

Host species Rabbit

Tested applications Suitable for: WB

Unsuitable for: Flow Cyt,ICC/IF,IHC or IP

Species reactivity Reacts with: Human

Predicted to work with: Mouse, Rat

Immunogen Synthetic peptide within Human LDB3 aa 50-150 (N terminal) (Cysteine residue). The exact

sequence is proprietary. Database link: **Q75112**

Positive control A431 and HT 1080 cell line lysates

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

- High batch-to-batch consistency and reproducibility

Improved sensitivity and specificityLong-term security of supply

- Animal-free production

For more information **see here**.

Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**[®] **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: 7.20

Preservative: 0.01% Sodium azide

Constituents: 9% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA, 50% Tissue culture

supernatant

Purity Protein A purified

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ClonalityMonoclonalClone numberEPR10126

Isotype IgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab171936 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 - 1/5000. Predicted molecular weight: 77 kDa.

Application notes Is unsuitable for Flow Cyt,ICC/IF,IHC or IP.

Target

Function May function as an adapter in striated muscle to couple protein kinase C-mediated signaling via

its LIM domains to the cytoskeleton.

Tissue specificity Expressed primarily in skeletal muscle and to a lesser extent in heart. Also detected in brain and

placenta.

Involvement in diseaseDefects in LDB3 are the cause of cardiomyopathy dilated type 1C (CMD1C) [MIM:601493].

Dilated cardiomyopathy is a disorder characterized by ventricular dilation and impaired systolic function, resulting in congestive heart failure and arrhythmia. Patients are at risk of premature

death.

Defects in LDB3 are the cause of left ventricular non-compaction type 3 (LVNC3) [MIM:601493]. Left ventricular non-compaction is characterized by numerous prominent trabeculations and deep

intertrabecular recesses in hypertrophied and hypokinetic segments of the left ventricle.

Defects in LDB3 are the cause of myopathy myofibrillar ZASP-related (MFM-ZASP)

[MIM:609452]. A neuromuscular disorder characterized by distal and proximal muscle weakness

with signs of cardiomyopathy and neuropathy.

Sequence similaritiesContains 3 LIM zinc-binding domains.

Contains 1 PDZ (DHR) domain.

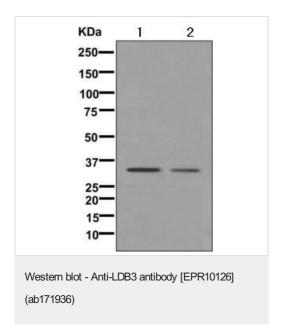
Cytoplasm > perinuclear region. Cell projection > pseudopodium. Cytoplasm > cytoskeleton.

Cytoplasm > myofibril > sarcomere > Z line. Localized to the cytoplasm around nuclei and

pseudopodia of undifferentiated cells and detected throughout the myotubes of differentiated

cells. Colocalizes with ACTN2 at the Z-lines.

Images



All lanes : Anti-LDB3 antibody [EPR10126] (ab171936) at 1/1000 dilution

Lane 1: A431 cell line lysate
Lane 2: HT 1080 cell line lysate

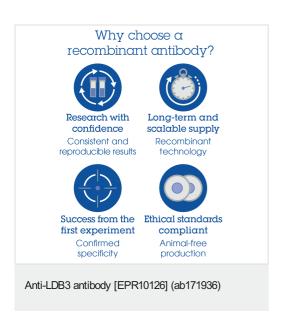
Lysates/proteins at 10 µg per lane.

Secondary

All lanes : Goat ant-rabbit HRP conjugated antibody at 1/500 dilution

Developed using the ECL technique.

Predicted band size: 77 kDa



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

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