abcam

Product datasheet

Anti-ENO3 antibody ab126259

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Overview

Product name Anti-ENO3 antibody

Description Rabbit polyclonal to ENO3

Host species Rabbit

Tested applications Suitable for: WB, ICC/IF

Species reactivity Reacts with: Human

Immunogen Recombinant fragment, corresponding to a region within amino acids 1-371 of Human ENO3.

Positive control A549, HeLa, HepG and HCT116 whole cell lysates; HeLa cells.

General notes

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle.

Storage buffer pH: 7.00

Preservative: 0.01% Thimerosal (merthiolate)

Constituents: 78.99% PBS, 1% BSA, 20% Glycerol (glycerin, glycerine)

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise quarantee Our Abpromise quarantee covers the use of ab126259 in the following tested applications.

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

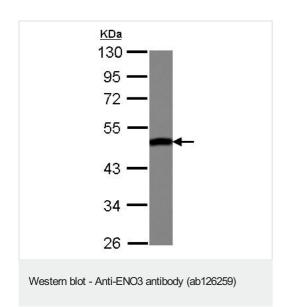
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Application	Abreviews	Notes
WB	*****(1)	1/5000 - 1/20000. Predicted molecular weight: 47 kDa.
ICC/IF		1/100 - 1/1000.

Target

Function Tissue specificity	Appears to have a function in striated muscle development and regeneration. The alpha/alpha homodimer is expressed in embryo and in most adult tissues. The alpha/beta heterodimer and the beta/beta homodimer are found in striated muscle, and the alpha/gamma heterodimer and the gamma/gamma homodimer in neurons.
Pathway	Carbohydrate degradation; glycolysis; pyruvate from D-glyceraldehyde 3-phosphate: step 4/5.
Involvement in disease	Defects in ENO3 are the cause of glycogen storage disease type 13 (GSD13) [MIM:612932]. A metabolic disorder that results in exercise-induced myalgias, generalized muscle weakness and fatigability. It is characterized by increased serum creatine kinase and decreased enolase 3 activity. Dramatically reduced protein levels with focal sarcoplasmic accumulation of glycogenbeta particles are detected on ultrastructural analysis.
Sequence similarities	Belongs to the enclase family.
Developmental stage	During ontogenesis, there is a transition from the alpha/alpha homodimer to the alpha/beta heterodimer in striated muscle cells, and to the alpha/gamma heterodimer in nerve cells.
Cellular localization	Cytoplasm. Localized to the Z line. Some colocalization with CKM at M-band.

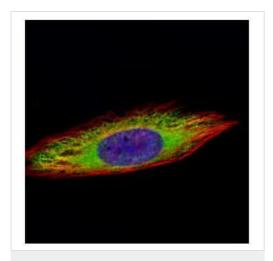
Images



Anti-ENO3 antibody (ab126259) at 1/10000 dilution + HepG2 cell lysate at 30 μg

Predicted band size: 47 kDa

10% SDS-PAGE



Immunocytochemistry/ Immunofluorescence - Anti-ENO3 antibody (ab126259)

ab126259 at 1/500 dilution staining ENO3 in methanol-fixed HeLa cells by Immunofluorescence (green). Alpha tubulin filaments are stained red.

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