abcam

Product datasheet

Anti-ERAB antibody [5F3] ab10260

6 References 1 Image

Overview

Product name Anti-ERAB antibody [5F3]

Description Mouse monoclonal [5F3] to ERAB

Host species Mouse

Specificity This antibody detects the recombinant human ERAB protein (27kDa) and recognizes the

endogenous ERAB protein in cell extracts with virtually no crossreactivity with other human

proteins.

Tested applications

Suitable for: IHC-P

Species reactivity

Reacts with: Human

Immunogen Recombinant, full-length Human ERAB protein (fusion protein).

General notesThe 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 and store at -20°C or -80°C. Avoid repeated freeze / thaw

cycles.

Storage buffer Constituent: PBS

no BSA

Purity Affinity purified

Purification notesAmmonium sulfate precipitated and dialyzed tissue culture supernatant.

Clonality Monoclonal

Clone number 5F3 lsotype lgG1

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Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab10260 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
IHC-P		1/100 - 1/1000.

Target

Function

Functions in mitochondrial tRNA maturation. Part of mitochondrial ribonuclease P, an enzyme composed of MRPP1/RG9MTD1, MRPP2/HSD17B10 and MRPP3/KIAA0391, which cleaves tRNA molecules in their 5'-ends. By interacting with intracellular amyloid-beta, it may contribute to the neuronal dysfunction associated with Alzheimer disease (AD).

Tissue specificity

Involvement in disease

Expressed in normal tissues but is overexpressed in neurons affected in AD.

Defects in HSD17B10 are the cause of 2-methyl-3-hydroxybutyryl-CoA dehydrogenase deficiency (MHBD deficiency) [MIM:300438]. MHBD deficiency leads to neurological abnormalities, including psychomotor retardation, and, in virtually all patients, loss of mental and motor skills. Defects in HSD17B10 are the cause of mental retardation syndromic X-linked type 10 (MRXS10) [MIM:300220]. MRXS10 is characterized by mild mental retardation, choreoathetosis and abnormal behavior.

A chromosomal microduplication involving HSD17B10 and HUWE1 is the cause of mental retardation X-linked type 17 (MRX17) [MIM:300705]; also known as mental retardation X-linked type 31 (MRX31). Mental retardation is characterized by significantly sub-average general intellectual functioning associated with impairments in adaptative behavior and manifested during the developmental period. In contrast to syndromic or specific X-linked mental retardation which also present with associated physical, neurological and/or psychiatric manifestations, intellectual deficiency is the only primary symptom of non-syndromic X-linked mental retardation.

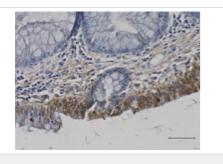
Sequence similarities

Belongs to the short-chain dehydrogenases/reductases (SDR) family.

Cellular localization

Mitochondrion.

Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ERAB antibody [5F3] (ab10260)

IHC staining of purifiedab10260 on formalin-fixed paraffinembedded human colon tissue. The tissue was incubated with 10 µg/ml of the primary antibody for 60 minutes at room temperature. A HRP kit was used for detection followed by hematoxylin counterstaining, according to the protocol provided. The image was captured with a 40X objective. Scale bar: 50 µm

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