abcam

Product datasheet

Anti-GAMT antibody [EPR6925] ab126736



RabMAb

4 Images

Overview

Product name Anti-GAMT antibody [EPR6925]

Description Rabbit monoclonal [EPR6925] to GAMT

Host species Rabbit

Tested applications Suitable for: WB, IHC-P

Unsuitable for: Flow Cyt,ICC/IF or IP

Species reactivity Reacts with: Human

Predicted to work with: Rat

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

Positive control Human fetal liver, HepG2 and SH-SY5Y lysates; Human liver tissue.

General notesThis 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.

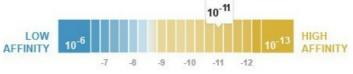
Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**[®] **patents**.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

Dissociation constant (K_D) $K_D = 7.40 \times 10^{-11} M$



Learn more about K_D

Storage buffer pH: 7.20

Preservative: 0.01% Sodium azide

1

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

supernatant

Purity Protein A purified

ClonalityMonoclonalClone numberEPR6925

Isotype IgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab126736 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 - 1/10000. Detects a band of approximately 26 kDa (predicted molecular weight: 26 kDa).
IHC-P		1/100 - 1/250. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

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

Target

Tissue specificity Expressed in liver.

Pathway Amine and polyamine biosynthesis; creatine biosynthesis; creatine from L-arginine and glycine:

step 2/2.

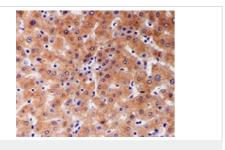
Involvement in diseaseDefects in GAMT are the cause of quanidinoacetate methyltransferase deficiency (GAMT

deficiency) [MIM:612736]. GAMT deficiency is an autosomal recessive disorder characterized by developmental delay/regression, mental retardation, severe disturbance of expressive and cognitive speech, intractable seizures and movement disturbances, severe depletion of creatine/phosphocreatine in the brain, and accumulation of guanidinoacetic acid (GAA) in brain

and body fluids.

Sequence similaritiesBelongs to the RMT2 methyltransferase family.

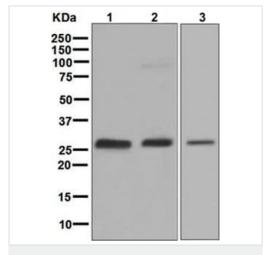
Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GAMT antibody
[EPR6925] (ab126736)

ab126736 at 1/100 dilution, staining GAMT in Formalin-fixed, Paraffin-embedded Human liver tissue by Immunohistochemistry.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Western blot - Anti-GAMT antibody [EPR6925] (ab126736)

All lanes : Anti-GAMT antibody [EPR6925] (ab126736) at 1/1000 dilution

Lane 1: Human fetal liver Lysate

Lane 2 : HepG2 lysate

Lane 3 : SH-SY5Y lysate

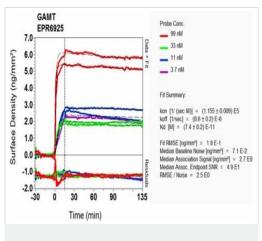
Lysates/proteins at 10 µg per lane.

Secondary

All lanes: Goat-anti-rabbit HRP at 1/2000 dilution

Developed using the ECL technique.

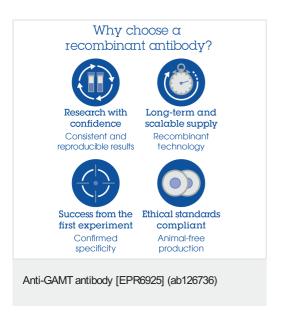
Predicted band size: 26 kDa



Ol-RD Scanning - Anti-GAMT antibody [EPR6925] (ab126736)

Equilibrium disassociation constant (K_D) Learn more about K_D

Click here to learn more about K_D



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