

A Reliable Research Partner in Life Science and Medicine

Recombinant CPS1 Monoclonal Antibody

catalog number: AN301906L

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description

Reactivity Human;Rat;Mouse

Immunogen Recombinant human CPS1 fragment

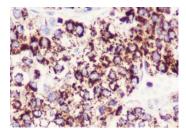
Host Rabbit Isotype lgG, κ Clone A622

Purification Protein Apurified

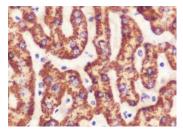
Buffer PBS, 50% glycerol, 0.05% Proclin 300, 0.05% protein protectant.

Applications Recommended Dilution

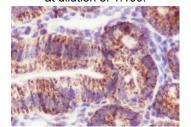
1:1000-1:5000 **WB** 1:50-1:100 **IHC**



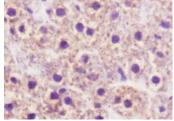
Immunohistochemistry of paraffin-embedded Human hepatocellular carcinoma using CPS1 Monoclonal Antibody at dilution of 1:100.



Immunohistochemistry of paraffin-embedded Human liver using CPS1 Monoclonal Antibody at dilution of 1:100.



intestine using CPS1 Monoclonal Antibody at dilution of 1:100.



Immunohistochemistry of paraffin-embedded Mouse small Immunohistochemistry of paraffin-embedded Rat liver using CPS1 Monoclonal Antibody at dilution of 1:100.

Preparation & Storage

Storage Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles.

Shipping Ice bag

Background

For Research Use Only

Toll-free: 1-888-852-8623 Fax: 1-832-243-6017 Tel: 1-832-243-6086 Web: www.elabscience.com Email: techsupport@elabscience.com Rev. V1.0

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Carbamoyl phosphate synthetase 1 (CPS1), a rate-limiting enzyme in the urea cycle, catalyzes the conversion of ammonia and bicarbonate to carbamoyl phosphate in mitochondria. Non-small cell lung carcinoma (NSCLC) cells with oncogenic KRAS and loss of the tumor suppressor LKB1 express CPS1 and depend on this enzyme for growth. CPS1 maintains pyrimidine/purine balance in these cancer cells. Silencing CPS1 reduces the pyrimidine to purine ratio and stalls DNA synthesis, leading to DNA damage and cancer cell death. In addition, the tumor suppressor p53 represses the expression of urea cycle enzymes CPS1, OTC, and ARG1 and, therefore, causes the accumulation of ammonia, suppressing cancer growth. Furthermore, research studies suggest that hypermethylation-mediated downregulation of CPS1 expression may contribute to the progression of normal hepatocytes to hepatocellular carcinoma (HCC).

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