Elabscience Biotechnology Co., Ltd.



A Reliable Research Partner in Life Science and Medicine

S100B Polyclonal Antibody

catalog number: D-AB-10118L

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

Description

Reactivity Human; Mouse; Rat

Immunogen Recombinant Human S100B protein expressed by E.coli

Host Rabbit Isotype IgG

Purification Antigen Affinity Purification

Conjugation Unconjugated

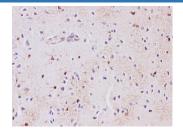
Buffer PBS with 0.05% Proclin 300, 1% protective protein and 50% glycerol, pH7.4

Applications Recommended Dilution

WB 1:500-1:1000 **IHC** 1:200-1:500

Data

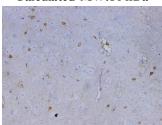




Western blot with anti-S100B polyclonal Antibody at dilution

of 1:500.lane 1: Rat brain tissue

Observed-MW:10 kDa Calculated-MW:10 kDa



Immunohistochemistry of paraffin-embedded Rat brain using S100B Polyclonal Antibody at dilution of 1:500

Immunohistochemistry of paraffin-embedded Mouse brain using S100B Polyclonal Antibody at dilution of 1:500

Preparation & Storage

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

Shipping The product is shipped with ice pack, upon receipt, store it immediately at the

Web: www.elabscience.cn

temperature recommended.

Background

For Research Use Only

Elabscience®

Elabscience Biotechnology Co., Ltd.

A Reliable Research Partner in Life Science and Medicine

The protein encoded by this gene is a member of the S100 family of proteins containing 2 EF-hand calcium-binding motifs. S100 proteins are localized in the cytoplasm and/or nucleus of a wide range of cells, and involved in the regulation of a number of cellular processes such as cell cycle progression and differentiation. S100 genes include at least 13 members which are located as a cluster on chromosome 1q21; however, this gene is located at 21q22.3. This protein may function in Neurite extension, proliferation of melanoma cells, stimulation of Ca2+ fluxes, inhibition of PKC-mediated phosphorylation, astrocytosis and axonal proliferation, and inhibition of microtubule assembly. Chromosomal rearrangements and altered expression of this gene have been implicated in several neurological, neoplastic, and other types of diseases, including Alzheimer's disease, Down's syndrome, epilepsy, amyotrophic lateral sclerosis, melanoma, and type I diabetes.

Web: www.elabscience.cn