Elabscience Biotechnology Co., Ltd.



A Reliable Research Partner in Life Science and Medicine

TRPM5 Polyclonal Antibody

catalog number: E-AB-12915

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

Description

Reactivity Human; Mouse

Immunogen Synthetic peptide of human TRPM5

Host Rabbit Isotype IgG

PurificationAffinity purificationConjugationUnconjugated

Buffer Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer and 50% glycerol.

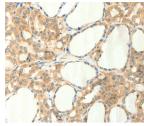
Applications Recommended Dilution

WB 1:500-1:2000 **IHC** 1:50-1:200

Data

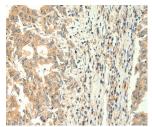
100a 130— 130— 95— 72— 55— 36—

Western Blot analysis of Mouse heart tissue using TRPM5 Polyclonal Antibody at dilution of 1:1200



Immunohistochemistry of paraffin-embedded Human thyroid cancer using TRPM5 Polyclonal Antibody at dilution of 1:60

Calculated-MW:131 kDa



Immunohistochemistry of paraffin-embedded Human gastric cancer using TRPM5 Polyclonal Antibody at dilution of 1:60

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

This gene encodes a member of the transient receptor potential (TRP) protein family, which is a diverse group of proteins with structural features typical of ion channels. This protein plays an important role in taste transduction, and has characteristics of a calcium-activated, non-selective cation channel that carries Na+, K+, and Cs+ ions equally well, but not Ca(2+) ions. It is activated by lower concentrations of intracellular Ca(2+), and inhibited by higher concentrations. It is also a highly temperature-sensitive, heat activated channel showing a steep increase of inward currents at temperatures between 15 and 35 degrees Celsius. This gene is located within the Beckwith-Wiedemann syndrome critical region-1 on chromosome 11p15.5, and has been shown to be imprinted, with exclusive expression from the paternal allele.

Web: www.elabscience.cn

For Research Use Only

Tel: 400-999-2100