

Elabscience®

A Reliable Research Partner in Life Science and Medicine

WIPF1 Polyclonal Antibody

catalog number: E-AB-91677

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

Description

Reactivity Human; Mouse; Rat

Immunogen A synthetic peptide of human WIPF1

Host Rabbit Isotype IgG

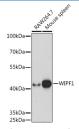
Purification Affinity purification

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

Recommended Dilution Applications

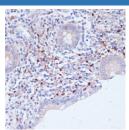
WB 1:500-1:2000 1:100-1:200 IHC

Data

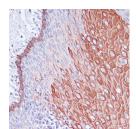


Western blot analysis of extracts of various cell lines using WIPF1 Polyclonal Antibody at1:1000 dilution.

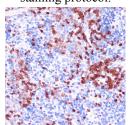
Observed-MW:51 kDa



Immunohistochemistry of paraffin-embedded human appendix using WIPF1 Polyclonal Antibody at dilution of 1:100 (40x lens). Perform microwave antigen retrieval with 10 mM PBS buffer pH 7.2 before commencing with IHC staining protocol



Immunohistochemistry of paraffin-embedded human esophageal using WIPF1 Polyclonal Antibody at dilution of 1:100 (40x lens). Perform microwave antigen retrieval with 10 lens). Perform microwave antigen retrieval with 10 mM PBS mM PBS buffer pH 7.2 before commencing with IHC staining protocol.



Immunohistochemistry of paraffin-embedded mouse spleen using WIPF1 Polyclonal Antibody at dilution of 1:100 (40x buffer pH 7.2 before commencing with IHC staining protocol.

Preparation & 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

temperature recommended.

Background

For Research Use Only

Toll-free: 1-888-852-8623 Web:www.elabscience.com

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Elabscience Bionovation Inc.



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This gene encodes a protein that plays an important role in the organization of the actin cytoskeleton. The encoded protein binds to a region of Wiskott-Aldrich syndrome protein that is frequently mutated in Wiskott-Aldrich syndrome, an X-linked recessive disorder. Impairment of the interaction between these two proteins may contribute to the disease. Two transcript variants encoding the same protein have been identified for this gene.

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