

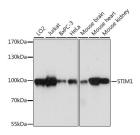
Tel:240-252-7368(USA) Fax: 240-252-7376(USA) techsupport@elabscience.com Website: www.elabscience.com

# **STIM1 Polyclonal Antibody**

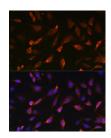
Catalog No.E-AB-67867ReactivityH,M,RStorageStore at -20°C. Avoid freeze / thaw cycles.HostRabbitApplicationsWB,IFIsotypeIgG

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

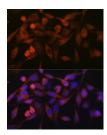
# **Images**



Western blot analysis of extracts of various cell lines using STIM1 Polyclonal Antibody at dilution of 1:1000.



Immunofluorescence analysis of C6 cells using STIM1 Polyclonal Antibody at dilution of 1:100. Blue: DAPI for nuclear staining.



Immunofluorescence analysis of HeLa cells using STIM1 Polyclonal Antibody at dilution of 1:100. Blue: DAPI for nuclear staining.

# **Immunogen Information**

**Immunogen** Recombinant fusion protein of human STIM1

(NP\_003147.2).

**GeneID** 6786 **Swissprot** Q13586

Synonyms STIM1,D11S4896E,GOK,IMD10,STRMK,TAM,TA

M1

#### **Product Information**

**Calculated MW** 62kDa/77kDa **Observed MW** 100kDa

**Buffer** PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

**Purify** Affinity purification

**Dilution** WB 1:200-1:2000 IF 1:50-1:200

# **Background**

This gene encodes a type 1 transmembrane protein that mediates Ca2+ influx after depletion of intracellular Ca2+ stores by gating of storeoperated Ca2+ influx channels (SOCs). It is one of several genes located in the imprinted gene domain of 11p15.5, an important tumor-suppressor gene region. Alterations in this region have been associated with the Beckwith-Wiedemann syndrome, Wilms tumor, rhabdomyosarcoma, adrenocrotical carcinoma, and lung, ovarian, and breast cancer. This gene may play a role in malignancies and disease that involve this region, as well as early hematopoiesis, by mediating attachment to stromal cells. Mutations in this gene are associated with fatal classic Kaposi sarcoma, immunodeficiency due to defects in store-operated calcium entry (SOCE) in fibroblasts, ectodermal dysplasia and tubular aggregate myopathy. This gene is oriented in a head-to-tail configuration with the ribonucleotide reductase 1 gene (RRM1), with the 3' end of this gene situated 1.6 kb from the 5' end of the RRM1 gene. Alternative splicing of this gene results in multiple transcript variants.

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