

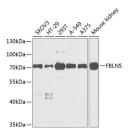
Tel:240-252-7368(USA) Fax: 240-252-7376(USA) techsupport@elabscience.com Website: www.elabscience.com

# FBLN5 Polyclonal Antibody

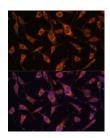
Catalog No. E-AB-63037 Reactivity H,M Store at -20°C. Avoid freeze / thaw cycles. Rabbit **Storage** Host **Applications** WB.IF **Isotype IgG** 

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

# **Images**



Western blot analysis of extracts of various cell lines using FBLN5 Polyclonal Antibody at dilution of 1:1000.



Immunofluorescence analysis of L929 cells using FBLN5 Polyclonal Antibody at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.

# **Immunogen Information**

**Immunogen** Recombinant fusion protein of human FBLN5

(NP 006320.2).

GeneID 10516 **Swissprot** Q9UBX5

FBLN5,ADCL2,ARCL1A,ARMD3,DANCE,EVEC,F **Synonyms** 

IBL-5,HNARMD,UP50,fibulin-5

#### **Product Information**

Calculated MW 50kDa **Observed MW** 72kDa

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

**Purify** Affinity purification

Dilution WB 1:500-1:2000 IF 1:50-1:100

# **Background**

The protein encoded by this gene is a secreted, extracellular matrix protein containing an Arg-Gly-Asp (RGD) motif and calcium-binding EGF-like domains. It promotes adhesion of endothelial cells through interaction of integrins and the RGD motif. It is prominently expressed in developing arteries but less so in adult vessels. However, its expression is reinduced in balloon-injured vessels and atherosclerotic lesions, notably in intimal vascular smooth muscle cells and endothelial cells. Therefore, the protein encoded by this gene may play a role in vascular development and remodeling. Defects in this gene are a cause of autosomal dominant cutis laxa, autosomal recessive cutis laxa type I (CL type I), and age-related macular degeneration type 3 (ARMD3).