

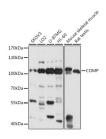
Tel:240-252-7368(USA) Fax: 240-252-7376(USA) techsupport@elabscience.com Website: www.elabscience.com

# **COMP Polyclonal Antibody**

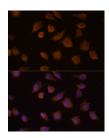
E-AB-66177 Reactivity Catalog No. H,M,R Storage Store at -20°C. Avoid freeze / thaw cycles. Rabbit Host **Applications** WB.IF **Isotype IgG** 

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

# **Images**



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



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

# **Immunogen Information**

Recombinant fusion protein of human COMP **Immunogen** 

GeneID 1311 P49747 **Swissprot** 

**Synonyms** COMP,EDM1,EPD1,MED,PSACH,THBS5,TSP5

#### **Product Information**

Calculated MW 77kDa/82kDa **Observed MW** 105kDa

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

**Purify** Affinity purification

Dilution WB 1:500-1:2000,IF 1:50-1:200

# **Background**

The protein encoded by this gene is a noncollagenous extracellular matrix (ECM) protein. It consists of five identical glycoprotein subunits, each with EGF-like and calcium-binding (thrombospondin-like) domains. Oligomerization results from formation of a five-stranded coiled coil and disulfides. Binding to other ECM proteins such as collagen appears to depend on divalent cations. Contraction or expansion of a 5 aa aspartate repeat and other mutations can cause pseudochondroplasia (PSACH) and multiple epiphyseal dysplasia (MED).