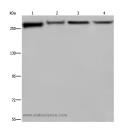
# **FLNA Polyclonal Antibody**

Catalog Number:E-AB-11240

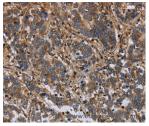


*Note:* Centrifuge before opening to ensure complete recovery of vial contents.

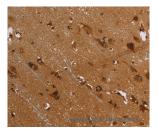
| Description  |                                                     |
|--------------|-----------------------------------------------------|
| Reactivity   | Human,Mouse                                         |
| Immunogen    | Recombinant protein of human FLNA                   |
| Host         | Rabbit                                              |
| Isotype      | IgG                                                 |
| Purification | Affinity purification                               |
| Conjugation  | Unconjugated                                        |
| Formulation  | PBS with 0.05% sodium azide and 50% glycerol, PH7.4 |
| Applications | Recommended Dilution                                |
| WB           | 1:500-1:2000                                        |
| IHC          | 1:50-1:200                                          |
| Data         |                                                     |



Western Blot analysis of PC3, Hela, NIH/3T3 and HUVEB cell using FLNA Polyclonal Antibody at dilution of 1:800 Calculated Mw:281kDa



Immunohistochemistry of paraffin-embedded Human liver cancer using FLNA Polyclonal Antibody at dilution of 1:40



Immunohistochemistry of paraffin-embedded Human brain using FLNA Polyclonal Antibody at dilution of 1:40

### **Preparation & Storage**

Storage

Store at -20°C. Avoid freeze / thaw cycles.

#### Background

The protein encoded by this gene is an actin-binding protein that crosslinks actin filaments and links actin filaments to membrane glycoproteins. The encoded protein is involved in remodeling the cytoskeleton to effect changes in cell shape and migration. This protein interacts with integrins, transmembrane receptor complexes, and second messengers. Defects

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in this gene are a cause of several syndromes, including periventricular nodular heterotopias (PVNH1, PVNH4), otopalatodigital syndromes (OPD1, OPD2), frontometaphyseal dysplasia (FMD), Melnick-Needles syndrome (MNS), and X-linked congenital idiopathic intestinal pseudoobstruction (CIIPX). Two transcript variants encoding different isoforms have been found for this gene.

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