

## FLNA Antibody

Catalog No: #36484

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## Description

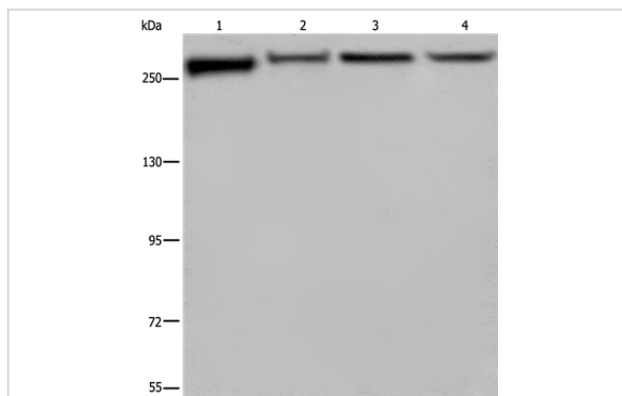
|                       |  |
|-----------------------|--|
| Product Name          | FLNA Antibody  |
| Host Species          | Rabbit   |
| Clonality             | Polyclonal   |
| Purification          | Antigen affinity purification.   |
| Applications          | WB IHC   |
| Species Reactivity    | Hu Ms  |
| Specificity           | The antibody detects endogenous levels of total FLNA protein.                            |
| Immunogen Type        | Recombinant Protein  |
| Immunogen Description | Fusion protein corresponding to residues near the C terminal of human filamin A, alpha   |
| Target Name           | FLNA   |
| Other Names           | FLN; FMD; MNS; OPD; ABPX; CSBS; CVD1; FLN1; NHBP; OPD1; OPD2; XLVD; XMVD; FLN-A; ABP-280 |
| Accession No.         | Swiss-Prot#: P21333NCBI Gene ID: 2316Gene Accssion: NP_001104026                         |
| Uniprot               | P21333   |
| GeneID                | 2316;  |
| SDS-PAGE MW           | 281kd  |
| Concentration         | 1.6mg/ml   |
| Formulation           | Rabbit IgG in pH7.4 PBS, 0.05% NaN <sub>3</sub> , 40% Glycerol.                          |
| Storage               | Store at -20°C   |

## Application Details

Western blotting: 1:500-1:2000

Immunohistochemistry: 1:50-1:200

## Images



Gel: 6%SDS-PAGE

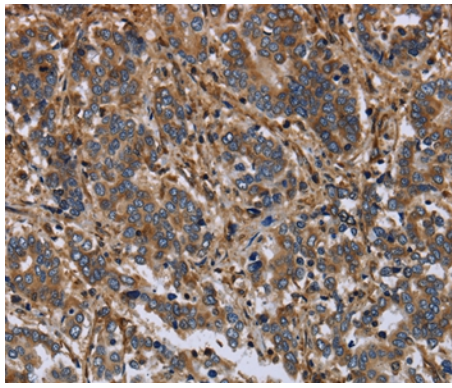
Lysates (from left to right): PC3, HeLa, NIH/3T3 and HUVEB cell

Amount of lysate: 40ug per lane

Primary antibody: 1/800 dilution

Secondary antibody dilution: 1/8000

Exposure time: 1 minute



Immunohistochemical analysis of paraffin-embedded Human liver cancer tissue using #36484 at dilution 1/40.

## 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 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.

Note: This product is for in vitro research use only