

Filamin 1 (phospho Ser2152) Polyclonal Antibody

Catalog No: YP0600

Reactivity: Human; Mouse; Rat; Monkey

P21333

Q8BTM8

Applications: WB;IHC;IF;ELISA

Target: Filamin 1

Fields: >>MAPK signaling pathway;>>Focal adhesion;>>Salmonella

infection;>>Proteoglycans in cancer

Gene Name: FLNA

Protein Name: Filamin-A

Human Gene Id: 2316

Human Swiss Prot

No:

Mouse Gene Id: 192176

Mouse Swiss Prot

No:

Immunogen: The antiserum was produced against synthesized peptide derived from human

Filamin A around the phosphorylation site of Ser2152. AA range:2121-2170

Specificity: Phospho-Filamin 1 (S2152) Polyclonal Antibody detects endogenous levels of

Filamin 1 protein only when phosphorylated at S2152.

Formulation : Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

Dilution: WB 1:500 - 1:2000. IHC 1:100 - 1:300. ELISA: 1:10000.. IF 1:50-200

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.



Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 280kD

Cell Pathway: MAPK_ERK_Growth;MAPK_G_Protein;Focal adhesion;

Background: filamin A(FLNA) Homo sapiens 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.[provided by RefSeg, Mar 2009],

Function: disease:Defects in FLNA are associated with cerebrofrontofacial syndrome

[MIM:608578]. This syndrome consists of a phenotype of male PVNH, with relatively normal development, no epilepsy or other neurological abnormality, severe constipation, and facial dysmorphism and without a discernible skeletal phenotype., disease:Defects in FLNA are the cause of frontometaphyseal dysplasia (FMD) [MIM:305620]. FMD is a congenital bone disease characterized by supraorbital hyperostosis, deafness and digital anomalies., disease:Defects in FLNA are the cause of Melnick-Needles syndrome (MNS) [MIM:309350]. MNS is a severe congenital bone disorder characterized by typical facies (exophthalmos, full cheeks, micrognathia and malalignment of teeth), flaring of the metaphyses of long bones, s-like curvature of bones of legs, irregular constrictions in the ribs,

and sclerosis of base of skull., disease: Defects i

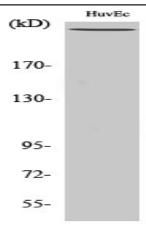
Subcellular Location:

Cytoplasm, cell cortex. Cytoplasm, cytoskeleton. Perikaryon. Cell projection, growth cone. Colocalizes with CPMR1 in the central region of DRG neuron growth cone (By similarity). Following SEMA3A stimulation of DRG neurons,

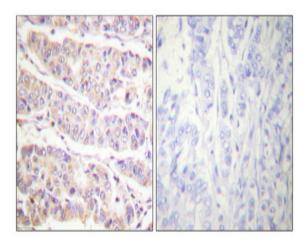
colocalizes with F-actin (By similarity). .

Expression: Ubiquitous.

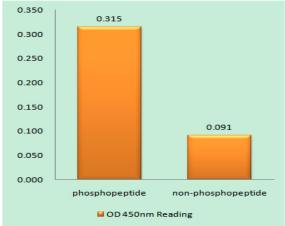
Products Images



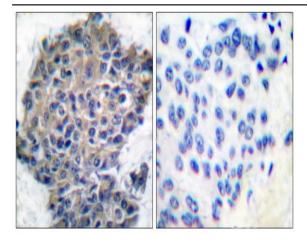
Western Blot analysis of various cells using Phospho-Filamin 1 (S2152) Polyclonal Antibody diluted at 1:2000



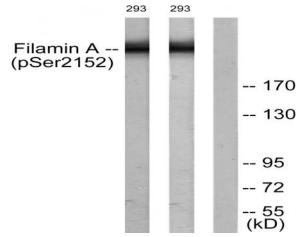
Immunohistochemical analysis of paraffin-embedded Human breast cancer. Antibody was diluted at 1:100(4° overnight). Highpressure and temperature Tris-EDTA,pH8.0 was used for antigen retrieval. Negetive contrl (right) obtaned from antibody was preabsorbed by immunogen peptide.



Enzyme-Linked Immunosorbent Assay (Phospho-ELISA) for Immunogen Phosphopeptide (Phospho-left) and Non-Phosphopeptide (Phospho-right), using Filamin A (Phospho-Ser2152) Antibody



Immunohistochemistry analysis of paraffin-embedded human breast carcinoma, using Filamin A (Phospho-Ser2152) Antibody. The picture on the right is blocked with the phospho peptide.



Western blot analysis of lysates from 293 cells treated with EGF 200ng/ml 5', using Filamin A (Phospho-Ser2152) Antibody. The lane on the right is blocked with the phospho peptide.

4/4