

## FGD1 rabbit pAb

Cat No.: ES16484

For research use only

## Overview

Product Name FGD1 rabbit pAb

Host species Rabbit Applications WB

Species Cross-Reactivity Human; Mouse Recommended dilutions WB 1:500-2000

Immunogen Synthesized peptide derived from human FGD1 AA

range: 508-558

**Specificity** This antibody detects endogenous levels of FGD1 at

Human/Mouse

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and

0.02% sodium azide.

Storage Store at -20°C. Avoid repeated freeze-thaw cycles.

Protein Name FGD1

Gene Name FGD1 FGDY ZFYVE3

**Cellular localization** Cytoplasm . Cell projection, lamellipodium . Cell

projection, ruffle . Cytoplasm, cytoskeleton . Associated with membrane ruffles and

lamellipodia...

**Purification** The antibody was affinity-purified from rabbit

antiserum by affinity-chromatography using

epitope-specific immunogen.

Clonality Polyclonal
Concentration 1 mg/ml
Observed band 105kD
Human Gene ID 2245
Human Swiss-Prot Number P98174

Alternative Names FYVE, RhoGEF and PH domain-containing protein 1

(Faciogenital dysplasia 1 protein) (Rho/Rac guanine nucleotide exchange factor FGD1) (Rho/Rac GEF) (Zinc finger FYVE domain-containing protein 3) This gene encodes a protein that contains Dbl (DH)

and pleckstrin (PH) homology domains and is similar to the Rho family of small GTP-binding proteins. The



**Background** 

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1 kDa 180 --140 --100 --75 --60 --45 --35 --25 --15 --

10 --

encoded protein specifically binds to the Rho family GTPase Cdc42Hs and can stimulate the GDP-GTP exchange of the isoprenylated form of Cdc42Hs. It also stimulates the mitogen activated protein kinase cascade leading to c-Jun kinase SAPK/JNK1 activation. Defects in this gene are the cause of faciogenital dysplasia and X-linked mental retardation, syndromatic 16.[provided by RefSeq, Mar 2011],

Western blot analysis of lysates from A431 cells, primary antibody was diluted at 1:1000, 4° over night



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