

# Rabbit Anti-FGD1 antibody

## SL16077R

Product Name:	FGD1
Chinese Name:	FGD1蛋白抗体
Alias:	AAS; Faciogenital dysplasia 1 protein; FGD1; FGD1_HUMAN; FGDY; FYVE RhoGEF and PH domain containing protein 1; FYVE, RhoGEF and PH domain-containing protein 1; Rho/Rac GEF; Rho/Rac guanine nucleotide exchange factor FGD1; ZFYVE3; Zinc finger FYVE domain containing protein 3; Zinc finger FYVE domain-containing protein 3.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Dog, Pig, Cow, Horse, Rabbit, Sheep,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=1:100-500IF=1:100-500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	107kDa
Cellular localization:	cytoplasmic
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human FGD1:601-700/961
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Storage:	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20 °C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed:	<u>PubMed</u>
Product Detail:	Activates CDC42, a member of the Ras-like family of Rho-and Rac proteins, by exchanging bound GDP for free GTP. Plays a role in regulating the actin cytoskeleton and cell shape.

## **Function:**

Activates CDC42, a member of the Ras-like family of Rho-and Rac proteins, by exchanging bound GDP for free GTP. Plays a role in regulating the actin cytoskeleton and cell shape.

### **Subcellular Location:**

Cytoplasm. Cell projection > lamellipodium. Cell projection > ruffle. Cytoplasm > cytoskeleton. Associated with membrane ruffles and lamellipodia.

## Tissue Specificity:

Expressed in fetal heart, brain, lung, kidney and placenta. Less expressed in liver; adult heart, brain, lung, pancreas and skeletal muscle.

## **DISEASE:**

Defects in FGD1 are the cause of Aarskog-Scott syndrome (AAS) [MIM:305400]. This faciogenital dysplasia is a rare multisystemic disorder characterized by disproportionately short stature, and by facial, skeletal, and urogenital anomalies. Note=Defects in FGD1 are found in a pateint with non-syndromal X-linked mental retardation.

## Similarity:

Contains 1 DH (DBL-homology) domain.

Contains 1 FYVE-type zinc finger.

Contains 2 PH domains.

## **SWISS:**

P98174

#### Gene ID:

2245

## Database links:

Entrez Gene: 2245 Human

Omim: 305400 Human

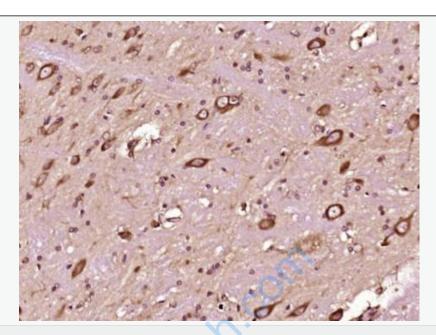
SwissProt: P98174 Human

Unigene: 631767 Human

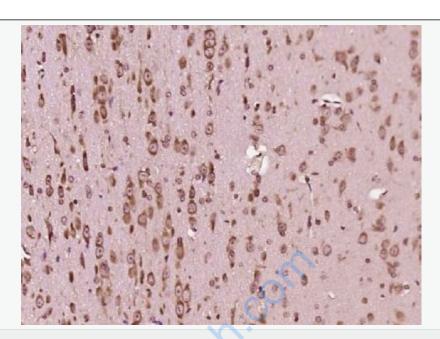
Unigene: 709201 Human

## **Important Note:**

	therapeutic or diagnostic applications.
Picture:	Sample: Cerebrum (Mouse) Lysate at 40 ug Primary: Anti-FGD1 (SL16077R) at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 107 kD Observed band size: 107 kD



Paraformaldehyde-fixed, paraffin embedded (Rat brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (FGD1) Polyclonal Antibody, Unconjugated (SL16077R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (Mouse brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (FGD1) Polyclonal Antibody, Unconjugated (SL16077R) at 1:500 overnight at 4°C, followed by a conjugated secondary (sp-0023) for 20 minutes and DAB staining.