

# Rabbit Anti-FGD3 antibody

## SL16079R

<b>Product Name:</b>	FGD3
Chinese Name:	FGD3蛋白抗体
Alias:	Faciogenital dysplasia 3; FGD1 family, member 3; FGD3; FGD3_HUMAN; FYVE; FYVE, RhoGEF and PH domain containing 3; FYVE, RhoGEF and PH domain containing protein 3; RhoGEF and PH domain-containing protein 3; ZFYVE5; Zinc finger FYVE domain containing protein 5; Zinc finger FYVE domain-containing protein 5.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat,
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:	79kDa
Cellular localization:	cytoplasmic
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human FGD3:501-600/725
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:	PubMed
Product Detail:	FGD1 gene mutations result in faciogenital dysplasia (FGDY, Aarskog-Scott syndrome), an X-linked developmental disorder that adversely affects the formation of multiple skeletal structures. FGD1 maps to human chromosome Xp11.21 and shares a

high degree of sequence identity with the FGD2 (6p21.2) and the FGD3 (9q22.31) proteins. FGD1 encodes a guanine nucleotide exchange factor that specifically activates the Rho GTPase Cdc42. FGD2 is present in several diverse tissues during embryogenesis, suggesting a role in embryonic development. FGD3 stimulates fibroblasts to form filopodia, which are Actin microspikes formed upon the stimulation of Cdc42. All FGD family members contain equivalent signaling domains and a conserved structural organization, which strongly suggests that these signaling domains form a canonical core structure for members of the FGD family of RhoGEF proteins. These proteins control essential signals required during embryonic development.

#### Function:

Promotes the formation of filopodia. May activate 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. Cytoplasm > cytoskeleton.

#### Similarity:

Contains 1 DH (DBL-homology) domain.

Contains 1 FYVE-type zinc finger.

Contains 2 PH domains.

### **SWISS:**

Q5JSP0

#### Gene ID:

89846

#### Database links:

Entrez Gene: 89846 Human

Entrez Gene: 30938 Mouse

Entrez Gene: 361223 Rat

SwissProt: Q5JSP0 Human

SwissProt: O88842 Mouse

Unigene: 411081 Human

Unigene: 291089 Mouse

Unigene: 214480 Rat

	Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.
Picture:	135— 100— 75— 63— 48— 35— 25—
	Sample: 293T Cell (Human) Lysate at 40 ug  Primary: Anti-FGD3 (SL16079R) at 1/300 dilution  Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution  Predicted band size: 79 kD
	Observed band size: 75 kD