

# **Product datasheet for TA386559M**

### OriGene Technologies, Inc.

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### **FLOT1 Rabbit Polyclonal Antibody**

#### **Product data:**

**Product Type:** Primary Antibodies

**Applications:** IF, IHC, IP, WB

Recommended Dilution: Recommended dilution: WB:1:500-1:5000, IHC:1:20-1:200, IF:1:50-1:200, IP:1:200-1:2000

**Reactivity:** Rat, Mouse, Human

Host: Rabbit Isotype: IgG

Clonality: Polyclonal

Immunogen: Recombinant Human Flotillin-1 protein (169-251AA)

Formulation: Preservative: 0.03% Proclin 300

Constituents: 50% Glycerol, 0.01M PBS, pH 7.4

**Concentration:** lot specific

**Purification:** >95%, Protein G purified

**Conjugation:** Unconjugated

**Storage:** Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.

**Stability:** 1 year from dispatch.

Database Link: <u>075955</u>

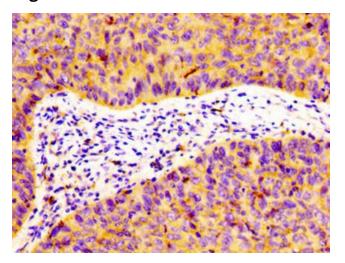
**Background:** May act as a scaffolding protein within caveolar membranes, functionally participating in

formation of caveolae or caveolae-like vesicles.

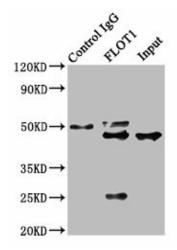




## **Product images:**



IHC image of [TA386559] diluted at 1:100 and staining in paraffin-embedded human cervical cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



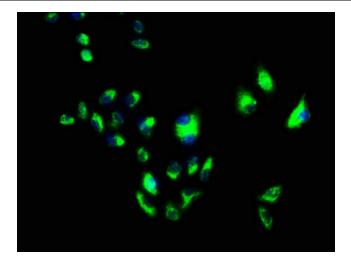
Immunoprecipitating FLOT1 in Hela whole cell lysate

Lane 1: Rabbit control IgG instead of [TA386559] in Hela whole cell lysate. For western blotting, a HRP-conjugated Protein G antibody was used as the secondary antibody (1/2000)

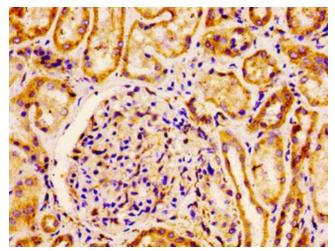
Lane 2: [TA386559] (8µg) + Hela whole cell lysate (500µg)

Lane 3: Hela whole cell lysate (10µg)

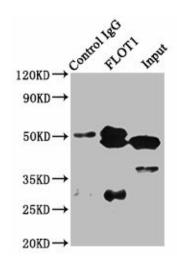




Immunofluorescence staining of Hela cells with [TA386559] at 1:100, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).



IHC image of [TA386559] diluted at 1:100 and staining in paraffin-embedded human kidney tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated ABC system.



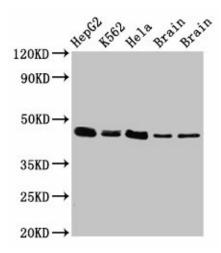
Immunoprecipitating FLOT1 in K562 whole cell lysate

Lane 1: Rabbit control IgG instead of [TA386559] in K562 whole cell lysate. For western blotting, a HRP-conjugated Protein G antibody was used as the secondary antibody (1/5000)

Lane 2: [TA386559] (8 $\mu$ g) + K562 whole cell lysate (500 $\mu$ g)

Lane 3: K562 whole cell lysate (20µg)





Western Blot

Positive WB detected in: HepG2 whole cell lysate, K562 whole cell lysate, Hela whole cell lysate, Rat brain tissue, Mouse brain tissue

All lanes: FLOT1 antibody at 3µg/ml

Secondary

Goat polyclonal to rabbit IgG at 1/50000 dilution

Predicted band size: 48, 43 kDa Observed band size: 48 kDa