

## [KD Validated] Anti-ARHGEF2 Rabbit mAb

Purified Recombinant Rabbit Monoclonal Antibody

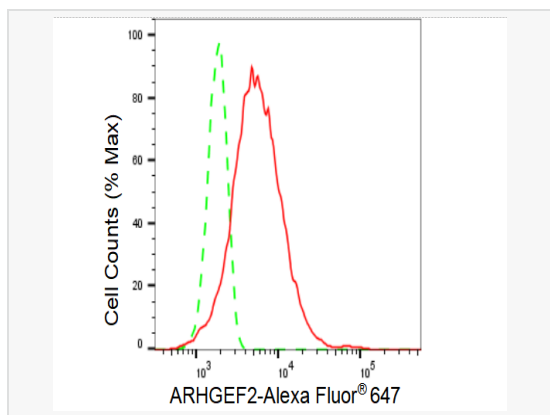
Catalog # R021629

### Product Information

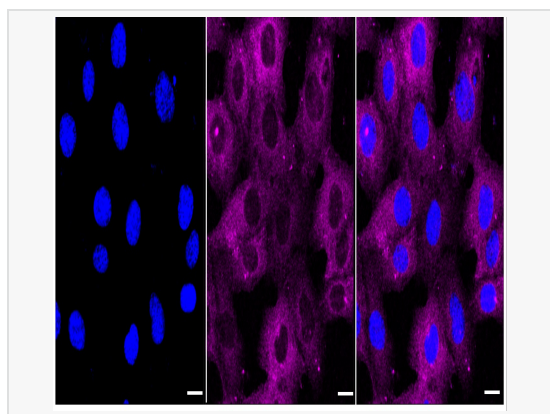
Application	WB, FC, IF (Cell)/ICC
Reactivity	Human, Mouse, Rat
Dilution	WB 1:1,000~1:5,000; FC 1:200~1:2,000; IF 1:100~1:1,000
Host	Rabbit
Clonality	Monoclonal
Clone No.	57E18J84
Isotype	IgG
Label	Unconjugated
Immunogen	A synthesized peptide derived from human GEF H1
Format	Affinity purified monoclonal antibody supplied in PBS with 0.02% sodium azide and 50% glycerol, pH 7.3.
Storage	Shipped on wet ice. Store at -20°C. Stable for 12 months from date of receipt. Aliquoting is unnecessary for -20°C storage.
Precautions	[KD Validated] Anti-ARHGEF2 Rabbit mAb [57E18J84] is for research use only and not for use in diagnostic or therapeutic procedures.

### Protein Information

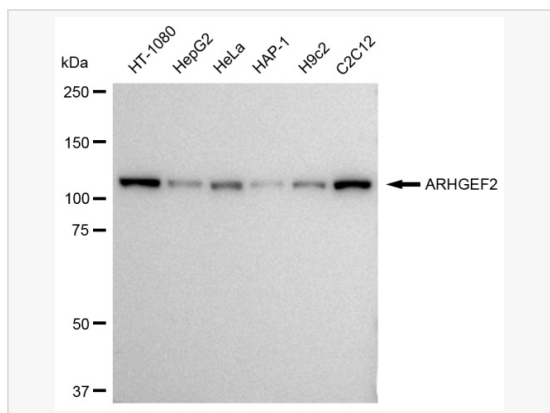
Synonyms	ARHGEF2; Rho/Rac Guanine Nucleotide Exchange Factor 2; GEF-H1; LFP40; Rho/Rac Guanine Nucleotide Exchange Factor (GEF); KIAA0651; GEFH1; P40; Rho Guanine Nucleotide Exchange Factor; Proliferating Cell Nucleolar Antigen P40; Guanine Nucleotide Exchange Factor H1; Microtubule-Regulated Rho-GEF; Lfc; NEDMHM; GEF; LFC.
Calculated MW	Calculated MW: 111 kDa, Observed MW: 112 kDa
Uniprot ID	Q92974
Gene ID	9181
Background	Rho GTPases play a fundamental role in numerous cellular processes that are initiated by extracellular stimuli that work through G protein coupled receptors. The encoded protein may form complex with G proteins and stimulate rho-dependent signals. Alternatively spliced transcript variants encoding different isoforms have been identified.[provided by RefSeq, Jun 2009].
Cellular Location	Cytoplasm. Cell junction > tight junction. Golgi apparatus. Cytoplasm > cytoskeleton > spindle. Cell projection > ruffle membrane. Localizes to the tips of cortical microtubules of the mitotic spindle during cell division, and is further released upon microtubule depolymerization. Recruited into membrane ruffles induced by <i>S.flexneri</i> at tight junctions of polarized epithelial cells.



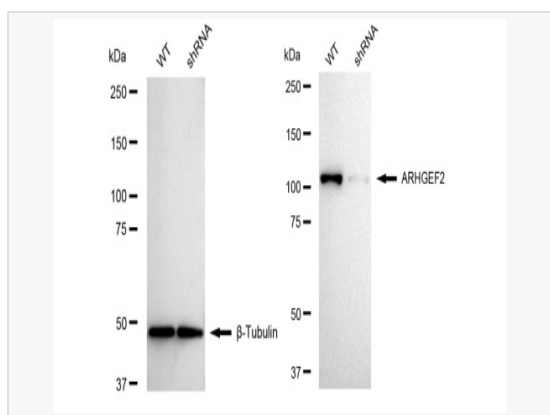
Flow cytometric analysis of ARHGEF2 expression in HT-1080 cells using ARHGEF2 antibody (R021629, 1:2,000). Green, isotype control; red, ARHGEF2.



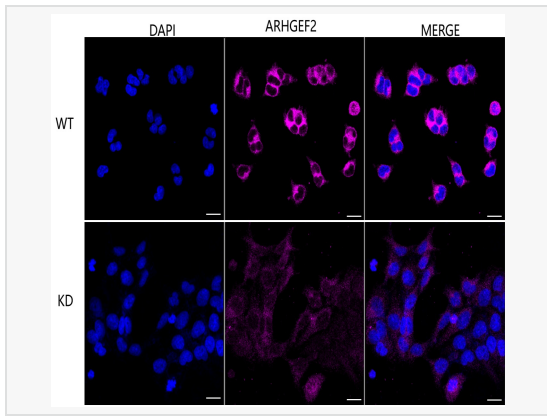
Immunocytochemical staining of HT-1080 cells with ARHGEF2 antibody (R021629, 1:1,000). Nuclei were stained blue with DAPI; ARHGEF2 was stained magenta with Alexa Fluor® 647. Images were taken using Leica stellaris 5. Protein abundance based on laser Intensity and smart gain: Medium. Scale bar, 20  $\mu$ m.



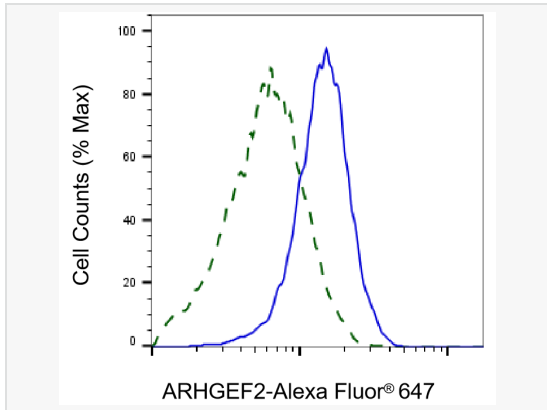
Western blotting analysis using ARHGEF2 antibody (R021629). Total cell lysates (30  $\mu$ g) from various cell lines were loaded and separated by SDS-PAGE. The blot was incubated with ARHGEF2 antibody (R021629, 1:5,000) and HRP-conjugated goat anti-rabbit secondary antibody (1:20,000) respectively. Image was developed using ECL Substrate Kit.



Western blotting analysis using ARHGEF2 antibody (R021629). ARHGEF2 expression in wild type (WT) and ARHGEF2 shRNA knockdown (KD) HeLa cells with 30  $\mu$ g of total cell lysates.  $\beta$ -Tubulin serves as a loading control. The blot was incubated with ARHGEF2 antibody (R021629, 1:5,000) and HRP-conjugated goat anti-rabbit secondary antibody (1:20,000) respectively. Image was developed using ECL Substrate Kit.



Immunocytochemical staining of HeLa cells using ARHGEF2 antibody (R021629, 1:1,000), Top panel: wild-type (WT); Bottom panel: ARHGEF2 shRNA knockdown (KD). Nuclei were stained blue with DAPI; ARHGEF2 was stained magenta with Alexa Fluor® 647. Scale bar, 20  $\mu$ m.



Validation of ARHGEF2 knockdown using flow cytometry. Wild-type(WT, Blue) and knockdown(KD, Green) HeLa cells were stained with ARHGEF2 antibody (R021629, 1:2,000) and analyzed using BD flow cytometer.