

## **ARHGEF9 Polyclonal Antibody**

Catalog No: YT0324

**Reactivity:** Human; Mouse; Rat

**Applications:** IHC;IF;ELISA

Target: ARHGEF9

**Gene Name:** ARHGEF9

**Protein Name:** Rho guanine nucleotide exchange factor 9

O43307

Q3UTH8

Human Gene Id: 23229

**Human Swiss Prot** 

No:

Mouse Gene ld: 236915

**Mouse Swiss Prot** 

No:

Rat Gene ld: 66013

Rat Swiss Prot No: Q9QX73

Immunogen: The antiserum was produced against synthesized peptide derived from human

ARHGEF9. AA range:399-448

**Specificity:** ARHGEF9 Polyclonal Antibody detects endogenous levels of ARHGEF9 protein.

**Formulation :** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

**Dilution :** IHC 1:100 - 1:300. ELISA: 1:40000.. IF 1:50-200

**Purification:** The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.



**Concentration**: 1 mg/ml

**Storage Stability:** -15°C to -25°C/1 year(Do not lower than -25°C)

Molecularweight: 61kD

**Cell Pathway:** Regulation of Actin Dynamics; AMPK

**Background:** The protein encoded by this gene is a Rho-like GTPase that switches between

the active (GTP-bound) state and inactive (GDP-bound) state to regulate CDC42 and other genes. Defects in this gene are a cause of startle disease with epilepsy (STHEE), also known as hyperekplexia with epilepsy. Three transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq,

Mar 2010],

**Function:** disease:Defects in ARHGEF9 are a cause of startle disease with epilepsy

(STHEE) [MIM:300607]; also known as hyperekplexia with epilepsy. Startle

disease is a genetically heterogeneous neurologic disorder. STHE is

characterized by muscular rigidity of central nervous system origin, particularly in the neonatal period, and by an exaggerated startle response to unexpected acoustic or tactile stimuli.,function:Acts as guanine nucleotide exchange factor (GEF) for CDC42. Promotes formation of GPHN clusters.,similarity:Contains 1 DH (DBL-homology) domain.,similarity:Contains 1 PH domain.,similarity:Contains 1 SH3 domain.,subunit:Interacts with GPHN.,tissue specificity:Detected in brain.

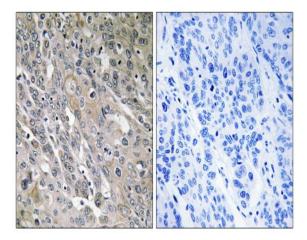
Detected at low levels in heart.,

Subcellular Location:

Cytoplasm . Cell junction, synapse, postsynaptic density .

**Expression:** Detected in brain. Detected at low levels in heart.

## **Products Images**



Immunohistochemistry analysis of paraffin-embedded human lung carcinoma tissue, using ARHGEF9 Antibody. The picture on the right is blocked with the synthesized peptide.