



PIPK I γ Monoclonal Antibody

Catalog No	YP-mAb-14921
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	PIP5K1C
Immunogen	The antiserum was produced against synthesized peptide derived from human PIP5K1C. AA range:305-354
Specificity	PIPK I γ Monoclonal Antibody detects endogenous levels of PIPK I γ protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	$\geq 90\%$
Storage Stability	-20°C/1 year
Synonyms	PIPK I γ ;PIP5K1C;Phosphatidylinositol 4-phosphate 5-kinase type-1 gamma; KIAA0589;PIP5K1-gamma;PtdIns(4)P-5-kinase 1 gamma;Phosphatidylinositol 4-phosphate 5-kinase type I gamma;PIP5KIgamma
Observed Band	80kD
Calculated Molecular Weight	80kD
Cell Pathway	Cell membrane; Peripheral membrane protein; Cytoplasmic side . Endomembrane system . Cytoplasm . Cell junction, focal adhesion . Cell junction, adherens junction . Cell projection, ruffle membrane . Cell projection, phagocytic cup . Cell projection, uropodium . Detected in plasma membrane invaginations. Isoform 3 is detected in intracellular vesicle-like structures.; [Isoform 2]: Cytoplasm. Nucleus.
Tissue Specificity	[Isoform 1]: Isoform 1 is strongly expressed in brain and also detected in heart and lung. ; [Isoform 2]: Isoform 2 is strongly expressed in pancreas and liver and in lesser quantities in brain, heart, lung and kidney. ; [Isoform 3]: Isoform 3 is detected in large amounts in heart and large intestine, is also present in lung, pancreas and thyroid, and to a lesser extent in brain, stomach and kidney.



Function

catalytic activity:ATP + 1-phosphatidyl-1D-myo-inositol 4-phosphate = ADP + 1-phosphatidyl-1D-myo-inositol 4,5-bisphosphate.,disease:Defects in PIP5K1C are the cause of lethal congenital contractural syndrome type 3 (LCCS3) [MIM:611369]; also known as multiple contractural syndrome Israeli Bedouin type B. LCCS is an autosomal recessive disorder characterized by early fetal hydrops and akinesia, the Pena-Shokeir phenotype, specific neuropathology with degeneration of anterior horn neurons and extreme skeletal muscle atrophy. LCCS3 patients present at birth with severe multiple joint contractures with severe muscle wasting and atrophy, mainly in the legs. LCCS3 can be distinguished from the original LCCS by the absence of hydrops, fractures, and multiple pterygia.,enzyme regulation:Activated by interaction with TLN2., function:Plays a role in membrane ruffling and assembly of clathrin-coate

Background

phosphatidylinositol-4-phosphate 5-kinase type 1 gamma(PIP5K1C) Homo sapiens This locus encodes a type I phosphatidylinositol 4-phosphate 5-kinase. The encoded protein catalyzes phosphorylation of phosphatidylinositol 4-phosphate, producing phosphatidylinositol 4,5-bisphosphate. This enzyme is found at synapses and has been found to play roles in endocytosis and cell migration. Mutations at this locus have been associated with lethal congenital contractural syndrome. Alternatively spliced transcript variants encoding different isoforms have been described.[provided by RefSeq, Sep 2010],

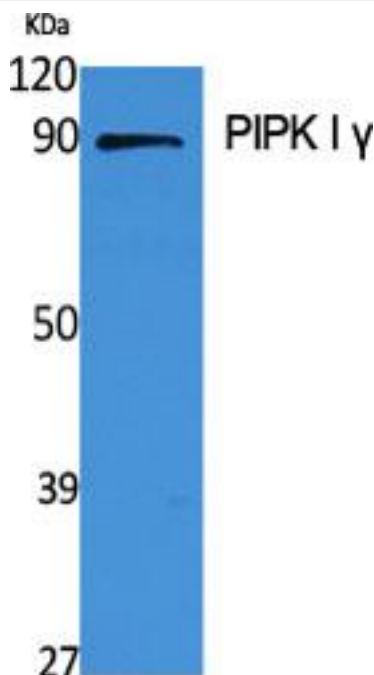
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western Blot analysis of various cells using PIPK I γ Monoclonal Antibody