

Rabbit Anti-FRYL antibody

SL16184R

Product Name:	FRYL
Chinese Name:	FRYL蛋白抗体
Alias:	2010313D22Rik; 2310004H21Rik; 2510002A14Rik; 9030227G01Rik; AF4P12; AI930088; ALL fused gene from chromosome 4p12; ALL1-fused gene from chromosome 4p12 protein; DKFZp686E205; FLJ161772; FRYL_HUMAN; FRY like; Furry homolog like (Drosophila); Furry homolog like; Furry like; KIAA0826; mKIAA0826; Protein furry homolog-like.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat, Cow, Horse, Sheep,
Applications:	ELISA=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:	339kDa
Cellular localization:	cytoplasmic
Form:	Lyophilized or Liquid
Concentration:	lmg/ml
immunogen:	KLH conjugated synthetic peptide derived from human FRYL:301-400/3013
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:	<u>PubMed</u>
Product Detail:	FRYL plays a key role in maintaining the integrity of polarized cell extensions during morphogenesis. It regulates the actin cytoskeleton and plays a key role in patterning sensory neuron dendritic fields by promoting avoidance between homologous dendrites

as well as by limiting dendritic branching. FRYL May function as a transcriptional activator.

Function:

Plays a key role in maintaining the integrity of polarized cell extensions during morphogenesis, regulates the actin cytoskeleton and plays a key role in patterning sensory neuron dendritic fields by promoting avoidance between homologous dendrites as well as by limiting dendritic branching (By similarity). May function as a transcriptional activator.

Tissue Specificity:

Widely expressed with higher expression in colon, placenta, brain and cells of lymphoid origin.

DISEASE:

Note=A chromosomal aberration involving FRYL is found in treatment-related acute lymphoblastic leukemia (ALL). Translocation t(4;11)(p12;q23) that forms a KMT2A/MLL1-FRYL fusion protein.

Similarity:

Belongs to the furry protein family.

SWISS:

O94915

Gene ID:

285527

Database links:

Entrez Gene: 285527 Human

Entrez Gene: 72313 Mouse

SwissProt: O94915 Human

Important Note:

This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.