

产品名称: ALMS1 Rabbit Polyclonal Antibody
产品货号: APRab06789



产品概述 (Summary)

产品名称 (Production Name)	ALMS1 Rabbit Polyclonal Antibody
描述 (Description)	Rabbit polyclonal Antibody
宿主 (Host)	Rabbit
应用 (Application)	WB,IHC,ICC/IF,ELISA
种属反应性 (Reactivity)	Human,Rat,Mouse

产品性能 (Performance)

偶联物 (Conjugation)	Unconjugated
修饰 (Modification)	Unmodified
同种型 (Isotype)	IgG
克隆 (Clonality)	Polyclonal
形式 (Form)	Liquid
存放说明 (Storage)	Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles.
储存溶液 (Buffer)	Liquid in PBS containing 50% glycerol, 0.5% protective protein and 0.02% New type preservative N.
纯化方式 (Purification)	Affinity purification

免疫原信息 (Immunogen)

基因名 (Gene Name)	ALMS1
别名 (Alternative Names)	ALMS1; KIAA0328; Alstrom syndrome protein 1
基因 ID (Gene ID)	7840.0
蛋白 ID (SwissProt ID)	Q8TCU4.Synthesized peptide derived from ALMS1 . at AA range: 1530-1610

产品应用 (Application)

稀释比 (Dilution Ratio)	WB 1:500-1:2000,IHC 1:100-1:300,ICC/IF 1:50-1:200,ELISA 1:20000-1:40000
蛋白分子量 (Molecular Weight)	460kDa

研究背景 (Background)

This gene encodes a protein containing a large tandem-repeat domain as well as additional low complexity regions. The

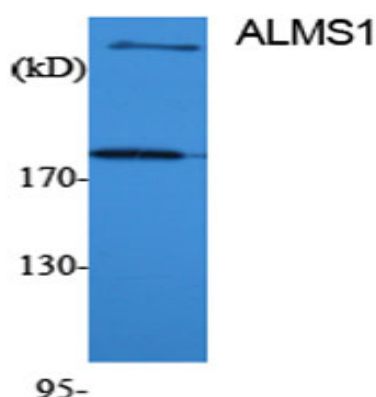
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encoded protein functions in microtubule organization, particularly in the formation and maintenance of cilia. Mutations in this gene cause Alstrom syndrome. There is a pseudogene for this gene located adjacent in the same region of chromosome 2. Alternative splice variants have been described but their full length nature has not been determined. [provided by RefSeq, Apr 2014],developmental stage:Widely expressed in fetal tissues. Detected in fetal pancreas, skeletal muscle, liver, kidney and brain (at protein level). Expressed in fetal aorta and brain.,disease:Defects in ALMS1 are the cause of Alstrom syndrome (ALMS) [MIM:203800]. Alstrom syndrome is a rare autosomal recessive disorder characterized by progressive cone-rod retinal dystrophy, neurosensory hearing loss, early childhood obesity and type 2 diabetes mellitus. Dilated cardiomyopathy, acanthosis nigricans, male hypogonadism, hypothyroidism, developmental delay and hepatic dysfunction can also be associated with the syndrome.,function:Possible role in intracellular trafficking.,PTM:Phosphorylated upon DNA damage, probably by ATM or ATR.,subcellular location:Associated with centrosomes and basal body at the base of primary cilia. During mitosis localizes to both spindle poles.,tissue specificity:Expressed in all tissues tested including adipose and pancreas. Expressed by beta-cells of the islets in the pancreas (at protein level),,

研究领域 (Research Area)

图片 (Image Data)



Western Blot analysis of various cells using ALMS1 Polyclonal Antibody

注意事项 (Note)

For research use only .