abcam

Product datasheet

Recombinant human CASK protein ab131707

5 图像

描述

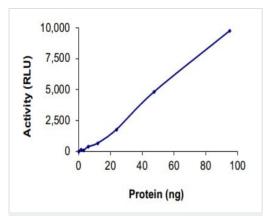
产 品名称	重组人CASK蛋白
生物活性	The specific activity of ab131707 was determined to be 3 nmol/min/mg.
纯 度	> 95 % SDS-PAGE. Assessed by densitometry. Affinity purified.
表达系 统	Baculovirus infected Sf9 cells
Accession	<u>014936</u>
蛋白长度	Protein fragment
无 动 物成分	No
性质	Recombinant
种属	Human
预测分子量	95 kDa including tags
氨基酸	1 to 564
标签	GST tag N-Terminus
技术指标	

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

应用	Functional Studies
	SDS-PAGE
	Western blot
形式	Liquid
补 充 说 明	ab204874 (PKI-alpha peptide) can be utilized as a substrate for assessing kinase activity
制备和贮存	
稳定性和存储	Shipped on dry ice. Upon delivery aliquot and store at -80℃. Avoid freeze / thaw cycles.
	pH: 7.50
	Constituents: 0.31% Glutathione, 0.002% PMSF, 0.005% DTT, 0.79% Tris HCI, 0.003% EDTA,
	25% Glycerol (glycerin, glycerine), 0.88% Sodium chloride

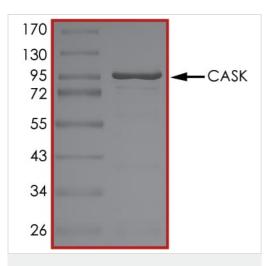
常规信息	
功能	Multidomain scaffolding protein with a role in synaptic transmembrane protein anchoring and ion channel trafficking. Contributes to neural development and regulation of gene expression via interaction with the transcription factor TRB1. Binds to cell-surface proteins, including amyloid precursor protein, neurexins and syndecans. May mediate a link between the extracellular matrix and the actin cytoskeleton via its interaction with syndecan and with the actin/spectrin-binding protein 4.1.
组织 特异性	Ubiquitous. Expression is significantly greater in brain relative to kidney, lung, and liver and in feta brain and kidney relative to lung and liver.
疾病相关	Defects in CASK are the cause of mental retardation X-linked CASK-related (MRXCASK) [MIM:300749]. Mental retardation is characterized by significantly below average general intellectual functioning associated with impairments in adaptative behavior and manifested during the developmental period. Patients with mental retardation X-linked CASK-related can manifest a severe phenotype consisting of severe intellectual deficit, congenital or postnatal microcephaly, disproportionate brainstem and cerebellar hypoplasia (MICPCH Syndrome). A milder phenotype consists of mental retardation alone or associated with nystagmus. Defects in CASK are the cause of FG syndrome type 4 (FGS4) [MIM:300422]. FG syndrome (FGS) is an X-linked disorder characterized by mental retardation, relative macrocephaly, hypotonia and constipation.
序列相似性	In the N-terminal section; belongs to the protein kinase superfamily. CAMK Ser/Thr protein kinase family. CaMK subfamily. Belongs to the MAGUK family. Contains 1 guanylate kinase-like domain. Contains 2 L27 domains. Contains 1 PDZ (DHR) domain. Contains 1 protein kinase domain. Contains 1 SH3 domain.
结 构域	The first L27 domain binds DLG1 and the second L27 domain probably binds LIN7. The protein kinase domain mediates the interaction with FCHSD2.
	Nucleus. Cytoplasm. Cell membrane.

图片

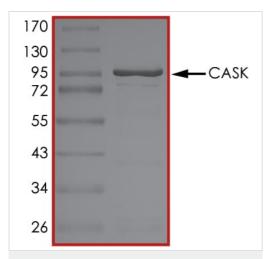


The specific activity of CASK (ab131707) was determined to be 0.24 nmol/min/mg as per activity assay protocol and was equivalent to 3.5 nmol/min/mg as per radiometric assay

Functional Studies - Recombinant human CASK protein (ab131707)



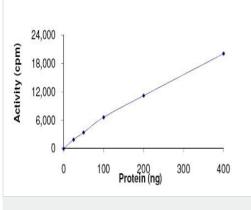
SDS-PAGE - Recombinant human CASK protein (ab131707)



SDS-PAGE - Recombinant human CASK protein (ab131707)

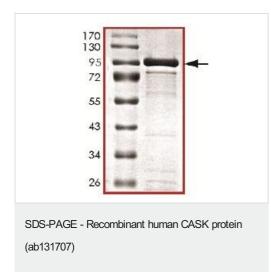
SDS PAGE analysis of ab131707

SDS PAGE analysis of ab131707



The specific activity of ab131707 was determined to be 3 nmol/min/mg.

Functional Studies - Recombinant human CASK protein (ab131707)



The purity of ab131707 was determined to be >95% by densitometry.

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