产品名称：FA58A Rabbit Polyclonal Antibody
产品货号：APRab10753

产品概述（Summary）

产品名称（Production Name）	FA58A Rabbit Polyclonal Antibody
描述（Description）	Rabbit polyclonal Antibody
宿主（Host）	Rabbit
应用（Application）	WB,ELISA
种属反应性（Reactivity）	Human,Mouse,Rat

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, 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）	FAM58A
别名（Alternative Names）	
基因 ID（Gene ID）	92002.0
蛋白 ID（SwissProt ID）	Q8N1B3.Synthesized peptide derived from human protein . at AA range: 30-110

产品应用（Application）

稀释比（Dilution Ratio）	WB 1:500-1:2000,ELISA 1:5000-1:20000
蛋白分子量（Molecular Weight）	27kDa

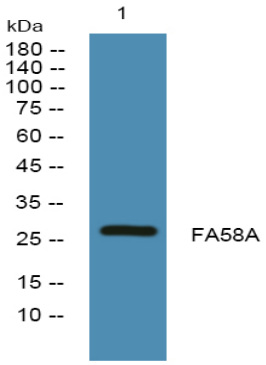
研究背景（Background）

Mutations in this gene have been shown to cause an X-linked dominant STAR syndrome that typically manifests syndactyly,

telecanthus and anogenital and renal malformations. The protein encoded by this gene contains a cyclin-box-fold domain which suggests it may have a role in controlling nuclear cell division cycles. Alternative splicing results in multiple transcript variants encoding distinct isoforms. [provided by RefSeq, Oct 2008],disease:Defects in FAM58A are the cause of toe syndactyly, telecanthus, and anogenital and renal malformations (STAR) [MIM:300707]; also known as STAR syndrome or syndactyly with renal and anogenital malformations.,function:May have a role in cell proliferation.,similarity:Belongs to the cyclin family. Cyclin-like FAM58 subfamily.,subunit:Interacts with SALL1.,

研究领域（Research Area）

图片（Image Data）



Western blot analysis of lysates from PC12 cells, primary antibody was diluted at 1:1000, 4°over night

注意事项（Note）

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