产品名称：GPR98 Rabbit Polyclonal Antibody
产品货号：APRab11711

产品概述（Summary）

产品名称（Production Name）	GPR98 Rabbit Polyclonal Antibody
描述（Description）	Rabbit polyclonal Antibody
宿主（Host）	Rabbit
应用（Application）	IHC,ICC/IF
种属反应性（Reactivity）	Human,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, 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）	GPR98
别名（Alternative Names）	KIAA0686 KIAA1943 MASS1 VLGR1
基因 ID（Gene ID）	84059.0
蛋白 ID（SwissProt ID）	Q8WXG9.Synthesized peptide derived from part region of human protein

产品应用（Application）

稀释比（Dilution Ratio）	IHC 1:50-1:300,ICC/IF 1:50-1:200
蛋白分子量（Molecular Weight）	693kDa

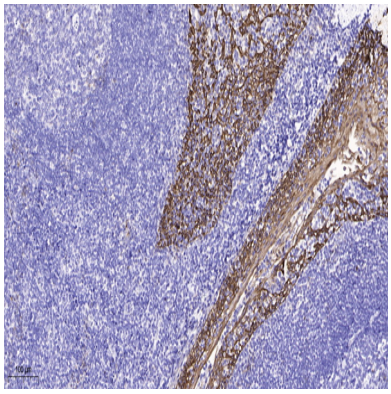
研究背景（Background）

This gene encodes a member of the G-protein coupled receptor superfamily. The encoded protein contains a 7-transmembrane receptor domain, binds calcium and is expressed in the central nervous system. Mutations in this gene are

associated with Usher syndrome 2 and familial febrile seizures. Several alternatively spliced transcripts have been described. [provided by RefSeq, Jul 2008],developmental stage:Isoform 1 is 4 times more abundant than isoform 2 in most tissues tested, despite wide variations in absolute levels of expression. Isoform 3 is expressed at about 1.5 times isoform 1 levels in most tissues examined. In fetal testis, isoform 3 is expressed almost exclusively.,disease:Defects in GPR98 are the cause of Usher syndrome type 2C (USH2C) [MIM:605472]. USH is a genetically heterogeneous condition characterized by the association of retinitis pigmentosa with sensorineural deafness. Age at onset and differences in auditory and vestibular function distinguish Usher syndrome type 1 (USH1), Usher syndrome type 2 (USH2) and Usher syndrome type 3 (USH3). USH2 is characterized by congenital mild hearing impairment with normal vestibular responses.,disease:Defects in GPR98 may be a cause of familial febrile convulsions type 4 (FEB4) [MIM:604352]; also known as familial febrile seizures 4. Febrile convulsions are seizures associated with febrile episodes in childhood without any evidence of intracranial infection or defined pathologic or traumatic cause. It is a common condition, affecting 2-5% of children aged 3 months to 5 years. The majority are simple febrile seizures (generally defined as generalized onset, single seizures with a duration of less than 30 minutes). Complex febrile seizures are characterized by focal onset, duration greater than 30 minutes, and/or more than one seizure in a 24 hour period. The likelihood of developing epilepsy following simple febrile seizures is low. Complex febrile seizures are associated with a moderately increased incidence of epilepsy.,function:Receptor that may have an important role in the development of the central nervous system.,miscellaneous:By far is the largest known cell surface protein.,similarity:Belongs to the G-protein coupled receptor 2 family. LN-TM7 subfamily.,similarity:Contains 1 GPS domain.,similarity:Contains 35 Calx-beta domains.,similarity:Contains 6 EAR repeats.,subunit:Interacts with WHRN.,tissue specificity:Expressed at low levels in adult tissues.,

研究领域（Research Area）

图片（Image Data）



Immunohistochemical analysis of paraffin-embedded human tonsil. 1, Antibody was diluted at 1:200（4° overnight）. 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200（room temperature, 45min）.

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注意事项（Note）

For research use only .