

MaxiKa Polyclonal Antibody

Catalog No: YT2666

Reactivity: Human; Mouse; Rat

Applications: WB;IHC;IF;ELISA

Target: MaxiKa

Fields: >>cGMP-PKG signaling pathway;>>Vascular smooth muscle

contraction;>>Insulin secretion;>>Renin secretion;>>Salivary

secretion;>>Pancreatic secretion

Gene Name: KCNMA1

Protein Name: Calcium-activated potassium channel subunit alpha-1

Q12791

Q08460

Human Gene Id: 3778

Human Swiss Prot

No:

Mouse Gene Id: 16531

Mouse Swiss Prot

No:

Rat Gene Id: 83731

Rat Swiss Prot No: Q62976

Immunogen: The antiserum was produced against synthesized peptide derived from human

MaxiKalpha. AA range:721-770

Specificity: MaxiKa Polyclonal Antibody detects endogenous levels of MaxiKa protein.

Formulation: Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, lgG

Dilution : WB 1:500 - 1:2000. IHC 1:100 - 1:300. ELISA: 1:40000.. IF 1:50-200

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Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 137kD

Cell Pathway: Vascular smooth muscle contraction;

Background: potassium calcium-activated channel subfamily M alpha 1(KCNMA1) Homo

sapiens MaxiK channels are large conductance, voltage and calcium-sensitive potassium channels which are fundamental to the control of smooth muscle tone and neuronal excitability. MaxiK channels can be formed by 2 subunits: the poreforming alpha subunit, which is the product of this gene, and the modulatory beta subunit. Intracellular calcium regulates the physical association between the alpha and beta subunits. Alternatively spliced transcript variants encoding different isoforms have been identified. [provided by RefSeq, Jul 2008],

Function: alternative products:May be partially controlled by hormonal stress. Additional

isoforms seem to exist, disease: Defects in KCNMA1 are the cause of generalized epilepsy and paroxysmal dyskinesia (GEPD) [MIM:609446]. Epilepsy is one of the most common and debilitating neurological disorders. Paroxysmal dyskinesias are neurological disorders characterized by sudden, unpredictable, disabling attacks of involuntary movement often requiring life-long treatment. The coexistence of epilepsy and paroxysmal dyskinesia in the same individual or family is an increasingly recognized phenomenon. Patients manifest absence seizures, generalized tonic-clonic seizures, paroxysmal nonkinesigenic dyskinesia, involuntary dystonic or choreiform movements. Onset is usually in

childhood and patients may have seizures only, dyskinesia only, or both.,domain:The calcium bowl constitutes one of the Ca(2+) sensors an

Subcellular
Location:

Cell membrane ; Multi-pass membrane protein .

Expression: Widely expressed. Except in myocytes, it is almost ubiquitously expressed.

Sort : 9424

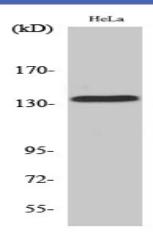
No4: 1

Host: Rabbit

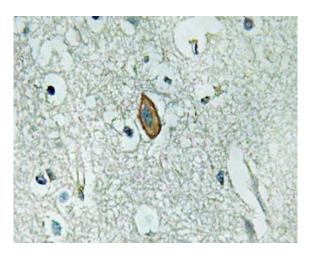
Modifications: Unmodified

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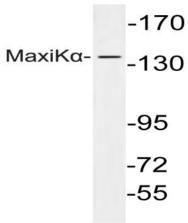
Products Images



Western Blot analysis of various cells using MaxiKa Polyclonal Antibody



Immunohistochemistry analysis of MaxiK α antibody in paraffinembedded human brain tissue.



Western blot analysis of lysate from HeLa cells, using MaxiK α antibody.