



NHE-6 Monoclonal Antibody

Catalog No	YP-mAb-16482
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	SLC9A6
Immunogen	The antiserum was produced against synthesized peptide derived from human SLC9A6. AA range:551-600
Specificity	NHE-6 Monoclonal Antibody detects endogenous levels of NHE-6 protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	3732426M5;64352C2Rik;KIAA267;mKIAA267;MRSA;Na(+)/H(+) exchanger 6; NHE-6;NHE6;OTTHUMP2489;OTTHUMP249;RGD1563582;RP11-274K13.1; RP23-15E2.4;SL9A6_HUMAN;SLC9A6;Sodium/hydrogen exchanger 6;Solute carrier family 9 (sodium/hydrogen exchanger);isoform 6;member 6;Solute carrier family 9 member 6;solute carrier family 9;subfamily A (NHE6;cation proton antiporter 6);KIAA0267
Observed Band	75kD
Calculated Molecular Weight	75kD
Cell Pathway	Endosome membrane ; Multi-pass membrane protein . Is present in the recycling compartments including early and recycling endosomes, and only appears transiently on the plasma membrane.; [Isoform 2]: Recycling endosome membrane ; Multi-pass membrane protein .
Tissue Specificity	Ubiquitous; but is most abundant in mitochondrion-rich tissues such as brain, skeletal muscle and heart.
Function	caution:Was initially identified as a mitochondrial inner membrane protein (PubMed:9507001), but was later shown to be localized in early and recycling endosomes and not mitochondria (PubMed:11940519).,disease:Defects in SLC9A6 are the cause of mental retardation syndromic X-linked Christianson



type (MRXSC) [MIM:300243]; also known as MRXS-Christianson or X-linked Angelman-like syndrome. The phenotype is characterized by profound mental retardation, epilepsy, ataxia, and microcephaly, and showed phenotypic overlap with Angelman syndrome. function: Electroneutral exchange of protons for Na(+) and K(+) across the early and recycling endosome membranes. Contributes to calcium homeostasis. similarity: Belongs to the monovalent cation:proton antiporter 1 (CPA1) transporter (TC 2.A.36) family. subcellular location: Is present in the recycling compartments including early and recycling endosomes,

Background

This gene encodes a sodium-hydrogen exchanger that is a member of the solute carrier family 9. The encoded protein localizes to early and recycling endosomes and may be involved in regulating endosomal pH and volume. Defects in this gene are associated with X-linked syndromic mental retardation, Christianson type. Alternate splicing results in multiple transcript variants. [provided by RefSeq, Apr 2010],

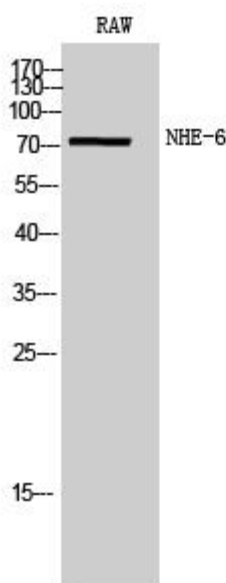
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western Blot analysis of various cells using NHE-6 Monoclonal Antibody