SLC9A1 (Human) Recombinant Protein (Q01)

Catalog # H00006548-Q01 Size 25 ug, 10 ug

Applications



Specification	
Product Description	Human SLC9A1 partial ORF (AAH12121, 31 a.a 130 a.a.) recombinant protein with GST-tag at N -terminal.
Sequence	VLRSHGLQLSPTASTIRSSEPPRERSIGDVTTAPPEVTPESRPVNHSVTDHGMKPRKAFPVLGIDY THVRTPFEISLWILLACLMKIGFHVIPTISSIVP
Host	Wheat Germ (in vitro)
Theoretical MW (kDa)	36.41
Interspecies Antigen Sequence	Mouse (81); Rat (79)
Preparation Method	in vitro wheat germ expression system
Purification	Glutathione Sepharose 4 Fast Flow
Quality Control Testing	12.5% SDS-PAGE Stained with Coomassie Blue.
Storage Buffer	50 mM Tris-HCI, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.
Note	Best use within three months from the date of receipt of this protein.



Applications

- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

Gene Info — SLC9A1	
Entrez GenelD	<u>6548</u>
GeneBank Accession#	<u>BC012121</u>
Protein Accession#	AAH12121
Gene Name	SLC9A1
Gene Alias	APNH, FLJ42224, NHE1
Gene Description	solute carrier family 9 (sodium/hydrogen exchanger), member 1
Omim ID	<u>107310</u>
Gene Ontology	Hyperlink
Gene Summary	The Na+/H+ antiporter (SLC9A1) is a ubiquitous membrane-bound enzyme involved in pH regulat ion of vertebrate cells. It is specifically inhibited by the diuretic drug amiloride and activated by a v ariety of signals including growth factors, mitogens, neurotransmitters, tumor promoters, and other s (Mattei et al., 1988 [PubMed 2846238]).[supplied by OMIM
Other Designations	Na+/H+ antiporter, amiloride-sensitive Na-Li countertransporter OTTHUMP00000004468 sodium/ hydrogen exchanger 1 solute carrier family 9 (sodium/hydrogen exchanger), isoform 1 (antiporter, Na+/H+, amiloride sensitive) solute carrier family 9 (sodium/hydroge

Pathway

- <u>Cardiac muscle contraction</u>
- Regulation of actin cytoskeleton

😵 Abnova

Product Information

Disease

- Cerebrovascular Accident
- Genetic Predisposition to Disease
- <u>Seizures</u>