bs-19821R

[Primary Antibody]

SLC32A1 Rabbit pAb



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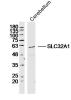
- DATASHEET	
	/
(IONALITY' POlyclonal	(1:100-500)
GenelD: 140679 SWISS: Q9H598 IF (1:10	1:100-500) 00-500)
Target: SLC32A1 Reactivity: Mouse,	Pat
	ted: Human, Rabbit,
Purification: affinity purified by Protein A	
Concentration: 1mg/ml Predicted MW.: 57 kDa	
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Subcellular Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated Subcellular freeze/thaw cycles. Location:	embrane ,Cytoplasm
Background: Synaptic transmission involves the controlled exocytosis of vesicles containing specific neurotransmitters. Usually, neurotransmitters are synthesized in the cytoplasm of the cell and must be transported into synaptic vesicles for release. The vesicular GABA transporter (VGAT) is responsible for loading γ-aminobutyric acid (GABA), an inhibitory neurotransmitter, from neuronal cytoplasm into synaptic vesicles and is expressed only in the nerve endings of inhibitory neurons that contain GABA and/or glycine. During neocortical development, VGAT expression barely precedes the maturation of inhibitory synaptogenesis, suggesting that it may contribute to the development of neocortical GABAergic circuitry. VGAT may also play a role in epileptogenesis and the recovery mechanisms that occur after a spontaneous seizure.	
- VALIDATION IMAGES	

189 = 100 -100 -63 - - - SLC32A1 48 -

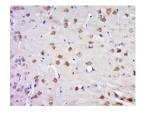
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Sample:heart (Mouse) Lysate at 40 ug Primary: Anti-SLC32A1(bs-19821R)at 1/300 dilution Secondary: IRDye800CW Goat Anti-RabbitIgG at 1/20000 dilution Predicted band size: 57kD Observed band size: 60kD



Sample:Cerebellum (Mouse) Lysate at 40 ug Primary: Anti-SLC32A1(bs-19821R)at 1/300 dilution Secondary: IRDye800CW Goat Anti-RabbitlgG at 1/20000 dilution Predicted band size: 57kD Observed band size: 57kD



Tissue/cell: rat brain tissue; 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-SLC32A1 Polyclonal Antibody, Unconjugated(bs-19821R) 1:500, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining