

SNAT1/SLC38A1 (D9L2P) Rabbit mAb



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Applications: W, IP	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 50-70	Source/Isotype: Rabbit IgG	UniProt ID: #Q9H2H9	Entrez-Gene Id: 81539
Product Usage Information		Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:50	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		SNAT1/SLC38A1 (D9L2P) Rabbit mAb recognizes endogenous levels of total SNAT1/SLC38A1 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gly37 of human SNAT1/SLC38A1 protein.				
Background		SNAT1/SLC38A1 belongs to the system A transporters that mediate Na ⁺ -dependent transport of short-chain neutral amino acids such as alanine, serine, and glutamine. SNAT1/SLC38A1 mediates the uptake of glutamine in neurons and plays a crucial role in glutamate-glutamine cycle. Steep concentration gradients across the plasma membrane are achieved by coupling of the electrochemical sodium gradient to amino acid transport. This allows a unidirectional mode of transport for SNAT1/SLC38A1. Upregulation of SNAT1/SLC38A1 by neurotrophic factors is key to dendritic growth and branching of cortical neurons. High expression of SNAT1/SLC38A1 is found in cerebral cortex primarily in neurons and to a lesser extent in astrocytes (1-4). Elevated SNAT1/SLC38A1 expression is prominent in human solid tumors including gliomas, hepatocellular carcinomas and human breast cancer (5-8). Research studies show that an aberrant SNAT1/SLC38A1 expression profile correlates with solid tumor recurrence and poor prognosis in patients with cholangiocarcinoma (9).				
Background References		1. Yao, D. et al. (2000) <i>J Biol Chem</i> 275, 22790-7. 2. Mackenzie, B. et al. (2003) <i>J Biol Chem</i> 278, 23720-30. 3. Chaudhry, F.A. et al. (2002) <i>J Cell Biol</i> 157, 349-55. 4. Yu, W.L. et al. (2011) <i>J Surg Res</i> 171, 663-8. 5. Melone, M. et al. (2004) <i>Cereb Cortex</i> 14, 562-74. 6. Kondoh, N. et al. (2007) <i>Int J Oncol</i> 31, 81-7. 7. Sidoryk, M. et al. (2004) <i>Neuroreport</i> 15, 575-8. 8. Wang, K. et al. (2013) <i>BMC Cancer</i> 13, 343. 9. Burkhalter, J. et al. (2007) <i>J Biol Chem</i> 282, 5152-9.				

Species Reactivity Species reactivity is determine

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X

TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key W: Western Blotting IP: Immunoprecipitation

Cross-Reactivity Key H: Human M: Mouse R: Rat

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