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SNAT1/SLC38A1 (D9L2P) Rabbit



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For Research Use Only. Not for Use in Diagnostic Procedures.							
Applications: WB, 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	Ар	plication		Dilution			
	We	stern Blotting		1:1000			
	Imr	munoprecipitation			1:50		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA, 50% glycerol and less 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				erol and less than	
Specificity / Sensi	itivity SNA	SNAT1/SLC38A1 (D9L2P) Rabbit mAb recognizes endogenous levels of total SNAT1/SLC38A1 protein.					
Source / Purificati		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gly37 of human SNAT1/SLC38A1 protein.					
Background	chai /SLC cycl elec SNA brar neu hum Res /SLC	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 Refe	2. M 3. C 4. Y 5. M 6. K 7. S 8. W	lackenzie, B. et al. (haudhry, F.A. et al. u, W.L. et al. (2011) lelone, M. et al. (200 ondoh, N. et al. (200 idoryk, M. et al. (200 /ang, K. et al. (2013	D. et al. (2000) <i>J Biol Chem</i> 275, 22790-7. enzie, B. et al. (2003) <i>J Biol Chem</i> 278, 23720-30. dhry, F.A. et al. (2002) <i>J Cell Biol</i> 157, 349-55. L. et al. (2011) <i>J Surg Res</i> 171, 663-8. le, M. et al. (2004) <i>Cereb Cortex</i> 14, 562-74. oh, N. et al. (2007) <i>Int J Oncol</i> 31, 81-7. or, M. et al. (2004) <i>Neuroreport</i> 15, 575-8. or, K. et al. (2013) <i>BMC Cancer</i> 13, 343. alter, J. et al. (2007) <i>J Biol Chem</i> 282, 5152-9.				

Species Reactivity

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

WB: Western Blotting IP: Immunoprecipitation

Cross-Reactivity Key

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster

X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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1/1/24, 9:32 AM **Limited Uses**

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