

## KCC2 (D1R2R) Rabbit mAb



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## For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> W, IP, IF-F	<b>Reactivity:</b> H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 130-260	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #Q9H2X9	Entrez-Gene Id: 57468
Product Usage Information		Application Western Blotting Immunoprecipitation Immunofluorescence (Frozen)			<b>Dilution</b> 1:1000 1:100 1:100	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
		For a carrier free (BSA and azide free) version of this product see product #56440.				
Specificity/Sensitivity		KCC2 (D1R2R) Rabbit mAb recognizes endogenous levels of total KCC2 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ala970 of human KCC2 protein.				
Background		The potassium/chloride cotransporter 2 (KCC2, SLC12A5) is a neuron-specific transport protein responsible for regulating the cotransport of potassium and chloride ions. KCC2 uses the energy of the electrochemical potassium gradient to export chloride ions from cells, therefore maintaining intracellular chloride ion concentrations in mature neurons (1,2). The intracellular concentration of chloride ions determines the neuronal response to the inhibitory neurotransmitter GABA and glycine. As a result, KCC2 can play a critical role in regulating neuronal excitability in mature central nervous system neurons (3-5). Altered KCC2 expression and reduced KCC2 activity can result in an increase in intracellular chloride ion concentrations and subsequent hyperexcitability of neuronal systems. Cases of aberrant KCC2 function are associated with neurological disorders, such as multiple forms of epilepsy, neuropathic pain, and schizophrenia (6-10).				
Background References		<ol> <li>Payne, J.A. et al. (1996) J Biol Chem 271, 16245-52.</li> <li>Delpire, E. (2000) News Physiol Sci 15, 309-12.</li> <li>Rivera, C. et al. (1999) Nature 397, 251-5.</li> <li>Woo, N.S. et al. (2002) Hippocampus 12, 258-68.</li> <li>Zhu, L. et al. (2005) J Neurophysiol 93, 1557-68.</li> <li>Arion, D. and Lewis, D.A. (2011) Arch Gen Psychiatry 68, 21-31.</li> <li>Hyde, T.M. et al. (2011) J Neurosci 31, 11088-95.</li> <li>Kaila, K. et al. (2014) Curr Opin Neurobiol 26, 34-41.</li> <li>Mòdol, L. et al. (2014) Pain 155, 1577-90.</li> <li>Kahle, K.T. et al. (2014) EMBO Rep 15, 766-74.</li> </ol>				

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

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 IF-F: Immunofluorescence (Frozen)

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

**Western Blot Buffer** 

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