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MDR1/ABCB1 Antibody

Applications: W, IP	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 130-180	Source/Isotype: Rabbit	UniProt ID: #P08183	Entrez-Gene Id: 5243
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 and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		MDR1/ABCB1 Antibody recognizes endogenous levels of total MDR1 protein. This antibody also cross- reacts with a 47 kDa protein of unknown origin in some cell lines.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Glu643 of human MDR1 protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		MDR1/ABCB1 belongs to the Mdr/Tap subfamily of the ATP-binding cassette transporter superfamily (1). Multidrug resistance 1 (MDR1) serves as an efflux pump for xenobiotic compounds with broad substrate specificity. MDR1 substrates include therapeutic agents such as actinomycin D, etoposide, imatinib, and doxorubicin, as well as endogenous molecules including β-amyloids, steroid hormones, lipids, phospholipids, cholesterol, and cytokines (2). Research studies have shown that MDR1 reduces drug accumulation in cancer cells, allowing the development of drug resistance (3-5). On the other hand, MDR1 expressed in the plasma membrane of cells in the blood-brain, blood-cerebral spinal fluid, or blood-placenta barriers restricts the permeability of drugs into these organs from the apical or serosal side (6,7). MDR1 is also expressed in normal tissues with excretory function such as small intestine, liver, and kidney (7). Intracellular MDR1 has been detected in the ER, vesicles, and nuclear envelope, and has been associated with cell trafficking machinery (8). Other reported functions of MDR1 include viral resistance, cytokine trafficking (9,10), and lipid homeostasis in the peripheral and central nervous system (11-13).				
Background References		 Furuya, K.N. et al. (1997) <i>Cancer Res</i> 57, 3708-16. Litman, T. et al. (1997) <i>Biochim Biophys Acta</i> 1361, 169-76. Chen, C.J. et al. (1986) <i>Cell</i> 47, 381-9. Kartner, N. et al. (1983) <i>Cancer Res</i> 43, 4413-9. Chen, G. et al. (1997) <i>J Biol Chem</i> 272, 5974-82. Brinkmann, U. and Eichelbaum, M. (2001) <i>Pharmacogenomics J</i> 1, 59-64. Fromm, M.F. (2004) <i>Trends Pharmacol Sci</i> 25, 423-9. Miller, D.S. et al. (1999) <i>Annu Rev Pharmacol Toxicol</i> 39, 361-98. Raviv, Y. et al. (2000) <i>FASEB J</i> 14, 511-5. Meijer, O.C. et al. (2003) <i>J Endocrinol</i> 178, 13-8. Karssen, A.M. et al. (2009) <i>Clin Chim Acta</i> 403, 198-202. 				
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		W: Western Blotting IP: Immunoprecipitation				
Cross-Reactivity Key		H: Human				
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