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**Patents** 

## MATE1/SLC47A1 (D4C6Z) Rabbit



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

Applications: WB, IP	Reactivity: H	Sensitivity: Endogenous	<b>MW (kDa):</b> 48-52	Source/Isotype: Rabbit IgG	UniProt ID: #Q96FL8	Entrez-Gene Id 55244	
Product Usage Information	Ap	plication		Dilution			
	We	stern Blotting		1:1000			
	Imr	nunoprecipitation			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. <i>Do not aliquot the antibody.</i>					
Specificity / Sensit	t <b>ivity</b> MAT	MATE1/SLC47A1 (D4C6Z) Rabbit mAb recognizes endogenous levels of total MATE1 protein.					
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Leu505 of human MATE1 protein.					
Background	antij of h end age SLC ben corr	The multidrug and toxin extrusion protein 1 (MATE1, SLC47A1) is a proton-coupled, organic cation antiporter located at the apical membrane of proximal kidney epithelial cells and the canalicular membrane of hepatocytes (1). MATE1 mediates the secretion of organic cations including drugs, toxins, and endogenous metabolites, into bile and urine (2,3). Substrates of MATE1 include multiple therapeutic agents, including metformin, cisplatin, acyclovir, and cephalexin (4,5). Polymorphisms in the corresponding <i>SLC47A1</i> gene may affect the rate of renal clearance of certain cationic drugs, limiting the therapeutic benefits of these agents (6). Specifically, research studies demonstrate that <i>SLC47A1</i> allelic variation correlates with differences in renal clearance rates of metformin (7), which may have an effect on the therapeutic impact of this drug in individuals diagnosed with type 2 diabetes (8).					
Background Refer	2. O 3. M 4. Ta 5. H 6. S 7. C	<ol> <li>Otsuka, M. et al. (2005) Proc Natl Acad Sci U S A 102, 17923-8.</li> <li>Omote, H. et al. (2006) Trends Pharmacol Sci 27, 587-93.</li> <li>Motohashi, H. and Inui, K. (2013) AAPS J 15, 581-8.</li> <li>Tanihara, Y. et al. (2007) Biochem Pharmacol 74, 359-71.</li> <li>Hume, W.E. et al. (2013) Bioorg Med Chem 21, 7584-90.</li> <li>Staud, F. et al. (2013) Int J Biochem Cell Biol 45, 2007-11.</li> <li>Christensen, M.M. et al. (2013) Pharmacogenet Genomics 23, 526-34.</li> <li>Becker, M.L. et al. (2009) Diabetes 58, 745-9.</li> </ol>					

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

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

milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

**Applications Key** WB: Western Blotting IP: Immunoprecipitation

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

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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