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Integrin α4 Antibody

Applications: W, IP	Reactivity: H	Sensitivity: Endogenous	<b>MW (kDa):</b> 70, 140, 150, (180)	<b>Source/Isotype:</b> Rabbit	<b>UniProt ID:</b> #P13612	Entrez-Gene Id: 3676	
Product Usage Information	3	Application Western Blotting Immunoprecipitatio	n		<b>Dilution</b> 1:1000 1:100		
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/Ser	sitivity	Integrin α4 Antibody detects endogenous levels of integrin α4 mature protein (150kDa), α4 precursor protein (140kDa), as well as 70 kDa cleaved C-terminal α4 fragment.					
Source / Purifi	cation	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser1027 of human integrin α4. Antibodies are purified by peptide affinity chromatography.					
Background		Integrins are α/β heterodimeric cell surface receptors that play a pivotal role in cell adhesion and migration, as well as in growth and survival (1,2). The integrin family contains at least 18 α and 8 β subunits that form 24 known integrins with distinct tissue distribution and overlapping ligand specificities (3). Integrins not only transmit signals to cells in response to the extracellular environment (outside-in signaling), but also sense intracellular cues to alter their interaction with the extracellular environment (inside-out signaling) (1,2). A pair of important α4 integrins, α4β1 and α4β7, interact with VCAM-1, fibronectin, and MAdCAM-1 at cell adhesions (3). Gene knockout and antibody blocking research reveal that α4 integrins play important roles in embryonic liver and heart development and in fetal lymphocyte homing (4-6). Phosphorylation at Ser988 within the cytoplasmic tail of integrin α4 blocks binding to paxillin and promotes leading edge migration (7,8). On SDS-PAGE, integrin α4 can migrate at several different apparent molecular sizes, a 150 kDa mature protein and a 140 kDa precursor protein (a 180 kDa protein also exists under mild non-reducing conditions) (9). Integrin α4 has a cleavage site at Arg558, which results in a small portion of the protein as either an 80 kDa N-terminal or 70 kDa C-terminal fragment (10).					
Background R	eferences	<ol> <li>Hood, J.D. and Cheresh, D.A. (2002) Nat Rev Cancer 2, 91-100.</li> <li>Liu, S. et al. (2000) J Cell Sci 113 ( Pt 20), 3563-71.</li> <li>Plow, E.F. et al. (2000) J Biol Chem 275, 21785-8.</li> <li>Bonder, C.S. et al. (2005) Immunity 23, 153-63.</li> <li>Arroyo, A.G. et al. (1999) Immunity 11, 555-66.</li> <li>Yang, J.T. et al. (1995) Development 121, 549-60.</li> <li>Nishiya, N. et al. (2005) J Cell Biol 7, 343-52.</li> <li>Alon, R. et al. (2005) J Cell Biol 171, 1073-84.</li> <li>Teixidó, J. et al. (1992) J Biol Chem 267, 1786-91.</li> <li>Pujades, C. et al. (1996) Biochem J 313 ( Pt 3), 899-908.</li> </ol>					
Species Reacti	vity	Species reactivity is o	determined by testing	in at least one approve	ed application (e.g.,	western blot).	
Western Blot E	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 K	ey	W: Western Blotting IP: Immunoprecipitation					
Cross-Reactivi	ty Key	H: Human					
Trademarks ar	nd Patents	Cell Signaling Technology is a trademark of Cell Signaling Technology, Inc.					

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