L/1/24, 12:55 PM Pho Revision 1	spho-Myosir	Ayosin IIa (Ser1943) (D7Z7T) Rabbit mAb (#14611) Datasheet Without Images Cell Signaling Tec					
Phospho-Myosin IIa (Ser1943) (D7Z7T) Rabbit mAb					Cell Signaling		
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For Research Use Only. Not for Use in Applications: Reactivity: WB, IF-IC H		Sensitivity: Endogenous	MW (kDa): 230	Source/Isotype: Rabbit IgG	UniProt ID: #P35579	Entrez-Gene Id: 4627	
Product Usage	Арр	Application				Dilution	
Information	Wes	Western Blotting				1:1000	
	Imm	Immunofluorescence (Immunocytochemistry)				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. Do not aliquot the antibody.					
Specificity / Sensitivi		Phospho-Myosin IIa (Ser1943) (D7Z7T) Rabbit mAb recognizes endogenous levels of myosin IIa protein only when phosphorylated at Ser1943.					
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser1943 of human myosin II protein.					
Background Nonmuscle myosin is an actin-based motor protein essential to cell motility, cell division, migration, adhesion, and polarity. The holoenzyme consists of two identical heavy chains and two sets of light chains (MLCs) regulate myosin II activity and stability. The heavy chains (NMHCs) are enco by three genes, <i>MYH9</i> , <i>MYH10</i> , and <i>MYH14</i> , which generate three different nonmuscle myosin II is light lia, IIb, and IIc, respectively (reviewed in 1). While all three isoforms perform the same enzymatic task binding to and contracting actin filaments coupled to ATP hydrolysis, their cellular functions do not ap to be redundant and they have different subcellular distributions (2-5). The carboxy-terminal tail doma myosin II is important in isoform-specific subcellular localization (6). Research studies have shown the phosphorylation of myosin IIa at Ser1943 contributes to the regulation of breast cancer cell migration.						two sets of light chains. MHCs) are encoded scle myosin II isoforms, me enzymatic tasks, nctions do not appear terminal tail domain of es have shown that	
Background References 1. Conti, M.A. and Adelstein, R.S. (2008) J Cell Sci 121, 11-18. 2. Sandquist, J.C. et al. (2006) J Biol Chem 281, 35873-83. 3. Even-Ram, S. et al. (2007) Nat Cell Biol 9, 299-309. 4. Vicente-Manzanares, M. et al. (2007) J Cell Biol 176, 573-80. 5. Wylie, S.R. and Chantler, P.D. (2008) Mol Biol Cell 19, 3956-68. 6. Sandquist, J.C. and Means, A.R. (2008) Mol Biol Cell 19, 5156-67. 7. Dulyaninova, N.G. et al. (2007) Mol Biol Cell 18, 3144-55.							
Species Reactivity	Specie	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 nonfat dry milk, 1X TBS, 0.1% Tween $^{\circ}$ 20 at 4°C with gentle shaking, overnight.					
Applications Key	WB:	WB: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)					
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 						

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 **Trademarks and** Cell Signaling Technology is a trademark of Cell Signaling Technology, Inc. All other trademarks are the property of their respective owners. Visit cellsignal.com/trademarks for more Patents information. Except as otherwise expressly agreed in a writing signed by a legally authorized representative of CST, the Limited Uses following terms apply to Products provided by CST, its affiliates or its distributors. Any Customer's terms and

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