

## Phospho-Lamin A/C (Ser22) (D2B2E) Rabbit mAb



Orders: 877-616-CELL (2355) orders@cellsignal.com

Support: 877-678-TECH (8324)

Web: info@cellsignal.com

cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

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

<b>Applications:</b> W, IP	Reactivity: H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 69,78	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P02545	Entrez-Gene Id: 4000
Product Usage Information	2	<b>Application</b> Western Blotting Immunoprecipitation			<b>Dilution</b> 1:1000 1:100	
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/Sensitivity		Phospho-Lamin A/C (Ser22) (D2B2E) Rabbit mAb recognizes endogenous levels of lamin A/C protein only when phosphorylated at Ser22.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser22 of human lamin A/C protein.				
Background		Lamins are nuclear membrane structural components that are important in maintaining normal cell functions such as cell cycle control, DNA replication, and chromatin organization (1-3). Lamin A/C is cleaved by caspase-6 and serves as a marker for caspase-6 activation. During apoptosis, lamin A/C is specifically cleaved into a large (41-50 kDa) and a small (28 kDa) fragment (3,4). The cleavage of lamins results in nuclear dysregulation and cell death (5,6).~Phosphorylation of lamin A/C at Ser22 was identified <i>in vivo</i> in several cell lines by mass spectrometry analysis in proteomic screens. The surrounding sequence is a typical MAPK/CDK phosphorylation motif, which implicates a role in the cell cycle and mitosis (7-11).				
Background References		<ol> <li>Gruenbaum, Y. et al. (2000) J Struct Biol 129, 313-23.</li> <li>Yabuki, M. et al. (1999) Physiol Chem Phys Med NMR 31, 77-84.</li> <li>Goldberg, M. et al. (1999) Crit Rev Eukaryot Gene Expr 9, 285-93.</li> <li>Orth, K. et al. (1996) J Biol Chem 271, 16443-6.</li> <li>Oberhammer, F.A. et al. (1994) J Cell Biol 126, 827-37.</li> <li>Rao, L. et al. (1996) J Cell Biol 135, 1441-55.</li> <li>Lowery, D.M. et al. (2007) EMBO J 26, 2262-73.</li> <li>Molina, H. et al. (2007) Proc Natl Acad Sci U S A 104, 2199-204.</li> <li>Beausoleil, S.A. et al. (2006) Nat Biotechnol 24, 1285-92.</li> <li>Nousiainen, M. et al. (2006) Proc Natl Acad Sci U S A 103, 5391-6.</li> <li>Beausoleil, S.A. et al. (2004) Proc Natl Acad Sci U S A 101, 12130-5.</li> </ol>				

**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 M: Mouse R: Rat

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