

## Phospho-Connexin 43 (Ser368) (D6W8P) Rabbit mAb



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## 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> 42, 44, 46	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P17302	Entrez-Gene Id: 2697
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation		<b>Dilution</b> 1:1000 1:50		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Phospho-Connexin 43 (Ser368) (D6W8P) Rabbit mAb recognizes endogenous levels of connexin 43 protein only when phosphorylated at Ser368.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser368 of human connexin 43 protein.				
Background		Connexin 43 (Cx43) is a member of the large family of gap junction proteins. Connexins assemble as a hexamer and are transported to the plasma membrane to create a hemichannel that can associate with hemichannels on nearby cells to create cell-to-cell channels. Clusters of these channels assemble to make gap junctions. Gap junction communication is important in development and regulation of cell growth. Phosphorylation of Cx43 is important in regulating assembly and function of gap junctions (1,2). Ser368 of Cx43 is phosphorylated by protein kinase C (PKC) after activation by phorbol esters, which decreases cell-to-cell communication (3). Src can interact with and phosphorylate Cx43 to alter gap junction communication (4,5).				
Background References		<ol> <li>Musil, L.S. et al. (1990) J Cell Biol 111, 2077-88.</li> <li>Musil, L.S. and Goodenough, D.A. (1991) J Cell Biol 115, 1357-74.</li> <li>Lampe, P.D. et al. (2000) J Cell Biol 149, 1503-12.</li> <li>Giepmans, B.N. et al. (2001) J Biol Chem 276, 8544-9.</li> <li>Lin, R. et al. (2001) J Cell Biol 154, 815-27.</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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