## Acidic FGF (11H11) 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

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<b>Applications:</b> W, IP	Reactivity: H	<b>Sensitivity:</b> Recombinant protein	<b>MW (kDa):</b> 17	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #P05230	Entrez-Gene Id 2246
Product Usage Information		ApplicationDilutionWestern Blotting1:1000Immunoprecipitation1:50				
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/Sensitivity		Acidic FGF (11H11) Rabbit mAb detects recombinant human aFGF proteins at various concentration. This antibody does not cross-react with human bFGF.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with recombinant human aFGF proteins.				
Background		Fibroblast growth factors are a family of broad-spectrum growth factors influencing a plethora of cellular activities. The interaction of at least 23 ligands, four receptors, and multiple coreceptors provides a dramatic complexity to a signaling system capable of effecting a multitude of responses (1,2). Basic fibroblast growth factor (bFGF or FGF2), initially identified as a mitogen with prominent angiogenic properties, is now recognized as a multifunctional growth factor (3). It is clear that bFGF produces its biological effects in target cells by signaling through cell-surface FGF receptors. bFGF binds to all four FGF receptors. Ligand binding induces receptor dimerization and autophosphorylation, allowing binding and activation of cytoplasmic downstream target proteins, including FRS-2, PLC, and Crk (4,5). The FGF signaling pathway appears to play a significant role not only in normal cell growth regulation but also in tumor development and progression (6). Acidic FGF (aFGF or FGF1) is another extensively investigated protein of the FGF family. aFGF shares 55% DNA sequence homology with bFGF. These two growth factors are ubiquitously expressed and exhibit a wide spectrum of similar biological activities with quantitative differences likely due to variations in receptor affinity or binding (7).				
Background References		<ol> <li>Powers, C.J. et al. (2000) Endocrine-Related Cancer 7, 165-197.</li> <li>Bansal, R. (2002) Dev. Neurosci. 24, 35-46.</li> <li>Morrison, R.S. et al. (1994) J. Neurooncol. 18, 207-216.</li> <li>Kouhara, H. et al. (1997) Cell 89, 693-702.</li> <li>Mohammadi, M. et al. (1991) Mol. Cell. Biol. 11, 5068-5078.</li> <li>Bikfalvi, A. (1995) Eur. J. Cancer 31A, 1101-1104.</li> <li>Ledoux, D. et al. (1992) Prog. Growth Factor Res. 4, 107-120.</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				
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