



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

#80994 store at +4C

Annexin A1 (D5V2T) XP[®] Rabbit mAb (PE Conjugate)

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

Applications: FC-FP	Reactivity: H	Sensitivity: Endogenous	Source/Isotype: Rabbit IgG	UniProt ID: #P04083	Entrez-Gene Id: 301
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Product Usage Information	Application Flow Cytometry (Fixed/Permeabilized)	Dilution 1:50
Storage	Supplied in PBS (pH 7.2), less than 0.1% sodium azide and 2 mg/ml BSA. Store at 4°C. Do not aliquot the antibody. Protect from light. Do not freeze.	
Specificity/Sensitivity	Annexin A1 (D5V2T) XP [®] Rabbit mAb (PE Conjugate) recognizes endogenous levels of total annexin A1 protein.	
Source / Purification	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of human annexin A1 protein.	
Description	This Cell Signaling Technology antibody is conjugated to phycoerythrin (PE) and tested in-house for direct flow cytometric analysis in human cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated Annexin A1 (D5V2T) XP [®] Rabbit mAb #32934.	
Background	The annexin superfamily consists of 13 calcium or calcium and phospholipid binding proteins with high biological and structural homology (1). Annexin-1 (ANXA1) is the first characterized member of the annexin family of proteins and is able to bind to cellular membranes in a calcium-dependent manner, promoting membrane fusion and endocytosis (2-4). Annexin A1 has anti-inflammatory properties and inhibits phospholipase A2 activity (5,6). Annexin A1 can accumulate on internalized vesicles after EGF-stimulated endocytosis and may be required for a late stage in inward vesiculation (7). Phosphorylation by PKC, EGFR, and Chak1 results in inhibition of annexin A1 function (8-10). Annexin A1 has also been identified as one of the 'eat-me' signals on apoptotic cells that are to be recognized and ingested by phagocytes (11). Annexin A1, as an endogenous anti-inflammatory mediator, has roles in many diverse cellular functions, such as membrane aggregation, inflammation, phagocytosis, proliferation, apoptosis, and tumorigenesis and cancer development (12-14).	
Background References	<ol style="list-style-type: none"> 1. Raynal, P. and Pollard, H.B. (1994) <i>Biochim Biophys Acta</i> 1197, 63-93. 2. Blackwell, G.J. et al. (1980) <i>Nature</i> 287, 147-9. 3. Rothhut, B. et al. (1983) <i>Biochem Biophys Res Commun</i> 117, 878-84. 4. Hirata, F. et al. (1981) <i>Proc Natl Acad Sci USA</i> 78, 3190-4. 5. Kim, K.M. et al. (1994) <i>FEBS Lett</i> 343, 251-5. 6. Kim, S.W. et al. (2001) <i>J Biol Chem</i> 276, 15712-9. 7. White, I.J. et al. (2006) <i>EMBO J</i> 25, 1-12. 8. Varticovski, L. et al. (1988) <i>Biochemistry</i> 27, 3682-90. 9. Dorovkov, M.V. and Ryazanov, A.G. (2004) <i>J Biol Chem</i> 279, 50643-6. 10. Wang, W. and Creutz, C.E. (1994) <i>Biochemistry</i> 33, 275-82. 11. Arur, S. et al. (2003) <i>Dev Cell</i> 4, 587-98. 12. Perretti, M. and Gavins, F.N. (2003) <i>News Physiol Sci</i> 18, 60-4. 13. Parente, L. and Solito, E. (2004) <i>Inflamm Res</i> 53, 125-32. 14. Lim, L.H. and Pervaiz, S. (2007) <i>FASEB J</i> 21, 968-75. 	
Species Reactivity	Species reactivity is determined by testing in at least one approved application (e.g., western blot).	
Applications Key	FC-FP: Flow Cytometry (Fixed/Permeabilized)	
Cross-Reactivity Key	H: Human	
Trademarks and Patents	Cell Signaling Technology is a trademark of Cell Signaling Technology, Inc. XP is a registered trademark of Cell Signaling Technology, Inc.	

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