Phospho-Akt1 (Ser473) (B9) rabbit mAb FITC conjugate

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Applications	Detection	Clonality	Isotype
Flow Cytometry	y N/A	Monoclonal	Rabbit IgGk
Format:	FITC		
Cross Reactivity:	Predicted to work with mouse, rat and other	er homologues.	
Formulation:	1X PBS, 0.09% NaN3, 0.2% BSA		
Preparation:	Protein A+G		
Reactivity:	Human,Mouse		
Recommended Usage:	For flow cytometric staining, the suggester μ L of staining volume. It is recommended application. See product image legends for	d use of this reagent is 5 μL per mill that the reagent be titrated for optim r additional information.	ion cells or 5 μL per 100 nal performance for each
Immunogen:	A synthetic phospho-peptide corresponding to residues surrounding Ser473 of human phospho Akt1		
Description:	Akt also known as PKB (Protein kinase B) or RAC-PK (Related to the A and C kinases) is a serine/threonine kinases that contains a pleckstrin homology (PH) domain. This protein kinase is activated by insulin and various growth and survival factors to function in a wortmannin sensitive pathway involving PI3 kinase. Akt is activated by phospholipid binding and activation loop phosphorylation at Thr308 by PDK1 and by phosphorylation within the carboxy terminus at Ser473. Phospho-Akt promotes cell survival by inhibiting apoptosis. Specifically, phospho-Akt1 has been shown to phosphorylate Bad, a member of the Bcl-2 family that promotes cell death. This phosphorylation results in the inactivation of the proapoptotic function of Bad. The Akt/phospho Akt molecule is thus considered to link extracellular survival signals (growth factors) with the apoptotic machinery (BAD). Akt is also a key mediator of the metabolic effects of insulin. Additionally, Akt has been referred to as an oncogene because it has increased activity in a number of tumors. This antibody recognizes phospho Akt. The homologous phosphorylation sites in Akt2 and Akt3 are S474 and S472, respectively.		
References:	 Franke TF, et al. (1997) Cell 88: 435-7. Burgering, B.M. and Coffer, P.J. (1995) Franke TF, et al. (1995) Cell 81: 727-36. Alessi DR, et al. (1996) EMBO J 15: 65- 	Nature 376: 599-602. 41-51.	

- 5. Sarbassov DD, et al. (2005) Science 307: 1098-101.
- 6. Jacinto E, et al. (2006) Cell 127: 125-37.
- 7 Cardona MH at al (1008) Science 282-1218 21



- 8. Brunet A, et al. (1999) Cell 96: 857-68.
- 9. Zimmermann S and Moelling K, (1999) Science 286: 1741-4.
- 10. Cantley LC and Neel BG (1999) Proc Natl Acad Sci USA 96: 4240-5.

Flow cytometric analysis of NIH3T3 cells treated with imatinib and unstained (blue) as

negative control or treated with imatinib (red) or pervadanate (PV) (green) and stained

using Phospho-Akt (Ser473) antibody AKTS473-B9 FITC conjugate. Cat. #2258.

- 11. Vlahos CJ, et al. (1994) J Biol Chem 269: 5241-8.
- 12. Hajduch E, et al. (2001) FEBS Lett 492: 199-203.
- 13. Cross DA, et al. (1995) Nature 378: 785-9.
- 14. Diehl JA, et al. (1998) Genes Dev 12: 3499-511.
- 15. Gesbert F, et al. (2000) J Biol Chem 275: 39223-30.
- 16. Zhou BP, et al. (2001) Nat Cell Biol 3: 245-52.
- 17. Navé BT, et al. (1999) Biochem J 344: 427-31.
- 18. Inoki K, et al. (2002) Nat Cell Biol 4: 648-57.
- 19. Manning BD, et al. (2002) Mol Cell 10: 151-62.



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