## FoxO1 (C29H4) Rabbit mAb (PE Conjugate)



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<b>Applications:</b> FC-FP	Reactivity: H M R Mk	<b>Sensitivity:</b> Endogenous	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q12778	Entrez-Gene Id: 2308
Product Usage Information		<b>Application</b> Flow Cytometry (Fixed/Po	ermeabilized)		<b>Dilution</b> 1:50
Storage		Supplied in PBS (pH 7.2), antibodies. Protect from		zide and 2 mg/ml BS/	A. Store at 4°C. Do not aliquot the
Specificity/Sensitivity		FoxO1 (C29H4) Rabbit mAb (PE Conjugate) detects endogenous levels of total FoxO1 protein. The antibody does not detect exogenously expressed family members FoxO3a or FoxO4.			
Source / Purification		Monoclonal antibody is produced by immunizing animals with a GST-fusion protein corresponding to carboxy-terminal residues of human FoxO1.			
Description		This Cell Signaling Technology antibody is conjugated to phycoerythrin (PE) and tested in-house for direct flow cytometry analysis in human cells. The antibody is expected to exhibit the same species cross-reactivity as the unconjugated FoxO1 (C29H4) Rabbit mAb #2880.			
Background		The Forkhead family of transcription factors is involved in tumorigenesis of rhabdomyosarcoma and acute leukemias (1-3). Within the family, three members (FoxO1, FoxO4, and FoxO3a) have sequence similarity to the nematode orthologue DAF-16, which mediates signaling via a pathway involving IGFR1, PI3K, and Akt (4-6). Active forkhead members act as tumor suppressors by promoting cell cycle arrest and apoptosis. Increased expression of any FoxO member results in the activation of the cell cycle inhibitor p27 Kip1. Forkhead transcription factors also play a part in TGF-β-mediated upregulation of p21 Cip1, a process negatively regulated through PI3K (7). Increased proliferation results when forkhead transcription factors are inactivated through phosphorylation by Akt at Thr24, Ser256, and Ser319, which results in nuclear export and inhibition of transcription factor activity (8). Forkhead transcription factors can also be inhibited by the deacetylase sirtuin (SirT1) (9).			
Background References		1. Anderson, M.J. et al. (1998) <i>Genomics</i> 47, 187-99. 2. Galili, N. et al. (1993) <i>Nat Genet</i> 5, 230-5. 3. Borkhardt, A. et al. (1997) <i>Oncogene</i> 14, 195-202. 4. Nakae, J. et al. (1999) <i>J Biol Chem</i> 274, 15982-5. 5. Rena, G. et al. (1999) <i>J Biol Chem</i> 274, 17179-83. 6. Guo, S. et al. (1999) <i>J Biol Chem</i> 274, 17184-92. 7. Seoane, J. et al. (2004) <i>Cell</i> 117, 211-23. 8. Arden, K.C. (2004) <i>Mol Cell</i> 14, 416-8. 9. Yang, Y. et al. (2005) <i>EMBO J</i> 24, 1021-32.			

**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 M: Mouse R: Rat Mk: Monkey

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