

Phospho-Tyrosine Mouse mAb (P-Tyr-100) (HRP Conjugate)



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Applications: W, E-P	Reactivity: All	Sensitivity: Endogenous	Source/Isotype: Mouse IgG1
Product Usage Information	Application Western Blotting Peptide ELISA (DELFI)		Dilution 1:2000 1:1000
Storage	Supplied in 140 mM NaCl, 3 mM KCl, 10 mM sodium phosphate (pH 7.4) dibasic, 2 mM potassium phosphate monobasic, 2 mg/mL BSA, and 50% glycerol. Store at -20°C. <i>Do not aliquot the antibody.</i>		
Specificity/Sensitivity	Phospho-Tyrosine Mouse mAb (P-Tyr-100) (HRP Conjugate) is a high affinity antibody. ELISAs against a wide variety of phosphopeptides indicate that P-Tyr-100 binds phospho-Tyr in a manner largely independent of the surrounding amino acid sequence. 2D gel western blot analysis of pervanadate-treated cell extracts also shows that P-Tyr-100 interacts with a broad range of tyrosine-phosphorylated proteins. P-Tyr-100 does not cross-react with peptides containing phospho-Ser or phospho-Thr. (U.S. Patent No's.: 6,441,140; 6,982,318; 7,259,022; 7,344,714; U.S.S.N. 11,484,485; and all foreign equivalents.)		
Source / Purification Description	Monoclonal antibody is produced by immunizing animals with phospho-tyrosine containing peptides. This Cell Signaling Technology® antibody is conjugated to the carbohydrate groups of horseradish peroxidase (HRP) via its amine groups. The HRP conjugated antibody is expected to exhibit the same species cross-reactivity as the unconjugated Phospho-Tyrosine Mouse mAb (P-Tyr-100) #9411.		
Background	Tyrosine phosphorylation plays a key role in cellular signaling (1). Research studies have shown that in cancer, unregulated tyrosine kinase activity can drive malignancy and tumor formation by generating inappropriate proliferation and survival signals (2). Antibodies specific for phospho-tyrosine (3,4) have been invaluable reagents in these studies. The phospho-tyrosine monoclonal antibodies developed by Cell Signaling Technology are exceptionally sensitive tools for studying tyrosine phosphorylation and monitoring tyrosine kinase activity in high throughput drug discovery.		
Background References	1. Schlessinger, J. (2000) <i>Cell</i> 103, 211-25. 2. Blume-Jensen, P. and Hunter, T. (2001) <i>Nature</i> 411, 355-65. 3. Ward, S.G. et al. (1992) <i>J Biol Chem</i> 267, 23862-9. 4. Glenney, J.R. et al. (1988) <i>J Immunol Methods</i> 109, 277-85.		

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 E-P: Peptide ELISA (DELFI)
Cross-Reactivity Key	All: All Species Expected
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