

**PTP-PEST (D4W7W) Rabbit mAb**

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

<b>Applications:</b> W, IP	<b>Reactivity:</b> H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 110 to 125	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q05209	<b>Entrez-Gene Id:</b> 5782
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**Product Usage Information****Application**

Western Blotting  
Immunoprecipitation

**Dilution**

1:1000  
1:50

**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**

PTP-PEST (D4W7W) Rabbit mAb recognizes endogenous levels of total PTP-PEST protein.

**Source / Purification**

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Arg747 of human PTP-PEST protein.

**Background**

PTP-PEST is a ubiquitously expressed cytosolic protein tyrosine phosphatase with multiple proline-rich regions that appear to be the docking sites for PTP-PEST binding partners or substrates (1). PTP-PEST regulates fibroblast adhesion, migration, and cytokinesis through its association with and dephosphorylation of p130 Cas, paxillin, PSTPIP1, WASP, and other adhesion molecules (1-5). By modulating phosphorylation states of Shc, Pyk2, Fak, and WASP, PTP-PEST negatively regulates lymphocyte activation (1,6). In mammary epithelial cells, EGF facilitates the dephosphorylation of Jak2 by PTP-PEST, thereby interfering with lactogenic hormone PRL signaling (7). PTP-PEST dephosphorylates c-Abl as well, which affects the phosphorylation states of PTP-PEST substrates such as paxillin, p130 Cas, Crk, and PSTPIP1 (8). PTP-PEST regulates adhesion and motility of cultured epithelial cells through modulation of Rho GTPase activity (9), and is required for integrin-mediated endothelial cell adhesion and migration (10).

**Background References**

1. Davidson, D. and Veillette, A. (2001) *EMBO J* 20, 3414-26.
2. Garton, A.J. and Tonks, N.K. (1999) *J Biol Chem* 274, 3811-8.
3. Shen, Y. et al. (2000) *J Biol Chem* 275, 1405-13.
4. Angers-Loustau, A. et al. (1999) *J Cell Biol* 144, 1019-31.
5. Côté, J.F. et al. (2002) *J Biol Chem* 277, 2973-86.
6. Badour, K. et al. (2004) *J Exp Med* 199, 99-112.
7. Horsch, K. et al. (2001) *Mol Endocrinol* 15, 2182-96.
8. Cong, F. et al. (2000) *Mol Cell* 6, 1413-23.
9. Espejo, R. et al. (2010) *Am J Physiol Cell Physiol* 299, C454-63.
10. Souza, C.M. et al. (2012) *J Biol Chem* 287, 43180-90.

**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 nonfat dry milk, 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 **M:** Mouse **R:** Rat

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