## Phospho-Ezrin (Thr567)/Radixin (Thr564)/Moesin (Thr558) (48G2) Rabbit mAb



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Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
WB, IHC-P, IF-IC	HMRMk	Endogenous	75 Moesin. 80	Rabbit IgG	#P15311, #P35241,	7430, 5962, 4478
		ŭ	Ezrin, Radixin.	· ·	#P26038	

**Product Usage** Application Dilution Information 1:1000 Western Blotting Immunohistochemistry (Paraffin) 1:200 - 1:800 Immunofluorescence (Immunocytochemistry) 1:100 - 1:400 Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than Storage 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody. For a carrier free (BSA and azide free) version of this product see product #32600. Phospho-Ezrin (Thr567)/Radixin (Thr564)/Moesin (Thr558) (48G2) Rabbit mAb is phospho-specific by Specificity / Sensitivity peptide-ELISA and exhibits a large signal to noise window. The antibody does not recognize the nonphosphorylated peptide in peptide based ELISA. Species predicted to **Bovine** react based on 100% sequence homology: Source / Purification Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Thr567 of human ezrin protein. **Background** The ezrin, radixin, and moesin (ERM) proteins function as linkers between the plasma membrane and the actin cytoskeleton and are involved in cell adhesion, membrane ruffling, and microvilli formation (1). ERM proteins undergo intra or intermolecular interaction between their amino- and carboxy-terminal domains, existing as inactive cytosolic monomers or dimers (2). Phosphorylation at a carboxy-terminal threonine residue (Thr567 of ezrin, Thr564 of radixin, Thr558 of moesin) disrupts the amino- and carboxy-terminal association and may play a key role in regulating ERM protein conformation and function (3,4). Phosphorylation at Thr567 of ezrin is required for cytoskeletal rearrangements and oncogene-induced transformation (5). Ezrin is also phosphorylated at tyrosine residues upon growth factor stimulation. Phosphorylation of Tyr353 of ezrin transmits a survival signal during epithelial differentiation (6).

**Background References** 

- 1. Tsukita, S. and Yonemura, S. (1999) J Biol Chem 274, 34507-10.
- 2. Mangeat, P. et al. (1999) Trends Cell Biol 9, 187-92.
- 3. Matsui, T. et al. (1998) J Cell Biol 140, 647-57.
- 4. Gautreau, A. et al. (2000) J Cell Biol 150, 193-203. 5. Tran Quang, C. et al. (2000) EMBO J 19, 4565-76.
- 6. Gautreau, A. et al. (1999) Proc Natl Acad Sci U S A 96, 7300-5.

**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** 

WB: Western Blotting IHC-P: Immunohistochemistry (Paraffin)

IF-IC: Immunofluorescence (Immunocytochemistry)

**Cross-Reactivity Key** 

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse GP: Guinea Pig Rab: rabbit All: all species expected

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