

**EPLIN (D1A7A) Rabbit mAb**

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

| Applications: | Reactivity: | Sensitivity: | MW (kDa): | Source/Isotype: | UniProt ID: | Entrez-Gene Id: |
|---------------|-------------|--------------|-----------|-----------------|-------------|-----------------|
| W, IF-IC      | H M R       | Endogenous   | 85, 110   | Rabbit IgG      | #Q9UHB6     | 51474           |

**Product Usage Information****Application**

Western Blotting  
Immunofluorescence (Immunocytochemistry)

**Dilution**

1:1000  
1:200

**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**

EPLIN (D1A7A) Rabbit mAb recognizes endogenous levels of total EPLIN protein, including EPLIN-α and EPLIN-β isoforms. Based on the absence of signal in NCI-H28 cells, which appear to express low levels of β-Eplin, this antibody may have a preference for α-Eplin by immunofluorescence.

**Source / Purification**

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gly455 of human EPLIN protein.

**Background**

Epithelial Protein Lost in Neoplasm (EPLIN) is an actin-binding protein that regulates actin filament dynamics and cross-linking (1). Alpha and beta isoforms are generated from alternate promoters, with the EPLIN-β isoform representing the full-length protein and the EPLIN-α isoform lacking the amino-terminal 160 amino acids (2). Increased expression of EPLIN protein results in more abundant and larger actin stress fibers due to stabilizing of cross-links and inhibition of actin depolymerization. EPLIN protein inhibits Rac1-promoted membrane ruffling and Arp2/3-associated actin filament branching (1).

Research studies demonstrate reduced EPLIN-α expression in tumor tissues, and correlate this reduction with increased invasiveness and poor clinical outcomes (3). The EPLIN protein is an important negative regulator of the epithelial-mesenchymal transition (EMT)(4). While EMT is a critical process during normal embryonic development, dysregulation in transformed cells is a key step in the transition to metastasis (5).

**Background References**

1. Maul, R.S. et al. (2003) *J Cell Biol* 160, 399-407.
2. Chen, S. et al. (2000) *Gene* 248, 69-76.
3. Liu, Y. et al. (2012) *Anticancer Res* 32, 1283-9.
4. Zhang, S. et al. (2011) *Oncogene* 30, 4941-52.
5. Tsai, J.H. and Yang, J. (2013) *Genes Dev* 27, 2192-206.

**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 **IF-IC:** Immunofluorescence (Immunocytochemistry)

**Cross-Reactivity Key**

**H:** Human **M:** Mouse **R:** Rat

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