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## Product Information

### Anti-Profilin 1 (N-terminal)

produced in rabbit, affinity isolated antibody

Catalog Number **P7749**

#### Product Description

Anti-Profilin 1 (N-terminal) is developed in rabbit using as immunogen a synthetic peptide corresponding to amino acid residues 2-17 of human profilin 1, conjugated to KLH. The corresponding rat and mouse sequence differs by one amino acid. This sequence is 70% similar to the corresponding sequence in Profilin 2. The antibody is affinity-purified using the immunizing peptide immobilized on agarose.

Anti-Profilin 1 (N-terminal) recognizes human, mouse, and rat profilin 1. Applications include immunoblotting (~15 kDa) and immunofluorescence. Detection of the profilin 1 band by immunoblotting is specifically inhibited by the immunizing peptide.

Profilin 1 is a ubiquitous actin monomer-binding protein involved in actin polymerization in response to extracellular signals.<sup>1</sup> Three profilin genes have been identified: profilin 1, 2, and 3. Profilin 1 is the most ubiquitous and abundant, and is highly expressed throughout development and adulthood in most tissues including brain. Profilin 2 is the neuronal specific isoform and profilin 3 is a testis specific isoform.<sup>2,3</sup> Profilins of eukaryotic cells are small (12-15 kDa) cytoplasmic proteins that bind to actin monomers, polyphosphoinositides and polymers of L-proline.<sup>1</sup> Profilins were shown to be important for normal cell proliferation, differentiation and motility.<sup>4</sup> Deletion of profilin 1 gene leads to an embryonic lethal phenotype.<sup>5</sup>

Profilin 1 is a potent regulator of actin filament dynamics. Although profilin 1 prevents spontaneous actin polymerization by complexing with unpolymerized actin *in vivo*, actin-profilin complexes can be added to free barbed ends, thereby stimulating actin polymerization. When bound to actin, profilin functions as an ATP nucleotide exchange factor recharging ADP-actin with ATP.<sup>6,7</sup> Dissociation of the profilin-actin complex is caused by binding of profilin to phosphatidylinositol 4,5-bisphosphate (PIP<sub>2</sub>), which liberates actin for polymerization.<sup>8</sup> Profilin binds with high affinity to poly-L-proline stretches found in many cellular proteins associated with the cytoskeleton.

Profilin is recruited to sites of active cytoskeletal assembly through its interaction with proteins such as N-WASP, the ARP2/3 complex, p140mDia, VASP and Mena.<sup>7</sup> The interactions of profilin 1 with actin, proline-rich proteins and PIP<sub>2</sub> influences neuronal differentiation of PC12 cells.<sup>9</sup> Profilin 1 was suggested to act as a tumor suppressor protein based on its reduced expression in several types of invasive cancers and its ability to suppress tumorigenicity when over-expressed in breast cancer cells.<sup>4,10</sup> Deletion of profilin 1 is associated with Miller-Dieker syndrome.<sup>11</sup>

#### Reagent

Supplied as a solution in 0.01 M phosphate buffered saline, pH 7.4, containing 15 mM sodium azide as a preservative.

Antibody concentration: ~1 mg/mL

#### Precautions and Disclaimer

This product is for R&D use only, not for drug, household, or other uses. Please consult the Material Safety Data Sheet for information regarding hazards and safe handling practices.

#### Storage/Stability

For continuous use, store at 2-8 °C for up to one month. For extended storage, freeze in working aliquots. Repeated freezing and thawing, or storage in "frost-free" freezers, is not recommended. If slight turbidity occurs upon prolonged storage, clarify the solution by centrifugation before use. Working dilutions should be discarded if not used within 12 hours.

#### Product Profile

Immunoblotting: a working concentration of 1-2 µg/mL is recommended using whole extracts of mouse NIH-3T3 and human HeLa cells, applying a chemiluminescent detection reagent.

Indirect immunofluorescence: a working concentration of 10-20 µg/mL is recommended using rat NRK cells.

**Note:** In order to obtain the best results using various techniques and preparations, we recommend determining the optimal working dilutions by titration.

## References

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