

#13347 Store at -20°C

**FABP7 (D8N3N) Rabbit mAb****Cell Signaling**  
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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: |
|---------------|-------------|--------------|-----------|-----------------|-------------|-----------------|
| WB, IF-IC     | H R         | Endogenous   | 15        | Rabbit IgG      | #O15540     | 2173            |

**Product Usage Information****Application**Western Blotting  
Immunofluorescence (Immunocytochemistry)**Dilution**1:1000  
1:2000**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**

FABP7 (D8N3N) Rabbit mAb recognizes endogenous levels of total FABP7 protein. Species cross-reactivity for IF-IC is human only.

**Source / Purification**

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val84 of human FABP7 protein.

**Background**

Fatty acid binding proteins (FABPs) bind to fatty acids and other lipids to function as cytoplasmic lipid chaperones (1). They participate in the transport of fatty acids and other lipids to various cellular pathways (2). Differential expression of FABPs is found in several types of tumors and their normal-cell counterparts (3). FABP7 is abundantly expressed in fetal brain and may be essential for development (4). Expression is required for the establishment of the radial glial fiber system, a system that is necessary for the development of cortical layers (5). Increased expression of FABP7 is associated with reduced survival in patients with glioblastoma (6), and is also found in glial cells following nerve injury (7). Investigators have found loss of FABP7 may be involved in the development and progression of breast cancer and expression of FABP7 has been shown to induce mammary differentiation and to inhibit growth of breast cancer cells (8,9).

**Background References**

1. Storch, J. and Thumser, A.E. (2010) *J Biol Chem* 285, 32679-83.
2. Haunerland, N.H. and Spener, F. (2004) *Prog Lipid Res* 43, 328-49.
3. Khan, S.H. and Sorof, S. (1994) *Proc Natl Acad Sci U S A* 91, 848-52.
4. Shimizu, F. et al. (1997) *Biochim Biophys Acta* 1354, 24-8.
5. Feng, L. and Heintz, N. (1995) *Development* 121, 1719-30.
6. Liang, Y. et al. (2005) *Proc Natl Acad Sci U S A* 102, 5814-9.
7. Miller, S.J. et al. (2003) *Mol Cell Biol* 23, 2213-24.
8. Shi, Y.E. et al. (1997) *Cancer Res* 57, 3084-91.
9. Wang, M. et al. (2000) *Cancer Res* 60, 6482-7.

**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****WB:** Western Blotting **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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