

BACKGROUND

Fatty acid-binding proteins (FABPs) are members of the superfamily of lipid-binding proteins (LBP). So far 9 different FABPs, with tissue-specific distribution, have been identified: L (liver), I (intestinal), H (muscle and heart), A (adipocyte), E (epidermal), Il (ileal), B (brain), M (myelin) and T (testis). The primary role of all the FABP family members is regulation of fatty acid uptake and intracellular transport. The structure of all FABPs is similar – the basic motif characterizing these proteins is beta-barrel, and a single ligand (e.g. a fatty acid, cholesterol, or retinoid) is bound in its internal water-filled cavity. Despite the wide variance in the protein sequence, the gene structure is identical. The FABP genes consist of 4 exons and 3 introns and a few of them are located in the same chromosomal region. For example, A-FABP, E-FABP and M-FABP create a gene cluster.¹ Physiological roles of these proteins are not only involved in FA transport, but also in regulation of cell growth and differentiation, cellular signalling, gene transcription and cytoprotection.² Because of their physiological properties some FABP genes were tested in order to identify mutations altering lipid metabolism and relating with other diseases.

Heart-type FABP3 (H-FABP), which is most abundantly expressed in heart, skeletal muscle, brain, and other tissues, is induced in brown adipose tissue (BAT) in hibernating squirrel and bat. It is also induced by acute cold exposure in rat. The physiological roles of FABP3 have been investigated with knockout mouse models. Characterization of FABP3-deficient mice has focused largely on effects on fatty acid metabolism in heart and skeletal muscle, issues in which this protein is abundantly expressed. *In vivo* labeling studies revealed reduced palmitic and arachidonic acid uptake in heart of *Fabp3*^{-/-} mice (17), and reduced palmitic acid uptake and utilization in isolated cardiomyocytes. There was a compensatory reliance on glucose oxidation in the heart of *Fabp3*^{-/-} mice. *Fabp3*^{-/-} and *Fabp3*^{+/-} mice also exhibit increased insulin sensitivity, perhaps related to the increased reliance on glucose rather than fatty acid fuels. Thus, FABP3 has important roles in fatty acid metabolism in heart and skeletal muscle, with effects on systemic glucose homeostasis. The induction of FABP3 in BAT during cold exposure described above raised the possibility that this protein may have a role in fatty acid trafficking in adaptive thermogenesis. Moreover, it was also shown that *Fabp3*^{+/-} mice exhibit dopamine D₂ receptor (D2R) dysfunction in the striatum. FABP3/H-FABP regulates functions of the dopamine D2R in the brain, through neuronal D2LR/FABP3 interaction.³ In addition, it was reported that FABP3/H-FABP inhibits growth of mammary epithelial cells. FABP3/H-FABP is a candidate tumor suppressor for human breast cancer. Furthermore, FABP3/H-FABP has been used as a serological biomarker of cardiac and skeletal muscle injury.⁴

References:

1. Merkel, M. Et al: J. Biol. Chem. 277:7405-11, 2001
2. Hamilton, M.T. et al: Am J Physiol Endocrinol Metab 275: E1016-E1022, 1998
3. Shioda, N. et al: J. Neurosci. 30:3146-55, 2010
4. Pritt, M.L. et al: Toxicol. Sci. 103:382-96, 2008

TECHNICAL INFORMATION

Source:

FABP3 Antibody is a rabbit antibody raised against a short peptide from human FABP3 sequence.

Specificity and Sensitivity:

This antibody detects endogenous levels of FABP3 proteins without cross-reactivity with other related proteins.

Storage Buffer: PBS and 30% glycerol

Storage:

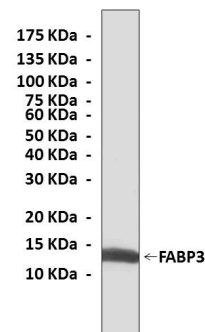
Store at -20°C for at least one year. Store at 4°C for frequent use. Avoid repeated freeze-thaw cycles.

APPLICATIONS

Application:	*Dilution:
WB	1:1000
IP	n/d
IHC	1:50-200
ICC	n/d
FACS	n/d

**Optimal dilutions must be determined by end user.*

QUALITY CONTROL DATA



Western Blot detection of FABP3 proteins in rat heart tissue lysate using FABP3 Antibody.

