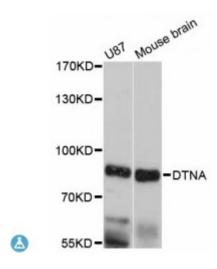


Anti-DTNA Antibody



Description

The protein encoded by this gene belongs to the dystrobrevin subfamily of the dystrophin family. This protein is a component of the dystrophin-associated protein complex (DPC), which consists of dystrophin and several integral and peripheral membrane proteins, including dystroglycans, sarcoglycans, syntrophins and alpha- and beta-dystrobrevin. The DPC localizes to the sarcolemma and its disruption is associated with various forms of muscular dystrophy. Mutations in this gene are associated with left ventricular noncompaction with congenital heart defects. Multiple alternatively spliced transcript variants encoding different isoforms have been identified for this gene.

Model STJ113966

Host Rabbit

Reactivity Human, Mouse

Applications WB

Immunogen Recombinant fusion protein containing a sequence corresponding to amino

acids 240-480 of human DTNA (NP_001381.2).

Gene ID 1837

Gene Symbol DTNA

Dilution range WB 1:500 - 1:2000

Tissue Specificity Highly expressed in brain, skeletal and cardiac muscles, and expressed at

lower levels in lung, liver and pancreas, Isoform 2 is not expressed in cardiac

muscle, Isoform 7 and isoform 8 are only expressed in muscle

Purification Affinity purification

Note For Research Use Only (RUO).

Protein Name Dystrobrevin alpha DTN-A Alpha-dystrobrevin Dystrophin-related protein 3

Molecular Weight 83.901 kDa

Clonality Polyclonal

Conjugation Unconjugated

Isotype IgG

Formulation PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

Storage Instruction Store at -20C. Avoid freeze / thaw cycles.

Database Links HGNC:3057OMIM:601239

Alternative Names Dystrobrevin alpha DTN-A Alpha-dystrobrevin Dystrophin-related protein 3

Function May be involved in the formation and stability of synapses as well as being

involved in the clustering of nicotinic acetylcholine receptors

Cellular Localization Cytoplasm, Cell junction, synapse, Cell membrane

Post-translational Phosphorylation of DTN-1 on tyrosine kinase substrate domain present in the

Modifications C-terminus,

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