**BACKGROUND**

Dystrobrevins are protein components of the dystrophin complex, whose disruption leads to Duchenne muscular dystrophy and related diseases. α-Dystrobrevin is a dystrophin-related and -associated protein that is involved in synapse maturation and is required for normal muscle function. α-Dystrobrevin is a component of the dystrophin glycoprotein complex. It is localized to the cytoplasmic side of the sarcolemma and is highly concentrated at the neuromuscular junctions in skeletal muscle. The insertion of 57 amino acids by alternative splicing accounts for the increase in molecular mass of α-Dystrobrevin 1 in skeletal and cardiac muscle compared with brain and lung. α-Dystrobrevin containing complexes are found in endothelial and smooth muscle cells, while β-Dystrobrevin containing complexes are present at the basal region of renal epithelial cells. Additionally, β-Dystrobrevin is found in neurons and is highly enriched in postsynaptic densities. Alternative splicing of α-Dystrobrevin produces γ-Dystrobrevin (isoform 5), δ-Dystrobrevin (isoform 7), ε-Dystrobrevin (isoform 6) and ζ-Dystrobrevin (isoform 8). Additional isoforms may also exist.

**REFERENCES**


**CHROMOSOMAL LOCATION**

Genetic locus: DTNA (human) mapping to 18q12.1; Dtna (mouse) mapping to 18 A2.

**SOURCE**

α-Dystrobrevin (E-12) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 351-371 within an internal region of α-Dystrobrevin of human origin.

**STORAGE**

Store at 4°C. **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

**PRODUCT**

Each vial contains 200 µg IgG2b kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin. Blocking peptide available for competition studies, sc-365289 P, (100 µg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

**APPLICATIONS**

α-Dystrobrevin (E-12) is recommended for detection of all isoforms of α-Dystrobrevin of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:300).

α-Dystrobrevin (E-12) is also recommended for detection of all isoforms of α-Dystrobrevin in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for α-Dystrobrevin siRNA (h): sc-43321, α-Dystrobrevin siRNA (m): sc-43322, α-Dystrobrevin shRNA Plasmid (h): sc-43321-SH, α-Dystrobrevin shRNA Plasmid (m): sc-43322-SH, α-Dystrobrevin shRNA (h) Lentiviral Particles: sc-43321-V and α-Dystrobrevin shRNA (m) Lentiviral Particles: sc-43322-V.

Molecular Weight of α-Dystrobrevin non-muscle α type: 78 kDa.

Molecular Weight of α-Dystrobrevin muscle α type: 94 kDa.

Positive Controls: α-Dystrobrevin (h): 293T Lysate: sc-177157, mouse skeletal muscle extract: sc-364250 or SK-N-SH cell lysate: sc-2410.

**RECOMMENDED SUPPORT REAGENTS**

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgGκ BP-HRP: sc-516102 or m-IgGκ BP-FITC: sc-516140 or m-IgGκ BP-PE: sc-2333 and Western Blotting Luminol Reagent: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

**DATA**

![α-Dystrobrevin (E-12): sc-365289. Western blot analysis of α-Dystrobrevin expression in non-transfected: sc-117752 (A) and human α-Dystrobrevin transfected: sc-117757 (B) 293T whole cell lysates.](image1)

![α-Dystrobrevin (E-12): sc-365289. Western blot analysis of α-Dystrobrevin expression in mouse skeletal muscle tissue extract.](image2)

**RESEARCH USE**

For research use only, not for use in diagnostic procedures.