

δ-sarcoglycan (H-55): sc-28281

BACKGROUND

The sarcoglycan transmembrane proteins are members of the dystrophin complex. Sarcoglycans cluster together to form a complex, which is localized in the cell membrane of skeletal, cardiac and smooth muscle fibers. Four sarcoglycan subunit proteins, designated α -, β -, γ - and δ -sarcoglycan, form a complex on the skeletal muscle cell surface membrane. A genetic defect in any one of these proteins causes the loss or marked decrease of the whole sarcoglycan complex, which is observed in the autosomal recessive muscular dystrophy, sarcoglycanopathy. In smooth muscle, β - and δ -sarcoglycans are associated with ϵ -sarcoglycan, a glycoprotein homologous to α -sarcoglycan. Additionally, a complete deficiency in δ -sarcoglycan is the cause of the Syrian hamster BIO.14 cardiomyopathy.

REFERENCES

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- Enigk, R.E., et al. 2001. Cellular and molecular properties of α -dystrobrevin in skeletal muscle. *Front. Biosci.* 6: D53-D64.
- Politano, L., et al. 2001. Evaluation of cardiac and respiratory involvement in sarcoglycanopathies. *Neuromuscul. Disord.* 11: 178-185.
- Ueda, H., et al. 2001. δ - and γ -sarcoglycan localization in the sarcoplasmic reticulum of skeletal muscle. *J. Histochem. Cytochem.* 49: 529-538.
- Wakabayashi-Takai, E., et al. 2001. Identification of myogenesis-dependent transcriptional enhancers in promoter region of mouse γ -sarcoglycan gene. *Eur. J. Biochem.* 268: 948-957.
- Anastasi, G., et al. 2004. Sarcoglycan and integrin localization in normal human skeletal muscle: a confocal laser scanning microscope study. *Eur. J. Histochem.* 48: 245-252.

CHROMOSOMAL LOCATION

Genetic locus: SGCD (human) mapping to 5q33.3; Sgcd (mouse) mapping to 11 B1.1.

SOURCE

δ -sarcoglycan (H-55) is a rabbit polyclonal antibody raised against amino acids 206-260 mapping within an extracellular domain of δ -sarcoglycan of human origin.

PRODUCT

Each vial contains 200 μ g IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

STORAGE

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

APPLICATIONS

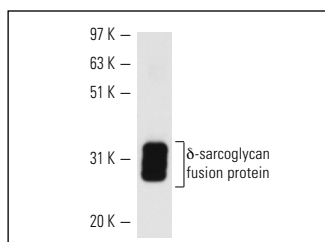
δ -sarcoglycan (H-55) is recommended for detection of δ -sarcoglycan of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

δ -sarcoglycan (H-55) is also recommended for detection of δ -sarcoglycan in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for δ -sarcoglycan siRNA (h): sc-43420, δ -sarcoglycan siRNA (m): sc-43421, δ -sarcoglycan shRNA Plasmid (h): sc-43420-SH, δ -sarcoglycan shRNA Plasmid (m): sc-43421-SH, δ -sarcoglycan shRNA (h) Lentiviral Particles: sc-43420-V and δ -sarcoglycan shRNA (m) Lentiviral Particles: sc-43421-V.

Molecular Weight of δ -sarcoglycan: 35 kDa.

DATA



δ -sarcoglycan (H-55): sc-28281. Western blot analysis of human recombinant δ -sarcoglycan fusion protein.

SELECT PRODUCT CITATIONS

- Li, D., et al. 2009. Sub-physiological sarcoglycan expression contributes to compensatory muscle protection in mdx mice. *Hum. Mol. Genet.* 18: 1209-1220.
- Daicho, T., et al. 2009. Alterations in dystrophin-related glycoproteins in development of right ventricular failure in rats. *J. Pharmacol. Sci.* 111: 405-415.
- Kakarla, S.K., et al. 2010. Possible molecular mechanisms underlying age-related cardiomyocyte apoptosis in the F344XBN rat heart. *J. Gerontol. A Biol. Sci. Med. Sci.* 65: 147-155.

RESEARCH USE

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



Try **δ -sarcoglycan (3G10): sc-517013**, our highly recommended monoclonal alternative to δ -sarcoglycan (H-55).