

MYOT polyclonal antibody

Catalog: BS8217

Host: Rabbit

Reactivity: Human, Mouse, Rat

BackGround:

Myotilin, a sarcomeric protein that is encoded by the gene mapping to human chromosome 5q31, binds to α -actinin and is localized in the Z-line of myofibrils. Myotilin is expressed in skeletal and cardiac muscle, and it co-localizes with α -actinin in the sarcomeric I-bands where it directly interacts with α -actinin. Defects in the myotilin gene are reported to cause a form of autosomal dominant limb-girdle muscular dystrophy (LGMD) (1). Symptoms of adult onset LGMD are progressive weakness of the hip and shoulder girdles as well as a distinctive dysarthric pattern of speech. The muscle of affected individuals with LGMD shows degeneration of myofibers, variations in fiber size, fiber splitting, centrally located myonuclei and an enhanced number of autophagic vesicles.

Product:

Rabbit IgG, 1mg/ml in PBS with 0.02% sodium azide, 50% glycerol, pH7.2

Molecular Weight:

~ 55 kDa

Swiss-Prot:

Q9UBF9

Purification&Purity:

The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific im-

munogen and the purity is > 95% (by SDS-PAGE).

Applications:

WB 1:500 - 1:2000

IHC 1:50 - 1:200

IF 1:10 - 1:100

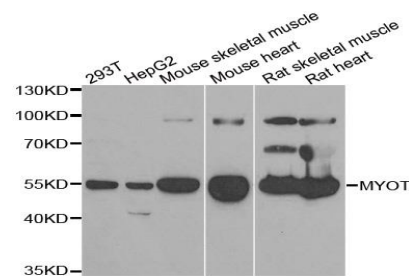
Storage&Stability:

Store at 4 °C short term. Aliquot and store at -20 °C long term. Avoid freeze-thaw cycles.

Specificity:

MYOT polyclonal antibody detects endogenous levels of MYOT protein.

DATA:



WesternBlot (WB) analysis of MYOT polyclonal antibody

Note:

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

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