

# **Dystrophin polyclonal antibody**

Catalog: **BS90437**  Host:

Rabbit

Reactivity: Human, Mouse, Rat

## **BackGround:**

Dystrophin-glycoprotein complex (DGC) connects the F-Actin cytoskeleton on the inner surface of muscle fibers to the surrounding extracellular matrix, through the cell membrane interface. A deficiency in this protein contributes to Duchenne (DMD) and Becker (BMD) muscular dystrophies. The human dystrophin gene measures 2.4 megabases, has more than 80 exons, produces a 14 kb mRNA and contains at least eight independent tissue-specific promoters and two poly A sites. The dystrophin mRNA can undergo differential splicing and produce a range of transcripts that encode a large set of proteins. Dystrophin represents approximately 0.002% of total striated muscle protein and localizes to triadic junctions in skeletal muscle, where it is thought to influence calcium ion homeostasis and force transmission.

## **Product:**

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

**Molecular Weight:** 

427 kDa

**Swiss-Prot:** 

P11532(Human) P11531(Mouse) P11530(Rat)

**Purification&Purity:** 

ProA affinity purified

**Applications:** 

WB:1:1,000

IHC:1:50-1:200

Storage&Stability:

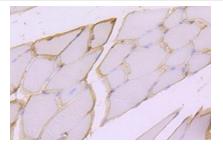
Store at +4 °C after thawing. Aliquot store at -20 °C or

-80 °C. Avoid repeated freeze / thaw cycles.

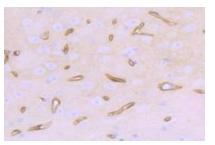
## **Specificity:**

Dystrophin polyclonal antibody detects endogenous levels of Dystrophin protein.

### **DATA:**



Immunohistochemical analysis of paraffin-embedded mouse skeletal muscle tissue using anti-Dystrophin antibody. Counter stained with hematoxylin.



Immunohistochemical analysis of paraffin-embedded mouse brain tissue using anti-Dystrophin antibody. Counter stained with hematoxylin.

#### Note:

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

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