

Dystrophin Antibody / DMD [clone DMD/8773R] (V5340)

Catalog No.	Formulation	Size
V5340-100UG	0.2 mg/ml in 1X PBS with 0.1 mg/ml BSA (US sourced), 0.05% sodium azide	100 ug
V5340-20UG	0.2 mg/ml in 1X PBS with 0.1 mg/ml BSA (US sourced), 0.05% sodium azide	20 ug
V5340SAF-100UG	1 mg/ml in 1X PBS; BSA free, sodium azide free	100 ug

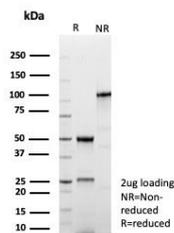
Recombinant **RABBIT MONOCLONAL**

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Availability	1-3 business days
Species Reactivity	Human
Format	Purified
Host	Rabbit
Clonality	Recombinant Rabbit Monoclonal
Isotype	Rabbit IgG, kappa
Clone Name	DMD/8773R
Purity	Protein A/G affinity
UniProt	P11532
Localization	Cell surface, Cytoplasm
Applications	Immunohistochemistry (FFPE) : 1-2ug/ml for 30 min at RT
Limitations	This Dystrophin antibody is available for research use only.



IHC staining of FFPE human skeletal muscle tissue with Dystrophin antibody (clone DMD/8773R). HIER: boil tissue sections in pH 9 10mM Tris with 1mM EDTA for 20 min and allow to cool before testing.



SDS-PAGE analysis of purified, BSA-free Dystrophin antibody (clone DMD/8773R) as confirmation of integrity and purity.

Description

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 8 independent tissue-specific promoters and 2 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.

This DMD antibody can be compared with our [Dystrophin Antibody](#) (clone DMD/3241) for analysis of dystrophin expression across muscle structure and neuromuscular disease studies.

Application Notes

Optimal dilution of the Dystrophin antibody should be determined by the researcher.

Immunogen

A recombinant partial protein sequence (within amino acids 1700-2300) from the human protein was used as the immunogen for the Dystrophin antibody.

Storage

Aliquot the Dystrophin antibody and store frozen at -20oC or colder. Avoid repeated freeze-thaw cycles.