

Product datasheet for TA353791

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Dystrophin (DMD) Mouse Monoclonal Antibody [Clone ID: SPM 499]

Product data:

Product Type: Primary Antibodies

Clone Name: SPM 499

Applications: IHC

Recommended Dilution: IHC-P

Reactivity: Human

Host: Mouse

Isotype: IgG1, kappa
Clonality: Monoclonal

Immunogen: Synthetic peptide corresponding to C-terminus of human dystrophin.

Formulation: PBS/1% BSA buffer pH 7.6 with less than 0.1% sodium azide.

Purification: protein A/G

Conjugation: Unconjugated

Storage: Store at -20°C as received.

Stability: Stable for 12 months from date of receipt.

Predicted Protein Size: 427 kDa

Gene Name: dystrophin

Database Link: NP 004006

Entrez Gene 1756 Human

P11532

Background: Dystrophin is a member of the spectrin/alpha-actinin family of actin-binding, triple helix

rodcontaining proteins. It is absent or greatly reduced in individuals with the X-linked recessive Duchennes muscular dystrophy disorder, as well as in mice with the mdx (murine muscular dystrophy) mutation. This antibody is highly specific to dystrophin and shows no cross-reaction with C-protein (an isoform of β -actinin), β -actin, or human muscle spectrin.

Synonyms: BMD; CMD3B; DXS142; DXS164; DXS206; DXS230; DXS239; DXS268; DXS269; DXS270; DXS272;

MRX85



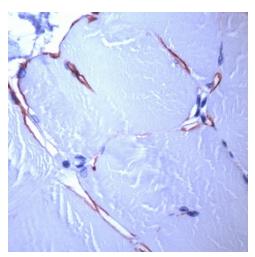


Protein Pathways:

Arrhythmogenic right ventricular cardiomyopathy (ARVC), Dilated cardiomyopathy,

Hypertrophic cardiomyopathy (HCM), Viral myocarditis

Product images:



Human Skeletal Muscle stained with anti-Dystrophin antibody.