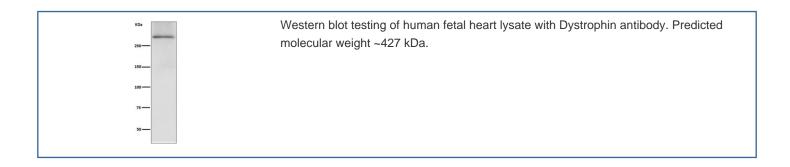


Dystrophin Antibody / DMD [clone AOGG-4] (RQ5369)

Catalog No.	Formulation	Size
RQ5369	Antibody in PBS with 0.02% sodium azide, 50% glycerol and 0.4-0.5mg/ml BSA	100 ul

Bulk quote request

Availability	1-2 weeks
Species Reactivity	Human
Format	Purified
Clonality	Rabbit Monoclonal
Isotype	Rabbit IgG
Clone Name	AOGG-4
Purity	Affinity purified
UniProt	P11532
Applications	Western Blot : 1:500-1:2000
Limitations	This Dystrophin antibody is available for research use only.



Description

The DMD gene spans a genomic range of greater than 2 Mb and encodes a large protein containing an N-terminal actin-binding domain and multiple spectrin repeats. The encoded protein forms a component of the dystrophin-glycoprotein complex (DGC), which bridges the inner cytoskeleton and the extracellular matrix. Deletions, duplications, and point mutations at this gene locus may cause Duchenne muscular dystrophy (DMD), Becker muscular dystrophy (BMD), or cardiomyopathy. Alternative promoter usage and alternative splicing result in numerous distinct transcript variants and protein isoforms for this gene. [RefSeq]

Application Notes

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

Immunogen

A synthetic peptide specific to human Dystrophin / DMD was used as the immunogen for the Dystrophin antibody.

Storage

Store the Dystrophin antibody at -20oC.