

## **Product datasheet for TA355287**

## OriGene Technologies, Inc.

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## **DNAH2 Mouse Monoclonal Antibody [Clone ID: 34C5]**

## **Product data:**

**Product Type:** Primary Antibodies

Clone Name: 34C5
Applications: IHC

Recommended Dilution: 1:40 - 1:80

Reactivity: Human
Host: Mouse
Isotype: IgG1

Clonality: Monoclonal

**Immunogen:** Prokaryotic recombinant protein corresponding to amino acids 321 to 494 of the dystrophin

molecule

**Specificity:** Human dystrophin molecule

**Formulation:** Lyophilized tissue culture supernatant containing 15 mM sodium azide as a preservative.

**Reconstitution Method:** The user is required to reconstitute the contents of the vial with the correct volume of sterile

distilled water as indicated on the vial label

Conjugation:UnconjugatedStorage:Store at 2-8°CStability:12 months

**Gene Name:** dynein, axonemal, heavy polypeptide 2

**Background:** Duchenne Muscular dystrophy (DMD) is the most common of the muscular dystrophies

resulting in progressive muscular wasting and death. Dystrophin is the 427kD protein product of the DMD gene located on the X chromosome at position Xp21. Abnormalities in protein expression occur in patients with DMD/BMD and dystrophin analysis may be used to distinguish these conditions from other neuromuscular diseases. Severe Duchenne muscular dystrophy is associated with a marked dystrophin deficiency, whereas patients with the milder form of Becker muscular dystrophy show less pronounced abnormalities of protein expression. The immunolabeling patterns for DYS1, DYS2 and DYS3 are similar; however, the use of all three antibodies is recommended to avoid the possibility of occasional false

negative results.

