



# Mouse Anti-Bovine Myosin Hybridoma [CG-56] (CSC-H1441)

This product is for research use only and is not intended for diagnostic use.

## PRODUCT INFORMATION

<b>Product Overview</b>	This hybridoma produces mAbs (IgG1) against bovine Myosin
<b>Target</b>	Myosin
<b>Immunogen</b>	Fetal bovine muscles
<b>Isotype</b>	IgG1
<b>Species</b>	Bovine
<b>Clone</b>	CG-56
<b>Storage</b>	Liquid nitrogen vapor phase. Freezing medium: to complete growth medium(20%FBS), add 10%(v/v) DMSO
<b>Ship</b>	Dry Ice
<b>Immunological Donor</b>	BALB/c mouse spleen
<b>Cell Line Description</b>	Described as secreting a mouse monoclonal antibody (igg1) that reacts with rat embryonic fibers in immunofluorescence and with embryonic myosin heavy chains in Western blotting; established by fusion of NSO myeloma cells with spleen cells from a BALB/c mouse immunized with purified myosin from fetal bovine muscles confirmed as mouse with IEF of NP, PEP B and by species PCR. Cytogenetics: murine flat-moded hypertetraploid karyotype with 5% polyploidy 78-93<4n> 1-2 dmin present in most cells. Viruses: ELISA: reverse transcriptase negative; PCR: SMRV
<b>Fusion Species</b>	Mouse X Mouse Hybridoma
<b>Growth Properties</b>	suspension

<b>Morphology</b>	single, round cells in suspension (some cells adhere partly, but trypsin is not necessary to detach these cells)
<b>Propagation</b>	Complete culture medium: 90% Dulbecco's MEM (4.5 g/L glucose) + 10% horse serum + 2 mM L-glutamine; at 37 centigrade with 5-10% CO <sub>2</sub>
<b>Culture Medium</b>	DMEM with 4.5 g/L glucose 2 mM L-glutamine, supplemented with 10%(v/v) horse serum
<b>Subculturing</b>	Split ratio: 1: 3 to 1: 4, every 2 days; seed out at 0.25 x 10 <sup>6</sup> cells/ml; Doubling time: 45 hours; Harvest: maximal density of about 1.0 x 10 <sup>6</sup> cells/ml.
<b>Mycoplasma</b>	Negative in DAPI, microbiological culture, RNA hybridization assays
<b>Preservation</b>	Frozen with 70% medium, 20% horse serum, 10% DMSO at about 2 x 10 <sup>6</sup> cells/ampoule
<b>Safety Considerations</b>	The following safety precautions should be observed. <ol style="list-style-type: none"> <li>1. Use pipette aids to prevent ingestion and keep aerosols down to a minimum.</li> <li>2. No eating, drinking or smoking while handling the hybridoma.</li> <li>3. Wash hands after handling the hybridoma and before leaving the lab.</li> <li>4. Decontaminate work surface with disinfectant or 70% ethanol before and after working with hybridoma.</li> <li>5. All waste should be considered hazardous.</li> <li>6. Dispose of all liquid waste after each experiment and treat with bleach.</li> </ol>

## GENE INFORMATION

<b>References</b>	1.Schiaffino,S., Gorza,L., Pitton,G., Saggin,L., Ausoni,S., Sartore,S. & Lomo,T. (1988). Embryonic and neonatal myosin heavy chain in denervated and paralyzed rat skeletal muscle. Dev.Biol 127(1):1-11.
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