

myogenin (5FD): sc-52903



The Power to Question

BACKGROUND

Differentiation of myogenic cells is regulated by multiple positively and negatively acting factors. One well characterized family of helix-loop-helix (HLH) proteins known to play an important role in the regulation of muscle cell development includes MyoD, myogenin, Myf-5 and Myf-6 (also designated Mrf-4 or herculin). Of interest, most muscle cells express either MyoD or Myf-5 in the committed state, but when induced to differentiate, all turn on expression of myogenin. MyoD transcription factors form heterodimers with products of a more widely expressed family of bHLH genes, the E family, which consists of at least three distinct genes: E2A, IF2 and HEB. MyoD-E heterodimers bind avidly to consensus (CANNTG) E box target sites that are functionally important elements in the upstream regulatory sequences of many muscle-specific terminal differentiation genes.

CHROMOSOMAL LOCATION

Genetic locus: MYOG (human) mapping to 1q32.1; Myog (mouse) mapping to 1 E4.

SOURCE

myogenin (5FD) is a mouse monoclonal antibody raised against amino acids 144-158 of myogenin of rat origin.

PRODUCT

Each vial contains 200 µg IgG₁ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

myogenin (5FD) is available conjugated to HRP (sc-52903 HRP), 200 µg/ml, for WB, IHC(P) and ELISA; and to either phycoerythrin (sc-52903 PE), fluorescein (sc-52903 FITC) or Alexa Fluor® 488 (sc-52903 AF488) or Alexa Fluor® 647 (sc-52903 AF647), 200 µg/ml, for IF, IHC(P) and FCM.

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APPLICATIONS

myogenin (5FD) is recommended for detection of myogenin of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500).

Suitable for use as control antibody for myogenin siRNA (h): sc-29402, myogenin siRNA (m): sc-35992, myogenin shRNA Plasmid (h): sc-29402-SH, myogenin shRNA Plasmid (m): sc-35992-SH, myogenin shRNA (h) Lentiviral Particles: sc-29402-V and myogenin shRNA (m) Lentiviral Particles: sc-35992-V.

Molecular Weight of myogenin: 34 kDa.

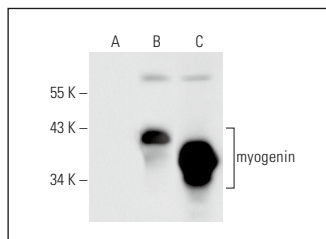
Positive Controls: SJRH30 cell lysate: sc-2287, myogenin (h3): 293T Lysate: sc-177592 or RD whole cell lysate: sc-364791.

RESEARCH USE

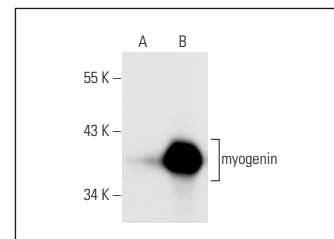
For research use only, not for use in diagnostic procedures.

STORAGE

Store at 4° C, ****DO NOT FREEZE****. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

DATA

myogenin (5FD): sc-52903. Western blot analysis of myogenin expression in non-transfected 293T: sc-117752 (A), human myogenin transfected 293T: sc-177592 (B) and SJRH30 (C) whole cell lysates.



myogenin (5FD): sc-52903. Western blot analysis of myogenin expression in non-transfected: sc-117752 (A) and human myogenin transfected: sc-116551 (B) 293T whole cell lysates.

SELECT PRODUCT CITATIONS

- Strle, K., et al. 2006. c-Jun N-terminal kinase mediates tumor necrosis factor- α suppression of differentiation in myoblasts. *Endocrinology* 147: 4363-4373.
- Cho, Y.W., et al. 2009. Histone methylation regulator PTIP is required for PPAR γ and C/EBP α expression and adipogenesis. *Cell Metab.* 10: 27-39.
- Galatioto, J., et al. 2010. CLP-1 associates with MyoD and HDAC to restore skeletal muscle cell regeneration. *J. Cell Sci.* 123: 3789-3795.
- Matheny, R.W. and Nindl, B.C. 2011. Loss of IGF-IEa or IGF-IEb impairs myogenic differentiation. *Endocrinology* 152: 1923-1934.
- Matheny, R.W., et al. 2012. Enhanced Akt phosphorylation and myogenic differentiation in PI3K p110 β -deficient myoblasts is mediated by PI3K p110 α and mTORC2. *Growth Factors* 30: 367-384.
- Mohamed, J.S., et al. 2013. Ankyrin repeat domain protein 2 and inhibitor of DNA binding 3 cooperatively inhibit myoblast differentiation by physical interaction. *J. Biol. Chem.* 288: 24560-24568.
- Zhang, W.W., et al. 2015. Identification of miR-2400 gene as a novel regulator in skeletal muscle satellite cells proliferation by targeting MYOG gene. *Biochem. Biophys. Res. Commun.* 463: 624-631.
- Wei, D.W., et al. 2017. Characterization of the promoter region of the bovine SIX1 gene: roles of MyoD, PAX7, CREB and MyoG. *Sci. Rep.* 7: 12599.
- Bi, P., et al. 2018. Fusogenic micropeptide Myomixer is essential for satellite cell fusion and muscle regeneration. *Proc. Natl. Acad. Sci. USA* 115: 3864-3869.
- Liu, D., et al. 2019. Podocan affects C2C12 myogenic differentiation by enhancing Wnt/ β -catenin signaling. *J. Cell. Physiol.* 234: 11130-11139.

PROTOCOLS

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