

## MyoD1 (D8G3) XP® Rabbit mAb



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## For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> W, IF-IC	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 45	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P15172	Entrez-Gene Id: 4654
Product Usage Information	2	<b>Application</b> Western Blotting Immunofluorescence	(Immunocytochem	istry)		<b>Dilution</b> 1:1000 1:400
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. <i>Do not aliquot the antibody.</i>				
Specificity/Sensitivity		MyoD1 (D8G3) XP <sup>®</sup> Rabbit mAb recognizes endogenous levels of total MyoD1 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gly190 of human MyoD1 protein.				
Background		Myoblast determination protein 1 (MyoD1), also called myogenic factor 3 (Myf3), is a member of the MyoD family of muscle specific bHLH transcription factors (1). This family is responsible for controlling specification of the muscle cell lineage and members are expressed only in skeletal muscle and its precursors. MyoD1 is considered a master regulator of skeletal myogenesis as its expression can induce myogenic differentiation in myoblasts, fibroblasts, and a variety of other cell types (2,3). Through ChIP-sequencing experiments, researchers have discovered that MyoD is associated with the promoters of many genes in muscle cells, but it only regulates a subset of those genes. These research studies point to regulation of MyoD transcriptional activity via epigenetic mechanisms involving SWI/SNF complexes and Polycomb and Trithorax Group proteins (4-6). Additional influences on muscle development include signal transduction through MAPK, PI3K/Akt, myostatin, NF-κB, and mTOR signaling pathways (5-7).				
1. Berkes, C.A. and Tapscott, S.J. (2005) Semin Cell Dev Biol 16, 2. Tapscott, S.J. (2005) <i>Development</i> 132, 2685-95. 3. Davis, R.L. et al. (1987) <i>Cell</i> 51, 987-1000. 4. de la Serna, I.L. et al. (2001) <i>Nat Genet</i> 27, 187-90. 5. Aziz, A. et al. (2010) Epigenetics 5, 691-5. 6. Guttridge, D.C. (2004) <i>Curr Opin Clin Nutr Metab Care</i> 7, 443 7. Ge, Y. and Chen, J. (2012) <i>J Biol Chem</i> 287, 43928-35.				2685-95. 0.	5-95.	

**Species Reactivity** Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X

TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key W: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)

Cross-Reactivity Key H: Human

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