Muscle-specific expression of the cardiac α-actin gene requires MyoD1, CArG-box binding factor, and Sp1

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Expression of the human cardiac α-actin gene (HCA) depends on the interactions of multiple transcriptional regulators with promoter elements. We report here that the tissue-specific expression of this promoter is determined by the simultaneous interaction of at least three specific protein-DNA complexes. The myogenic determinant gene MyoD1 activated the transcription of transfected HCA-CAT promoter constructs in nonmuscle cells, including CV-1 and HeLa cells. Gel mobility-shift and footprinting assays revealed that MyoD1 specifically interacted with a single consensus core sequence, CANNTG, at ~50. Previously characterized sites interact with a protein identical with or related to the serum response factor (SRF) at ~100 and Sp1 at ~70. All three elements must be intact to support transcription in muscle cells: site-specific mutation within any one of these three elements eliminated transcriptional expression by the promoter. Furthermore, expression of the promoter in embryonic Drosophila melanogaster cells that lack MyoD1 and Sp1 is strictly dependent on all three sites remaining intact and on the presence of exogenously supplied Sp1 and MyoD1. These experiments suggest that the presence of three sequence-specific binding proteins, including MyoD1, and their intact target DNA sequences are minimal requirements for muscle-specific expression of the HCA gene.

[Key Words: Transcription, MyoD1, serum response factor, Sp1, cardiac actin, muscle; differentiation]

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Transcriptional regulation requires the interaction of proteins with sequence-specific DNA elements. DNA-binding proteins may be expressed ubiquitously in eukaryotic cells, they may be restricted to a particular cell or tissue type, and their abundance or binding activity may be regulated by extracellular signals or inducers. Post-translational modifications may also contribute to regulation of the binding activity of a factor. Transcriptional activation appears to involve complex and often cooperative interactions of these factors [Andrisani et al. 1988; Davis et al. 1990; Simmons 1990; for review, see Jones 1990] and gene expression is probably determined at least in part by the availability or activity of factors in a particular cell type and by the unique organization of the DNA target elements within the gene promoter [Guarente 1988; Prashne 1988].

Previously, we analyzed the mechanisms governing the tissue-specific expression of the human cardiac α-actin [HCA] gene (Minty and Kedes 1986; Miwa and Kedes 1987; Miwa et al. 1987; Gustafson et al. 1988; Gustafson and Kedes 1989). Cardiac α-actin is the major α-actin in adult cardiac muscle [Gunning et al. 1983], is expressed in both skeletal muscle and heart during embryogenesis [Minty et al. 1982], and is the dominant α-actin gene expressed during differentiation of both established cell lines [Bains et al. 1984] and primary myogenic cells [Minty et al. 1986a; Gunning et al. 1987]. The HCA gene promoter is expressed exclusively in myogenic cells [Minty and Kedes 1986a,b] and contains numerous nuclear protein-binding domains upstream of the TATA box [Gustafson and Kedes 1989], several of which display cooperative transcriptional activating interactions [Miwa and Kedes 1987]. A promoter domain starting at ~117 bp from the start of transcription is sufficient to convey muscle-specific expression to a heterologous reporter gene [Minty et al. 1986]. Deletion and clustered mutagenesis analyses demonstrated that a decameric sequence at ~100 bp from the transcription start site, CC[AT]6GG, designated as a CArG box, is required for both tissue-specific and vigorous expression from this promoter [Miwa et al. 1987]. Subsequent analysis unexpectedly demonstrated that a protein that binds to this CArG box appears to be expressed in all cell types [Boxer et al. 1989a,b] and is immunologically and biochemically indistinguishable from the ubiquitously expressed serum response factor (SRF) [Boxer et al. 1989b]. Furthermore, by modifying sequences between the CArG element and the TATA box we have shown...
that the HCA promoter can be activated in nonmuscle cells [Webster and Kedes 1990, T. Miwa and L. Kedes, unpubl.]. In these instances, expression of the modified promoter in nonmuscle cells was still dependent on an intact CArG element. These data, along with the report of a muscle-specific protein binding to the chick skeletal α-actin CArG box [Walsh 1989] suggested that both muscle and nonmuscle proteins can bind to the CArG box. Analogous examples are reported in the literature, including the octamer binding proteins (Tanaka and Herr 1990). An alternative possibility is that CArG box-binding factors in the HCA gene are not tissue specific, and additional properties of the promoter are required to determine muscle-specific expression.

The recent finding that several members of a muscle determination gene family [Davis et al. 1987; Pinney et al. 1988; Braun et al. 1989; Wright et al. 1989] are able to activate the myogenic program in a variety of cells as well as promote expression of some muscle-specific genes [Weintraub et al. 1989] prompted us to investigate the role of one of those muscle determinant genes, MyoD1, in controlling the HCA gene expression. Preliminary studies [Sartorelli et al. 1989], as well as an evaluation of a chicken homolog of MyoD1, CDM1 [Lin et al. 1989], have shown that muscle determinate genes can indeed activate expression of the cognate cardiac actin gene.

Here, we report that the transcriptional regulatory activity of the complex formed by the CArG box and its binding protein(s) appears to involve additional interactions with immediately adjacent protein–DNA assemblies. To activate transcription the CArG-box complex at position −100 requires the presence of the recognition sites for both the ubiquitous transcription regulatory factor Sp1 at −75 and the myogenic determinant, MyoD1, bound at −50. All three DNA elements must be intact to support transcription in muscle cells: When site-specific mutation of any one of these three elements precludes binding of its sequence-specific protein, it also foils transcriptional expression of the promoter. Furthermore, expression in nonmuscle cells of transfected DNA constructs that carry the intact promoter is strictly dependent on the presence of an intact CArG box element and the presence of transcription factors Sp1 and MyoD1. Thus the combination of three protein–DNA complexes consisting of at least two nonmuscle-specific proteins plus a cell type-specific regulatory protein, MyoD1, is required for muscle-specific expression of the HCA gene.

Results

The MyoD1 protein activates transcription from the HCA promoter in nonmyogenic cell lines

Previous studies from this laboratory [Gustafson and Kedes 1989, Webster and Kedes 1990; T. Miwa and L. Kedes, unpubl.] have suggested that factors in addition to the CArG-binding factor may be involved in transcriptional activation of the HCA gene specifically in muscle cells. Because members of the MyoD1 family are obvious candidates for such a role, we assayed for a trans-activation function of the HCA promoter by MyoD1. To do this we cotransfected nonmuscle cells with plasmids carrying the HCA gene promoter linked to the chloramphenicol acetyltransferase gene (HCA–CAT) and with plasmids that express MyoD1. Because trans-activation of the HCA–CAT transcription could arise by activation of the endogenous myogenic program of the test cells, we examined several cell lines in this way including NIH-3T3, HeLa, and Green monkey kidney cells (CV-1). Unlike a variety of other cell types including NIH-3T3, liver, fat, or nerve cells, both CV-1 and HeLa cells have proven refractory to MyoD1-induced myogenic conversion [Weintraub et al. 1989]. The results of such a cotransfection experiment are presented in Figure 1. In both CV-1 and HeLa cell lines (as well as in NIH-3T3 cells, data not shown), MyoD1 trans-activated the whole HCA promoter (HCA485-CAT) and the shorter construct containing the CArG element at position −100 (HCA117-CAT) but not the HCA construct truncated at −47 (HCA47-CAT). Furthermore, the HCA construct LS6, bearing a mutated CArG box, was not trans-activated by MyoD1. A map of the DNA sequences of the HCA promoter from −26 to −118 is presented in Figure 2A, which also depicts the sites of protein–DNA interactions described previously [Gustafson et al. 1988; Gustafson and Kedes 1989] and in this discussion.

Expression of the ubiquitous β-actin promoter was unaffected by the presence of MyoD1, and the tissue-specific β-globin gene promoter remained repressed. Although both HeLa and CV-1 cells sustain MyoD1-driven HCA expression, HCA485-CAT expresses three- to four-fold more than HCA117-CAT in HeLa cells but at approximately the same levels in CV-1 cells. The HeLa response mimics the relative transcriptional response of these two promoter constructs in myogenic cell lines [Minty and Kedes 1986].

Optimal trans-activation occurred when 1 μg of the MyoD1 cDNA vector was cotransfected with 5 μg of the HCA/CAT constructs (approximately a 1 : 5 molar ratio). Increasing the amount of the expression vector to 5 μg (approximately a 1 : 1 ratio) inhibited transcription from both the HCA and β-actin promoters [Fig. 1]. This phenomenon might be ascribed either to “transcriptional interference” [Meyer et al. 1989] or “squelching” [Gill and Ptashne 1988]. This initial result suggested that MyoD1 protein might physically interact with the cardiac actin gene promoter as a positively acting regulatory transcription factor and that the site(s) of interaction were located between −117 and +68 of the gene. To test this possibility further we examined the ability of this region of the gene to physically interact with MyoD1.

The glutathione–MyoD1 fusion protein and muscle MyoD-like nuclear factors bind to the HCA promoter

DNA fragments representing segments of the HCA promoter encompassing nucleotides −485 to +68 were derived by restriction endonuclease digestion and assayed
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Figure 1. MyoD1 activates transcription from the HCA promoter in different nonmyogenic cell lines. A constant amount (5 µg) of different HCA-CAT chimera constructs (pHCA485CAT, pHCA118CAT, and pHCA47CAT) were mixed with varying amounts of the LTR–MyoD1 expression vector and transfected into CV-1 [A] and HeLa [B] cells. The ubiquitously expressed β-actin and the erythroid-specific β-globin CAT constructs were included as controls. The LS6 construct has a mutated CArG box at -100 (see Fig. 2). (0, 1, and 5) Microgram amounts of the LTR–MyoD1 expression vector used in each experiment. After transfection, the cells were cultured for 48 hr and the determination of CAT activity was performed as described in Methods. The figure is an autoradiograph of thin-layer chromatographic plates after separation of the acetylated and nonacetylated forms of [14C]-chloramphenicol.

For interaction with a bacterially produced glutathione–MyoD1 fusion protein by gel electrophoretic mobility-shift (GEMS) assays. A DNA fragment representing nucleotides -70/+68 of the gene bound the MyoD1 fusion protein [Fig. 3A, lane 1] but the glutathione-S-transferase protein itself did not bind [data not shown]. The specificity of this interaction was challenged by adding different DNA fragments spanning the HCA promoter. A 100-fold molar excess of nonlabeled -70/+68 or -127/-47 fragment competed for the MyoD1 binding (lanes 2 and 3), whereas the remaining promoter fragments (lanes 4 and 5) used at the same molar excess failed to prevent the interaction. Furthermore, incubation of the glutathione–MyoD1 protein with HCA DNA fragment -47/+68 did not result in any specific shifted bands in the GEMS assay [lanes 6] and did not compete for binding [data not shown]. We conclude that a binding site for MyoD1 appears to lie between -70 and -47 of the HCA promoter. We sought to verify that a muscle cell nuclear factor interacts with the same HCA DNA sequence recognized by the glutathione–MyoD1 fusion protein. C2 and HeLa cell nuclear extracts were tested for the presence of DNA-binding proteins that would interact with the -70/-40 DNA element of the HCA gene. The data presented in Figure 3B demonstrate that several major protein–DNA complexes [right-hand symbols in Fig. 3B] are observed with C2 cell nuclear extracts [lane 5]. The intensity of the binding of the bands indicated by filled circles is diminished in the presence of excess unlabeled probe [lane 6]. The bands with faster mobility [indicated by the bracket] were variably affected and appear to be nonspecific. A mutant oligonucleotide [5 µg, see Fig. 2A] does not compete for any of the complexes [data not shown]. A different and less complex banding pattern was seen with HeLa cell nuclear extracts [open circle, left side of Fig. 3B] and this DNA–protein complex [lane 1] appears unaffected by the presence of an excess of unlabeled probe [lane 2]. We used monoclonal antibodies directed against the MyoD amino acid residues 66–99 of a trpE–MyoD1 fusion protein to test whether the DNA–protein complexes that we observed represented interactions with proteins antigenically related to MyoD1. The monoclonal antibody, αMyoD1, as well as a second monoclonal antibody [6F1], directed against the bovine histone 1 protein, which was used as a control serum, were generously provided by Stave Kohtz (Mount Sinai Medical Center, New York, in prep.). αMyoD1, but not the control antibody 6F1, eliminated the major DNA–protein complexes formed with C2 cell nuclear extracts [compare bands indicated by filled circles in lanes 7 and 8]. Neither antibody affected the nonspecific protein–DNA complexes formed with HeLa cell nuclear extracts [lanes 3 and 4].

These data suggest that one or more nuclear proteins that share antigenicity with MyoD are present in C2 muscle cells but not in HeLa cells and bind to the MyoD binding site we identified by use of the bacterial fusion protein. The inhibition of MyoD1 binding by αMyoD1 monoclonal antibody in a GEMS assay has been observed by others [A. Lassar and S. Kohtz, pers. comm.]. Because oligonucleotides often form multiple complexes in retardation gels, even with purified binding proteins, it is not clear whether the multiplicity of protein–DNA complexes we observed with C2 cell nuclear extracts represents more than one MyoD-related binding protein.

To define the nucleotides involved in the HCA–MyoD1 interaction we used a synthetic DNA oligonucleotide spanning HCA nucleotides -70 to -40 and the glutathione–MyoD1 fusion product in methylation interference assays. Both sense and antisense DNA strands were investigated and, as shown in Figure 4, A and B, the methylation of two guanine residues at positions -56 and -49 on the antisense strand and of a guanine at po-
Figure 2. The HCA gene proximal promoter nucleotide sequences, nuclear protein–DNA interaction sites, mutations and comparison of the human and mouse sequences. (a) The nucleotide sequence of the HCA gene from -26 to -113. Boldface nucleotides and shaded areas indicate the locations of the binding domains for the transcriptional regulators, SRF, Sp1, and MyoD1, TUBF and the TATA box-binding TFID. Circles above and below some bases represent sites of complete (●) and partial (○) interference with factor binding, as determined by guanine methylation interference experiments (Gustafson and Kedes 1989) [SRF, Sp1, and TUBF; and this work (MyoD1)]. The sequences of the coding strand of SRF-, Sp1-, and MyoD1-binding site mutations (LS-6, μSp1, and μ6, respectively) are shown below the wild-type sequence. Mutated bases are shown in underlined lowercase. (b) Nucleotide sequence comparison (coding strand) of the human (Minty and Kedes 1986) and mouse HCA sequences (Garner et al. 1986). Capitalized, shaded base pairs represent the location in the human sequence and the putative locations in the mouse sequence of the transcription regulatory factors SRF, Sp1, and MyoD1. The two dots just upstream of the human Sp1 site represent the minimal gap required to maintain colinearity of the two sequences for comparative purposes. Dots between the two sequences are located at the mismatches.

position -51 on the sense strand interfered with the MyoD1 protein binding.

Densitometric scanning [data not shown] confirmed strong interference of the antisense guanine at position -56, weak interference of the guanine at -49 and weak interference of the guanine on the sense strand at -51. No additional guanine methylation appeared to interfere with binding of the fusion protein. This region contains a hexameric sequence [CANNTG] [see Fig. 2] that has been proposed as the core MyoD1-binding sequence in the mouse MCK gene (Lassar et al. 1989). Interestingly, a GCTGC motif involved in the MyoD1–MCK interaction and located 3' with respect to the CANNTG sequence [Lassar et al. 1989] is located 5' to the HCA interaction site but apparently does not participate in the MyoD1 binding to the HCA gene.

The trans-activation induced by the MyoD1 protein in nonmyogenic cell lines requires at least three intact regulatory modules in the HCA promoter

To test whether MyoD1 activates transcription by binding to the HCA promoter, several promoter mutants were generated by introduction of site-specific mutations at or adjacent to the nucleotides shown to interact with the MyoD1 protein. Because previous experiments [Gustafson and Kedes 1989, T. Miwa, unpubl.] suggested that the transcriptional activator Sp1 may participate in regulating cardiac α-actin gene expression, we assessed the functional role of an Sp1 binding site located at -70, just upstream from the MyoD1 binding site [Gustafson and Kedes 1989] by constructing a mutant at this site as well. The relevant sequences of SRF, Sp1, and MyoD1 site mutants are shown in Figure 2A. As demonstrated previously, the LS-6 mutant [Miwa and Kedes 1987] fails to bind SRF or a related protein and does not compete for its binding [Gustafson et al. 1988]. The HCA -70/–47 μ6 MyoD1-site mutant does not compete for MyoD1 binding to the nonmutated promoter [Fig. 5A] and does not complex with the protein [data not shown]. Similarly the HCA oligomer -87/–65 μSp1 mutant [see Methods] does not compete for Sp1 binding to the HCA -87/–65 oligonucleotide [Fig. 5B], but does compete for MyoD1 binding [data not shown] and the nonmutated DNA self-competes for Sp1 binding [Fig. 5B, lane 3] [Gustafson and Kedes 1989]. The appearance of a doublet when purified Sp1 binds in the GEMS assay has been described previously [Dawson et al. 1988; Gustafson and Kedes 1989].

The HCA mutants HCA177μ6-CAT, HCA177μSp1-CAT, and LS6 [HCA177CAT with mutations in the CArG element] were cotransfected into CV-1 cells with the MyoD1 cDNA expression vector. The mutants μ6 and μSp1 [Fig. 6A] and LS6 [data not shown] showed little or no transcriptional induction by the MyoD1 suggesting that trans-activation by MyoD1 correlates not only with an intact MyoD1 binding site but also with intact binding sites for Sp1 and CArG binding factor.

Intact binding sites for MyoD, Sp1, and SRF are indispensable for HGA activation in myogenic cell lines

Additional muscle-specific factors that are not required for in vitro DNA–protein interaction, may be
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Figure 3. The glutathione–MyoD1 fusion protein and factors present in C2 cell line nuclear extracts bind the HCA promoter. (a) The glutathione–MyoD1 fusion protein (0.1 μg) was incubated with the end-labeled -70/+68 HCA fragment (lane 1). Poly[d(I-C)] (0.1 μg) was added to each reaction as nonspecific competitor. A 100-fold molar excess of various unlabeled HCA DNA fragments spanning the whole promoter region from -485 to +68 was employed as competitor (lanes 2–5). The glutathione–MyoD1 protein complex formation is shown by the appearance of a major shifted band in the gel mobility-shift assay. In lane 6, the HCA fragment -47/+68 was analyzed for protein interaction. (b) The same end-labeled HCA oligomer employed in a was used in GEMS assays with nuclear extracts from either HeLa cells (lanes 1–4) or C2 myotubes (lanes 5–8). All lanes in the photograph were run on the same gel. The monoclonal antibodies 6F1 and αMyoD (lanes 3, 4, 7, and 8) were preincubated with the nuclear extracts before adding the end-labeled HCA -70/-40 probe. A 50-fold molar excess of nonlabeled HCA -70/-40 (lanes 2 and 6) served as DNA competitor. (O) Nonspecific retarded band detected apparently only with the HeLa cell extract; (●) specific shifted complexes formed when nuclear extract from C2 myotubes was employed; (bracket) complexes that were unaffected by the presence of αMyoD antibody. After electrophoresis, the gels were dried and autoradiographed over night.

required for MyoD1 to promote HCA gene expression in vivo. Lack of such factors in the “nonphysiological” nonmuscle environment of CV-1 could be responsible for the failure of MyoD1 to trans-activate the HCA mutants. To distinguish this possibility, different HCA promoter–CAT constructs including μ6 and μSp1 were transiently transfected in the myogenic C2 cell line and assayed for expression. The lack of expression by the Sp1 and MyoD1 binding site mutants is consistent with the result in CV-1 cells (Fig. 6B). Relative to the activity of the wild-type HCA177CAT promoter, HCA177μ6-CAT and HCA177μSp1-CAT expressed at 3% and 23%, respectively. The LS6 mutant (HCA177μCARG-CAT) has been shown previously (Miwa and Kedes 1987) to express at 12% wild type. Taken together, these results suggest that the MyoD1 gene product and its intact target element are required for the expression of the cardiac α-actin gene promoter. Furthermore, muscle-specific expression does not appear to require either the presence of other muscle-specific auxiliary factors or tissue-specific isoforms or modifications of the CARG binding factor. It does however appear to require unaltered elements for CARG binding factor and Sp1.

The HCA transcription is reconstituted in Drosophila melanogaster cells on complementation with MyoD1 and Sp1

We tested the functional activity of the promoters and the transcription regulatory proteins in the Schneider
cell line derived from the D. melanogaster embryos, which are devoid of many mammalian transcriptional activator homologs including Sp1 [Courey and Tjian 1988; Santoro et al. 1988]. Preliminary experiments established that the Moloney sarcoma virus (MSV) LTR segment driving MyoD1 expression in the LTR–MyoD1 construct was active in Schneider cells [data not shown]. Accordingly, we transfected Schneider cells with pHCA177-CAT and with either or both the LTR–MyoD1 expression vector and the plasmid pPACSp1. pPACSp1 carries the Drosophila nonmuscle actin gene promoter driving human Sp1 cDNA transcription [Courey and Tjian 1988]. As demonstrated in the results presented in Figure 7A, Sp1 and MyoD1 are required for transcription and have a synergistic effect [24-fold]. The LTR–MyoD1 vector alone has no effect on HCA177-CAT expression; the pPACSp1 plasmid alone engenders but weak expression [less than twofold over its effect on the HCA177μSp1-CAT mutant]. To demonstrate further the requirement of the CArG binding factor, Sp1, and MyoD1 in activating the HCA transcription, the LS6, HCA177μ6-CAT and HCA177μSp1-CAT mutants were cotransfected in Schneider cells with different combinations of Sp1 and MyoD1 expression vectors [Fig. 7B]. The results of these experiments reveal that Sp1, the CArG binding factor, and the tissue-specific factor MyoD1 are all simultaneously required for expression of the HCA gene. Several constructs showed low level [two- to fourfold] increases of expression in the presence of exogenously supplied Sp1 alone [Fig. 7]. Although these effects remain unexplained, we note that they were not observed in every construct or in the presence of combinations of exogenously supplied Sp1 and MyoD1.

Figure 4. Methylation interference footprinting of the MyoD1 protein on the HCA promoter. The coding [A] and noncoding [B] strands of an oligonucleotide representing the –70/-40 HCA promoter region were methylated partially and analyzed after binding with the glutathione–MyoD1 protein. [F and B] Free and bound probes, respectively. Analysis included densitometric scanning of the autoradiographs [data not shown]. The coding strand shows a single, partially interfered guanine residue [arrowhead in A]. The noncoding strand shows one complete and one incomplete guanine interference residue [arrows in B]. In B, twice as many decays per minute were applied in the bound probe lane [B] such that the degree of interference should be judged by comparing the intensity of bands above and below the contacted nucleotide as well as between the different lanes. [C] Summary of the methylation footprint data. [●] Partial interference; [●] strong interference.

Discussion

Here, we report that the MyoD1, Sp1, and SRF or SRF-related proteins bind specifically to the human cardiac α-actin gene promoter and are involved in its activation. The organization of the proximal promoter of the HCA gene is summarized in Figure 2A. The mechanisms by which the muscle determinant factors of the MyoD1 family [Davis et al. 1987] activate expression of muscle-specific genes are not well understood. The muscle creatine kinase [MCK] gene and the human cardiac actin gene are apparently activated by the MyoD1 product [Lassar et al. 1989] (and this work) but it is not clear how other genes interact with MyoD1. For example, a chicken congener of MyoD1, CMD1, promotes transcription of the chicken skeletal α-actin gene [and the myosin light chain 1/3 gene [Lin et al. 1989]], but neither MyoD1 fusion protein [Lassar et al. 1989] nor in vitro translation products [Lin et al. 1989] appear to bind to the skeletal α-actin DNA. We have not investigated the role of other members of the MyoD1 muscle-determinant gene family, such as myogenin [Wright et al. 1989] and Myf5 [Braun et al. 1989] in activating HCA gene expression. These two myogenic determinant gene products have been shown recently to interact with the same DNA sequence recognized by MyoD1 [Braun et al. 1990, Brennan and Olson 1990]. Nevertheless, it is unlikely that myogenin is involved directly in the effects we observe in CV-1 cells because CV-1 cells stably expressing transfected MyoD1 fail to activate the endogenous myogenin gene [A. Lassar, pers. comm.].

The MyoD1 product belongs to a super-family of proteins having in common an amphipathic helix–loop–helix domain [Murre et al. 1989a; Kingston 1990]. Among those proteins are the product of the proto-oncogene c-myc [Alitalo et al. 1983], the lyl-1 gene product [Mellentin et al. 1989], the we enhancer-binding proteins E12 and E47 [Murre et al. 1989a, b], the immunoglobulin enhancer-binding protein TFE3 [Beckman et al. 1990] and the products of the Drosophila genes daughterless [Caudy et al. 1988a, b], twist [Thisse et al. 1988], enhancer of split [Klaembt et al. 1989], and achaete-scute T4/T5 [Alonso and Cabrera 1988]. The MyoD1 protein can form heterodimers in vitro with the E12 protein [Murre et al. 1989b], and heterodimerization of MyoD1 and E12 or related proteins takes place also in vivo.
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1988; Halozonetis et al. 1988; Kouzarides and Ziff 1988; Nakabeppu et al. 1988; Rauscher et al. 1988) and usually increases the affinity of the heterodimer complex for the DNA.

Binding of MyoD1 alone (Lassar et al. 1989) or MyoD1-E12 complexes (Murre et al. 1989b) to DNA requires a defined nucleotide sequence. The CANNTG motif is the minimal region in the MCK enhancer recognized by the MyoD1, MyoD1/E12, and MEF 1 factors (Buskin and Hauschka 1989) and is present (Donoghue et al. 1988) in the MyoD1 responsive enhancer of the mouse MLC 1/3 gene (Lin et al. 1989). An identical, and functional, consensus element lies at the core of the MyoD1 binding site in the HCA promoter. Abolition of the consensus CANNTG sequence prevented MyoD1

Figure 5. The HCA mutant μ6 does not bind MyoD1 and base changes in the Sp1 site of the HCA −87/−65 oligomer prevent binding of Sp1. [a] The HCA μ6 mutant (see Fig. 2a) was assayed for its binding property by use of GEMS assay with purified glutathione–MyoD1 fusion protein. [Lane 1] The end-labeled HCA −70/−47 oligonucleotide, incubated with glutathione–MyoD1 protein, generated a complex visible as a shifted band. A 50-fold molar excess of unlabeled HCA −70/−47 [lane 2] or HCA −70/−47 μ6 [lane 3] oligonucleotides were included in the protein–DNA binding reaction. [b] The end-labeled HCA −87/−65 oligomer [0.1 ng] was incubated with homogeneously purified Sp1 [1 ng] and analyzed by use of GEMS assay [lane 1]. The arrows point to the two shifted complexes. Competition experiments were performed adding a 50-fold molar excess of either unlabeled HCA −87/−65 (wild type, lane 2) or HCA −87/−65 μSp1 [lane 3] oligonucleotides. The figure is an autoradiograph of the dried GEMS assays.

Figure 6. The HCA μ6 and μSp1 mutations no longer function as promoters and cannot respond to MyoD1. [A] Various chimeric plasmid constructs [5 μg each] were transfected in the absence [−] or presence [+] of 1 μg of the LTR–MyoD1 expression vector in the CV-1 cell line: [lanes 1 and 2] HCA177 −CAT; [lanes 3 and 4] HCA177μ6-CAT; [lanes 5 and 6] HCA177μSp1-CAT. [B] HCA177-CAT [5 μg] [lane 1], HCA177μ6-CAT [lane 2], HCA177μSp1-CAT [lane 3], β-actin-CAT [lane 4] and β-globin CAT [lane 5] were transfected into the C2 myogenic cell line. After 48 hr, the cells were harvested and the CAT assay performed as described in Methods.

[Davis et al. 1990]. These findings, along with the ubiquitous expression of E12 (Murre et al. 1989b), the increased affinity of MyoD1/E12 heterodimers versus homodimers for specific DNA interaction, and the indirect evidence for a biological activity of the MyoD1–E12 complex in vivo (Davis et al. 1990), support the hypothesis that the combination of MyoD1 and E12 is critical for activation for the myogenic program. Heterodimerization has been reported also for other transcription regulatory factors (Chiu et al. 1988; Curran andFranza
were harvested after 48 hr and assayed for CAT activity as described in Methods. MyoD1. Various CAT expression vectors (5 μg each) were transfected into Drosophila Schneider cells in the absence (−) or presence of different combinations of the pPacSp1 and LTR–MyoD1 expression vectors as indicated in the figure. (a) pHCA177-CAT or pHCA177-LS6-CAT. (b) pHCA177μ6-CAT, pHCA177μSp1-CAT, or pSV40-CAT. pSV40-CAT is inactive in the absence of Sp1 (Courey and Tjian 1988). The cells were harvested after 48 hr and assayed for CAT activity as described in Methods.

Figure 7. Drosophila Schneider cells express the HCA gene when complemented with exogenously introduced Sp1 and MyoD1. Various CAT expression vectors (5 μg each) were transfected into Drosophila embryo Schneider cells in the absence (−) or presence of different combinations of the pPacSp1 and LTR–MyoD1 expression vectors as indicated in the figure. (a) pHCA177-CAT or pHCA177-LS6-CAT. (b) pHCA177μ6-CAT, pHCA177μSp1-CAT, or pSV40-CAT. pSV40-CAT is inactive in the absence of Sp1 (Courey and Tjian 1988). The cells were harvested after 48 hr and assayed for CAT activity as described in Methods.

binding and eliminated the functional response of the promoter to MyoD1 protein. The HCA promoter is strongly trans-activated when cotransfected with a MyoD1 cDNA expression vector in both CV-1 and HeLa cells. This is in contrast to the observations that neither CV-1 nor HeLa cells support efficient trans-activation of the MCK or the chicken cardiac α-actin promoters [Lin et al. 1989; Weintraub et al. 1989]. These experiments are consistent with a model for HCA promoter expression in which MyoD1 is the only required muscle specific protein. However it is not clear why HCA but neither MCK nor chick cardiac α-actin promoter are trans-activated by MyoD1 or MyoD1-related factors in CV-1 [Lin et al. 1989; Weintraub et al. 1989]. One possible explanation relies on different amounts of cotransfected plasmid employed by us and by others [Lin et al. 1989; Weintraub et al. 1989] and/or on different transfection protocols [Weintraub et al. 1989].

The HCA promoters containing the mutant sequences LS6, μ6, or μSp1 were not trans-activated by MyoD1 in either CV-1 or HeLa cells. The same mutant elements were totally inactive (μ6) or showed only a vestigial inducibility (LS6 and μSp1) when transfected in the C2 myogenic cell line. Because MyoD1 does not interact either with the CArG motif at −100 or the Sp1 site at −75 [data not shown], we conclude that an SRF-related factor, Sp1, and MyoD1 cooperate in insuring the tissuespecific transcription of the HCA gene and that the muscle-specific factor MyoD1 by itself is not sufficient to do so. The results presented here confirm the hypothesis that the CArG box motif is a necessary element for the cardiac α-actin transcription in concert with other cis-acting elements [Minty and Kedes 1986; Miwa and Kedes 1987; Phan-Dinh-Tuy et al. 1988; Mohun et al. 1989; Taylor et al. 1989; Webster and Kedes 1990], but make unlikely the possibility that the CArG box itself determines the tissue-specific expression of the human cardiac actin gene. That role appears to be reserved for MyoD1 or a member of the MyoD1 family.

Experiments using Schneider cells [Fig. 7] support the conclusion that the HCA expression can be induced by the simultaneous presence of Sp1, MyoD1, and Drosophila SRF [Norman et al. 1988]. In addition, the daughteless gene product, which appears to be a homolog of the mammalian El2 protein [Caudy et al. 1988a,b], may be playing a role in cooperation with MyoD1 in these cells. The existence of both functional and positional relationships among the Sp1, MyoD1, and SRF-related factor binding sites is supported strongly by the structural and sequence conservation within this region when the human [Minty and Kedes 1986] and mouse sequences [Garner et al. 1986] are compared [Fig. 2B]. The sites are identical between the two species, and the whole promoter region is highly conserved. The few base mismatches that do occur cluster between the functional domains that we have defined. The pentameric GCTGC sequence associated with the MyoD1 core consensus (see above) is also conserved. The cardiac α-actin gene is strongly expressed in skeletal muscle during embryogenesis in man [Minty et al. 1986] and mouse [Minty et al. 1982]. The gene is also coexpressed in adult human skeletal muscle [Gunning et al. 1983], in immortalized myogenic cell lines [Bains et al. 1984], and in human primary muscle cells [Minty et al. 1986].

As its name implies, the cardiac α-actin mRNA is a prominent transcript in heart muscle but, as of this writing, no proteins or cDNAs homologous to MyoD1 have been reported in heart muscle at any stage of vertebrate development [Sassoon et al. 1989]. Does the absolute reliance of HCA transcription on the presence of MyoD1 in skeletal muscle mean that the HCA gene must be subjected to an entirely different set of transcriptional controls in heart muscle? And that the MyoD1 binding site is not a critical DNA regulatory element in the heart? It would seem possible, therefore, that a heart-specific regulatory factor, possibly analogous to MyoD1 but sufficiently diverged so that it does not share easily detectable nucleotide sequences or epitopes, binds to the MyoD1 element of the HCA promoter. Experiments of the kinds reported here but performed in cardiocytes and developing embryos can ad-
dress these questions and unravel the mechanisms governing the disparate patterns of α-actin gene expression during cell differentiation and development.

**Methods**

**Plasmids, site-directed mutagenesis, and synthetic oligonucleotides**

The HCA–CAT constructs employed have been described previously (Minty and Kedes 1986; Miwa and Kedes 1987). In brief, a promoterless plasmid was constructed consisting of the HindIII–BamHI restriction endonuclease fragment of pSV2CAT (Gorman et al. 1982) containing the bacterial chloramphenicol acetylase gene coupled to the SV40 small t intron and early region polyadenylation signals. This DNA fragment was inserted in pBR322 to create pHCA0CAT. A DNA segment of the human cardiac actin gene from −177 bp from the start of transcription (EcoRI restriction endonuclease linker end) to +68 (HindIII restriction endonuclease linker end) was used to replace the EcoRI–HindIII fragment of pHCA0CAT to create pHCA177CAT. Gel retardation assays and methylation interference footprinting studies have defined seven sites of binding by nuclear proteins. These seven binding sites are: CArG box 2, which is distinct from that of SRF; CArG box 1, a site for SRF binding at −100 (Gustafson et al. 1988; Miwa et al. 1987); three Sp1 sites at −80, +20, and +33; the site for TATA-box upstream binding factor (TUBF) at −40; that binds just upstream of the TATA box and is otherwise uncharacterized (Gustafson and Kedes 1989); and a MyoD1 binding site at −50 (this work). In addition the TATA box presumably is the site for interaction with the constitutive transcription initiation complex but this has not been evaluated directly in the cardiac actin promoter. The sequences and binding sites critical to this study are diagrammed in Figure 2.

Mutations in a number of these binding sites were created. Mutation of CArG box 1, μ6, and the mutation of the Sp1 site at −80, μSp1, were generated by use of the polymerase chain reaction (PCR) (Higuchi et al. 1988). Briefly, different synthetic oligonucleotides complementary to the HCA wild-type promoter sequence except for the base substitutions to be introduced and an oligomer complementary to the CAT gene were annealed, after heat denaturation, to pHCA177CAT. Taq polymerase-mediated elongation in the presence of dATP, dGTP, dCTP, dTTP and heat denaturation of the PCR products were performed for a total of 25 cycles using the DNA Thermal Cycler (Perkin Elmer Cetus). The PCR products were cleaved with restriction endonucleases Apal and HindIII and were used to replace the unique Apal–HindIII fragment in the promoter. The resulting plasmids were designated as pHCA177 μ6-CAT and pHCA177 μSp1-CAT. The integrity of the base substitutions was verified by DNA sequencing of each construct. The sequences of the coding strand of the oligomers employed in the methylation interference assays and for site-directed mutagenesis are shown below. The Sp1 and MyoD1 binding sites are shown in shaded boxes; mutated bases are shown in underlined lowercase.

The sequence of the oligomer complementary to the CAT gene is: 5′-GTGACCTATGGTGTCG-3′.

The plasmids used in transfection experiments were purified by an alkaline lysis and by two rounds of CsCl density gradient sedimentation in the presence of ethidium bromide (Maniatis et al. 1982). The LTR-MyoD1 eukaryotic expression vector was a kind gift of Andrew Nassar and Harold Weintraub (Fred Hutchinson Cancer Center, Seattle, WA) and is described in Davis et al. (1987). The plasmid pPAcSp1 was generously provided by Robert Tjian (Courery and Tjian 1988).

**Production and purification of the glutathione–MyoD1 fusion protein**

The bacterial vector expressing the glutathione-MyoD1 fusion protein was provided by Andrew Nassar, and the purification of the protein from bacterial lysates was performed according to Lassar et al. (1989).

**GEMS and methylation interference assays**

DNA fragments and synthetic oligonucleotides used in GEMS assays were end-labeled with T4 polynucleotide kinase and γ-[32P]ATP and purified by electrophoresis in a polyacrylamide gel. The binding reactions for the GEMS were performed in the presence of end-labeled DNA fragments or synthetic oligonucleotides [6000 cpm], purified glutathione–MyoD1 fusion protein [0.1 μg] with the poly[d(I-C)] [0.1 μg] as nonspecific competitor. The binding buffer is described in Nassar et al. (1989). Purified Sp1 (kindly provided by R. Tjian) [1 ng] was employed in the GEMS assay with different synthetic oligonucleotides. The binding reactions containing Sp1 were carried out in 20 mM HEPES [pH 7.9], 1.5 mM MgCl2, 0.2 mM EDTA, 100 mM KC1, and 20% [vol/vol] glycerol. No nonspecific competitor was added. GEMS assays with HeLa and C2 myotubes nuclear extracts [10 μg] were performed in the same buffer employed with purified Sp1 [see above] in the presence of the poly[d(I-C)] [2 μg] and 0.1 ng of end-labeled HCA-70/40 probe. The monoclonal antibodies 6F1 and αMyoD1 [2.5 μg each] [a generous gift of Stave Kohetz, Mount Sinai Medical Center, NY] were preincubated at room temperature for 15 min with HeLa and C2 myotubes nuclear extracts before adding end-labeled DNA probe and continuing the reaction for an additional 15 min. For the methylation interference assay, the end-labeled synthetic oligonucleotides [200,000 cpm] were partially methylated [Sturm et al. 1987] prior to complex formation with purified glutathione MyoD1 fusion protein [2 μg]. The complexed and free DNA were localized on the gel by autoradiography, eluted, cleaved with 10% piperidine, dried twice, and analyzed by autoradiography of the samples after electrophoresis in 12% polyacrylamide gels containing 7 M urea. Densitometric analyses of the autoradiographs were performed with an LKB Ultrascan laser densitometer.

**Cell culture, nuclear extracts, transfections, and CAT assays**

C2 myogenic cells [Yaffe and Saxel 1977a,b] were grown in Dulbecco's modified Eagle medium [DMEM] enriched with 20% fetal calf serum [FCS], with 2 mM Glutamine and differentiation was induced culturing confluent myoblasts in DMEM with 2% horse serum. Both HeLa and CV-1 cell lines were grown in DMEM with 10% FCS, 2 mM glutamine. Drosophila Schneider cells [Schneider 1972] were kindly provided by Deborah Johnson [University of Southern California] and were maintained at 25°C in Schneider medium [GIBCO] supplemented with 10% FCS in 25-ml flasks and transferred to
60-mm plates 12 hr before transfection. HeLa cell nuclear extracts were made according to the method of Dignam et al. [1983]. C2 cells were harvested 5 days after induction of myogenic differentiation with DMEM containing 2% horse serum, and nuclear extracts were made according to Dignam et al. [1983] with additional protease inhibitors. Leupeptin (2 μg/ml), and aprotinin (2 μg/ml) were added to buffers A, C, and D. Sucrose [10%] was included in buffer C. The extracts were aliquoted and stored at -80°C in buffer D.

DNA transfections by the calcium phosphate precipitation method were performed as described previously [Minty et al. 1986]. The reporter constructs [HCA-CAT series] were transfected at concentrations of 5 μg per dish, the LTR-MyoD1 expression vector at concentrations varying from 1 to 5 μg and the pPAcSpl expression vector at 1 μg. The final DNA concentration was brought to 20 μg per dish using pUC118 plasmid DNA. Transfected C2 cells were incubated for 24 hr at 37°C, 10% CO₂ in 20% FCS, 2 mM glutamine. DNA was then removed and the cells switched to DMEM with 2% horse serum, 2 mM glutamine to induce differentiation. HeLa and CV-1 cells were transfected for 12 hr at 37°C, 10% CO₂ in 10% FCS, 2 mM glutamine before removing the DNA and cultured for an additional 24 hr. Drosophila Schneider cells were transfected by use of the same procedure employed for the mammalian cells except that they were kept at 25°C and left in the presence of the DNA for 48 hr. The transfection experiments were repeated at least twice with two independent plasmid preparations. Cells were harvested and CAT assays were performed as described by Gorman et al. [1982]. CAT assays were quantitated by excision of spots and counted in a liquid scintillation counter. Results are expressed as percent activity relative to that of the wild-type promoter CAT construct, pHCA177CAT. The variability between independently performed CAT assays was <15% and usually <5%.

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