Combinatorial Control of Smooth Muscle–Specific Gene Expression

Meena S. Kumar, Gary K. Owens

Abstract—Alterations in the differentiated state of vascular smooth muscle cells (SMCs) are known to play a key role in vascular diseases, yet the mechanisms controlling SMC differentiation are still poorly understood. In this review, we discuss our present knowledge of control of SMC differentiation at the transcriptional level, pointing out some common themes, important paradigms, and unresolved issues in SMC-specific gene regulation. We focus primarily on the serum response factor–CArG box–dependent pathway, because it has been shown to play a critical role in regulation of multiple SMC marker genes. However, we also highlight several other important regulatory elements, such as a transforming growth factor β control element, E-boxes, and MCAT motifs. We present evidence in support of the notion that SMC-specific gene regulation is not controlled by a few SMC-specific transcription factors but rather by complex combinatorial interactions between multiple general and tissue-specific proteins. Finally, we discuss the implications of chromatin remodeling on SMC differentiation.

Key Words: smooth muscle cells • differentiation • transcriptional regulation • serum response factor • CArG box

Abnormal growth and proliferation of vascular smooth muscle cells (SMCs) is a key feature of vascular diseases such as atherosclerosis, restenosis, and hypertension. Differentiated SMCs of mature animals express a unique repertoire of contractile proteins, ion channels, and signaling molecules that are necessary for their principle function of contraction. However, in response to various injurious stimuli, SMCs can dedifferentiate, converting from a quiescent, contractile phenotype to a highly proliferative, synthetic phenotype. Despite the importance of changes in the differentiated state of SMCs in vascular disease, the molecular mechanisms controlling SMC differentiation are still largely unknown. An understanding of the normal regulation of SMC development and differentiation will not only provide the foundation for elucidating how these processes may be disrupted in vascular disease but will also be critical to understanding congenital defects in vascular development and vascular development within solid tumors.

Control of cellular differentiation is ultimately regulated at the level of gene transcription, although cell-specific alternative splicing and posttranslational modifications also play important roles. This review will focus on our present understanding of the transcriptional pathways that normally control SMC differentiation as well as discuss the important, but still poorly understood, role of chromatin modifications on SMC-specific gene expression. But before reviewing transcriptional control in SMCs, we will first briefly discuss the developmental origins of SMCs, SMC differentiation/dedifferentiation markers, and several recently described in vitro SMC differentiation models.
Developmental Origin of SMCs

One of the challenges to studying mechanisms of SMC development and differentiation is the fact that unlike cardiac and skeletal muscle (which are derived from specific populations of mesodermal precursors), SMCs are derived from multiple and, in some cases, unknown cell types. For example, SMCs of the ascending aorta and branchial arches are derived from neural crest (ectodermal) cells. SMCs of the coronary vessels arise from epithelial-to-mesenchymal transition of cells in the proepicardial organ. Most of the remaining SMCs, including visceral SMCs, are thought to be derived from condensation of local mesenchyme under the influence of local environmental cues.\(^{3,5,6}\) The diverse origins of SMCs likely contribute to their heterogeneity in structure, function, and SMC marker gene expression. Moreover, the signals and molecular pathways leading to SMC differentiation may prove to be somewhat different depending on the cell type of origin.

Markers of SMC Differentiation and Dedifferentiation

To study cell-specific transcriptional regulation, it is necessary to first identify a set of cell-specific target genes and then determine how these genes may be coordinately regulated at the transcriptional level. This too has proven challenging in the SMC field, because most (if not all) SMC markers, although highly selective for SMCs in adult animals, are expressed, at least transiently, in other cell types. These include the following SM isoforms of contractile apparatus proteins: SM a-actin, SM myosin heavy chain (SM MHC), calponin, SM 22a, h-caldesmon, and smoothelin. A detailed description of these proteins and other potentially useful SMC differentiation markers and their expression patterns has been reviewed elsewhere.\(^{3,5,6}\) Of these factors, the most SMC-restrictive protein seems to be SM MHC.\(^{3}\) Although we cannot rule out that SM MHC may be expressed, at least transiently or under certain circumstances, in cells other than SMCs in vivo, to our knowledge, conclusive evidence of SM MHC expression (by immunostaining with SM MHC-specific antibodies or reverse transcriptase–polymerase chain reaction followed by sequencing) in a non-SMC in vivo does not exist. In addition, two proteins with unknown function, aortic preferentially expressed gene 1 (APEG-1)\(^{7}\) and Crp2/SmLim,\(^{8,9}\) have been found to be preferentially expressed in arterial SMCs and are likely to be useful markers of this subtype of SMCs. Finally, a secreted protein, aortic carboxypeptidase-like protein (ACLP),\(^{10,11}\) whose function is also not well-defined, has been shown to be upregulated during SMC differentiation but is by no means SMC-specific. Because no individual marker is exclusive to the entire SMC lineage (with the possible exception of SM MHC), identification of SMCs and investigations into SMC-specific transcriptional regulation requires the use of multiple markers.

When SMCs undergo phenotypic modulation, as occurs in vascular injury, virtually every SMC differentiation marker (with the exception of ACLP\(^{10}\)) is downregulated.\(^{3,5,6}\) In addition, certain proteins, which may be used as markers of dedifferentiated VSMCs, are upregulated, including osteopontin (a multifunctional adhesive glycoprotein)\(^{13}\) and the embryonic/nonmuscle isoform of myosin heavy chain, SMemb/NMHC-B.\(^{14}\) Although we will not be discussing the regulation of these SMC dedifferentiation markers in this review, some of the factors that have been shown to upregulate these markers can be found in the Table. Although it is widely accepted that SMCs retain a high degree of plasticity, which allows them to modulate their phenotype when neces-

### Transcription Factors Known or Likely to be Involved in Regulation of SMC Differentiation/Dedifferentiation

<table>
<thead>
<tr>
<th>Transcription Factor</th>
<th>Target Gene(s)</th>
</tr>
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<tbody>
<tr>
<td>MADS Box Proteins</td>
<td></td>
</tr>
<tr>
<td>SRF</td>
<td>(\uparrow) numerous SMC differentiation markers(^{4,6})</td>
</tr>
<tr>
<td>MEF2B</td>
<td>(\uparrow) SM MHC(^{101})</td>
</tr>
<tr>
<td>MEF2C</td>
<td>SMC specific targets unknown(^{102})</td>
</tr>
<tr>
<td>Homeodomain Proteins</td>
<td></td>
</tr>
<tr>
<td>Phox1/Mhox</td>
<td>(\uparrow) SM a-actin(^{44})</td>
</tr>
<tr>
<td>Banx1b</td>
<td>(\uparrow) (\beta)-Tropomyosin(^{40})</td>
</tr>
<tr>
<td>Banx2b</td>
<td>SMC specific targets unknown(^{46})</td>
</tr>
<tr>
<td>Nkx3.1</td>
<td>(\uparrow) SM (\gamma)-actin(^{46})</td>
</tr>
<tr>
<td>Nkx3.2</td>
<td>(\uparrow) (\alpha_1) integrin, SM22a, and caldesmon(^{10})</td>
</tr>
<tr>
<td>Hox B7</td>
<td>(\uparrow) SM22a and calponin(^{103})</td>
</tr>
<tr>
<td>Hex</td>
<td>(\uparrow) SMemb/NMHC-B(^{104})</td>
</tr>
<tr>
<td>Gax (Mox2)</td>
<td>(\uparrow) p21; (\downarrow) proliferation(^{106})</td>
</tr>
<tr>
<td>GATA family (GATA 4/5/6)</td>
<td>(\uparrow) p21; (\downarrow) proliferation(^{106,108})</td>
</tr>
<tr>
<td>Myocardin</td>
<td>(\uparrow) numerous SMC differentiation markers(^{10,51})</td>
</tr>
<tr>
<td>Kruppel-like Zinc Finger Proteins</td>
<td></td>
</tr>
<tr>
<td>Sp1/3</td>
<td>(\uparrow) ACLP(^{12}); (\downarrow) SM MHC and SM22a(^{26,107})</td>
</tr>
<tr>
<td>GKF/KLF4</td>
<td>(\downarrow) SM22a and SM a-actin(^{105})</td>
</tr>
<tr>
<td>BTEB2/KLF5</td>
<td>(\uparrow) SMemb/NMHC-B(^{106}; (\uparrow) SM22a(^{10})</td>
</tr>
<tr>
<td>Helix-Loop-Helix (HLH) Proteins</td>
<td></td>
</tr>
<tr>
<td>USF</td>
<td>(\uparrow) osteopontin(^{111}); SM (\alpha)-actin(^{105}), APEG-1(^{174})</td>
</tr>
<tr>
<td>ChF/HRT/gridlock</td>
<td>SM specific targets unknown(^{112,114})</td>
</tr>
<tr>
<td>eHAND/HAND2</td>
<td>SM specific targets unknown(^{115})</td>
</tr>
<tr>
<td>capsulin/epicardin</td>
<td>(\uparrow) SM (\alpha)-actin(^{117})</td>
</tr>
<tr>
<td>Id2/3</td>
<td>(\downarrow) p21; (\uparrow) proliferation(^{116,118,119})</td>
</tr>
<tr>
<td>Twist</td>
<td>(\downarrow) SM22a(^{91})</td>
</tr>
<tr>
<td>Single-Stranded DNA Binding Proteins</td>
<td></td>
</tr>
<tr>
<td>Pur(\alpha), Pur(\beta), and MSY1</td>
<td>(\uparrow) SM (\alpha)-actin(^{83,84})</td>
</tr>
<tr>
<td>Others</td>
<td></td>
</tr>
<tr>
<td>Egr-1</td>
<td>(\uparrow) proliferation (numerous targets)(^{100})</td>
</tr>
<tr>
<td>TEF-1</td>
<td>(\uparrow) SM (\alpha)-actin(^{112,114})</td>
</tr>
<tr>
<td>SRRP1</td>
<td>(\uparrow) SM22a(^{47})</td>
</tr>
<tr>
<td>Mrt2δ/β</td>
<td>(\uparrow) SM (\alpha)-actin and SM22a; (\downarrow) proliferation(^{10})</td>
</tr>
<tr>
<td>AP-1</td>
<td>(\uparrow) osteopontin(^{111})</td>
</tr>
<tr>
<td>YY1</td>
<td>(\uparrow) SM22a(^{27})</td>
</tr>
<tr>
<td>p53</td>
<td>(\uparrow) SM (\alpha)-actin(^{120})</td>
</tr>
<tr>
<td>c-myb</td>
<td>(\uparrow) SM (\alpha)-actin(^{121})</td>
</tr>
<tr>
<td>CRP1/2</td>
<td>SM specific targets unknown(^{112})</td>
</tr>
<tr>
<td>NFATc1</td>
<td>(\uparrow) SM MHC(^{122})</td>
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</table>
necessary to perform diverse functions, including contraction, extracellular matrix protein synthesis, migration, and replication, an alternate (although not necessarily mutually exclusive) view is that the SMCs that make up the vessel wall are heterogeneous and that the diverse functions of SMCs are carried out by specific subpopulations with different functional capabilities as opposed to modulation of any SMC into functionally different phenotypes. For additional reading on this topic, see reports by Bochaton-Piallat and colleagues.15,16 Frid et al,17 and references therein. In addition, an emerging body of evidence now suggests that bone marrow stem cells may be a potential source of neointimal vascular SMCs in conditions such as transplant arteriopathy and atherosclerosis.18,19

**In Vitro SMC Differentiation Models**

Although cultured SMCs have proven invaluable for addressing many questions related to control of SMC-specific gene expression, inducible SMC lineage systems are likely to prove more powerful for identifying nodal regulatory factors and pathways leading to SMC differentiation. Such a system for the skeletal muscle lineage (based on treatment of 10T1/2 cells with the DNA hypomethylating agent 5-azacytidine) has led to the elucidation of critical regulatory factors and mechanisms governing skeletal muscle differentiation.20,21 However, until recently, such systems were lacking for SMCs. Fortunately, there are now several SMC-inducible systems, each with its own advantages and disadvantages.

One of the first systems, based on treatment of P19 embryonal carcinoma cells with retinoic acid (RA), was described by our laboratory in 1995.22 We recently improved on this system by generating stable lines of P19 cells carrying a SM α-actin promoter/puromycin cassette to enrich for cells that have differentiated into the SMC lineage using puromycin selection.23 One such stable line, designated as A404, behaves like a SMC precursor/progenitor line in that it shows a high propensity for differentiation into SMC-like cells with RA treatment (even before puromycin selection). As described below, we have already begun to use this system to elucidate SMC-specific chromatin modifications and their effect on transcription factor accessibility and gene expression.23 For a more extensive review on the effects of retinoids on vascular SMC differentiation and growth, see the study by Miano and Berk.24

A second system is an embryoid body model described by Drab et al in 199725 in which totipotent mouse embryonic stem (ES) cells are cultured in such a way that they form embryo-like aggregates containing numerous differentiated cell types, including SMCs. This is potentially a very powerful system, and because ES cells with various candidate genes knocked out can be used, it provides a rapid and high throughput method of screening effects of knockouts of various proteins on SMC differentiation. However, to our knowledge, such a study has not yet been performed.

A third system, described by Jain et al in 1998,26 involves culturing Monc-1 cells, pluripotent neural crest cells first generated and characterized by Rao and Anderson,27 in a medium that promotes differentiation into the SMC lineage. Using screens to identify factors upregulated in SMC differentiated versus undifferentiated Monc-1 cells, Jain and colleagues28 have identified several factors that may play important roles in SMC differentiation, including Mrf2α/β (see Table 1) and a Kruppel-like zinc finger protein (M. Jain, personal communication, 2002).

A fourth system, described by Hirschi et al in 1998,29 involves coculture of 10T1/2 cells with endothelial cells or treatment of 10T1/2 cells with transforming growth factor β (TGFβ). A limitation of this system is that, unlike the previous 3 systems in which induction of SM MHC (the most definitive marker for SMCs) has been well documented, it is controversial as to whether SM MHC is induced in the 10T1/2 system. For example, we have been unable to detect SM MHC expression in TGFβ-treated 10T1/2 cells by either reverse transcriptase–polymerase chain reaction or Western analysis (Manabe and Owens, unpublished observations, 2000). Although Hirschi et al29 attempted to document its induction in this system using a SM MHC antibody, it is unclear whether the antibody they used specifically recognizes the SM isoform of MHC. We would like to emphasize here the importance of carefully evaluating the specificity of SM MHC antibodies when interpreting previous studies or designing future studies, because we have found that at least one of the principal antibodies used in many studies cross-reacts with a nonmuscle form of MHC (Thompson and Owens, unpublished observations, 1992). This issue is discussed more fully in the study by Owens.3 Regardless, the TGFβ-10T1/2 system may still be a useful model of, at least, the early stages of SMC differentiation (ie, before the induction of SM MHC, a relatively late-stage SMC marker) and has been used to investigate mechanisms of TGFβ-dependent regulation of SMC marker genes.30,31

To the extent that these in vitro models mimic SMC differentiation in vivo, they may prove to be very powerful tools for dissecting mechanisms of SMC differentiation. However, because SMC differentiation seems to be highly dependent on the integration of numerous complex local environmental cues3 that cannot be adequately recapitulated in culture, it is imperative that findings from these systems be verified in vivo using transgenic or knockout/knockin approaches.

**Transcriptional Regulation of SMC Differentiation Markers**

Tremendous progress has been made in the last decade in identifying transcriptional regulators of SMC marker genes, most of which are summarized in the Table. Because of space limitations, we will not be able to discuss each of these factors in detail in this review. Rather, we will focus primarily on the serum response factor (SRF)-CArG–dependent pathway, because it seems to play a central and critical role in SMC differentiation, and will draw on several additional factors and regulatory elements as examples to illustrate some common themes, important paradigms, and unresolved issues in SMC-specific gene transcription.

**SRF-CArG Dependent Pathway**

SRF is a MADS (MCM1, Agamous, Deficiens, SRF) box transcription factor that was first identified and named be-
cause of its ability to confer serum inducibility to the growth-responsive gene c-fos through binding to a sequence known as the serum response element or CArG box (CC[A/T]6 GG). SRF binds CArG boxes as a dimer, with dimerization and DNA binding occurring through the MADS box domain. In addition to regulating growth-responsive genes such as c-fos, SRF binding to CArG boxes has been shown to regulate numerous muscle-specific genes. In particular, the SRF-CArG-dependent pathway has been demonstrated to play a central and critical role in regulating virtually every SMC-specific gene, with the possible exception of ACLP, whose promoter does not seem to contain a CArG box. (However, because ACLP is also expressed at high levels in nonmuscle tissues such as bone and dermis, it can be argued that ACLP is not a true SMC-specific gene.) Of major significance, transgenic mouse studies have shown that CArG box mutations in the SMC selective promoters SM MHC, SMα-actin, and SM22α inactivate the respective promoter in one or more SMC subtypes. Because growth and differentiation are generally exclusive processes, a longstanding paradox and unresolved issue in this field is how SRF, a ubiquitously expressed transcription factor, can regulate both growth-responsive and muscle-specific genes. Although SRF levels seem to be higher in cardiac and skeletal muscle and SM compared with most other tissues, this expression pattern alone cannot explain how SRF exerts its selective effects. Several mechanisms have been demonstrated to regulate SRF activity, including association with cofactors, posttranscriptional modifications of SRF, variations in SRF binding affinity among different CArG boxes, and number, position, and spacing of CArG boxes. Of course, these mechanisms are not mutually exclusive, and it is likely that a combination of these factors is important in modulating the diverse functions of SRF.

**SRF Cofactors**

The c-fos CArG box is adjacent to a binding site for a subfamily of ETS domain transcription factors known as ternary complex factors, which, as their name suggests, form a ternary complex with SRF and the CArG box to activate transcription of the c-fos gene. However, because muscle-specific CArG boxes are generally not flanked by such ternary complex factor binding sites, other factors are likely to play a role in muscle-specific gene expression. Interestingly, Grueneberg et al demonstrated that the homeodomain
protein, \textit{Phox1}, enhanced SRF binding to the \textit{c-fos} \textit{CArG} box without formation of a higher order/tertiary complex with SRF-\textit{CArG} DNA. Hautmann et al\textsuperscript{44} subsequently showed that the mouse homologue of \textit{Phox1}, \textit{Mhox}, was able to increase SRF binding to the promoter distal \textit{CArG} box, \textit{CArG} B, of the SM \textit{\alpha-actin} promoter as well as bind itself to an \textit{ATTA} element, a homeodomain binding site, \textit{5'} to \textit{CArG} B. Moreover, \textit{Mhox} expression, \textit{Mhox} binding to the \textit{ATTA} element, and enhanced SRF binding to \textit{CArG} boxes were all increased by treatment of cultured SMCs with the contractile agonist, angiotensin II (see the Figure). Additional factors, including the homeodomain proteins Barx2\textsuperscript{45} and Nkx3.1\textsuperscript{46} and the high-mobility group protein SSRP1,\textsuperscript{47} have subsequently been shown to function similarly to \textit{Mhox} in that they enhanced the DNA binding activity of SRF but did not form a detectable ternary complex with SRF-\textit{CArG} DNA. This suggests that if such a complex exists, either it is transient or unstable or other factors or posttranslational modifications are required to stabilize it. Although interaction of SRF with one or more of the above cofactors may be an important mechanism for regulating SRF activity, it is unlikely on its own to contribute to SMC-specific gene expression, because these cofactors are not SMC selective, nor is there evidence that they can differentiate between differentiation genes and growth-responsive genes (such as \textit{c-fos}).\textsuperscript{43–47}

Interestingly, the homeodomain proteins Barx1\textsuperscript{b48} and Nkx3.2\textsuperscript{49} and the zinc finger protein GATA6\textsuperscript{50} have each been shown to form a stable, detectable ternary complex with SRF-\textit{CArG} DNA. A recent elegant study from Nishida et al\textsuperscript{51} showed that the triad of SRF, Nkx3.2, and GATA6 cooperatively activated several SMC marker genes, including SM22 and caldesmon, but not \textit{c-fos}. Thus, this combination of factors provides a mechanism by which SMC-specific genes can be regulated selectively and independently of growth-responsive genes. Moreover, although each individual factor was not SMC-specific, this triad of factors was found to be coexpressed only in vascular SMCs, illustrating an important paradigm whereby SMC-specific gene transcription can be achieved in the absence of SMC-specific transcription factors through combinatorial interactions between multiple transcription factors whose presence, at the appropriate levels and stoichiometry, may be SMC-specific.

A major breakthrough in the study of SRF cofactors and cardiac/SMC differentiation came in 2001, when Wang et al\textsuperscript{52} reported the identification of a novel SAP (SAF-A/B, Acinus, Pias) domain containing cardiac and SM-specific SRF cofactor, which they named myocardin for its critical role in cardiac-specific gene expression and cardiac development. However, it is becoming increasingly clear that myocardin may also play a critical role in SMC differentiation. Myocardin has been shown to activate several SMC-specific or -selective promoters, including SM-calponin, SM MHC, SM \textit{\alpha-actin}, and SM22, while failing to activate the \textit{c-fos} promoter.\textsuperscript{50,51} Thus, like the triad of SRF/GATA6/Nkx3.2, myocardin provides a mechanism by which SRF can discriminate between muscle-specific promoters and growth-responsive promoters. In addition to physically interacting with SRF, additional evidence that myocardin acts through the SRF-\textit{CArG}-dependent pathway comes from the following findings:

1. Myocardin forms a stable, ternary complex with SRF-\textit{CArG} box DNA\textsuperscript{50,52}; (2) myocardin-mediated transactivation is \textit{CArG} box-dependent\textsuperscript{50–52}; and (3) myocardin fails to activate SRF-dependent reporters (including SM22) in SRF-null ES cells but can do so when both SRF and myocardin are cotransfected into these cells.\textsuperscript{52} Interestingly, when overexpressed in L6 skeletal myoblast cells, myocardin is sufficient to induce the expression of several endogenous smooth and cardiac muscle genes.\textsuperscript{53} Recently, two myocardin-related transcription factors, MRTF-A and MRTF-B, have been identified whose expression is much more widespread than that of myocardin.\textsuperscript{52} Nevertheless, these factors may be important in modulating the activity of SRF or SRF-dependent genes in SMCs. A key question now is what are the mechanisms by which myocardin distinguishes between SM and cardiac muscle genes in vivo. Possibilities include additional cell-specific cofactors, posttranslational modifications, or the presence of tissue-specific alternative spliced forms of myocardin. Interestingly, the mechanism of myocardin-induced activation does seem to be somewhat different between the SM22 and the cardiac ANF promoter in that mutations in the SAP domain of myocardin, although having modest effects on myocardin-induced transactivation of the SM22 promoter, profoundly diminished activation of the cardiac ANF promoter.\textsuperscript{50} It remains to be determined whether this observation extends to other smooth versus cardiac selective promoters and how the SAP domain of myocardin discriminates between these different target genes.

\textbf{Posttranscriptional Modifications of SRF}

In addition to interacting with cofactors, SRF has been shown to be regulated by several posttranscriptional mechanisms, including alternative splicing,\textsuperscript{40,53,54} phosphorylation,\textsuperscript{55,56} and regulated nuclear translocation.\textsuperscript{57,58} In fact, several alternatively spliced isoforms of SRF have been reported, including one isoform that seems to function as a dominant-negative inhibitor of SRF.\textsuperscript{53} Phosphorylation of SRF by casein kinase II\textsuperscript{53} or ribosomal S6 kinase (pp90\textsuperscript{rsk})\textsuperscript{56} has been shown to increase its DNA-binding activity.

\textbf{Variations in SRF-Binding Affinity Among Different \textit{CArG} Boxes}

Interestingly, \textit{CArG} boxes from muscle-specific promoters, in general, have a reduced binding affinity for SRF compared with \textit{CArG} boxes from growth-responsive genes such as \textit{c-fos} and \textit{egr1}.\textsuperscript{59–61} The determinants of SRF binding affinity seem to lie both within the central A/T-rich region of the \textit{CArG} box and flanking sequences.\textsuperscript{51,62} Because muscle tissues contain higher levels of SRF than most other cell types, it has been proposed that the decreased affinity of muscle \textit{CArG} boxes serves to restrict SRF binding and therefore expression of muscle-specific genes to muscle cells.\textsuperscript{60,61,63} In support of this hypothesis, Chang et al\textsuperscript{61} demonstrated that in 11.5-day-old mouse embryos, a transgene containing multimerized \textit{c-fos} \textit{CArG} boxes upstream of a minimal promoter showed widespread activity whereas a transgene containing multimerized SM22 \textit{CArG}-near boxes upstream of a minimal promoter was active primarily in muscle cells. However, they reported that at later developmental time points and postnatally, the SM22 \textit{CArG}-near box...
reporter was also active throughout the embryo, suggesting that SRF levels may only be rate limiting very early in embryonic development and that other mechanisms subsequently take over to restrict muscle gene expression. Although this study used artificial multimerized CArG box reporters, several studies have actually replaced one or more muscle CArG boxes with c-fos CArG boxes and studied the effect of this substitution in the context of the intact muscle promoter. The results of these types of studies have shown either no effect on muscle specificity, increased basal expression with reduced cell specificity, or lack of expression. Possible reasons for these discrepancies include the model system used (i.e., cell culture versus in vivo systems), the promoter context, and whether CArG flanking sequences were included in the substitution. Of particular note, many investigators failed to document whether the resultant chimeric CArG region created by the substitution truly increased SRF binding affinity as intended. For example, substitution of C-fos CArG boxes for the two SM22 CArG boxes abolished activity of the SM22 promoter in transgenic mouse embryos. It was subsequently shown that such a substitution (at least for one of the SM22 CArG boxes [CArG-near]) actually does not increase, and may even decrease, SRF binding affinity. Another consideration is that SRF binding affinity in vitro may be very different from SRF binding affinity in vivo. Nevertheless, it is intriguing that such weak SRF binding CArG boxes have been conserved throughout evolution in muscle-specific genes, and clearly additional work is needed in this area to elucidate the reasons and implications of this finding.

**Number, Position, and Spacing of CArG Boxes**

In contrast to the C-fos promoter, which contains a single CArG box, many muscle-specific promoters contain 2 or more CArG elements. Although the significance of this is unclear, it is thought that this may be one mechanism by which myosin distinguishes between muscle promoters and the c-fos promoter. Position and spacing of CArG boxes also seem to be important determinants of promoter activity. Studies using the SM α-actin promoter have revealed that the position of the 2 CArG boxes in the 5′ promoter region are not interchangeable and that changes in the spacing/phasing of these 2 CArG boxes profoundly affects the activity of this promoter in cultured SMCs.

**Other cis Elements and trans Binding Factors**

Although the SRF-CArG–dependent pathway has been shown to be critically important in the regulation of multiple SMC marker genes, CArG elements are clearly not sufficient for SMC-specific gene regulation. Strong evidence that other cis elements and trans binding factors are important comes from numerous transgenic studies showing that deletions or mutations of various SMC marker gene promoters, despite having intact CArG boxes, are not properly expressed in vivo (and Kumar et al., unpublished observations, 2002). In this section, we will highlight 3 additional cis elements, a transforming growth factor β (TGFβ) control element (TCE), E-boxes, and MCAT elements, and discuss what is known about their binding factors and their role in SMC marker gene expression. Although we will focus on the SM α-actin promoter, these elements are highly conserved in multiple SMC marker genes and are likely to contribute to coordinate regulation of SMC differentiation.

**TGFβ and the TCE**

TGFβ has been shown to increase the expression of several SMC differentiation marker genes, including SM α-actin, SM MHC, h1 calponin, SM22α, and SM γ-actin, in cultured SMCs or the 10T1/2 model. Studies on the SM α-actin promoter revealed that 3 cis elements, the 2 5′ CArG boxes and a novel TGFβ control element (TCE), were required for TGFβ inducibility (see the Figure). TGFβ enhanced SRF binding to the CArG boxes, and this effect was at least partly attributable to increased SRF expression. The TCE element, G(A/C)GT(T/G)GG(T/G)GA, is found in several SMC-selective promoters and mutation of this element in the SM22α promoter inactivated the promoter in vivo in transgenic mice. TGFβ induced the binding of a factor to the TCE element whose identity is unknown but which appears to belong to the Kruppel-like factor (KLF) family of zinc finger proteins that includes GKLF, BTEB2/KLF5, and Sp1 (See Table). Interestingly, although GKLF repressed the SM α-actin and SM22α promoters, it appears that another related KLF member, possibly KLF5, may activate transcription through the TCE, suggesting a reciprocal role for KLF factors in control of SMC differentiation through the TCE. Additional work is needed to identify the key KLF members that activate or repress SMC marker gene expression in vivo.

**E-boxes**

E-boxes (CAAnTG motifs) bind to homodimers or heterodimers of basic helix-loop-helix (bHLH) proteins and have been found in many cell-specific promoters, including the SM-specific or -selective promoters SM MHC, SM22α, Crp2/Smlim, APEG-1, and SM α-actin. The discovery in 1989 that MyoD, a bHLH protein, functions as a master regulatory gene for skeletal myogenesis, sufficient to activate the skeletal muscle differentiation program in a variety of nonmuscle cell types, led to intense interest in identifying similar master regulatory bHLH proteins in other cells. Although this search has resulted in the identification of bHLH proteins critical to the differentiation of numerous cell types, including neurons, pancreatic β-cells, hematopoietic cells, and cardiac muscle, no SMC-specific master regulatory bHLH protein has yet been identified. Although several bHLH proteins have now been found in SMCs (see Table), their role in SMC differentiation is still largely undefined.

We previously showed that 2 E-boxes in SM α-actin promoter (designated E1 and E2) functioned as positive regulatory elements in cultured SMCs and that members of the upstream stimulatory factor (USF) family of bHLH proteins could bind the E1 E-box and regulate SM α-actin transcription in an E1-dependent manner. Whether USF proteins are the relevant E1 binding factors in vivo and what the nature of the E2 binding factors is remain to be determined. Interestingly, Chen et al recently demonstrated that USF factors can bind to an E-box in exon 1 of the APEG-1 gene and transactivate the APEG-1 promoter. More work is needed to determine the role of E-boxes on the expression of...
SMC marker genes in vivo, to additionally define the nature of E-box binding proteins in SMCs, and to determine the role of the known SMC-expressed bHLH proteins (see Table) on SMC differentiation. Because myogenic bHLH proteins have been shown to interact with SRF, it is tempting to speculate that interactions between E-box bound bHLH proteins and SRF will prove to be important in the regulation of SMC marker genes (see the Figure).

MCAT Elements
The SM α-actin promoter contains 2 MCAT elements (designated MCAT-1 and MCAT-2) with the sequence AG-GAATG. MCAT elements bind the transcription enhancer factor 1 (TEF-1) family of transcription factors and have been implicated in the control of several cardiac, skeletal, and nonmuscle genes. However, relatively little is known about their role in SMC marker gene expression. Swartz et al found that not only do Purα, Purβ, and MSY1 maintain the MCAT-containing region in a single-stranded conformation and prevents binding by TEF-1, but to some extent in SMCs, have demonstrated that the MCAT-1-containing region can be recognized by the single-stranded DNA-binding proteins Purα, Purβ, and MSY1 and have provided evidence in support of at least 2 potential models of cryptic MCAT enhancer regulation.

One model is a competition model in which binding by Purα, Purβ, and MSY1 can be recognized by the single-stranded DNA-binding proteins Purα, Purβ, and MSY1 and have provided evidence in support of at least 2 potential models of cryptic MCAT enhancer regulation.

Role of Chromatin Modifications in SMC Differentiation
Up to now, we have discussed the role of interactions between transcriptional regulators and their cognate DNA-binding sites in SMC marker gene expression. However, it must be kept in mind that endogenous genes are not naked DNA sequences but rather components of a complex, higher-order, dynamically changing structure called chromatin whose building block is the nucleosome. Nucleosomes are composed of ~146 base pairs of DNA wrapped around a histone octamer containing two molecules each of histone H2A, H2B, H3, and H4. Studies from the emerging field of chromatin regulation have revealed that the N-terminal tails of histones are the target of numerous posttranslational modifications, including acetylation, methylation, and phosphorylation, and that these modifications affect numerous biological processes, including transcription. In particular, histone acetylation has been shown to be associated with transcriptional activation. Because transcription factors generally recognize 6 to 10 base pair sequences, their selectivity of action in vivo is remarkable. Although combinatorial interactions with other promoter-bound factors (as discussed in this review) are likely to restrict the activity of certain transcription factors, DNA accessibility attributable to chromatin configuration is also likely to play a key role.

Using chromatin immunoprecipitation (ChIP) assays, Swartz et al found that SRF was strongly associated with the SM MHC, but not skeletal actin, promoter in SMCs. In contrast, ChIP assays in skeletal myoblasts and myotubes revealed SRF binding to the skeletal actin, but not SM MHC, promoter in these cells. Additional studies by Manabe et al using the A404 system revealed that SRF accessibility to the SM22 promoter in differentiated versus undifferentiated A404 cells. ChIP analyses revealed a preferential association of SRF with the SM α-actin and SM MHC promoters in differentiated and undifferentiated A404 cells. Moreover, differentiation resulted in changes in local chromatin structure, with enhanced H3 or H4 acetylation associated with the SM22 promoter in differentiated A404 cells. Although these studies demonstrate that chromatin remodeling is likely to play an important role in SMC differentiation, the question remains as to what factors are responsible for these changes in chromatin structure.

Interestingly, several transcription factors that we have discussed in this review, including SRF, GATA6, and myocardin, have been implicated either directly or indirectly in chromatin remodeling functions. For example, SRF has been shown to interact with the histone acetyltransferase protein CBP. Qiu et al showed that SRF and CBP were recruited to the CArG containing region of the SM22 promoter in cultured SMCs. GATA6 was shown to interact with the related histone acetyltransferase protein p300, and this combination of factors synergistically activated the SM MHC promoter. Myocardin contains a protein motif called the SAP domain, which has been implicated in diverse aspects of chromatin remodeling. Intriguingly, SRF may also be involved in gene silencing under some circumstances, because it has been shown to interact with SMRT, a protein that recruits histone deacetylases.

Although our knowledge of the complex interplay between transcriptional regulators and chromatin modifying enzymes and their role in SMC differentiation is still clearly in its infancy, this area of research holds great promise for the future in terms of understanding and possibly even manipulating changes in the differentiation state of SMCs.
Conclusion and Perspectives

Although changes in the differentiated state of SMCs are known to play a key role in vascular diseases such as atherosclerosis and hypertension, the mechanisms controlling SMC differentiation are not fully understood. Studies of SMC development and differentiation are challenging because of the diverse, and sometimes unknown, origin of SMCs in vivo as well as the finding that many (if not all) SMC differentiation markers are not exclusive to the SMC lineage. Nevertheless, the last decade has resulted in tremendous progress and advances in our understanding of the factors controlling SMC differentiation. Although numerous environmental cues, including growth factors, cell-cell contacts, extracellular matrix components, and neuronal input, affect the differentiation state of SMCs, in this review we focused on the final common pathway of most of these influences—the transcriptional regulators and how they interact with each other and with critical cis elements in the promoters of SMC marker genes to regulate SMC differentiation.

A common theme that has emerged from these studies is that SRF interacts with multiple proteins, likely forming a macromolecular complex in vivo containing both chromatin-modifying and transcription-activating functions. Although much work has been done investigating mechanisms of SRF-mediated regulation of SMC marker genes, our understanding of control by other cis elements (such as the TCE, E-boxes, and MCAT elements) is still in its infancy. Table summarizes most of the transcription factors known or likely to play an important role in SMC differentiation or dedifferentiation. The challenge for the future is to not only add to this list of key players but to continue to draw connections between them, mapping out transcriptional regulatory networks governing SMC-specific gene expression. Hopefully the use of inducible SMC lineage systems will help make this task possible.

Equipped with an understanding of some of the basic molecular pathways that normally control SMC differentiation, the next step is to elucidate how such pathways are disrupted in vascular injury and disease. We now have evidence that decreases in SMC marker gene expression that occur after vascular injury are at least partly transcriptionally mediated. And although we are beginning to elucidate transcriptional regulatory pathways in vascular injury, including downregulation of differentiation promoting/growth inhibitory factors such as GATA-6 and Gax and upregulation of dedifferentiation/growth promoting factors such Id2 and Egr-1, our understanding of the molecular mechanisms of SMC phenotypic modulation in more complex atherosclerotic lesions is much farther behind. Hopefully, with the combination of mouse models of atherosclerosis, SMC-specific promoter constructs, and conditional inducible gene expression systems, we will soon be able to not only address these issues but also deliver therapeutic genes in a controlled and precise manner to sites of vascular injury and disease.

In summary, what makes SMCs unique is not the presence of one specific marker protein but a whole family of factors expressed at the appropriate levels and stoichiometry necessary to carry out the differentiated functions of SMCs.

Similarly, it seems that what controls SMC differentiation is likely not one unique SMC-specific transcription factor but rather a combination of ubiquitous and tissue-restricted proteins expressed at the appropriate levels and stoichiometry necessary to initiate, maintain, and, occasionally, modulate the SMC phenotype.

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Combinatorial Control of Smooth Muscle–Specific Gene Expression
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