Agricultural Sciences. In the article “Genes from mycorhiza-otic fungi as a source for improving plant resistance to fungal pathogens” by Matteo Lorito, Sheridan L. Woo, Irene Garcia Fernandez, Gabriella Colucci, Gary E. Harman, José A. Pintor-Toro, Edgardo Filippone, Simona Muccifora, Christopher B. Lawrence, Astolfo Zoina, Sadik Tuzun, and Felice Scala, which appeared in number 14, July 7, 1998, of Proc. Natl. Acad. Sci. USA (95, 7860–7865), the following correction should be noted. The third author’s name was erroneously written as Irene Garcia Fernandez. It should be Irene Garcia.

Immunology. In the article “Differential thymic selection outcomes stimulated by focal structural alteration in peptide/major histocompatibility complex ligands” by Yoseph Gendler, Mai-kun Teng, Jin-huan Liu, Torsten Witte, Ju Liu, Ki Seok Kim, Petra Kern, Hsiu-Ching Chang, Jia-huai Wang, and Ellis L. Reinherz, which appeared in number 17, August 18, 1998, of Proc. Natl. Acad. Sci. USA (95, 10061–10066), the following correction should be noted. In Fig. 1C, the bars for L4 should be striped instead of black. The figure and legend appear below.

FIG. 1. The L4 peptide induces positive selection of N15tg thy-mocytes. FTOC was performed by using N15tg/RAG-2+/−/β2m−/− (H-2b) thymic lobes in media containing 5 μg/ml human β2m with or without 10 μM of the indicated peptides. After 7 days, thymocytes were released from the lobes by pressing through a steel mesh, counted, and triple-stained with PE-conjugated anti-CD4, Red613-conjugated anti-CD8, and mAb R53 (anti-N15 β chain clonotype) plus FITC-conjugated anti-rat IgG (19). (A) The CD8 versus CD4 staining profile of total thymocytes is shown. In this representative experiment the yield of CD8+ SP cells after L4 incubation was 13.5 × 10^4 cells compared with 0.3–2.2 × 10^4 cells in FTOC incubated in the absence of exogenous peptides or in the presence of the other indicated peptides. The total thymocyte number recovered from VSV8-exposed FTOC was significantly lower (10^5 cells per lobe) than from FTOC incubated with any of the other peptides (4–6 × 10^5 cells per lobe). (B) The histograms of the N15 TCRβ chain expression on DP and CD8+ SP thymocytes derived from FTOC incubated with K1 or L4 peptides are shown. Note that the K1 histogram represents data similar to that obtained with the other peptides (except VSV8). The CD8+ SP thymocytes that mature on L4 express a higher level of the TCR than the DP thymocytes harvested from the same lobe. (C) Thymocytes selected on L4 are functionally responsive to VSV8. Thymocytes from the organ cultures described above [cultured with (+) or without (−) L4] or fresh splenocytes from N15tg/RAG-2+/− mouse were assayed for their proliferative response to irradiated EL-4 cells, in the present of rIL-2 and 10 nM VSV8 or 10 μM L4 or no peptide. After 48 h, each well was pulsed for 18 h with [3H]thymidine, harvested on filter discs, and counted. The proliferative responses for the peptides are shown. Results are mean values of triplicate samples with SD noted.
Medical Sciences. In the article “CREB binding protein is a required coactivator for Smad-dependent, transforming growth factor β transcriptional responses in endothelial cells” by James N. Topper, Maria R. DiChiara, Jonathan D. Brown, Amy J. Williams, Dean Falb, Tucker Collins, and Michael A. Gimbrone, Jr., which appeared in number 16, August 4, 1998, of Proc. Natl. Acad. Sci. USA 95, 9506–9511), the following correction should be noted. The graphics, but not the corresponding legends, of Figs. 4 and 6 have been inadvertently transposed. The corrected figures and corresponding legends are shown below.

**FIG. 4.** Smads 2 and 4 can interact with CBP in vivo. Cos-7 cells were transfected with the indicated combinations of epitope-tagged activated TBF-β type-1 receptor, Smad expression construct, or empty expression vector. The cells were then lysed and immunoprecipitated with an anti-CBP/p300 antisera. Coprecipitating Smad proteins were detected by Western blot. The upper two panels demonstrate that significant amounts of both Smad2 and Smad4 protein coimmunoprecipitate with CBP/p300 in the presence of the activated receptor. These interactions are inhibited by the simultaneous expression of either Smad6 or Smad7 but not by cotransfection of an empty expression vector (pCIneo). The bottom panels confirm comparable Smad2, Smad4, and activated receptor expression by immunoprecipitation and Western blotting with antisera against the epitopes fused to these proteins.

**FIG. 6.** 12S E1A inhibits the association of activated Smad2 and Smad4 with CBP in vivo. Cos-7 cells were transfected with expression constructs expressing epitope-tagged (HA, Flag) versions of Smad2 and Smad4 and the indicated combinations of activated TBF-β type-1 receptor and 12S E1A expression vector. The cells were subsequently lysed and subjected to immunoprecipitation with anti-CBP/p300 antisera, and levels of coimmunoprecipitating Smad proteins were determined by Western blot. The bottom panels confirm comparable Smad2, Smad4, and activated receptor expression.
CREB binding protein is a required coactivator for Smad-dependent, transforming growth factor β transcriptional responses in endothelial cells

(vascular endothelium/P300/transcription factors/signal transduction)

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Contributed by Michael A. Gimbrone, Jr., May 28, 1998

ABSTRACT The transforming growth factor-β (TGF-β) superfamily of growth factors and cytokines has been implicated in a variety of physiological and developmental processes within the cardiovascular system. Smad proteins are a recently described family of intracellular signaling proteins that transduce signals in response to TGF-β superfamily ligands. We demonstrate by both a mammalian two-hybrid and a biochemical approach that human Smad2 and Smad4, two essential Smad proteins involved in mediating TGF-β transcriptional responses in endothelial and other cell types, can functionally interact with the transcriptional coactivator CREB binding protein (CBP). This interaction is specific in that it requires ligand (TGF-β) activation and is mediated by the transcriptional activation domains of the Smad proteins. A closely related, but distinct endothelial-expressed Smad protein, Smad7, which does not activate transcription in endothelial cells, does not interact with CBP. Furthermore, Smad2,4–CBP interactions involve the COOH terminus of CBP, a region that interacts with other regulated transcription factors such as certain signal transduction and transcription proteins and nuclear receptors. Smad–CBP interactions are required for Smad-dependent TGF-β-induced transcriptional responses in endothelial cells, as evidenced by inhibition with overexpressed 12S E1A protein and reversal of this inhibition with exogenous CBP. This report demonstrates a functional interaction between Smad proteins and an essential component of the mammalian transcriptional apparatus (CBP) and extends our insight into how Smad proteins may regulate transcriptional responses in many cell types. Thus, functional Smad–coactivator interactions may be an important locus of signal integration in endothelial cells.

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Abbreviations: BAEC, bovine aortic endothelial cells; CBP, CREB binding protein; STAT, signal transduction and transcription; TGF-β, transforming growth factor β.

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and that this correlates with their ability to interact with the conserved nuclear transcriptional coactivator CREB binding protein (CBP). Furthermore, specific ligand-activated Smad–CBP interactions seem essential for Smad-mediated transcriptional effects. CBP (and its homologue P300) have been demonstrated to play essential coactivator roles for a growing number of regulated transcription factors including CREB, nuclear receptors, myogenic helix-loop–helix factors, signal transduction and transcription (STAT) proteins, and members of the Rel (NF-κB) family (26–34). Because these coactivators have been proposed as an important locus of integration for signaling pathways in EC and other cell types (26, 27), our data suggest a novel mechanism whereby signals derived from the TGF-β superfamily of cytokines modulate a variety of cellular effects in endothelial cells.

**METHODS**

**Cell Culture.** Primary bovine aortic endothelial cells (BAEC) were isolated as described (22) and cultured in low-glucose-DMEM supplemented with 10% heat-inactivated bovine calf serum, 2 mM L-glutamine, 250 units/ml penicillin G, and 250 µg/ml streptomycin. These were used at passages 3–12. Cos-7 cells were maintained in the same medium.

**Expression Constructs and Transfections.** For transient transfections, cells were seeded at 50–70% confluency and transfected by using Lipofectamine (GIBCO/BRL) for 5 h. The cells were allowed to recover overnight in media containing 0.2% serum. Cells were then incubated in the absence or presence of 5 ng/ml human TGF-β1 (Genzyme). After approximately 18 h of incubation, luciferase, and β-galactosidase activity were measured (Tropix). All results are reported as luciferase activity (RLU) normalized to cotransfected (Tropix). All results are reported as luciferase activity (RLU) normalized to cotransfected, constitutive expression constructs, e.g., cytoskeletal α-actin. c-Jun-Gal4 and Elk-Gal4 fusion constructs (pMJun, pMEik) consist of the activation domains of c-Jun (amino acids 1–223) and Elk-1 (amino acids 307–427) fused to Gal4 DB.

**Immunoprecipitation and Western Blotting.** Antibodies directed against the epitopes used in the immunoprecipitations and Western blots were obtained from commercial suppliers (Boehringer Mannheim, Santa Cruz Biotechnology). The anti-CBP/P300 antisera were obtained from Santa Cruz Biotechnology. Cell lysates were made approximately 24 h after transfection in 1% Triton X-100, 150 mM NaCl, 50 mM Tris-HCl, pH 8.0, with protease inhibitors, and immunoprecipitations were performed overnight at 4°C. Proteins were resolved on 10–12% SDS/PAGE denaturing gels, transferred to nitrocellulose by electroblotting and probed with appropriate antisera at 1/1,000 to 1/2,000 as indicated in the figures. Individual proteins were detected with a secondary antibody coupled to peroxidase and visualized with chemiluminescence (ECL).

**RESULTS**

**Smad-Dependent Transcriptional Activation Is Enhanced by CBP.** An increasing number of regulated transcription factors have been shown to interact with the coactivator CBP (35). To investigate whether transcriptional events modulated by TGF-β involve CBP as a coactivator, we tested the ability of overexpressed recombinant CBP to stimulate two TGF-β responsive promoters in cultured endothelial cells. Both the plasminogen activator inhibitor-1 promoter (P800) and the p3TP promoter constructs are responsive to TGF-β, and this response is augmented approximately 2-fold by cotransfection of a plasmid expressing CBP. An expression plasmid for the closely related coactivator P300 gave similar results, although the magnitude of induction was consistently lower than that seen with CBP (data not shown).

Recently, the conserved family of Smad proteins has been shown to transduce TGF-β superfamily transcriptional signals in species ranging from *Drosophila* to humans (6, 7, 9). To elucidate whether the Smad proteins involved in mediating TGF-β responses in endothelial cells are using CBP as a coactivator, we used a Gal4-based system to examine the ability of various Smad proteins to enhance transcription in these cells. Fusion proteins between the Gal4 DNA binding domain and either full-length or the COOH-terminal MH2 domains of Smads 2, 4, 6, and 7 were constructed and cotransfected with a luciferase reporter construct containing a minimal promoter coupled to five tandem Gal4 binding sites. As shown in Fig. 2, full-length Smad2 can
TGF-β receptor-mediated transcription in endothelial cells, CBP can function as a coactivator for Smad2, Smad4-mediated transcription.

**Smad2 and Smad4 Specifically Interact with CBP in a Ligand-Dependent Fashion in Vivo.** To determine whether Smad2 and 4 can physically interact with CBP in vivo, we used both two-hybrid and coimmunoprecipitation strategies. In the mammalian two-hybrid assay, an interaction between two proteins is detected by fusing one protein to the Gal4 DNA binding domain, fusing the second protein to the powerful transcriptional activation domain of the VP16 protein, and coexpressing both constructs in a cell type of interest. An interaction between the two test proteins results in activation of the Gal4 luciferase reporter via the VP16 activation domain. Using the Smad–Gal4 fusion proteins described above, we tested the ability of these proteins to interact in endothelial cells with all of the domain of the CBP protein fused to VP16. As shown in the two representative experiments displayed in Fig. 3, both Smad2 and Smad4 interact with only the COOH-terminal 549 amino acids of CBP in a TGF-β dependent manner. All of the Smad2 and Smad4 constructs examined (i.e., pM2, 2C, 2*P, 2C*P, 4, 4C) interacted with CBP, and these interactions were limited to the single COOH-terminal domain of CBP (data not shown). We also tested the ability of Smad1, a distinct human Smad protein closely related to Smad2 but which mediates bone morphogenic protein signaling (19), to interact with CBP. Smad1, or its carboxyl terminal MH2 domain, also interacted specifically with the COOH-terminal domain of CBP. However, this process required the presence of an activated bone morphogenic protein type-1 receptor and was not observed in the presence of an activated TGF-β receptor alone (data not shown).

To demonstrate Smad–CBP interactions in cell extracts, we performed a series of communoprecipitations. Preliminary data demonstrated the communoprecipitation of recombinant, full-length CBP (epitope tagged) with tagged Smad2 and Smad4 proteins, but not Smad7, consistent with the mammalian two-hybrid results (data not shown). In the experiment displayed in Fig. 4, we used a polyclonal antisera to CBP/P300 to communoprecipitate the endogenous molecule and determine whether Smad proteins were associated in endothelial cells. As shown in lane 1, even when Smad2 and Smad4 proteins were overexpressed as epitope-tagged species, there is essentially no detectable interaction with CBP in the absence of TGF-β receptor stimulation. In contrast, if the cells were cotransfected with a constitutively active form of the TGF-β type 1 receptor, significant amounts of both Smad2 and Smad4 proteins communoprecipitated with CBP/P300 (Fig. 4, lane 2). To confirm that these Smad–coactivator interactions were dependent on activation of the Smad pathway, we coexpressed two recently identified inhibitory, endothelial-expressed Smads, Smad6 and Smad7 (22–24).
As shown in Fig. 4, lane 3, the expression vector alone (pCIneo) had no effect, whereas both Smad6 and Smad7 completely inhibited the interactions of Smads 2 and 4 with CBP. These results thus confirm that activation of the Smad proteins is required for the CBP interactions seen and that these effects are not solely caused by overexpression of these proteins.

**Smad-Mediated Transcriptional Activation Requires Smad-CBP Interactions.** The COOH-terminal domain of CBP, which interacts with activated Smads in the mammalian two-hybrid system, is also the site at which this coactivator interacts with a number of other important regulatory proteins (29, 31, 37, 38). One of these, the viral protein E1A, is thought to sequester CBP and thus modulate the transcriptional effects of many effector proteins (34, 39, 40). We used expression of 12S E1A to test whether CBP was required for Smad-mediated transcriptional events. Preliminary experiments demonstrated potent inhibition of the TGF-β responsive promoters P800 and p3TP by 12S E1A (data not shown), so we examined the effect of 12S E1A expression in the Smad–Gal4 system. As shown in Table 1, 12S E1A can almost completely inhibit the stimulated activity of all of the Smads and Smad constructs in endothelial cells. As a control for nonspecific effects, we also expressed a mutant form of E1A (mutE1A). This molecule harbors an NH2-terminal deletion that renders it unable to interact significantly with CBP or P300. As shown in Table 1, this protein did not inhibit Smad-dependent transcription in this system. As a control for the assay, we also looked at the effects of these E1A proteins on transcription mediated by the activation domains of c-Jun and Elk-1 fused to Gal4 (c-Jun–Gal4, Elk–Gal4), as both of these transcriptional activators have been demonstrated previously to interact functionally with CBP. The results for these constructs were virtually identical to those of the Smad constructs (Table 1), in that they are inhibited by 12S E1A but not by the mutant protein.

E1A can interact with a variety of cellular proteins. To ascertain if the inhibitory effect on Smad-mediated transcription observed was caused by sequestration of limiting amounts of CBP within the cell, we attempted to rescue the E1A-mediated inhibition by titrating in increasing amounts of CBP expression plasmid. As shown in Fig. 5, for both Smad2 and Smad4, as well as the c-Jun activation domain, we could effectively restore transcriptional activation in the presence of 12S E1A by increasing amounts of CBP expression. In fact, all of the Smad constructs inhibited by 12S E1A expression were effectively rescued by coexpression of CBP (data not shown). To determine whether these results correlated with the biochemical association of Smads and CBP, we performed immunoprecipitations in the presence and absence of 12S E1A expression. Fig. 6 is a coimmunoprecipitation experiment demonstrating that in the presence of 12S E1A, the TGF-β stimulated association of Smads 2 and 4 with the coactivator CBP was effectively inhibited, whereas the association of Smads 2 and 4 with the mutant E1A form was not. As a control, we also examined the association of Smads 2 and 4 with the viral protein E1A, which is unable to bind to CBP. These results confirm that Smad-mediated transcription is inhibited by the simultaneous expression of either Smad6 or Smad7 but not by cotransfection of an empty expression vector (pCIneo). The bottom panels confirm comparable Smad2, Smad4, and activated receptor expression by immunoprecipitation and Western blotting with antisera against the epitopes fused to these proteins.

**Table 1.** 12S E1A, but not an amino-terminal truncated mutant of E1A, can inhibit Smad-dependent transcription in BAEC

<table>
<thead>
<tr>
<th>Vector</th>
<th>12S E1A</th>
<th>mut-E1A</th>
</tr>
</thead>
<tbody>
<tr>
<td>pMSmad2</td>
<td>100</td>
<td>8 ± 1.2</td>
</tr>
<tr>
<td>pMSmad2C</td>
<td>100</td>
<td>10 ± 2.1</td>
</tr>
<tr>
<td>pMSmadC*P</td>
<td>100</td>
<td>3 ± 1.1</td>
</tr>
<tr>
<td>pMSmad4C</td>
<td>100</td>
<td>3 ± 0.4</td>
</tr>
<tr>
<td>pMJun</td>
<td>100</td>
<td>7 ± 2.3</td>
</tr>
<tr>
<td>pMElk</td>
<td>100</td>
<td>2 ± 1.0</td>
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</tbody>
</table>

The left side of the table lists the various Smad–Gal4 constructs that were assayed for their ability to activate the Gal4 reporter. The first column represents the level of transcriptional activation seen with maximal stimulation (TGF-β) normalized to 100%. 12S E1A markedly inhibits the transcriptional activity of all of the Smad constructs tested. The mutant form of E1A, which is unable to bind to CBP, does not inhibit the Smad-dependent transcription. Gal4–Jun and Gal4–Elk were also assayed as positive controls for proteins known to require CBP as a coactivator.
FIG. 5. Overexpression of CBP can rescue 12S E1A-mediated inhibition of Smad transcriptional activation. pM2, pM4C, and pMJun were cotransfected into BAEC with 12S E1A and increasing amounts of CBP. Total DNA was kept constant by the addition of empty vector. In the presence of increasing amounts of CBP expression vector, the inhibition observed in the presence of 12S E1A is reversed.

CBP is absent (compare lanes 3 and 4). Taken together, these results indicate that CBP–Smad interactions seem to be required for Smad-mediated transcriptional activation to occur in endothelial cells.

DISCUSSION

The identification of Smad proteins as critical intracellular mediators of TGF-β superfamily transcriptional responses has been a major advance in understanding how this important family of cytokines and/or growth factors elicit their protean biologic effects (4–6). In this paper, we demonstrate that Smad2 and Smad4, two mediators of TGF-β1 signaling in endothelial and other cells, can act as ligand (TGF-β)–activated transcriptional activators in endothelial cells, whereas two distinct, recently identified endothelial-expressed Smads, Smad6 and Smad7, do not. Furthermore, transcriptional activation by Smad proteins in endothelial cells requires a ligand-induced interaction with the transcriptional coactivator CBP.

The molecular mechanisms by which certain Smad proteins can function as transcriptional activators and/or coactivators are just now being elucidated. Work in the Xenopus system has identified a specific cis-acting activin response element that interacts with a complex consisting of two distinct Xenopus Smad proteins and a forkhead-containing DNA binding protein known as Fast-1 (41). Alone, the individual components of this complex are thought to be unable to activate transcription efficiently, until the activin pathway is activated by receptor–ligand interactions and the active complex is formed. Xenopus Fast-1 can interact with certain mammalian Smad proteins and activate the activin response element when introduced into mammalian cells, suggesting that human homologues of Fast-1 exist (23, 41). In contrast to these Xenopus Smad proteins, a Smad protein from Drosophila has been reported to bind DNA directly, and in vitro, human Smad3 and Smad4, but not Smad2, can interact directly with specific DNA sequences (42, 43). In the studies reported here, we have demonstrated that Smad2 and Smad4, two Smad proteins that can synergize as activators of transcription in response to TGF-β, selectively interact with the COOH-terminal domain of the conserved mammalian transcriptional coactivator CBP. This interaction was demonstrated by both a two-hybrid approach in endothelial cells as well as a biochemical approach using antisera against endogenous CBP. In addition, this interaction seems specific, in that only certain Smad proteins (or truncated forms of these proteins) that contain potential transcriptional activation domains, display any detectable interaction. A closely related endothelial-expressed Smad, Smad7, which does not possess transcriptional activation activity, did not demonstrate any detectable interaction with CBP. Furthermore, both Smad2 and Smad4 interact with CBP in a ligand-dependent manner. In the absence of receptor (TGF-β) stimulation, or in the presence of the inhibitory Smads 6 and 7, no significant interaction with CBP is observed. These results argue strongly that the interactions observed are specifically a result of activation of the TGF-β signaling pathway and are not merely a result of overexpression of recombinant Smad proteins.

The interactions that we have observed between Smads and CBP closely resemble the interactions previously described between nuclear receptors such as the retinoic acid receptor and CBP (31, 38, 44, 45). These nuclear receptors also require ligand activation to interact with CBP and have been demonstrated to bind the same COOH-terminal domain of CBP as the Smads. This domain is adjacent to, but physically distinct from, the C/H3 domain of CBP that interacts with 12S E1A and STAT-1α. It is interesting to note that several members of the nuclear receptor superfamily are capable of mediating both negative and positive transcriptional regulation. It will be interesting to see whether Smad-mediated transcriptional regulation demonstrates similar complexity.

In addition to demonstrating selectivity, Smad–CBP interactions seem to be critical for TGF-β transcriptional responses. In the presence of 12S E1A, a viral molecule that is capable of binding and sequestering limiting amounts of CBP present within the cell, Smad-dependent, TGF-β transcriptional responses were completely inhibited. This response was not seen with a mutant form of E1A that does not interact with CBP,
and more important, the 12S E1A-mediated inhibition could be reversed by the addition of exogenous CBP. Furthermore, in the presence of 12S E1A, the biochemical association of Smad2 and Smad4 with endogenous CBP was blocked. These results indicate that 12S E1A can inhibit Smad function by interfering with the assembly of Smad–coactivator complexes and that interactions between CBP and activated Smads are required for Smad-dependent transcriptional responses.

Transcriptional regulation requires at least three general classes of proteins: (i) proteins that recognize specific cis-acting DNA motifs, (ii) proteins that are recruited to promoters by protein–protein interactions and act as coactivators or corepressors, and (iii) proteins that alter the structure of chromatin. CBP, which possesses histone acetyltransferase activity (46), has been demonstrated to interact directly with elements of the basal transcriptional apparatus such as TFIIB and RNA polymerase II, as well as several classes of sequence-specific transcription factors including c-Jun, c-Fos, Sap-1a, Elk-1, STAT-1α, CREB, NF-κB, and members of the nuclear receptor superfamily (26, 27, 29, 31–33, 37, 44, 47, 48). In this capacity, CBP is thought to function as a critical bridge between these sequence-specific factors and the basal transcriptional machinery. Moreover, competition for limiting pools of intracellular CBP is thought to mediate the functional interactions between diverse signaling pathways such as the Ras/AP-1, Jak/STAT, and nuclear receptor-mediated pathways (27, 45, 48). Thus, CBP may function as an important locus of signal integration for diverse signals within the cell.

Our results suggest that Smad proteins such as Smad2 and Smad4 may function in a manner analogous to many of the above-mentioned signal-dependent transcription factors. Namely, either alone or in combination with yet-to-be defined sequence-specific DNA binding molecules (e.g., Fast-1 like factors), these proteins interact with specific cis-acting DNA regulatory elements and modulate transcription by interacting in a specific manner with CBP to facilitate formation of productive transcription initiation complexes. Such a model would predict that there may be complex functional interactions between Smad proteins and other regulated transcription factors that utilize these coactivators.

In summary, we have demonstrated that Smad2 and Smad4 may function in a manner analogous to many of the above-mentioned signal-dependent transcription factors. Namely, either alone or in combination with yet-to-be defined sequence-specific DNA binding molecules (e.g., Fast-1 like factors), these proteins interact with specific cis-acting DNA regulatory elements and modulate transcription by interacting in a specific manner with CBP to facilitate formation of productive transcription initiation complexes. Such a model would predict that there may be complex functional interactions between Smad proteins and other regulated transcription factors that utilize these coactivators. Indeed, it has recently been independently reported that Smad-dependent activation of a TGF-β dependent promoter required a series of TRE or AP-1 sites (49) and that a dominant negative form of Smad3 could inhibit transcriptional responses elicited by diverse stimuli. In addition, we have recently demonstrated that overexpression of Smad proteins can modulate endothelial gene expression in response to fluid mechanical (flow) stimuli (22). These complex interactions between potentially disparate stimuli, the transcription factors they regulate, and Smad proteins likely involve shared interactions with common coactivators such as CBP.

In summary, we have demonstrated that Smad2 and Smad4, two proteins involved in mediating TGF-β transcriptional responses in endothelial and other cell types, can functionally interact with the conserved transcriptional coactivator CBP. Specific Smad–CBP interactions seem to be essential for Smad-mediated transcriptional activation in endothelial cells. This demonstrates a functional interaction between this newly identified class of signaling molecules (Smads) and an essential component of the mammalian transcriptional apparatus (CBP) and extends our insight into how Smad proteins may mediate transcriptional responses in cells. Furthermore, our observations suggest that functional integration at the level of required trans-criptional coactivators such as CBP/P300 may play an important role in the many biologic effects characteristic of the TGF-β superfamily of cytokines and growth factors.

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