Conserved motifs in Fos and Jun define a new class of activation domain

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Fos and Jun form a tight heterodimeric complex that activates transcription by AP1 sites. We have recognized that two adjacent regions of the Jun A1 activation domain are conserved in the Fos protein, and we refer to these two homologous regions as homology box 1 (HOB1) and homology box 2 (HOB2). Using GAIA chimeras, we show that the HOB1/HOB2 region of Fos and Jun is an independent activation domain in which HOB1 and HOB2 act cooperatively to activate transcription. This cooperativity is retained after the replacement of Fos HOB1 or HOB2 with the equivalent domain of Jun or when duplicated HOB1/HOB1 and HOB2/HOB2 combinations are generated. In the Fos protein, HOB1 or HOB2 can also cooperate with a distinct domain at the carboxyl terminus of the protein. Using the HOB2 consensus sequence as a guide, we identified a HOB2-containing activation domain in the CCAAT/enhancer binding protein (C/EBP) protein. This HOB2 motif can cooperate with as yet undefined sequences in C/EBP and will function even when linked to Jun HOB1. Thus, HOB1 and HOB2 represent inert “cooperating modules” that are combined to generate a functional activation domain. Each of these modules has the potential to cooperate with both distinct and identical domains. The presence of HOB-like modules in three different transcription factors indicates that the HOB motifs characterize a new class of activation domain. These motifs can be used now to identify other transcription factors with such modular characteristics.

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The transcription factors Fos and Jun can regulate the expression of genes bearing AP1 sites. These two proteins share a common basic leucine zipper (bZIP) DNA-binding domain [Landschulz et al. 1988a] that allows them to dimerize by the leucine zipper and bind DNA by a basic motif [Halazonetis et al. 1988; Kouzarides and Ziff 1988; Nakabeppu et al. 1988; Sassone-Corsi et al. 1988; Gentz et al. 1989; Schuermann et al. 1989, Turner and Tjian 1989]. Each of these two proteins belongs to a different family of either Fos- or Jun-related proteins. Members of each family share a number of conserved domains, but the only recognized homology between the Fos family (Fos, Fra-1, Fra-2, Fos-B) and the Jun family (Jun, Jun-B, Jun-D) is within their DNA-binding domain. The leucine zipper structure within this domain provides the specificity of interaction between Fos and Jun family members [Kouzarides and Ziff 1989a; Sellers and Struhl 1989]. All Fos-related proteins can form functional, DNA-binding heterodimers with any of the Jun-related proteins, suggesting that a variety of Fos- and Jun-related heterodimers, with potentially different effector domains, may be present in the cell [for review, see Kouzarides and Ziff 1989b]. The Jun family members can also form homodimers and heterodimers with each other, whereas the Fos family members can only heterodimerize with members of the Jun family. The role of the Fos- and Jun-related proteins and how this role differs from that of Fos and Jun is unclear, but some clues do exist. For example, unlike Jun, Jun-B is incapable of activating the transcription of a promoter bearing a single AP1 site and can inhibit the activation of such a promoter by the Jun protein [Chiu et al. 1989; Schütte et al. 1989].

Once bound to DNA, the Fos–Jun complex can regulate gene expression by virtue of specific activation domains carried by each protein [Angel et al. 1989; Abate et al. 1990]. Structure–function analysis of the Jun protein has identified several transcriptional activation domains in assays both in vitro [Bohmann and Tjian 1989] and in vivo [Angel et al. 1989; Baichwal and Tjian 1990]. A major determinant appears to be at the amino terminus of the protein [Angel et al. 1989; Baichwal and Tjian 1990], whereas a relatively weak activation domain is present toward its carboxyl terminus [Baichwal and Tjian 1990]. Various lines of evidence suggest that the ability of the amino-terminal domain to activate is regulated both positively and negatively. Evidence for negative regulation comes from in vivo competition experiments that show that a cell type-specific inhibitor interacts with this domain and represses its activity [Baichwal and Tjian 1990]. The target for this down-regulation appears to be [at least partly] a 27-residue domain, called the $\delta$ region,
which is found to be deleted in the v-jun oncogene. Positive regulation of this activation domain is mediated by certain activated oncogene products. Cotransfection of H-ras or v-src expression vectors stimulates the activation potential of this Jun domain [Baichwal et al. 1991; Binétruy et al. 1991]. The stimulatory effect of H-ras may be a direct consequence of Jun phosphorylation, as two serine residues within the Jun activation domain are specifically phosphorylated in response to H-ras [Binétruy et al. 1991; Smeal et al. 1991].

Here we show that Fos and Jun have previously unrecognized homology outside their bZIP DNA-binding domain. This homology represents the conservation of two adjacent regions, homology box 1 (HOB1) and homology box 2 (HOB2), that fall within their activation domain. Using GAL4 DNA-binding domain chimeras, we show that HOB1 and HOB2 function cooperatively to activate transcription. In addition, they each cooperate with a distinct domain in the Fos carboxyl terminus and can function even when repeated in a HOB1/HOB1 or HOB2/HOB2 combination. Our results establish the existence of inert "cooperating modules" that activate transcription when combined with a second [identical or distinct] module.

**Results**

**Fos shows homology to a Jun activation domain**

Given that Fos and Jun cooperate to activate transcription, we reasoned that Fos and Jun may possess common trans-activating domains, in addition to a common DNA-binding domain. Because comparison of the primary amino acid sequence did not reveal any obvious homology outside their DNA-binding domain, we compared the secondary structure predictions of the two proteins. This type of analysis focused our attention on two adjacent, potentially helical regions of the Fos protein [Figs. 1 and 2]. We refer to these two homologous regions as HOB1 and HOB2. In the Jun protein, HOB1 and HOB2 are present near the amino terminus, within the A1 activation domain [Baichwal and Tjian 1990]. Deletion analysis of Jun has shown that domains covering precisely the HOB1 and HOB2 sequences are essential for Jun-induced transcriptional activation [Angel et al. 1989; Fig. 1]. In Fos, HOB1 and HOB2 are present after the leucine zipper region, near the carboxyl terminus of the protein [Fig. 1].

**Transcriptional activation domains of Fos**

We set out to establish whether the HOB1/HOB2 region of Fos possesses transcriptional activation functions, as in the case of the HOB1/HOB2 region of Jun. Because Fos is unable to bind DNA on its own, we fused the Fos protein to the DNA-binding domain of the yeast transcription factor GAL4 and used this chimera (GF 1-380) to activate transcription of a promoter-bearing GAL4-binding site [Fig. 3]. Deletion analysis of GF 1-380 indicates that a region containing both HOB1 and HOB2 (amino acids 210–308) is a major contributor to Fos transcriptional activity [cf. lanes 2 and 5] and that sequences at the amino terminus of the protein also possess transcriptional activation functions [cf. lanes 1 and 3]. The HOB1/HOB2 region (amino acids 210–308) appears to have independent trans-activating ability in the absence of the rest of the Fos protein, as it can activate transcription very efficiently when fused to the GAL4 DNA-binding domain [lane 6]. When a fragment containing only

**Figure 1.** The HOB1 and HOB2 motifs of Fos lie within a trans-activating domain. Diagrammatic representation of the Fos and Jun proteins shows the position and alignment of HOB1 and HOB2. The previously described in vivo activation regions of Jun are shown below the Jun protein. Regions I, II, and III were described by Angel et al. (1989) and regions A1 and A2 were reported by Baichwal and Tjian (1990). The HOB1 and HOB2 motifs of Jun lie within trans-activating regions II and III, respectively. The position of the Fos and Jun basic DNA-binding motif (B) and leucine zipper dimerization motif (Z) is indicated.
A HOB1                 HOB2

| JUN     | S P D V G L K L A E R L V I S S N G E I T C E R | T P Q F L L P R N V T D E G E D F A R G F V A L A K |
| FOS     | D L T G L P R A T P D E E A F T L L W D P E S L E P V E N I S K E L K A R D C D L P F A S S R |
| 65      | 75     | 85     | 95     | 105    | 115    |
| FOS mutants: |       |        |        |        |        |
| GF 210-308 EE |   | R     | R     |       |        |
| GF 210-308 FD |   |       |       | A     | A      |
| GF 210-308 EE/FD | A     |       |       | A     | A      |
| GF 210-308 TS | A     | A     |       |
| GF 210-308 NP |       |       | A     |       |

Figure 2. Residues within HOB1 and HOB2 contribute to Fos-induced transcriptional activation. (A) Alignment of the rat c-Fos (Curran et al. 1987) and the human c-Jun (Angel et al. 1988) regions that span HOB1 and HOB2, showing the precise conservation of distance between the HOB1 and HOB2 in Fos and Jun. Below the alignment, the mutations introduced within the Fos sequence are shown. (B) The GF 210-308 plasmid (wt) and the various mutants of this [shown in A] were introduced into 1 BR cells along with the pUAS10CAT reporter plasmid as described in Fig. 3. CAT activity of the GF 210-308 plasmid (wt) was arbitrarily set at 100, and the values for the various mutants are shown relative to this.

HOB1 (GF 210-244, lane 7) or only HOB2 (GF 250-308, lane 8) is used, however, no activation is detected. This result is consistent with the notion that HOB1 and HOB2 are interdependent for transcriptional activity.

Our deletion analysis also suggests the presence of an additional trans-activating domain carboxy-terminal to HOB2. A chimera that contains the HOB1/HOB2 region plus the remaining Fos carboxyl terminus (GF 210-380, lane 9) shows higher activity than GF 210-308, which contains the HOB1/HOB2 region alone. This carboxy-terminal domain is unable to function in the context of a larger protein (cf. lanes 2 and 4), however, probably owing to the presence of a repressor function elsewhere in the protein. A repressor function must also affect the activity of the HOB1/HOB2 region as this domain shows higher activity when amino-terminal sequences are removed (cf. lanes 4 and 6). The presence of repressive domains within the Fos protein is consistent with recent in vitro results from Abate et al. [1991]. These studies also show that the region covering the Fos HOB1/HOB2 domain can function in vitro in the context of the full-length Fos-Jun complex.

Residues within Fos HOB1 and HOB2 contribute to trans-activation

Having shown that a 99-amino-acid region [210–308] encompassing the Fos HOB1 and HOB2 motifs can function as an independent trans-activating domain, we sought to verify that residues within HOB1 and HOB2 contribute to the activity exhibited by this region. First, we chose to mutate two glutamic acid residues [E-234 and E-236] in Fos HOB1 (Fig. 2A). Substitution of the corresponding residues [E-75 and E-77] in Jun to arginine (R) results in a reduction of Jun trans-activating ability (Angel et al. 1989). Figure 2B shows that mutating Fos E-234 and E-236 (EE) to arginine reduces trans-activation by over sevenfold. In contrast, mutating two residues within the nonconserved region between HOB1 and HOB2 (NP) has no effect on trans-activation. This supports the argument that homologous residues within the Fos and Jun HOB1 motif are functionally related. Inspection of the HOB1 motif suggested to us that these two E residues may form part of a recognition site for a protein kinase, as an S/T residue [73 in Jun and 232 in Fos] is conserved in the vicinity. We therefore mutated Fos T-232 and an additional serine residue, S-235, to alanine and found that this double mutation, TS, also affects trans-activation. This result is consistent with the possibility that a phosphorylation event may regulate the activity of this region.

To assess the contribution of the HOB2 motif, we mutated two conserved residues, F-269 and D-271, to alanine. As seen in Figure 2B, the FD mutant is severely defective in trans-activating ability compared with the wild-type GF 210-308 (20-fold lower activity). This
HOB motifs of Fos and Jun

Figure 3. Identification of transcriptional activation domains within the Fos protein. (A) Schematic representation of the Fos deletion analysis. Deletions of the Fos protein are fused to the GAL4 DNA-binding domain (DBD). All constructs are in an SV40 early promoter expression vector. The activity of each fusion is shown as an increase in CAT activity above the level induced by the GAL4 DBD alone. Values represent an average of several independent transfections. The HOB I and HOB2 motifs (black and stippled boxes) as well as the basic (B) and zipper (Z) regions are shown. The GAL4 DBD is shown as a box with diagonal lines. (B) Activity of the GAL4-Fos deletions. Four micrograms of the target reporter plasmid pUAS10CAT (containing GAL4-binding sites; Cousens et al. 1989) was cotransfected with 1 gg of the SV40 promoter-expressed GAL4-Fos chimeras (shown in A) into 1 BR cells. Extracts from these cells were used for CAT assays and Western blots. The Western blots were probed with an antibody against the GAL4 DBD to verify that each plasmid expressed equivalent levels of protein.

HOB2 mutant shows less activity than the HOB1 mutant EE, whereas the double mutant EE/FD shows marginally less activity than either. These results verify that both HOB1 and HOB2 contribute to the transcriptional activity exhibited by this region.

Fos and Jun have functionally interchangeable HOB1 and HOB2 domains

Having established that Fos HOB1 and HOB2 only function in combination [Fig. 3], we asked whether either of these motifs could be replaced by the corresponding HOB motif of Jun. Figure 4 shows that a domain of Jun that contains both HOB1 and HOB2 (lane 4) can activate transcription when linked to the GAL4 DNA-binding domain, whereas a region containing HOB1 (lane 5) or HOB2 (lane 6) alone is essentially inactive. This parallels the results obtained with Fos, where HOB1 and HOB2 together activate transcription (lane 1), whereas HOB1 or HOB2 alone do not (lanes 2,3). If we now make a chimera containing Fos HOB1 linked to Jun HOB2 (GFH1/JH2, lane 8) we can restore activity. The cooperativity between these Fos and Jun domains is dependent on an intact HOB motif, as mutagenesis of Fos HOB2 (EE) or Fos HOB1 (FD) severely reduces this cooperativity. These data suggest that the HOB1/HOB2 motifs of Fos and Jun are operationally equivalent and provide strong evidence that the HOB1 and HOB2 domains activate transcription cooperatively.

Fos HOB1 or HOB2 can cooperate with carboxy-terminal sequences

Our analysis of the HOB1/HOB2 domain of Fos [see Fig. 3] revealed that an additional activation domain may be present within the carboxyl terminus of the Fos protein [Fig. 5, cf. lanes 1 and 2]. Having established that HOB1 and HOB2 are cooperating units, we asked whether this additional carboxy-terminal activity was reliant on the presence of HOB1 or HOB2 or both. Figure 5 shows that the carboxyl terminus of Fos [sequences precisely carboxy-terminal to HOB2, GF 276-380, lane 6] has no independent trans-activating ability. When the inactive...
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Figure 4. The HOB1 or HOB2 domains of Jun can be functionally replaced by the equivalent HOB domain of Fos. (A) Schematic representation of the GAL4 DBD fusions containing various combinations of Fos HOB1 (FH1), Fos HOB2 (FH2), Jun HOB1 (JH1), and Jun HOB2 (JH2). The EE and FD mutations are as shown in Fig. 2. The values on the right represent the increase in CAT activity induced by each construct, above the level exhibited by the GAL4 DBD alone. (B) Activity of “domain-swapped” chimeras. The chimeric constructs shown in A expressed under an SV40 promoter were cotransfected with the pUAS10CAT reporter into 1 BR cells as described in Fig. 3.

HOB1 or HOBB2 domain is linked to the carboxyl terminus, however, very efficient trans-activation is observed (GF Δ210-380, lane 4 and GF 250-380, lane 3). This indicates that HOB1 and HOB2 can cooperate equally well with a distinct domain present at the carboxyl terminus. The presence of both HOB1 and HOB2 along with the carboxyl terminus does not increase cooperativity but shows additive behavior (GF 210-380, lane 1). Thus, the higher activity of GF 210-380 compared with GF 210-308 is due to cooperation between both HOB1 and HOB2 with the carboxy-terminal domain.

The domain cooperating with HOB2 appears to lie within the carboxy-terminal 48 residues of Fos. Deletion of these sequences (GF 250-332) severely reduces the trans-activating potential of GF 250-380. Because these 48 carboxy-terminal residues do not show significant trans-activating potential (GF 314-380, lane 8), we conclude that the very carboxyl terminus of Fos must contain an inert domain that has the capacity to activate transcription in cooperation with HOB2.

A functional HOB2 domain in C/EBP

The presence of the HOB1 and HOB2 motifs in members of two different transcription factor families prompted us to look for the presence of these motifs in the trans-activating domains of other transcription factors. The search for HOB1 was not fruitful, but the search for HOB2 homologies proved more conclusive (Fig. 6). This motif was found within the activation domain of the CCAAT/enhancer binding protein (C/EBP) transcription factor. Friedman and McKnight (1990) have shown that a 29-residue region of C/EBP (which contains the HOB2 motif) is unable to activate transcription when linked to the GAL4 DNA-binding domain but can do so very efficiently when amino-terminal sequences with no intrin-
Figure 5. Fos HOB2 cooperates with the carboxyl terminus. (A) Schematic representation of GAL4 DBD fusions containing portions of the Fos carboxyl terminus (210–380). The values on the right represent the increase in CAT activity induced by each construct, above the level exhibited by the GAL4 DBD alone. (B) CAT activity of each construct shown in A. The chimeras were expressed under an SV40 promoter and cotransfected with the pUAS10CAT reporter into 1 BR cells as described in Fig. 3.

Discussion

Sequence homology between a Jun activation domain and an undefined region of the Fos protein has led to the identification of a novel class of activation domain. In vivo assays with GAL4 DBD chimeras have allowed us to compare directly the activation potential of Fos and Jun sequences containing the homologous regions HOB1 and HOB2. The results clearly show that [1] domains containing HOB1 or HOB2 have almost no activation capacity but can work cooperatively to activate transcription; [2] the HOB motifs in Fos and Jun are functionally interchangeable; therefore, chimeras consisting of Fos HOB1/Jun HOB2 or Jun HOB1/Fos HOB2 form functional activating domains; [3] HOB1 and HOB2 can activate transcription synergistically even when present in a repeating HOB1/HOB1 or HOB2/HOB2 combination; [4] Fos HOB1 or HOB2 can cooperate with a distinct sequence at the Fos carboxyl terminus; and [5] a functional HOB2 motif, capable of cooperating with Jun HOB1, is present in the C/EBP protein.

These properties, exhibited by HOB1 and HOB2, demonstrate the existence of inert “activation modules.” Such modules can activate transcription cooperatively with a second activation module, which can be either identical to itself or of a distinct class. The use of activation modules may reflect a widely used principle for the generation of activation domains. We have identified three different transcription factors that possess such

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### Table: Schematic representation of GAL4 DBD fusions containing portions of the Fos carboxyl terminus (210–380)

| Construct | CAT Activity |
|-----------|-------------|
| GF 210-380 | 70          |
| GF 210-308 | 37          |
| GF 250-380 | 34          |
| GFA 210-380 | 38        |
| GF 250-332 | 3           |
| GF 276-380 | 3           |
| GF 276-332 | 1           |
| GF 314-380 | 2           |

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Figure 6. The HOB2 motif shows homology to a trans-activating domain of C/EBP. The homologous residues are boxed. The residues predicted to be within an α-helical conformation are marked by an x under the amino acid sequence. The phosphorylation of Ser residues (S-63 and S-73) in Jun (marked by an *) is stimulated by H-ras and is carried out in vitro by MAP kinase [Binétry et al. 1991; Pulverer et al. 1991]. Mutagenesis of these two serine residues drastically reduces Jun-mediated trans-activation [Pulverer et al. 1991]. The sequence of C/EBP is from Landschulz et al. [1988b].

Figure 7. C/EBP contains a cooperating HOB2 domain. (A) Schematic representation of the GAL4 DBD fusions containing various combinations of C/EBP HOB2 (CH2) and Jun HOB1 (JH1) domains. The activity of each chimera is shown on the right as the increase in CAT activity above the level exhibited by the GAL4 DBD alone. (B) The CAT activity of each chimera shown in A. The constructs, expressed under an SV40 promoter, were cotransfected with the pUAS10CAT reporter into 1 BR cells as described in Fig. 3.

Figure 8. HOB1 or HOB2 function when duplicated. (A) Schematic representation of GAL4 DBD fusions containing various combinations of Fos HOB1 (FH1) and Fos HOB2 (FH2). The values on the right represent the increase in CAT activity induced by each construct, above the level exhibited by the GAL4 DBD alone. (B) CAT activity of each construct shown in A. The chimeras were expressed under the SV40 promoter and cotransfected with pUAS10CAT reporter into 1 BR cells, as described in Fig. 3.
phosphorylation events within HOB1. The transcriptional activity of the Jun A1 domain is augmented by the H-ras protein. This “superactivation” process is reliant on the phosphorylation of two serine residues, S-63 and S-73, which is induced by H-ras in vivo and can be carried out by MAP kinase in vitro (Binétruy et al. 1991; Pulverer et al. 1991; Smeal et al. 1991). Interestingly, Jun S-73 falls within Jun HOB1 and is conserved in Fos HOB1 as a threonine (T-232; Fig. 2). Fos T-232 may also be the target for MAP kinase activity, because, as with Jun S-63 and S-73, it is followed by a proline residue and, therefore, conforms to the consensus sequence for MAP kinase (S/TP) (Payne et al. 1991). Preliminary observations suggest that, indeed, T-232 is a substrate for MAP kinase and that the Fos HOB1/HOB2 domain is responsive to superactivation by H-ras [J.A. Sutherland and A.J. Bannister, unpubl.]. It is therefore possible that H-ras-induced phosphorylation of HOB1 regulates the activation functions of both Fos and Jun. This provides an attractive model by which external mitogenic signals, acting through Ras, affect the activation potential of nuclear transcription factors.

The HOB1 and HOB2 motifs may represent binding sites for proteins required to mediate the activation process. Phosphorylation by MAP kinase may potentiate the binding of such proteins to HOB1 (but not HOB2). Regulating one of the activation modules [HOB1] would then leave the second module [HOB2] free to cooperate with a distinct module within the same protein such as that present at the carboxyl terminus. Interestingly, the carboxy-terminal domain of Fos that cooperates with HOB2 is phosphorylated by protein kinase A (Tratner et al. 1992). Regulation by phosphorylation may turn out to be a common feature of activation modules that cooperate with HOB2. The use of such regulation may explain why HOB2 is found cooperating with distinct, rather than identical, modules within a given protein.

Pertinent to this discussion is the conservation of the HOB1 and HOB2 motifs within other members of the Jun, Fos, and C/EBP families. Both motifs are conserved in the two Jun family members Jun-B and Jun-D. In Jun-B, however, the proline residue that follows the MAP kinase-phosphorylated residues S-63 and S-73 is not conserved. This may reflect the fact that a kinase, other than the “proline-specific” MAP kinase is responsible for phosphorylating these serines in Jun-B. This would then allow Jun-B to be regulated independently of Jun and Jun-D.

In contrast to Jun family members, the Fos-related proteins, Fra-1 and Fos-B, do not have a conserved HOB1 and HOB2 motif. The HOB1/HOB2 region is unique to the Fos protein, suggesting that the activity residing within this domain may distinguish Fos function from that of other Fos-related proteins. The HOB2 motif is highly conserved in all members of the C/EBP family (Williams et al. 1991). This is particularly striking as this family does not show extensive conservation outside its DNA-binding domain. It is interesting to note that the three proteins shown here to possess a HOB2 activation domain, Fos, Jun, and C/EBP, are all members of the bZIP family of DNA-binding proteins. Although this sample is too small to be significant, the possibility that this domain is specific to members of the bZfP family cannot be excluded.

The HOB2-containing activation domains of Fos, Jun, and C/EBP show some interesting similarities that may reflect common function. Deletion analysis of these proteins indicates that the HOB2-containing activation domain is less active in the context of the full-length proteins (Friedman and McKnight 1990; Baichwal et al. 1991; Fig. 3). This suggests that a repressive activity may regulate the function of these domains. In the case of Jun, a cell-specific repressor has been implicated in the regulation of the Jun HOB1/HOB2 domain. It has been proposed that H-ras-induced superactivation of this domain is a result of relief from repressor activity (Baichwal et al. 1991). The results of the Fos deletion analysis imply that the cooperative activity between the HOB2 and the Fos carboxy-terminal domain is masked completely in the presence of amino-terminal sequences (Fig. 3, lanes 2,4). This may be an indication that in 1 BR cells, Fos HOB2 cooperates with HOB1 but not with the carboxy-terminal domain, owing to the presence of an inhibitor that specifically prevents cooperation with the carboxy-terminal domain.

Analysis of the secondary structure predictions for Fos and Jun reveal that the HOB1 motif is present partly within an α-helical region. An α-helix is predicted for residues 75–82 of Jun and 234–241 in Fos (Fig. 6). This helix falls just after the S/TP sequence that forms part of the recognition site for MAP kinase. Within this potential helix lie the two conserved glutamic acid residues (E-234 and E-236 in Fos, E-75 and E-77 in Jun) that are required for activation. In addition, the general character of the residues within this helix is conserved between Fos and Jun.

The HOB2 motif is also predicted to lie within an α-helix (Fig. 6). In the case of Jun and C/EBP, the entire region is α-helical, whereas in Fos, a central core [bounded by two prolines] is predicted to be within an α-helical structure. This central core contains the most conserved residues in this motif and has the consensus F, X, D/E, X, F/L, F/L. The first and fifth positions of the core motif are hydrophobic and are predicted to fall on the same face of the potential α-helix.

Activation domains characterized so far appear to be related merely by the prevalence of certain residues, and they are categorized as such. Families of activators are exemplified by the glutamine-rich domain of SP1 (Courey and Tjian 1988), the acidic domain of VP16 (Triezenberg et al. 1988), the proline-rich domain of CTF (Mermod et al. 1989), and the serine–threonine-rich domain of Oct-1 (Tanaka and Herr 1990). Transcription factors that fall into each one of these categories do not show absolute sequence identity. Furthermore, there is no evidence that any of these loosely related activation domains are related in function. In contrast, the HOB2 domain of Fos, Jun, and C/EBP shows precise sequence conservation, is similar in predicted secondary structure, and functions in a cooperative manner. Using the HOB2...
homology as a basis, it has been possible to identify successfully a transcription factor (C/EBP) with a cooperating HOB2 domain. The HOB2 motif therefore defines a new class of modular activation domain that can be identified through sequence homology. This motif can now be used to identify other transcription factors with modular activation domains. Cooperation assays with the prototype Fos or Jun HOB motifs could then be used to establish the validity of the observed homologies.

Materials and methods

Recombinant DNA

The Fos protein and the various deletions were cloned in-frame with the carboxyl terminus of the GAL4 DNA-binding domain present within the plasmid pHKG, which has the SV40 promoter and polyadenylation site, a polylinker 3' to the carboxyl terminus of the GAL4 DBD and carries on F1 origin of replication (C. Hagemeier and T. Kouzarides, in prep.). The rat c-fos sequence was used (Curran et al. 1987). The deleted versions S3–380 and 103–380 were generated using the naturally occurring BglII and HincII sites, respectively. To generate the remaining deletions, restriction sites were introduced into the c-fos gene. Three mutagenic oligonucleotides (TK67, TK68, and TK69) were used to introduce the following restriction sites: a BamHI–EcoRI site at positions 749–754 and 756–761, respectively (TK67), a BamHI–SacI site at positions 1061–1066 and 1068–1073, respectively (TK68), and a BamHI–EcoRI site at positions 869–874 and 876–881, respectively (TK69). Mutagenesis was carried out as described previously (Kouzarides et al. 1991). The deletions were generated in the following way: GF 53–308 with SacI of TK68, GF 53–205 with EcoRI of TK67, GF 210–380 with EcoRI of TK67, GF 314–380 with BamHI of TK68. GF 210–308 contains sequences from EcoR–SacI of TK67 and TK68, GF 210–244 contains sequences from EcoR–EcoRI of TK67 and TK69, GF 250–308 contains sequences from EcoR–SacI of TK69 and TK68. To separate the HOB1 and HOB2 motifs of Jun, two adjacent restriction sites were introduced [a BamHI site at 669–674 and an EcoRI site at 676–681] using oligonucleotide TK90 (Angel et al. 1988). These two sites are at precisely the same relative position as the sites described by oligonucleotide TK69 (Fig. 1), which were used to separate the HOB1 and HOB2 motifs of Fos. Consequently, the distance between the two motifs in GFH1/H2 and GH1/H2 is identical to that found in Fos and Jun. The GFH2/H2 construct has Fos sequences from the EcoRI site of TK67 to the BamHI site of TK69 and Jun sequences from the BamHI site of TK90 to a Nael site after residue 193. The GH1/H2 clone has Jun sequences from residue 1 of Jun to the EcoRI site of TK90 and Fos sequences from the EcoRI site of TK69 to the EcoRI site of TK68.

To generate the carboxy-terminal truncations of the Fos protein, a mutagenic oligonucleotide, TK104, was used to introduce a Nael site at positions 749–754 and 756–761, respectively. Constructs containing sequences from residue 267–380 were generated using PCR, and deletion at residue 332 was achieved using the XbaI site of TK104. Domain analysis of C/EBP was accomplished using specific PCR primers to amplify the appropriate domains. To generate the GH1/CH2 chimera, a mutagenic oligonucleotide (TK100) was used to introduce a BamHI and an EcoRI site at positions 712–719 and 720–726 in c-jun, respectively. The 29-residue HOB2 domain of C/EBP (amplified by PCR) was fused to Jun at the BamHI site of Jun/TK100. Thus, the distance between Jun HOB1 and C/EBP HOB2 is the same as the distance between Jun HOB1 and HOB2.

Dimerization of HOB1 was accomplished by introducing the BamHI fragment described by TK67–TK69 into pHKG 210–244. Dimerization of HOB2 was accomplished by introducing the BamHI fragment described by TK69 and TK68 into pHKG 250–308.

Transfections

Approximately 1 × 10⁶ 1 BR cells [SV40 T antigen-transformed human skin fibroblasts (Mayne et al. 1986)] grown at 37°C [5% CO₂] in Dulbecco’s Eagle medium supplemented with 10% fetal calf serum were transfected with a total of 7.5 µg of DNA using the calcium–phosphate coprecipitation technique. The precipitate was washed 6 hr after transfection and the cells were harvested 24 hr after transfection. Extracts from these cells were then used for CAT assays and for Western blots. The CAT assays were quantitated by liquid scintillation counting. The Western blots were probed with an antibody against the GAL4 DNA-binding domain [gift of M. Ptashne] to check the level of protein expressed by each vector. All plasmids expressed proteins at relatively equivalent levels. Each transfection was repeated a minimum of two times.

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