Yeast GCN4 as a probe for oncogenesis by AP-1 transcription factors: transcriptional activation through AP-1 sites is not sufficient for cellular transformation

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The Jun and Fos oncoproteins belong to the AP-1 family of transcriptional activators and are believed to induce cellular transformation by inappropriately activating genes involved in cell replication. To determine whether transcriptional activation through AP-1 sites is sufficient for transforming activity, we examined the properties of an autonomous and heterologous AP-1 protein, yeast GCN4, in rat embryo fibroblasts. GCN4 induces transcriptional activation through AP-1 sites but, unlike Jun and Fos, fails to induce cellular transformation, in cooperation with Ha-ras. Jun-GCN4 and Fos-GCN4 homodimers independently induce cellular transformation indicating that the amino-terminal regions of Jun and Fos each contain regulatory functions that are required for oncogenesis but are distinct from generic transcriptional activation domains. In addition, these observations have implications for the nature of the oncogenically relevant target genes that respond to Jun and Fos.

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v-Fos and v-Jun are oncogenes carried by murine osteosarcoma and avian fibrosarcoma retroviruses, respectively (Curran et al. 1982; Miller et al. 1984; Maki et al. 1987). The cellular homologs of these oncogenes, c-Jun and c-Fos, belong to the AP-1 family of transcription factors that bind specific DNA sequences (consensus TGAC-TCA), termed AP-1 sites (Bohmann et al. 1987; Distel et al. 1987; Struhl 1987; Angel et al. 1988a; Curran and Franza 1988; Ransone and Verma 1990; Boyle et al. 1991). AP-1 sites are found in many eukaryotic genes and are often associated with transcriptional stimulation in response to a variety of extracellular stimuli, including tumor promoters such as phorbol esters (Angel et al. 1987; Lee et al. 1987; Brenner et al. 1989; Stein et al. 1989; Jonat et al. 1990; Schule et al. 1990; Yang-Yen et al. 1990). AP-1 proteins contain a bZIP DNA-binding domain (Landschulz et al. 1988) consisting of a leucine zipper that mediates dimerization (Kouzarides and Ziff 1988; Neuberger et al. 1989a; Sellers and Struhl 1989) and symmetrically positions the adjacent basic regions for specific interactions with DNA (Agre et al. 1989; Pu and Struhl 1991a,b). AP-1 proteins can form homodimeric and/or heterodimeric complexes with DNA, with individual members of the family having distinct dimeriza-

tion specificity. Jun forms weak DNA-binding homodimers and Fos cannot dimerize, whereas Fos-Jun heterodimers bind AP-1 sites with high affinity.

It is generally believed that Fos and Jun contribute to the oncogenic state by binding to AP-1 sites and inappropriately activating genes involved in the control of cell proliferation. In this regard, several mutational analyses indicate a strong correlation between AP-1 site-dependent transcriptional activation, cellular transformation (often in cooperation with the activated Ha-ras), and tumor formation (Bohmann and Tjian 1989; Neuberger et al. 1989b; Schuermann et al. 1989; Baichwal and Tjian 1990; Alani et al. 1991; Binetruy et al. 1991; Lucibello et al. 1991; Wisdom et al. 1992). However, Jun and Fos are complex transcriptional regulators that activate or repress transcription, depending on the promoter and cellular factors with which they interact. Such complexities include the existence of multiple Jun-like and Fos-like proteins with different functional properties (Chiu et al. 1989; Li et al. 1990; Castellazzi et al. 1991; Doucas et al. 1991; Mumberg et al. 1991; Nakabeppu and Nathans 1991; Wisdom et al. 1992), the ability of the Jun and Fos proteins to regulate their own expression (Angel et al. 1988b; König et al. 1989), and the potential of Jun

and Fos to interact with other proteins involved in transcriptional regulation (Diamond et al. 1990; Jonat et al. 1990; Schule et al. 1990; Yang-Yen et al. 1990; Bengal et al. 1992; Li et al. 1992). It is also unclear whether Fos and/or Jun encode oncogenic-specific functions or whether they cause cellular transformation more indirectly by associating with proteins containing the leucine zipper motif such as other members of the AP-1 family or activating transcription factor–cAMP response element binding (ATF–CREB) proteins. Finally, the physiologically relevant target genes regulated by the Jun and Fos oncoproteins have not yet been identified. Thus, the functional complexities of mammalian AP-1 factors that allow for the exquisite regulation of gene expression make it difficult to determine the mechanisms by which Jun and Fos cause malignant transformation.

In this paper we investigate the role of AP-1 transcriptional activation in the process of oncogenic transformation using a novel approach involving the yeast transcription factor GCN4. GCN4, Jun, and Fos bind to identical DNA sequences (Hill et al. 1986; Bohmann et al. 1987; Distel et al. 1987; Struhl 1987; Angel et al. 1988a; Halazonetis et al. 1988; Nakabeppu et al. 1988), and Jun homodimers or Jun–Fos heterodimers can activate transcription in yeast cells and functionally substitute for GCN4 (Struhl 1988; Oliviero and Struhl 1991). However, GCN4 cannot interact with Jun or Fos (Kouzarides and Ziff 1989; Sellers and Struhl 1989), and it activates transcription primarily, and probably exclusively, by binding DNA as a homodimer (Pu and Struhl 1991b). Given the long evolutionary distance between yeast and mammals, it seems unlikely that GCN4 will be able to carry out Jun- or Fos-specific regulatory interactions beyond DNA-binding and transcriptional activation. Thus, GCN4 can be viewed as an autonomous and heterologous AP-1 transcription factor whose function should not be subject to the vast regulatory complexities of the Jun–Fos family. By analyzing the oncogenic properties of GCN4 derivatives, as well as GCN4–Jun and GCN4–Fos chimeric proteins, we demonstrate that transcriptional activation through AP-1 sites is not sufficient for cellular transformation. Moreover, we show that the amino-terminal regions of Fos and Jun each encode oncogenic functions that are distinct from generic transcriptional activation domains.

Results

GCN4 homodimers activate transcription from AP-1 sites in mammalian cells

To investigate whether yeast GCN4 can activate AP-1-dependent transcription and induce neoplastic transformation in mammalian cells, the coding region of GCN4 and several derivatives were cloned into the expression vector pMTpn (Fig. 1). To achieve high expression of the protein in mammalian cells, the GCN4 translational initiation region was replaced by an optimal mammalian translational initiation sequence (Kozak 1987). The GCN4 3′-untranslated region was substituted with the

Figure 1. Schematic diagram of GCN4 and GCN4 mutants. GCN4 wild type (GCN4) was modified and/or the specific regions were deleted to generate the mutants G1–G5 (Material and methods). (□) Activation domain; (■) basic domain; (▨) dimerization domain; (▤) 7-amino-acid insertion; (S) optimal translation start.

untranslated region of the human growth hormone, which increases message stability. Transcriptional activities of the various GCN4 derivatives were assessed in rat embryo fibroblasts (REFs) by transient cotransfection with chloramphenicol acetyltransferase (CAT)–reporter constructs (Fig. 2).

GCN4 stimulates transcription from a promoter containing five copies of the AP-1 site from the collagenase gene upstream of the TATA element of the thymidine kinase gene (5× TRE/tk–CAT). The level of transcription is comparable to that achieved by Jun and/or Fos. Deletion mutants that remove the activation (G2), DNA-binding (G3), or dimerization (G4) domains prevent AP-1 site-dependent transcriptional activation as does the N235E substitution in the GCN4 basic region (G5) that severely reduces DNA-binding activity (Pu and Struhl 1991a). Similarly, GCN4 activates transcription of the natural aP2 promoter, which contains one AP-1 binding site, as well as a CCAAT/enhancer-binding protein (C/EBP)–binding site (Herrera et al. 1989). As expected, transcriptional activation by GCN4 requires functionally intact bZIP and activation domains, and it is inhibited by a point mutation in the AP-1 recognition sequence. Thus, GCN4 activates AP-1 site-dependent transcription efficiently in mammalian cells with the same functional domains required for activity in yeast cells.

GCN4 functions primarily (and probably exclusively) as a homodimer in yeast cells (Pu and Struhl 1991b), and it does not form heterodimers with Jun or Fos (Kouzarides and Ziff 1989; Sellers and Struhl 1989). To determine whether GCN4 activity in mammalian cells is the result of homodimer formation, we examined the G1 derivative, which contains a 7 amino acid insertion between the leucine zipper and the basic region (Fig. 1). G1 efficiently binds AP-1 sites as a homodimer but it is unable to form DNA-binding heterodimers with wild-type GCN4; presumably, any heterodimeric interactions with GCN4 or G1 would involve a different (and probably mutually exclusive) set of bZIP proteins (Pu and Struhl 1991b). The observation that G1 activates tran-

5xTRE-tkCAT

aP2-CAT

Figure 2. AP-1-dependent transcription occurs in natural or hybrid promoter CAT constructs. (Top) REFs were transiently transfected with the 5× TRE/tk-CAT construct and 10 μg of each plasmid construct, respectively. Cellular extracts were analyzed for CAT activity as described in Material and methods. Three or more independent experiments were completed, and a representative chromatograph is shown. When the CAT assay is carried out for less time (to ensure linearity), the G1 derivative shows 50–70% of the activity of GCN4. (Bottom) REFs were transiently transfected with Ap2 (WT)-CAT, a construct containing one AP-1 site, or with a construct where the AP-1 consensus sequence has been mutated (mutant). The cellular extracts were assayed, and the transcriptional activity for each is listed.

scription to levels comparable to those achieved by GCN4 (Fig. 2) indicates that transcriptional activity is mediated primarily (and probably solely) by homodimers.

Finally, we tested whether GCN4 could activate transcription from AP-1 sites when stably expressed in REFs. We generated several G418-resistant pools of cell lines expressing GCN4, Jun, or Fos and measured transcriptional activity upon transfection with the 5× TRE/tk-CAT plasmid (Table 1). Cell lines containing GCN4, Jun, and Fos DNAs displayed comparable levels of AP-1 site-dependent activity, whereas control cell lines containing the expression vector showed background levels. As expected, we were unable to recover clones from cells transfected with a vector expressing C/EBP, a transcription factor that inhibits cell replication (Umek et al. 1991). The fact that GCN4, Jun, and Fos can activate transcription in stable cell lines indicates that expression of these proteins is not cytostatic or cytotoxic.

AP-1 transcriptional activation is not sufficient to induce transformation

We then examined whether GCN4, like Fos and Jun, could induce cellular transformation with the standard assay of c-Ha-ras cotransfection into REFs (Schutte et al. 1989a,b) (Fig. 3). When cotransfected with activated *ras*,

Fos and Jun yielded refractile, transformed foci, with Fos being most active. In contrast, GCN4 (and all derivatives tested) gave essentially no transformed foci despite its ability to activate AP-1 site-dependent transcription in transient or stably transfected REFs. Thus, it is clear that cellular transformation cannot depend simply on the overall level of AP-1 transcriptional stimulation.

The amino-terminal regions of Jun and Fos stimulate transformation independently

To determine the functional differences between GCN4 and the Jun and Fos oncogenes, we generated Jun-GCN4 and Fos-GCN4 hybrid proteins by reciprocal exchange of their activation domains, basic regions, or leucine zippers (Fig. 4). For each derivative, transcription and transformation assays were carried out in the presence of *v-ras* on equivalent populations of transfected cells. After normalizing for transfection efficiency, the relative transcription and transformation activities of each derivative were determined.

We first considered the possibility that the functional distinctions might reflect differences in DNA-binding activities especially because Fos/Jun and Jun/Jun complexes induce topological changes in the DNA (Kerppola and Curran 1991), whereas GCN4 does not (Gartenberg et al. 1990; Weiss et al. 1990). The substitution of the

Table 1. Stable clones of Fos, Jun, and GCN4 activate AP-1 CAT constructs

A. Construct	Number of independent G418 colonies
Control	140
Fos	220
Jun	180
GCN4	190
C/EBP	0
B. Stable pools	CAT conversion (%)
Control pool 1	5
Control pool 2	6
Control pool 3	3
Control pool 4	9
GCN4 pool 1	43
GCN4 pool 2	75
GCN4 pool 3	49
GCN4 pool 4	67
Fos pool 1	29
Fos pool 2	77
Fos pool 3	40
Fos pool 4	83
Jun pool 1	68
Jun pool 2	45
Jun pool 3	32
Jun pool 4	54

(A) G418-resistant colonies stably expressing the respective DNAs were generated as described in Materials and methods by use of a DNA/neo ratio of 15 : 1. Colonies were scored and either subcloned or pooled 14–21 days post-transfection. (B) G418-resistant colonies were isolated and transfected with the 5× TRE/tk-CAT construct. Transcriptional activation was assayed as described in Materials and methods.

GCN4 basic region into Jun and Fos (molecules f1 and j2; Fig. 4) should generate chimeric molecules with the binding specificity of GCN4 but with the activation and dimerization specificities of Jun or Fos. Examination of these hybrid proteins for transcriptional activity and oncogenic transformation (Figs. 4 and 5) indicates that the GCN4 basic region can functionally substitute for the basic regions of Jun and Fos in both assays. This indicates that the failure of GCN4 to transform cells is probably not related to subtle discrimination of AP-1 sequences in target genes.

To address the potential role of heterodimeric interactions, we analyzed chimeric molecules f2 and j3 in which the Fos and Jun leucine zipper are replaced by the GCN4 leucine zipper. These molecules, which should retain their own sequence specificity but be able to bind DNA only as homodimers, also induce cellular transformation. This observation indicates that interactions between Fos and Jun are not necessary for oncogenesis, and it suggests that Fos and Jun independently have transforming activity.

Although functional, molecules f1, f2, j2, and j3 showed relatively low transcriptional and transforming activities (Figs. 4 and 5), presumably reflecting reduced

DNA-binding affinities of the hybrid bZIP domains. We therefore generated additional Jun or Fos chimeric molecules in which the intact bZIP domains were substituted as a single unit. The resulting chimeras containing either the Jun or Fos amino terminus and the GCN4 bZIP domain (j4 and f3) induced both transcriptional activation and cellular transformation to levels comparable or even better than the wild-type molecules (Figs. 4 and 5). Conversely, a chimera containing the GCN4 activation domain fused to the bZIP domain of Jun (j1) activates AP-1-dependent transcription but fails to induce cellular transformation. To further map the Fos region(s) involved in cellular transformation, we fused amino-terminal (f4) and carboxy-terminal (f5) regions to the GCN4 bZIP domain. f5 is more effective at activation of AP-1 transcription than f4, although f4 is a more effective inducer of focus formation. The chimeric molecules f3, f4, and j4 also permit the formation of colonies in soft agar, and REFs stably transfected with these molecules cause aggressive tumor formation upon injection in athymic nude mice (data not shown).

To demonstrate that the GCN4 chimeras described above induced cellular transformation as homodimers, we substituted the GCN4 bZIP domain of molecule j4 with the bZIP domain derived from the G1 derivative, which has 7 extra amino acids between the basic region and the leucine zipper. Although the resulting molecule, j5, has a different dimerization specificity from j4, these derivatives induce transcriptional and transforming activities to comparable levels. These activities are inhibited by the introduction of an amino acid substitution (N235E) in the basic domain that inhibits DNA binding (j6). Taken together, these results demonstrate that the stimulation of AP-1-dependent transcription is not sufficient for cellular transformation. Both c-Jun and c-Fos carry information in their amino termini that do not stimulate generic AP-1 transcription better than the activation domain of GCN4, but these domains of Fos and (independently Jun) have the special property of inducing cellular transformation.

Discussion

Transcriptional activation through AP-1 sites is not sufficient for cellular transformation

The complex interactions that occur within and between different families of transcription factors are necessary for multicellular organisms to respond to the vast array of hormonal and developmental stimuli. A few specific domains are used by multiprotein families to promote the dimerization that is often necessary to bind to target DNA sequences with high affinity. In addition, interactions between different transcription factor families can permit “cross talk” between different regulatory circuits. However, this great complexity raises a significant challenge in the mechanistic interpretation of data obtained after introducing transcription factors into cells. It is extremely difficult to trace the multiple protein–protein and protein–DNA interactions in which a single factor might participate.

Figure 3. GCN4 does not transform REFs. REFs were transfected with 10 μ g of activated *ras* and each DNA construct, respectively. Stable foci were produced after 10–14 days in culture. The cells were fixed and stained with methylene blue. A representative field of cells for each DNA construct is shown.

To circumvent these problems, we have expressed a heterologous and autonomous AP-1 transcription factor, yeast GCN4, in mammalian cells to address three critical issues relating to the function of the mammalian AP-1 protein family. First, is there an obligatory relationship between AP-1-dependent transcription and AP-1-dependent malignant transformation? Second, what is the inherent transforming potential of c-Jun and c-Fos under situations in which they cannot interact? Third, what is the nature of the oncogenically relevant regulatory regions of Fos and/or Jun?

Our results clearly demonstrate that increased levels

of AP-1-dependent gene activation do not necessarily lead to cell transformation. When expressed in REFs, GCN4 stimulates transcription from artificial and natural promoters with an efficiency comparable to that of Jun and/or Fos. In striking contrast to Jun and Fos, expression of GCN4 does not yield transformed foci to a detectable extent, despite the facts that GCN4 does not appear to be cytostatic or cytotoxic and can chronically activate target genes in recovered G418-resistant colonies (Table 1). This leads to the conclusion that strong AP-1 site-dependent transcription does not necessarily lead to transformation.

name	construct	relative transcriptional activity	relative transforming activity
GCN4		100	< 2
jun		53	50
fos		65	100
f1		38	50
f2		16	20
f3		45	70
f4		22	80
f5		40	25
j1		37	4
j2		26	15
j3		27	20
j4		75	70
j5		66	60
j6		14	< 3
control		12	< 2

Figure 4. Amino-terminal regions of Fos and Jun are necessary for cellular transformation. A schematic diagram of the different Fos/GCN4 and Jun/GCN4 chimeric molecules is shown. Each chimeric construct was tested in the presence of c-Ha-ras for both AP-1-dependent transcriptional activity (5 $\times$ TRE/tk-CAT) and foci formation. Each construct was prepared in duplicate and double banded on a cesium chloride gradient. These numbers represent at least three to six independent transfection.

Figure 5. Homodimerization of Fos/GCN4 and Jun/GCN4 chimeric molecules can induce stable transformation. REFs were transfected with each chimeric construct (10 μ g) and activated *ras* (10 μ g). Stable transformants were visible after 10–14 days. Cells were fixed and stained with methylene blue. A representative field of cells transfected with each chimeric construct is shown.

ras plays an important role in AP-1-dependent transformation, and it also affects the phosphorylation of the amino-terminal region of Jun, which increases its transcriptional activity (Binetruy et al. 1991). However, our assays of transcriptional activation and cellular transformation are carried out in the presence of *ras* on identical populations of transfected cells. Thus, the role of *ras* in cellular transformation cannot be explained simply as increased AP-1-dependent transcriptional activity. Instead, it seems likely that *ras*-dependent phosphorylation might be necessary for the oncogenic regulatory function of Jun (and perhaps Fos).

The amino-terminal regions of Fos and Jun have inherent transformation potential

AP-1 family members interact with each other and with other transcription factor families, making it difficult to know whether Jun or Fos individually has transforming activity or whether the heterotypic interactions of each are necessary for transformation. Because the GCN4 bZIP domain strongly favors homodimer formation, chimeras between this domain and other portions of Jun or Fos may reasonably be assumed to result in homodimers *in vivo*. Furthermore, transcription and transformation activities are not affected by the insertion of 7 residues between the leucine zipper and the basic region of GCN4, which retains wild-type DNA-binding affinity as a homodimer but alters potential heterodimeric partners (Pu and Struhl 1991b). Thus, from the observation that Jun–GCN4 and Fos–GCN4 homodimers transform cells efficiently, we conclude that c-Jun and c-Fos each contain inherent transforming activity.

As the various members of the AP-1 transcription factor family might subtly discriminate between genomic AP-1 sites, it is possible that GCN4 fails to induce cellular transformation because it binds poorly to that subset of genes involved in cell proliferation that are capable of being activated by AP-1. However, this hypothesis is excluded by the properties of the Fos and Jun chimeras containing the GCN4 bZIP domain. In these cases (f3, f4, j4), all of the information needed to recognize AP-1 sites is provided by GCN4, yet cellular transformation is induced with an efficiency comparable to that of Jun or Fos. Transformation by these GCN4 chimeras requires AP-1 DNA-binding activity because foci are not observed upon transfection of a derivative containing a mutated bZIP domain (j6).

Mechanistic implications

Our results clearly indicate that the amino-terminal regions of Jun and Fos each contain special regulatory functions that are required for cellular transformation but are mechanistically distinct from generic transcriptional activation domains. Although these Fos and Jun regions are no better than the GCN4 activation domain at stimulating transcription from the promoters tested, it is conceivable that the oncogenically relevant AP-1 site containing a target gene has special requirements for tran-

scriptional activation by the AP-1 family. For example, activation of the key transforming gene might require a protein–protein interaction between Fos or Jun and some other specific transcription factor that binds (directly or indirectly) to the promoters. Presumably, such interactions would involve the amino termini of Fos or Jun and would not occur with the GCN4 activation domain. Another possibility is that Fos and Jun might differ from GCN4 in interacting with the basal transcription machinery at specific promoters; in this regard, certain TATA elements respond differentially to upstream activator proteins (Struhl 1986; Homa et al. 1988; Simon et al. 1988; Harbury and Struhl 1989; Wefald et al. 1990). All of the above models are consistent with the mutational analyses of Jun and Fos in which transcriptional activity correlates well with transformation ability (Bohmann and Tjian 1989; Neuberger et al. 1989b; Schuermann et al. 1989; Baichwal and Tjian 1990; Alani et al. 1991; Binetruy et al. 1991; Lucibello et al. 1991; Wisdom et al. 1992). However, although AP-1 transcriptional activity may be important or even necessary for transformation, it is not sufficient.

Alternatively, transformation by Fos and Jun might require AP-1-dependent repression of one or more tumor suppressor genes, especially as Fos and Jun can each act negatively at certain promoters containing AP-1 sites. In this view, GCN4 would function well as a positive activator but be incapable of acting negatively. A very recent study of Jun derivatives with mutations in the amino-terminal δ region shows an inverse correlation between transforming ability and transcriptional activity, leading to the suggestion of a negative role for Jun in transformation (Havarstein et al. 1992). Although our data show no correlation between the relative transcriptional activities of the various derivatives and their transforming activities, the possibility of transformation by a negative function of Jun and Fos is not excluded.

Our results are also useful in addressing which of the various functions of Jun and Fos might be relevant to cellular transformation. For example, the bZIP domains of Jun and Fos can interact directly with the glucocorticoid receptor (Schule et al. 1990; Yang-Yen et al. 1990) and with myoD (Bengal et al. 1992), generally (though not always) resulting in transcriptional repression. However, such interactions seem unlikely to be relevant for oncogenesis because the heterologous GCN4 and G1 bZIP domains are very compatible with cellular transformation. On the other hand, the amino terminus of Jun can mediate repression of transcriptional activation by the estrogen receptor (Doucas et al. 1991) and by myogenin or MyoD (Li et al. 1992); this suggests the possibility that transformation involves similar mechanisms to those utilized in suppressing the function of transcription factors that are necessary for establishing or maintaining the differentiated states of the cell.

Finally, it should be noted that the various constructs used here may be helpful in identifying AP-1-regulated genes that are involved in cellular transformation. Whatever mechanisms distinguish wild-type GCN4 from Fos or Jun, there must be differences in their abilities to

activate or repress certain crucial genes. By cloning mRNAs differentially regulated by GCN4 versus a GCN4 derivative with transforming activity, it might be possible to identify target genes that are centrally involved in AP-1-dependent malignant transformation.

Materials and methods

DNA constructs

Human c-Jun cDNA, rat c-Fos cDNA, and activated cHa-ras (pEJ6.6) were obtained respectively, from R. Tjian (University of California, Berkeley), T. Curran (Roche Institute of Molecular Biology, Nutley, NJ), and R. Weinberg (MIT, Cambridge, MA). All constructs were cloned into the *Bam*HI site of the mammalian expression vector pMTpn, which contains the SV40 early/late enhancer, the human metallothionein IIIa promoter, and the human growth hormone 3'-untranslated region (Greene et al. 1986). 5× TRE/tk-CAT (obtained from M. Karin, University of California, San Diego) contains five copies of the collagenase AP-1 sequence linked to the tk-CAT. The GCN4-coding sequence contained a *Bam*HI restriction site 5' of the basic domain and a *Xho*I restriction site at the first leucine of the dimerization domain (Pu and Struhl 1991). To permit efficient translational initiation in mammalian cells, we modified the region surrounding the GCN4 translational initiation codon by cloning TGCACGCCACCATGGTACCAAGTTTATTTGCT at the *Dra*I site; the native GCN4 gene was poorly translated in mammalian cells. The GCN4 mutant G1 was obtained by subcloning the *Bam*HI-*Sac*I fragment, which contains the 7 amino-acid insertion at amino acid 252. G2 was obtained by an in-frame deletion of the *Bam*HI restriction fragment containing the entire activation domain. G3 was obtained by restricting pMTpn-GCN4 with *Xba*I and *Xho*I, blunting the ends, and religating. G4 was obtained by deleting the DNA between the *Xho*I site and the *Sac*I site, leaving the TGA stop codon in-frame. G5 was generated by subcloning the *Bam*HI-*Sac*I restriction fragment derived from the GCN4-Glu-235 derivative (Pu and Struhl 1991) into the GCN4-pMTpn.

Chimeric constructs

The different chimeric Jun-GCN4 and Fos-GCN4 constructs were generated by inserting in-frame restriction sites without altering the amino acids sequence. DNA fragments were amplified by use of the polymerase chain reaction (PCR) and subcloned into the specific regions of each gene. All chimeric molecules were confirmed by DNA sequencing.

Jun chimeras j1 was obtained by amplifying the region containing the Jun bZIP domain with oligonucleotides A (CCCTCTAGAGCCTCAGACAGTGCCGAG) and B (CCCGAGCTCAAAATGTTTGCAACTGCTGCGTTAGCATG), cleaving the resulting product with *Xba*I and *Sac*I, and ligating the fragment into the GCN4 pMTpn plasmid. j4 was obtained by amplifying the Jun amino-terminal domain with oligonucleotides C (CCCGTTCGACCATGGAAACGACCTTCTATGACGATGCC) and D (CCCGGATCCCTCCTGGGACTCCATGTC), cleaving the resulting product with *Sal*I and *Bam*HI, and inserting the fragment into pMTpn-GCN4. The j2 molecule was obtained by amplification of the Jun dimerization region with oligonucleotide E (CCCCTCGAGGAAAAAGTGAAAA-CCTTGAAAGC) and B, and the fragment was subcloned into the chimeric molecule j4. The j3 molecule was obtained by subcloning the amplified Jun dimerization domain into the pMTpn-GCN4 molecule. The j5 molecule was obtained by sub-

cloning the *Bam*HI-*Sac*I fragment from the mutant G2 into the same sites of the chimeric j3 molecule. j6 was cloned as j5; however, the *Bam*HI-*Sac*I fragment was obtained from the mutant G3.

Fos chimeras The f1 chimera was obtained by substitution of the GCN4 basic region with the Fos basic domain in two successive subcloning reactions. The Fos carboxy-terminal domain was amplified with F (CCCAAGCTTCTCGAGGCGGAGACAGATCAACTTGAAGAC) and G (CCAAGGCCAGGAGAGGCACT); the resulting product was cleaved with *Hind*III and *Sac*I and subcloned into the vector pBluescript II KS to generate intermediate molecule SO68. The GCN4 basic domain was amplified with oligonucleotide H (CCCGTTCGACGGATCCTCGTGCTCTAAAACGTGCTGC), and the resulting *Sal*I-*Bam*HI fragment was cloned into SO68. Finally, the *Sal*I-*Sac*I fragment containing the GCN4 basic and Fos dimerization domain was subcloned into the pMTpn-Fos vector. The f2 chimera was obtained by amplifying the amino terminus of the Fos molecule with oligonucleotides I (AGAGCGCCCCATCCTTACGG) and J (GCCTCGAGCGTATCTGTGTCAGTCCCT); and following digestion with *Sal*I and *Xho*I, the fragment was subcloned into the pGEM vector (SO75). The carboxyl terminus of Fos in this reaction was obtained with oligonucleotides M (CCCAAGC-TTGTGGGTGAGCGTGAAAACTGGAGTTTATTTGGC-AGCC) and G, and the resulting DNA fragment was subcloned into an intermediate pMTpn-GCN4 vector (SO74). The complete fragment containing the GCN4 dimerization domain fused to the Fos carboxyl terminus was subcloned into the *Sal*I-*Sac*I restriction site of the pMTpn-Fos vector. The f3 chimera was obtained by subcloning the *Xho*I-*Sac*I fragment derived from the f2 chimera into the same sites of the f1 chimera. The f4 molecule was obtained by subcloning the GCN4 dimerization domain into the vector (SO68). The GCN4 basic and dimerization regions were then subcloned into the pMTpn-Fos molecule as a *Sal*I-*Sac*I fragment. The f5 molecule was obtained by cloning the amplified Fos carboxyl terminus used to create the f2 chimera downstream of a synthetic traditional start site containing an in-frame ATG, the amino acids for the influenza epitope, and a *Hind*III site in the same reading frame as the Fos carboxyl terminus (SO73). The *Sac*I site of the Fos carboxy-terminal and the *Bam*HI site in the GCN4 molecule were blunt ended, and the GCN4 basic and zipper domains were then subcloned in-frame 3' of the Fos molecule.

Cell culture

REFs were isolated as described previously (Schutte et al. 1989a). Briefly, 14- to 15-day Fisher rat embryos were sacrificed, rinsed, and trypsinized for 30 min at 37°C. Culture medium [Dulbecco's modified Eagle (DME) supplemented with glutamine, penicillin-streptomycin, and 10% fetal calf serum] was added, and the cells were dispersed, counted, and plated on 100-mm tissue culture dishes (Costar) at a density of 2×10^6 to 4×10^6 /dish. The cells were refed with culture medium after 12 hr, and trypsinized after 24–36 hr, and aliquots were frozen in liquid nitrogen. For experiments, REFs were plated at 1×10^5 to 2×10^5 /100-mm tissue culture dish and allowed to reach 80–90% confluence (1–2 days). All DNAs used in cell transfections were doubly banded on cesium chloride gradients followed by ethanol precipitation, and at least two different DNA preparations of each construct were independently transfected.

CAT assays

Confluent REFs (80–90%) were fed with culture medium 1–2 hr before transfection. For the experiment in Figure 2, 10 µg of

each construct was cotransfected with 2 μ g of 5 $\times$ TRE/tk-CAT and 1.5 μ g of β -actin-*lacZ* with standard calcium phosphate precipitation techniques. After 6–8 hr the cells were washed with PBS and refed with culture medium. The cells were harvested 40–48 hr after transfection, and the extract was prepared as described by Gorman et al. (1982). β -Galactosidase activity was determined for each extract, and all CAT assays were normalized for equivalent transfection efficiency. For the experiment in Figure 4, the same procedure was followed, except that 10 μ g of activated Ha-*ras* was also cotransfected (see below).

Foci formation

Confluent REFs (80–90%) were fed with culture medium 1–2 hr before to transfection. Ten micrograms of each construct was cotransfected with 10 μ g of activated Ha-*ras*, 2 μ g of 5 $\times$ TRE/tk-CAT, and 1.5 μ g of β -actin-*lacZ* with the standard calcium phosphate techniques. The cells were rinsed with PBS and refed with culture medium 6–8 hr post-transfection. The cells were trypsinized 48 hr after transfection and split 1 : 3. One of the plates was used to measure both the β -galactosidase activity and the levels of AP-1 transcriptional activation, as measured by CAT conversion. The remaining cells were left to reach confluence, at which time they were refed with culture medium containing 5% fetal calf serum and refed every 2 days thereafter. Stable cell transformants (foci) were visible after 7–10 days, and the numbers of foci were normalized for transfection efficiency by β -galactosidase activity. Foci generated were subjected to soft agar growth test. A pool of foci (plate) for several constructs (j4, f3, f4) were injected into nude mice, and tumors were observed from 8 to 25 days later.

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