TFIID can be rate limiting in vivo for TATA-containing, but not TATA-lacking, RNA polymerase II promoters

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We have studied the effect of exogenous expression of the basal transcription factor TFIID on the activities of several different TATA-containing and TATA-lacking promoters. Overexpression of TFIID from a transfected plasmid in Drosophila Schneider cells resulted in substantial concentration-dependent increases in expression from a cotransfected minimal TATA-containing promoter. Overexpression of TFIID activated expression from all TATA-containing promoters tested, with the maximum level of activation being inversely proportional to the strength of the promoter. In contrast, expression from TATA-less promoters was not enhanced, and could in fact be reduced, by increased expression of TFIID. Consistent with these findings overexpression of TFIID had opposite effects on Sp1-mediated activation observed from minimal synthetic promoters consisting of Sp1-binding sites and either a TATA box or initiator element. We discuss the significance of these results in terms of the role of TFIID in the initiation of transcription and as a possible regulatory target for expression from TATA-containing promoters, as well as the role TFIID may play in expression from TATA-less promoters.

[Key Words: TFIID–promoter interactions; transcriptional regulation; transcription initiation; Drosophila]

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Initiation of transcription by RNA polymerase II involves interactions between the polymerase, several essential transcription factors, and promoter DNA. To understand this critical process, the characteristics and functions of the factors necessary for the initiation reaction must be determined. Through fractionation of nuclear extracts from mammalian cells, six factors, designated TFIIB, D, E, F, and G, have been identified as being required in addition to the multisubunit RNA polymerase for initiation from a minimal class II promoter (Reinberg and Roeder 1987, Reinberg et al. 1987, Nakajima et al. 1988, Flores et al. 1990, Ohkuma et al. 1990; Sumimoto et al. 1990, for review, see Mermelstein et al. 1989, Sawadogo and Sentenac 1990). Counterparts to some of these essential factors have been shown to exist in Drosophila (e.g., Wampler et al. 1990) as well as in yeast (Buratowski et al. 1988, Cavallini et al. 1988, Hahn et al. 1989c), and the functional properties required for interactions between these essential components appear to have been well conserved during evolution, as certain factors can function in place of their counterparts in vitro in heterologous reconstituted systems (Buratowski et al. 1988, Cavallini et al. 1988, Hahn et al. 1989c, Flanagan et al. 1990, Wampler et al. 1990). The promoters of genes transcribed by RNA polymerase II commonly contain a conserved sequence element known as the TATA box [Breathnach and Chambon 1981], although a subset, referred to as TATA-less promoters, lack sequences that bear any significant homology to this element [for review, see Dynan 1986]. Kinetic and template commitment studies using TATA-containing promoters have indicated that two factors, TFIIB and TFIID, act at an early step in the initiation pathway [Davison et al. 1983, Fire et al. 1984; Reinberg et al. 1987]. Analysis of factor–template interactions in vitro have identified TFIID as the basal factor responsible for TATA box binding [Davison et al. 1983, Nakajima et al. 1988], which is required to promote transcription complex formation [Van Dyke et al. 1988, Buratowski et al. 1989] as well as to prevent repression of transcription by nucleosome assembly [Workman and Roeder 1987]. Studies defining the events leading to initiation from TATA-less promoters have yet to be reported, but it appears that TFIID and the other essential factors are required for this process as well [Pugh and Tjian 1990, Smale et al. 1990, Carcamo et al. 1991].

The cloning of the TFIID gene from yeast [Cavallini et al. 1989, Eisenmann et al. 1989, Hahn et al. 1989, Hörkosti et al. 1989, Schmidt et al. 1989] led to the isolation of cDNA clones encoding TFIID from a host of organisms, including plant [Gasch et al. 1990], human [Hofmann et al. 1990, Kao et al. 1990, Peterson et al. 1990],...
and Drosophila (Hoey et al. 1990; Muhich et al. 1990). Comparison of the primary amino sequence derived from TFIID cDNAs from these and other organisms has revealed that the TFIID protein is bipartite, composed of a highly conserved carboxy-terminal domain of ~180 amino acid residues and an amino-terminal region that is variable with respect to length and amino acid content. Analysis of recombinant TFIID proteins has shown that only the carboxy-terminal core is required for TATA box binding and support of basal transcription in vitro, whereas the amino-terminal region has been implicated in mediating the response of TFIID to certain upstream activator proteins [Hoey et al. 1990; Horikoshi et al. 1990; Peterson et al. 1990; Pugh and Tjian 1990]. Gene replacement studies done in yeast have shown that the carboxy-terminal domain alone is sufficient for growth and have also revealed that species-specific differences exist within this region, despite its high degree of conservation [Cormack et al. 1991; Gill and Tjian 1991; Reddy and Hahn 1991].

TFIID has been suggested to be a likely target for regulation by transcriptional activator proteins [for review, see Mitchell and Tjian 1989; Johnson and McKnight 1990; Ptashne and Gann 1990] in large part because it functions as the initiator of transcription complex formation. A number of in vitro studies have provided evidence consistent with this idea, as several factors have been shown to influence the DNA-binding characteristics of TFIID [Sawadogo and Roeder 1985; Horikoshi et al. 1988a,b; Sawadogo 1988, Meisterernst and Roeder 1991], while others appear to affect the rate and/or stability of TATA box binding under certain conditions, including during competition with chromatin assembly [Abamyr et al. 1988; Workman et al. 1988, 1990, 1991; Meisterernst et al. 1990]. The functional contacts, if any, that occur between activator proteins and TFIID have not yet been identified, but two viral activator proteins have been shown to bind to TFIID directly [Stringer et al. 1990; Horikoshi et al. 1991; Lee et al. 1991]. It also appears that at least some upstream activator proteins can influence the activity of TFIID through a class of intermediary molecules known as coactivators or adaptors [Berger et al. 1990; Kelleher et al. 1990; Martin et al. 1990; Pugh and Tjian 1990; for review, see Lewin 1990], examples of which have been partially purified from mammalian and Drosophila nuclear extracts [Dynlacht et al. 1991; Meisterernst et al. 1991].

The idea that TFIID–promoter interactions are promoted by certain activator proteins suggests that these factors may serve to recruit TFIID to the template which, in turn, allows for higher promoter activity. The importance of such interactions for transcriptional regulation would be heightened if TFIID were present in limiting quantities in vivo. Here, we present evidence that TFIID is maintained at a lower effective concentration relative to the other essential transcription factors within Drosophila cells and that TFIID can be rate limiting in vivo for expression from TATA-containing promoters. In contrast, we also show that TFIID is not limiting for expression from TATA-lacking promoters but appears to associate with another factor that is limiting for such promoters.

Results
Wild-type and mutant TFIID protein can be produced in transfected cells

Before embarking on an analysis of the possible effects of increased TFIID expression on promoter activity, we wished to verify that wild-type and mutant TFIID proteins could be transiently overexpressed in transfected cells. To this end, a cDNA encoding the Drosophila TFIID protein [kindly provided by T. Hoey and R. Tjian] was fused at the 5’ end of the coding region to a fragment encoding an 11-amino-acid sequence from the influenza virus HA1 [Wilson et al. 1984; Field et al. 1988] and then inserted into an expression vector containing the Drosophila actin 5C distal promoter. The resulting plasmid (Act flu–TFIID) was introduced into Drosophila Schneider cells by transient cotransfection as described previously [Han et al. 1989]. Western blot analysis of whole-cell extracts from cells transfected with Act flu–TFIID using a monoclonal antibody that recognizes the flu epitope [Niman et al. 1983] revealed a protein of ~38 kD [Fig. 1A], a molecular mass identical essentially to that of the Drosophila TFIID protein [Hoey et al. 1990]. When extracts prepared from cells transfected with increasing concentrations of Act flu–TFIID were analyzed, the amount of flu–TFIID protein detected increased in a manner directly proportional to the amount of expression vector [Fig. 1A].

We also analyzed the expression of two mutant forms of TFIID. Western blotting of extracts from cells transfected with Act flu–TFIIDΔNde (Fig. 1A), which should express the evolutionarily conserved carboxy-terminal core of TFIID joined to the flu epitope, and Act flu–TFIIDΔBgl (Fig. 1A), which encodes a flu–TFIID fusion protein with 17 amino acids deleted from the carboxyl terminus, confirmed that both of these proteins were also expressed in transfected cells. The accumulation of these two mutant forms, however, was somewhat lower than observed for the full-length protein, suggesting that the mutant proteins may be somewhat unstable. As with Act flu–TFIID, the amount of each mutant protein detected was dependent on the amount of TFIID expression plasmid transfected [Fig. 1A]. Therefore, we conclude that both wild-type and mutant TFIID proteins can be stably expressed from a plasmid in Drosophila Schneider cells. Furthermore, the amount of these proteins produced can be modulated by altering the amount of expression plasmid.

Increased expression of TFIID activates a minimal TATA promoter

The above experiments suggested that it would be feasible to determine whether increasing the level of TFIID could influence the expression of a cotransfected re-
porter gene. We chose to analyze initially the effect of increasing the concentration of TFIID on transcription from a very weak minimal promoter. For this, a reporter plasmid containing only the adenovirus E1b–TATA se-

quence [a gift of K. Martin and M. Green] inserted upstream of the bacterial chloramphenicol acetyltransferase (CAT) gene was cotransfected in the presence or absence of increasing concentrations of Act flu–TFIID. Alone, the level of CAT activity detected was barely above background. However, in the presence of Act flu–TFIID, expression was strikingly enhanced, reaching a maximum increase of ~135-fold at the highest amount of expression vector tested [Fig. 2A]. That this enhanced expression was dependent on the TATA element was verified by deletion of the TATA sequence from the reporter plasmid, which not only eliminated the weak basal CAT activity but also abolished completely the response of the plasmid to any amount of Act flu–TFIID tested [Fig. 2A].

Plasmids producing the truncated forms of TFIID were also tested for their ability to increase CAT gene expression from the E1b–TATA element. Cotransfection of Act flu–TFIIDΔNde with E1b–TATA CAT also showed concentration-dependent activation of CAT activity [Fig. 2B]. Somewhat lower levels of activation were observed with Act flu–TFIIDΔNde [50-fold vs. 135-fold], most likely owing, at least in part, to the reduced accumulation of the truncated protein in transfected cells relative to the full-length protein. This finding is consistent with previous experiments showing that only the carboxy-terminal core of TFIID is required to support basal transcription [Hoey et al. 1990; Horikoshi et al. 1990; Peterson et al. 1990; Cormack et al. 1991; Gill and Tjian 1991; Reddy and Hahn 1991]. The mutant form of TFIID lacking the carboxy-terminal 18 amino acids [Act flu–TFIIDΔBgl] failed to activate CAT expression from E1b–TATA CAT at any concentration tested [Fig. 2B]. This suggests that the exogenously expressed TFIID must bind DNA to activate CAT expression, as previous studies using the cloned yeast gene demonstrated that the entire conserved region of TFIID is required for both TATA box binding and initiation of transcription in vitro [Horikoshi et al. 1990].

These experiments establish that increased expression of functional TFIID molecules can enhance significantly

![Figure 1. Expression of wild-type and mutant TFIID proteins. (A) Western blot analysis of extracts from cells transfected with 4 μg of Act–PPA (M) or increasing concentrations of the indicated flu–TFIID expression plasmids as described in Materials and methods. The predicted sizes of the expressed proteins are 39 kD for Act flu–TFIID, 22 kD for Act flu–TFIIDΔNde, and 38 kD for Act–TFIIDΔBgl. The mobility of molecular mass marker proteins is indicated in kilodaltons (left). (B) Schematic depicting the structure of TFIID and showing the location of restriction sites used to create expression vectors encoding wild-type and mutant forms of TFIID fused to the flu epitope as described in Materials and methods. The filled box represents the amino-terminal domain, the hatched box represents the conserved carboxy-terminal core domain of TFIID.](image)

![Figure 2. TFIID can activate a minimal TATA box promoter. (A) Plot of relative CAT values obtained from cotransfections containing the E1b–TATA CAT or the E1bΔ–TATA CAT reporter plasmid with increasing concentrations of the TFIID expression plasmid. (B) Plot of relative CAT values obtained from cotransfections containing the E1b–TATA CAT reporter with increasing concentrations of the indicated TFIID expression plasmid. The total amount of expression vector in each sample was adjusted to 4 μg by the addition of Act–PPA as needed (see Materials and methods).](image)
transcription from a transfected reporter construct in a TATA box-dependent fashion. The results also indicate that TFIID is present in limiting quantities within the transfected cells because increased production of this protein allows an increase in the rate of transcription from a minimal TATA-containing promoter.

**Relative activation of TATA-containing promoters by increased TFIID is dependent on promoter strength**

Expression from the Elb-TATA CAT plasmid was extremely weak in the absence of exogenous TFIID. To determine whether the “strength” of a promoter might influence its response to TFIID, a variety of different TATA-containing promoters were tested for activation by increased levels of TFIID. These experiments were performed with a TFIID expression vector that lacked the flu sequences [Act–TFIID], but preliminary experiments indicated that these sequences did not affect TFIID activity (results not shown). We first tested two very strong promoters. The 2.6-kb actin 5C promoter, used to express TFIID from transfected plasmids owing to its high rate of transcription in Schneider cells, showed only slight increases in activity (approximately twofold) when fused to CAT and cotransfected with Act–TFIID (Table 1). The modest response of the actin 5C promoter to increased TFIID expression was not the result of some other limitation (e.g., exceeding the translational capacity of the transfected cells), as significantly higher CAT activities (>2000 relative units; see Table 1) could be obtained from transfections that included increased concentrations of the actin 5C reporter plasmid (data not shown). A reporter construct containing a 137-bp promoter fragment [−47 to +90], lacking heat shock factor [HSF]-binding sites, taken from the *Drosophila* 70K heat shock gene (*hsp70*; Ingolia et al. 1980), another relatively strong promoter, was also activated to a maximum of two- to threefold by the highest amount of Act–TFIID tested (Table 1). Transcription from both promoters was stimulated by Act–TFIID in a concentration-dependent manner, indicating that increased TFIID expression was responsible for the observed activation.

Several weaker promoters gave rise to intermediate levels of activation. *hsp70ΔAcc* CAT, which contains a 43-bp promoter fragment from the *hsp70* gene (−49 to −6), and thus the same TATA box and surrounding sequences found in *hsp70* CAT, displayed a much lower basal level than *hsp70* CAT (0.5 vs. 133; see Table 1). This difference is presumably the result of the presence of activator protein-binding sites in the *hsp70* CAT promoter that are absent in *hsp70ΔAcc* CAT (Parker and Topol 1984; Gilmour et al. 1989, 1990). Comparison of TFIID-mediated activation of *hsp70* and *hsp70ΔAcc* (Table 1) indicates that the weaker promoter showed significantly higher relative activation levels, reaching a maximum of 13-fold induction over basal expression. Two other weakly expressed TATA-containing fragments, taken from the *Drosophila* metallothionein promoter (Bunch et al. 1988) and the SV40 early promoter (SV2 CAT; Gorman et al. 1982), also showed relatively strong activation (~15-fold and 30-fold, respectively; see Table 1). These experiments indicate that an inverse relationship exists between promoter strength and relative activation by increased TFIID expression.

### Table 1. TFIID can increase TATA box promoter activity

| Act–TFIID (µg) | Actin 5C | *hsp70* | *hsp70ΔAcc* | Met | SV40 E |
|---------------|---------|---------|-------------|-----|-------|
| 0             | 30.6    | 132     | 0.5         | 1.5 | 1.0   |
| 0.5           | 39.6    | 156     | 3.1         | 10.6| 15.1  |
| 1.0           | 53.2    | 262     | 4.6         | 19.1| 30.1  |
| 4.0           | 64.3    | 280     | 6.6         | 16.3| 28.0  |
| Max activation| 2.1×    | 2.1×    | 13.2×       | 12.7×| 30.1× |

Cotransfections with the indicated plasmids were performed as described in Materials and methods, except the amount of the Act CAT reporter plasmid used was only 0.05 µg. Values in the upper part are displayed as the ratio between CAT activity and the internal control (β-gal activity) for each promoter in the presence of the indicated amount of Act–TFIID; the lower part gives the maximum fold activation observed in the presence of Act–TFIID relative to the promoter activity obtained in its absence.

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Increasing levels of TFIID do not enhance basal expression from TATA-less promoters

A plasmid expressing β-galactosidase from the *copia* long terminal repeat (LTR) [Mount and Rubin 1985], a TATA-less promoter, was used in the above experiments to monitor transfection efficiencies (Han et al. 1989). Somewhat unexpectedly, no variations in p-galactosidase activity were observed between extracts from cells cotransfected with Act–TFIID and those cotransfected with the control expression vector, Act–PPA (Han et al. 1989; data not shown). Although these findings validated the use of this plasmid as an internal control, they also prompted us to determine the effect of increased TFIID expression on other TATA-less promoters.

We first tested a relatively strong TATA-less promoter fragment (−38 to +172) from the segment polarity gene *en* (Poole et al. 1985; Soeller et al. 1988), which gives rise to approximately the same CAT expression as the *hsp70* promoter described above (132 vs. 158). In contrast to the *hsp70* promoter, which was activated in a concentration-dependent fashion by cotransfection with Act–TFIID, the TATA-less *en* promoter showed essentially no response to the addition of any amount of TFIID expression vector (Table 2). This was not the result of the strength of the *en* promoter, as expression levels from a second reporter containing a smaller *en* promoter fragment (+2 to +117; *enΔHinc* CAT) and another TATA-less promoter fragment, from the homeotic gene *Ultrabithorax* (*Ubx*; Hogness et al. 1985), were also not enhanced by TFIID despite their significantly decreased basal activities (158 vs. −7; see Table 2). CAT expression from a *copia* LTR reporter plasmid, which by
Colgan and Manley

Table 2. TFIID does not enhance TATA-less promoter activity

| Act–TFIID (μg) | en  | copia | Ubx | en Δ Hinc |
|---------------|-----|-------|-----|-----------|
| —            | 148 | 12.3  | 7.5 | 7.1       |
| 0.2          | 164 | 11.0  | 5.4 | 5.8       |
| 1.0          | 150 | 7.7   | 5.5 | 6.2       |
| 4.0          | 170 | 8.9   | 6.2 | 6.0       |

Max repression: 1.0 × 0.6 × 0.7 × 0.8 ×

Cotransfections with the indicated plasmids were performed as described in Materials and methods. Values in the upper part are displayed as the ratio between CAT activity and the internal control (β-gal activity) for each promoter in the presence of the indicated amount of Act–TFIID; the lower part gives the maximum fold repression observed in the presence of Act–TFIID relative to the promoter activity obtained in its absence.

Itself was also relatively weak, likewise showed no activation when cotransfected with Act–TFIID. Slight decreases in expression from all three of these weaker promoters were reproducibly observed (Table 2). Therefore, in contrast to the behavior of all TATA-containing promoters tested, all of the TATA-less promoters used here showed no increases in expression upon cotransfection with Act–TFIID; this indicates that TFIID is not rate limiting for transcription from TATA-less promoters under conditions where it is for TATA-containing promoters.

Overexpression of TFIID has opposite effects on TATA-containing and TATA-lacking promoters during competition with excess TATA sequences

As described above, addition of exogenous TFIID allowed increased expression from all TATA-containing promoters tested, indicating that the endogenous TFIID was present in concentrations that limited their activity. In contrast, expression from TATA-less promoters was not enhanced by overexpression of TFIID, which suggests that TFIID is not involved in expression from promoters of this type or, alternatively, that TFIID is not a limiting factor for TATA-less promoter activity. To test whether the endogenous TFIID is limiting for TATA-containing promoters and to determine whether TFIID is required for TATA-less promoter expression, we attempted to create competition for the endogenous TFIID. For this, a set of “competitor” plasmids was constructed by inserting either TATA-containing or TATA-less promoter fragments into a plasmid lacking a CAT gene. Each of these was then cotransfected with hsp70 CAT or en CAT plasmids under conditions in which the competitor was present in a large molar excess (40-fold) over the reporter.

We initially tested a competitor plasmid containing the same promoter fragment as the TATA-containing reporter itself (GEM hsp) and found that expression from hsp70 CAT was reduced by a factor of ~2.5 (Table 3). This reduction proved to be largely TATA dependent, as deletion of the TATA element from the hsp70 promoter fragment severely impaired its ability to serve as a competitor (data not shown). TATA-specific inhibition was also observed using the plasmid GEM Met–TATA, which bears a 33-bp TATA-containing fragment (~41 to ~8) from the metallothionein promoter. This competitor was able to reduce CAT expression from hsp70 by a factor of ~5 (Table 3). In contrast, a plasmid containing a TATA-less promoter fragment from the en gene produced, at most, only slight reductions in expression from the hsp70 reporter plasmid.

To determine whether the endogenous TFIID was actually the limiting factor being competed away in the above experiments, we tested whether expression from the TATA-containing reporter could be restored by adding Act–TFIID to transfection mixtures in place of Act–PPA (see Materials and methods) during competition. This resulted in significantly increased activity from the hsp70 reporter during competition with GEM hsp or GEM Met–TATA (Table 3) competitor plasmids. Collectively, these experiments indicate that excess TATA-containing promoter fragments can inhibit transcription from the hsp70 reporter by competing for the endogenous TFIID and that increased expression of TFIID can counteract such inhibition.

As with the hsp70 reporter, cotransfection of the competitor plasmids with the TATA-less en reporter plasmid brought about reductions in CAT activity. The TATA-containing competitor GEM hsp gave rise to a modest inhibition of CAT expression (~1.5-fold, see Table 3). Cotransfection with GEM Met–TATA or GEM en, which carries the same TATA-less promoter fragment as the reporter gene tested, also brought ~1.5-fold reduction in CAT expression. Although the effects on the en promoter were small, they have been observed consistently in multiple, independent experiments. In contrast to the results obtained with hsp70 CAT, when Act–

Table 3. TFIID can restore activity from TATA-containing, but not TATA-less, promoters during competition

| Promoter           | Competitor | Expression vector |
|--------------------|------------|-------------------|
|                    | ActpBR     | Act–TFIID         |
| hsp70 (TATA)       | GEM Met–TATA | 0.20 | 0.76 |
|                    | GEM hsp    | 0.40 | 1.02 |
|                    | GEM en     | 0.79 | 1.57 |
|                    | GEM3       | 1.0  | 2.3  |

| en CAT (TATA-less) | GEM Met–TATA | 0.66 | 0.54 |
|--------------------|---------------|------|------|
|                    | GEM hsp      | 0.67 | 0.51 |
|                    | GEM en       | 0.62 | 0.66 |
|                    | GEM3         | 1.0  | 1.0  |

Competition experiments were performed with the indicated reporter plasmids and competitor plasmids as described in Materials and methods. Transfections contained either 4 μg of Act–PPA or 4 μg of Act–TFIID, as indicated at the top. To determine the effect of Act–TFIID on promoter activity during competition, CAT activities were normalized so that the values from cotransfections with GEM3 and Act–PPA were set at 1.0.
TFIID was added during competition between the en CAT reporter and the two TATA-containing competitor plasmids, no increase in expression from the en promoter was detected. Instead, a further decrease in CAT activity was observed consistently with both [Table 3]. These results indicate that increased expression of TFIID alone cannot restore activity from a TATA-less promoter during competition with TATA-containing [or TATA-lacking] promoter fragments, suggesting that another, unidentified protein that can associate with both kinds of promoters is limiting for TATA-lacking promoters. The further decrease brought about by the TATA-containing competitor plasmids in the presence of excess TFIID suggests that TFIID can increase the association of this factor with the competing plasmid.

**Comparison of activated expression from minimal TATA-containing and TATA-less promoters in the presence of higher levels of TFIID reveals opposite effects**

The results from the competition experiments suggest that a TFIID-associated factor[s] may be limiting for expression from the TATA-less promoters tested. These findings are consistent with previous in vitro studies demonstrating that transcription from a TATA-less promoter composed of the initiator elements from the murine terminal deoxynucleotidyltransferase [TdT] gene or the adenovirus major late [AdML] promoter [Smale and Baltimore 1989] and the GC boxes constituting Sp1-binding sites from the SV40 early promoter [Dyan and Tjian 1983] requires the human TFIID fraction, but is not supported by purified yeast TFIID or recombinant hTFIID (Pugh and Tjian 1990; Smale et al. 1990). We therefore wanted to test the effect of increased TFIID levels on expression from this promoter in vivo to compare the results with those obtained with a similar Sp1-responsive, TATA-containing promoter. In addition, with such synthetic constructs the effects of “promoter strength” on TFIID responsiveness can be assayed systematically simply by altering the concentration of Sp1 expression vector included in the transfections.

We first constructed reporter plasmids containing the GC boxes alone [G CAT] or in combination with either the AdML promoter TATA box [GT CAT] or the TdT initiator [GI CAT] inserted upstream of the CAT gene. Comparison of the CAT activities obtained from these reporters in the presence of Sp1 [produced from Act Sp1] confirmed that the initiator element was recognized as a functional element in *Drosophila* cells, as its removal caused a significant decrease [10- to 30-fold] in Sp1-mediated activation [data not shown]. As was the case with other TATA-less promoters tested, the weak basal expression [i.e., in the absence of Sp1] from GI CAT was not enhanced by increased TFIID expression. In contrast, expression from GT CAT was activated ~20-fold under the same conditions [data not shown]. Increasing concentrations of Act Sp1 resulted in strong activation of expression from both GI CAT and GT CAT, but overexpression of TFIID with Sp1 had strikingly different effects on the activation levels observed. Activity from the TATA-containing reporter GT CAT was further activated 4- to 14-fold by overexpression of TFIID at all Act Sp1 concentrations tested. [Note that some of this increase may reflect elevated Sp1 concentrations, as the exogenous TFIID may have increased its expression. However, as shown in Table 1, activation of the strong Actin 5C promoter by TFIID overexpression was at most twofold, which is insufficient to account for the GT activation levels observed.] These findings are entirely consistent with the results obtained with natural TATA-containing promoters described above, including the observation that the lower the promoter activity is, the greater the response to TFIID. In contrast, expression from the TATA-less reporter GI CAT was inhibited [by a factor of ~3] at all Act Sp1 concentrations tested. These findings are also consistent with our observations with natural TATA-lacking promoters, although the inhibitory effect was significantly greater, perhaps reflecting the minimal nature of the synthetic GI CAT promoter. In addition, the approximately threefold reductions observed may well underestimate the actual inhibition if, as mentioned above, TFIID overexpression enhances Sp1 expression somewhat. Together, these results provide strong support for the notion that TFIID can be limiting for expression in vivo from TATA-containing, but not TATA-lacking, promoters. However, TFIID is able to functionally interact with a factor required for TATA-less promoter expression.

**Discussion**

The isolation of a cDNA encoding *Drosophila* TFIID [Hoe et al. 1990; see also Muhich et al. 1990] allowed us to create expression plasmids encoding full-length and truncated forms of this protein. We have shown that wild-type and mutant TFIID can be produced in a concentration-dependent manner from plasmids transfected into cultured *Drosophila* cells and that overexpression of the full-length protein can enhance expression dramatically from a weak minimal TATA-containing promoter. Expression from several different TATA-containing promoters was also activated by increased synthesis of TFIID, and the level of activation observed was inversely proportional to the strength of the promoter. Activity from TATA-less promoters, however, was not enhanced and, in some cases, was inhibited by increased expression of TFIID. In addition, competition experiments showed that the presence of excess TATA sequences could cause reductions in expression from TATA-containing, as well as TATA-lacking, promoters. Increased expression of TFIID restored activity from the TATA-containing, but not the TATA-less, promoter during such competition.

**Overexpression of TFIID can increase basal transcription**

The enhanced expression from TATA-containing promoters brought about by the production of exogenous
TFIID is a strong indication that the endogenous TFIID is maintained at lower levels in Schneider cells relative to other essential transcription factors. We found that expression increased as a function of TFIID concentration, demonstrating that the amount of functional TFIID present can be a factor that determines the transcription rate from a TATA-containing promoter. If a basal transcription factor other than TFIID was present in limiting concentrations, then increasing the TFIID concentration would have been without effect. Although it is conceivable that the increase in template concentration brought about by transfection significantly decreases the transcription factor–template ratio, our results nonetheless provide strong support for the notion that the available concentration of TFIID is significantly less than other essential transcription factors, which supports the view that TFIID itself, and not some other factor with which it interacts, is limiting for transcription from TATA-containing promoters.

It is conceivable that the level of functionally competent TFIID available within Schneider cells is controlled not by limiting its accumulation, but by maintaining TFIID in a conformation or complex that alone is incapable of associating with other essential transcription factors and/or binding productively to TATA sequences. This view is supported by the observation that TFIID characteristically behaves as a factor significantly larger than its molecular mass during purification (Reinberg et al. 1987; Pugh and Tjian 1990; Dynlacht et al. 1991) and is capable of associating with multiple factors that can inhibit basal transcription, and whose influence can be counteracted by the activator proteins USF (upstream stimulating factor) and Sp1 (Meisterernst et al. 1991; Meisterernst and Roeder 1991). The TFIID-mediated activation of basal transcription observed from minimal TATA-containing promoters may therefore be the result of a mechanism analogous to the previously described phenomenon of squelching (Gill and Ptashne 1988). Squelching, or transcriptional interference (Meyer et al. 1989), is an inhibition of transcription that can result from overexpression of a DNA-binding activator protein. Presumably, this reflects an interaction between the excess activator and a limiting essential transcription factor, resulting in sequestration of the factor in nonproductive complexes. Assuming that TFIID not bound to TATA sequences can interact stably with one or more other essential transcription factors, then the fact that overexpression of TFIID, or the DNA-binding defective mutant TFIIDΔBgl, did not reduce expression from any promoter tested supports the view that TFIID itself, and not some other factor with which it interacts, is limiting for transcription from TATA-containing promoters.

Figure 3. Overexpression of TFIID has opposite effects on the Sp1-mediated activation of minimal TATA-containing and TATA-less promoters. (Solid bar) Act pBR; (hatched bar) Act flu–TFIID. (A) Bar graph showing the CAT activity observed from transfections including the TATA-containing reporter plasmid GT CAT, which is represented schematically above the plot, the indicated amount of Sp1 expression vector, and 4 μg of either Act–PPA or Act flu–TFIID plus additional Act–PPA as necessary to bring the total amount of expression vector added to 4.6 μg. Transfections were performed as described in Materials and methods, except that each precipitate containing a total of 4.6 μg of expression vector. (B) Bar graph showing the CAT activity observed from transfections containing the TATA-less reporter plasmid GI CAT, which is represented schematically above the plot, the indicated amount of Sp1 expression vector, and 4 μg of either Act–PPA or Act flu–TFIID plus additional Act–PPA as necessary to bring the total amount of expression vector added to 4.6 μg. Transfections were performed as described in Materials and methods, except that each precipitate contained a total of 4.6 μg of expression vector.
of an imbalance in the stoichiometry between such negatively acting factors and TFIID brought about by the overexpression of TFIID.

Response to increased TFIID expression is inversely proportional to promoter strength

It has long been proposed that some upstream activator proteins function during transcription complex formation to facilitate TFIID–TATA box interactions, and several in vitro studies have provided support for this idea. Binding experiments have shown that certain factors such as USF or GAL4 can bind cooperatively with TFIID (Sawadogo and Roeder 1985; Sawadogo 1988), alter its DNase I footprint (Horikoshi et al. 1988a,b), and/or facilitate binding of TFIID during competition with nucleosome assembly (Workman et al. 1988, 1990, 1991; Meisternerst et al. 1990). Several classes of transcriptional activators appear to function through TFIID using a class of intermediary molecules known as coactivators or adaptors (Berger et al. 1990; Kelleher et al. 1990; Martin et al. 1990; Pugh and Tjian 1990), some of which have been partially purified (Dylnacht et al. 1991; Meisternerst et al. 1991). Intriguingly, the activation domains of the viral proteins VP16 and E1a are capable of binding directly to TFIID (Stringer et al. 1990; Horikoshi et al. 1991; Ingles et al. 1991, Lee et al. 1991). Thus, a number of activator proteins appear to affect, directly or indirectly, the rate and/or stability of the TFIID binding.

If the interaction of TFIID with the TATA box is a factor that influences the rate of transcription, then strong expression from a TATA-containing promoter must represent, to some degree, the ability of the promoter to recruit TFIID. Our experiments examining the effect of TFIID accumulation on promoters of different strength extend these in vitro studies by providing evidence that activator proteins can influence the ability of a promoter to recruit TFIID. The strong hsp70 promoter fragment (−37 to +90) that was only slightly affected by overexpression of TFIID contains binding sites presumably recognized by activator proteins (Parker and Topol 1984; Gilmour et al. 1989, 1990). Removal of some of these binding sites (hsp70ΔAcc, −39 to −11) not only caused substantially decreased promoter activity but also increased the sensitivity of the promoter to the level of TFIID. Likewise, the reduced enhancement of expression by exogenous TFIID from the synthetic GT promoter observed at elevated Sp1 concentrations suggests that TFIID may be recruited more efficiently to the promoter under these conditions. These results also suggest that at least some of the TFIID-associated factors apparently required for Sp1-mediated activation (Dylnacht et al. 1991; Meisternerst et al. 1991) must not be present in limiting concentrations or the exogenously expressed TFIID presumably would have been unable to function.

Overexpression of TFIID can reduce expression from TATA-less promoters

Because TFIID can function to initiate transcription upon binding to sequences that are divergent from the TATAAA consensus (Nakajima et al. 1988; Hahn et al. 1989a; Singer et al. 1990; Wobbe and Struhl 1990) or to TATA sequences located downstream of the transcription start site (Carcamo et al. 1990), it is difficult to determine whether a given promoter truly lacks a functional TATA box. Mammalian promoters considered to be TATA-less commonly contain GC-rich sequences upstream of the start site that can be bound by Sp1 (Dynan et al. 1986; Swick et al. 1989) and by another mammalian factor, known as ETF, which activates expression only from promoters lacking TATA boxes (Kageyama et al. 1989). In yeast, a binding site for the transcriptional activator GCN4, which appears to interact specifically with yeast RNA polymerase II (Brandl and Struhl 1989), can substitute for a TATA element in a naturally occurring, or hybrid, promoter (Zalkin et al. 1984; Chen and Struhl 1989). Together, these results suggest that TATA-less promoters may use a mechanism of initiation that does not require TFIID or one that uses TFIID in a manner unique to these promoters. Consistent with this latter proposal, results from reconstitution experiments have suggested that transcription in vitro from a synthetic TATA-less promoter requires a protein complex containing human TFIID associated with (an) auxiliary protein, designated the tethering factor, in addition to the other basal transcription factors and Sp1 (Pugh and Tjian 1990, 1991; Smale et al. 1990).

Our experiments have shown that expression from TATA-less promoters, unlike all TATA-containing promoters tested, cannot be increased by overexpression of TFIID, which suggests that TFIID is not limiting for TATA-lacking promoters. The observed inhibition of TATA-less promoter activity in vivo by TATA-containing segments and/or overexpression of TFIID suggests that a factor able to interact with TFIID is limiting for TATA-less promoter expression. This was demonstrated most clearly in our experiments where overexpression of TFIID resulted in substantial reductions of Sp1-mediated activation of the synthetic TATA-lacking promoter GI. The simplest explanation for these findings is that overexpressed TFIID can squelch (Gil and Prashne 1988) expression from the TATA-lacking promoters by sequestering a limiting factor. Therefore, the existence of limiting amounts of a factor[s], the function of which involves interaction with TFIID, could make TFIID de facto limiting for expression from TATA-less, as well as TATA-containing, promoters.

Materials and methods

Recombinant plasmids

All expression vectors were derivatives of the plasmid Act–PPA, which contains the Drosophila actin SC promoter and poly[A] site inserted into pBR322, as described in Han et al. (1989). Act–TFIID was created with the plasmid pFX29-6 (provided by T. Hoc and R. Tjian), which contains a 1.4-kb Drosophila TFIID (dTFIID) cDNA fragment inserted into the polylinker of Bluescript SK. This plasmid was cut with HphI, and the 3′ overhang was removed by digestion with T4 DNA poly-
merase and then digested with BamHI. The resulting 1.2-kb fragment was inserted into Act–PPA that had been cut with BamHI, filled in with Klenow, and digested with BglII. A precursor to Act flu–TFIIDBgl, known as Act–TFIIDABgl, was also constructed from pFX29-6 by isolating a 1-kb fragment generated by cutting with HphI, removing the 3' ends with T4 DNA polymerase, and digesting with BglII followed by fill-in with Klenow fragment. This 1-kb fragment was then inserted into Act–PPA that had been digested with BamHI, filled in with Klenow and digested with EcoRV. Act P-flu (provided by K. Han), used to create the flu epitope–TFIID fusion expression vectors, was constructed by cutting Act–PPA with BamHI and then partially with SalI and ligating the isolated 0.2-kb BamHI–SalI fragment with annealed complementary synthetic oligonucleotides encoding a 10-amino-acid peptide from influenza virus (Field et al. 1988). Actin flu–TFIID was created using pFX–Nde–Lo (provided by T. Hoey and R. Tjian), which contains the dT–TFIID cDNA with an NdeI site introduced at nucleotide 183 of the cDNA. This plasmid was cut with Ndel, filled in with Klenow, and then digested with BstXI, which generated an 840-bp fragment that was ligated with a 350-bp HindIII–Neol fragment from Act P-flu and a 7-kb Neo–BstXI fragment from Act–TFIID. Actin flu–TFIIDANde was constructed in the same way, except for the 363-bp Ndel–Klenow–BstXI fragment from the plasmid pFX–Nde–SH (provided by T. Hoey and R. Tjian), which contains an Ndel site introduced at nucleotide 660 of the TFIID cDNA, was used in place of the 840-bp fragment from pFX–Nde–Lo. Act flu–TFIIDABgl was constructed by ligating the 1.2-kb Ncol–BstXI fragment from Act flu–TFIID with the 7.1-kb Ncol–BstXI fragment from Act–TFIIDABgl. Act SpI (gift of T. Hoey and R. Tjian) is described in Courje and Tjian (1988).

All reporter plasmids except Ubx CAT, GT CAT, and GI CAT were constructed with the plasmid pUC–CAT, which is a derivative of pSV0CAT (Gorman et al. 1982). The plasmid E1b CAT (provided by K. Martin and M. Green; see Martin et al. 1990) contains a 17-bp oligonucleotide encoding the E1b–TATA element inserted immediately upstream of the CAT gene and was activated to a maximum of ~60-fold when cotransfected with Act–TFIID. When the TATA-containing oligonucleotide was removed from E1b CAT by digestion with XhoI and BamHI, followed by fill-in with Klenow and religation, the resulting plasmid, E1bΔCAT, was also activated (~35-fold) by cotransfection with Act–TFIID. This response was suppressed by mediating a cryptic TATA element located ~80 bp away from the 5' end of the CAT gene that was encoded in the SP27 vector [Promega] used to construct the original plasmid. Analysis of the sequence and construction of pUC–CAT showed that no such cryptic TATA elements had been placed near the 5' end of the CAT gene in this plasmid, therefore, the 570-bp Neol–Hindlll fragment that included the E1b–TATA element was removed from E1b CAT and inserted into pUC–CAT that had been digested with Neol and HindIII, generating the E1b TATA CAT plasmid used in this study. The plasmid E1bΔTATA CAT was created in a similar manner by inserting the 570-bp Neol–HindIII fragment from E1b CAT into pUC–CAT that had been digested with Neol and HindIII and showed no response to increasing concentrations of TFIID. ActSC CAT contains the 2.6-kb promoter fragment from Act–PPA inserted into pUC–CAT. hsp70 CAT contains a 137-bp NruI–PstI fragment from the Dro sophila 70K heat shock gene (Ingolia et al. 1980) inserted into pUC–CAT, whereas Hsp70ΔAcc contains a 43-bp MnlI–AccI fragment from the same promoter inserted into pUC–CAT. The plasmids Met–TATA CAT and en CAT (containing the en promoter fragment from bases ~39 to +172) have been described in Han et al. (1989), the construction of SV2 CAT (SV40 E) is detailed in Gorman et al. (1982). The reporter enΔHinc CAT was created by removing the 665-bp Hincll–Neol fragment from en CAT and inserting it into E1bΔTATA CAT that had been digested with SmaI and Ncol. copia CAT was constructed by inserting the 240-bp HindIII–XbaI fragment from the copia–lacZ internal control plasmid (Han et al. 1989) into HindIII and XbaI-digested pUC–CAT. Ubx CAT contains a promoter fragment from the Drosophila homeotic gene Ubx (Hogness et al. 1985) inserted into the plasmid vector p4 CAT (Thummel et al. 1988). GI CAT (provided by T. Hoey) contains the SV40 GC boxes inserted 42 bp upstream of the murine TdT initiator sequence (Smale and Baltimore 1989), which were taken from the plasmid pSP42 Ins (Smale et al. 1990) and inserted into the CAT vector pBLCAT3ΔHN (Luckow and Schutz 1987). A similar plasmid, known as GTI CAT (provided by T. Hoey), contains the SV40 GC boxes, the AdML promoter TATA sequence, and the TdT initiator sequences (all from plasmid "VII" as described in Smale et al. 1990) inserted upstream of the CAT gene in pBL3CATΔAHN. GT CAT was constructed by removing a Ndel–SalI fragment containing the GC boxes and AdML TATA box from GTI CAT and reinserting it upstream of the CAT gene in pBL3CATΔAHN.

All competitor plasmids contained promoter fragments inserted into the polylinker of the plasmid vector GEM3 (Promega). GEM hsp was constructed by removing the 400-bp HindIII–EcoRI fragment (which contained 137 bp of hsp70 promoter sequence and the first 246 bp of the CAT gene) from hsp70 CAT reporter plasmid and inserting it into GEM3 that had been digested with HindIII and EcoRI. GEM hspΔTATA was constructed by digesting GEM hsp with AccI, filling in with Klenow, and then religating. This removed a ~40-bp fragment from GEM hsp that encodes the hsp70 TATA box and surrounding sequences [bases ~47 to ~6]. GEM en was constructed by ligating the 475-bp HindIII–EcoRI fragment (which contained 221 bp of en promoter sequence and the first 246 bp of the CAT gene) from en CAT with GEM3 digested with HindIII and EcoRI. GEM Met–TATA consisted of the 40-bp SalI–MscI fragment from Met–TATA CAT inserted into GEM3 that had been digested with SalI and SmaI.

DNA transfection and transient expression assays

Transient cotransfection assays were performed essentially as described in Han et al. (1989). Drosophila Schneider L2 cells were maintained at 24°C in M3 media supplemented with 10% fetal bovine serum (GIBCO) and were plated at 2–4 x 10^5 cells per 60-mm tissue culture dish one day before transfection, and plasmid DNA was transfected with calcium phosphate coprecipitation. Except for the competition experiments or unless indicated otherwise, each precipitate contained 0.2–4 μg of the appropriate TFIID expression vector, 2 μg of reporter plasmid, 2 μg of copia–lacZ internal control plasmid, and GEM3 as needed to reach a total of 10 μg. For competition experiments, each precipitate contained 4 μg of Act–PPA or Act–TFIID, 0.2 μg of reporter plasmid, 2 μg of copia–lacZ, and 5 μg of GEM3 or the indicated competitor plasmid for a total of 11.2 μg. In all experiments, the total amount of actin 5C promoter in each sample was kept constant by the addition of Act–PPA as required. All experiments were repeated at least three times and many (e.g., the competitions) up to six times. Transfected cells were incubated at 24°C for 48–50 hr, removed from tissue culture dishes by agitation, washed twice with PBS, resuspended in 0.1 ml of 0.25 M Tris–HCl [pH 8.0], and frozen at ~70°C. Whole-cell extracts were prepared by thawing and then sonicating for 2 min in an ultrasonic sonifier (Branson), followed by centrifugation at full speed for 10 min in a microcentrifuge. Transfection efficiencies were determined by assaying for β-galactosidase activi-
ity as described in Han et al. (1989). Variations in transfection efficiencies within a given experiment were almost always less than twofold. CAT assays for promoter activity were also performed as described in Han et al. (1989).

**Western blot analysis**

For preparation of whole-cell extracts for Western blot analysis, Schneider cells were transfected and harvested exactly as described above, except that the cell pellet was resuspended in 1 ml of PBS instead of 0.25 ml Tris-Cl (pH 8.0). A 0.25-ml aliquot was removed from each resuspended sample and the cells pelleted, resuspended in 30 μl of 0.25 ml Tris-Cl (pH 8.0), and frozen at −70°C. Extracts were then prepared from these aliquots as described above and assayed for β-galactosidase activity.

The remaining cells were pelleted, and the packed cell volume [PCV] for each sample was determined. [Cell pellets were resuspended in 1× PCV of PBS plus 2× PCV Laemmli loading buffer, and the samples were boiled for 5 min and centrifuged for 5 min in a microcentrifuge.] Aliquots were taken from each, with the amount removed determined by the β-galactosidase activity observed for the sample, and the volumes of the samples were made equal by the addition as needed of identically prepared extracts from mock-transfected cells. The proteins in these samples were resolved on a 10% SDS–polyacrylamide gel, transferred to nitrocellulose, and blotted with the mouse monoclonal antibody 12CA5 [Berkeley Antibody Company, Niman et al. 1983]. A goat anti-mouse IgG antibody, conjugated to alkaline phosphate (Sigma Chemical Company), was used as the clonal antibody 12CA5 (Berkeley Antibody Company; Niman et al. 1983). For preparation of whole-cell extracts for Western blot analysis, Schneider cells were transfected and harvested exactly as described above, except that the cell pellet was resuspended in 1 ml of PBS instead of 0.25 ml Tris-Cl (pH 8.0). A 0.25-ml aliquot was removed from each resuspended sample and the cells pelleted, resuspended in 30 μl of 0.25 ml Tris-Cl (pH 8.0), and frozen at −70°C. Extracts were then prepared from these aliquots as described above and assayed for β-galactosidase activity.

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