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TATA-Binding Protein Recognition and Bending of a Consensus Promoter Are Protein Species Dependent

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Abstract
The structure and behavior of full-length human TBP binding the adenovirus major late promoter (AdMLP) have been characterized using biophysical methods. The human protein induces a 97° bend in DNA_{AdMLP}. The high-resolution functional data provide a quantitative energetic and kinetic description of the partial reaction sequence as native human TBP binds rapidly to a consensus promoter with high affinity. The reaction proceeds with successive formation of three bound species, all having strongly bent DNA, with the concurrence of binding and bending demonstrated by both fluorescence and anisotropy stopped flow. These results establish the protein species dependence of the TBP–DNA_{AdMLP} structure and recognition mechanism. Additionally, the strong correlation between the DNA bend angle and transcription efficiency demonstrated previously for yeast TBP is shown to extend to human TBP. The heterologous NH₂-terminal domains are the apparent source of the species-specific differences. Together with previous studies the present work establishes that TBP_{wt}–DNA_{TATA} function and structure depend both on the TATA box sequence and on the TBP species.

Abbreviations
TBP TATA-binding protein, with the y and h prefixes denoting yeast and human TBP, respectively
AdMLP adenovirus major late promoter
PIC preinitiation complex
CTD COOH-terminal domain
NTD NH₂-terminal domain
FRET fluorescence resonance energy transfer
TAMRA N,N,N',N'-tetramethyl-6-carboxyrhodamine
T*AdMLP_{dp} F 14 bp DNA duplex (5'-CGCTATAAAAGGGC-3') bearing the eight base AdMLP TATA sequence, with 5'-TAMRA and 3'-fluorescein on the top strand
TFIIA and -B class II general transcription initiation factors A and B
Table 1. Optimal Rate Constants and Corresponding Enthalpy and Entropy Changes for hTBP−DNA\textsubscript{AdMLP} Partial Reaction Steps in Accord with Equation 2 Together with Relative Quantum Yields for Each of the Three Bound Species

<table>
<thead>
<tr>
<th>(i)</th>
<th>(k_i \text{ (s}^{-1})</th>
<th>(\Delta H_i^{\ddagger} \text{ (kcal mol}^{-1})</th>
<th>(\Delta S_i^{\ddagger} \text{ (cal K}^{-1} \text{ mol}^{-1})</th>
<th>QYI(_1)</th>
<th>QYI(_2)</th>
<th>QY\text{final}</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>(5.5(5.2,5.6)\mu\text{M}^{-1})</td>
<td>(7.4(6.2,8.6))</td>
<td>(-4.3(-8.5,-0.3))</td>
<td>0.52(0.42,0.59)</td>
<td>0.52(0.49,0.54)</td>
<td>0.52(0.49,0.54)</td>
</tr>
<tr>
<td>2</td>
<td>(2.75(2.6,3.1))</td>
<td>(3.0(3.0,4.1))</td>
<td>(-48(-52,-45))</td>
<td>0.52(0.42,0.59)</td>
<td>0.52(0.49,0.54)</td>
<td>0.52(0.49,0.54)</td>
</tr>
<tr>
<td>3</td>
<td>(0.41(0.39,0.43))</td>
<td>(9.5(7.4,11))</td>
<td>(-30(-37,-23))</td>
<td>0.52(0.42,0.59)</td>
<td>0.52(0.49,0.54)</td>
<td>0.52(0.49,0.54)</td>
</tr>
<tr>
<td>4</td>
<td>(0.017(0.013,0.022))</td>
<td>(3.2(3.0,4.9))</td>
<td>(-58(-63.5,-52))</td>
<td>0.52(0.42,0.59)</td>
<td>0.52(0.49,0.54)</td>
<td>0.52(0.49,0.54)</td>
</tr>
<tr>
<td>5</td>
<td>(0.045(0.025,0.066))</td>
<td>(26(22,28))</td>
<td>(23(8.4,38))</td>
<td>0.52(0.42,0.59)</td>
<td>0.52(0.49,0.54)</td>
<td>0.52(0.49,0.54)</td>
</tr>
<tr>
<td>6</td>
<td>(0.022(0.018,0.026))</td>
<td>(9.2(6.1,12))</td>
<td>(-37(-47,-26))</td>
<td>0.52(0.42,0.59)</td>
<td>0.52(0.49,0.54)</td>
<td>0.52(0.49,0.54)</td>
</tr>
</tbody>
</table>

a The values shown are at 20 °C and 1 M standard state for each step, \(i\), along the reaction pathway. The assumption was made that \(\Delta C_p^{\ddagger} = 0\); therefore, there is no temperature dependence for \(\Delta H_i^{\ddagger}\) or \(\Delta S_i^{\ddagger}\).

b Parameter errors, which derive from a joint confidence region, were obtained in the following manner: the 15 × 15 variance matrix was calculated from the inverse of the approximate Hessian matrix following the method of Bard (44), using a 1% change in the various parameters to calculate the required partial derivatives. The partial derivatives were taken over 50 time points for each of the 10 kinetic curves and over 50 fractional saturation points for each of the 3 binding curves. In other studies, we have used simulated noisy data with appropriately scaled random noise that corresponded to the experimental errors to obtain the distribution of fitted parameters, a widely recommended and common procedure (45). That approach yields results that are in excellent agreement with a third approach: knowing the global optimum from 25 random starting positions in parameter space, the 25 sets of parameter values obtained when the Simplex search crossed the 68% joint confidence boundary were used to obtain the distribution of parameter values. The latter two procedures provided estimates that were not dependent on an assumed hyperelliptical joint confidence region. The reported parameter error estimates are averages of those obtained from the variance matrix and the latter procedure employing the joint confidence boundary. Despite the care taken in determining the error estimates, these values must be considered with caution since they are the extreme limits for each individual parameter as projections of the joint confidence region onto a given parameter axis. Because the volume of the joint confidence region is markedly less than that of the hyperrectangle described by all of the parameter errors (46), one cannot assume that any parameter vector with projections jointly within the error estimates provides an adequate description of the data. Rather, the valid values are those that adequately represent the satisfactory fits to the experimental data based on the reduced \(\chi^2\) values for the stopped flow and relaxation curves of 0.015 and 0.025 (normalized from 1 to 0) and an average 6% error for the equilibrium constant.

c \(\Delta S_i^{\ddagger}\) values were calculated from \(k_i\) and \(\Delta H_i^{\ddagger}\) using transition state theory.
Figure 1. Donor fluorescein emission in the steady state (A) and resolved in the ns time regime (C, upper curve) for unbound T*AdMLP
\(_{dp}^*F\) and the corresponding spectra following human TBP binding (B and C, lower curve). The relatively straight and rigid unbound
duplex maintains maximum separation of the 3′ FRET donor, fluorescein (F, 518 nm peak), and the 5′ acceptor, TAMRA (T, 578 nm peak). (A) Since the rate of energy
transfer from donor to acceptor depends on the inverse sixth power of the distance between these dyes, the intensity of the fluorescein emission peak is high: the excited state fluorescein population relaxes back to the
ground state primarily by photon emission rather than by the transfer of energy to the TAMRA population. (B) \(h\text{TBP}\) binding and bending result in a much decreased interdye distance, greatly increasing the efficiency of energy transfer and thus decreasing the donor emission. We have shown previously that both the labeled duplex and the protein are stable for at least 1 h under our solution conditions with no measurable change in the spectra over that time period (22). (C) The upper decay curve was generated using the free duplex and is the time-resolved counterpart of (A) with a 1.49 ns average donor lifetime. The increased transfer rate following protein-induced bending shortens the lifetime of the observed fluorescein emission (lower curve) to yield the time decay equivalent of (B) and a 0.726 ns average donor lifetime. The protein activity was determined as described to be 26%.
Figure 2. Normalized fluorescence stopped flow kinetics curves of hTBP associating with T*AdMLP*$F (observed, open circles, and calculated, solid line) compared to the corresponding yTBP reaction (broken line, panel A) and in panel B to the hTBP–AdMLP anisotropy stopped flow kinetics trace (open circles). Shown are the time-dependent changes in the donor fluorescein emission as the T*AdMLP*$F population binds to human and yeast TBP, with the time axis interrupted to clarify the differences (panel A). Although both interactions are biphasic, the human protein binds with an initial relatively very fast phase not seen with the yeast protein and reaches completion significantly faster than does yTBP. Additionally, this fast phase accounts for only 10–22% of the total amplitude change observed with hTBP with these conditions, whereas the faster eigenvalue dominates the yTBP reaction, with the corresponding amplitude ranging from 52% to 75% of the overall change. This difference arises because the equilibrium in the first partial hTBP reaction is strongly toward dissociation, overwhelming the contribution of $k_3$ and yielding a much less stable complex than with yTBP. The curves shown for both proteins were obtained using 109 nM protein reacting with 20 nM duplex at 15 °C. The hTBP experiments were conducted identically to those using yTBP except for the presence of 10% glycerol in the hTBP buffer, shown previously to have no effect on yTBP kinetics (22). The yTBP association curve was constructed for identical conditions using previously collected data (18). The hTBP–DNA AdMLP reaction monitored by stopped flow fluorescence anisotropy (open circles, panel B) is compared with the corresponding stopped flow FRET curve (solid line, panel B). The former was obtained using T*AdMLP$dp∗, with the normalized change in anisotropy ($r$) equal to $(r_t - r_0)/(r_\infty - r_0)$ and is the average of five replicate curves. Both traces are biphasic and show clearly the initial fast phase reflecting $h_1$ formation. Because the TAMRA emission changes only slightly, the anisotropy change accurately tracks the kinetics of any process yielding a change in rotational correlation time of the labeled oligonucleotide, which in this case is hTBP–DNA AdMLP binding. Replacing the numerator of the above anisotropy expression with $(r_\infty - r_t)$ shows that the two processes are proceeding in tandem (inset), confirming the concurrence of DNA binding and bending. The average signal/noise at $t_{1/2}$ was ~86 for the FRET data and ~8 for the anisotropy data; the two curves agree within error.
Figure 3. Comparative relaxation kinetics of the TBP−T*AdMLP_dpx*F complex at 20 °C following a challenge with unlabeled duplex for the human (solid line) and yeast (broken line) proteins. The hTBP complex responds to addition of ~1 µM unlabeled DNA_{AdMLP} with monophasic relaxation with $R = 0.00417$ s$^{-1}$. In contrast, the yTBP complex yields sharply biphasic decay with $R_1 = 0.0766$ s$^{-1}$ (11%) and $R_2 = 0.00154$ s$^{-1}$ (89%), published previously (18, 47). Neither the hTBP nor yTBP (18) complex is sensitive to the concentration of challenge DNA from 1 to 10 µM. The full hTBP relaxation curve from the global analysis (solid line) is shown in the inset together with the raw data (open circles). The reaction went to ~95% completion, and the observed amplitude change was consistent with that from the equilibrium experiments.
Figure 4. The energetic progression of the reaction from TBP + DNA_{AdMLP} on the left to the most stable binary complex on the right for the human (A) and yeast (B) proteins at 25 °C. Differences in the energetics of the hTBP and yTBP (18) reaction progressions are readily apparent from a graphic representation of $\Delta H^{\circ\dagger}$ (solid line) and $T \Delta S^{\circ\dagger}$ (dashed line). Both reactions are endothermic overall, with $\Delta H^c$ and $\Delta S^c$ values for hTBP of 27.8 (23.8, 31.8) kcal mol$^{-1}$ and 131.3 (118, 145) cal K$^{-1}$ mol$^{-1}$ and for yTBP, 13.4 (11.6, 15.2) kcal mol$^{-1}$ and 81.3 (76.6, 86.6) cal K$^{-1}$ mol$^{-1}$ (18). These “overall” values represent the difference between the thermodynamic parameters for the final complex and free TBP + DNA. Because all three conformers are present at equilibrium, the observed change in such parameters is the difference between a weighted average of the three bound species and the corresponding value for the reactants; $\Delta H^c$ obtained from the van’t Hoff analysis thus differs from the “overall” value shown in the figure [Table 1 (18)]. The modest activation energy of 7 kcal mol$^{-1}$ required for the first hTBP transition (TBP + DNA$\rightarrow$I$_1$) differs markedly from that of the yTBP pathway, for which this step presents the largest energetic barrier with $\Delta H^{\circ\dagger} = 35.1$ kcal mol$^{-1}$. The latter is overcome by a commensurate increase in entropy with $\Delta S^{\circ\dagger} = 87.8$ cal K$^{-1}$ mol$^{-1}$, in contrast to the decrease in entropy in the course of hI$_1$ formation with $\Delta S^{\circ\dagger} = -4$ cal K$^{-1}$ mol$^{-1}$. In the second partial reaction (I$_1$ $\rightarrow$ I$_2$) the hTBP transformation repeats its pattern for the first step with a similar entropic gain and enthalpic loss. In contrast, the yTBP partial reaction is strongly exothermic with a decrease in entropy, with $\Delta H^{\circ\dagger} = 3$ kcal mol$^{-1}$ and $\Delta S^{\circ\dagger} = -55$ cal K$^{-1}$ mol$^{-1}$. The result is that, whereas the hI$_2$ conformer is 6.3 kcal mol$^{-1}$ higher in energy and 27.9 cal K$^{-1}$ mol$^{-1}$ higher in entropy than hI$_1$, the corresponding thermodynamic changes in the yTBP reaction are $-26$ kcal mol$^{-1}$ and $-94$ cal K$^{-1}$ mol$^{-1}$, respectively. In the final transformation, hI$_2$ surmounts a substantial activation energy to become hTBP−AdMLP$\rightarrow$final, aided by an accompanying increase in entropy, with $\Delta H^{\circ\dagger} = 25.7$ kcal mol$^{-1}$ and $\Delta S^{\circ\dagger} = 23.1$ cal K$^{-1}$ mol$^{-1}$, to achieve the largest energetic changes, with $\Delta H^c$ and $\Delta S^c$ values of 17.1 kcal mol$^{-1}$ and 59.6 cal K$^{-1}$ mol$^{-1}$. The final yTBP step is likewise entropically driven as the energetic losses associated with yI$_2$ formation are overcome to form the final complex. The largest partial free energy change for hTBP, $-8.7$ kcal mol$^{-1}$, occurs in the initial binding step. hI$_2$ is 2.0 kcal mol$^{-1}$ lower in free energy than hI$_1$, whereas yI$_2$ is significantly less stable than yI$_1$. The corresponding equilibrium constant is thus 100× larger for the complex incorporating the human protein. Transition states are denoted (‡), and arrows show progressive changes in $\Delta H^c$. 
Figure 5. Comparison of the mole fractions of the hTBP and yTBP species at optimal in vivo temperatures. The time-dependent populations of the three TBP–DNA$_{AdMLP}$ species formed during association with the human protein at 37 °C (A) and the yeast protein at 30 °C (B) were simulated using 10 µM DNA$_{AdMLP}$ and 10 µM protein and previously collected yTBP data (18). The species-dependent differences are apparent in the rates of complex formation overall and in the comparative evolution of I$_1$, I$_2$, and the final conformer (F) for each protein.
Figure 6. Comparative T*AdMLP_{dp{x}*F bending by yTBP_{wt} (dotted line) and hTBP_{wt} (broken line). Time-resolved FRET provides a rigorous approach to the determination of the structure of TBP–DNA_{TATA} complexes in solution, yielding emission decays from which the probability distribution of the 5'TAMRA–3'fluorescein distance can be precisely determined (5, 14-16, 23, 24). Further, the sensitivity of measurements at ~60 Å with this dye pair is such that a 1° change in the bend angle results in a 1% change in the observed emission intensity. A high degree of confidence is thus ascribed to the difference measured for the hTBP and yTBP induced bends, with the latter redetermined herein under identical conditions using yTBP prepared as described (22). yTBP-bound T*AdMLP_{dp{x}*F has an R = 52.2 Å with σ = 8.9 Å. Bend angles (α) were obtained from these data using a simple two-kink bending model (5, 14) and the method of moments (24).

Figure 7. In vitro transcription efficiency (38) is strongly correlated with the degree of the TBP-induced bend for both the human and yeast proteins. Experimentally measured bends induced by yTBP in AdMLP (solid square) and the A3, T6, C7, G6, and T5 AdMLP variant TATA sequences (open circles in order from bottom left) have been shown previously to correlate with both in vitro and in vivo transcription activity (5). The high relative transcription activity of hTBP–DNA_{AdMLP}, 172% (38), and the mean hTBP-induced AdMLP bend angle of 97° (solid triangle) extend this trend ($R^2 = 0.957$). The yTBP–DNA_{AdMLP} bend angle was redetermined herein to ensure comparability.