Molecular basis of the recognition of the ap65-1 gene transcription promoter elements by a Myb protein from the protozoan parasite *Trichomonas vaginalis*

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ABSTRACT

Iron-inducible transcription of the ap65-1 gene in *Trichomonas vaginalis* involves at least three Myb-like transcriptional factors (tvMyb1, tvMyb2 and tvMyb3) that differentially bind to two closely spaced promoter sites, MRE-1/MRE-2r and MRE-2f. Here, we defined a fragment of tvMyb2 comprising residues 40–156 (tvMyb240–156) as the minimum structural unit that retains near full binding affinity with the promoter DNAs. Like c-Myb in vertebrates, the DNA-free tvMyb240–156 has a flexible and open conformation. Upon binding to the promoter DNA elements, tvMyb240–156 undergoes significant conformational re-arrangement and structure stabilization. Crystal structures of tvMyb240–156 in complex with promoter element-containing DNA oligomers showed that 5'-a/gACGAT-3' is the specific base sequence recognized by tvMyb240–156, which does not fully conform to that of the Myb binding site sequence. Furthermore, Lys49, which is upstream of the R2 motif (amino acids 52–102) also participates in specific DNA sequence recognition. Intriguingly, tvMyb240–156 binds to the promoter elements in an orientation opposite to that proposed in the HADDOCK model of the tvMyb135–141/MRE-1-MRE-2r complex. These results shed new light on understanding the molecular mechanism of Myb–DNA recognition and provide a framework to study the molecular basis of transcriptional regulation of myriad Mybs in *T. vaginalis*.

INTRODUCTION

The Myb proteins consist of a large family of proteins with diverse functions. These proteins have been demonstrated to play key roles in various important biological functions, including roles as regulators of stem and progenitor cells in the bone marrow, colonic crypts and a neurogenic region of the adult brain; they have also been identified as oncogenes involved in some human leukemias (1). In plants, Mybs were found to be involved in the control of plant-specific processes including metabolism, cell fate and identity, developmental processes and responses to biotic and abiotic stresses (2). Most Myb proteins function as transcription factors by binding to DNA via highly conserved Myb domains that recognize the t/cAACt/gG sequence, known as the Myb binding site (MBS) (3,4). The Myb domain generally consists of up to four imperfect amino acid sequence repeats (R) of about 50 amino acids, which form three α-helices. The second and third helices of each repeat form a helix turn helix (HTH) structure. The third helix of the second (R2) and third repeat (R3) are the recognition helices that bind to the major groove cooperatively and recognize the specific DNA sequence (3,5).
Trichomonas vaginalis is a protozoan parasite that causes trichomoniasis, one of the most common sexually transmitted diseases in humans (6). The ap65-1 gene from T. vaginalis encodes a 65-kDa protein, which upon iron repletion has been reputed to be one of the surface adhesins that mediate cytoadherence of the parasite to vaginal epithelial cells (7–9). We have shown that transcription of the ap65-1 gene is regulated in cis by two neighboring MBS, MRE-1/MRE-2r and MRE-2f, along with several other neighboring DNA regulatory elements in the ap65-1 promoter. The MRE-1/MRE-2r site comprises three overlapping DNA elements, which are the binding sites for three Myb-like proteins: tvMyb1, tvMyb2 and tvMyb3 (Figure 1A) (10–12). tvMyb1 and tvMyb2 each comprises a R2R3 class, whereas tvMyb3 comprises a R1R2R3 class of the MBS domain and shorter but highly variable N- and C-termini (Figure 1B) (12,13). The constituatively expressed tvMyb1 may repress basal or iron-inducible, while enhance growth related, ap65-1 transcription through differential bindings to the entire MRE-1/MRE-2r and/or MRE-2f (13). tvMyb2, which shares 33% identity with tvMyb1, may enhance ap65-1 transcription by bindings to the MRE-2r moiety of MRE-1/MRE-2r and MRE-2f. It may preferentially target the ap65-1 promoter in iron-replete cells to iron-depleted cells at an early growth stage when its expression is low, but the preference shifts to iron-depleted cells at a later growth stage when its expression is high (14). tvMyb3, which binds only to the MRE-1 moiety of MRE-1/MRE-2r, may display a differential promoter selection antagonistic to that of tvMyb2, to activate transcription (12). It is intriguing why and how this simple protozoan without noticeable differentiation uses so many distinct Myb proteins for transcription.

The solution structure of a truncated form of tvMyb1 (tvMyb135–141) was previously solved by nuclear magnetic resonance (NMR), and the structural basis for its interaction with the promoter. The MRE-1/MRE-2r DNA duplex was modeled by a data-driven HADDOCK software (15). However, unambiguous intermolecular nuclear overhauser effects (NOE) between tvMyb135–141 and DNA could not be assigned; thus, the detailed interaction between tvMyb1 and the promoter DNA, as well as the molecular basis of sequence-specific recognition, remains elusive. Here we report the identification and dynamics characterization of a tvMyb2 fragment comprising amino acid residues 40–156 (tvMyb240–156) as the minimum structural unit that retains near full binding affinity with the promoter DNA. We also report the X-ray crystal structures of tvMyb240–156 in complex with DNA promoter elements. Structural comparison showed a remarkable similarity between c-Myb and tvMyb240–156, as well as features unique to tvMyb2.

MATERIALS AND METHODS

Cloning, protein expression and purification

A DNA fragment containing the coding region of the full-length T. vaginalis Myb2 (designated tvMyb2) gene was cloned by polymerase chain reaction (PCR). The amplified DNA was digested with BamHI and SmaI and was inserted into the pGEX-6p3 vector (GE) containing a removable Glutathione-S-transferase (GST) tag at the N-terminus. The DNA encoding the truncated fragment comprising amino acid residues from 40 to 156 (designated tvMyb240–156) was amplified by PCR, and subcloned for ligation into the pET22b+ vector (Novagen), which contains a hexahistidine tag at the C-terminus. The forward primer was designed to contain an NdeI site and the reverse one a stop codon followed by an XhoI site.

The plasmids were each transformed into Escherichia coli strain BL21 star (DE3) competent cells (Invitrogen). The transformed cells were grown in 1 liter LB media supplemented with 50 μg/ml ampicillin in a 2.81 Fernbach baffled flask with vigorous shaking (~200 rpm) at 37°C. The cells expressing tvMyb2 were induced at OD600 ~0.8 by adding IPTG to a final concentration of 0.1 mM, and the incubation was continued for another 4 h. tvMyb240–156 was similarly expressed except for a final concentration of IPTG to be 1 mM and post-induction incubation to be 6 h at 30°C.

The cells from 11 culture were harvested by centrifugation, washed twice with 50 ml of phosphate buffered saline (PBS) containing 1 mM EDTA and 1 mM DTT, resuspended in 25 ml of a lysis buffer (50 mM Na3PO4, 500 mM NaCl, 1 mM EDTA) containing 1x Complete Protease inhibitor cocktail (Roche) and lysed with a microfluidizer (Microfluidics, MA, USA). To purify the GST–tvMyb2 fusion protein, the supernatant was incubated with gentle agitation at 4°C for 1 h in 10 ml of Glutathione Sepharose 4B resin (GE) in PBS and washed with 200 ml of a washing buffer (PBS containing 500 mM NaCl, 1 mM EDTA). The GST–tvMyb2 protein was cleaved from the resin by adding 20 μl of the Precision Protease to the resin in 25 ml of a cleavage buffer (50 mM Tris–HCl, pH 7, 150 mM NaCl, 1 mM EDTA, 1 mM DTT) at 4°C for 24 h.

To purify the His–tvMyb240–156, the supernatant was loaded to a column packed with 10 ml of Profinity Ni-charged IMAC resin (Bio-rad) equilibrated in a lysis buffer (50 mM Na3PO4, 500 mM NaCl, 0.1 mM EDTA, pH 8.0). After 1 h incubation at 4°C, the column was washed with 200 ml of a washing buffer (50 mM Na3PO4, 500 mM NaCl, 20 mM imidazole, pH 8.0) to remove unbound proteins. The His–tvMyb240–156 protein was eluted with 50 ml of an elution buffer (50 mM Na3PO4, 300 mM NaCl, 250 mM imidazole, pH 8.0).

The eluted proteins were further purified through a Superdex75 size exclusion column (SEC) connected to an AKTA Explorer system (GE), and the protein authenticities were verified by peptide mass fingerprinting (PMF). To prepare the uniformly 15N,13C-labeled protein for NMR experiments, cells were grown in M9 minimal medium supplemented with 15NH4Cl (1 g/l), 13C-glucose (2 g/l) and 15N,13C-isogro (Sigma-Aldrich). DNA oligonucleotides containing MRE-1/MRE-2r or MRE-2f promoters were purchased from Genomics Inc. (Taiwan) (Figure 1C). The concentrations of proteins and double-stranded DNAs (dsDNA) were determined by absorbance at 280 and 260 nm, respectively, with an
Figure 1. (A) The promoter organization of the \textit{ap}65-1 gene. (B) Amino acid sequence alignment of the R2R3 motif regions of \textit{tv}Myb1, \textit{tv}Myb2, \textit{tv}Myb3 and mouse \textit{c-myb} (SwissProt ID number P06876). The sequence is numbered according to that of \textit{tv}Myb2. The R2 (amino acids 52–102) and R3 (amino acids 103–154) motif regions are underlined. The stars identify the conserved aromatic residues and the dotted residues are the conserved residues participating in specific DNA sequence recognition. The helical regions of \textit{tv}Myb240–156 are also shown. (C) DNA 20-mer duplexes used for ITC studies. The sequences span the −81 or −100 and −31 or −50 regions of the promoter containing the MRE-1/MRE-2r and MRE-2f recognition sites, respectively. The red-colored bases are the 12-mer DNA sequences (MRE-1-12 and MRE-2-12) used for NMR studies. The red-colored bases plus the blue-colored bases in MRE-2-20 are the DNA sequences of the 13-mer DNA (MRE-2-13) used for crystallography studies. For the convenience of structural comparison, the bases in both MRE-1-20 and MRE-2-20 are numbered such that the specific sequence recognized by \textit{tv}Myb2 (i.e. A1C2G3A4T5A6) is numbered the same in the two oligomers. (D) Superposition of the $^{15}$N-HSQC spectra of \textit{tv}Myb240–156 (red) and \textit{tv}Myb240–156/MRE-1-12 complex (black).
Isothermal titration calorimetry

Isothermal titration calorimetry (ITC) was performed with an ITC200 calorimeter (MicroCal Inc.) at 25°C. Protein and DNA solution were dialyzed overnight against the same reaction buffer (20 mM Na3PO4, 150 mM NaCl, 1 mM EDTA, pH 7.4). The titration was carried out by injecting 1 μl (first injection) or 2.5 μl (2nd to 15th injection) of rTMyb240-156 solution at 100 μM concentration into the sample cell filled with 10 μM DNA solution. The initial delay was 300 s, with a 120 s interval between two successive injections. Since the $T_m$ values of MRE-1-12 and MRE-2-12 are near room temperature, the longer DNA duplexes of MRE-1-20 and MRE-2-20 were used for ITC studies. ITC binding curves were fitted to the single-site binding equation in the Microcal Origin software package, with the last two points taken as dilution heat.

NMR experiments

The NMR spectra were acquired on Bruker AV500, AV600 or AV800 spectrometers equipped with 5-mm triple resonance cryoprobe and a single-axis pulsed field gradient at 310 K. NMR data were acquired in Shigemi tubes on 0.7 mM protein samples in 20 mM Na3PO4, 50 mM NaCl, 1 mM EDTA, 1 mM DTT, 10% (v/v) D$_2$O at pH 6. The $^{15}$N,$^{13}$C-labeled rTMyb240-156-DNA complex samples were prepared at a protein–DNA molar ratio of 1:1.2. As 12-mer DNA duplexes gave the best NMR spectra, they were used for all NMR studies. Protein backbone resonance assignments were achieved by standard triple resonance experiments, including HNCA, HN(CA)CO, CBCA(CO)NH, HNCO and HN(CA)CO (16–20). Aliphatic side chain assignments primarily involved HCCH-COSY and HCCH-TOCSY, with the help of complementary HCC(CO)NH, (H)CC(CO)NH,(H)CB(CGCD)HE experiments (21–23). $^{13}$N-edited TOCSY-HSQC acquired with 60-ms mixing time was used to resolve ambiguity due to spectral overlap. Aromatic side chains were assigned with the use of proton homo-nuclear TOCSY and NOESY experiments, with additional verification from $^{13}$C-HMQC, (HB)CB(CGCD)HD and (HB)CB(CGCDCE)HE experiments (24,25). $^1$H chemical shifts were externally referenced to 0 ppm methyl resonance of 2,2-dimethyl-2-silapentane-5-sulfonate (DSS), whereas $^{13}$C and $^{15}$N chemical shifts were indirectly referenced according to the recommendations of the International Union of Pure and Applied Chemistry (26). The NMR spectra were processed using Bruker TOPSPIN 2.0 and analyzed by Sparky (27) and CARA (Available from http://www.nmr.ch).

$^{15}$N-T$_1$, $^{15}$N-T$_2$ and [{H$^{13}$N}] NOE were determined by using standard pulse sequences (28). Ten $^{15}$N spin–lattice/longitudinal relaxation rate constant (T$_1$) experiments were performed in random order, with relaxation delays of 0, 195.6, 301.0, 496.6 (duplicate), 707.3, 1098.5 (in duplicate), 1610.2 and 2197.1 ms. Similarly, spin–spin/transverse relaxation rate constant (T$_2$) experiments were performed randomly with relaxation delays of 0 (in duplicate), 15.8, 31.5, 47.3 (duplicate), 63.0, 78.8, 94.5 and 126.0 ms. Both rate constants were determined using the program Curvefit, assuming mono-exponential decay of the peak intensities. The errors in peak intensities were calculated from two duplicate experiments. The steady-state heteronuclear [{H$^{13}$N}] NOE experiment was carried out in duplicate in an interleaved manner, with and without proton saturation. The NOE was calculated as the error-weighed average ratio of peak intensities, with error estimate by propagating the base-plane noise. The reduced spectral density analysis was performed as previously described (29–35).

To measure one-bond $^1$H–$^{15}$N residual dipolar couplings (RDC), rTMyb240-156 was partially aligned in diluted liquid crystalline phase of 5% C12E5 polyethylene-glycol/n-hexanol (r = 0.96) (36). The rTMyb240-156/MRE-1-12 complex was partially aligned in liquid crystalline phase containing 15 mg/ml Pf1 bacteriophage (Asla Biotech Ltd, Latvia). Changes in splitting relative to the isotropic $^{1}$J$_{NH}$ values were measured using DSSE-HSQC experiments to obtain one bond $^1$H–$^{15}$N RDC (37). The measured RDCs were analyzed using the program PALES (38).

Crystallization of protein–DNA complex

Samples of protein–DNA complex were prepared by mixing rTMyb240–156 or Se-Met–rTMyb240–156 with DNAs at 1:1 molar ratio at a final concentration of 15 mg/ml in a buffer containing 20 mM Tris–HCl (pH 7.5) and 100 mM NaCl. The mixtures were further purified through Superdex75 size exclusion column. The 12-mer duplex DNA with a sequence of 5'-ATAACGATTT-3' (MRE-1-12) and the 13-mer duplex DNA with a sequence of 5'-CTGTATCGTCTTG-3' (MRE-2-13) (Figure 1C) were purchased from Genomics, Inc. (Taiwan). The rTMyb240–156/MRE-1-12 and rTMyb240–156/MRE-2-13 complexes produced best quality crystals for structural analysis. The crystallization experiments were carried out at 22°C by the vapor-diffusion method. The crystals of the rTMyb240–156/MRE-2-13 complex were obtained by mixing 1 μl of the protein–DNA complex solution and 1 μl of a crystallization solution composed of 0.1 M sodium cacodylate (pH 6.0), 15% isopropanol, 25% PEK4000 and 10% glycerol. The crystals of the Se-Met rTMyb240–156/MRE-2-13 complex were obtained by mixing 1 μl of the protein solution with 1 μl of a solution contain 0.2 M ammonium nitrate and 2.2 M ammonium sulfate and the crystals of the rTMyb240–156/MRE-1-12 complex were obtained by mixing 1 μl protein solution with 1 μl of a solution contain 0.1 M HEPES (pH 7.0) and 1.5–1.7 M ammonium sulfate.

Data collection and structure determination

The crystals of the Se-Met rTMyb240–156/MRE-2-13 complex were flash-cooled in liquid nitrogen without a cryo-protectant, whereas the crystals of the rTMyb240–156/MRE-1-12 complex were flash-cooled in the liquid nitrogen using 20% glycerol as a cryo-protectant. Data
were collected on a quantum 315 CCD detector at beamlines 13B and 13C of the National Synchrotron Radiation Research Center (NSRRC) in Taiwan, the Republic of China and the Mar22SHE detector at beamline 44XU of the Super Photon Ring-8 Synchrotron Radiation Center in Japan. Data were processed and scaled with HKL2000 (39). The structure of the Se-Met-Myb240–156/MRE-2-13 complex was determined by Se-MAD phasing. Four of the expected selenium sites were found, and the initial phase was calculated by the use of SOLVE (40). Density modification and auto-model building were performed with RESOLVE (40). The structure of the tvMyb240–156/MRE-1-12 complex was determined by molecular replacement with the program Phaser in PHENIX (41,42), with the solved structure of the tvMyb240–156/MRE-2-13 complex as a model. Iterative cycles of manual model building in COOT (43) and refinement in PHENIX yielded the final model. Table 2 gives the refinement statistics. Overall geometry of the model was assessed by Molprobity (44,45). The structures were analyzed by the use of PyMol (PyMOL Molecular Graphics System, v1.3, Schrödinger, LLC.) The structure factors and coordinates were deposited in the Protein Data Bank under (PDB) under PDB ID codes 3OSF (tvMyb240–156/MRE-2-13) and 3OSG (tvMyb240–156/MRE-1-12).

RESULTS

Mapping the minimum structured fragment of tvMyb2

Full-length tvMyb2 (tvMyb2) is most likely to exist as monomer, as predicted from analytical ultracentrifugation (Supplementary Figure S1A). However, the molecular weight of 36.1 kDa deduced from size exclusion chromatogram is much larger than the 21 kDa calculated from the sequence (Supplementary Figure S1B), suggesting that the protein may not be folded as a compact globular shape. To map the structured region essential for DNA binding, we performed a Predictor of Natural Disordered Regions (PONDR) prediction of the order-disorder profile (46) and showed that the N-terminal 50 amino acid and the C-terminal 20 amino acid are disordered (Supplementary Figure S1C). Gene fragments encoding various lengths of the tvMyb2 were constructed and expressed. The foldings of these fragments were assessed by 15N-HSQC NMR. The spectrum of tvMyb2 contains many well-dispersed peaks, (red peaks in Supplementary Figure S2), the characteristic of a folded protein, but many resonances were crowded in the 7.6–8.6 ppm region in the proton dimension (red peaks in the boxed region in Supplementary Figure S2), characteristics of a disordered protein. In contrast, the peaks in the spectrum of tvMyb240–156 are well dispersed (black peaks in Figure 1D). The addition of MRE-1-12 (red peaks in Figure 1D) or MRE-2-12 (data not shown) to tvMyb240–156 resulted in the disappearance of a set of resonances with the concurrent appearance of another set of resonances, the characteristics of a high-affinity DNA binding. Extending towards N- or C-terminus did not give more resonances outside of the 7.6–8.6 ppm region (data not shown), suggesting that tvMyb240–156 is the minimal fragment with an ordered structure. Circular dichroism studies showed that tvMyb240–156 is a helical protein (Supplementary Figure S1D).

NMR characterization of tvMyb240–156–promoter interaction

To gain insight into the structure and interaction with promoter DNAs, heteronuclear 3D NMR spectra of DNA-free tvMyb240–156 and tvMyb240–156 bound to MRE-1-12 or MRE-2-12 were obtained (Figure 1D), and the resonances were assigned and deposited in the Biological Magnetic Resonance Data Bank (accession number 17170). The amide resonances of Lys91–Ser97 in the 15N-HSQC spectrum were observed in the DNA–protein complexes, but not in the DNA-free protein form (Figure 1D). Further analysis of the relaxation data revealed that these residues reside in a flexible linker region with the structure of the DNA-free form much more flexible than that of the DNA-bound form (to be discussed later). The secondary structure of tvMyb240–156 was deduced from the consensus chemical shift index derived from the spectra of the promoter DNA-bound forms (47). The results showed that tvMyb240–156 consists of six helices, designated to be H1 (Glu55–Gly56), H2 (Trp71–Ala76), H3 (Ala83–Leu94), H4 (Ala105–Tyr118), H5 (Trp122–Phe127) and H6 (Ile137–Gly149) (Supplementary Figure S3).

To probe the Myb binding site, we further determined the changes of DNA binding-induced chemical shift, Δδ, as calculated from the equation \[\delta_{\text{NH}}^2 + (0.154\Delta\delta_{\text{N}})^2\] \((1/2)\), where δ\text{NH} is the chemical shift change of the amide proton and δ\text{N} is the chemical shift change of the amide 15N (Figure 2) (48). The chemical shift perturbation patterns induced by binding of MRE-1-12 or MRE-2-12 are nearly identical. The residues with Δδ > 0.2 ppm clustered around the N-terminus; H2–H3 linker, H3–H4 linker and the H5 and H6. To ensure that the 12-mer DNAs were sufficient for protein recognition, we also obtained 1H,15N-HSQC spectra of (u-15N)-tvMyb240–156 in complex MRE-1-20 or MRE-2-20 (data not shown). Although the spectral quality of these two complexes were not as good, nevertheless, their chemical shift perturbation patterns were practically identical to those observed for the corresponding 12-mers, suggesting that the bases beyond the 12-mer region are not essential for interaction with tvMyb240–156.

Promoter DNA-binding affinity and thermodynamics of tvMyb2 variants

MRE-1-12 and MRE-2-12 are not suitable for tvMyb2 binding study by ITC since their thermal melting temperatures are close to room temperature. Thus, MRE-1-20 and MRE-2-20 were used for ITC studies. As shown in Figure 3 and Table 1, tvMyb2 and tvMyb240–156 bind to MRE-1-20 of similar affinities with \(K_D = 100 \pm 35\) nM and 110 ±23 nM, corresponding to binding free energy of ΔГ = −9.7 and −9.6 kcal/mol, respectively. In contrast, the binding affinity towards MRE-2-20 was higher for tvMyb2 (\(K_D = 12.4 \pm 2.6\) nM) than tvMyb240–156 (KD = 32.3 ± 6.1 nM), suggesting that amino acid
residues flanking tvMyb240–156 also contribute to the binding of tvMyb2 to MRE-2-20. Nonetheless, the difference in their binding energies is small compared with the overall binding energy (−10.8 and −10.3 kcal/mol for tvMyb2 and tvMyb240–156, respectively). Therefore, tvMyb240–156 was used for structural studies by NMR and X-ray crystallography. This fragment encompasses the R2 and R3 motifs corresponding to vertebrate c-Myb. Interestingly, both entropy and enthalpy changes at room temperature contributed to a similar free energy of ~4.8 kcal/mol toward the binding of tvMyb2 and tvMyb240–156 to MRE-1-20, whereas the binding to MRE-2-20 was primarily enthalpy driven, with a small unfavorable entropy change.

Crystal structure of tvMyb240–156 in complex with MRE-1-12

The crystal structure of tvMyb240–156 in complex with MRE-1-12 was determined by molecular replacement, using the structure of the tvMyb240–156/MRE-2-13 complex as a template and refining it to 2.0 Å resolution (Figure 4A and B). Table 2 shows the diffraction parameters and refinement statistics. The protein was crystallized in P2₁ symmetry. Each asymmetric unit consisted of two tvMyb240–156–DNA complexes of practically identical structures, with an rmsd of 0.464 Å for the two tvMyb240–156 proteins. The structure of tvMyb240–156 consists of six α-helices, denoted H1 (Pro54–His67), H2...
Table 1. Thermodynamic data of tvMyb240–156- and mutant tvMyb240–156-promoter DNA binding deduced from ITC

| Sample                        | ΔH (cal/mol) | −TΔS (cal/mol) | K_D (or K_D^m) (nM)^a | K_D^m / K_D^b | ΔG (kcal/mol) |
|-------------------------------|--------------|----------------|------------------------|---------------|---------------|
| tvMyb2/MRE-1-20               | −4819 ± 63   | −4792 ± 213    | 100 ± 35               | –             | −9.61         |
| tvMyb2/MRE-2-20               | −11 730 ± 28 | 939 ± 102      | 12.4 ± 2.6             | –             | −10.79        |
| tvMyb240–156/MRE-1-20        | −4485 ± 250  | −5040 ± 360    | 110 ± 23               | –             | −9.53         |
| tvMyb240–156/MRE-2-20        | −11 240 ± 160| 1017 ± 45      | 32.3 ± 6.1             | –             | −10.22        |
| K49A/MRE-1-20                 | −3571        | −2795          | 21 459                 | 195           | −6.37         |
| K49A/MRE-2-20                 | −9495        | 1743           | 2079                   | 64            | −7.75         |
| K51A/MRE-1-20                 | −4014        | −3129          | 5680                   | 52            | −7.14         |
| K51A/MRE-2-20                 | −11 680      | 3457           | 962                    | 30            | −8.22         |
| R84A/MRE-1-20                 | n/a          | n/a            | n/a                    | n/a           | n/a           |
| R84A/MRE-2-20                 | −1628        | −4669          | 21 142                 | 654           | −6.3          |
| R87A/MRE-1-20                 | −4740        | −1833          | 15 200                 | 138           | −6.57         |
| R87A/MRE-2-20                 | −9694        | 1842           | 1750                   | 54            | −7.85         |
| K138A/MRE-1-20                | −1464        | −6020          | 3247                   | 33            | −7.48         |
| K138A/MRE-2-20                | −5588        | −2175          | 2033                   | 63            | −7.76         |
| N139A/MRE-1-20                | −5004        | −1910          | 8547                   | 78            | −6.91         |
| N139A/MRE-2-20                | −10 250      | 2314           | 1550                   | 48            | −7.94         |
| F52A/MRE-1-20                 | −17 290      | 11 115         | 30 700                 | 279           | −6.18         |
| F52A/MRE-2-20                 | −21 620      | 14 215         | 3970                   | 123           | −7.41         |

^K_D^m, the dissociation constants of wild-type (K_D) tvMyb240–156; K_D^m, the dissociation constant of the mutant tvMyb240–156.

|^aK_D|^b, the ratio of the dissociation constants of mutant tvMyb240–156 (K_D^m) and the wild-type tvMyb240–156 (K_D). It represents the factor in reduced binding affinity.

^cNo binding of R84A to MRE-1-20 can be detected by ITC.

Figure 3. ITC studies of the binding of tvMyb240–156 with MRE-1-20 (A) or MRE-2-20 (B). ITC traces are shown on the upper panels, and the binding isotherms are fit with a single exponential shown on the corresponding lower panels.
(Trp71–Thr77), H3 (Ala83–Leu94), H4 (Ala105–Tyr118), H5 (Trp122–Phe128) and H6 (Asp134–Leu148). The secondary structure assignments are almost identical to those determined by NMR in solution, with the exception of H6, which starts at Ile137 in contrast to Asp134 in solution. The six helices are arranged in two triple-helix bundles, with H1–H3 in the R2 motif forming the first bundle and H4–H6 within the R3 motif forming the second bundle. The two triple-helix bundles connected by a 10-residue linker comprising residues Ala95–Thr104 (designated the R2–R3 linker) assume a similar structure fold. The conserved aromatic groups are buried inside the hydrophobic cores of the helix bundles (Figure 4C and D). tvMyb240-156 interacts with DNA bases by positioning the third long helix of each motif, H3 and H6, on the major groove in a HTH fashion. The DNA binding pocket is lined with numerous positively charged residues, including residues Arg120–Asn124 at the N-terminal side of H5 (Figure 4E and F). Helices H1, H2 and H4 make no contact with the DNA molecule.

Figure 5A depicts molecular interactions between tvMyb240-156 and MRE-1-12. The interactions can be categorized into four types. First, there are the direct base interactions: Lys49 (NZ)-T3(O2) and T5(O2), Arg84-
Table 2. Statistics for data collection and structural refinement

| Data collection | Myb2/MRE-1-12 | Myb2/MRE-2-13 | Se-Met-Myb2/MRE-2-13 |
|-----------------|---------------|---------------|---------------------|
| Wavelength (Å)  | 0.97622       | 0.97622       | 0.97884             |
| Space group     | P2₁           | P2₁          | P2₁                 |
| Unit cell (Å)   | 40.1, 127.5, 41.4 | 73.9, 73.3, 84.4 | 73.4, 77.1, 84.1     |
| β = 100.2       |               |               |                     |
| Resolution range (Å) | 30–2.0 (2.07–2.0) | 50–2.03 (2.07–2.03) | 30–2.24 (2.32–2.24) |
| Unique reflections | 26911 (2556)  | 31 586 (1442) | 23 625 (2241)       |
| Redundancy      | 3.5 (3.2)     | 4.9 (4.7)     | 4.7 (4.5)           |
| Completeness (%) | 97.3 (93.7)   | 99.1 (93.1)   | 98.3 (95.7)         |
| Rmerge (%)b     | 5.0 (37.8)    | 4.7 (26.9)    | 6.7 (53.3)          |

Rigidity

| Reflections (F > 0 σF) | 26037 (2191) | 20.6 (25.7) |
|------------------------|--------------|-------------|
| Rcryst (%) for 95% data | 19.0 (23.9)  | 23.3 (32.3) |
| Rfree (%) for 5% data  | 20.6 (25.7)  | 25.8 (31.6) |

RMSD

| Bond lengths (Å) | 0.008 | 0.007 |
|------------------|-------|-------|
| Bond angles (°)  | 1.40  | 1.39  |

Average B-factors (Å²)

| Protein       | 39.9  | 38.0  |
|---------------|-------|-------|
| DNA           | 29.8  | 37.9  |
| Water         | 36.7  | 42.2  |
| Isopropanol   | –     | 42.9  |

Values in parentheses are for the highest resolution shell.

Rmerge = Σ|Fo|−|Fc|/Σ|Fc|.

The crystal structure of Se-Met-νMyb240–156 in complex with MRE-2-13 was determined by the multiple-wavelength anomalous diffraction method and refining it to 2.03 Å resolution (Table 2). The protein was crystallized in P2₁,2₁,2₁ symmetry with each asymmetric unit also consisting of two νMyb240–156/MRE-2-13 complexes with indistinguishable structures. Although the recognition sequence of MRE-2f (5'-TATCGT-3') differs from that of MRE-1/MRE-2r (5'-ACGATA-3'), its complementary strand is identical to that of MRE-1/MRE-2r, and vice versa. Thus, the structures of νMyb240–156 in the two complexes are practically identical, with an rmsd of 0.322 Å. Furthermore, the residues involved in specific DNA sequence recognition, including Lys89 are identical. However, there is an additional solvent-mediated base interaction, i.e. Lys89(NZ)-C4(O2) observed in this complex (Figure 5C and D), which together with the results of MRE-1-12 indicates that 5'-gACGAT-3' is the specific base sequence recognized by νMyb240–156.

Nonetheless, a closer inspection of the structures showed that νMyb240–156 has more extensive interactions with MRE-2-13 than with MRE-1-12, consistent with the observed binding affinity by ITC. Three solvent-mediated interactions beyond the promoter region, namely Lys87(NZ)-C4(O2), Arg87(NH2)-A3(N7) and Arg87(NH2)-G3(O4) were observed in the MRE-2-13 complex, whereas only the Arg87(NH2)-A3(N7) interaction was observed in MRE-1-12/νMyb240–156 complex. The Asn139(OD1)-T4(O4) interaction was replaced by the Lys138(NZ)-T4(O4) interaction. Moreover, we observed five protein-DNA backbone phosphate interactions in the MRE-2-13 complex but only three in the MRE-1-12 complex.

Crystal structure of νMyb240–156 in complex with MRE-2-13

The crystal structure of Se-Met-νMyb240–156 in complex with MRE-2-13 was determined by the

G5(O6 and N7), Lys138(NZ)-G4(O6) and Asn139(ND2)-A3(N7) (Figure 5B). These interactions are responsible for the sequence-specific recognition between νMyb240–156 and MRE-1/MRE-2r. The interactions of Lys49(NZ) with T3(O2) and T4(O2), as well as Lys85(N) and Glu50(N) with the backbone phosphate groups stabilize the N-terminal loop. These interactions involving residues at the N-terminal side of H1 are unique to the νMyb2 protein. Second are the solvent-mediated base interaction: Arg87(NH2)-A-1(N7), Asp88(OD2)-C4(O2) and Lys49(N) and Gln50(N)-Gln50(NE2) with the backbone phosphate groups stabilize the recognition sequence of MRE-2f (5'-TATCGT-3'), its complementary strand is identical to that of MRE-1/MRE-2r, and vice versa. Thus, the structures of νMyb240–156 in the two complexes are practically identical, with an rmsd of 0.322 Å. Furthermore, the residues involved in specific DNA sequence recognition, including Lys89 are identical. However, there is an additional solvent-mediated base interaction, i.e. Lys89(NZ)-C4(O2) observed in this complex (Figure 5C and D), which together with the results of MRE-1-12 indicates that 5'-gACGAT-3' is the specific base sequence recognized by νMyb240–156.

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Energetic contribution to specific DNA sequence recognition of the base-recognition residues

To corroborate the structural studies and to further determine the energetic contribution of residues responsible for specific DNA sequence recognition we generated six alanine mutants (K49A, K51A, R84A, R87A, K138A and R139A) and determined their promoter DNA-binding affinities by ITC (Table 1). The mutation of any of these residues to alanine drastically reduced the binding affinity and in general the mutation has greater effect on the binding to MRE-1-20 than it does to MRE-2-20. The reduction in the binding affinity of mutant tvMyb240-156 follows the following order: R84A > K49A > R87A > N139A > K138A > K51A. In particular, the mutation of Arg84 to alanine completely abolished protein binding to MRE-1-20 and reduced the

Figure 5. Schematic representation of the detailed interactions between tvMyb240–156 and MRE-1-12 (A) or MRE-2-13 (C). The corresponding pair-wise interactions of tvMyb240–156 with MRE-1-12 and MRE-2-13 are shown in (B) and (D), respectively. There are two monomers (a and b) in an asymmetric unit. The blue, red and green dotted lines indicate interactions observed in both monomers, monomer a and monomer b, respectively. W indicates a water molecule.

of any of these residues to alanine drastically reduced the binding affinity and in general the mutation has greater effect on the binding to MRE-1-20 than it does to MRE-2-20. The reduction in the binding affinity of mutant tvMyb240–156 follows the following order: R84A > K49A > R87A > N139A > K138A > K51A. In particular, the mutation of Arg84 to alanine completely abolished protein binding to MRE-1-20 and reduced the
binding affinity to MRE-2-20 by a factor of 654. Arg84 interacts with the T6-A1 and G5-C2 base pairs. The corresponding residue, Lys128 in c-Myb is also the most important residue in promoter recognition as the K128A mutant has lost the ability to bind cognate DNA (49). The significant role of the N-terminal residue Lys49 not located on the third helix of each motif is surprising and is not previously reported in other Myb proteins (49).

The amide resonance of Phe52 experienced the largest chemical shift perturbation upon binding to the promoter elements with its amide nitrogen and carbonyl group contacting the phosphate group between A(2) and T(3). In c-Myb, the indole ring of the corresponding Trp95 significantly shifted toward the cavity of the R2 motif on DNA binding (49). We found that the F52A mutant reduced the binding affinity toward MRE-1-20 and MRE-2-20 by a factor of 279 and 123, respectively, corresponding to ΔΔGs of 3.68 and 2.82 kcal/mol. The drastic reduction in binding energy illustrates the pivotal role of this hydrophobic residue in the structural integrity and DNA binding of the Myb family of proteins.

**Dynamics of tvMyb240–156**

Figure 6A–C show the backbone amide ^15^N-R1 (A), ^15^N-R2 (B) and [^1H-^15^N] NOE (C) of tvMyb240–156 (blue), the tvMyb240–156/MRE-1-12 complex (red) and the tvMyb240–156/MRE-2-13 complex (green). The calculated reduced spectral density functions J(0), J(60) and J(522) are shown in (D–F), respectively.

Figure 6. Sequence variation of ^15^N-R1 (A), ^15^N-R2 (B) and [^1H-^15^N] NOE (C) of tvMyb240–156 (blue), the tvMyb240–156/MRE-1-12 complex (red) and the tvMyb240–156/MRE-2-13 complex (green). The calculated reduced spectral density functions J(0), J(60) and J(522) are shown in (D–F), respectively.
functions, reflected in the low NOE values. For the DNA-free form residues in both complexes, probably due to disorder as homologous (an rmsd of 2.81 Å) (15,51). DNA binding features of c-Myb match quite well, but the H6 on R3 showed the relative orientation of R2 and R3 with little effect on the induced a significant structural rearrangement in the R2R3 region is now fully immobilized except for some of the residues in the loop regions connecting the helices.

For a globular protein, the rotational correlation time can be predicted from the empirical Stokes–Einstein relation, \( \tau_r = \frac{1}{k_B T/8\pi\eta^3} \), where \( k_B \) is the Bolzmann’s constant, \( T \) is the absolute temperature and \( \eta \) is solvent viscosity. The predicted \( \tau_r \) values are \( \sim 8.0 \) and \( \sim 4.1 \) ns for tvMyb240–156 and the isolated R2 or R3 motif, respectively (50). For a rigid globular molecule, the overall rotational correlation time, \( \tau_c \), can be obtained from spectral density functions, \( \tau_c = \frac{\alpha_N}{J(0) - J(o_N)}/J(o_N) \), as previously described (29,32). For DNA-free tvMyb240–156, we obtained a mean \( \tau_c = 5.3 \) ns for the more rigid region, Thr53 to Ala63, of the R2 motif and \( \tau_c = 6.0 \) ns for the R3 motif. These values indicate that the R2 and R3 motifs move with a correlation time of a globular protein of \( \sim 78 \) and \( 87 \) residues, respectively. Thus, the two motifs move with a considerable degree of independence. In contrast, we observed \( \tau_c \) of 10.3 ns and 10.4 ns for tvMyb240–156/MRE-1-12 and tvMyb240–156/MRE-2-13, respectively, consistent with the predicted correlation times of their sizes. In addition, the two motifs in the DNA complexes move with the same rotational correlation times. Thus, tvMyb240–156/MRE-1-12 and tvMyb240–156/MRE-2-13 move as monomeric rigid bodies.

**DISCUSSION**

**Solution structure of DNA-free Myb proteins**

TvMyb240–156 shares 26% sequence identity with the R2/R3 regions of vertebrate c-Myb and 40% sequence identity with the R2/R3 region of TvMyb1 (13) (Figure 1B). The solved structures of DNA-free c-Myb (PDB ID: 1GV2) and DNA-free tvMyb135–141 (PDB ID: 2K9N) are highly homologous (an rmsd of 2.813 Å) (15,51). DNA binding induced a significant structural rearrangement in the relative orientation of R2 and R3 with little effect on the folding of the two motifs. In fact, when superimposed, the R2 domains of the DNA free and DNA-bound structures of c-Myb match quite well, but the H6 on R3 showed a \( \sim 90^\circ \) rotation about the H3 (51). A similar change was observed in the DNA-free structure versus and the HADDOCK modeled DNA-bound structure of tvMyb135–141, but with a 50° reorientation of the two motifs (15).

We were unable to solve the solution structure of DNA-free tvMyb240–156 to high resolution by NMR due to insufficient NOE data, likely due to the high flexibility of the protein. To gain further insights on the structure transition upon DNA binding to tvMyb240–156, we have measured the \( ^1\text{H,}^{15}\text{N} \) RDC of DNA-free (u\(^{15}\text{N} \)) tvMyb240–156 and (u\(^{15}\text{N} \))tvMyb240–156/MRE-1-12 complex. Totally, 59 measured \( ^1\text{D}_{\text{NH}} \) RDCs for the helical region of the tvMyb240–156/MRE-1-12 complex were fit against the calculated values from the crystal structure of the tvMyb240–156/MRE-1-12 complex (Figure 7A). A low value of 0.21 RDC quality factor (Q-factor) was obtained, suggesting that the solution structure of the DNA-bound tvMyb240–156 is very similar to that in the crystal structure. For the DNA-free tvMyb240–156, the measured \( ^1\text{D}_{\text{NH}} \) values for the R2 motif (25 \( ^1\text{D}_{\text{NH}} \) RDCs) and R3 motif (32 \( ^1\text{D}_{\text{NH}} \) RDCs) individually fit well with the calculated values from the crystal structure of the individual motifs in the tvMyb240–156/MRE-1-12 complex (Q-factors of 0.23 and 0.32 for R2 and R3, respectively) (Figure 7B), suggesting that the structures of R2 and R3 are similar in the presence and absence of DNA. On the other hand, the measured \( ^1\text{D}_{\text{NH}} \) RDCs from R2 and R3 of tvMyb240–156 together fit poorly with the calculated RDCs from the crystal structure of the tvMyb240–156/MRE-1-12 complex (Q-factor = 1.04, Figure 7C). This suggests that the relative orientation of R2 and R3 in tvMyb240–156 changes substantially upon DNA binding. We also compared the measured RDC values with those predicted from two homology models of DNA-free tvMyb240–156, based on DNA-free c-Myb or DNA-free tvMyb135–141 (Figure 7D). The fit is also rather poor, suggesting that the structure of DNA-free tvMyb240–156 is very different from that of the other two DNA-free Myb proteins. Alternatively, because the DNA-free tvMyb240–156 is highly flexible, it is possible that the R2 and R3 motifs in DNA-free tvMyb240–156 are undergoing significant structural fluctuation so that the RDC values are further averaged.

To further corroborate the structural information, we conducted small angle X-ray scattering (SAXS) experiments to assess the structure of DNA-free tvMyb240–156. The radius of gyration (Rg) values, estimated from experimental curves using Guinier analysis (52), are 22.5 ± 0.09 Å and 18.6 ± 0.05 Å for DNA-free tvMyb240–156 and the tvMyb240–156/MRE-1-12 complex, respectively (Supplementary Figure S4). Thus, the Rg value of the DNA-free tvMyb240–156 is considerably larger than that in the DNA-bound form, suggesting that the DNA-free tvMyb240–156 assumes a much more open conformation. This conclusion is consistent with the poor fit of RDC data of DNA-free tvMyb240–156 with that predicted from the structures of tvMyb240–156/MRE-1-12.
Roles of protein flexibility in specific DNA sequence recognition in Myb proteins

Sarai et al. (53) have shown that the R2 motif of c-Myb is the least stable motifs with a thermal melting temperature of 43°C compared with 57°C for R3. The flexibility of the isolated motifs was further confirmed by NMR relaxation studies (3,51) (For a lack of better term, the word ‘flexibility’ here indicate a protein, or a portion of it, is not rigid and shows motion at various time scales as indicated by NMR relaxation parameters). However, the dynamics of the R2–R3 linker was not studied in c-Myb. Recently Lou et al. (15) also reported the presence of considerable flexibility in DNA-free tvMyb135–141, especially the R2 motif and the linker region. Here, we demonstrated that in the DNA-free state tvMyb240–156 is even more flexible than that of either c-Myb or tvMyb135–141, as indicated by the lower 1H-15NOE for a larger number of residues. Consistently, the linker region is the most flexible part of the molecule, and R2 is more flexible than R3. Upon binding to the cognate DNA, all three Myb proteins lost their flexibility.

The structure and dynamics information obtained so far suggest a unique molecular mechanism of Myb–DNA recognition. The DNA-binding pocket of the Myb proteins is lined with many positively charged residues that interact with numerous negatively charged backbone phosphate groups. Since Coulomb charge–charge interaction is a long range interaction, we suggest that the open and flexible structure of a Myb protein behaves much like an open net to catch the DNA molecule as the first step in DNA recognition. The intrinsic flexibility of the two
motifs and the linker region allow the protein to adapt to the optimal conformation in placing the H3 and H6 on the major groove in an induced-fit manner. Upon first contact, the properly folded protein could presumably slide along the DNA molecule until the conserved residues on the H3 and H6 make proper contact with the specific DNA bases without altering the conformation of the R2 and R3 motifs. Thus, the intrinsic flexibility of the Myb proteins is essential for DNA binding and sequence specific recognition in a manner resembling that proposed for the intrinsically disordered proteins (IDP) in molecular recognition (54–56). The importance of protein flexibility in DNA recognition has also been demonstrated recently in a papillomavirus E2 proteins (57).

Ogata attributed the source of the flexibility to the presence of a cavity in the center of the hydrophobic core of the R2 motif due to the presence of a small hydrophobic residue, Val103. The presence of flexibility in R2 allows the side chain of Trp95 to rearrange its packing for optimal DNA binding (3,51,53). The flexibility can be greatly reduced upon mutating the small Val residue to a larger Leu. Sequence alignment shows that Val103 in c-Myb is substituted by a larger Leu in tvMyb240–156. However, such a substitution does not reduce the flexibility of the R2 motif in the DNA-free tvMyb240–156, perhaps due to the simultaneous substitution of Trp95 in c-Myb by Phe252 in tvMyb240–156. In c-Myb the indole ring is significantly shifted toward the cavity of R2 on DNA binding. Such a rearrangement appears to be hampered by the absence of the cavity in the V103L of c-Myb, but it may not be a problem when Trp is simultaneously mutated to a smaller Phe in tvMyb2. Nonetheless, the important role played by the hydrophobic aromatic residue, Trp95 in c-Myb, was substantiated by our F52A study in which mutation of Phe252 to Ala reduced the binding affinity by two orders of magnitude.

Specific DNA sequence recognition in Myb proteins

In spite of the low sequence identity, the tertiary folds of the DNA-bound forms of tvMyb240–156 and c-Myb (PDB ID: 1MSE) are very similar (an rmsd of 2.3 Å) (49). Here, we showed that tvMyb240–156 recognizes the a/gACGAT sequence, which does not fully conform to the MBS sequence recognized by c-Myb. Nonetheless, the four conserved residues in the H3 and H6, namely Arg84, Asp138, Lys139 and Asn139 (as the Lys128, Glu132, Lys182 and Asn183 in c-Myb) and two unique residues, the N-terminal Lys49 and Arg87 in H3, are the major contributors in tvMyb240–156 for recognizing the specific DNA sequence. In c-Myb, specific base sequence recognition involves four other non-conserved residues: Asn136, Asn139, Asn186 and Ser187. The participation of residues outside of the R2R3 motifs in specific DNA sequence recognition is unique among the Myb proteins. It is likely that these unique base-recognition residues contribute to the differential recognition of the MBS by a particular Myb protein. The variation in recognition of MBS is conceivably important in transcription regulation of T. vaginalis since the parasite genome encodes more than 400 distinct Myb proteins (58).

Using the program HADDOCK (59,60), Lou et al. (15) also proposed a structural model of tvMyb135–141/MRE-1/ MRE-2f complex, which revealed the binding region in tvMyb135–141 to be similar to that identified in the crystal structures of tvMyb240–156–DNA complex reported here as well as that in c-Myb (49). However, there are two significant differences. First, tvMyb135–141 and tvMyb240–156 bind to the promoter element in opposite orientations (Supplementary Figure S5). Specifically, in the tvMyb135–141–DNA complex, both the H3 and H6 align in parallel to the 5′-A1C2G3A4T5A6-3′ orientation. In contrast, in tvMyb135–141/DNA complex the H3 and H6 are aligned anti-parallelly to the 5′-A1C2G3A4-3′ sequence orientation. As a result, tvMyb135–141 and tvMyb240–156 use different residues to recognize the same bases. Second, in tvMyb240–156 the N-terminal residues, Lys49 and Lys51, participate in DNA-specific sequence recognition but no interaction between the corresponding Lys residues with promoter DNA was observed in the tvMyb135–141/DNA complex. The HADDOCK structure was generated based on RDC, chemical shift perturbation and DNA specificity data, but without specific distance constraints, such as NOEs. It is known that two pairs of 1H–15N dipoles in opposite orientation have the same RDC and that chemical shift perturbation can only locate the binding site, but not the specific interacting pairs. Thus, two HADDOCK structures that bind to DNA in opposite orientations may both satisfy the HADDOCK constraints. In spite of the 40% sequence homology between tvMyb135–141 and tvMyb240–156, it is probable that tvMyb1 and tvMyb2 may bind to the MRE-1/MRE-2 in opposite orientations (21). This difference may have biological implications that are yet to be exploited. Intriguingly, examination of the structures shown on Figure 5A and Supplementary Figure S5A, revealed that DNA sequence-specific recognition of tvMyb1 involves two Asn residues (Asn110 with A2′ and Asn122 with C4′) that are not conserved and not involved in DNA sequence-specific recognition in tvMyb2 (The corresponding residues in tvMyb2 are Ala123 and Ile135, respectively) (Figure 1B). It is tempting to speculate that these specific recognition pairs might affect the binding orientation.

Biological implications

The genome of T. vaginalis encodes more than 400 Myb proteins, most of which share conserved R2R3 Myb domains with highly variable N- and C-termini (58). It is conceivable that they are the major regulators of gene transcription in T. vaginalis. To better understand the transcription regulation of the parasite, it is desirable to know how myriad tvMyb proteins select their target genes. Intriguingly, the DNA-binding specificity and nuclear translocation of tvMyb2 rely on a contiguous region spanning the entire R2R3 domain. A single point mutation at Ile74 to Ala, which slightly changes the helical content and tertiary folding of the R2R3 domain resulted in the inhibition of tvMyb2 nuclear import and
loss of DNA-binding activity. This suggests that the R2R3 domain is likely a common module for myriad Myb proteins in the parasite to regulate both DNA-binding specificity and nuclear import (Tai, J.H., unpublished data). Moreover, the nuclear import of Myb2 could be greatly facilitated in the presence of hydrogen peroxide (Tai, J.H., unpublished observation), suggesting an important role of this transcription factor in protecting the parasite from oxidative stress produced from host-immune surveillance. Consistent with this speculation, the Myb2 recognition sequence, MRE-1/MRE-2r, could be found in the promoter regions of several antioxidant genes, such as the genes encoding superoxide dismutase (accession number Gi123485668) and thioredoxin peroxidase (accession number Gi123459140).

It is therefore important to study the structural basis of tvMyb2 in both DNA-free form and DNA-bound forms. In summary, the structures of tvMyb240–156 in free form and in DNA-bound form were elucidated in this report, which reveals conserved and unique features of a particular tvMyb contributing to its recognition of specific DNA sequences. These characteristics are conceivably important in transcription regulation of T. vaginalis since the parasite genome encodes more than 400 distinct Myb proteins (58). Information derived from current study expands our understanding of the molecular basis of Myb protein function in general and for further study on the structure-functional relationship of Myb-regulated gene transcription in T. vaginalis in particular.

ACCESSION NUMBERS
3OSF, 3OSG, BMRB accession no. 17170.

SUPPLEMENTARY DATA
Supplementary Data are available at NAR Online.

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REFERENCES
1. Ramsay,R.G. and Gonda,T.J. (2008) MYB function in normal and cancer cells. Nat. Rev. Cancer, 8, 523–534.
2. Dubos,C., Stracke,R., Grotewold,E., Weisshaar,B., Martin,C. and Lepinec,L. (2010) MYB transcription factors in Arabidopsis. Trends Plant Sci., 15, 573–581.
3. Ogata,K., Kane-Ishii,C., Sasaki,M., Hatanaka,H., Nagadoi,A., Enari,M., Nakamura,H., Nishimura,Y., Ishii,S. and Sarai,A. (1996) The cavity in the hydrophobic core of Myb DNA-binding domain is reserved for DNA recognition and trans-activation. Nat. Struct. Mol. Biol., 3, 178–187.
4. Biedenkapp,H., Borgmeyer,U., Sippel,A.E. and Klenpmuier,K.-H. (1988) Viral myb oncogene encodes a sequence-specific DNA-binding activity. Nature, 335, 835–837.
5. Jia,L., Clegg,M.T. and Jiang,T. (2004) Evolutionary dynamics of the DNA-binding domains in putative R2R3-MYB genes identified from rice subspecies indica and japonica genomes. Plant Physiol., 134, 575–585.
6. Sorvillo,F., Smith,L., Kerndt,P. and Ash,L. (2001) Trichomonas vaginalis, HIV, and African-Americans. Emerg. Infect. Dis., 7, 927–932.
7. Alderete,J.F. and Garza,G.E. (1988) Identification and properties of Trichomonas vaginalis Myb proteins involved in cytoadherence. Infect. Immun., 56, 28–33.
8. Alderete,J.F., O’Brien,J.L., Arroyo,R., Engbring,J.A., Musatovova,O., Lopez,O., Lauriano,C. and Nguyen,J. (1995) Cloning and molecular characterization of two genes encoding adhesion proteins involved in Trichomonas vaginalis cytoadherence. Mol. Microbiol., 17, 69–83.
9. Arroyo,R., Engbring,J. and Alderete,J.F. (1992) Molecular basis of host epithelial cell recognition by Trichomonas vaginalis. Mol. Microbiol., 6, 853–862.
10. Tsai,C.-D., Liu,H.-W. and Tai,J.-H. (2002) Characterization of an iron-responsive promoter in the protozoan pPathogen Trichomonas vaginalis. J. Biol. Chem., 277, 5153–5162.
11. Ong,S.-J., Huang,S.C., Liu,H.W. and Tai,J.H. (2004) Involvement of multiple DNA elements in iron-inducible transcription of the ap65-1 gene in the protozoan parasite Trichomonas vaginalis. Mol. Microbiol., 52, 1721–1730.
12. Huo,H.-M., Ong,S.-J., Lee,M.-C. and Tai,J.-H. (2009) Transcriptional regulation of an iron-inducible gene by differential and alternate promoter entries of multiple Myb proteins in the protozoan parasite Trichomonas vaginalis. Eukaryot. Cell, 8, 362–372.
13. Ong,S.-J., Hsu,H.-M., Liu,H.M., Chu,C.H. and Tai,J.H. (2006) Multifarious transcriptional regulation of adhesion protein gene ap65-1 by a novel Myb1 protein in the protozoan parasite Trichomonas vaginalis. Eukaryot. Cell, 5, 391–399.
14. Ong,S.-J., Hsu,H.-M., Liu,H.W., Chu,C.H. and Tai,J.H. (2007) Activation of multifarious transcription of an adhesion protein gene ap65-1 by a novel Myb2 protein in the protozoan parasite Trichomonas vaginalis. J. Biol. Chem., 282, 6716–6725.
15. Lou,Y.C., Wei,S.Y., Rajasekaran,M., Chou,C.C., Hsu,H.M., Tai,J.H. and Chen,C. (2009) NMR structural analysis of DNA recognition by a novel Myb1 DNA-binding domain in the protozoan parasite Trichomonas vaginalis. Nucleic Acids Res., 37, 2381–2394.
16. Gardner,K.H. and Kay,L.E. (1998) The use of 2H, 13C, 15N multidimensional NMR to study the structure and dynamics of proteins. Annu. Rev. Biophys. Biomol. Struct., 27, 357–406.
17. Kanelis,V., Forman-Kay,J.D. and Kay,L.E. (2001) Multidimensional NMR methods for protein structure determination. J. Biomol. NMR, 25, 291–302.
18. Bax,A. (1994) Multidimensional nuclear magnetic resonance methods for protein studies. Curr. Opin. Struct. Biol., 4, 738–744.
33. Ishima, R. and Nagayama, K. (1995) Quasi-spectral density functions. Methods in Enzymology, Vol. 239. Elsevier, Waltham, MA, USA, pp. 79–105.

22. Lyons, B.A., Tashiro, M., Cedergren, L., Bilsson, B. and Montelione, G.T. (1996) An efficient triple resonance experiment using carbon-13 isotopic mixing for determining sequence-specific resonance assignments of isotopically-enriched proteins. J. Am. Chem. Soc., 118, 10974–10975.

4. Small, J. (1983) Sparky.

28. Arrowsmith, C.H. (2001) Latent and active p53 are identical in structure and function in protein interaction networks. Nat. Struct. Mol. Biol., 8, 363–392.

23. Montelione, G.T., Lyons, B.A., Emerson, S.D. and Tashiro, M. (1992) Two-dimensional NMR experiments for correlating positions of aromatic residues in 13C-labeled proteins via scalar couplings. J. Magn. Chem., 115, 11054–11055.

19. Cavanagh, J., Fairbrother, W.J., Palmer, A.G. and Skelton, N.J. (1996) Protein NMR Spectroscopy - Principles and Practice. Academic Press, San Diego.

24. Logan, T.M., Olejniczak, E.T., Xu, R.R. and Fesik, S.W. (1992) Side chain and backbone assignments in isotopically labeled proteins from two heteronuclear triple resonance experiments. FEBS Lett., 314, 413–418.

22. Lyons, B.A., Tashiro, M., Cedergren, L., Bilsson, B. and Montelione, G.T. (1993) An improved strategy for determining resonance assignments for isotopically enriched proteins and its application to an engineered domain of staphylococcal protein A. Biochemistry, 32, 7839–7845.

20. Sattler, M., Schleucher, J. and Griesinger, C. (1999) Heteronuclear multidimensional NMR experiments for the structure determination of proteins in solution employing pulsed field gradients. Prog. Nucl. Mag. Res. Sp., 34, 93–158.

32. Farrow, N.A., Zhang, O., Szabo, A., Torchia, D.A. and Kay, L.E. (1988) Determination of 15N chemical shift tensor and T2 values in proteins. J. Magn. Res., 78, 39–47.

25. Montelione, G.T., Lyons, B.A., Emerson, S.D. and Tashiro, M. (1992) An efficient triple resonance experiment using carbon-13 isotopic mixing for determining sequence-specific resonance assignments of isotopically-enriched proteins. J. Am. Chem. Soc., 114, 10974–10975.

20. Sattler, M., Schleucher, J. and Griesinger, C. (1999) Heteronuclear multidimensional NMR experiments for the structure determination of proteins in solution employing pulsed field gradients. Prog. Nucl. Mag. Res. Sp., 34, 93–158.

24. Logan, T.M., Olejniczak, E.T., Xu, R.R. and Fesik, S.W. (1992) Side chain and backbone assignments in isotopically labeled proteins from two heteronuclear triple resonance experiments. FEBS Lett., 314, 413–418.

22. Lyons, B.A., Tashiro, M., Cedergren, L., Bilsson, B. and Montelione, G.T. (1993) An improved strategy for determining resonance assignments for isotopically enriched proteins and its application to an engineered domain of staphylococcal protein A. Biochemistry, 32, 7839–7845.

20. Sattler, M., Schleucher, J. and Griesinger, C. (1999) Heteronuclear multidimensional NMR experiments for the structure determination of proteins in solution employing pulsed field gradients. Prog. Nucl. Mag. Res. Sp., 34, 93–158.

24. Logan, T.M., Olejniczak, E.T., Xu, R.R. and Fesik, S.W. (1992) Side chain and backbone assignments in isotopically labeled proteins from two heteronuclear triple resonance experiments. FEBS Lett., 314, 413–418.

22. Lyons, B.A., Tashiro, M., Cedergren, L., Bilsson, B. and Montelione, G.T. (1993) An improved strategy for determining resonance assignments for isotopically enriched proteins and its application to an engineered domain of staphylococcal protein A. Biochemistry, 32, 7839–7845.

20. Sattler, M., Schleucher, J. and Griesinger, C. (1999) Heteronuclear multidimensional NMR experiments for the structure determination of proteins in solution employing pulsed field gradients. Prog. Nucl. Mag. Res. Sp., 34, 93–158.
58. Carlton, J. M., Hirt, R. P., Silva, J. C., Deleher, A. L., Schatz, M., Zhao, Q., Wortman, J. R., Shelby, L. B., Alsmark, U. C. M., Bestiero, S., et al. (2007) Draft genome sequence of the sexually transmitted pathogen Trichomonas vaginalis. *Science*, **315**, 207–212.

59. Dominguez, C., Boelens, R., and Bonvin, A. M. (2003) HADDOCK: a protein-protein docking approach based on biochemical or biophysical information. *J. Am. Chem. Soc.*, **125**, 1731–1737.

60. van Dijk, M., van Dijk, A. D., Hsu, V., Boelens, R., and Bonvin, A. M. (2006) Information-driven protein-DNA docking using HADDOCK: it is a matter of flexibility. *Nucleic Acids Res.*, **34**, 3317–3325.