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To cite this version:

Yang-Xue Dai, Hai-Lei Guo, Na-Nv Liu, Wei-Fei Chen, Xia Ai, et al.. Structural mechanism underpinning Thermus oshimai Pif1-mediated G-quadruplex unfolding. EMBO Reports, 2022, 23 (7), pp.e53874. 10.15252/embr.202153874 . hal-03718467

HAL Id: hal-03718467

https://hal.science/hal-03718467v1

Submitted on 10 Oct 2022

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Structural mechanism underpinning *Thermus oshimai* Pif1-mediated G-quadruplex unfolding

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ABSTRACT

G-quadruplexes (G4s) are unusual stable DNA structures that cause genomic instability. To overcome the potential barriers formed by G4s, cells have evolved different families of proteins that unfold G4s. Pif1 is a DNA helicase from Superfamily 1 (SF1) conserved from bacteria to humans with high G4 unwinding activity. Here, we present the first X-ray crystal structure of the *Thermus oshimai* Pif1 (*To*Pif1) complexed with a G4. Our structure reveals that *To*Pif1 recognizes the entire native G4 via a cluster of amino acids at domains 1B/2B which constitute a G4-Recognizing Surface (GRS). The overall structure of the G4 maintains its three-layered propeller-type G4 topology, without significant reorganization of G-tetrads upon protein binding. The three G-tetrads in G4 are recognized by GRS residues mainly through electrostatic, ionic interactions, and hydrogen bonds formed between the GRS residues and the ribose-phosphate backbone. Compared with previously solved structures of SF2 helicases in complex with G4, our structure reveals how helicases from distinct superfamilies adopt different strategies for recognizing and unfolding G4s.

Key words: G-quadruplexes; *To*Pif1, G4-Recognizing Surface, X-ray, structures

INTRODUCTION

G-quadruplexes (G4s) are higher-order DNA/RNA structures formed from nucleic acid sequences that contain four stretches of three or more guanines interspaced by at least one random nucleotide (Burge *et al*, 2006; Castillo Bosch *et al*, 2014). The G-tetrad module, composed of four guanines, forms a planar structure stabilized by non-canonical Hoogsteen hydrogen bonds in the presence of monovalent cations, such as sodium or potassium (Castillo Bosch *et al.*, 2014; Griffin & Bass, 2018). Genome-wide bioinformatics analyses have revealed the prevalence of G4 motifs in key regulatory regions of the human genome, such as promoters, gene bodies, and untranslated regions (Eddy & Maizels, 2006, 2008; Huppert & Balasubramanian, 2005; Todd *et al*, 2005; Varshney *et al*, 2020). Following this prediction, a tremendous increase in cell experimental data has demonstrated that G4 structures indeed exist under physiological conditions and are involved in a variety of biological processes, including telomere maintenance, gene expression, epigenetic regulation, and DNA replication (Maizels & Gray, 2013; Schierer & Henderson, 1994). The high thermodynamic stability of G4s in cells can erect significant barriers to replication fork progression. Therefore, failing to resolve the folded G4s during DNA transactions can cause replication to stall and thus may trigger the rampant genomic instability characteristic of certain cancer cells (De & Michor, 2011; Eddy *et al*, 2015; Edwards *et al*, 2014; Fontana & Gahlon, 2020). Cells have evolved the capability of resolving stable G4s with a group of helicases that play important roles in the maintenance of genomic stability in many organisms (Mendoza *et al*, 2016). Pif1 family helicases, belonging to the superfamily 1 (SF1), are conserved across evolution from bacteria to humans and have been shown to unfold G4, being critical for DNA replication, telomere maintenance, and genome stability (Dahan *et al*, 2018; Li *et al*, 2016; Wang *et al*, 2018). For instance, a recent study has demonstrated that Pif1 helicase promotes break-induced replication in mammalian cells (Li *et al*, 2021). DNA G4-resolving helicases are also found in superfamily 2 (SF2) in groups such as the XPD family (FANCJ and RTEL1) and the RecQ family of helicases (BLM, WRN, RecQ). FANCJ bearing G4-resolving activity is necessary for DNA replication and has shown a unique substrate specificity for G4 architecture (Bharti *et al*, 2013; Lee *et al*, 2021). RecQ helicases have been found to efficiently unfold a variety of G4 DNA structures and play essential roles in DNA repair and telomere maintenance (Singh *et al*, 2012). Furthermore, mutations in FANCJ, BLM, or WRN are associated with human diseases (Budhathoki *et al*, 2014; Levitus *et al*, 2005; Zimmer *et al*, 2020). Though G4 resolvases have attracted much attention, few solved co-crystal structures in which helicase complexed with G4 are available. The X-ray crystal structure of DHX36 in complex with a Myc-promoter-derived G4 was firstly reported and shows that G4 is captured by a specific DHX36-specific motif (DSM) which bears a signature peptide AKKQ found in FANCJ (Wu & Spies, 2016). Based on DHX36-G4 crystal structures, it was suggested that DSM, in coordination with the helicase core, unfolds the G4 one base at a time in an ATP-hydrolysis-independent manner. Recently, a crystal structure (Voter *et al*,

2018) solved from a helicase core of a RecQ family helicase (*Cs*RecQ) complexed with oligonucleotides harboring a human telomere sequence revealed that a guanine base within the putative G-tetrad is located at the 5'-gate of the electropositive channel, where the G-base is flipped and sequestrated in a guanine specific pocket (GPS) (Voter *et al.*, 2018). This structure is interpreted to represent a post-catalytic complex of *Cs*RecQ bound to unfolded G4. Therefore, SF2 helicases may use a DSM/AKKQ motif or GSP site for G4 binding/unfolding, but these motifs are not found in Pif1.

Though crystal structures of Pif1 helicases from *Homo sapiens*, *Saccharomyces cerevisiae, Bacteroides sp*, and *Thermus oshimai* have been recently solved with different DNA substrates such as ssDNA or replication fork mimic, little is known about how Pif1 helicases interact with G4 (Dai *et al*, 2021; Dehghani-Tafti *et al*, 2019; Lu *et al*, 2018a; Zhou *et al*, 2016). Elucidating the molecular mechanism by which Pif1 helicase recognizes and unfolds G4 molecules largely depends on the knowledge of crystal structures. Access to structures that represent a "substrate" complex, intermediates, and a "product" complex, provides very valuable mechanistic information. Although the helicase-G4 DNA complex structures of DHX36 (Chen *et al*, 2018) and *Cs*RecQ (Voter *et al.*, 2018) can be considered as representations of an intermediate substrate complex (Chen *et al.*, 2018) and a final product complex of helicases (Voter *et al.*, 2018), respectively, the remaining challenge is to obtain the structure in which the helicases interact with an intact G4 (initial substrate complex) or to capture the structure of the very first steps leading to the transition state.

In this study, we report the X-ray crystal structures solved from the Pif1-family helicase from *T. oshimai* (*To*Pif1) bound to a pre-folded G4 bearing 6-8 nt on both sides of the G-tetrads, thereby mimicking the physiological G4 state formed during DNA replication. This is the first structural snapshot of a helicase complexed with an integral G4. In combination with bulk and single molecular fluorescent resonance energy transfer assays (smFRET), our studies provide new mechanistic insight into how a Pif1-family helicase recognizes, binds, and unfolds G4s.

RESULTS

*To*Pif1 unfolds G-quadruplexes in an ATP-dependent manner without topological preference

Parallel and antiparallel intramolecular G4s, 12 nt-ssDNA, and 24 bp blunt-end DNA were titrated with increasing concentrations of *To*Pif1 under equilibrium conditions. The apparent dissociation constant ($K_{d,app}$) determined from the titration curves (Fig 1A and Table 1) demonstrated that *To*Pif1 binds both antiparallel ($G4^{Tel}$) and parallel G4s ($G4^{CEB}$) (Fig EV1A) with similar affinity ($K_{d,app}{}^{antipara} \approx 29.63$ nM ($G4^{Tel}$); $K_{d,app}{}^{para} \approx 23.27$ nM ($G4^{CEB}$)) without any topological preference. Furthermore, although both the G4 and ssDNA (T_{12}) were bound with essentially the same affinities ($K_{d,app}{}^{antipara} \approx 29.63$ nM versus $K_{d,app}{}^{ssDNA} \approx 29.12$ nM) (Fig 1A (insert) and Table 1), blunt-end DNA bound *To*Pif1 very weakly ($K_{d,app} > 250.00$ nM; Fig 1A (insert) and Table 1). The fact that the apparent dissociation constant of G4 and ssDNA are nearly identical suggests that *To*Pif1 possesses two independent binding sites for ssDNA and G4, respectively. Indeed, competitive binding assays between the labeled G4 and unlabeled ssDNA demonstrated that labeled G4 binding was not influenced by increasing unlabeled ssDNA and vice versa (Fig EV1B). We then investigated *To*Pif1-mediated G4-unfolding activities by stopped-flow assay according to the previous report (Liu *et al*, 2020) with both parallel and antiparallel G4 substrates (Cheng *et al*, 2018) in the absence or the presence of ATP. The kinetic data were fit to Equation 2 to determine the unfolding amplitude (A_m) and unfolding rate (k_{cat}) measured with different G4 DNAs (Fig 1B and Table 1**)**. Although unfolding activity is undetectable in the absence of ATP or the presence of the non-hydrolyzable ATP analog (Fig EV1C), the values of unfolding amplitudes (A_m) and unwinding rates (k_{cat} (s^{-1})) determined with antiparallel G4s were 1.31-fold and 1.39-fold higher than those determined with parallel G4s, respectively (Fig 1B and Table 1). The calculated p-value (0.001) from the above data is less than 0.01, indicating that the observed differences between parallel and anti-parallel are statistically significant.

To gain mechanistic insight into G4 unfolding and understand the molecular basis underlying the differences in G4 unfolding between the parallel and antiparallel G4s, smFRET assays were performed with G4 DNAs in which the bases near the 5' and 3' sides of G4 were fluorescently labeled. Therefore, its unfolding activity could be sensitively recorded by smFRET time-traces. The smFRET time-trajectories recorded with antiparallel G4s comprising three or four G-tetrads ($smG4^{AP-424}$ and $sm4G4^{AP-333}$, respectively) were characterized by a stepwise unfolding procedure, in which the smFRET level successively decreased from E ≈ 0.90 to E ≈ 0.12, and then finally disappeared due to the escape of the unwound DNA from the coverslip (Fig 1C). This phenomenon has been observed previously in studies of *Sc*Pif1 by two different laboratories (Hou *et al*, 2015; Zhou *et al*, 2014), and was interpreted as that Pif1 helicase first unfolds G4 from the 5'-lateral G-column, resulting in a G-triplex (G3), then further transforms G3 into a G-hairpin structure and finally completely unwinds the duplex. Interestingly, for parallel G4 and under the same experimental conditions, although 30-35% time-traces (type I) recorded were essentially the same as those observed with antiparallel G4, ~65% smFRET time-traces (type II) for parallel G4 oscillated between E ≈ 0.90 to E ≈ 0.60 without further decrease before the fluorescence signal bleaching (Fig 1D). Thus, the smFRET assays confirmed that *To*Pif1 unfolds both parallel and antiparallel G4 using the same mechanism, but with different G4 unfolding efficiencies. The reduced unfolding efficiency in parallel G4 may simply result from the higher thermal stability in the

parallel G4 compared with the antiparallel G4 (Figs EV1D and EV1E). We then experimentally address this possibility by determining T_m values of antiparallel ($G4^{AP-424}$) and parallel ($G4^{P-214}$) G4 DNAs using a CD melting assay. The results (Figs EV1D and EV1E) show that T_m values determined with antiparallel G4 ($G4^{AP-424}$) and parallel G4 ($G4^{P-214}$) are 40.1 °C and 43.8 °C, respectively, indicating that the reduced unfolding efficiency in parallel G4 may simply result from the higher thermal stability of parallel G4.

The overall structure of *To*Pif1 in complex with G4

To probe the structural basis of *To*Pif1-mediated G4 unfolding, *To*Pif1 was complexed with parallel and antiparallel G4 DNAs flanked at their 5'- and 3'- ends with 6 and 8 nt poly(dT), respectively, mimicking G4s formed at a replication fork (Lee *et al.*, 2021; Lemmens *et al*, 2015). Only parallel G4 ($G4^{T6/T8}$) in complex with *To*Pif1 in a 1:2 ratio in the presence of ADP·AlF_4 was crystallized and diffracted to 2.58 Å. The asymmetric unit of the crystal structure contained one G4 that was sandwiched between two *To*Pif1 molecules (named as molecule *a* and *b*) bound to ssDNAs with 5'-3' polarity. The two protein molecules in the asymmetric unit were linked through the DNA substrate and related by a rotation of 102.8˚, forming a dumbbell-sharped structure (Fig 2A and B). Molecule *a* anchored close to the 5'-most G-tetrad and molecule *b* was bound downstream of the 3'-most G-tetrad. Considering the distance between the binding/unwinding surface constituted by domains 1B/2B and the nearest G-tetrad, in the molecule *a,* this distance is ≈ 3.00 Å with the 5'- most G-tetrad; while in molecule *b,* the distance with the last G-tetrad at the 3' end is greater than 12.0 Å (Fig 2A and B). Therefore, molecule *a* in 5' is in an active unwinding state during G4 processing, establishing many interactions with G4, while molecule *b* in 3' is not positioned for G4 unwinding and establishes few contacts with G4 (Fig 2A and B). The domain folding and the spatial arrangement of the modules in *To*Pif1 adopt a similar architecture as those observed in the solved crystal structures of *To*Pif1 (Dai *et al.*, 2021) and other Pif1 family helicases from yeast (Lu *et al.*, 2018a) and *Bacteroides* species (Zhou *et al.*, 2016). Domains 1A and 2A constitute a deep cleft where ADP·AlF_4 is bound, and domain 2B is composed of an SH3-like-domain and a prominent β-hairpin (loop3) (Figs 2A and B and EV1F-I)**.** Each *To*Pif1 molecule binds a 6 nt poly(dT) stretch. Molecule *a* binds T1 to T6, and molecule *b* binds T22 to T27. The conformation of T2-T5 and T24-T27 are conserved, and they superpose with a root mean square deviation (RMSD) of 1.20 Å, as already shown for *To*Pif1 complexed with several ligands, ssDNA and ss/dsDNA (Dai *et al.*, 2021), thus domain 2B exhibits high conformational flexibility. Structural superposition between molecule *a* and the previously determined *To*Pif1 complexed with ss/dsDNA (Dai *et al.*, 2021) on domain 1A demonstrated that the center of mass of domain 2B moves upwards by 5.40 Å and rotates 25.70˚ upon G4 binding. The domains 2A/1B, the configurations of bound ssDNA and the amino-acid residues involved in ssDNA binding overlap substantially (RMSD = 0.28 Å over 103 residues) (Fig EV2A-F), indicating that domain 2B assumes different conformations to accommodate G4 and double-stranded DNA (dsDNA) binding, respectively. Similarly, superposing molecules *a* and *b* on domain 1A shows that G4 binding induced a stable β-hairpin (loop3) formation, and the center of mass of domain 2B in the molecule *a* moved upwards 5.70 Å, accompanied by 22.90˚ rotation (Fig 2C), suggesting that domain 2B undergoes a significant conformational adjustment before and after G4 unfolding. G4 is complexed with three K^+ ions: two are located inside G4 between the tetrads and an extra K^+ is on top of the 3' side of G4. This K^+ ion is stabilized by the last tetrad (tetrad III:

G9·G13·G17·G21), T22, and R448 of *To*Pif1 molecule *b*. The overall topology of the G4 structure solved in this work is consistent with G4s previously reported in complexed with other proteins. They are characterized by a stack of three G-tetrads, and the three short loops containing single thymine cross the grooves of the parallel G4 helix, joining the top and bottom G-tetrads in the stack (Fig 2D). *To*Pif1-G4$^{T6/T8}$ crystal structure exhibits a good fit with SAXS data (Figs EV3A-E and Table EV3) with a χ^2 of 1.91, confirming that the conformation observed in the crystal state is also the one found in the solution.

Structural basis of recognitions and interactions of the entire G-quadruplex and the presumed G-base from the unwound G4

*To*Pif1 recognizes the entire G4 through a cluster of amino acids at domains 1B/2B, constituting G4-Recognizing Surfaces (GRS) (Fig 2E). Although R419 and R392 interact with G11 and G19 in the 5'-most G-tetrad (G7·G11·G15·G19) through cation-π interactions, respectively, the sidechain amido group of Q327 interacts with the deoxyribose group of G11 through nonbonded contacts (Figs 3A and EV2D). Intriguingly, the negatively charged carboxyl group of E397 against the negatively charged oxygen atom on the ribose-phosphate backbone near G7 will exert a repulsion force, which may influence the global conformation of the G4 (Fig 3B). Furthermore, the phosphate group of G20 in the middle tetrad (G8·G12·G16·G20) is stabilized by R135 (Fig 3C). Two positively charged residues, K329 and R150, interact with G9 and G21 in 3'-G-tetrad (G9·G13·G17·G21) through ionic and cation-π interactions (Fig 3D). Thus, the residues of GRS form structural clusters which differentially recognize the guanine bases from three different tetrads in the G4. It is therefore plausible that the spatial conformation of G4 is destabilized by these interactions, and G4 is efficiently unfolded in coordination with translocation forces derived from ATP hydrolysis.

In our *To*Pif1-G4 complex structure (*To*Pif1- G4$^{T6/T8}$), the T6 just upstream of the 5'-tetrad is interacting with a cluster of amino acids (Fig 4A). T6 also stacks against G19 in the flat 5'-G-tetrad (G7·G11·G15·G19) through a π-π interaction. R392 and V395 are specifically involved in thymine base recognition through hydrogen bonds (R392-T6 = 2.50 Å and V395-T6 = 2.80 Å). Furthermore, the base configuration is further stabilized by π-π stacking between F394 and T6. The residues R355 and R135 interact with the ribose and the phosphodiester of T6, but not its thymine base. To determine whether the binding of a guanine base at the T6 position can remodel the conformation of GRS to accommodate G-base-specific binding, an ssDNA bearing a G4-forming sequence (GR_{17}) was directly complexed with *To*Pif1 without pre-folding (Dai *et al.*, 2021). The crystal structure at a resolution of 2.21 Å demonstrated that the global conformation of *To*Pif1-GR_{17} (Dai *et al.*, 2021) is similar to the complex *To*Pif1-G4$^{T6/T8}$ (Fig 4B), except that the G4 structure is absent. Structural superposition between the two structures revealed three remarkable features: i) the G7 in binary complex (*To*Pif1-GR_{17}) occupies the equivalent spatial position of T6 in *To*Pif1-G4$^{T6/T8}$ (Fig 4C); ii) both G7 and T6 flip to the same degree (Fig 4D, ≈113.0°); iii) both the bases and ribose groups of G7 and T6 are bound by the same residues (R392, V395, and R335) with essentially the same conformation, except that residue N356 establishes an additional hydrogen bond with the phosphate group of G7 upon its binding (Fig 4D). Therefore, if base G7 is considered as the first G base from an unfolded G-tetrad, there is no striking structural reorganization to

accommodate the binding of G rather than T. Although G7 in GR_{17} described here provides a conformation for G4 being unwound in the first base, other mechanisms may also exist.

Mutational analysis of the residues involved in G4 binding and unfolding

According to the potential functions in G4 binding or/and unfolding, structurally guided single alanine substitution variants of *To*Pif1 were purified to homogeneity, including i) the mutants R392A, R135A, and R355A, which interact with the T6/G7 base and/or the ribose/phosphodiester moieties; ii) four variants (Q327A, E397A, R419A, and K329A) involved in the 5'-most G-tetrad; and iii) R135 and R150 involved in the interactions with middle G-tetrad and the 3'-most G-tetrad, respectively (Table 2). To make sure that the each given residue mutation does not result in a global conformational change, which will affect ascertain the distinct role of the given residue in G4 binding/unfolding, we first performed circular dichroism (CD) spectroscopy and dynamic light scattering (DLS) assays, which will provide the secondary structure information and hydrodynamic behavior (monomer or aggregation) of the variants. The results show that the variants displayed the same CD spectra as *To*Pif1, and the determined DLS parameters are essentially the same between *To*Pif1 and variants (Fig EV4A and Table EV4). The DNA-dependent ATPase activities of the *To*Pif1 variants were measured using a coupled spectrophotometric ATPase assay. Fig EV4B and Table EV4 demonstrated the determined K_m and k_{cat} values with the variants are not significantly different from those determined with the wild-type *To*Pif1.

To ascertain qualitatively how the modified residues affect the unfolding activities, the $K_{d,app}$ values, and the steady-state kinetic parameters, including the unfolding magnitude (A_m) and the unfolding rate (k_{cat}), were determined with fluorescent-labeled ssDNA (T_{12}), partial duplex DNA ($S_{26}D_{17}$), and G4 DNA substrates as described above. Several representative G4 unfolding kinetic curves are shown in Fig 5A and the kinetic parameters of all variants are summarized in Table 2. The parameters ($K_{d,app}$, A_m, and k_{cat}) determined with G4 and ssDNA substrates showed a range of variability. To best understand the mutation results, the ratio of $k_{cat}/K_{d,app}$, a useful index for comparing the catalytic efficiency (Barlow *et al*, 2009), was determined (Table 2). Arranging the G4 $k_{cat}/K_{d,app}$ values in ascending numerical order shows that the relative increase in G4 $k_{cat}/K_{d,app}$ values can be roughly classified into three categories (Fig 5B): i) group I (R355A, R135A, and R150A) is characterized by zero values in $k_{cat}/K_{d,app}$ for G4 unfolding while their $k_{cat}/K_{d,app}$ values determined from dsDNA unwinding are reduced by 33% (R150A) and 75% (R355A); ii) group II (R392A, Q327A, K329A, and R419A) is marked by about 44-69% reduction in $k_{cat}/K_{d,app}$ values for G4 unfolding and the corresponding values determined with dsDNA are essentially the same as these determined with *To*Pif1, except that $k_{cat}/K_{d,app}$ value determined with R419A for dsDNA is reduced to 50%; iii) group III (E397A, E397L, and E397H) displays surprising results in which $k_{cat}/K_{d,app}$ values determined with G4 are increased to 2-2.6 folds while the $k_{cat}/K_{d,app}$ values for dsDNA are inversely reduced to 50% compared with that of *To*Pif1. Taken together, these results indicate that alteration of the residues in group I-II, particularly the residues in group I, significantly impair G4 unfolding, but only moderately reduces dsDNA unwinding. Consistently, smFRET time trajectories recorded with the mutants in group III revealed that in sharp contrast to wt*To*Pif1, the curves only oscillate between 0.90 and 0.60, but

scarcely attain a completely unfolded level, indicating that the variants can just release the 5'-most lateral G column from G4 DNA, but is unable to unfold the integral G4 DNA completely (Fig 5C-F).

In group III, the kinetic parameters determined with E397A were significantly higher in terms of unfolding amplitude (A_m) and rate (k_{cat}), respectively (Figs 5B and 6A and Table 2). Compared with *To*Pif1, the $k_{cat}/K_{d,app}$ value of E397A determined for G4 unfolding increased by 2.5-fold, although that for dsDNA remained unchanged (Table 2). This stimulation effect was also observed with parallel G4s (Fig 6B). To confirm this interpretation, residue E397 was therefore replaced with a hydrophobic (E397L), a positively charged (E397H), and a negatively charged (E397D) residue. As expected, G4 unfolding activities determined with the first two variants were systematically higher than those of *To*Pif1. E397D bearing a negative charge displayed activity comparable to the wild type (Fig 6A). To gain in-depth mechanistic insight into how residue E397 affects G4-unfolding activity, all of the variants were further studied with smFRET techniques. Analysis of the smFRET time-traces demonstrated that the unfolding times ($t(s)$) determined with the three typical variants were significantly reduced, except those for E397D, whose unfolding time remained essentially the same as that of *To*Pif1 (Fig 6C-F). The mutation-stimulating G4-resolution activity is surprising and raises the question of whether this may be a general autoregulatory mechanism by which the activity of G4 unfolding is regulated both *in vitro* and *in vivo*.

DISCUSSION

While increasing evidence has demonstrated that helicase-mediated G-quadruplex unfolding plays an essential role in preserving genome stability, the lack of structural information on G4-processing helicases still hampers our mechanistic understanding of G4 resolution. With a wide array of approaches ranging from bulk/smFRET assays to structural biology analyses, our results reported here provide in-depth insight into the currently largely unknown, but mechanistically important issue of how the Pif1 helicase family recognizes the integral G4 structure.

To our best knowledge, the structure reported here is the first co-crystal structure of SF1 helicase bound to G4, while crystal structures of SF2 helicase-G4 complexes available are DHX36 and *Cs*RecQ proteins. Thus, a structural comparison between SF1 and SF2 helicase-mediated G4 binding/unfolding may provide the common and distinct mechanism underlying G4 unfolding.

DHX36 protein, one of the most extensively studied G4-unfolding helicases, is characterized by a DHX36-specific motif (DSM), which specifically binds parallel G4s (Hossain *et al*, 2021). Interestingly, the X-ray crystal structure of DHX36 (Chen *et al.*, 2018) in complex with a Myc-promoter-derived G4 demonstrates a rearrangement of the 5'-G-tetrad with a one-base translocation upon DSM binding (Chen *et al.*, 2018). Because an NMR study shows that G4 keeps its integral state upon the binding of an isolated DSM (Heddi *et al*, 2015), it was, therefore, suggested that DSM, in coordination with the helicase core, unfolds the G4 one base at a time in an ATP- hydrolysis-independent manner (Chen *et al.*, 2018). Recently, a crystal structure solved from a helicase core of a RecQ family helicase (*Cs*RecQ) complexed with

oligonucleotides harboring a human telomere sequence (Voter *et al.*, 2018) revealed that a guanine base within the putative G-tetrad is located at the 5'-gate of the electropositive channel where the G-base is flipped and sequestrated (Voter *et al.*, 2018). This structure is interpreted to represent "a product complex of *Cs*RecQ-unwound G4 rather than a folded quadruplex" (Voter *et al.*, 2018) although the complexed ssDNA was not pre-folded to form a stable G4 before the preparation of the G4-*Cs*RecQ complex.

DHX36 uses DSM to bind and unfold G4s, and some G4-resolving helicases as FANCJ also possess a motif equivalent to DSM (Chen *et al.*, 2018; Lee *et al.*, 2021). However, not all the SF2 helicases bear DSM. An intriguing and unresolved question is how the G4-resolving helicases generally recognize and unfold G4s. We demonstrated that the 1B/2B domains in *To*Pif1 constitute a G4-Recognizing Surface (GRS) in which the residues recognize the whole G4 structure, not just a tetrad, and differentially interact with the ribose and phosphate units in different tetrads. Superposition of the structural conformation of G4 complexed with *To*Pif1 with that of the DHX36-Myc complex (Chen *et al.*, 2018) demonstrated that, when the G4s well superposed, loop3 in *To*Pif1 occupies the spatial position of the DSM motif in DHX36, binding on the 5' side of the G4. Given that Pif1 and DHX36 have opposite polarity for ssDNA translocation, it appears that helicase binding on the 5' side of G4 is a critical feature for G4 unwinding. Furthermore, it appears that the previously identified GSP in the RecQ helicase family is not conserved in *To*Pif1, because the residues involved in the first base downstream of 5'-G-tetrad are not selective for guanine and do not form the equivalent hydrogen bonds that stabilize the guanines within G4s as observed in GSPs (Figs 4A-D and EV4C-E). If base G7 is considered as the first G base from an unfolded G-tetrad, there is no striking structural reorganization to accommodate the binding of G rather than T. Compared with the previously identified GSP, the cluster of residues surrounding the base (T6/G7) upstream of the G-tetrad in *To*Pif1 are not bound with the equivalent residues identified in the GSP in CsRecQ (Fig EV4C-E). Of note, the G-rich ssDNA (GR_{17} (Dai *et al.*, 2021) and resolved G4 DNA (Voter *et al.*, 2018)) were not pre-folded to form stable G4 DNA in the *To*Pif1 and *Cs*RecQ structures; it, therefore, remains to be determined whether these structures (*To*Pif1-GR_{17} (Dai *et al.*, 2021) and *Cs*RecQ-ssDNA (Voter *et al.*, 2018)) represent an intermediate or/and a product complex of helicases bound to unwound G4 DNA or just ssDNA bound to the proteins.

Our binding studies demonstrated that *To*Pif1 possesses strong G4 binding activity without any topological preference. This is not surprising because *To*Pif1 does not bear a DSM motif which was reported exclusively bind parallel G4 (Hossain *et al.*, 2021). The smFRET time-trajectory feature of *To*Pif1-mediated G4 unfolding is mainly characterized by a stepwise unfolding procedure, without a long oscillation period, which is consistent with the previously recorded smFRET time-trajectory with *Sc*Pif1 (Hou *et al.*, 2015; Zhou *et al.*, 2014). These results imply that *To*Pif1 translocates along the DNA lattice and sequentially unfolds G4 into G-triplexes and then into G-hairpins. *To*Pif1-mediated G4

processing is radically different from previous reports on DHX36 (Chen *et al.*, 2018) and the RecQ (Voter *et al.*, 2018) helicase family, which were shown to use a common mechanism of ATP-independent repetitive unfolding to resolve G4s: binding of these helicases triggers the reorganization and destabilization of G4 with a one-at-a-time type base translocation, and ATP is only needed for releasing the unfolded G4 (Chen *et al.*, 2018; Voter *et al.*, 2018). Therefore it appears that SF1 helicases unfold G4 in a stepwise manner while SF2 helicases unfold G4 repetitively; however, establishing these observations as a general rule needs further studies. Another interesting finding is that two negative charges between E397 and the oxygen atom on the DNA backbone near G7 repel one another. A series of mutations at E397 demonstrated that replacement of the negative residue by any non-negative residue stimulates G4 unfolding activity, raising the question that whether an auto-inhibitory mechanism applies to other *To*Pif1 helicases. A sequence alignment of Pif1 family helicases revealed that a negative residue (E/D) is highly conserved in thermophilic gram-negative bacteria (Fig EV5). However, there is a tendency towards replacing negative residue (E/D) with no-negative residues (S/G) in evolutionarily advanced species (Fig EV5). Therefore it appears that *To*Pif1 has not evolved with an optimum efficiency, but this could be due to growth conditions, G4 frequency in the genome, or a yet unknown regulatory mechanism stimulating *To*Pif1 activity.

The relation between growth temperature, G4 frequency, and Pif1 efficiency is not straightforward. Analysis of the occurrence of putative G4 sequences (PQS) in bacteria has recently been conducted. Though thermophile bacteria have genomes with higher GC content, the distribution of PQS is higher than expected by random chance in bacteria groups *Deinococcales* and *Thermales* while using either quadparser (Ding *et al*, 2018) or G4hunter (Bartas *et al*, 2019) algorithms. In later study, it was shown that *T. oshimai* had the highest frequency of PQS among all bacterial genomes explored (Table EV5). Surprisingly, the same study showed that *Thermotogae* which are also extremely thermophilic gram-negative bacteria, have the lowest frequency of PQS (Table EV5). Therefore, G4 occurrence does not seem to be linked to thermal conditions of growth. Bacteria from *Thermotoga* group possess also a Pif1 helicase which exhibits G4 activity (Andis *et al*, 2018). The biochemical activities of Pif1 from *Thermotoga elfi* (*Te*Pif1) have been measured, and *Te*Pif1 exhibits a C-terminal WYL domain with ssDNA binding function (Andis *et al.*, 2018) necessary for substrate binding and helicase activity of *Te*Pif1. WYL domain is shared by Pif1 from other groups, such as *Deferribacteres*, but *To*Pif1 does not have one. Since Pif1 from other thermophilic bacteria such as *Thermotoga elfii* possessed an extra WYL domain (Andis *et al.*, 2018), modulation of Pif1 activity in cis (WYL domain) and a regulatory effect on Pif1 in trans by unidentified co-factor proteins can be hypothesized.

It was recently reported that the proteins Mgs1 and Mms1 in *S. cerevisiae* binding to G4 motifs *in vivo* partially depends on the helicase *Sc*Pif1 (Paeschke & Burkovics, 2021; Schwindt & Paeschke, 2018). Using the sequence similarity search tool, we identified the Mgs1-like protein in *Thermus oshimai* genome (*To*Mgs1 like, Genebank: WP_018462088.1) which share shares 29% identity with *Sc*Mgs1 across their lengths. *To*Mgs1 could be a co-factor regulating *To*Pif1 in trans.

Though Pif1 is conserved from bacteria to humans, Pif1 has very diverse functions in bacteria, yeast, and human. Pif1 sequence exhibits variations with the insertion domain 2C in budding yeast (Lu *et al.*, 2018a) (Lu *et al*, 2021) and accessory domains as WYL (Andis *et al.*, 2018). Furthermore, sequence alignments show that some of the key residues identified in GRS are shared with other bacteria but not with Pif1 from eukaryotes (for an alignment, see Fig S1 in Supplementary material in (Dai *et al.*, 2021)). We show here that *To*Pif1 can rely only on its helicase core domain to recognize and unfold G4 DNA while the activity of *Te*Pif1 needs a WYL accessory domain (Andis *et al.*, 2018). Therefore, it appears that Pif1 from different species adopts different solutions for enhancing its activity. Thus, co-factors acting in cis or trans may enhance Pif1 activity for certain species, and the situation is far from being universal even in the Bacteria domain. Further studies, instigated by the work reported here, are needed to complete our understanding of the general mechanism underlying how G4-resolving helicases recognize and unfold G4s.

MATERIALS AND METHODS

Reagents and buffers

All chemicals were reagent grade and all buffers were prepared in high-quality deionized water from a Milli-Q ultrapure water purification system (Millipore) with a resistivity of >18.2 MΩ.cm and were filtered again on a 0.20 μm filter before use. All DNA unwinding and binding assays were performed in buffer A (20 mM Tris-HCl (pH 7.5 at 37 °C), 50 mM NaCl, 2 mM $MgCl_2$, 2 mM DTT) which was optimized previously by our group (Dai *et al.*, 2021).

DNA substrate preparation

All the DNA substrates used in this study were chemically synthesized and HPLC-purified by Sangon Biotech (Shanghai) as listed in Table EV1. The oligonucleotides used in binding and unwinding assays were prepared at a 2 μM working concentration. The duplex or G4 DNA used in the stopped-flow and smFRET assays were heated to 95 °C for 5 min in stocking buffer (20 mM Tris-HCl, pH 7.5, 100 mM NaCl) and annealed by slow cooling to room temperature. The DNA substrates in crystallization were dissolved in 20 mM Tris-HCl (pH 7.5) with 100 mM KCl, heated to 95 °C, and allowed to cool slowly to room temperature in a water bath. After purification on a Mono Q column, the formation of G4 structures was checked using circular dichroism (CD) spectropolarimetry.

Protein expression and purification

*To*Pif1 (residues 64-507) and its mutants (R355A, R135A, R150A, R419A, R392A, Q327A, K329A, E397A, E397L, E397H, and E397D) were all cloned into pET15b-SUMO and then transformed into the C2566H *E. coli* strain (New England Biolabs), respectively. When the culture reached the early stationary phase (OD_{600} = 0.55 - 0.6) at 37 °C, 0.3 mM IPTG was added and the protein expression was induced at 18 °C over 16 h. Cells were harvested by centrifugation (4500 g, 4 °C, 15 min) and pellets were suspended in lysis buffer (20 mM Tris-HCl pH 7.5, 500 mM NaCl, 10 mM Imidazole, and 5% glycerol (v/v)). Cells were broken with a French press and then further sonicated 2-3 times to shear DNA. After centrifugation at 12,000 rpm for 40 min, the supernatants were filtered through a 0.45-μm filter and loaded onto a Ni^{2+} charged IMAC column (GE Healthcare). After washing twice, the SUMO-*To*Pif1 was then eluted from the Ni^{2+} affinity column with elution buffer (20 mM Tris-HCl, pH 7.5, 500 mM NaCl, 300 mM Imidazole, and 5% glycerol (v/v)) at 4 °C. The eluted protein was treated with SUMO protease (Invitrogen, Beijing). Then the SUMO digested protein was further purified by a HiTrap Heparin column (GE Healthcare) to remove the SUMO-tag and other protein impurities. The eluted fraction containing *To*Pif1 was collected and concentrated. The final purified protein was dialyzed against the storage buffer (20 mM Tris-HCl, pH 7.5, 500 mM NaCl, 1 mM DTT) and concentrated to approximately 10

mg/mL for crystallization, and was about 95% pure as determined by SDS-PAGE. Mutations and truncations were engineered by PCR overlapping-PCR protocol.

Equilibrium binding assays

The isothermal binding curves were determined using a fluorescence polarization assay on an Infinite F200 plate reader (Tecan). FAM-labeled DNA substrates were used in this study. Varying amounts of protein were added to a 150 μL aliquot of buffer A containing 5 nM FAM-labeled DNA. Each sample was allowed to equilibrate in solution for 5 min at 37 °C, and then fluorescence polarization was measured. Less than 5% change was observed between the 5 and 10 min measurements, indicating that equilibrium was reached in 5 min. The equilibrium dissociation constants were determined by fitting the binding curves using Equation 1:

$$\Delta r = \Delta r_{\max} \times P / (K_{d,app} + P) \quad (1)$$

where $\Delta r_{\max}$ is the maximal amplitude of the anisotropy (i.e., $r_{\max} - r_{\text{free,DNA}}$), P is the helicase concentration, and $K_{\text{d,app}}$ is the midpoint of the curve corresponding to the apparent dissociation constant.

Crystallization of *To*Pif1-nucleic acid complexes

For crystallization, purified *To*Pif1 was incubated with the $G4^{T6/T8}$ after dissolving and folding in 20 mM Tris-HCl (pH 7.5) with 100 mM KCl at a molar ratio of 2:1 in the presence of ATP analog ADP·AlF_4. The resulting *To*Pif1-$G4^{T6/T8}$-ADP·AlF_4 complex was purified using size-exclusion chromatography and then concentrated to approximately 10 mg/mL. Crystallization trials on *To*Pif1 and its complexes with DNA (Table EV1) and ADP·AlF_4 were performed at 20 °C using the sitting-drop vapor diffusion method. Crystallization screening was carried out at 20 °C using commercial screening kits (Hampton Research, Molecular Dimensions, and Rigaku Reagents), where the *To*Pif1-DNA complex was mixed at a 1:1 ratio with the reservoir solution. Crystals of *To*Pif1-$G4^{T6/T8}$-ADP·AlF_4 were obtained using 0.1 M Bis-Tris (pH = 6.5), 10% PEG20000. All these conditions were optimized with a grid search using 48-well Linbro plates at 20 °C, where 1 μL of protein sample and 1 μL of precipitant were mixed and equilibrated with 60 μL of precipitant.

X-ray data collection, phasing, and refinement

All X-ray diffraction data were collected on beamline BL19U1 (Zhang *et al*, 2019) at the Shanghai Synchrotron Radiation Facility (China) using a Pilatus 6M detector (Dectris) and were processed using XDS (Kabsch, 2010). *To*Pif1-$G4^{T6/T8}$-ADP·AlF_4 structures were solved by molecular replacement, performed in the PHENIX software suite (Adams

et al, 2010) with Phaser (McCoy *et al*, 2007), using the *To*Pif1 apo structure (PDB: 6S3E (Dai *et al.*, 2021)) as the search model (Lovelace *et al*, 2005). Manual rebuilding was done using the Coot (Emsley *et al*, 2010) and further refinement was performed in PHENIX. Cell parameters and data collection statistics are reported in Table EV2.

Small-angle X-ray scattering assay

SEC-SAXS experiment was carried out at beamline SWING (SOLEIL Synchrotron, Saint-Aubin, France) with SEC-HPLC coupled to SAXS. The sample of *To*Pif1-G4 complex was injected at a concentration of 10 mg/ml on a Superdex 200 5/150 Increase column (Cytiva) at a flow rate of 0.2 ml/min equilibrated in buffer 25 mM Hepes (pH=7.5), 150 mM KCl, 5% glycerol. Scattering data were collected at 20 °C using a Pilatus 1M detector (Dectris), and the data reduction and processing of images were done with Foxtrot (David & Pérez, 2009). Further analysis was done with ATSAS 2.8 suite (Petoukhov *et al*, 2012). Experimental R_g, $I(0)$, and D_{max} values were calculated with PRIMUS and GNOM4 programs respectively. *Ab initio* envelopes for isolated complexes were determined using DAMMIF with the pair distance distribution function calculated with GNOM4. Full atomic models derived from crystal structure were modeled with MODELLER and adjusted to SAXS data with DADIMODO. In this modeling procedure, only the missing parts from the X-ray structure were kept flexible. Profiles of atomic models were calculated and fitted to the experimental data using CRYSOL and aligned on ab initio bead models with SUPCOMB. All the SAXS parameters are summarized in Table EV3.

Stopped-flow unwinding assay

Briefly, unwinding kinetics were measured in a two-syringe mode, where *To*Pif1 and fluorescently labeled DNA substrate were pre-incubated at 37 °C in one syringe for 5 min and the unwinding reaction was initiated by rapidly mixing with ATP from another syringe. Each syringe contained unwinding reaction buffer A (25 mM Tris-HCl (pH 7.5), 50 mM NaCl, 2 mM $MgCl_2$ and 2 mM DTT). All concentrations listed are after mixing, unless otherwise noted. For converting the output data from volts to percentage unwinding, a calibration experiment was performed in a four-syringe mode, where the helicase, the hexachlorofluorescein-labeled single-stranded oligonucleotides, the fluorescein-labeled single-stranded oligonucleotides, and ATP were in four syringes, respectively. The fluorescent signal of the mixed solution from the four syringes corresponded to 100% unwinding. The standard reaction was performed with 4 nM DNA substrates, 1 mM ATP, and 100 nM *To*Pif1 in buffer A.

All stopped-flow kinetic traces were averages of ≥10 individual traces. The kinetic traces were analyzed using Bio-Kine (version 4.26, Bio-Logic, France) using Equation 2:

$$A(t)=A_1(1-e^{-k_1(t-t_0)})+A_2(1-e^{-k_2(t-t_0)}) \quad (2),$$

where A(t) represents the fraction of DNA unwound at time t, A_1 and A_2 are the unwinding amplitudes, k_1 and k_2 are the unwinding rate constants of the two phases, t_0 is the time at which the fraction of DNA unwound starts to rise. From the four parameters obtained through the fitting, we can determine the total unwinding amplitude $A_m = A_1 + A_2$ and the initial unwinding rate (i.e., the slope of the kinetic unwinding curve at early times) $k_{cat} = k_1A_1 + k_2A_2$.

Single-molecule fluorescence data acquisition

50 pM fluorescently labeled DNA was added to the chamber containing imaging buffer composed of 20 mM Tris-HCl (pH 7.5), 1 mM $MgCl_2$, 50 mM NaCl, 1 mM DTT and an oxygen scavenging system (0.8% D-glucose, 1 mg/mL glucose oxidase, 0.4 mg/mL catalase and 1 mM Trolox). After immobilization for 10 min, free DNA molecules were removed by washing with the imagining buffer. We used an exposure time of 100 ms for all recordings at a constant temperature of 22 °C. FRET efficiency was calculated using $I_A/(I_D + I_A)$, where I_D and I_A represent the intensities of donor and acceptor, respectively. Basic data analysis was carried out by scripts written in MATLAB, and all data fitting were generated using Origin 9.0. Histograms were fitted to Gaussian distributions, with the peak positions unrestrained.

Circular dichroism spectropolarimetry

Circular dichroism (CD) experiments were performed with a Bio-Logic MOS450/AF-CD optical system (BioLogic Science Instruments, France) equipped with a temperature-controlled cell holder, using a quartz cell with a 1-mm path length. A 2.5 μM solution of G4 DNA was prepared in 25 mM Tris–HCl (pH 7.5), 100 mM NaCl. CD spectra were recorded in the UV (220–320 nm) regions in 0.75 nm increments with an averaging time of 2 s at 25°C.

Dynamic light scattering (DLS) assay

DLS measurements were performed at 37°C using a DynaPro NanoStar instrument (Wyatt Technology Europe GmbH, Germany) equipped with a thermostated cell holder and disposable cuvettes (UVette, Eppendorf). All solutions were filtered using 0.22 μm filters. Purified ToPif1 or its mutants (10 μM) was dialyzed in buffer A (final volume, 30 μl). The scattered light was collected at an angle of 90°. Recording times were typically between 3–5 min (20–30 cycles in average, 10 s in each cycle). The analysis was performed with the Dynamics 7.0 software using regularization method (Wyatt Technology). The molecular weight was calculated from the hydrodynamic radius using the following empirical Equation 1,

$$Mw = \left(1.68 \times R_H\right)^{2.34} \qquad (1)$$

where Mw and R_H represent the molecular weight (in kDa) and the hydrodynamic radius (in nm), respectively.

ATP hydrolysis measurement

The malachite green assay (MAK113; Sigma-Aldrich) was used to test the ATP hydrolysis activity of ToPif1 and its mutants by measuring the release of inorganic phosphate as previously described (Liu *et al*, 2021; Mahieu *et al*, 2020). Briefly, 100 nM purified protein and ATP (concentrations ranging from 0 μM to 400 μM) were incubated with or without 1 mM DNA effector ($S_{18}H_{11}$, sequence were shown in Table S1) using a VICTOR Nivo Multimode Microplate Reader (PerkinElmer, Waltham, MA, USA). All reactions were performed in triplicates of 150 μL each at 25℃ in buffer A. The parameters of K_m and k_{cat} were calculated by fitting the experimental data to the Michaelis–Menten equation using Origin 9.0 (OriginLab Corporation, Northampton, MA, USA).

DATA AVAILABILITY

Accession codes: The atomic coordinate and structure factors have been deposited in the Protein Data Bank (https://www.rcsb.org/) with the following accession numbers: 7OAR (*To*Pif1-G4$^{T6/T8}$-ADP·AlF_4).

Expanded View for this article is available online.

ACKNOWLEDGEMENTS

We are grateful for access to the SOLEIL (SWING) synchrotron radiation facility for SAXS data collections. We thank the staffs from BL17U1/BL18U1/BL19U1 beamline of National Facility for Protein Science in Shanghai (NFPS) at Shanghai Synchrotron Radiation Facility, for assistance during data collection. This work was supported by National Natural Science Foundation of China [31870788, 11574252, 11774407]; the Natural Science Basic Research Program of Shaanxi (2020JQ-251); CNRS LIA ('Helicase-mediated G-quadruplex DNA unwinding and genome stability'). Funding for open access charge: National Natural Science Foundation of China [31870788].

Author contributions

Conceived and designed study: XGX, XMH, and SR. Experiments: YXD, HLG, NNL, WFC, XA, and HHL. Data processing: YXD, NNL, WFC, and HLG. Discussion and interpretation of data: XGX, XMH, WFC, and SR. Manuscript writing and editing: XGX, YXD, NNL, BS, and SR, with input from all authors.

Disclosure and competing interest statement. None declared.

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TABLES

Table 1. Binding and unwinding activities of *To*Pif1 for different DNA substrates[1].

Substrate		Binding[2]	Unwinding[3]	
		$K_{d,app}$ (nM)	A_m	k_{cat} (s^{-1})
Parallel G4	$G4^{CEB}$	23.27 ± 3.12	ND[4]	ND
	$G4^{P-214}$	ND	0.55 ± 0.02	1.57 ± 0.07
	$G4^{P-241}$	ND	0.35 ± 0.02	1.23 ± 0.05
	$G4^{P-124}$	ND	0.22 ± 0.01	1.17 ± 0.04
Antiparallel G4	$G4^{Tel}$	29.63 ± 2.64	ND	ND
	$G4^{AP-424}$	ND	0.72 ± 0.06	2.18 ± 0.02
	$4G4^{AP-333}$	ND	0.70 ± 0.03	0.57 ± 0.01
ssDNA	T_{12}	29.12 ± 4.19	-	-
dsDNA	$S_{26}D_{17}$	ND	0.59 ± 0.02	3.53 ± 0.02
Blunt-end DNA	D_{24}	250~280	-	-

[1]Data were determined from 2-3 independent experiments under the experimental conditions as described in Materials and Methods.

[2]Data obtained from Fig 1A.

[3]Parameters were fitted from Figs 1B and 6B. The original curve of *To*Pif1 unwinding substrate $S_{26}D_{17}$ were not shown.

[4]ND, not determined.

Table 2. Parameters of binding and unwinding activities of *ToPif*1 and its variants[1].

	G4 binding ($G4^{Tel}$)	G-quadruplex unwinding ($AP\text{-}S_{16}\text{-}TelG4$)			ssDNA binding (T_{12})	dsDNA unwinding ($S_{26}D_{17}$)		
Variants	$K_{d,app}$ (nM)	A_m	k_{cat} (s^{-1})	$k_{cat}/K_{d,app}$[2]	$K_{d,app}$ (nM)	A_m	k_{cat} (s^{-1})	$k_{cat}/K_{d,app}$[3]
*To*Pif1	29.63 ± 2.64	0.53 ± 0.01	0.48 ± 0.01	0.0162 ± 0.0012	30.20 ± 1.41	0.59 ± 0.02	3.53 ± 0.02	0.1169 ± 0.0031
R392A	28.50 ± 2.20	0.40 ± 0.01	0.27 ± 0.02	0.0095 ± 0.0001	28.30 ± 1.80	0.32 ± 0.01	3.50 ± 0.02	0.1237 ± 0.0067
R355A	133.00 ± 18.04	0.38 ± 0.02	0.07 ± 0.01	0.0005 ± 0.0001	38.10 ± 1.15	0.34 ± 0.01	1.27 ± 0.01	0.0333 ± 0.0005
R135A	119.00 ± 10.67	0.34 ± 0.02	0.05 ± 0.01	0.0004 ± 0.0001	30.90 ± 0.47	0.47 ± 0.01	1.42 ± 0.02	0.0460 ± 0.0010
Q327A	51.50 ± 3.23	0.48 ± 0.01	0.41 ± 0.02	0.0080 ± 0.0002	63.40 ± 9.42	0.43 ± 0.03	6.14 ± 0.03	0.0968 ± 0.0121
R419A	38.50 ± 2.25	0.24 ± 0.02	0.21 ± 0.01	0.0055 ± 0.0001	38.20 ± 5.47	0.32 ± 0.02	2.46 ± 0.01	0.0644 ± 0.0016
K329A	48.70 ± 2.65	0.37 ± 0.01	0.29 ± 0.02	0.0060 ± 0.0001	30.00 ± 1.38	0.39 ± 0.02	3.38 ± 0.01	0.1127 ± 0.0040
R150A	152.00 ± 12.97	0.28 ± 0.01	0.06 ± 0.01	0.0004 ± 0.0001	31.30 ± 1.76	0.43 ± 0.03	2.50 ± 0.01	0.0799 ± 0.0039
E397A	28.30 ± 2.21	1.00 ± 0.01	1.19 ± 0.01	0.0420 ± 0.0027	29.40 ± 1.36	0.49 ± 0.02	3.79 ± 0.02	0.1289 ± 0.0050
E397L	33.00 ± 3.24	0.88 ± 0.01	1.18± 0.01	0.0358 ± 0.0020	36.00 ± 2.84	0.48 ± 0.01	2.36 ± 0.02	0.0656 ± 0.0043
E397H	44.60 ± 1.68	0.78± 0.03	1.32± 0.01	0.0296 ± 0.0009	50.90 ± 3.45	0.31 ± 0.02	2.64 ± 0.02	0.0519 ± 0.0030
E397D	38.80 ± 8.07	0.45± 0.01	0.35± 0.02	0.0090 ± 0.0011	28.00 ± 3.18	0.49 ± 0.01	3.27 ± 0.01	0.1168 ± 0.0116

[1]The reported data are the average values determined from 3-4 independent assays as performed under experimental conditions as described in 'Materials and Methods'.

[2] The values are the equivalent ratios of k_{cat} (G-quadruplex unwinding) and $K_{d,app}$ (G4 binding) comes from the data of Figs 1A, 5A and 6A.

[3] The values are the equivalent ratios of k_{cat} (dsDNA $S_{26}D_{17}$ unwinding) and $K_{d,app}$ (ssDNA T_{12} binding).

FIGURE LEGENDS

Figure 1. Binding and unwinding activities of *To*Pif1 for different configurations of antiparallel and parallel G4s.

A Comparison of the binding activity of *To*Pif1 for different configurations of antiparallel ($G4^{Tel}$, black points) and parallel ($G4^{CEB}$, red points) G4s. (Insert: Comparison of the binding activity of *To*Pif1 to single-stranded DNA (T_{12}, black points) with $G4^{Tel}$, red points), the blue points represent of the double-strand DNA (D_{24})). The binding activity was measured using steady-state fluorescence anisotropy assays; 5 nM of fluorescein-labeled DNA substrate and 0.5 mM ADP·AlF_4 were titrated with increasing protein concentrations at 37 °C. The different binding curves represent an average of 3-4 independent experiments for each substrate with standard deviations. The solid line represents the fit of data according to Eq. (1).

B Stopped-flow DNA unwinding kinetics of *To*Pif1 for different configurations of G4 DNA ($G4^{AP-424}$, $4G4^{AP-333}$, $G4^{P-214}$) under multiple turnover conditions. All curves represent the average of at least 10 individual traces and the plots are representative of three independent experiments. 4 nM G4 DNA and 100 nM *To*Pif1 were used under experimental conditions as described in 'Materials and Method'.

C, D The typical smFRET trajectories of antiparallel ($smG4^{AP-424}$, $sm4G4^{AP-333}$) and parallel ($smG4^{P-111}$) G4 unwinding catalyzed by 50 nM *To*Pif1 and 150 μM ATP. Black arrows in the solid style indicate the time of adding proteins. Dashed arrows in black represent the unwinding time of *To*Pif1 with different G4 DNA substrates.

Figure 2. Structure analysis of *To*Pif1 complexed with G-quadruplex (G4) DNA and ADP·AlF_4.

A Overall structure of *To*Pif1-$G4^{T6/T8}$-ADP·AlF_4 ternary complex in cartoon mode with the conserved domains 1A (green), 1B (orange), 2A (pink), 2B (blue), and C-terminal (CTD, brown). G4 DNA ($G4^{T6/T8}$) is colored in red and ADP·AlF_4 is shown as a black stick. *To*Pif1 molecules binding at the 5'- and 3'-ends of G4 DNA are labeled as molecule *a* and molecule *b*, respectively.

B Surface view after cutting the crystal structure *To*Pif1-$G4^{T6/T8}$-ADP·AlF_4 ternary complex and the domains are colored as in (A).

C Structural superposition for the structures of molecules *a* and *b* with the mass centers of domain 2B shown as a sphere (mass centers of molecules *a* colored in blue and that of molecules *b* colored in cyan).

D Comparison of $G4^{T6/T8}$ (red) in the *To*Pif1 ternary complex, $G4^{Myc}$ (PDB: 5VHE (Chen *et al.*, 2018)) colored in cyan and G4 'T-loops' (PDB: 6LDM (Traczyk *et al*, 2020)) colored in yellow.

E Residues of the molecule *a* that interact with G4 layers in the *To*Pif1-$G4^{T6/T8}$-ADP·AlF_4 ternary complex; G4 DNA ($G4^{T6/T8}$) is shown in red and the interacting residues are represented as sticks.

Figure 3. Atomic interactions between the integral G-quadruplex (G4) DNA and *To*Pif1.

A Interactions of *To*Pif1 and the G4 layer I.

B The negative-negative repulsion interaction between E397 and the G7 of $G4^{T6/T8}$ in the *To*Pif1 ternary complex structure. The electrostatic potential at ± 5 kTe^{-1} was colored in blue (basic/positive), white (neutral), and red (acidic/negative).

C, D Interaction of *To*Pif1 and G4 layers II and III, respectively. Molecular electrostatic potential map calculated with G4 DNA omitted from the co-crystal structure (blue to red, ± 5 kTe^{-1}).

Figure 4. Structural superposition of *To*Pif1 with the $G4^{T6/T8}$ and the GR_{17}.

A Interaction between *To*Pif1 and the T6 of $G4^{T6/T8}$ in the *To*Pif1-$G4^{T6/T8}$-ADP·AlF_4 ternary complex.

B Superposition on domain 1A of structures in *To*Pif1-$G4^{T6/T8}$ and *To*Pif1-GR_{17}.

C Interaction between *To*Pif1 and the G7 of GR_{17} in the *To*Pif1-GR_{17}-ADP·AlF_4 ternary complex.

D Structural superposition of interacting residues for *To*Pif1 with the T6 of $G4^{T6/T8}$ and the G7 of GR_{17}.

Figure 5. G-quadruplex (G4) unwinding activity of wild-type *To*Pif1 (*To*Pif1) and its mutants.

A Stopped-flow kinetics of the G4 (AP-S_{16}-TelG4) unwinding activity of *To*Pif1 and the various modified proteins. The different unwinding curves represent an average of 3-4 independent experiments for each substrate. The solid line represents the fit of data according to Eq. (2).

B Bar plot of $k_{cat}/K_{d,app}$ values in Table 2 (n=3 biological replicates). The $k_{cat}/K_{d,app}$ values of G4 in descending numerical order (front panel), and these are determined with the dsDNA ($S_{26}D_{17}$, back panel). Data are presented as means ± SD.

C-F The typical smFRET trajectories of antiparallel G4 ($smG4^{AP-424}$) unwinding catalyzed by different *To*Pif1 mutants based on single-molecular FRET assays. The dashed magenta, black, and red lines represent the states of G4: fully folded, incompletely folded, and fully unfolded, respectively. The concentrations of the R355A (C), R135A (D), R150A (E), and R419A (F) mutants were 50 nM with 150 μM ATP. The experimental conditions are described in Methods. Black arrows in the solid style indicate the time of adding proteins.

Figure 6. Stimulating effect upon mutation of E397.

A Stopped-flow kinetics of the antiparallel G4 (AP-S_{16}-TelG4) unwinding activity of *To*Pif1 and its mutants. The different curves represent an average of 3-4 independent experiments for each substrate. The solid line represents the fit of data according to Eq. (2).

B E397A mutant stimulating effect determined with parallel G4 DNA.

C-E The typical FRET trajectories of antiparallel G4 ($smG4^{AP-424}$) unwinding catalyzed by a different point mutations on E397 using single-molecular FRET assays. The dashed magenta and red lines represent the fully folded and unfolded states of G4, respectively. The different unwinding curves represent an average of 3-4 independent experiments for each substrate. Black arrows in the solid style indicate the time of adding proteins.

F Error bar plot of unwinding time of E397A, E397L, and E397H determined from unfolding curves of (C), (D), and (E), respectively (n=3 biological replicates). The experimental conditions are described in Materials and Methods and data are presented as means ± SD.

Expanded View FIGURE LEGENDS

Figure EV1. *To*Pif1 recognition for G4 and ssDNA.

A CD experiments of G4 DNA.

B Left panel: DNA binding of *To*Pif1 to $G4^{CEB}$ with 1 μM T_{12} (black line) or 1μM $G4^{CEB}$ (blue line) or none of them (red line). The binding of Fitting of the data to the Eq.1 yields a $K_{d,app}$ value for *To*Pif1-$G4^{CEB}$ of 30.30 ± 1.17 nM, a $K_{d,app}$ value for *To*Pif1-$G4^{CEB}$ (added with 1 μM T_{12}) of 30.91 ± 2.32 nM. Right panel: DNA binding of *To*Pif1 to T_{12} with 1 μM $G4^{CEB}$ (red line) or 1 μM T_{12} (blue line) or none of them (black line). Fitting of the data to the Eq.1 yields a $K_{d,app}$ value for *To*Pif1-T_{12} of 29.10 ± 0.53 nM, a $K_{d,app}$ value for *To*Pif1-T_{12} (added with 1 μM $G4^{CEB}$) of 26.50 ± 0.71 nM. DNA binding assays were carried out in buffer A (25 mM Tris-HCl (pH 7.5), 50 mM NaCl, 2 mM $MgCl_2$ and 2 mM DTT) with 0.5 mM ADP·AlF_4.

C Stopped-flow unwinding kinetic curves of *To*Pif1 unwinding G4 DNA ($G4^{AP-424}$, $G4^{P-214}$, $4G4^{AP-333}$) in buffer A with 1 mM ADP·AlF_4.

D CD analysis of $G4^{AP-424}$ and $G4^{P-214}$.

E Normalized melting curves from CD melting assays for $G4^{AP-424}$ and $G4^{P-214}$ at different temperatures (20-75℃).

F Cartoon representation of *To*Pif1 (molecule *a*) with $G4^{T6/T8}$ in the presence of ADP·AlF_4 shown as a black stick and loop3 highlighted in magenta.

G Cartoon representation of *To*Pif1-dT_{15}-ADP·AlF_4 (PDB: 6S3M (Dai *et al.*, 2021)) with ADP·AlF_4 shown as a black stick and loop3 highlighted in magenta.

H Cartoon representation of *Ba*Pif1-dT_{10}- ADP·AlF_4 (PDB: 5FHE (Zhou *et al.*, 2016)) with ADP·AlF_4 shown as a black stick and loop3 highlighted in magenta.

I Cartoon representation of *Sc*Pif1p-poly (G3T5)-ADP·AlF_4 (PDB: 5O6B (Lu *et al*, 2018b)) with ADP·AlF_4 shown as a black stick and loop3 highlighted in magenta.

Figure EV2. Comparison of domain 2B of the molecule *a* of the *To*Pif1-$G4^{T6/T8}$-ADP·AlF_4 ternary complex and the previously determined *To*Pif1-ss/dsDNA structure (PDB: 6S3H (Dai *et al.*, 2021)).

A Cartoon representation of molecule *a* in the *To*Pif1-$G4^{T6/T8}$-ADP·AlF_4 structure with domain 2B and the center mass (sphere) of domain 2B is highlighted in blue. G-quadruplex (G4) DNA is shown in red.

B Cartoon representation of the *To*Pif1-S_7D_{11}-ADP·AlF_4 structure (PDB: 6S3H (Dai *et al.*, 2021)) with domain 2B shown in magenta. The center mass of domain 2B is shown as a red sphere.

C Superposition on domain 1A of the structures in (A) and (B).

D Domain 2B and G4 DNA of molecule *a* in the *To*Pif1-$G4^{T6/T8}$-ADP·AlF_4 structure with the close-up view of interactions between *To*Pif1 and the G4 tetrad I.

E Domain 2B and modeled duplex DNA in the *To*Pif1-S_7D_{11}-ADP·AlF_4 structure (PDB: 6S3H (Dai *et al.*, 2021)).

F Superposition on domain 1A of the structure in (C) with only domain 2B and G4 DNA being shown.

Figure EV3. SAXS results of *To*Pif1-$G4^{T6/T8}$-ADP·AlF_4.

A Fit curve of the SAXS data of *To*Pif1- $G4^{T6/T8}$-ADP·AlF_4 with two modeled *To*Pif1 molecules bound to the substrate $G4^{T6/T8}$ calculated with Crysol.

B The model of *To*Pif1- $G4^{T6/T8}$-ADP·AlF_4 superimposed on the *ab initio* envelope calculated with DAMMIF.

C The SAXS Model of *To*Pif1-$G4^{T6/T8}$ -ADP·AlF_4.

D Crystal structure of *To*Pif1-$G4^{T6/T8}$ -ADP·AlF_4.

E Superposition for SAXS model and crystal structure of *To*Pif1-$G4^{T6/T8}$ -ADP·AlF_4.

Figure EV4. Functional validation of *To*Pif1 mutants and ligand interaction diagram of T6/G7 in the *To*Pif1 complex compared with the guanine-specific pocket (GSP) in the *Cs*RecQ structure.

A CD results of *To*Pif1 and its mutants.

B ATP hydrolysis of *To*Pif1 and its mutants with or without DNA effector ($S_{18}H_{11}$).

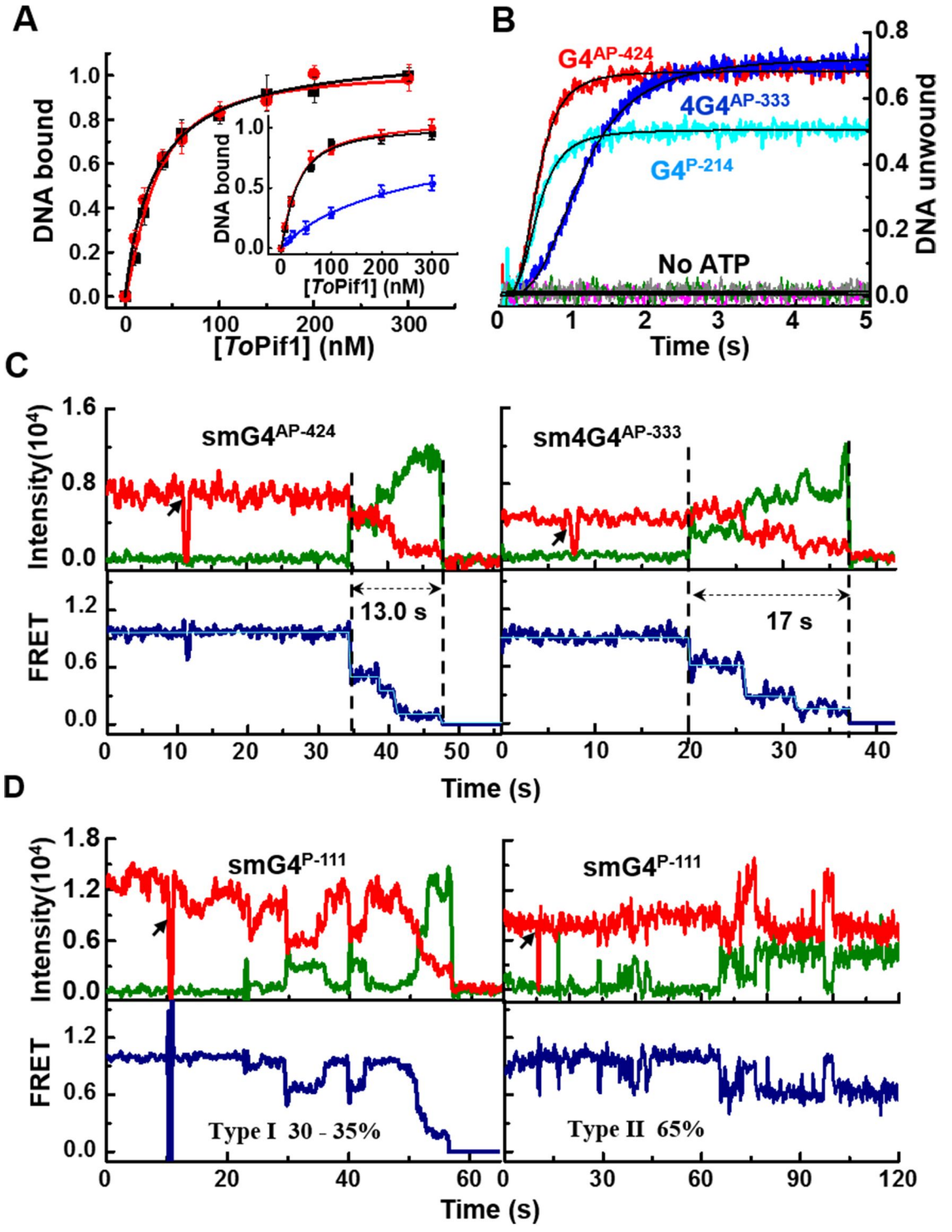
C Ligand interaction between T6 and *To*Pif1 in the *To*Pif1-$G4^{T6/T8}$- ADP·AlF_4 structure.

D Ligand interaction between G7 and *To*Pif1 in the *To*Pif1-GR_{17}- ADP·AlF_4 structure (PDB: 7BIL (Dai *et al.*, 2021)).

E Ligand interaction between G21 and *Cs*RecQ in *Cs*RecQ-ssDNA structure (PDB: 6CRM (Voter *et al.*, 2018)).

Figure EV5. Sequence alignment of Pif1 helicase from bacteria, humans, and yeast.

Thermus bacteria Pif1 (1-8) are from *Thermus oshimai* (*To*Pif1), *Thermodesulfovibrio yellowstonii* (*Ty*Pif1), *Sulfurihydrogenibium sp* (*Su*Pif1), *Anaerobaculum hydrogeniformans* (*Ah*Pif1), *Thermodesulfobacterium commune* (*Tc*Pif1), *Deferribacter desulfuricans* (*De*Pif1), *Mycolicibacterium thermoresistibile* (*Mt*Pif1), *Synergistales bacterium* (*Sb*Pif1). Bacteria at room temperature are from *Bacteroides sp* (*Bs*Pif1). Homo sapiens and *Saccharomyces cerevisiae* are labeled as *Hs*Pif1 and *Sc*Pif1. Residues involved in G4-Recognizing Surface (GRS) are shown with green stars.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[ToPif1] (nM)
B
G4AP-424
4G4AP-333
G4P-214
No ATP
DNA unwound
Time (s)
C
smG4AP-424
sm4G4AP-333
Intensity(10^4)
FRET
13.0 s
17 s
Time (s)
D
smG4P-111
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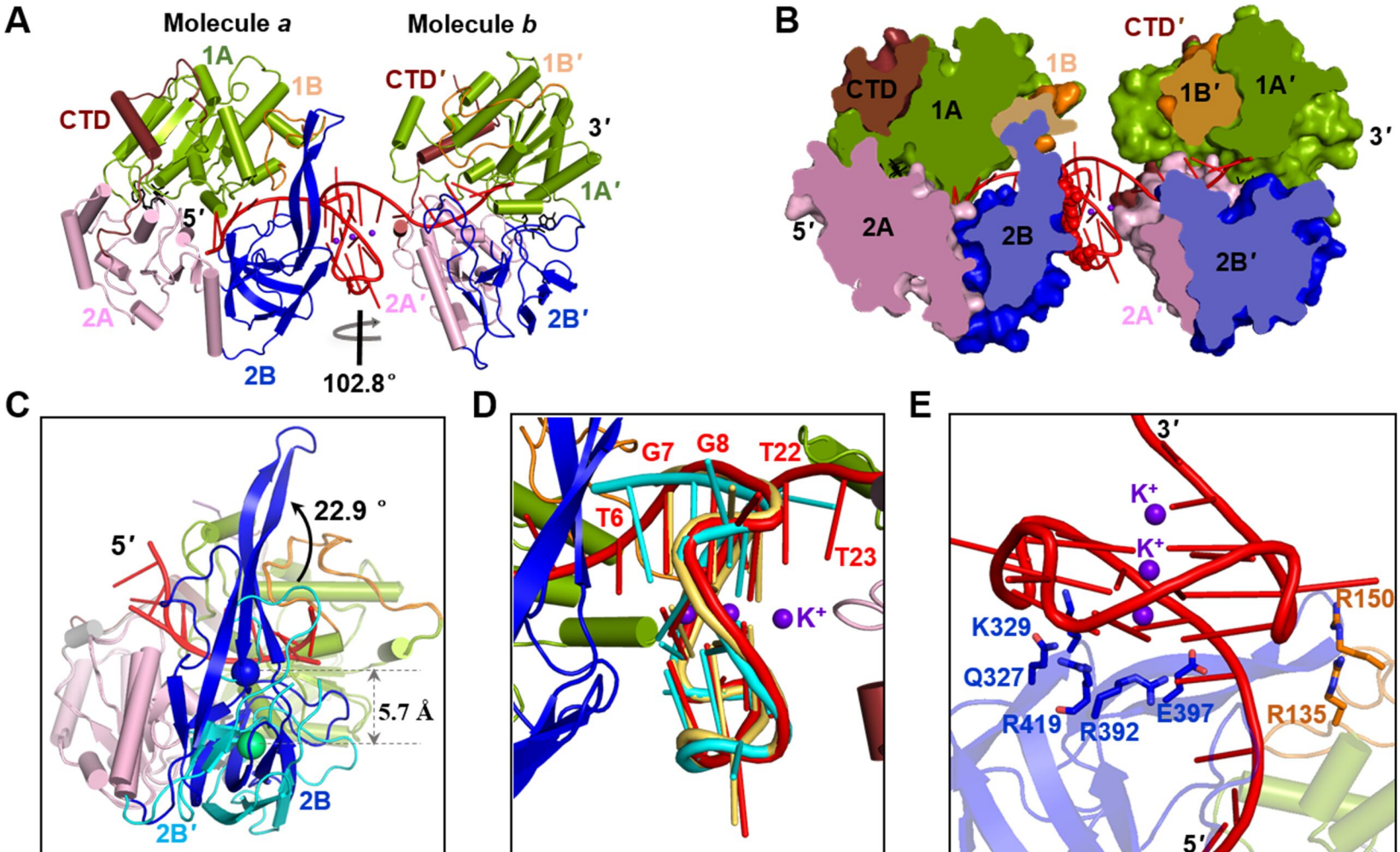
Intensity(10^4)
FRET
Type I 30 - 35%
Type II 65%
Time (s)


A
Molecule a
Molecule b
1A
1B
CTD
CTD′
1B′
3′
1A′
5′
2A
2A′
2B
2B′
102.8°
B
CTD
1A
1B
CTD′
1B′
1A′
3′
5′
2A
2B
2B′
2A′
C
22.9 °
5′
5.7 Å
2B
2B′
D
G7
G8
T22
T6
T23
K+
E
3′
K+
K+
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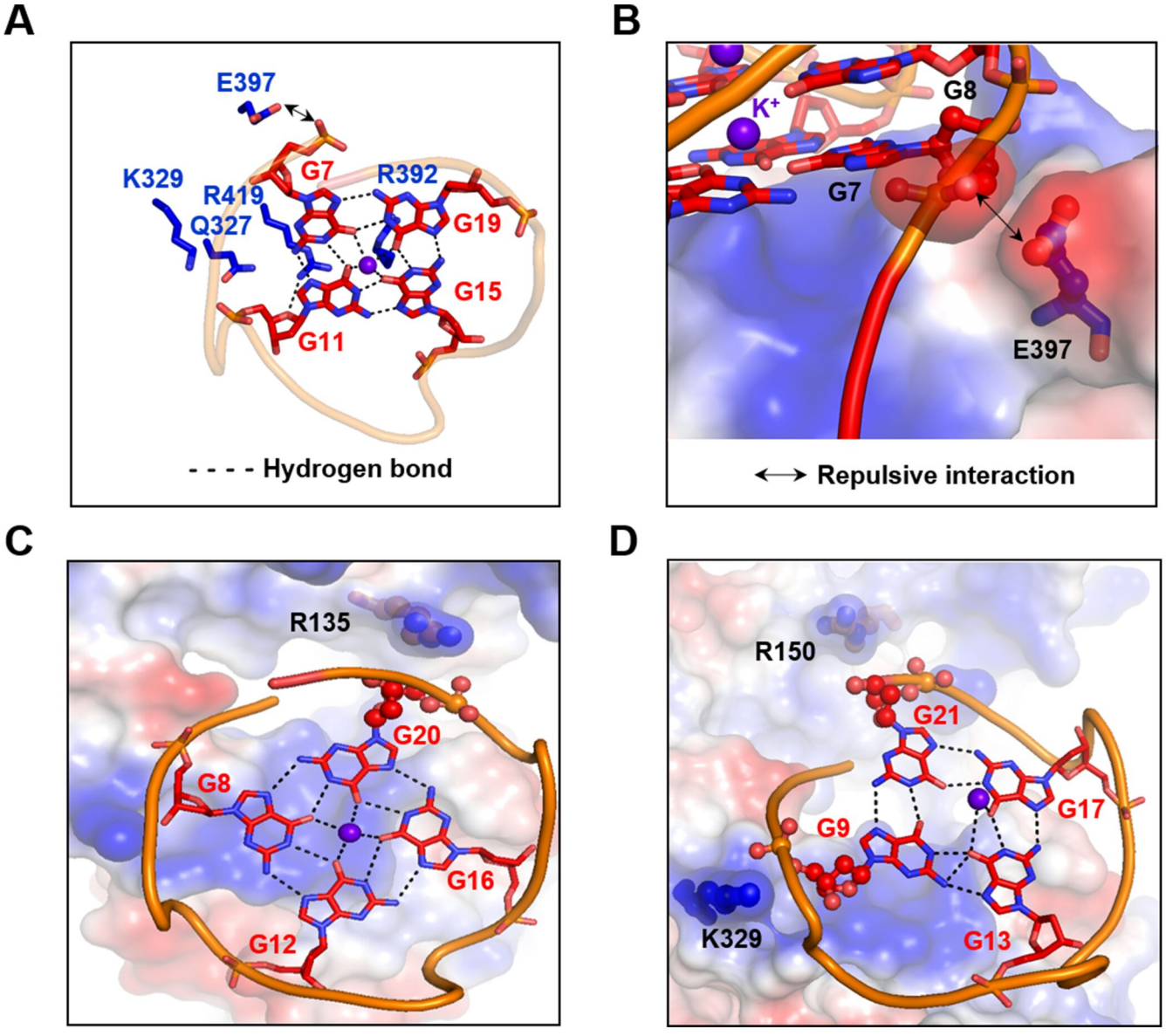
E397
R135
5′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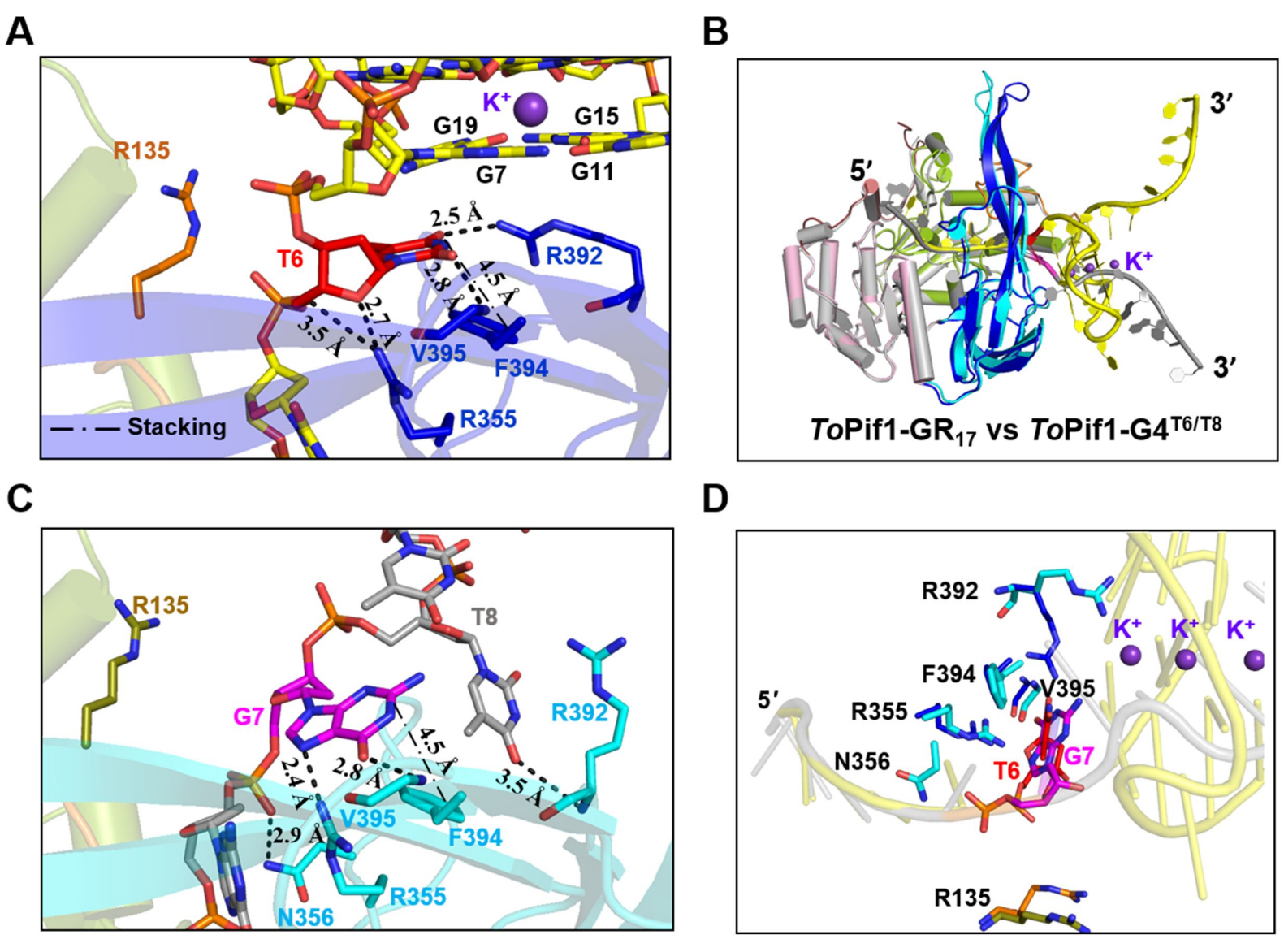
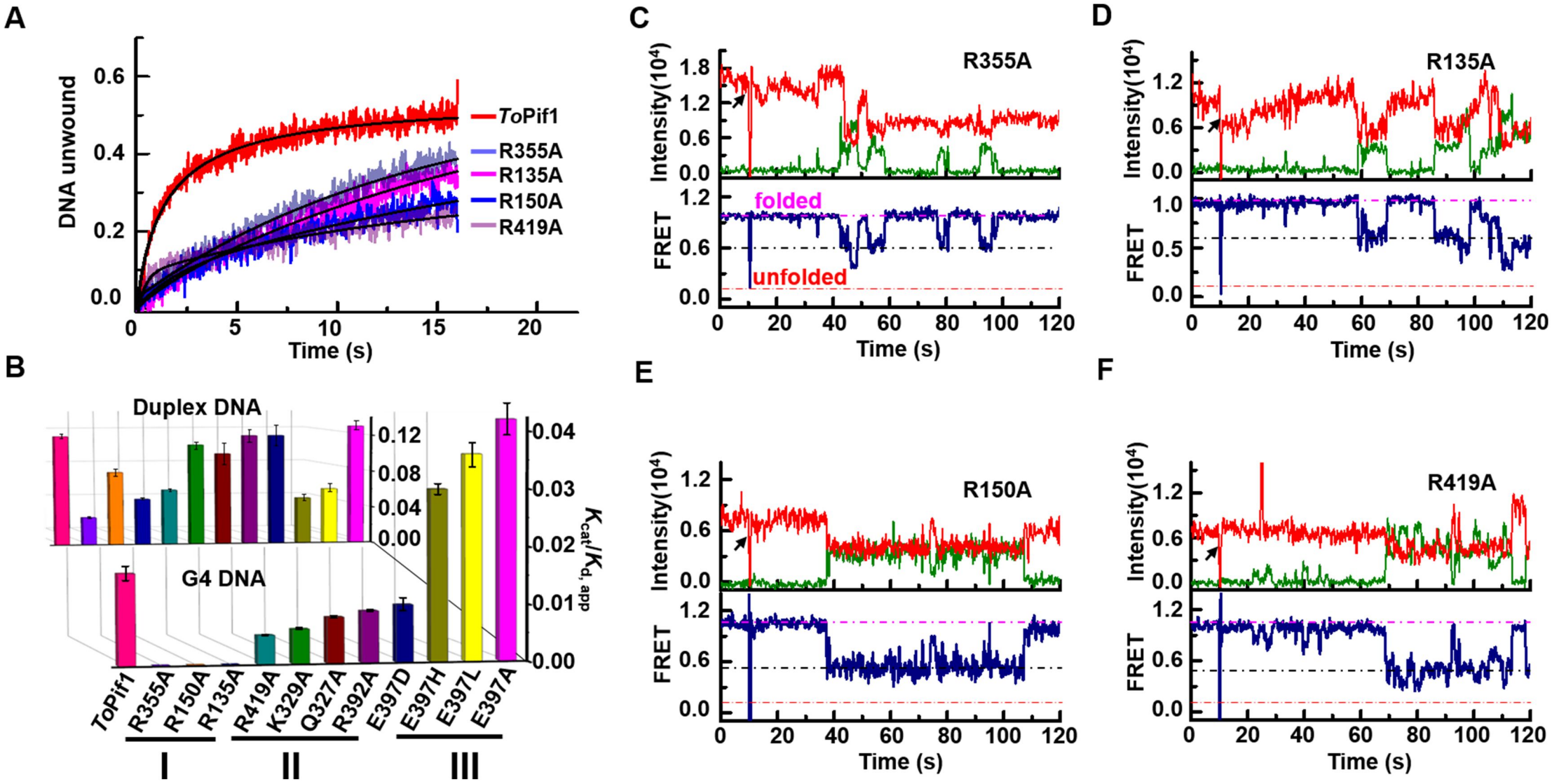
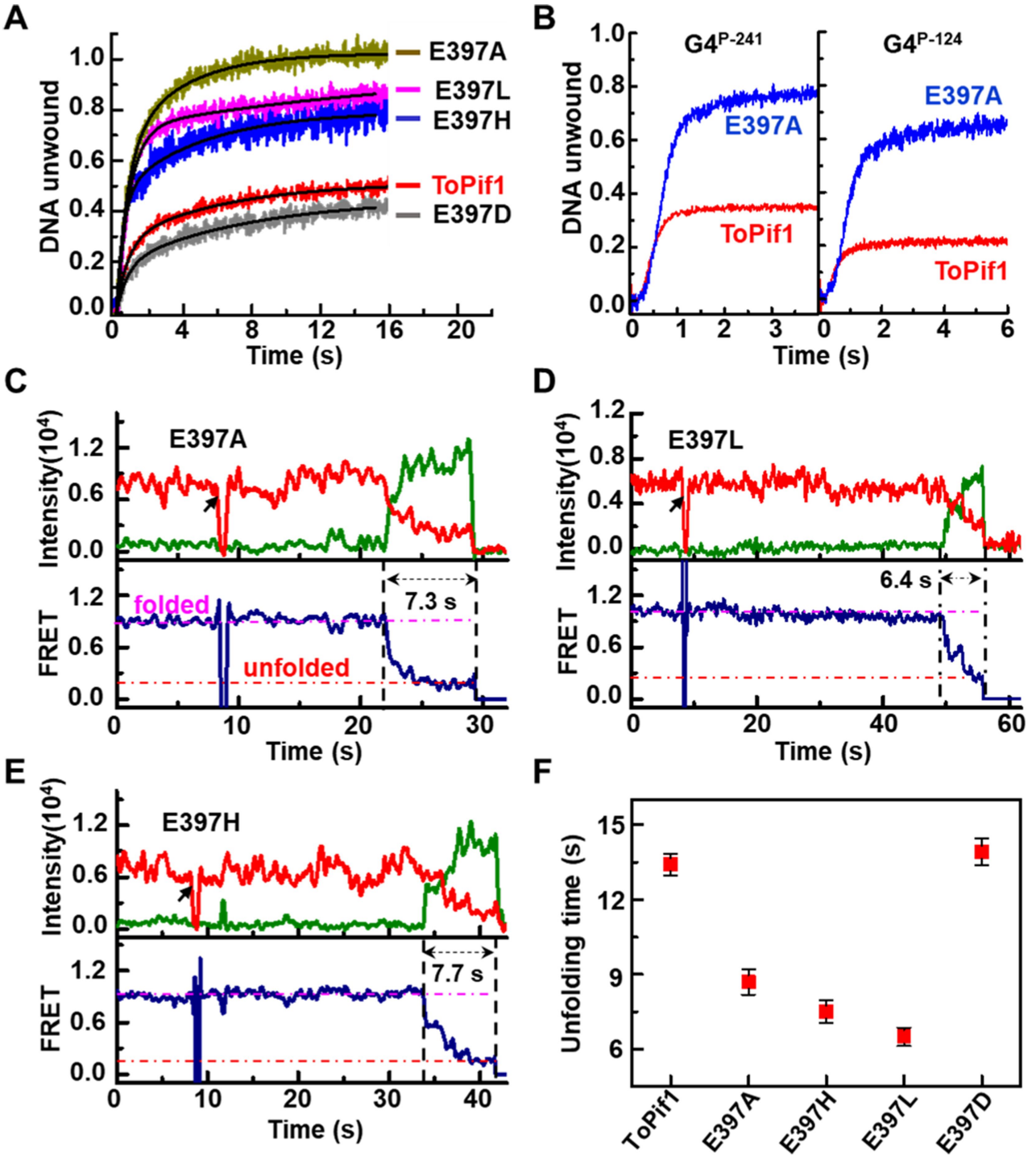
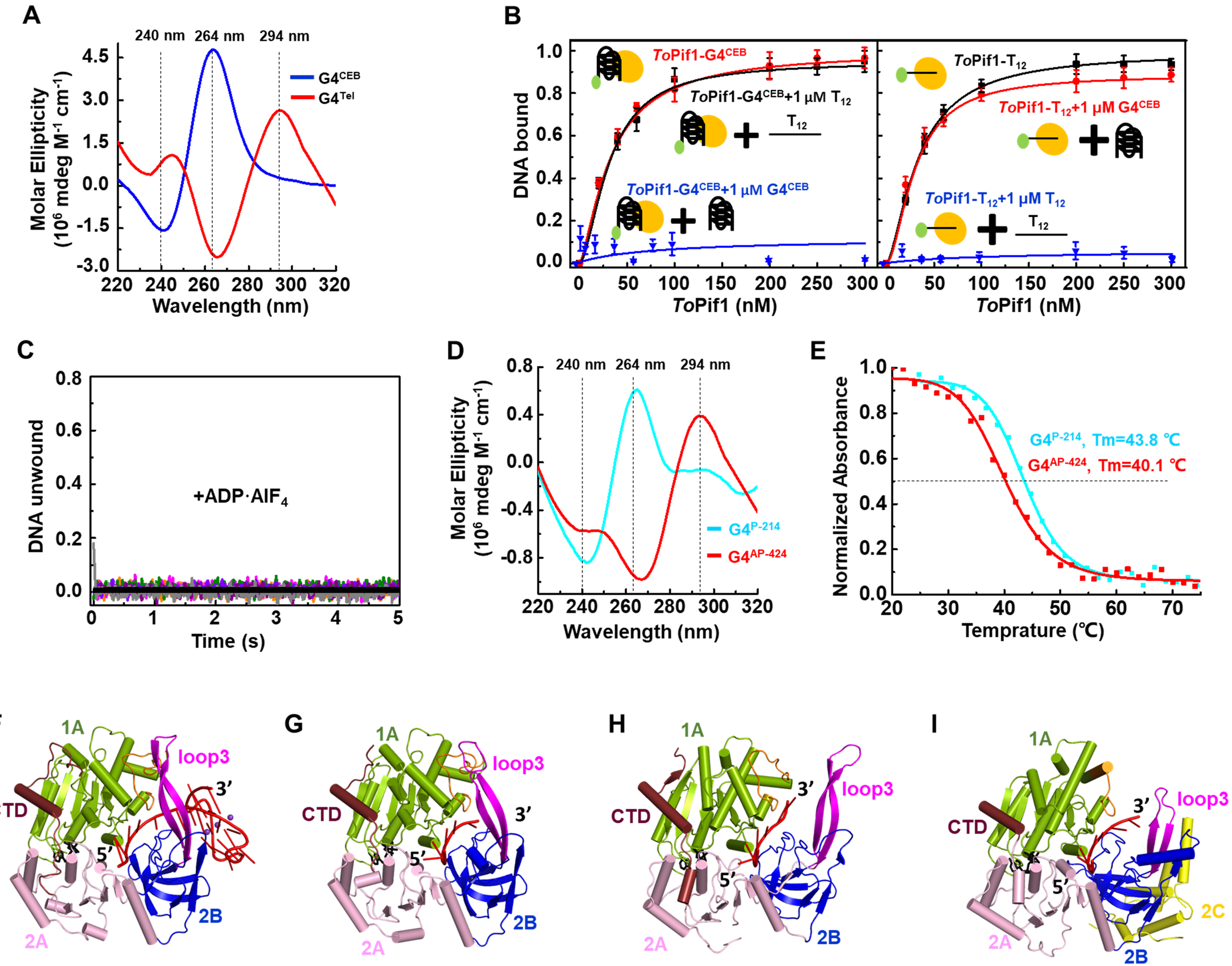
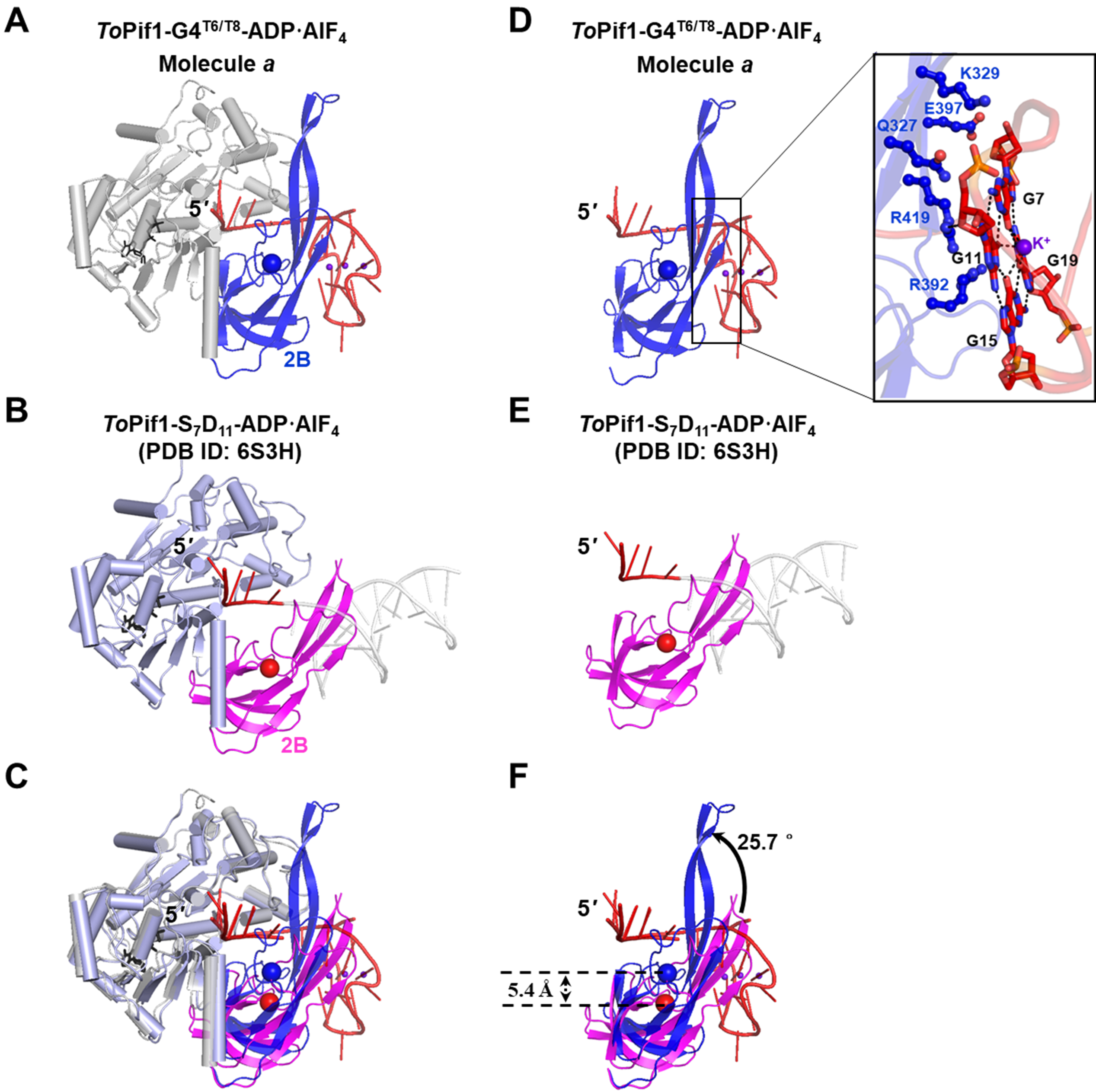
E397
K329
R419
Q327
G7
R392
G19
G15
G11
- - - - Hydrogen bond
B
K+
G8
G7
E397
Repulsive interaction
C
R135
G20
G8
G16
G12
D
R150
G21
G17
G9
K329
G13


A
K+
G19
G15
R135
G7
G11
2.5 Å
T6
R392
2.8 Å
4.5 Å
2.7 Å
3.5 Å
V395
F394
R355
Stacking
B
3′
5′
K+
3′
ToPif1-GR17 vs ToPif1-G4T6/T8
C
R135
T8
G7
R392
4.5 Å
2.4 Å
2.8 Å
3.5 Å
V395
F394
2.9 Å
R355
N356
D
R392
K+
K+
K+
F394
V395
5′
R355
N356
T6
G7
R135


A
DNA unwound
Time (s)
ToPif1
R355A
R135A
R150A
R419A
B
Duplex DNA
G4 DNA
$K_{cat}/K_{d,app}$
ToPif1
R355A
R150A
R135A
R419A
K329A
Q327A
R392A
E397D
E397H
E397L
E397A
I
II
III
C
R355A
Intensity(10^4)
FRET
folded
unfolded
Time (s)
D
R135A
E
R150A
F
R419A

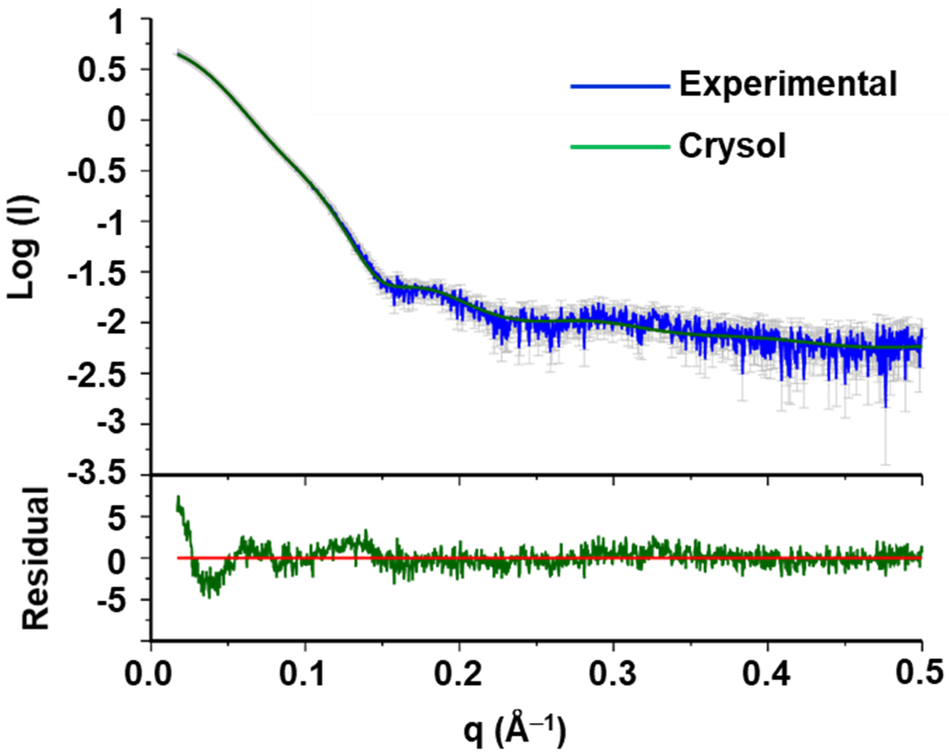

A
DNA unwound
Time (s)
E397A
E397L
E397H
ToPif1
E397D
B
G4P-241
G4P-124
E397A
ToPif1
DNA unwound
Time (s)
C
E397A
Intensity(10^4)
FRET
folded
unfolded
7.3 s
Time (s)
D
E397L
Intensity(10^4)
FRET
6.4 s
Time (s)
E
E397H
Intensity(10^4)
FRET
7.7 s
Time (s)
F
Unfolding time (s)
ToPif1
E397A
E397H
E397L
E397D


A
240 nm
264 nm
294 nm
G4CEB
G4Tel
Molar Ellipticity (10^6 mdeg M^-1 cm^-1)
Wavelength (nm)
B
ToPif1-G4CEB
ToPif1-G4CEB+1 μM T12
ToPif1-G4CEB+1 μM G4CEB
ToPif1-T12
ToPif1-T12+1 μM G4CEB
ToPif1-T12+1 μM T12
T12
DNA bound
ToPif1 (nM)
C
+ADP·AlF4
DNA unwound
Time (s)
D
240 nm
264 nm
294 nm
G4P-214
G4AP-424
Molar Ellipticity (10^6 mdeg M^-1 cm^-1)
Wavelength (nm)
E
G4P-214, Tm=43.8 ℃
G4AP-424, Tm=40.1 ℃
Normalized Absorbance
Temprature (℃)
F
G
H
I
1A
loop3
CTD
3'
5'
2A
2B
2C


A
ToPif1-G4T6/T8-ADP·AlF4
Molecule a
5′
2B
B
ToPif1-S7D11-ADP·AlF4
(PDB ID: 6S3H)
5′
2B
C
5′
D
ToPif1-G4T6/T8-ADP·AlF4
Molecule a
5′
K329
E397
Q327
R419
G7
K+
G11
G19
R392
G15
E
ToPif1-S7D11-ADP·AlF4
(PDB ID: 6S3H)
5′
F
25.7 °
5′
5.4 Å

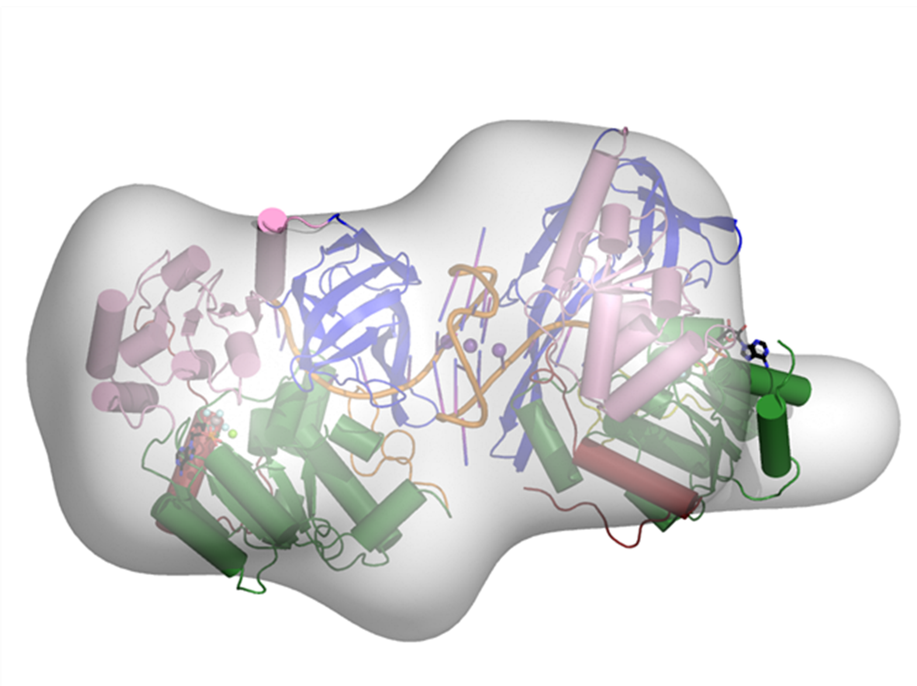
A

Crysol fit of *To*Pif1-G4$^{T6/T8}$-ADP·AlF$_4$, χ^2 = 1.905



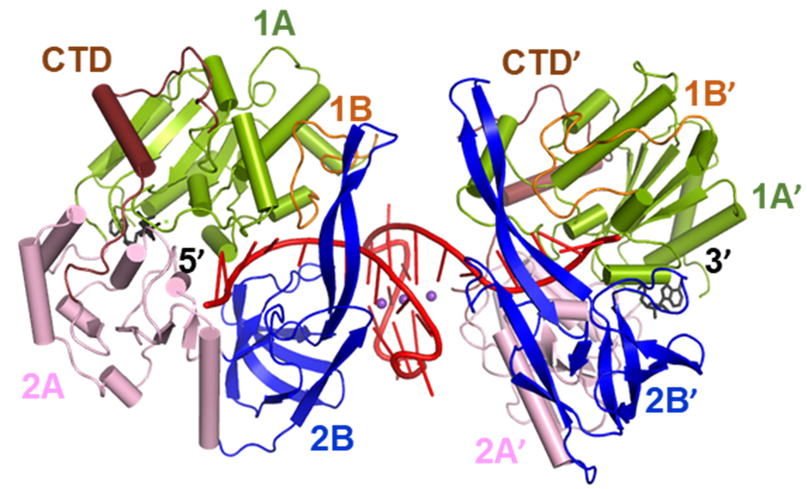
B

SAXS Model of *To*Pif1-G4$^{T6/T8}$-ADP·AlF$_4$



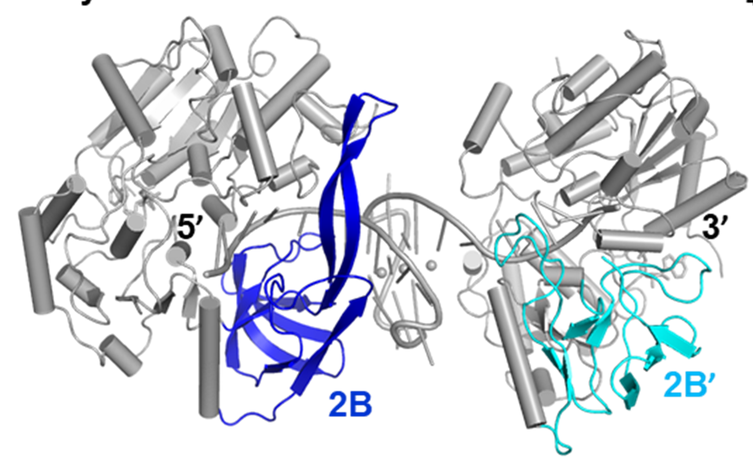
C

SAXS Model of *To*Pif1-G4$^{T6/T8}$-ADP·AlF$_4$

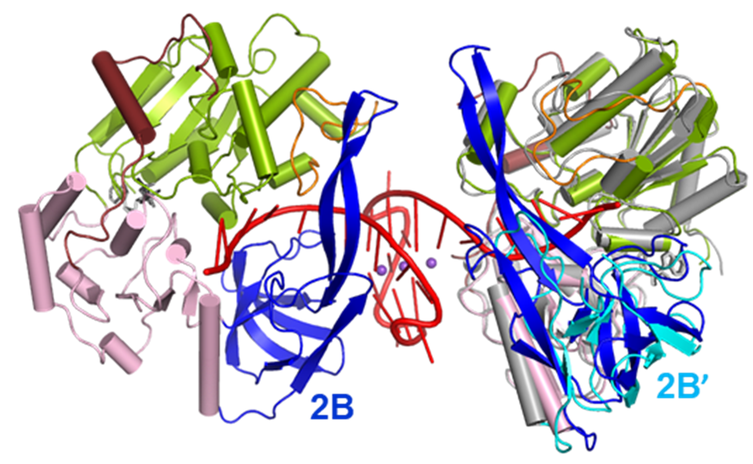


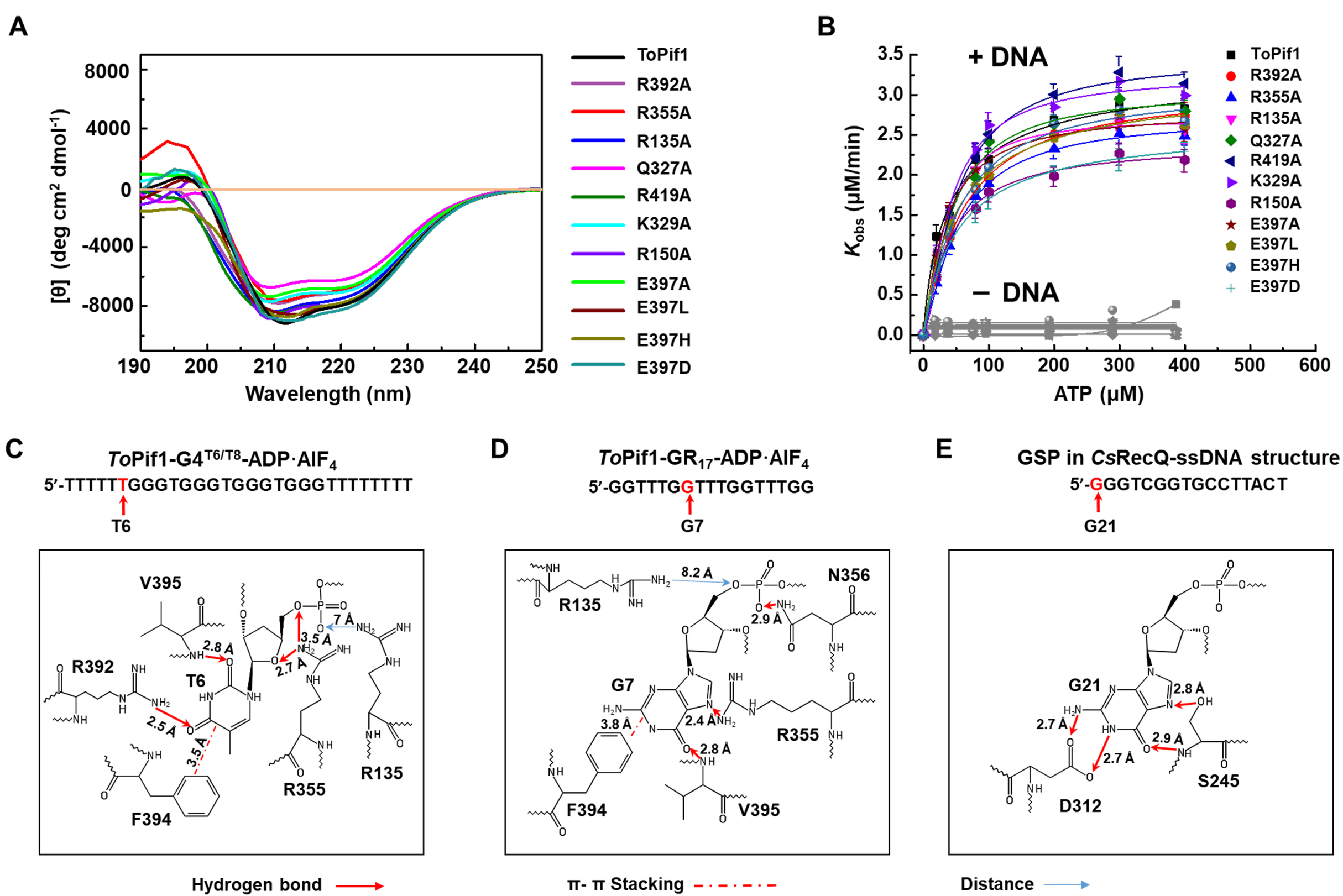
D

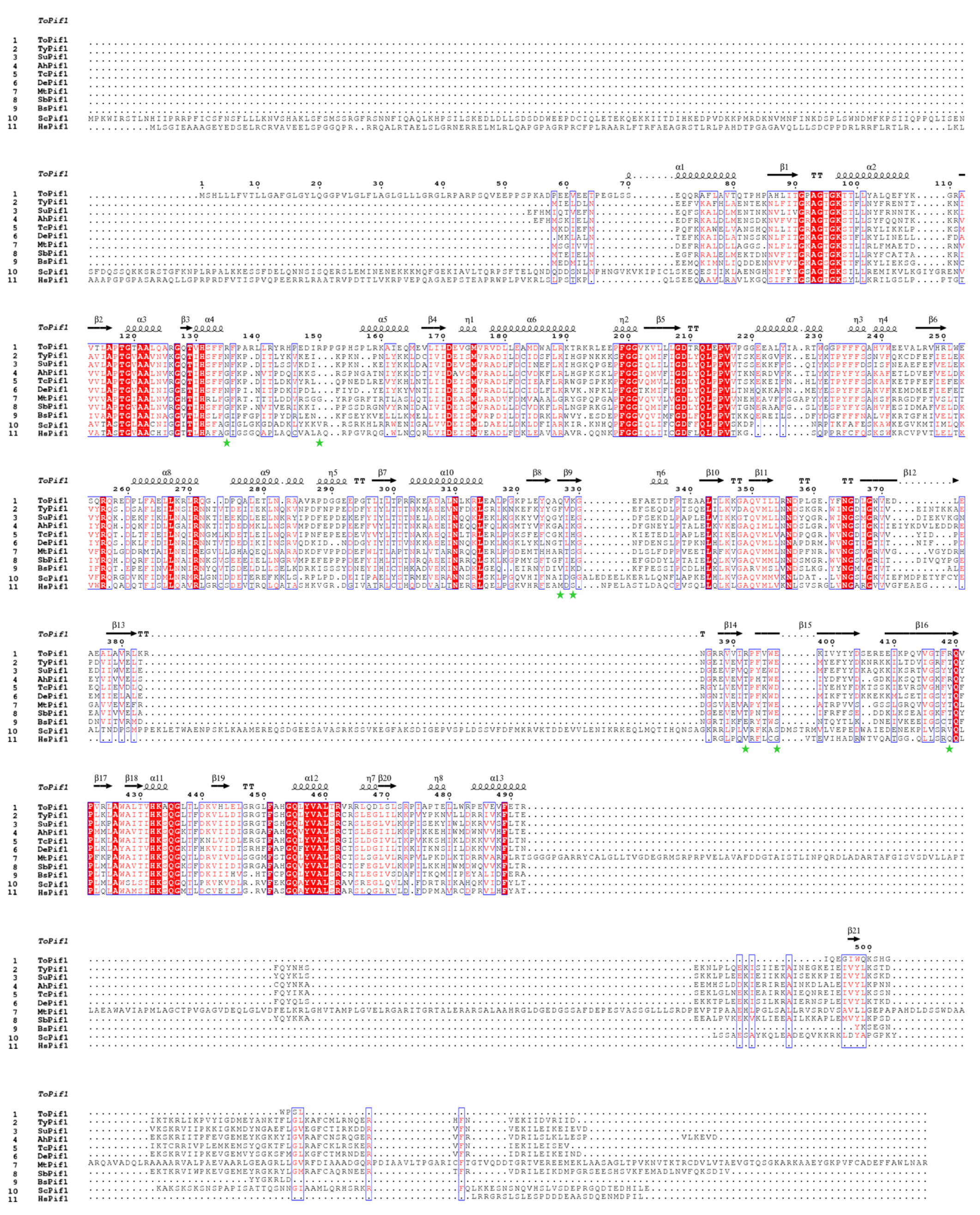
Crystal structure of *To*Pif1-G4$^{T6/T8}$-ADP·AlF$_4$



E Superposition for SAXS model and crystal structure of *To*Pif1-G4$^{T6/T8}$




A
8000
4000
0
-4000
-8000
[θ] (deg cm² dmol⁻¹)
190
200
210
220
230
240
250
Wavelength (nm)
ToPif1
R392A
R355A
R135A
Q327A
R419A
K329A
R150A
E397A
E397L
E397H
E397D
B
+ DNA
− DNA
3.5
3.0
2.5
2.0
1.5
1.0
0.5
0.0
Kobs (μM/min)
0
100
200
300
400
500
600
ATP (μM)
ToPif1
R392A
R355A
R135A
Q327A
R419A
K329A
R150A
E397A
E397L
E397H
E397D
C
ToPif1-G4T6/T8-ADP·AlF4
5′-TTTTTTGGGTGGGTGGGTGGGTTTTTTTT
T6
V395
R392
T6
2.8 Å
2.5 Å
3.5 Å
7 Å
3.5 Å
2.7 Å
R355
R135
F394
D
ToPif1-GR17-ADP·AlF4
5′-GGTTTGGTTTGGTTTGG
G7
R135
8.2 Å
N356
2.9 Å
G7
3.8 Å
2.4 Å
2.8 Å
R355
F394
V395
E
GSP in CsRecQ-ssDNA structure
5′-GGGTCGGTGCCTTACT
G21
G21
2.8 Å
2.7 Å
2.9 Å
2.7 Å
S245
D312
Hydrogen bond
π- π Stacking
Distance


ToPif1
TyPif1
SuPif1
AhPif1
TcPif1
DePif1
MtPif1
SbPif1
BsPif1
ScPif1
HsPif1