Bacterial DnaB helicase interacts with the excluded strand to regulate unwinding

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*Running title: *DnaB excluded strand interaction*

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Keywords: DNA helicase / DNA replication / DnaB / steric exclusion and wrapping / unwinding

ABSTRACT

Replicative hexameric helicases are thought to unwind duplex DNA by steric exclusion (SE), where one DNA strand is encircled by the hexamer and the other is excluded from the central channel. However, interactions with the excluded strand on the exterior surface of hexameric helicases have also been shown to be important for DNA unwinding, giving rise to the steric exclusion and wrapping (SEW) model. For example, the archaeal *Sso*MCM helicase has been shown to unwind DNA via a SEW mode to enhance unwinding efficiency. Using single molecule FRET (smFRET), we now show that the analogous *E. coli* DnaB helicase also interacts specifically with the excluded DNA strand during unwinding. Mutation of several conserved and positively-charged residues on the exterior surface of *Ec*DnaB resulted in increased interaction dynamics and states compared to wild-type. Surprisingly, these mutations also increased the DNA unwinding rate, suggesting that electrostatic contacts with the excluded strand act as a regulator for unwinding activity. In support, experiments neutralizing charge of the excluded strand with a morpholino substrate instead of DNA also dramatically increased the unwinding rate. Of note, although the stability of the excluded strand was nearly identical for *Ec*DnaB and *Sso*MCM, these enzymes are from different superfamilies and unwind DNA with opposite polarities. These results support the SEW model of unwinding for *Ec*DnaB that expands on the existing SE model of hexameric helicase unwinding to include contributions from the excluded strand to regulate the DNA unwinding rate.

INTRODUCTION

Hexameric helicases are structurally conserved toroidal enzyme complexes capable of translocating and separating double-stranded DNA (dsDNA) into two single-strands (ssDNA) providing templates for DNA replication. They utilize the inherent energy from ATP hydrolysis to translocate along an encircled strand physically displacing the opposing excluded strand. The translocation polarity of hexameric helicases differs among helicase superfamilies (SF), defined by the organization and conservation of various folds (1). SF4 helicases from bacteria and associated phages (T4 and T7) include RecA-like folds and have 5'-3' unwinding polarity, translocating on the lagging strand while SF6 helicases from archaea and eukaryotes have AAA^+ folds and 3'-5' unwinding polarity, translocating on the leading strand (2). Although these two well-studied helicase families have globally conserved structural features, their amino acid sequences, structural folds, and unwinding polarities are not. Moreover, the precise contacts with each DNA strand to facilitate duplex unwinding are not known.

The bacterial replicative helicase, DnaB, has been shown to encircle the 5' lagging strand in its central channel. Orientation of DnaB binding on ssDNA was shown to place the RecA motor C-terminal domain (CTD) adjacent to the duplex region and the N-terminal domain (NTD) towards the 5' end (3). DnaB unwinds dsDNA in a steric

exclusion (SE) mechanism, and can even translocate over two or three strands of DNA indicating plasticity within the central channel of the hexamer (4,5). Currently, various X-ray and EM structures of hexameric DnaB (with and without DNA or accessory proteins) show the hexamer as either a closed ring (6-8) or a split lock washer (9,10). ssDNA bound in the central channel adopts an A-form right-handed spiral conformation making contacts with multiple interior DNA binding loops from different subunits to pass DNA along in a hand-over-hand mechanism (9).

In addition to specific DNA contacts that exist within the central channel, further exterior contact sites are proposed to exist for hexameric helicases to aid in DNA loading and unwinding (2). Previously, we have identified an interaction path on the external surface of the archaeal (AAA^+) *Sso*MCM helicase with the excluded strand that both protects and stabilizes the complex in a forward unwinding mode (11,12). This interaction expanded the widely accepted SE model of unwinding to include contributions of the excluded strand in the mechanism. This new unwinding model was termed steric exclusion and wrapping (SEW). Recently, interactions with the excluded strand have been uncovered from a variety of hexameric helicase complexes in addition to archaeal MCM including: E1 (13), SV40 LargeT (14), T7gp4 (15,16), *Ec*DnaB (17,18), TWINKLE (19), and the eukaryotic Cdc45/MCM2-7/GINS (CMG) complex (20,21). It is hypothesized that external interactions with the excluded strand will not only protect ssDNA, but also stabilize the helicase/DNA complex, and modulate the unwinding rate.

In this study, we examined whether *Ec*DnaB has a similar specificity for exterior interactions with the excluded single-strand DNA. Single molecule FRET (smFRET) experiments were employed to directly detect *Ec*DnaB binding to the excluded strand and compared with *Sso*MCM binding. The absolute FRET states, transition probabilities, and dwell times were strikingly similar between *Ec*DnaB and *Sso*MCM, even though they reside in different superfamilies and have opposite unwinding polarities. Mutation of several conserved external positively charged residues on *Ec*DnaB differentially altered the observed FRET states and binding dynamics consistent with disruption of the excluded strand binding path. Notably, those external SEW mutations on *Ec*DnaB dramatically increased the dsDNA unwinding rate compared to wild-type (WT). Neutralizing the negative charge on the excluded stand with a morpholino substrate increased the unwinding rate for wild-type *Ec*DnaB consistent with electrostatic interactions regulating activity. This is the first molecular explanation for controlling/regulating the rate of DNA unwinding through specific external surface interactions on the helicase with the excluded strand.

RESULTS

EcDnaB Interacts with the Excluded-Strand - Previously, we have shown that interactions with the excluded strand exist for the 3'-5' hexameric *Sso*MCM helicase (11,22). Using a similar smFRET approach, we sought to examine whether analogous contacts on the exterior of *Ec*DnaB also interact with the excluded strand despite the opposing 5'-3' translocation polarity. Three separate model fork substrates, 30/30 (DNA43/DNA44), 40/30 (DNA111/DNA44), and 50/30 (DNA116/DNA44), composed of an 18 bp duplex with 30 (dT) on the 5'-lagging strand and 30, 40, and 50 (dT) on the 3'-leading strand, respectively, were used. DNA forks alone result in low FRET signals as a result of Cy3 and Cy5 on the termini of the fork arms not being in close proximity (**Fig. 1A**). Additional nucleotides on the leading strand 3'-arm of the fork further decrease the FRET efficiency as expected. Addition of *Ec*DnaB to each of these substrates shifts the signal to higher FRET states (**Fig. 1B**). *Ec*DnaB preferentially encircles the 5'-strand (3), and a titration of *Ec*DnaB onto 30/30 showed little to no variation in the resulting histogram profiles, suggesting that only one hexamer can be accommodated by the fork substrate over a large concentration range (**Fig. 1C**). Should a second hexamer encircle the opposing 3'-strand, the FRET values would decrease. The occurrence of a high FRET state is consistent with an interaction of the excluded Cy3 3'-strand on the external surface of *Ec*DnaB, analogous to *Sso*MCM binding to DNA and consistent with the proposed SEW model of helicase interaction and unwinding (11).

*Ec*DnaB loaded onto the 40/30 substrate produced an almost exclusively high FRET state (>0.9); the 50/30 fork produced a bimodal distribution of high (>0.9) and medium (~0.5) FRET states, and the 30/30 fork yielded a bimodal distribution of two high FRET (0.8 & >0.9) states in the presence of *Ec*DnaB. The interaction with the excluded strand likely differs from varying excluded strand lengths sampling slightly different external binding paths. However, in all cases the shifts to high FRET states correspond to a stably wrapped excluded strand that places the Cy3 dye near the Cy5 dye on the encircled strand.

ExPRT Analysis of Excluded Strand Dynamics - To better visualize the FRET states, explicit transition probabilities, and dwell times on a single plot to compare different conditions, mutants, and helicases, we developed Explicit Probability and Rate Transition (ExPRT) Plots (**Fig. 2A**). The positions of the circular markers correspond to transitions between specific FRET states: initial FRET state on the x-axis and the final FRET state on the y-axis. The initial and final FRET states for a particular transition refer to the observed FRET states immediately preceding and immediately following the transition of interest, respectively. The size and color of each marker correspond to the probability of that transition occurring within a measured trace and the average dwell time of the state preceding the transition, respectively.

A comparison of analyses between established programs, the HaMMY and Transition Density Plot (TDP) programs (**Fig. 2B**) (23) as well as the POpulation-weighted and Kinetically-Indexed Transition density (POKIT) program (**Fig. 2C**) (24), and the ExPRT program is shown. Each program analyzed and visualized identical data corresponding to *Ec*DnaB bound to the 30/30 substrate and can distinguish states (**Fig. 2D**). Each plot illustrates that the transitions between FRET states of ~0.8 and ~0.95 are the most frequent. The TDP program analysis works on a trace-by-trace basis and is able to reveal heterogeneities in the transition data that can be missed by programs that work on stitched datasets such as the POKIT and ExPRT programs. However, despite the TDP program's ability to gather probability and rate values, these values are not directly visualized by the resulting plot. The POKIT program bins the probabilities and rates of each transition into user-defined ranges and produces plots that allow for some level of quantitative comparison between experimental conditions. However, these plots fail to display explicit probabilities and rate values. Determining explicit transitions, probabilities, and rates without binning is an inherent advantage of single-molecule methods that allows for extensive insights into the dynamics and kinetics of molecular interactions and enzymatic activities. The ExPRT analysis program extracts these explicit values from the smFRET data and visualizes them directly in a single plot. This allows users and readers to easily make comparisons between datasets on the most detailed level. Therefore, the ExPRT plots provide a useful advance in the investigation and comparison of the probability and kinetics of smFRET dynamics.

***Ec*DnaB and *Sso*MCM Interact the Excluded Strand Similarly -** The bimodal distribution observed for *Ec*DnaB on 30/30 fork substrate was very similar to the distribution produced by the archaeal MCM helicase on the same substrate (**Fig. 3A**) (11). In fact, the single-molecule traces for *Ec*DnaB and *Sso*MCM exhibited strikingly similar dynamics between two high FRET states (~0.95 and ~0.8) (**Fig. 3B-C**). This is highlighted by the ExPRT plots for *Ec*DnaB compared to *Sso*MCM on the 30/30 fork substrate (**Fig. 3D-F**). In both cases, the excluded strand has a reversible transition between two high FRET states (~0.8 and ~0.95) that is exhibited by ~70% of the molecules analyzed. For both *Ec*DnaB and *Sso*MCM, there is a preference for the ~0.95 FRET state, indicated by the longer dwell times measured for that state (shades of orange vs. green/yellow). In addition, both data sets exhibit a reversible lower probability transition between each of high FRET states and a medium FRET state of ~0.55. Altogether, it is noteworthy that similarities in the FRET states, transitions, and dwell times exist for *Ec*DnaB and *Sso*MCM on the 30/30 fork even though these two helicases belong to different superfamilies with low sequence homology, exist in different domains of life (bacteria vs. archaea), and have opposite unwinding polarities. The similarities are suggestive of a common SEW unwinding mechanism across diverse replicative helicases.

Exterior Surface Mutations of *Ec*DnaB Alter Excluded-Strand Wrapping - As positively charged residues on the surface of *Sso*MCM had previously been shown to support an external interaction with the excluded strand (11), similar surface exposed and conserved residues were identified based on a homology model for *Ec*DnaB (**Fig. 4**). Four *Ec*DnaB surface positions (R74, R164, K180, and R328/R329) that exist in positively charged electrostatic patches were mutated to alanine, overexpressed, and purified. All mutant proteins were consistent with a hexamer as the major peak after gel filtration (data not shown). smFRET DNA fork binding assays were performed for the wild-type and each mutant on each of the three fork templates (30/30, 40/30, 50/30). The results were analyzed and compared using traditional histograms and ExPRT plots. The mutations gave rise to several important differences in the hexamer-excluded strand interactions and dynamics. Across all the excluded strand lengths tested, *Ec*DnaB (R74A) does not sample the highest observed FRET state (E_{app}= ~0.95) observed for the WT and the other mutants. This can be seen clearly in the histograms (**Fig. 5 B** vs. **A**) and the ExPRT plots (**Fig. 5 G, L, Q** vs. **F, K, P**). R74A also produces less transitions between FRET states compared to the WT for the 30/30 and 40/30 forks because the most frequent transitions observed for the WT data are those between the highest FRET state (E_{app}= ~0.95) and lower FRET states. The absence of the 0.95 FRET state for R74A across all substrates tested suggests that R74 is necessary to close the connection of the excluded strand to the NTD traversing the entire longitudinal length of *Ec*DnaB (9).

*Ec*DnaB (K180A) bound to the 30/30 fork produces similar FRET states and dynamics when compared to the WT *Ec*DnaB, however, there are now five states compared to three (**Fig. 5 I vs. F**). Examples traces for individual molecules for WT compared to K180A as well as other mutants are shown in **Figure 6**. As an example, for molecule 51, there are ten transitions between two states, and for molecule 177, there are seventeen transitions between three states. A greater number of FRET states and transitions are indicative of a less stable and less precise interaction between the exterior surface of the helicase and the excluded strand leading to alternative binding paths. Similarly, the R164A mutant also samples a greater number of states than WT on the 40/30 (5 vs. 2 states) and 50/30 substrates (4 vs. 2 states) (**Fig. 5 M & R** vs. **K & P**). The ExPRT plots for both K180A (**Fig. 5 I vs. F**) and R164A (**Fig. 5H & F**) on the 30/30 substrate somewhat resemble the WT on the same substrate. However, the histograms of K180A and R164A (**Fig. 5C & D**) show populations that are broader than those seen for the WT and R74A (**Fig. 5A & B**), which is indicative of less stable or precise interactions between the helicase and the excluded strand as visualized in the ExPRT plots. For the R164A mutation on the 30/30 substrate (**Fig. 5H**), the most probable transitions occur at the approximately the same rate reversibly. Similar to the WT, K180A on 40/30 and 50/30 substrates show less transitions to and from the high FRET state with and a significantly longer dwell time for the high FRET state (**Fig. 5N & S**). This may indicate greater stabilization of an interaction towards the NTD of the helicase. Therefore, K180 and R164 contribute to but do not solely mediate the helicase-excluded strand interactions that give rise to the FRET states we observe for the WT.

The *Ec*DnaB (R328A/R329A) mutant shows similar FRET state transitions, probabilities of those transitions, and dwell times on the 30/30 fork when compared to the WT (**Fig. 5 J** vs. **F**). However, there are extreme differences in the binding states and dynamics on the longer DNA strands compared to WT (**Fig. 5 T vs. P**). WT *Ec*DnaB bound to the 50/30 fork shows a small fraction of traces that transition between high and medium FRET states. In comparison, *Ec*DnaB (R328A/R329A) produces almost entirely medium FRET states that are very dynamic, with many transitions and relatively short dwell times, indicative of severe destabilization of binding. In contrast to WT, very little high FRET signal from the R328A/R329A mutant on the 30/50 substrate is observed. These results indicate that R328 and R329 may be required to stabilize longer excluded strands (40 and 50mers) along the waist of the hexamer and mediate interactions between the excluded strand and other regions, such as NTD where R74 is located and responsible for the highest FRET state (E_{app}= ~0.95).

Although we primarily tested the effect of eliminating positive charge on the exterior surface of *Ec*DnaB and its effect on excluded strand

binding, there may be also be additional noncovalent binding interactions defining a path. To directly test whether electrostatic interactions exclusively define the excluded strand binding path, we titrated NaCl into WT *Ec*DnaB prebound to a 30/30 fork in our smFRET experiments. Increased salt concentrations resulted in increased dynamics (shorter dwell times), but the FRET states were not significantly affected as visualized using ExPRT plots (**Fig. 7**). The decreased dwell times while keeping the FRET states and transition probabilities constant suggests that *Ec*DnaB utilizes electrostatic interactions to mediate wrapping but that other noncovalent interactions are also important.

SEW Mutants of *Ec*DnaB Have Enhanced DNA Unwinding Activity - Mutating positively charged residues involved in the excluded-strand interaction inhibited *Sso*MCM's unwinding activity (11,22). Gel based fluorescent DNA unwinding assays were performed to determine whether these *Ec*DnaB SEW mutants have any effect on activity. **Figure 8A** shows a representative six-minute time point; however, quantification of the steady-state unwinding rates occurred over multiple time points for each mutant (**Fig. 8B**). All mutants had increased unwinding rates compared to the WT *Ec*DnaB. Specifically, R74A (26 ± 1 nM s^{-1}) and R164A (53 ± 13 nM s^{-1}) have 3-fold and 6-fold increases, respectively, over WT (9 ± 1 nM s^{-1}); while K180A (199 ± 16 nM s^{-1}) and R328A/R329A (191 ± 13 nM s^{-1}) have more than 20-fold increases in unwinding activity.

ATPase assays were performed for WT and mutant *Ec*DnaB proteins, and the rates were quantified in the absence and presence of DNA (**Fig. 8C**). R74A, K180A, and R328A/R329A have similar basal rates to WT, however, R164A had a 2.5-fold enhancement over WT. Similar to WT *Ec*DnaB, both R74A and R164A were stimulated 1.5-2.0-fold in the presence of DNA consistent with previous results (25). Interestingly, the fastest unwinding mutants, K180A and R328A/R329A, were only weakly stimulated further in the presence of DNA. Previously, the R328A/R329A mutant was investigated for its potential role in a leucine zipper motif (26) (although later discounted) (27) and was also found not to have DNA stimulated ATPase activity. No stimulation in ATPase rate with DNA is sometimes indicative of a perturbation in DNA binding, however, these mutants show stimulated unwinding abilities and fluorescence anisotropy experiments showed no significant differences in K_d values measured for mutants binding to fork DNA compared with WT (data not shown).

To validate whether alteration of electrostatic interactions are responsible for the increased unwinding rates in the mutants compared to WT *Ec*DnaB, we performed DNA unwinding reactions with a 3'-morpholino (morph) strand. Morpholino nucleic acids have standard base pairing properties but instead have morpholine rings linked through phosphorodiamidate groups that lack negative charge and are as stable or more stable than an equivalent DNA duplex (28,29). Previously, the homologous hexameric T7 gp4 DNA helicase was shown to unwind excluded strand morpholino substrates with a greater rate and efficiency than for DNA (16). They attributed this enhanced unwinding activity to the disruption of the helicase's interaction with the displaced strand that limits its activity.

Interestingly, WT *Ec*DnaB also unwinds excluded strand morpholino substrates with a profoundly enhanced rate compared to a DNA/DNA duplex (**Fig. 9**). The rate of unwinding for the Morph/DNA is at least 1000-fold faster with a > 0.7 amplitude after 1 minute, while the DNA/DNA duplex is only ~0.4 unwound after 45 minutes. No unwinding or strand separation is seen when ATP is excluded from the experiment. Accurate quantification of the unwinding rate would require rapid quench experiments, but the point that *Ec*DnaB unwinds excluded strand morpholinos rapidly is apparent. The unwinding rates for the Morph-DNA are also significantly faster (~10-fold) than even those seen above for the fastest SEW mutant (K180A) on a DNA fork (**Fig. 8**). This may not be surprising as the SEW mutants only affects contact at one specific mutated site on the exterior surface, while the excluded strand morph eliminates electrostatic contacts throughout the longitudinal length of *Ec*DnaB.

DISCUSSION

Although hexameric DNA replication helicases have global structural conservation, their amino acid sequences are not conserved, allowing for the classification of these helicases into

different superfamilies. We have shown previously that the 5' excluded strand makes important external surface interactions that aid in the mechanism of unwinding for the SF6 archaeal *Sso*MCM helicase (3'-5') to develop the SEW of unwinding (11). In this report, we can now show that the bacterial replication helicase, *Ec*DnaB, with opposite unwinding polarity (5'-3') and of a different family (SF4) and organismal domain, has similar conformational states and dynamics of binding the excluded strand that also regulate DNA unwinding (**Fig. 10**). The combined results highlight the importance and conservation of the SEW model for hexameric helicase unwinding of DNA and reveal external surface residues required for regulating the activity of the *Ec*DnaB helicase. Importantly, the SEW effects on the mechanism of unwinding are opposing for *Sso*MCM and *Ec*DnaB.

It is striking that the absolute FRET states, transition probabilities, and dwell times visualized by the ExPRT plots are extremely similar between *Sso*MCM and *Ec*DnaB hexameric helicases bound to fork DNA. In both cases, the large increase in FRET observed is consistent with encircling of the translocating strand and exclusion of the other along the exterior surface. For *Ec*DnaB, there have been reports of hexamers loading on opposing strands in opposite orientations (3,30-32). The consequence of loading two hexamers would ultimately separate the strands further, resulting in a decrease in the FRET signal, which is opposite to what we observe in the smFRET experiments even at high concentrations of *Ec*DnaB (**Fig. 1B**). Both helicases are of similar size, oligomeric state, and are thought to engage their respective translocating strands in a similar way. So, the DNA bound states of each helicase may be structurally equivalent even with opposing translocation polarities. Therefore, *Ec*DnaB binding of DNA includes both the encircling of the 5'-strand and the exclusion and external interaction of the 3'-strand in a similar manner to *Sso*MCM and the SEW model for unwinding.

Without an appropriate DNA bound crystal structure of *Ec*DnaB, we had to infer binding positions for the excluded 3'-strand based on amino acid homology and electrostatics from crystal structures that represented a closed ring (6,7) or a split lock washer structure (9). Because ssDNA was contained in the central channel of the split lock washer structure, we used this conformation as a primary model to interpret interactions with the excluded strand. This restricted the definition of any precise or specific exterior binding path, and rather we can only conclude general binding to the CTD, the waist, and the NTD. That along with the specific residues that were mutated and the smFRET data informed our interpretation of excluded strand binding.

To test the specificity of this external interaction, mutations of conserved residues were found to both disrupt and alter the binding states. In particular, R74 was found to be necessary for stable interactions of the excluded strand at the NTD giving rise to the highest FRET state (~0.95). R164A and K180A exhibited somewhat different dynamics than WT, generally fitting to more FRET states, which may reflect alternate binding paths on the helicase exterior. It is likely that these two residues partially contribute to the excluded strand interaction. R328 and R329 both seem to be important for wrapping longer ssDNA substrates at the waist; where upon mutation, the values shift from a medium (~0.8) to a lower (~0.5) FRET state consistent with decreased wrapping. Altogether, these mutants individually alter the interaction between the excluded strand and the exterior of the helicase to varying degrees, and the amount of destabilization or altered external DNA binding paths can depend on excluded strand length. The data provide information on the contacts all along the longitudinal length of the hexamer defining a minimal binding path.

Previously, mutation of external positively charged residues on *Sso*MCM reduced DNA unwinding, presumably through a slippage mechanism where the mutant helicase was unable to stabilize forward unwinding steps (11). However, for *Ec*DnaB, mutation of external positively charged residues generally increased unwinding rates. The enzymatic effects of the mutations largely correlate with two classes of results. Two of the mutations (R74A and R164A) show slightly enhanced unwinding activity and their ATPase rates are stimulated with DNA similarly to WT. For R164A, the increased ATPase activity of this mutant could account for the DNA unwinding enhancement, but R74A has similar ATPase rates to WT. In both cases the mutation to alanine has disrupted or altered the external interaction as measured by smFRET. We

propose that releasing some of the electrostatic wrapping interaction frees the helicase to unwind faster.

The more intriguing class of mutations (K180A and R328A/R329A) exhibit more than 20-fold increases in DNA unwinding. Interestingly, they do not show as significant DNA-dependent stimulation of ATP hydrolysis rates. For these mutants, the most significant differences are an increase in the number of conformations for K180A with shorter fork arms and a global change in FRET states and an increase in dynamics for R328A/R329A with the longer excluded strand. Overall, a general trend of increased unwinding activity emerges as we neutralize positively charged residues found on the exterior of the hexamer. Therefore, exterior electrostatic interactions with the excluded strand restrict the unwinding activity of *Ec*DnaB.

By using excluded strand morpholino substrates instead, we have the added benefit of testing the total effect of exterior electrostatics on unwinding, instead of contributions at specific amino acid sties. Disruption of the electrostatic interaction of the excluded strand through this morpholino chemistry was strongly stimulatory to unwinding. A similar stimulation in unwinding of morpholino strands were also seen with the homologous T7 gp4 hexameric helicase (16). In both cases, interactions on the outer surface of the helicase with the excluded strand will act to regulate the unwinding rate. However, this external interaction is not entirely electrostatic as increasing ionic strength in the smFRET experiments resulted in decreased dwell times but did not significantly affect the FRET states.

Previous single-molecule work has detailed the single molecule force contributions of each DNA strand to unwinding by *Ec*DnaB using either a hairpin or fork substrate (33). In that study, it was concluded that the unwinding rate was controlled by both force-induced destabilization of the duplex as well as interactions of the excluded strand with the exterior surface. The main apparent discrepancy between our work and theirs is that when the excluded strand is sequestered because of constraints in the hairpin assay, the rate is slower than when it is allowed to interact with the exterior surface in the fork assay. This would imply that contacts of the excluded strand with the external surface of *Ec*DnaB increase unwinding; whereas we show that specific external contacts restrain unwinding. However, it is probable that the force applied to the excluded strand in the fork assay artificially alters the interaction with the exterior surface in a way analogous to the altered DNA binding paths and kinetics for the R74A and R164A mutants. Therefore, measured increases in unwinding in both studies can be explained by altered DNA binding paths on the exterior surface. It is notable that a variety of recent biophysical techniques monitoring *Ec*DnaB activity and binding have detected an elusive external interaction with the excluded strand (18,33). We can now conclude that this SEW interaction and the precise binding path regulates the speed of unwinding for *Ec*DnaB.

Clearly, interactions with the excluded strand are acting as a regulator to control the speed of unwinding. It remains to be seen whether this is because of a greater increased force applied by the motor domain for *Ec*DnaB that is modulated by the excluded strand or whether discrete external binding paths or polarity dictate the rate of unwinding. Further experiments will be needed to more specifically define the exterior binding path. For *Ec*DnaB, the excluded strand interaction may act as a 'molecular brake' to control the amount of exposed ssDNA or provide a platform for accessory helicases, *i.e.* Rep, to assemble and rescue stalled forks (34,35). Coupled DNA synthesis by the leading strand polymerase (Pol III) could sequester the excluded strand from the exterior surface of *Ec*DnaB and explain the increased rate of unwinding by the coupled replisome (36-38). In fact, a recent report has shown that when the bacterial helicase and polymerase become decoupled, the unwinding rate is reduced by 80% as a fail -safe 'dead man's switch' (39). This can be explained at the molecular level by our data in which decoupling of the polymerase engages the excluded strand with the exterior surface of *Ec*DnaB slowing its unwinding progression. In addition to controlling the unwinding rate, the external DNA binding sites on both helicases are likely to contribute during the loading mechanism for encircling of the translocating strand to maintain strand separation during the action and conformational changes induced by the initiation enzymes.

EXPERIMENTAL PROCEEDURES

Materials **-** Oligonucleotides used (**Supplemental Data Table S1**) were purchased from IDT (Coralville, IA). Fluorescently labeled DNA was HPLC purified (IDT), and non-labeled oligos were gel purified (40). Morpholinos were from GeneTools (Philomath, OR). *Sso*MCM was purified as previously described and reported as hexamer concentrations (11). All other chemicals were analytical grade or better.

Cloning and Protein Purification of EcDnaB **-** The R74A, R164A, K180A, and R328A/R329A mutations of *Ec*DnaB were created by overlap extension from pET11b-*Ec*DnaB. The DNA primers are listed in **Supplemental Table S1**. Mutations were confirmed by the DNA sequencing facility at the University of Pittsburgh. Mutants and WT *Ec*DnaB were expressed in Rosetta 2 cells (EMD Millipore, Billerica, MA) or C41 cells (Lucigen, Middleton, WI) using autoinduction (41) or by induction with 0.1 mM IPTG. Cells were pelleted, resuspended in *Ec*DnaB lysis buffer [10 % sucrose, 50 mM Tris-HCl (pH 7.5), 50 mM NaCl, 5 mM dithiothreitol (DTT)], and lysed using lysozyme and sonication. Ammonium sulfate was added to the resulting supernatant at 0.2 g/mL, pelleted, and then resuspended in *Ec*DnaB buffer A [10 % glycerol, 0.1 mM EDTA, 50 mM Tris-HCl (pH 7.5), 50 mM NaCl, 5 mM DTT]. The supernatant was purified using an AKTA Prime FPLC equipped with a HiTrap MonoQ column (GE Healthsciences, Sunnyvale, CA) and eluted with a stepwise gradient of *Ec*DnaB buffer A with 500 mM NaCl followed by a similar procedure using a HiTrap Heparin column (GE Healthsciences, Sunnyvale, CA). The purified fractions were combined and applied to a Superdex S-200 26/60 gel filtration column (GE Healthsciences, Sunnyvale, CA) with Buffer C [50 mM Tris-HCl (pH 7.5), 50 mM NaCl, 5 mM DTT] to isolate the hexamer. An extinction coefficient (185,000 cm^{-1} M^{-1}) was used to quantify the fractions containing purified hexameric *Ec*DnaB (42). All concentrations for *Ec*DnaB are indicated as hexamer throughout.

Single-molecule Fluorescence Resonance Energy Transfer - DNA substrates labeled with Cy3 and Cy5 fluorophores were immobilized on a PEGylated quartz slide utilizing biotin-streptavidin interactions (43). A prism-based total internal reflection microscope was used to collect all smFRET data (44,45). A 532 nm diode laser was used to excite Cy3 fluorophore, and subsequent Cy3 and Cy5 emission signals were separated by a 610 nm dichroic longpass mirror, a 580/40 nm bandpass filter, and a 660 nm longpass filter. An EM-CCD iXon camera (Andor, Belfast, UK) was used to image the signals. Data was acquired at 10 fps for ten or more regions with each region containing 50 – 250 molecules in the presence of an oxygen radical scavenging solution [1 mg/mL glucose oxidase, 0.4 % (w/v) D-glucose, 0.04 mg/mL catalase] and 2 mM trolox. *Ec*DnaB (250 nM) was added and given a five-minute equilibration period. All single-molecule experiments were performed in reaction buffer as described previously (11).

Single-molecule FRET signals were identified by fitting individual regions of signal intensity to a 2D Gaussian and measuring the goodness of fit. These peaks were corrected for thermal drift and local background intensity (46,47). The resulting signal was used to calculate the apparent FRET efficiency, E_{app}, according to

$$E_{app} = \frac{I_A}{I_A + I_D} \quad (1)$$

in which I_A and I_D are the intensity of the acceptor and donor signals respectively.

Single-molecule FRET data analysis and ExPRT Plots **-** Data analysis and visualization were performed using manually selected single-molecule traces that displayed anti-correlation between the donor and acceptor fluorophores and single-step fluorophore photobleaching. Traces collected under identical experimental conditions were stitched together, and fit to ideal states via Hidden Markov Modeling using the vbFRET software package (48). Stitched traces were fit to a given number of states based on those states being more than $E_{app} = 0.1$ apart from one another, and the variation of one state not overlapping with another. Traces were then unstitched and fed into the Explicit Probability and Rate Transition (ExPRT) analysis program. This MATLAB executable program produces transition plots where the markers are sized based on the probability of transition occurring within an observed single-molecule trace and colored based on the dwell time(s) of the state preceding the transition. The average of all dwell times for a

given transition was used to determine the color of the marker. Only dwell times that were both preceded and followed by transitions were included. Stitched data as fit by vbFRET were also analyzed by the POKIT analysis program, and the resulting plot contains a legend for the ranges of rates and probabilities (24). The data were also fit using the HaMMY program, allowing the program to fit the data to up to five states. The output of the HaMMY program was subsequently analyzed by the Transition Density Plot program (23).

EcDnaB Structural Homology Model - Global sequence alignments were performed using ClustalW2 analysis (http://www.ncbi.nlm.nih.gov/blast/bl2seq/wblast2.cgi). The homology model of *Ec*DnaB was created by threading the alignment on to the structure of *Geobacillus stearothermophilus* DnaB (PDBID: 4ESV) (9) using Swiss-MODEL (49).

ATPase Assay - *Ec*DnaB variants (350 nM) were incubated in the absence and presence of 4 μM forked DNA (DNA14/DNA15) as described previously (50). Briefly, 25 μL reactions were incubated at 37 °C for 5 minutes in unwinding buffer [50 mM HEPES (pH 8.0), 10 mM $Mg(OAc)_2$, 5 mM DTT, 0.2 mg/mL BSA], and 1 mM ATP with trace amounts of ^{32}P-γ-ATP was added to initiate the reaction. Samples were quenched at 2, 5, 10, and 15 min after initiation in equal volumes of 0.7 M formic acid. A total of 1 μL of quenched reaction was spotted on Millipore TLC PEI Cellulose F, allowed to dry, resolved in 0.6 M potassium phosphate (pH 3.5), phosphorimaged, and quantified for the linear ATPase rate (pmol/min).

Gel Based DNA Unwinding Assays - Helicase assays were assembled in unwinding buffer at 37 °C. 15 nM of fluorescent forked DNA (DNA14/F/DNA15) was incubated with 500 nM *Ec*DnaB at 37 ºC for 5 minutes before initiating with 5 mM ATP. Reactions were quenched with using an equal volume of quench solution [50% glycerol, 1% SDS, 100 mM EDTA (pH 8.0), and 300 nM ssDNA trap (unlabeled strand with same sequence as radiolabeled strand)] at various time points from 1-15 minutes. Reactions were kept on ice until loading and resolved on 20% (29:1 acryl:bisacryl) and TBE gels electrophoresed in TBE buffer. The gels were imaged on a Typhoon 9400 phosphorimager (GE Healthsciences, Sunnyvale, CA), and the fraction unwound was calculated after background subtraction using ImageQuant software according to:

$$F = \left(\frac{I_{s(t)}}{I_{s(t)}+I_{D(t)}} - \frac{I_{s(0)}}{I_{s(0)}+I_{D(0)}}\right) / \left(\frac{I_{s(b)}}{I_{s(b)}+I_{D(b)}} - \frac{I_{s(0)}}{I_{s(0)}+I_{D(0)}}\right) \quad (2)$$

where $I_{s(t)}$ and $I_{D(t)}$ are the intensities of the single and double-strand bands, respectively, at time *t*; subscript *0* and *b* indicate equivalent counts at t = 0 and the boiled sample, respectively. The fraction unwound or unwinding rate (nM min^{-1}) was calculated from a linear regression fit of the fraction unwound for each time point.

Single-Turnover DNA Unwinding Assays: Single-turnover unwinding experiments were also performed by initiating the reaction simultaneously with 5 mM ATP and 150 nM of a ssDNA trap (unlabeled strand with the same sequence as the radiolabeled strand). Experiments were performed with 250 nM *Ec*DnaB and 15 nM forked 3'-DNA (DNA163/DNA161) or 3'-morpholino (DNA163/DNA160m) substrates at 37 ºC. The 5'-end of DNA163 was labeled with ^{32}P-γ-ATP (Perkin Elmer, Waltham, MA) with Optikinase (Affymetrix, Cleveland, OH) according to manufacturer directions. Reaction was quenched at various times using an equal volume of quench solution [20% Ficoll, 1.0 %SDS, 200 mM EDTA (pH 8.0), 2 mg/mL Proteinase K] followed by incubation at 37 ºC for 10 min. After electrophoresis as above, the gels were imaged on a Storm 820 (GE Healthsciences, Sunnyvale, CA), and the fraction unwound was calculated using ImageQuant software according to Equation 2. Single turnover data were further quantified using:

$$F = C + A_1\left(1 - e^{-k_1 t}\right) + k_{ss}t \quad (3)$$

where A_1 is the amplitude associated with the initial burst rate increase (k_1), k_{ss} is a steady state rate, and *C* is a constant.

ACKNOWLEDGEMENTS

pET11b-*Ec*DnaB was a generous gift from Dan Kaplan. We thank the Walter lab for access to their POKIT analysis program. We thank Brian Graham, Heather McFarland, Grant Schauer, and Valerie O'Shea for helpful discussions. This work was supported by Baylor University, The University of Pittsburgh, a Research Scholar Grant (RSG-11-049-01-DMC to M.A.T.) from the American Cancer Society, NSF-MCB (NSF1613534 to M.A.T.), and the NIH (AI081571 and GM068406 to S.H.L.).

CONFLICT OF INTEREST

The authors declare that they have no conflicts of interest with the contents of this article.

AUTHOR CONTRIBUTIONS

S.M.C., S.H.L., and M.A.T. designed the experiments. S.M.C. performed the cloning, protein purification, biochemical assays, and single-molecule experiments. S.M.C. and S.G. performed DNA unwinding assays and protein purifications. M.A.T. directed the project, and together with S.M.C. wrote the manuscript. All authors edited the manuscript.

FOOTNOTES

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ABBREVIATIONS

AAA^+, ATPase associated with diverse cellular activities; ATP, adenosine 5'-triphosphate; BME, β-mercaptoethanol; bp, basepair; BSA, bovine serum albumin; Cdc45, cell division cycle 45 protein; CTD, C-terminal domain; Cy3, cyanine 3 dye; Cy5, cyanine 5 dye; dsDNA, double-stranded DNA; dT, deoxythymidine; E1, Papillomavirus E1; *Ec*DnaB, *Escherichia coli* DnaB; *Ec*DnaC, *Escherichia coli* DnaC; EDTA, ethylenediaminetetraacetic acid; EM, electron microscopy; EM-CCD, electron multiplying charge-coupled device; ExPRT plots, Explicit Rate and Probability Transition plots; FPLC, fast protein liquid chromatography; FRET, fluorescence resonance energy transfer; GINS, Go Ichi Nii San (5,1,2,3 in Japanese) complex; HEPES, 2-[4-(2-hydroxyethyl)piperazin-1-yl]ethanesulfonic acid; HPLC, high pressure liquid chromatography; IPTG, isopropyl β-D-1-thiogalactopyranoside; MCM, minichromosome maintenance protein; NTD, N-terminal domain; SE, steric exclusion; SEW, steric exclusion and wrapping; SF4, super family 4; SF6, super family 6; smFRET, single-molecule FRET; ssDNA, single-stranded DNA; *Sso*MCM, *Sulfolobus solfataricus* MCM; SV40 LargeT, Simian Vacuolating Virus 40 TAg; T4, T4 bacteriophage; T7, T7 bacteriophage; TBE, tris-borate-EDTA; TEV, tobacco etch virus protease; Tris, Tris(hydroxymethyl)aminomethane; WT, wild-type

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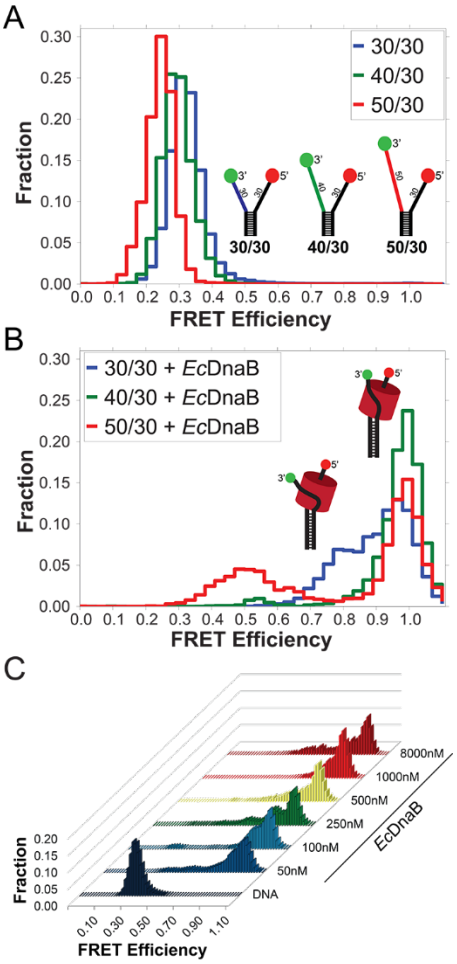


FIGURE 1. **Single-molecule FRET monitoring of *Ec*DnaB binding to DNA fork substrates.** *A*, histograms of the smFRET signal from the DNA fork substrates alone, colored to match cartoon models of the DNA forks with a static 30 base encircled 5' strand and variable excluded-strand 3' arm lengths (30, 40, and 50 nt) shown in blue, green and red, respectively. *B*, histograms of the three DNA substrates in the presence of 250 nM WT *Ec*DnaB. *C*, the histogram profiles from a titration of WT *Ec*DnaB onto the 30/30 fork substrate from 50 nM to 8 μM is shown. 30/30 alone exhibits low FRET (shown in dark blue). Adding WT *Ec*DnaB shifts the FRET signal to higher FRET values in all cases, without significantly altering the histogram profile.

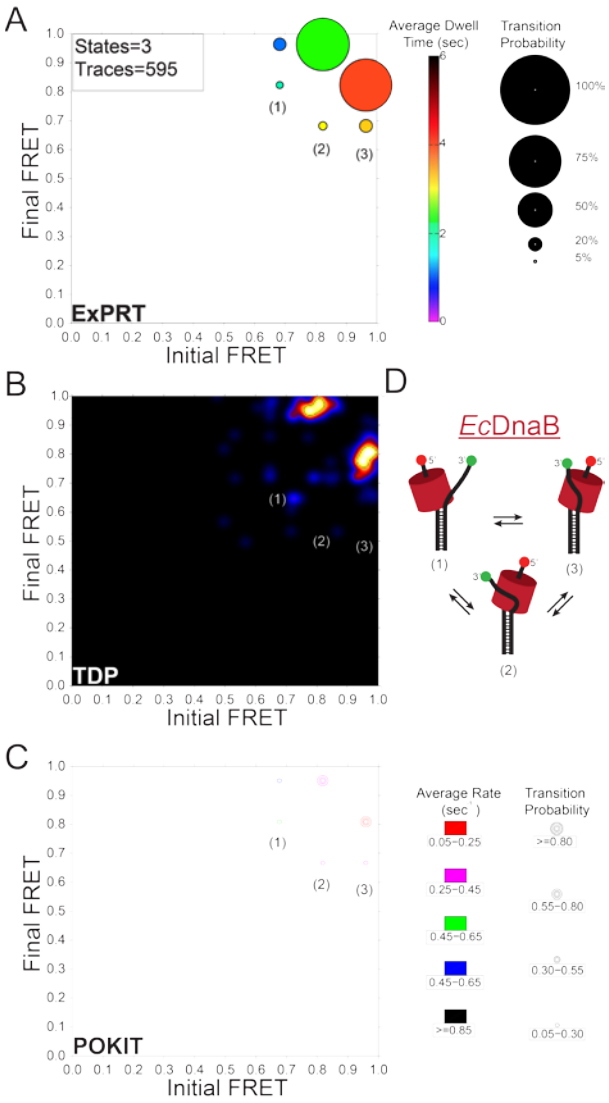


FIGURE 2. **Comparison of smFRET dynamics for excluded strand interactions on *Ec*DnaB.** *A*, ExPRT plot showing the probability (size) and dwell time (color) of transitions for *Ec*DnaB (250 nM) on 30/30 fork DNA. The number of states and traces fit by the data is in the upper left-hand corner for each plot. *B,* The smFRET data set from WT *Ec*DnaB on 30/30 was also analyzed and fit using HaMMY, and subsequently analyzed and visualized by the Transition Density Plot (TDP). *C,* Separately, the same data was stitched together and fit using vbFRET and visualized using the POKIT analysis program. *D,* Cartoon representation of the hypothesized three states (1, 2, 3) of bound helicase to DNA fork identified in the ExPRT plots and indicated on each plot.

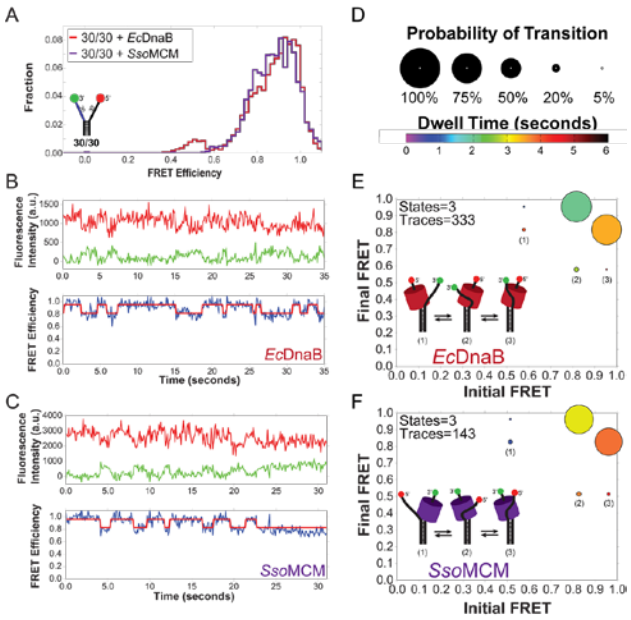
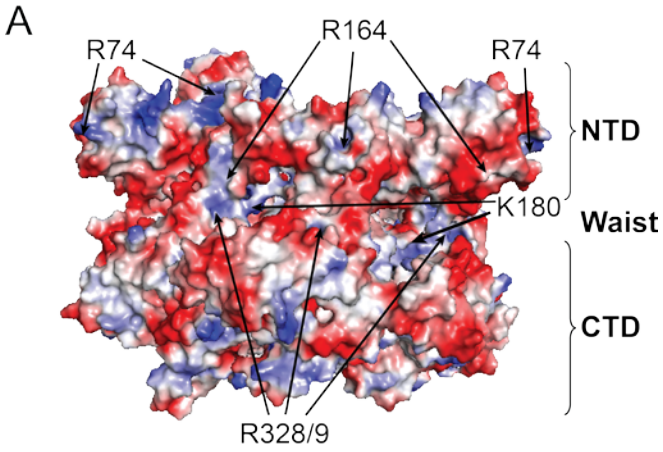


FIGURE 3. **Comparison of the excluded strand interactions of *Sso*MCM and *Ec*DnaB by smFRET.** *A*, shows the overlaid histograms of both *Sso*MCM and *Ec*DnaB on the 30/30 fork. Representative single-molecule traces for *B*, *Ec*DnaB and *C*, *Sso*MCM on the 30/30 DNA template. The top panels show the Cy3 (green) and Cy5 (red) signals. The bottom panels show the corresponding FRET signal (blue) with overlaid ideal states (red) for each trace as fit by vbFRET (see Materials and Methods). *D*, ExPRT plots showing the probability (size) and dwell time (color) of transitions for *E*, *Ec*DnaB (250 nM) and *F*, *Sso*MCM (1.3 μM), respectively, on 30/30 fork DNA. The number of states and traces fit by the data is in the upper left-hand corner for each plot. Inset is a carton representation of the hypothesized three states of bound helicase to DNA fork identified in the ExPRT plots.



B

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          R74      R164      K180     RR328/9
ECOLI  MARLQE  ESRVFKI  GPKNIA  RARRIA
GEOSE  MLRVAD  DRKIMEV  AFKNIK  KCRRL-
ECOBD  MARLQE  ESRVFKI  GPKNIA  RARRIA
BARGA  ISKIIQ  ENQLFEL  GFENFN  RARRLK
SALTY  MGRLQE  ESRVFQI  FPKSID  RARRIF
HELPY  ALKLHE  EREVYAL  GFRNIK  QLRKLK
ECO57  MARLQE  ESRVFKI  GPKNIA  RARRIA
SHIFL  MARLQE  ESRVFKI  GPKNIA  RARRIA
TREPS  LVELSD  QREIYDL  TFKLLK  VARRLC
          .: : : .:. :    .  :     *::
```

FIGURE 4. **Identification of Exterior Electrostatic SEW Sites for *Ec*DnaB.** *A*, Position of the SEW mutations (from multiple subunits) mapped onto the homology model for *Ec*DnaB colored with an electrostatic surface identifying the N-terminal domain (NTD) C-terminal RecA domain (CTD) and the waist. *B*, multiple amino acid alignment of DnaB helicases using CLUSTAL W2 (http://www.ebi.ac.uk/Tools/clustalw2). Identical (*), similar (:), and somewhat similar (.) residues are indicated. ECOLI - *Escherichia coli* strain (K-12); GEOSE - *Geobacillus stearothermophilus*; ECOBD - *Escherichia coli* strain (BL21-DE3); BARGA - *Bartonella grahamii* (strain as4aup) SALTY - *Salmonella typhimurium*; HELPY - *Helicobacter pylori* strain (26695); ECO57 - *Escherichia coli* O157:H7; SHIFL - *Shigella flexneri*; TREPS - *Treponema pallidum* SS14.

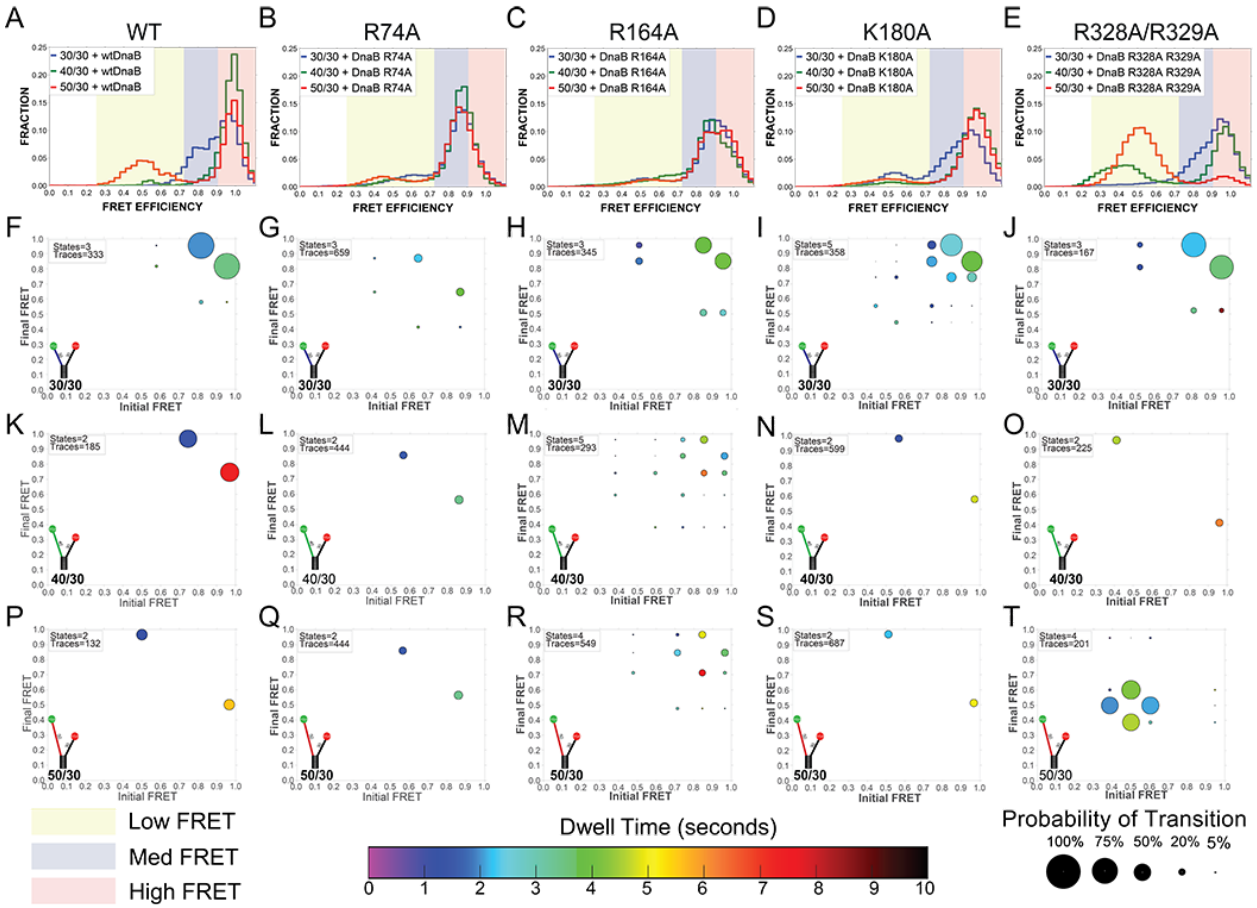

FIGURE 5. **Histograms and ExPRT Plots of WT *Ec*DnaB and mutants bound to DNA forks.** Histograms (*A-E*) report the population of molecules as a function of FRET states on DNA forks with a 30-base 5'-strand and a 30-base (blue), 40-base (green), or 50-base (red) 3'-strand for WT, R74A, R164A, K180A, and R328A/R329A, respectively. Yellow, blue, and red regions highlight low, medium, and high FRET populations, respectively. Corresponding ExPRT plots are shown for (*F-J*) 30/30, (*K-O*) 40/30, and (*P-T*) 50/30 forks for each of the respective *Ec*DnaB helicases.

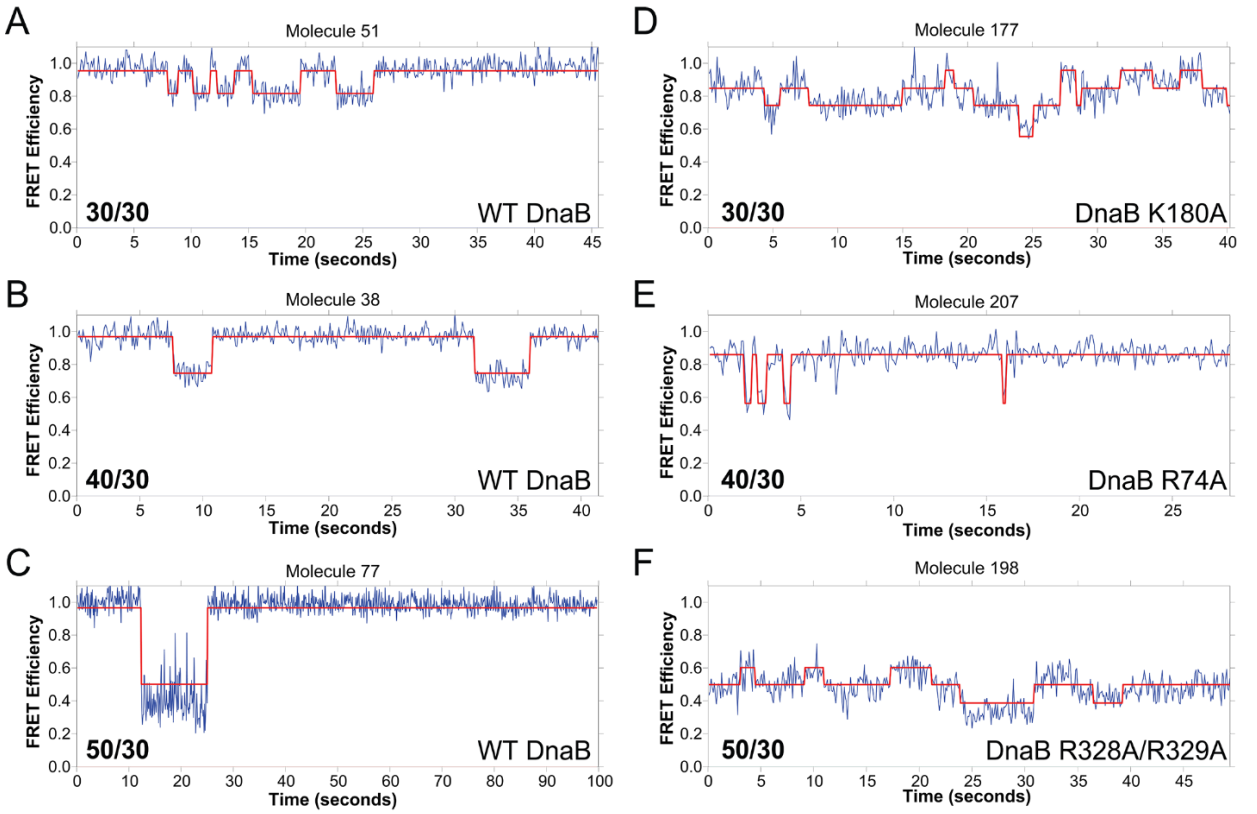

FIGURE 6. **Example smFRET kinetic traces.** Comparison of *A-C,* WT and *D,* K180A, *E,* R74A, and *F,* R328A/R329A *Ec*DnaB FRET efficiencies as a function of time on the 30/30, 40/30 and 50/30 forks, respectively. The calculated FRET values (blue) are overlaid with the ideal state fits (red).

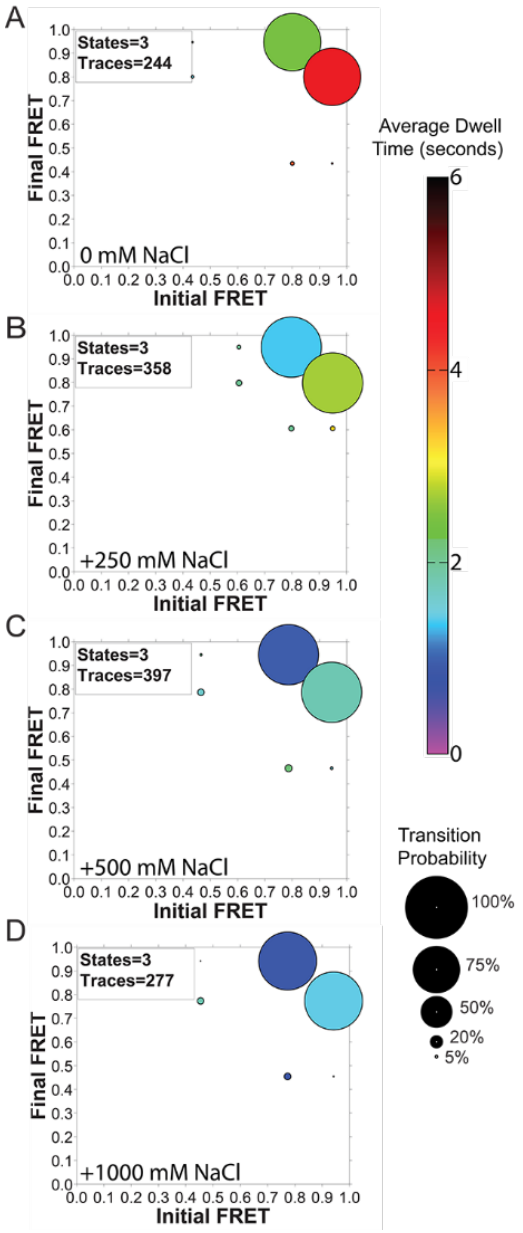

FIGURE 7. **Titration of NaCl onto *Ec*DnaB-bound 30/30 Fork.** *A-D,* ExPRT plots of [NaCl] titration onto *Ec*DnaB (250 nM) prebound to 30/30 DNA fork.

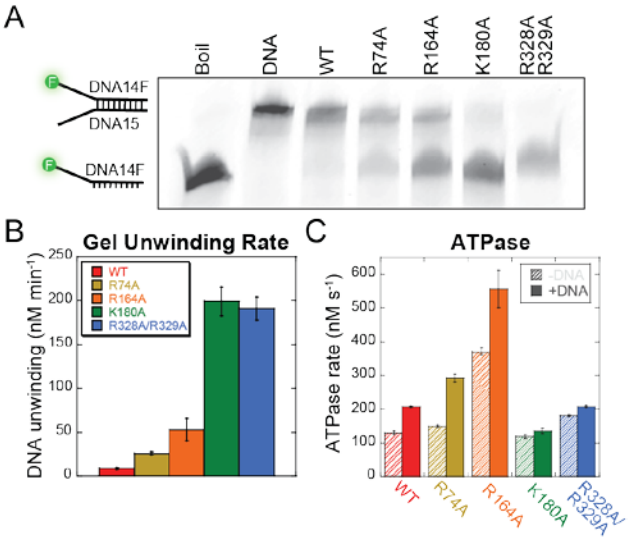

FIGURE 8. **Biochemical Properties of SEW Mutants of *Ec*DnaB.** *A*, Representative gel for six-minute time point is shown for *Ec*DnaB (WT and mutants, 3 □M) unwinding assays performed on a fluorescein labeled fork DNA (DNA14F/DNA15) (15 nM) and *B*, quantified over multiple time points. Throughout, data for *Ec*DnaB constructs are consistently colored (WT - red; R74A – ochre; R164A – orange; K180A – green; R328A/R329A – blue). Error bars represent the standard error from at least three independent experiments. *C*, Quantification of the ATP hydrolysis rate in the absence (diagonal hash) and presence (solid) of fork DNA (DNA14/DNA15).

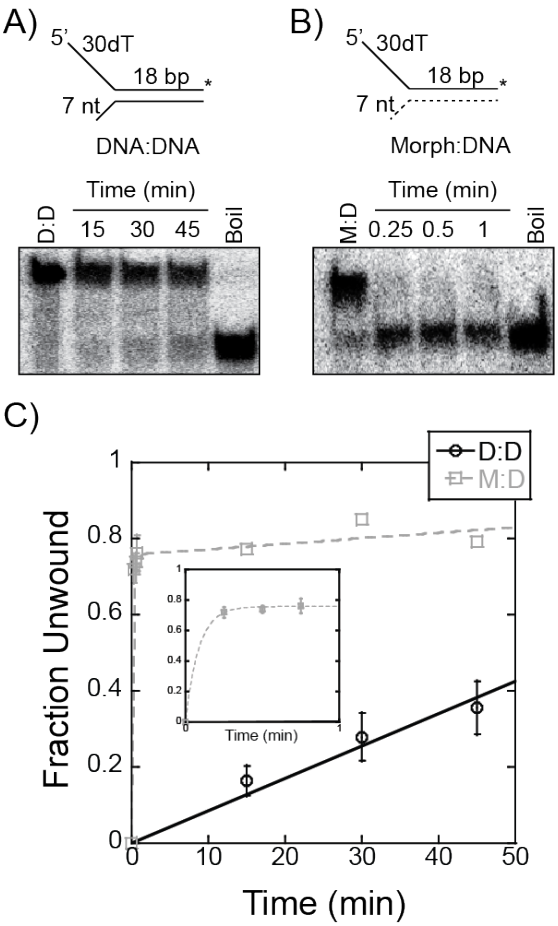

FIGURE 9. **Unwinding of Morpholino Forked Substrates.** A representative unwinding time course for 250 nM *Ec*DnaB on either an 18 bp A) DNA/DNA (D:D) or B) Morph/DNA (M:D) substrate (15 nM) with a 7 base 3'-excluded strand flap. Single turnover experiments were initiated with ATP and a single-strand trap identical to the radiolabeled strand as described in Materials and Methods. C) Averaged unwinding data for D:D (black ○) was plotted and fit with a linear regression to give $k = 0.009$ min^{-1} and for (M:D, grey □) was plotted and fit using **Equation 3** to give $k_1 = 11$ min^{-1}. The inset plot highlights data within the first minute. Error bars are the standard error from three independent experiments.

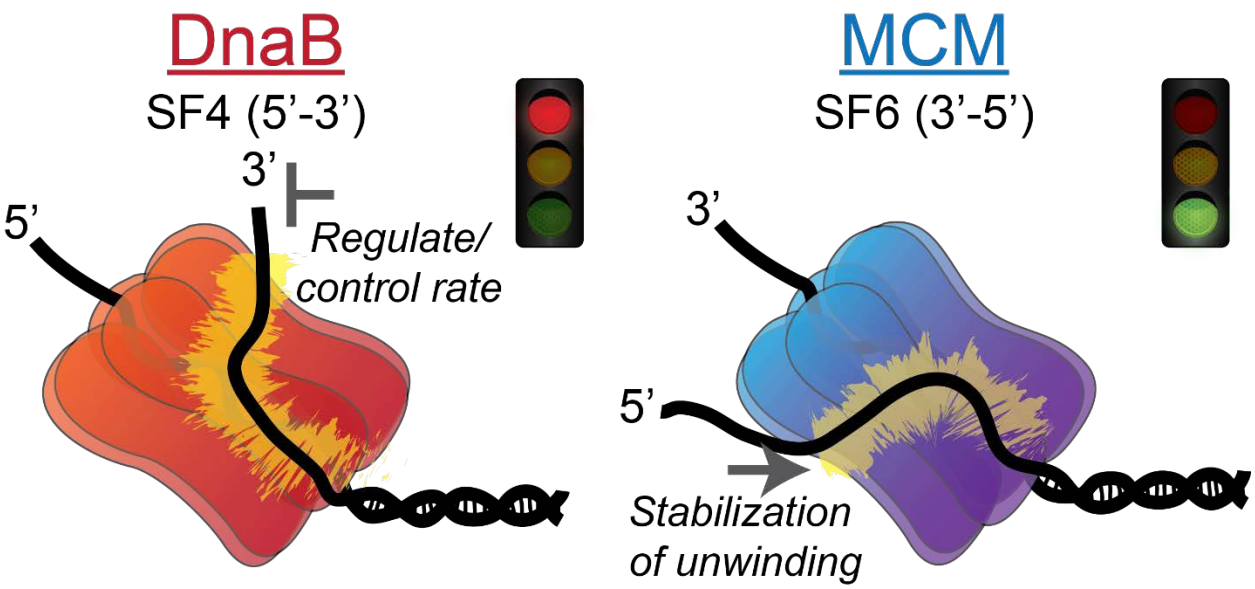

FIGURE 10. **SEW models for hexameric helicase unwinding.** Bacterial DnaB and archaeal MCM encircle the lagging or leading strand, respectively, and interact with the excluded strand on the exterior surface to either regulate the unwinding rate using this electrostatic brake or stabilize unwinding in a forward direction.