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Structure and primase-mediated activation of a bacterial dodecameric replicative helicase

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ABSTRACT

Replicative helicases are essential ATPases that unwind DNA to initiate chromosomal replication. While bacterial replicative DnaB helicases are hexameric, *Helicobacter pylori* DnaB (HpDnaB) was found to form double hexamers, similar to some archaeal and eukaryotic replicative helicases. Here we present a structural and functional analysis of HpDnaB protein during primosome formation. The crystal structure of the HpDnaB at 6.7 Å resolution reveals a dodecameric organization consisting of two hexamers assembled via their N-terminal rings in a stack-twisted mode. Using fluorescence anisotropy we show that HpDnaB dodecamer interacts with single-stranded DNA in the presence of ATP but has a low DNA unwinding activity. Multi-angle light scattering and small angle X-ray scattering demonstrate that interaction with the DnaG primase helicase-binding domain dissociates the helicase dodecamer into single ringed primosomes. Functional assays on the proteins and associated complexes indicate that these single ringed primosomes are the most active form of the helicase for ATP hydrolysis, DNA binding and unwinding. These findings shed light onto an activation mechanism of HpDnaB by the primase that might be relevant in other bacteria and possibly other organisms exploiting dodecameric helicases for DNA replication.

INTRODUCTION

DNA replication is an essential process in all organisms and consists of the faithful duplication of the genetic material to be transmitted to daughter cells. Initiation of DNA replication relies on helicases, key motor proteins that unwind DNA so that it can be used as a template for DNA polymerases (1). Replicative helicases are hexameric ring-shaped proteins that hydrolyse ATP and unwind double-stranded DNA (dsDNA), with one strand running through the central cavity of the hexamer and the other being excluded from the ring. In bacteria, the replicative helicase DnaB functions as a hexamer to unwind dsDNA in the 5’ to 3’ direction (1).

The DnaB monomer consists of an N-terminal domain (NTD, comprising a head and a helical hairpin) and a C-terminal RecA type ATPase domain (CTD) associated through a linker (Figure 1A) (2). The NTDs and CTDs assemble into separate rings (namely NTD- and CTD-rings) that have different symmetries and form a two-tiered hexamer (3,4). In the absence of a ligand, the NTD ring adopts a conformation compatible with interaction with the *H*. *pylori* subunit of the polymerase (5).

Because DnaB forms closed rings, threading of single-stranded DNA (ssDNA) through the central channel of DnaB requires ring-breaking or ring-making mechanisms during initiation of replication (6). In addition, because of their polarity, two hexamers of DnaB must be positioned in opposite directions at the origin on each of the DNA forks for replication to proceed bi-directionally (7). In some bacteria, loading of the helicase is assisted by AAA+ helicase loaders such as DnaC in *Escherichia coli* or DnaI in *Bacillus subtilis* (*Bst*DnaI) (7). E. *coli* DnaC forms a 6:6 complex with EcDnaB (8,9). The interaction with the loader opens the EcDnaB rings into a lock-washer conformation and triggers a helical conformation of the overall complex, thereby providing an entry point for ss-
DNA (10). A non-planar architecture has also been observed in the pre-priming complex composed of BstDnaI and Bacillus steaerothermophilus helicase (BstDnaB), as well as the helicase-binding domain (HBD) of the primase (BstDnaGHBD) (11), suggesting that helicase loading might depend on a transition of the helicase from planar to non-planar conformation induced or stabilized by the AAA+ helicase loader (10,11).

During replication, the active form of DnaB is a complex of the helicasewith up to three DnaG primases, named the primosome. Transition from a loaded helicase-helicase-primasecomplex to an active unwinding primosome requires additional conformational changes of the helicase. Closing of the helicase rings is likely facilitated by nucleotide and ssDNA binding since the translocation conformation of DnaB on ssDNA is also spiral (12). BstDnaGHBD interaction with the pre-primed BstDnaB-BstDnaI complex resulted in a ternary complex that might be loaded onto the replication fork (11). In E. coli, primer synthesis by DnaG in complex with DnaB releases DnaC (13) and stimulates DnaB helicase and ATPase activities (14). Therefore, bacteria appear to have developed different strategies for the formation of primosomes. Moreover, AAA+ loader-dependent primosomes are unlikely to represent a universal strategy since loader homologues have been identified in only a subset of bacterial species (~20%) (15). Concomitantly, the human pathogen Helicobacter pylori does not encode for a DnaC/I homologue. Moreover, H. pylori DnaB (HpDnaB) could complement two temperaturesensitive mutants of E. coli dnaBts and dnaCts, suggesting that HpDnaB was able to bypass DnaC in these cells (16). HpDnaB formed a double hexamer (17) with an architecture remarkably similar to head-to-head double hexamers of some replicative helicases of archaea, yeast and some viruses (7,17). However, the absence of higher resolution structures of dodecameric helicases and the limited resolution of our HpDnaB reconstruction (23 Å) prevented insightful structural comparison. It was also unclear whether such a double hexameric assembly could bind and unwind DNA and what its fate was during primosome formation.

Here, we have solved the crystal structure of the HpDnaB dodecamer at a resolution of 6.7 Å, which reveals that the helicase can assemble as a planar, stack-twisted double hexamer. Using a complementary set of methods, we found that HpDnaB interacts preferentially with ssDNA as a dodecamer in the presence of ATP but is poorly active for DNA unwinding. Interaction with HpDnaGHBD dissociates the helicase dodecamer into single-ringed primosomes that represent a more active form of the helicase for DNA unwinding. We propose that in the absence of a control mechanism by AAA+ loaders, dissociation of the double hexamer of the helicase by the primase represents a key switch to launch DNA replication in H. pylori and possibly other replication systems utilizing double hexamer replicative helicases.

**MATERIALS AND METHODS**

Gene cloning, protein expression and purification

His-HpDnaB was purified as previously described (17) with an N-terminal histidine-tag consisting of 28
amino-acids (MHHHHHHGBKPNPLGLDSTENTLYFQG). Selenomethionine-substituted His-HpDnaB (His-HpDnaB<sup>Se</sup>) was produced using the protocol described in (18) and purified as His-HpDnaB (17). For wild-type HpDnaB expression, the DNA sequence of hpdnaB gene was amplified by polymerase chain reaction (PCR) using the forward primer 5'-agctatgagatataaagactttgcaagt-g-3' containing a NdeI restriction site (underlined) and reverse primer 5'-atactgagatataaagactttgcaagt-g-3' containing a XhoI site (underlined). The fragment was inserted into the pACYCDuet-1 vector to produce the pACYChpdnaB expression vector. Single point mutations were introduced into pACYChpdnaB using the Quick change site directed mutagenesis kit (Stratagene) using manufacturer's protocol. An N-terminal deletion mutant change site directed mutagenesis kit (Stratagene) using forward primer 5'-caccgtctcttttcagcctttttaccc-3' (same reverse primer than for wild-type) and in-vitro DNA ligase and reverse primer 5'-caccatgttgcaaaacattgaaaggatcgtgc-3' inserted into pET101/D-TOPO vector (Invitrogen). The sequence encoding for the helicase binding domain of HpDnaG (HpDnaG<sup>HB</sup>, residues 415–559) was amplified from H. pylori genomic DNA (strain 26695) using the forward 5'-caccgtctcttttcagcctttttaccc-3' and reverse 5'-tcataagactttgcaagt-g-3' primers and inserted into pET151/D-TOPO (Invitrogen) to generate the plasmid pETHpdnaGHBD.

HpDnaB and mutants were expressed in E. coli BL21 Star (DE3) cells (Invitrogen) grown in LB medium with chloramphenicol at 50 μg·L<sup>-1</sup> (or ampicline at 100 μg·L<sup>-1</sup> for HpDnaB<sup>Se</sup>) at 37°C until OD<sub>600</sub> = 0.6. Protein expression was induced with 1 mM isopropyl β-D-thiogalactopyranoside (IPTG) during 16 h at 20°C. Harvested cells were resuspended in lysis buffer (10 mM phosphate pH 7.5, 200 mM NaCl) with protease inhibitor (complete EDTA-free, Roche), lysozyme (Roche) and DNase I (Sigma-Aldrich). The cells were lysed by sonication and centrifuged at 16 000 g for 20 min. The soluble fraction was diluted four-fold with buffer A and applied onto a Hi-TrapTM Heparin 5 ml column equilibrated with buffer A (10 mM phosphate pH 7.5, 2 mM βME). The protein was eluted at a 0–100% gradient of buffer B (10 mM phosphate pH 7.5, 1 M NaCl, 2 mM βME). Fractions containing the protein were pooled and applied on a HiTrap Q HP (GE Healthcare) column as a second purification step using the same elution protocol. The protein was concentrated and loaded on a Superdex 200 10/300 GL gel filtration column (GE Healthcare) eluted in 10 mM phosphate pH 7.5, 200 mM NaCl, 2 mM βME.

For HpDnaG<sup>HB</sup> expression, E. coli BL21 Star (DE3) cells (Invitrogen) carrying the pETHpdnaGHBD plasmid were grown in LB medium with ampicillin (100 μg·L<sup>-1</sup>) at 37°C until an OD<sub>600</sub> of 0.6 and protein expression was induced by adding 1 mM IPTG at 20°C overnight. The cells were harvested and resuspended in lysis buffer (50 mM Tris pH 8.0, 200 mM NaCl) with protease inhibitor (Roche), lysozyme (Roche) and DNase I (Sigma). The cells were sonicated and centrifuged at 16 000 g for 20 min at 4°C. The supernatant was applied on a HisTrapTM HP 5 ml (GE Healthcare) pre-equilibrated with buffer C (50 mM Tris pH 8.0, 200 mM NaCl, 1 mM DTT and 5% (v/v) glycerol). The protein was eluted by applying a linear gradient (10–100%) of buffer D (buffer C with 500 mM imidazole). Fractions containing HpDnaG<sup>HB</sup> were pooled and incubated with TEV protease with 1 mM DTT and 0.5 mM EDTA, and dialyzed overnight against buffer D at 4°C. After TEV cleavage the protein fraction was loaded on the HisTrapTM column and HpDnaG<sup>HB</sup> eluted with 10% of buffer B. The protein was injected on a Superdex 200 10/300 GL (GE Healthcare) gel filtration column and eluted with buffer C.

Crystallization, X-ray diffraction data collection and structure determination

Crystals of His-HpDnaB and His-HpDnaB<sup>Se</sup> were grown at 292 K in polyethylene glycol 3350 26%, 100 mM Tris pH 8.6 and 200 mM lithium sulphate with protein incubated with 1 mM ADP and 1 mM MgCl<sub>2</sub>. Drops were set-up by mixing 1.5 μl of protein solution (10 mg·ml<sup>-1</sup>) with 1.5 μl of reservoir solution. Hexagonal crystal forms were obtained after one to two months. Tantalum Bromide derivative crystals were obtained by incubating the crystals for 2 h in the presence of 0.01 mg of Ta<sub>2</sub>Br<sub>12</sub>·27 powder (Jena Biosciences Gmbh). Single crystals were harvested and flash frozen in liquid nitrogen (100 K) using the mother liquor supplemented with 10% sucrose for cryoprotection. Two single-wavelength X-ray diffraction data sets (Tant 1 and Tant 2) were collected from a single Tantalum Bromide crystal on beamline Proxima 2a at the synchrotron SOLEIL. His-HpDnaB<sup>Se</sup> X-ray data were collected on the beamline ID14EH4 at the European Synchrotron Radiation Facility. The diffraction data were indexed and integrated using XDS (19) and scaled with SCALa from the CCP4 program suite (20). To verify that the crystals contained the full-length protein His-HpDnaB, crystals were dissolved and analyzed on SDS-PAGE.

Structure determination, model building and refinement

The structure of HpDnaB was solved by the single anomalous dispersion method with Tant 1 data collected at the inflection wavelength from a single crystal. The positions of three Tantalum clusters were found using HYSS (21) and were then submitted to Phaser (22), which is integrated in Autosol in PHENIX (23). The experimental map at 7.5 Å resolution was of excellent quality and revealed that the asymmetric unit contained four molecules of HpDnaB. Four molecules of the NTD high-resolution (2.2 Å) crystal structure (residues 1 to 120, (24)) of HpDnaB (HpDnaB<sup>NTD</sup>) were placed into the electron density map and the second helices of the four NTD α-hairpins were built using COOT (25). These domains were used for NCS averaging using PARROT (26) which improved the electron density map. Two single helices of the second CTD-ring but the two subunits could not be placed reliably. Soaking the crystals in a solution containing...
ssDNA or co-crystallization experiments in various conditions (ssDNA lengths, nucleotides and concentration) did not improve either the resolution or the electron density map corresponding to this region. No ADP molecule could be placed in the nucleotide binding site since no density was observed in the Fo-Fc map with or without NCS averaging. The resulting model was subsequently refined against Tant 2 data set to extend the resolution to 6.7 Å using BUSTER TNT (27) and PHENIX (23). In the final rounds of refinement, a cluster of Ta6Br122+ could be placed in the density. During refinement, tight geometry constraints were maintained, group B-factors and NCS restraints were used. The final His-HpDnaB model was refined to a resolution of 6.7 Å with Rfactor/ Rfree of 0.25/0.29 and very good geometry. Data collection, phasing and refinement statistics are shown in Table 1. The model coordinates were deposited in the protein data bank (pdb code 4ZC0).

Small-angle X-ray scattering experiments
SAXS data were recorded on beamline Swing at SOLEIL Synchrotron (Gif sur Yvette, France) at a wavelength of 1.003 Å on a 17 cm × 17 cm low-noise Aviex charge-coupled device detector positioned at a distance of 1800 mm from the sample, with the direct beam off-centered. The useful Q-range was 0.004–0.61 Å⁻¹, where Q = 4πsinθ/λ is the scattering vector, and 2θ is the scattering angle. 50 μl of sample in buffer S (50 mM Tris pH 8.0, 200 mM NaCl, 5% (v/v) glycerol, 1 mM DTT) were injected into a size-exclusion column (Shodex KW405-4F) using an Agilent HPLC system cooled at 288 K and eluted directly into the SAXS flow through capillary cell at a flow rate of 200 μl.min⁻¹, as previously described (33). Protein and buffer samples were prepared exactly as described for the MALS experiments, i.e. for samples containing nucleotides, the proteins were first incubated with 5 mM ATP or AMPPNP and 5 mM MgCl2 and the buffer was supplemented with 0.5 mM of the corresponding nucleotides and 5 mM MgCl2 to minimize the effects due to ATP hydrolysis. SAXS data were collected online throughout the whole elution time, with a frame duration of 2 s and a dead time between frames of 1 s. A first data set of 90 frames, collected before the void volume, was averaged to account for buffer scattering. A second data set was collected for the sample, from which the 10 frames corresponding to the top of the elution peak were averaged and used for data processing after baseline subtraction (Supplementary Figure S1). Data were processed using the local application FOXTROT (http://www.synchrotron-soleil.fr/Recherche/LignesLumiere/SWING) and analyzed using PRIMUS (34). After processing, Kratky plots were generated to verify if proteins were correctly folded and P(r) functions were calculated (Supplementary Figure S2). Theoretical curves from the models were generated by FoXS (35). Minimal Ensemble Search was performed using MES program (36).

Chemical cross-linking
Cross-linking of HpDnaB and mutants L4A, Q8A, E80A and D9 was performed in 20 mM HEPES pH 7.6, 150 mM NaCl, 0.5 mM βME and 5% (v/v) glycerol. For HpDnaB with AMPPNP and ssDNA, the protein was incubated 5 min with AMPPNP (5 mM), MgCl2 (5 mM) and a 20mer polydT ssDNA oligonucleotide (75 μM final concentration) at room temperature before cross-linking. Crosslinking agent ethylene glycol bis(succinimidyl succinate) (EGS) (5 mM final concentration) was added to 20 μg of protein in 20 μl final volume. The mixture was then incubated at room temperature during 10 min and the reaction was quenched with 2 μl of 1M Tris pH 7.5. The samples were run on gradient SDS-PAGE (4–12%) and stained with Coomassie blue for analysis.

ATPase activity
The rate of ATP hydrolysis by HpDnaB (500 nM of monomer) in reaction buffer 20 mM Tris pH 8.0, 100 mM KCl, and 5 mM MgCl2 was measured at a concentration of 5 mM ATP and in the presence and absence of 500 nM single stranded 20mer polyT DNA and/or 500 nM

Size exclusion chromatography (SEC)-Multi-angle light scattering (MALS)
Size exclusion chromatography (SEC) combined with multi-angle light scattering (MALS) and refractometry (RI) experiments were performed with a Shodex KW405-4F size exclusion column equilibrated with 50 mM Tris pH 8.0, 200 mM NaCl, 1 mM DTT and 5% (v/v) glycerol. 12.5 μl of protein samples of HpDnaB or mutants (15 mg.ml⁻¹) and HpDnaGHBD (20 mg.ml⁻¹) were injected onto the column. For samples containing nucleotides the proteins were first incubated with 5 mM ATP or AMPPNP and 5 mM MgCl2 and the buffer was supplemented with 0.5 mM of the corresponding nucleotides and 5 mM MgCl2. For samples of the HpDnaB+HpDnaGHBD complex, the separated proteins were incubated at equal molar amounts at a final concentration of 15 mg.ml⁻¹. For ssDNA complex measurements the protein or complexes were first mixed with a 20mer polydT oligonucleotide at either 45 μM or 75 μM (final concentration). On-line MALS detection was performed with a miniDAWN-TREOS detector (Wyatt Technology Corp., Santa Barbara, CA) using a laser emitting at 690 nm and refractive index measurements were performed using an Optilab T-rex system (Wyatt Technology Corp., Santa Barbara, CA). Weight averaged molar masses were calculated using the ASTRA software (Wyatt Technology Corp., Santa Barbara, CA).

Negative staining electron microscopy and image analysis
HpDnaB sample from SEC-MALS (0.1 mg.ml⁻¹) was applied to the clear side of carbon on a carbon-mica interface and stained with 2% (w/v) uranyl acetate. Images were recorded with a JEOL 1200 EX II microscope at 100 kV and at a nominal 40 000x magnification. Negatives were digitized on a Zeiss scanner (Photoscan TD) to a pixel size of 3.5 Å at the specimen level. 9135 individual particles of HpDnaB were semi-automatically selected with the EMAN boxer routine (28), CTF-corrected with CTFIND3 (29) and bssoft (30), and low-path-filtered at 15 Å with Spider (31). This data set was subjected to multivariate statistical analysis and classification with Imagic-5 (32).
Table 1. Data collection, phasing and refinement statistics

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Values in parentheses refer to the indicated resolution shell.

**HpDnaG<sup>HBD</sup>**. ATP hydrolysis was measured using the previously described spectrophotometric method (37) based on the coupling of ATP hydrolysis to the oxidation of NADH with pyruvate kinase and lactate dehydrogenase. The hydrolysis rates presented correspond to the mean of three independent experiments performed with each sample.

**DNA binding**

Equilibrium DNA binding assays were performed on a Clarisostar (BMG Labtech) microplate reader, fitted with polarization filters to measure fluorescence anisotropy. The binding assays were conducted in 384-well plates at room temperature in 40 μl reaction volumes in 20 mM Hepes pH 7.5, 5 mM MgCl₂, 50 mM potassium acetate and 5% (v/v) glycerol supplemented with 0.2 mg.ml⁻¹ BSA and 0.5 mM AMPPNP or ATP. 0 to 100 nM HpDnaB, HpDnaB<sup>•</sup>HpDnaGHBD complex, or 0 to 15 μM HpDnaG<sup>HBD</sup> were titrated into 1 nM 5′-FAM-labeled 20mer dsDNA composed of oligonucleotides 5′-GACTACGTAACAGCTACGTCT-3′ and 5′-AGACGTACAGCTACGTAGTC-3′ or 1 nM 5′-FAM-labeled 20mer polydT ssDNA oligonucleotide. After subtracting the polarization values obtained for DNA alone, the mean data from three independent experiments were fitted to a standard binding equation (y = Bmax*xⁿ/(Kdⁿ+xⁿ)) assuming a single binding site with Hill slope (h) using GraphPad Prism6. The fits were very good, with R² values all above 0.98 (Supplementary Table S1).

**DNA unwinding assay**

Unwinding of forked DNA was carried out by first annealing a 5′-FAM-labeled oligonucleotide (5′-FAM-TACGTAACAGCTACGTCT(dT)₂₅–3′) to a 1.1 molar excess of a 3′ dabcyl quencher-labeled strand (5′-(dT)₂₅-GCAGGCTGTACGTAT-Dabcyl-3′). A capture oligo (5′-GCAGGCTCAGTTACGTA-3′) complementary to the base-paired region of the FAM-labeled strand was added to all reactions to prevent renamelling of the unwound substrate. 80 μl reactions containing 50 nM HpDnaB dodecamer (with or without 7.3 μM HpDnaG<sup>HBD</sup>) and 100 nM fork substrate in helicase buffer (10 mM Tris pH 8.0, 50 mM NaCl, 1% glycerol and 5 mM MgCl₂) were started by addition of 100 nM capture DNA and 2 mM ATP and were monitored at 37°C for 3 h using a Clarisostar (BMG Labtech) fluorescence plate reader. Reactions were performed in triplicate and presented data correspond to the average of at least three independent measurements.

**RESULTS**

**Crystal structure of HpDnaB**

Crystals of N-terminally 6-histidine-tagged HpDnaB (His-HpDnaB) diffracted to 6.7 Å resolution and belonged to
the high symmetry space group I213. The structure was solved by the single anomalous dispersion method and the final model was refined to an $R_{factor} / R_{free}$ of 0.25/0.29 with very good geometry (Supplementary Figure S3A, B and Table 1). Given the low resolution, SeMet-substituted protein crystals were grown and used to collect a single wavelength data set at the Se edge to ascertain the position of the methionines in the structure. An anomalous difference Fourier map confirmed the correct location of modeled methionines leaving no ambiguity regarding the position of the subdomains of His-HpDnaB (Supplementary Table S2 and Figure S3C).

The crystal structure consists of two His-HpDnaB homodimers (chains AB and CD) per asymmetric unit. While chains A and B comprise both the NTD and CTD, chains C and D display only the NTD. Two hexamers, each of which is composed of three homodimers, are generated with the operation of a crystallographic three-fold symmetry (Supplementary Figure S3A). As a result, hexamer 1 consisting of three copies of chains AB is complete with both NTD- and CTD-rings, while hexamer 2 consisting of three copies of chains CD has only the NTD-ring. The absence of the CTD-ring in hexamer 2 cannot be accounted for by protein degradation, since HpDnaB was intact in the crystals (Supplementary Figure S3D). Instead, analysis of the crystal packing indicates that the entire CTD-ring 2 is in a large solvent channel. A model of the complete dodecamer was generated by superimposing a copy of hexamer 1 onto the NTD-ring 2 (Figure 1B). The complete dodecamer measures 190 Å in length and 120 Å in diameter (Figure 1B). When this model is used to generate the crystal lattice, no clashes were observed between symmetry-related molecules and the subunits closest to the CTD-ring 2 are at 25 Å (Sup. Movie M1). This demonstrates that the solvent channel is sufficiently large to accommodate the CTD-ring 2. Given that no particular flexibility of the CTD was detected in EM studies (17), we concluded that the absence of density for the CTD-ring 2 is due to rigid-body movements of individual CTDs permitted by the high crystal solvent content (70%) and the lack of crystal contact in this area.

**HpDnaB forms a stack-twisted double hexamer**

The structure of His-HpDnaB reveals that the last two helices of the NTD fold into a helical hairpin that mediates NTD dimer formation (Figure 1C and Supplementary Figure S3B) as in all DnaB structures (3,4,38). The two NTD-rings of His-HpDnaB are identical and composed of a planar trimer of dimers delimiting a 50 Å wide channel, a state recently described as ‘dilated’, found in apo DnaBs (3,4,38,39) (Figure 1C). The NTD-rings stack on each other, and if the two hexamers are projected onto the same plane, the vectors through the center-of-mass are offset by 60 degrees (Figure 1C). These two hexamers can be described as a ‘stack-twisted’ assembly. Strikingly, the arrangement adopts an apparent six-fold symmetry when observed from the top thus explaining the six-fold symmetry detected in our previous negative staining electron microscopy (nSEM) reconstruction (17). The interactions of the NTD-rings are mediated by head domains associations, burying a total surface of ~1400 Å². Helices α1 and α5 of chain A engage α1 of chain D and α5 of chain C (NTD-ring 2), respectively, and chain B (NTD-ring 1) interacts with chain C via α1 – α1 interactions (Figure 2A). Although the low resolution of the structure does not allow for a precise description of the dodecamer interface, α1 – α1 interactions seem predominantly hydrophobic, involving the first residues of the helix.

The structural basis for the connection between the CTD subunits is the same as in other DnaB structures, with each linker helix inserted in the hexamerisation pocket of the adjacent ATPase domain (Figure 1B,C) (3–5). The CTD-ring adopts a pseudo six-fold symmetry with a closed-ring similar to GP40 (Figure 1C) (3,4). One feature that we identified previously in the structure of the isolated HpDnaBCTD is a large insertion (named HPI for Helicobacter pylori insertion) within the ATPase domain forming a helix-turn-helix composed of α15 and α16 (Figure 1B,C). In the dodecameric crystal structure, the HPI helices from chain A and chain B undergo important conformational changes resulting in different structures compared to the HPI of the HpDnaBCTD structure (17). In chain A, α16 and the N-terminal portion of α15 are ordered while the situation is inverted in chain B with α15 and the C-terminal part of α16 ordered (Figure 1C and Supplementary Figure S4). As anticipated from the nSEM structure, the HPIs form a collar extending the CTD-ring (Figure 1B,C). Interestingly, α15 of chain A (but not of chain B) interacts with the adjacent ATPase domain. This feature suggests a structural explanation for some of the unusual properties of HpDnaB. In contrast with other DnaBs, HpDnaB maintains both ATPase and helicase activities in the absence of the NTD head domain (40). The structure of the His-HpDnaB dodecamer suggests that interaction of the HPI collar with the ATPase domain could stabilize the CTD-ring in the absence of NTD during DNA unwinding and/or translocation experiments. This is also supported by the fact that a HPI-deleted HpDnaB mutant protein does not form hexamers and is inactive for ATP hydrolysis and DNA unwinding (40).

**Dynamics of HpDnaB in solution**

We observed that in the crystal structure, some residues of the N-terminal His-tag are located at the NTD-NTD interface (Figure 2A). Concerned by the fact that the Histag might influence the oligomeric state of HpDnaB as seen in other hexameric AAA+ ATPases (41), we purified the protein without the His-tag (named HpDnaB hereafter). Size-Exclusion Chromatography coupled with Multi-Angle Static Light Scattering (SEC-MALS) experiments showed that HpDnaB elutes as a single peak and has a mass of around 585 kDa. This molecular mass is lower than the expected HpDnaB dodecamer (669 kDa) suggesting that the protein assembly might be in equilibrium between hexamer and dodecamer. Online size exclusion chromatography coupled to small angle X-ray scattering (SAXS) was used to determine the radius of gyration (Rg) of the proteins. Rgs obtained for His-HpDnaB and HpDnaB are comparable (67.4 Å and 66.7 Å, respectively; Supplementary Table S3). Rgs calculated from the model of the HpDnaB hexamer or dodecamer based on our crystal structure are 46.8 Å and 65.5 Å, respectively (Table 2, see also Figures S1 and S2). Thus,
our SAXS data clearly demonstrate that His-HpDnaB and HpDnaB are both predominantly dodecameric in solution. We noticed that the scattering curves of the two proteins were nonetheless different, with the His-HpDnaB curve exhibiting a dip and hump between Q values of 0.07 to 0.09 (Figure 2B). This feature has previously been found in ring shape helicases in solution, including EcDnaB (5,42). This feature is absent in the HpDnaB SAXS curve, suggesting that HpDnaB is more dynamic and that the shapes of the two dodecameric particles are different. By comparing the SAXS profiles with a theoretical SAXS curve of the dodecamer model, we observed that the stack-twisted dodecamer fits better the His-HpDnaB SAXS curve ($\chi^2 = 11.8$) than the HpDnaB curve ($\chi^2 = 23.6$). Both fits were improved by using multi-ensemble searches (MES, i.e. a combination of different structures (36)). A better fit was obtained for His-HpDnaB data ($\chi^2 = 8.8$) using a combination of 91% of dodecamer and 9% of hexamer (Supplementary Figure S5A) and for HpDnaB data ($\chi^2 = 14.1$) using a combination of 75% of dodecamer and 25% of hexamer (Supplementary Figure S5B).

To gain further insights into the organization of HpDnaB dodecamers, nsEM images of a HpDnaB sample were compared with the previously obtained His-HpDnaB images (17). As in the case of His-HpDnaB, class averages of HpDnaB confirm the presence of four-layered particles, but the symmetry is less obvious than for His-HpDnaB (Figure 2C). No 3D reconstruction was attempted at this stage because, in contrast to His-HpDnaB, multivariate statistical analysis did not reveal any clear cyclic or dihedral symmetry. Nevertheless, the obtained classes suggest that the HpDnaB dodecamer also relies on NTD–NTD interactions.

To determine if the interface between the NTD-rings seen in the crystal structure of His-HpDnaB is also relevant in HpDnaB, we generated a series of structure-guided mutants. An N-terminal deletion mutant starting at residue 10 (HpDnaB$^{Δ9}$) and two point mutants L4A (HpDnaB$^{L4A}$) and Q8A (HpDnaB$^{Q8A}$) were constructed to disrupt the contacts between $\alpha_1$ helices. Another mutant, E80A (HpDnaB$^{E80A}$) was designed to target the $\alpha_5–\alpha_5$ interaction. SEC-MALS experiments performed on the purified HpDnaB mutants showed that the mass of the HpDnaB$^{Δ9}$ was 322 kDa, so closer to that of a hexamer (334 kDa) (Figure 2D). In comparison, single mutants HpDnaB$^{L4A}$, HpDnaB$^{Q8A}$ and HpDnaB$^{E80A}$ had a mass of 543, 486 and 368 kDa, respectively. So, the dodecamer–hexamer equilibrium previously observed in HpDnaB MALS measurement was clearly affected in mutants HpDnaB$^{Δ9}$, HpDnaB$^{Q8A}$, HpDnaB$^{E80A}$ but not in HpDnaB$^{L4A}$. The oligomeric states of wild-type and mutants HpDnaBs were then compared using chemical cross-linking (Supplementary Figure S6). HpDnaB in complex with AMPPNP and ssDNA, which form hexamers (see after), was used as a control. In the presence of cross-linking agent, HpDnaB and HpDnaB$^{L4A}$ formed assemblies of the same size and larger than a hexamer, while the mutants HpDnaB$^{Δ9}$, HpDnaB$^{Q8A}$ and to a less extent HpDnaB$^{E80A}$
formed predominantly hexamers (Supplementary Figure S6). These results are in agreement with MALS measurements. Collectively these experiments demonstrate that mutations targeting the interface seen in the crystal structure affect dodecamer formation of *HpDnaB*. We concluded that the structure of *HpDnaB* in solution is similar to the one seen in the crystal structure and that α1 is essential for hexamer–hexamer assembly. However, in the absence of the His-tag, *HpDnaB* probably adopts multiple conformations, some of which might trigger dissociation of the two hexamers. Our data suggest that the N-terminal His-tag reduces the dynamics of the protein, possibly explaining that only the His-tagged protein crystallized. This is also supported by the observation that the values of Rg derived from the SAXS are more stable across the peak of His-*HpDnaB* than across the peak of *HpDnaB* (Supplementary Figure S1).

**HpDnaB binds ssDNA but not dsDNA**

Since DnaBs are loaded onto ssDNA at replication forks and *HpDnaB* is able to translocate in the 5' to 3' direction (40), we investigated the fate of the double ring on DNA. Using fluorescence anisotropy, we found that *HpDnaB* interacts with a 20mer polydT (20dT) ssDNA oligonucleotide in the presence of ATP or the non-hydrolysable ATP analogue, AMPPNP, with dissociation constants (K_D) of 15 ± 1 nM and 3.1 ± 0.1 nM, respectively, but not in the absence of nucleotides (Figure 3A). In contrast, no or very weak binding was observed with dsDNA even in the presence of nucleotides (Figure 3A). Increasing the length of the ssDNA oligonucleotide (Figure 3B) barely increased the binding affinity in the presence of AMPPNP (K_D = 1.8 ± 0.2 nM) and instead reduced the binding affinity of *HpDnaB* in the presence of ATP (K_D = 70 ± 6 nM). The binding of *HpDnaB* to ssDNA was found to be cooperative in the presence of AMPPNP, but not in the presence of ATP (Supplementary Table S1), indicating that the DNA binding mode may be affected by the nucleotide-bound state of *HpDnaB*.

Next, the complexes formed by *HpDnaB* and ssDNA were analyzed by SEC-MALS in the presence of either ATP or AMPPNP. The elution profile and the estimated molecular weight of *HpDnaB* in the presence and absence of 75 µM ssDNA were very similar when 0.5 mM ATP was added to the buffer (Figure 3C). In both cases, the mass of *HpDnaB* derived from the MALS was around 550 kDa thus corresponding to the mostly dodecameric *HpDnaB* (Figure 3B). When SAXS data were collected on the same sample and in the same size exclusion chromatography conditions, the Rgs of *HpDnaB*•ATP and *HpDnaB*•ATP•20dT samples were 67.9 Å and 68.0 Å, respectively (Supplementary Table S2). Together these data suggest that the complex *HpDnaB*•ATP•20dT is a dodecamer. In contrast, in the presence of AMPPNP, the addition of 45 µM of 20dT to the sample resulted in the reduction of the peak corresponding to the *HpDnaB* dodecamer and the appearance of second and third peaks at later elution volumes. The second peak contained *HpDnaB* and had a molecular weight of 310 kDa (Figure 3D). The addition of 75 µM of 20dT to the *HpDnaB*•AMPPNP sample resulted in the complete disappearance of the dodecamer peak. Interestingly, the Rg derived from the SAXS analysis of the sample corresponding to the second peak containing the *HpDnaB*•AMPPNP•20dT was 47.0 Å, corresponding to a hexamer while in the presence of AMPPNP alone, the Rg of *HpDnaB* was 67.6 Å and corresponds to a dodecamer (Table 2).

### Table 2. Summary of the oligomeric states of *HpDnaB* and associated complexes studied

<table>
<thead>
<tr>
<th></th>
<th>MALS</th>
<th></th>
<th>SAXS</th>
<th></th>
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<tr>
<td></td>
<td>MW (kDa)</td>
<td>*HpDnaB oligomer</td>
<td>Rg (Å)</td>
<td>Dmax (Å)</td>
</tr>
<tr>
<td>His-<em>HpDnaB</em></td>
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<td>67.4</td>
<td>234</td>
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<tr>
<td><em>HpDnaB</em></td>
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<td>6–12</td>
<td>66.7</td>
<td>228</td>
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<tr>
<td><em>HpDnaB</em> + ATP</td>
<td>546</td>
<td>6–12</td>
<td>67.9</td>
<td>234</td>
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<tr>
<td><em>HpDnaB</em> + ATP + ssDNA</td>
<td>553</td>
<td>6–12</td>
<td>68.0</td>
<td>235</td>
</tr>
<tr>
<td><em>HpDnaB</em> + AMPPNP</td>
<td>500</td>
<td>6–12</td>
<td>67.6</td>
<td>234</td>
</tr>
<tr>
<td><em>HpDnaB</em> + AMPPNP + ssDNA</td>
<td>310</td>
<td>6</td>
<td>47.0</td>
<td>146</td>
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<tr>
<td><em>HpDnaB</em> + *HpDnaG^{HBD}</td>
<td>345</td>
<td>6 (+ x *HpDnaG^{HBD})</td>
<td>55.0</td>
<td>193</td>
</tr>
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</table>

**Theoretical values**

<table>
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<th>MW (kDa)</th>
<th>*HpDnaB hexamer</th>
<th>Rg (Å)</th>
<th>Diameter (Å)</th>
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<td>65.5</td>
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<tr>
<td><em>HpDnaB</em>•*HpDnaG^{HBD}</td>
<td>385</td>
<td>6 (+ x *HpDnaG^{HBD})</td>
<td>50.7</td>
<td>152</td>
</tr>
</tbody>
</table>

MW: Molecular weight; x = 1, 2 or 3; † Model obtained using BstDnaB• BstDnaG^{HBD} structure (pdb code 2R6A) as template. # Theoretical Rg and diameter values were obtained using CRYOSOL (49).

**HpDnaB interaction with *HpDnaG^{HBD}*** dissociates double rings into hexamers

We investigated the oligomeric state of *HpDnaB* during formation of the *HpDnaB•HpDnaG* complex. The crystal structure of the BstDnaB•BstDnaG^{HBD} revealed that each of the three DnaG^{HBD} engages the NTD dimer via interactions with the head domains of BstDnaB (4). To gain insight into the assembly of the primosome of *H. pylori* we analyzed the *HpDnaB•HpDnaG^{HBD} complex obtained by mixing the two proteins together using SEC-MALS and...
Figure 3. Effects of ATP and AMPPNP on the HpDnaB interaction with DNA. (A) Fluorescence anisotropy measurements of HpDnaB binding to either 5′-FAM labeled 20mer ssDNA (20dT) or dsDNA in the presence and absence of nucleotide (ATP and AMPPNP). The curves represent the mean of three independent experiments. (B) Similar experiments performed with 5′-FAM labeled 50mer ssDNA (50dT). (C) Size exclusion chromatograms (280 nm) of HpDnaB alone (light blue) or mixed with a 20dT oligonucleotide (75 μM) and 5 mM ATP (dark blue). MALS weight-averaged molar masses are indicated as dotted lines. The SDS-PAGE analysis shows the HpDnaB protein elution fractions. (D) SEC-MALS experiment performed as in C) except that HpDnaB was incubated with 5 mM AMPPNP and the running buffer contained 0.5 mM AMPPNP. Three samples were analyzed: HpDnaB (blue) and HpDnaB with 45 μM (orange) or 75 μM (red) 20dT. The SDS-PAGE analysis shows that HpDnaB elutes slightly later in the presence of ssDNA and AMPPNP.

SAXS. SEC experiments showed that this complex eluted as two peaks and SDS-PAGE showed that the first peak contained both HpDnaB and HpDnaG<sup>HBD</sup> <sup>HBD</sup>, while the second contained exclusively HpDnaG<sup>HBD</sup> (Figure 4A). Measurements of the molecular weight by MALS indicated that the peak corresponding to the HpDnaB•HpDnaG<sup>HBD</sup> complex had a molecular mass of approximately 345 kDa, suggesting that this peak corresponds to hexameric rather than dodecameric HpDnaB (the theoretical mass of the hexamer is 334 kDa) with one or perhaps more HpDnaG<sup>HBD</sup> (17 kDa) molecules bound. HpDnaG<sup>HBD</sup> alone had a mass of around 35 kDa corresponding to a dimer as previously observed (43).

SAXS data corroborated these observations with the complex HpDnaB•HpDnaG<sup>HBD</sup> having an Rg of 55.0 Å, much lower than a dodecamer (66.7 Å) but greater than a theoretical HpDnaB hexamer (46.8 Å) (Supplementary Table S3). We then modeled the HpDnaB<sub>6</sub>•HpDnaG<sub>HBD</sub><sub>3</sub> complex using the crystal structures of His-HpDnaB and HpDnaG<sup>HBD</sup> and the BstDnaB<sub>6</sub>•BstDnaG<sub>HBD</sub><sub>3</sub> crystal structure as a template (43). A very good fit of the experimental SAXS profile was obtained with the theoretical curve derived from the HpDnaB<sub>6</sub>•HpDnaG<sub>HBD</sub><sub>3</sub> model (χ<sup>2</sup> = 5.8) compared with the curve derived from the HpDnaB dodecamer (χ<sup>2</sup> = 36.8) (Figure 4B). When using MES, the fit was significantly improved with a mixture of 89% of the HpDnaB<sub>6</sub>•HpDnaG<sub>HBD</sub><sub>3</sub> complex and of 11% of HpDnaB dodecamer (χ<sup>2</sup> = 3.5; Figure 4B), suggesting that the peak observed in SEC-MALS contains both uncomplexed dodecameric HpDnaB and hexameric HpDnaG<sup>HBD</sup>-bound HpDnaB. Together, these data reveal that formation of the H. pylori primosome results in the dissociation of the helicase dodecamer into hexamers that can interact with up
to three HpDnaG<sup>HBD</sup> in a conformation similar to the crystal structure of BstDnaB<sub>6</sub>•BstDnaG<sup>HBD</sup><sub>3</sub> (Table 2).

**HpDnaG<sup>HBD</sup> interacts with HpDnaB and ssDNA to prime helicase and ATPase activities**

A major role of primosome formation is the modulation of primase and helicase activities (14,44–46). As for other DnaBs, the ATPase activity of HpDnaB has previously been shown to be stimulated by HpDnaG<sup>HBD</sup> or by ssDNA (17,40). We have thus investigated the effect of HpDnaG<sup>HBD</sup> on the ability of HpDnaB to hydrolyse ATP, bind DNA and unwind forked DNA duplexes. HpDnaB had an ATPase activity of 0.04 ± 0.03 µM ATPs<sup>−1</sup> at a concentration of 500 nM (Supplementary Table S4). As expected, addition of HpDnaG<sup>HBD</sup> or ssDNA increased the ATPase activity of HpDnaB approximately three-fold (Supplementary Table S4). The activity was increased almost eight-fold by adding both ssDNA and HpDnaG<sup>HBD</sup>, suggesting that the...
ternary complex is the most active for ATP hydrolysis with a hydrolysis rate of 0.35 ± 0.08 μM ATPs⁻¹ (Supplementary Table S4).

The binding affinity of these proteins to ssDNA was then measured by fluorescence anisotropy. The HpDnaB·HpDnaGHBD complex exhibited a higher affinity for ssDNA than HpDnaB alone in the presence of ATP (K_D = 8.3 ± 0.6 nM versus 15 ± 1 nM) or AMPPNP (K_D = 1.0 ± 0.1 versus 3.1 ± 0.1 nM) (Figure 5A). HpDnaGHBD thus increases the affinity of HpDnaB for ssDNA in the presence of nucleotides. Interestingly, HpDnaGHBD alone interacts with ssDNA, but with an affinity in the micromolar range (Supplementary Figure S7). Translocation of the double hexamer on ssDNA poses several mechanistic problems. (i) HpDnaB unwinds dsDNA in the 5' to 3' direction, so only one hexamer can be active; (ii) the second (inactive) hexamer would considerably hinder the first one’s activity by binding non-specifically to ssDNA via its DNA binding loops; (iii) an inactive hexamer would represent a significant inert mass to be translocated by the active one. This suggests that for efficient helicase activity, a double hexamer would have to be separated to release the correctly positioned HpDnaB hexamer. To test this hypothesis we analyzed the helicase activity of HpDnaB in the presence and absence of HpDnaGHBD. We used a fluorescence-based assay similar to the one used previously to measure EcDnaB·EcDnaC helicase activity (10). A forked DNA substrate was formed by annealing an oligonucleotide bearing a 3’ ssDNA tail and a fluorescein label at its 5’ end with a complementary oligonucleotide bearing a 5’ ssDNA tail and a quenching dye (Dabcyl) at its 3’ end. As seen in Figure 5B, double hexameric HpDnaB displayed weak DNA unwinding activity, rapidly reaching a maximum of only 10% DNA unwound. In contrast, in complex with HpDnaGHBD, HpDnaB was able to unwind dsDNA, reaching 60% DNA unwound. The initial rates of DNA unwinding were very similar in both cases, confirming that HpDnaB is indeed an active DNA helicase, but the activity of the enzyme was drastically affected by the presence of the primase HpDnaGHBD. No more activity was detected for HpDnaB alone after approximately 20 min, whereas in the presence of HpDnaGHBD, helicase activity could still be detected 2 h after the start of the reaction.

**DISCUSSION**

DNA replication relies on replicative helicases to initiate progression of replication forks and to anchor the primase. While most DnaBs characterized to date have been reported to be hexameric, we previously found that HpDnaB from *H. pylori* was able to form double hexamers, thus questioning how this structure would assemble and function on replication forks. In this study, we have addressed the structural mechanisms of HpDnaB activation during primosome formation on ssDNA.

The crystal structure of the HpDnaB dodecamer solved here provides significant information on HpDnaB dodecamer compared to our previous nsEM reconstruction (17). It reveals that the NTDs of the helicase are arranged in dilated collars that self-interact in a stack-twisted mode to generate a head-to-head double hexamer. The dodecamer assembly stabilizes the NTD-ring conformation but allows for the CTD-ring to remain sufficiently dynamic to accept structural changes associated with ATP hydrolysis or ssDNA binding. Our work establishes that if the topology of this bacterial dodecameric helicase (i.e. NTD–NTD in-
teractions) is reminiscent of Mcm2–7 and some MCMs, the structural and molecular determinants of the hexamers association are different. In Methanobacterium thermoautotrophicum MCM (MtMCM) (47), dodecamerization relies on interactions between loops that coordinate Zn atoms in the MtMCM B domain (Supplementary Figure S8). The structural basis for dodecamer formation is completely different in HpDnaB (Figure 2) and thus rules out a common evolutionary structural motif. Nevertheless, it is interesting to note that, despite diverse evolutionary routes, dodecameric replicative helicases might exist in all kingdoms of life, suggesting that some selective advantages might be associated with this organization (48).

It is as yet unclear how HpDnaB is loaded onto the replication forks and if an additional factor is required for this process (Figure 5C). In this regard, we found that the double hexameric HpDnaB does not interact efficiently with ds-DNA contrary to ss-DNA, suggesting that unwound origins of replication might be better substrates (Figure 3A). The HpDnaB dodecamer positions two hexamers in opposite directions but the two rings cannot unwind ds-DNA simultaneously in the 5′ to 3′ direction if a single strand passes through the double ring. Accordingly, the use of a longer ss-DNA oligonucleotide did not increase the affinity of the HpDnaB for DNA, suggesting that only one of the two rings of the dodecamer is efficiently bound to ss-DNA (Figure 3B).

We found that the HpDnaB dodecamer is indeed poorly active for DNA unwinding and that the protein unwinds ds-DNA as a hexamer, similar to other DnaBs (12). ATP hydrolysis and ss-DNA binding are not sufficient to separate the two rings (Figure 3C and Table 2), suggesting that an additional factor is required to initiate DNA unwinding at the replication fork. Our study suggests that the primase might be such a factor since the interaction with HpDnaG HBD separates the HpDnaB double hexamer (Figures 3D, 5C and Table 2). This can be explained by the crystal structure of His-HpDnaB in which the dodecamer interface mediated by NTD collars overlaps the primase interaction site described in BstDnaB (4) and conserved in HpDnaB (43). HpDnaG association with HpDnaB not only separates the dodecamer, but also increases its ATPase activity, its affinity for ss-DNA and its DNA unwinding capacity. These findings suggest that the separation of the dodecamer by HpDnaG drives DNA unwinding by one hexamer in the 5′ to 3′ direction allowing replication to initiate (Figure 5C). Mechanistically, the separation of the two helicase rings by interaction with the primase might also be relevant to other systems such as archaeal or eukaryotic systems, which also exploit double ring helicases during replication initiation.

SUPPLEMENTARY DATA
Supplementary Data are available at NAR Online.

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Conflict of interest statement. None declared.

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