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Solid-state NMR chemical-shift perturbations indicate domain reorientation of the DnaG primase in the primosome of *Helicobacter pylori*

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Abstract

We here investigate the interactions between the DnaB helicase and the C-terminal domain (CTD) of the corresponding DnaG primase of *Helicobacter pylori* using solid-state NMR. The difficult crystallization of this 387 kDa complex, where the two proteins interact in a six to three ratio, is circumvented by simple co-sedimentation of the two proteins directly into the MAS-NMR rotor. While the amount of information that can be extracted from such a large protein is still limited, we can assign a number of amino-acid residues experiencing significant chemical-shift perturbations upon helicase-primase complex formation. The location of these residues is used as a guide to model the interaction interface between the two proteins in the complex. Chemical-shift perturbation also reveals changes at the interaction interfaces of the hexameric *HpDnaB* assembly on *HpDnaG* binding. A structural model of the complex that explains the experimental findings is obtained.

Introduction

The DnaB helicase and the DnaG primase are proteins acting at the heart of the replication machinery which function is to produce identical copies of the genomic DNA of bacteria (Corn and Berger 2006). In order to do so, the DNA double helix is unwound by the DnaB helicase in the 5'-3' direction. DnaB also recruits the DnaG primase, which synthesizes RNA primers to be used as template by the DNA polymerase. The DnaB helicases generally function as hexamers, and the N-terminal domain displays trimers of dimers arranged in a head-to-tail manner. Structures from DnaB domains of different organisms (Kashav et al. 2009; Stelter et al. 2012) and the interacting DnaG C-terminal domain (CTD), (Oakley et al. 2005; Abdul Rehman et al. 2013) as well as a complex of DnaB/DnaG-CTD from *Bacillus stearothermophilus* (*Bst*) (Bailey et al. 2007) are available.

The structures of different DnaG-CTDs reveal a fold very similar to the N-terminal domain of the DnaB helicase, consisting of a globular domain and a helical hairpin region (Oakley et al. 2005; Syson et al. 2005; Su et al. 2006; Abdul Rehman et al. 2013). The DnaG-CTD helical hairpin has been described to adopt multiple orientations relative to the globular domain in the crystal structures of isolated DnaG-CTDs from different organisms, and is thus believed to have a flexible orientation with respect to the globular domain. This has also been revealed by NMR studies, which showed that the linker region between the two domains is flexible and dynamic (Oakley et al. 2005).

Helicobacter pylori is a common pathogen and can be at the origin of various diseases, from chronic gastritis to gastric cancers (Parsonnet 1995). In *H. pylori*, DnaG (*HpDnaG*) and DnaB (*HpDnaB*) show *in vitro* moderate interactions when compared to other organisms as *E. coli* and *B. stearothermophilus* (Abdul Rehman et al. 2013), which may be a reason why crystallization of the complex proved difficult. For solid-state NMR studies, sedimentation of proteins has recently emerged as a valuable sample-preparation venue and enables the study of large soluble protein assemblies (Mainz et al. 2009; Bertini et al. 2011; Gardiennet et al.

2012). This technique has been shown to yield high-resolution NMR spectra for several preparations, including of *HpDnaB* (Gardiennet et al. 2012). We here use this approach to study a complex formed by two different interacting proteins, in which the large molecular weight of one protein multimer allows to sediment the complex it forms with a smaller interacting protein. The analysis of the chemical-shift changes of *HpDnaB* allows to identify the interaction interfaces in the *HpDnaB/HpDnaG-CTD* complex, and we devise a model in which the hairpin in the DnaG-CTD of the primase undergoes a conformational change with respect to its X-ray structure in isolation (Abdul Rehman et al. 2013) in order to comply with the observed chemical-shift perturbations.

HpDnaB, in the presence of *HpDnaG-CTD*, forms a protein complex. In order to evaluate whether the two proteins cosediment, we analyzed the pellet and supernatant fractions after ultracentrifugation of a test sample (Fig. S1). We recorded 2D ^{13}C - ^{13}C solid-state NMR DARR spectra of the sedimented *HpDnaB* full-length protein (Gardiennet et al. 2012), as well as of the co-sedimented *HpDnaB/HpDnaG-CTD* sample. Spectra of the isolated N-terminal domain (NTD) of *HpDnaB*, which has been sequentially assigned (Wiegand et al. 2015), as well as the C-terminal domain (CTD) of *HpDnaB* (both in crystalline form), served as a reference in this work.

Results

An overlay of the DARR spectra from sedimented ^{13}C , ^{15}N labeled *HpDnaB* and the co-sedimented protein complex between ^{13}C , ^{15}N labeled *HpDnaB* and *HpDnaG-CTD* (at natural isotopic abundance) is shown in Fig. 1. The quality of both spectra is comparable, with linewidths (full-width at half maximum) of isolated signals of about 0.5 ppm. Spectra of sedimented *HpDnaB* share common features when *HpDnaB* is alone or in the presence of *HpDnaG-CTD*, which indicates, not surprisingly, that the overall conformation of *HpDnaB* is

conserved in the co-sediment. A closer look at the superposition of the two spectra, however, allows to identify clear spectral modifications due to the presence of *HpDnaG*-CTD (which itself is not visible as it is not isotopically labeled). This indicates that the co-sedimented sample represents a well-defined protein complex and that *DnaB* undergoes local conformational changes when interacting with *DnaG*.

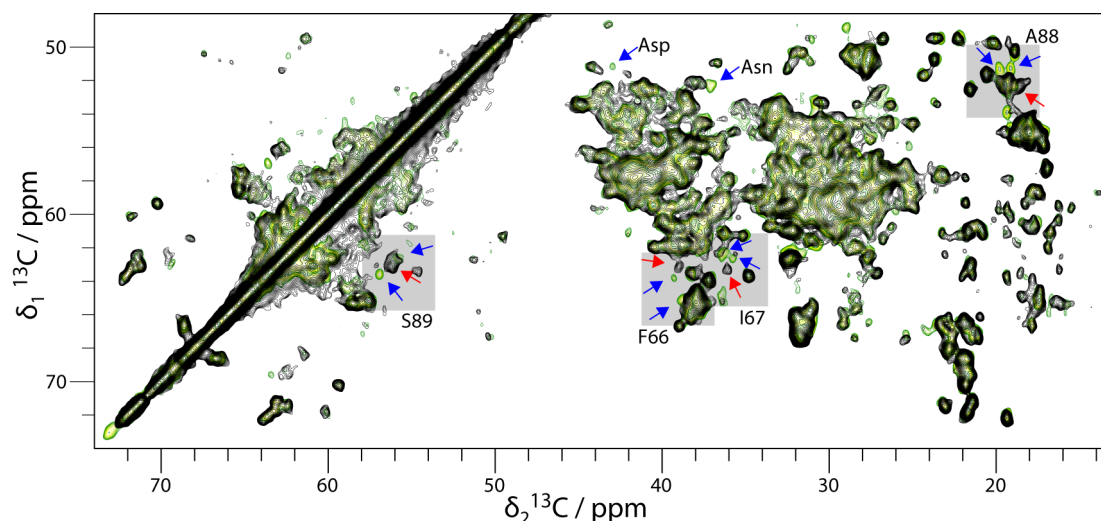


Figure 1. Superposition of the aliphatic regions of 20 ms DARR spectra recorded on the *HpDnaB/HpDnaG*-CTD co-sediment (green) and the *HpDnaB* sediment (black). Red arrows indicate signals which “disappear” in the spectra of the sedimented *HpDnaB*, and blue arrows those which are “newly” observed in the *HpDnaB/HpDnaG*-CTD co-sediment and which will be further discussed in the text. Extracts corresponding to the grey regions are shown in Fig. 2. Individual spectra are given in the Supporting Information (Fig. S2), as well as the full aliphatic region including new signals in the isoleucine side-chain region (Fig. S3).

The most important chemical shift changes concern resonances that appear only in one of the two spectra. These signals can be classified into two categories: resonances present in *HpDnaB* but no longer observed in the protein complex (red arrows in Fig.1), and resonances

which appear only in the spectra of the protein complex (blue arrows in Fig.1, see also Fig. S3). Most probably, these resonances belong to the same residues but are shifted, as a consequence of the complex formation inducing a considerable chemical-shift perturbation (CSP). For further resonances, small changes in chemical shift were observed. As they lie either within or very close to the experimental error which we estimate to about 0.3 ppm, we will not further discuss these. Other differences between the two spectra are explained by slight intensity changes in the corresponding resonances, e.g. resonances that drop below the lowest contour level in one of the spectra but can still be identified in both spectra upon detailed inspection.

Residues that disappear from NMR spectra are either no longer observed due to dynamical or static disorder, or shift so far that they can not be easily paired with the original resonance. The latter is in the current context more likely since, considering the appearance of narrow signals on interaction, spectral evidences for increasing disorder or dynamics on complex formation are not detected in the NMR spectra. Chemical shifts are a highly sensitive probe for conformational changes induced by interactions with partner molecules (for a recent review, see (Williamson 2013)). While solution-state protein-ligand interactions are often found to be in the fast exchange limit for NMR, with continuous titration curves for the CSP, the protein complexes observed here are not in equilibrium with the unbound *HpDnaB* and we observed only disappearing signals from the *HpDnaB* and new signals for the complex arising. For full-length *HpDnaB*, sequence specific assignments of the peaks of interest can be obtained by comparing the spectra of *HpDnaB* to spectra recorded on its isolated N-terminal and C-terminal domains. If the peaks overlap within 0.3 ppm with signals which were assigned in the N-terminal domain (Wiegand et al. 2015), and if no resonance from the C-terminal domain is found in vicinity, the assignment from the isolated N-terminal domain can be taken over. This is illustrated in Fig. 2 where an overlay, as well as the individual spectra,

are displayed of extracts from *HpDnaB*-NTD (Wiegand et al. 2015), *HpDnaB*, and also the *HpDnaB/HpDnaG*-CTD protein complex. Four of the disappearing peaks could be assigned this way, F66, I67, A88 and S89. One more Ala peak is disappearing, but it could not be assigned, since it does not correspond to a signal observed in the isolated N- or C-terminal domains.

Correspondingly, several peaks are only present in the *HpDnaB/HpDnaG*-CTD spectra, which are highlighted by blue arrows in Figs. 1 and S3. Tentative assignments of several appearing resonances can be proposed by pairing them with the most nearby disappearing resonances (grey boxes in Figs. 1 and S3). For F66, I67, A88 and S89, actually *two* newly appearing peaks are detected in close vicinity. This is shown in the right-most column of Fig. 2, where in the *HpDnaB/HpDnaG*-CTD spectra these appearing signals are labeled with red crosses. The CSP as identified under these assumptions are quantified in Table S1.

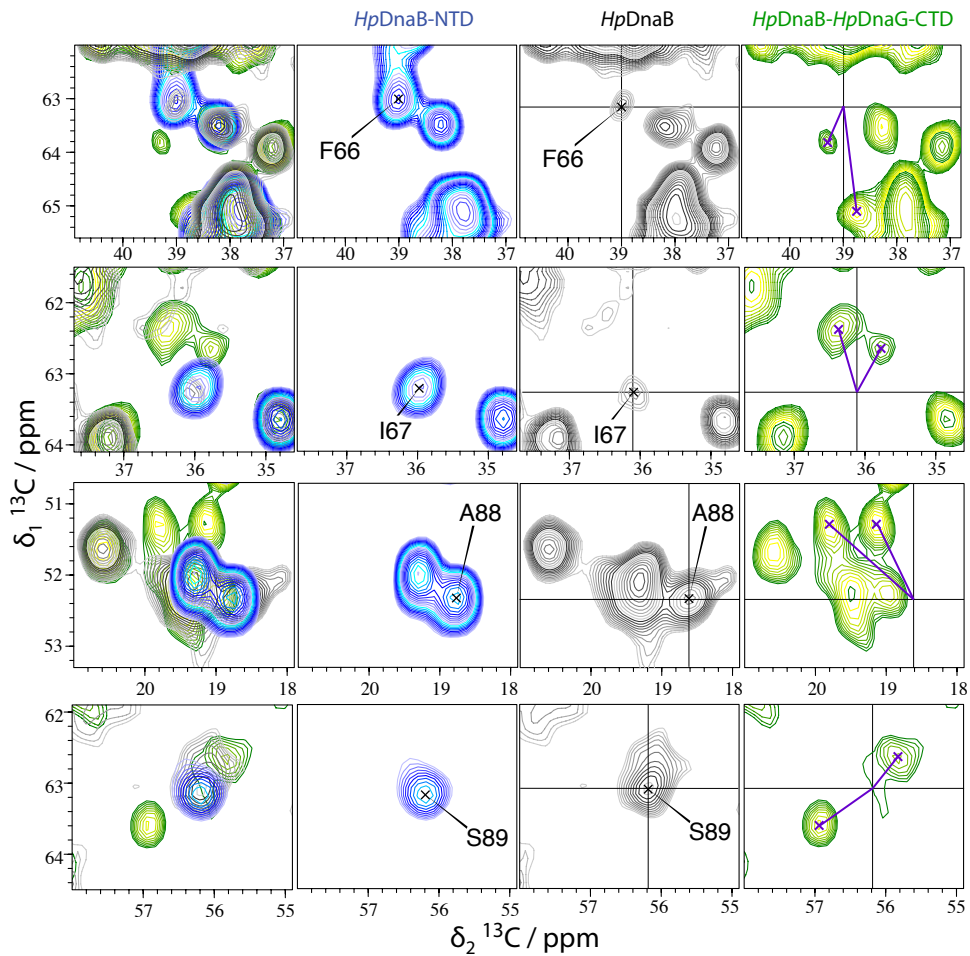


Figure 2: 2D extracts of 20 ms DARR spectra recorded on the *HpDnaB*-NTD (in blue), *HpDnaB* (in black) and the *HpDnaB*/*HpDnaG*-CTD complex (in green). The corresponding 1D traces are shown in Fig. S4. Black crosshairs mark assigned peaks that are observed in *HpDnaB* alone but not in the complex; purple crosses mark peaks that newly appear in the complex; and purple lines indicate the likely CSP shifts involved. Signal doubling is assigned to the asymmetric interaction with *HpDnaG* (see text).

Additionally, appearing resonances of one Asn, one Asp and at least two Ile residues can be identified amino-acid type specifically. These signals are highlighted outside the grey boxes with blue arrows in Figs. 1 and S3. For these residues, no nearby disappearing signals could be identified.

Discussion

The interaction between DnaB and DnaG-CTD has been shown to involve mainly the N-terminal domain of the DnaB helicase. Indeed, its interaction with the DnaG primase has been revealed for *B. stearothermophilus* by X-ray crystallography (Bailey et al. 2007). For the *H. pylori* protein complex, we built a homology model of the *HpDnaB/HpDnaG-CTD* protein complex based on the *HpDnaB* (Bazin et al. 2015) and *HpDnaG-CTD* (Abdul Rehman et al. 2013) crystal structures and their alignment on the crystal structure of the *BstDnaB/BstDnaG-CTD* complex (Bailey et al. 2007). To support the homology model, the sequence alignment of *Bst* vs. *Hp* for both DnaB and DnaG-CTD is shown in Figs. S5 and S6. The hexameric ring formed by the *HpDnaB*-NTDs is shown in grey in Fig. 3, with two monomers highlighted in green, and the *HpDnaG-CTD* primase is shown in pink. Residues F66, I67, A88 and S89 from *HpDnaB* for which disappearing signals were identified are highlighted in blue.

One primase is interacting with two monomers of the helicase, and it can be seen in Fig. 3A that for one of the monomers, the A88/S89 pair is located in this model right at the interaction interface, at the beginning of the loop contacting the *HpDnaG-CTD*. On the other monomer, the pair seems far from the interaction site. F66/I67 are, in one monomer, located at the interface between two *HpDnaB*-NTD molecules, and on the other monomer they belong to a solvent-exposed helix. From the model, we predict that the shifts for residues A88 and S89 can be interpreted by conformational changes caused by the direct interaction with the *HpDnaG* primase with the *HpDnaB* helicase, whereas F66/I67 experience a perturbation due to changes at the *HpDnaB*-NTD dimer interface upon binding. Still, the simple homology model does not account for all observations. It would predict that a signal with half the intensity would be left at the original position of the crosshairs in Fig. 2. Instead, this peak vanishes and a *pair* of new peaks (red crosses in Fig. 2) can be associated with the vanishing peak. Considering that the C-terminal hairpin of the DnaG primase has been shown to be flexible (Oakley et al. 2005; Syson et al. 2005), this behavior can however be predicted with a

modified model where the hairpin adapts on binding to maximize the interaction interface. Such a model has been put forward using energetical arguments combined with mutational studies (Abdul Rehman et al. 2013). When considering this alternative conformation of *HpDnaG*-CTD (Abdul Rehman et al. 2013), as shown in Fig. 3B/C in dark red, the A88/S89 residues, as well as the F66/I67 of the other *HpDnaB* monomer, come close to the intermolecular interface, and can make contact to *HpDnaG*-CTD. This would fully account for the shifting of the resonances to a pair of new positions, and our data thus strongly support an interaction in which the *HpDnaG*-CTD hairpin is making contact, as proposed (Abdul Rehman et al. 2013), with the second monomer of *HpDnaB*-NTD. This model also produces similar interaction interfaces as observed in the crystal structure of the *BstDnaB*/*BstDnaG*-CTD complex (Bailey et al. 2007). There, the hairpin of *BstDnaG*-CTD indeed interacts with the second monomer of *BstDnaB* as well. The conformation observed in the *HpDnaG*-CTD crystal structure might thus have been selected out of a variety of flexible conformations in solution for its ability to crystallize, but likely does not correspond to the one which is able to interact with *HpDnaB*.

The signals which are observed only in the spectra of the *HpDnaB*/*HpDnaG*-CTD protein complex, and for which no neighboring missing peaks can be identified in the isolated *HpDnaB* protein, must be located in regions in which only one monomer contacts *HpDnaG*, and the other remains untouched. This would support the assignment of these signals to the loop containing residues I91-D92-N93-I94, which are shown in grey in Fig. 3C. This region remained unassigned in the isolated *HpDnaB*-NTD domain due to the absence of signal in the 3D spectra (Wiegand et al. 2015), which is indicative of it being dynamic in isolation. In the complex, this loop is in intimate contact with *HpDnaG*-CTD for one *HpDnaB*-Nter monomer, but not for the other, which yields a rationale to the appearance of new resonances. Still, in

order to confirm this hypothesis, more information has to be obtained to assess the identity of the appearing peaks.

Conclusions

Well-resolved NMR spectra can be obtained from a DnaB sediment alone and from a cosediment of the DnaB/DnaG protein complex. A comparison of the spectra allows us to identify residues that undergo large chemical-shift changes. In the context of a previously proposed model, we can interpret these changes and add experimental support to a so far theoretical model. In addition, the study illustrates opportunities and challenges when working with large protein assemblies where a full sequential assignment is presently extremely demanding.

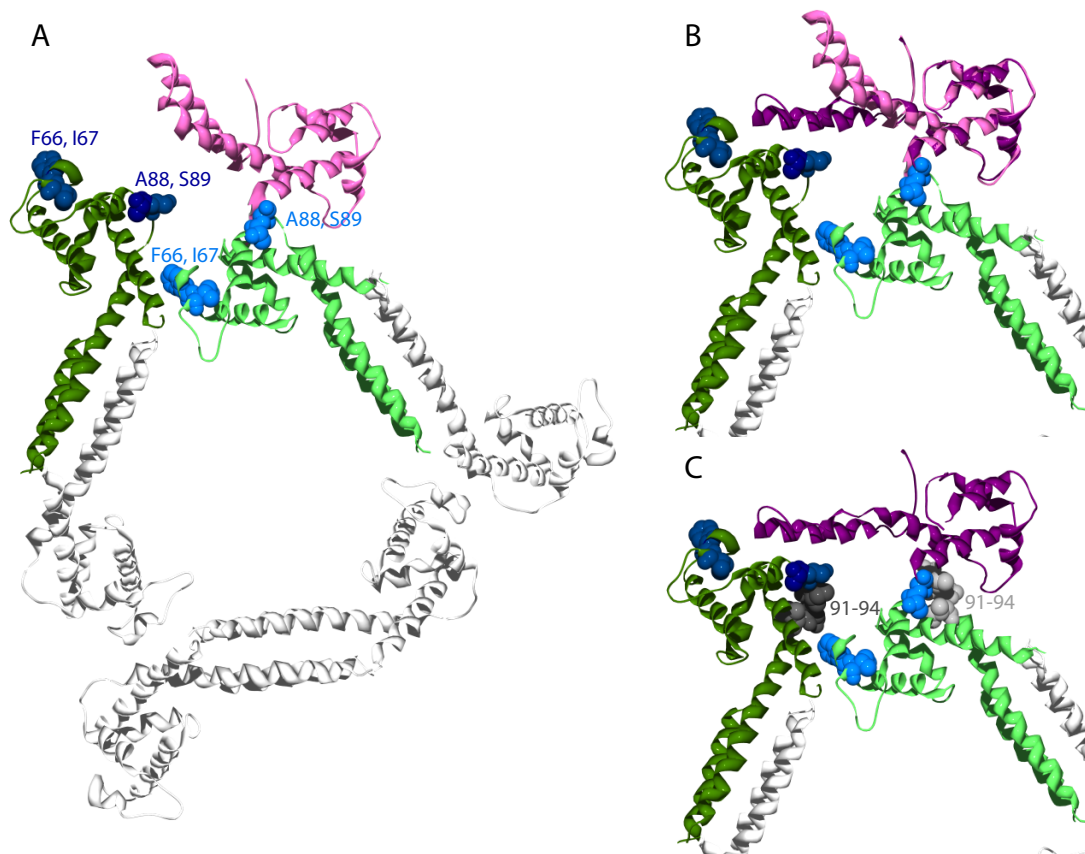


Figure 3: A: Homology model of the *HpDnaB/HpDnaG-CTD* protein complex based on the *HpDnaB* (Bazin et al. 2015) and *HpDnaG-CTD* (Abdul Rehman et al. 2013) crystal structures and their alignment on the crystal structure of the *BstDnaB/BstDnaG-CTD* complex (Bailey et al. 2007). Identified interacting residues F66, I67, A88 and S89 are highlighted in two different shades blue for the two non-equivalent monomers. B: An alternative conformation modeled for *HpDnaB-CTD* built in order to satisfy interactions singled out in mutational studies (Abdul Rehman et al. 2013) is shown in violet. The globular part of the *HpDnaG-CTD* is aligned for the two proteins. C: Putative interacting residues 91-94 are shown in grey for the two monomers.

We have shown here how solid-state NMR of co-sedimented *HpDnaB* helicase and *HpDnaG-CTD* primase can be used to reveal interaction interfaces in a large, multimeric and dynamic protein complex. Using the NMR derived restraints to guide homology-model building and

taking into account the flexibility of the *HpDnaG*-CTD helical hairpin leads to a structural model which fully explains the spectral features identified, and in which the two proteins form an extended interface involving two *HpDnaB* monomers. The identified interacting residues of *HpDnaB* open the way to mutational studies to gain further insights into the helicase-primase interaction central in replication.

Experimental Section

Protein expression and purification

Uniformly ^{13}C , ^{15}N enriched *HpDnaB* was expressed from plasmid pACYC-duet1 in *E. coli* BL21 Star (DE3) cells (Invitrogen) using minimal M9-type media (Studier 2005) supplemented with uniformly ^{13}C enriched glucose (2 g/L) and ^{15}N -labeled ammonium chloride (2 g/L) as sole carbon and nitrogen sources. The protein was purified using affinity chromatography (5 mL HiTrap Heparin HP column from GE Healthcare) followed by anion exchange chromatography (5 mL HiTrap Q HP from GE Healthcare). Purified protein was in 10 mM sodium phosphate buffer at pH 7.5 containing 150 mM NaCl, 1 mM DTT and 0.05% sodium azide. It was concentrated up to 30 mg/mL by centrifugation using Amicon Ultracel-10K (Millipore).

The gene coding for *HpDnaG*-CTD (residues 415-559) was cloned into the plasmid pET151/D-TOPOR (Invitrogen). Unlabeled *HpDnaG*-CTD was expressed in *E. coli* BL21 Star (DE3) cells (Invitrogen) using LB medium. The protein was purified using affinity chromatography (5 mL HisTrap HP from GE Healthcare). After TEV-cleavage of the N-terminal His-tag, the protein was further purified by size-exclusion chromatography (Superdex 200 10/300GL column, GE Healthcare) in a buffer containing 50mM Tris-HCl at pH 8.0, 200mM NaCl, 1mM DTT and 5 % glycerol. It was then concentrated up to 13 mg/mL by centrifugation using Amicon Ultracel-3K (Millipore).

A mixture of 10 mg *HpDnaB* and 10 mg *HpDnaG*-CTD was concentrated up to 42 mg/mL total protein concentration using Amicon Ultracel-5K (Millipore) and sedimented by ultracentrifugation at 200000 g, 4 °C during 10 hours.

NMR spectroscopy

Solid-state NMR experiments were carried out on Bruker Biospin AVANCE III spectrometers operating at 800 MHz and 850 MHz ¹H frequency using 3.2 mm triple-resonance (¹H, ¹³C, ¹⁵N) E-free probes (Bruker Biospin). Sample temperature was determined using the chemical shift of supernatant water (Böckmann et al. 2009) and was adjusted to 278 K. Dipolar Assisted Rotational Resonance DARR (Takegoshi et al. 2001; Scholz et al. 2008) experiments were recorded with a mixing time of 20 ms. Spectra were processed with Topspin using a shifted cos² function and analyzed using CcpNmr Analysis (Stevens et al. 2011). For full experimental parameters see Table S2.

Model building

HpDnaB/ HpDnaG-CTD models were built using the crystal structure of *HpDnaB* (pdb code 4ZC0). A hexamer of *HpDnaB* was superimposed to the structure of *BstDnaB/BstDnaG*-CTD complex (pdb code 2R6A (Bailey et al. 2007)). Then three subunits of *HpDnaG*-CTD (pdb code 4EHS (Abdul Rehman et al. 2013)) were superimposed to the three *BstDnaG*-CTD subunits to generate a *HpDnaB*₆/*HpDnaG*-CTD₃ complex. Minimal rearrangements were required to prevent small steric clashes between *HpDnaB* and *HpDnaG*-CTDs. The same method was used to generate the second *HpDnaB/ HpDnaG*-CTD model, but this time using three modified *HpDnaG*-CTD models as described in (Abdul Rehman et al. 2013) and provided by Dr. Gourinath.

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References

- Abdul Rehman SA, Verma V, Mazumder M, et al (2013) Crystal structure and mode of helicase binding of the C-terminal domain of primase from *Helicobacter pylori*. *J Bacteriol* 195:2826–2838. doi: 10.1128/JB.00091-13
- Bailey S, Eliason WK, Steitz TA (2007) Structure of Hexameric DnaB Helicase and Its Complex with a Domain of DnaG Primase. *Science* 318:459–463. doi: 10.1126/science.1147353
- Bazin A, Cherrier MV, Gutsche I, et al (2015) Structure and primase-mediated activation of a bacterial dodecameric replicative helicase. *Nucleic Acids Res* gkv792–13. doi: 10.1093/nar/gkv792
- Bertini I, Luchinat C, Parigi G, et al (2011) Solid-state NMR of proteins sedimented by ultracentrifugation. *Proceedings of the National Academy of Sciences* 108:10396–10399. doi: 10.1073/pnas.1103854108
- Böckmann A, Gardiennet C, Verel R, et al (2009) Characterization of different water pools in solid-state NMR protein samples. *J Biomol NMR* 45:319–327. doi: 10.1007/s10858-009-9374-3
- Corn JE, Berger JM (2006) Regulation of bacterial priming and daughter strand synthesis through helicase-primase interactions. *Nucleic Acids Res* 34:4082–4088. doi: 10.1093/nar/gkl363
- Gardiennet C, Schütz AK, Hunkeler A, et al (2012) A sedimented sample of a 59 kDa dodecameric helicase yields high-resolution solid-state NMR spectra. *Angew Chem Int Ed Engl* 51:7855–7858. doi: 10.1002/anie.201200779
- Kashav T, Nitharwal R, Abdulrehman SA, et al (2009) Three-dimensional structure of N-terminal domain of DnaB helicase and helicase-primase interactions in *Helicobacter pylori*. *PLoS ONE* 4:e7515. doi: 10.1371/journal.pone.0007515
- Mainz A, Jehle S, van Rossum BJ, et al (2009) Large Protein Complexes with Extreme

- Rotational Correlation Times Investigated in Solution by Magic-Angle-Spinning NMR Spectroscopy. *J Am Chem Soc* 131:15968–15969. doi: 10.1021/ja904733v
- Oakley AJ, Loscha KV, Schaeffer PM, et al (2005) Crystal and solution structures of the helicase-binding domain of *Escherichia coli* primase. *J Biol Chem* 280:11495–11504. doi: 10.1074/jbc.M412645200
- Parsonnet J (1995) Bacterial infection as a cause of cancer. *Environ Health Perspect* 103 Suppl 8:263–268.
- Scholz I, Huber M, Manolikas T, et al (2008) MIRROR recoupling and its application to spin diffusion under fast magic-angle spinning. *Chemical Physics Letters* 460:278–283. doi: 10.1016/j.cplett.2008.05.058
- Stelter M, Gutsche I, Kapp U, et al (2012) Architecture of a dodecameric bacterial replicative helicase. *Structure* 20:554–564. doi: 10.1016/j.str.2012.01.020
- Stevens TJ, Fogh RH, Boucher W, et al (2011) A software framework for analysing solid-state MAS NMR data. *J Biomol NMR* 51:437–447. doi: 10.1007/s10858-011-9569-2
- Studier FW (2005) Protein production by auto-induction in high density shaking cultures. *Protein Expression and Purification* 41:207–234.
- Su X-C, Schaeffer PM, Loscha KV, et al (2006) Monomeric solution structure of the helicase-binding domain of *Escherichia coli* DnaG primase. *FEBS J* 273:4997–5009. doi: 10.1111/j.1742-4658.2006.05495.x
- Syson K, Thirlway J, Hounslow AM, et al (2005) Solution structure of the helicase-interaction domain of the primase DnaG: a model for helicase activation. *Structure/Folding and Design* 13:609–616. doi: 10.1016/j.str.2005.01.022
- Takegoshi K, Nakamura S, Terao T (2001) dipolar-assisted rotational resonance in magic-angle spinning NMR. *Chemical Physics Letters* 344:631–637. doi: 10.1016/S0009-2614(01)00791-6
- Wiegand T, Gardiennet C, Ravotti F, et al (2015) Solid-state NMR sequential assignments of the N-terminal domain of HpDnaB helicase. *Biomol NMR Assign* 1–11. doi: 10.1007/s12104-015-9629-8
- Williamson MP (2013) Using chemical shift perturbation to characterise ligand binding. *Progress in Nuclear Magnetic Resonance Spectroscopy* 73:1–16. doi: 10.1016/j.pnmrs.2013.02.001

Supporting Information

Solid-state NMR chemical-shift perturbations indicate domain reorientation of the DnaG primase in the primosome of *Helicobacter pylori*

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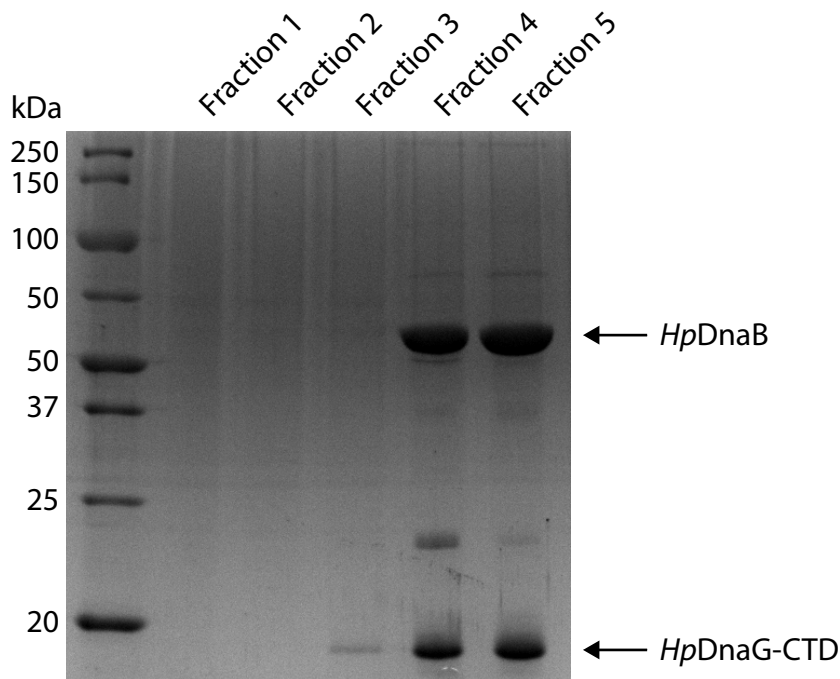
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A**B**

	F1	F2	F3	F4	F5
A280	0.47	0.54	0.83	5.54	8.96
[C] (mg/mL)	1.18	1.35	2.08	13.84	22.41
Total amount of protein (mg)	0.01	0.01	0.02	0.14	0.45

Figure S1: A: Coomassie-blue stained SDS-PAGE analysis of an analytical run of *HpDnaB/HpDnaG-CTD* co-sedimentation; the lanes correspond to the different fractions in the tube after ultracentrifugation at 200000 g overnight going from 1(top) to 5 (bottom of the tube). Fractions 1-4 were each 10 μ l and the pellet (fraction 5) was resuspended in 20 μ l Tris-HCl buffer at pH 8.0 for analysis. 5 μ l of each fraction were loaded on the Coomassie-blue stained gel. The initial total protein concentration of the sample (*HpDnaB+HpDnaG-CTD*) was 25 mg/ml. In B is shown the absorption measured at 280 nm of the different fractions, and the resulting concentrations. The total amount of protein in each fraction is calculated taking into account the total volume of each fraction. The 0.45 mg found in fraction 5 corresponds to 70 % of the total sample. In order to achieve even higher efficiency,

the protein solution used for sedimentation of the NMR sample was concentrated to 42 mg/ml (see Methods section in main text). Some DnaG-CTD is found in fraction 3 as it was added in excess to the theoretical 2:1 ratio of *HpDnaB*: *HpDnaG-CTD*.

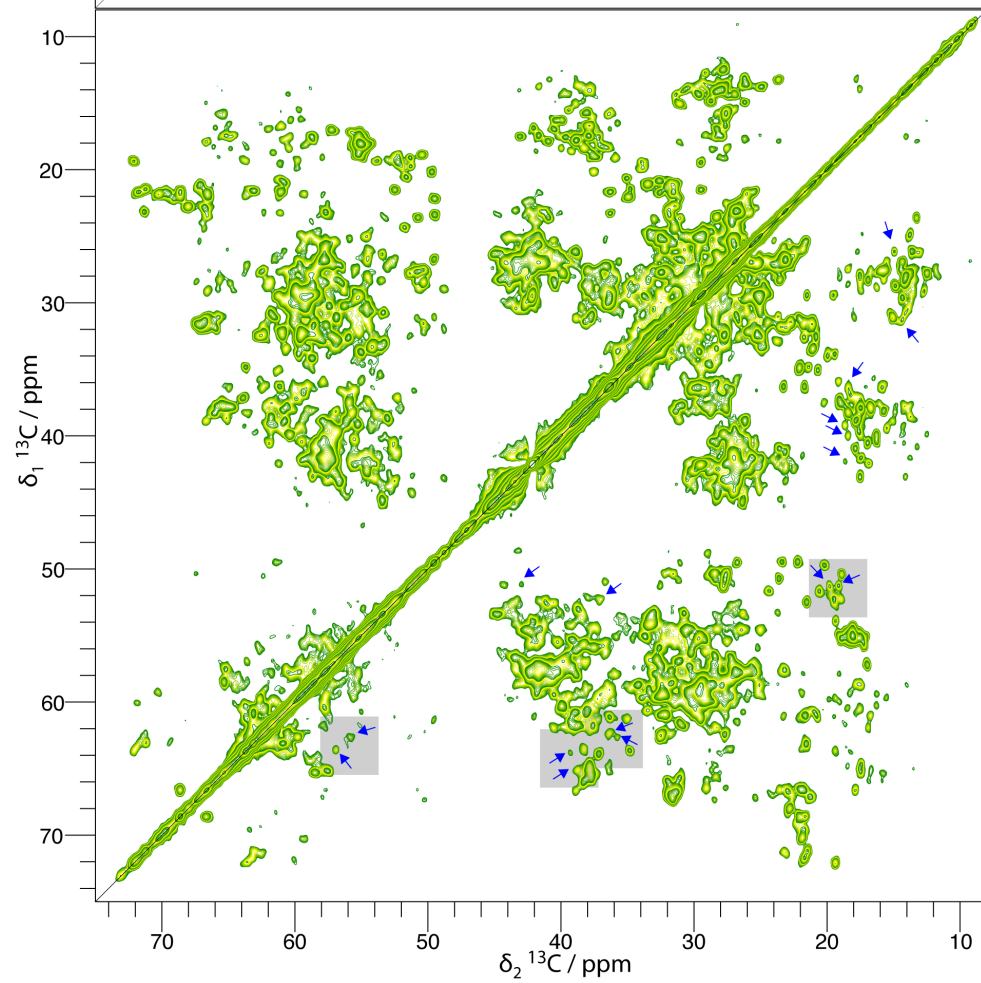
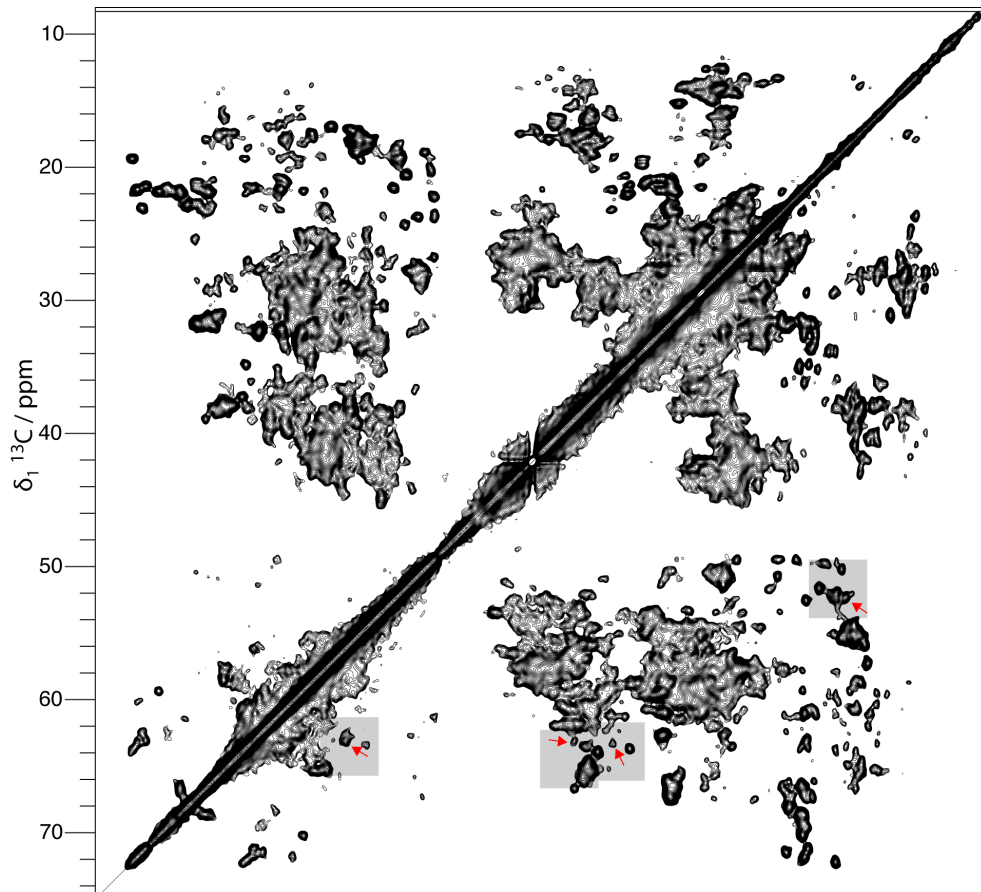


Figure S2: Aliphatic region of DARR spectra of *HpDnaB* (black, top) and *HpDnaB/HpDnaG-CTD* (green, bottom) recorded with 20 ms mixing time. Red arrows indicate signals only observed in the *HpDnaB*, and blue arrows indicate signals only observed in the protein complex.

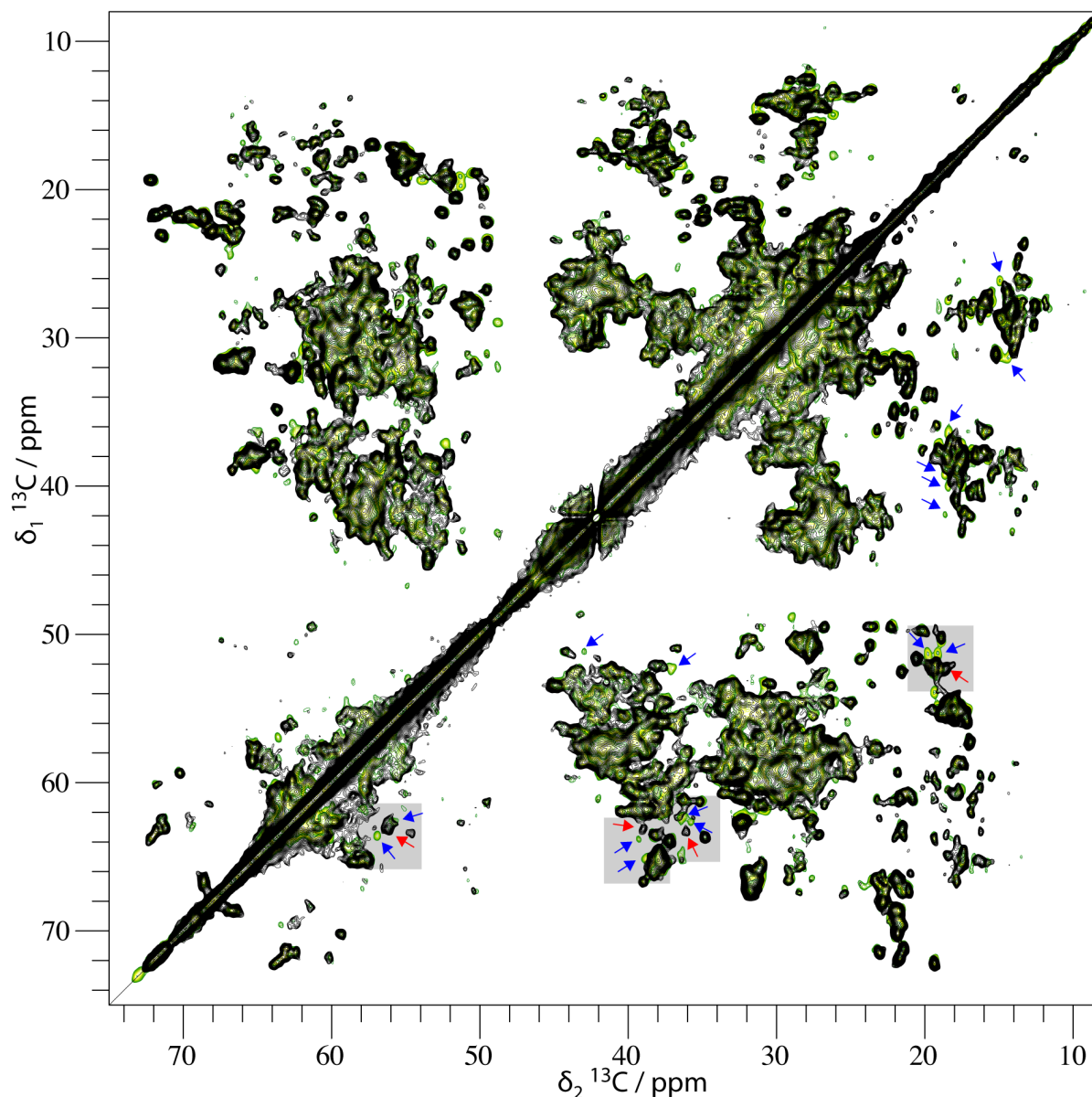


Figure S3: Overlay of the full aliphatic region of DARR spectra recorded with 20 ms mixing time. In green the spectrum of the *HpDnaB:HpDnaG-CTD* complex, and in black the spectrum of the *HpDnaB* protein. Red arrows indicate signals only observed in the *HpDnaB*, and blue arrows indicate signals only observed in the protein complex.

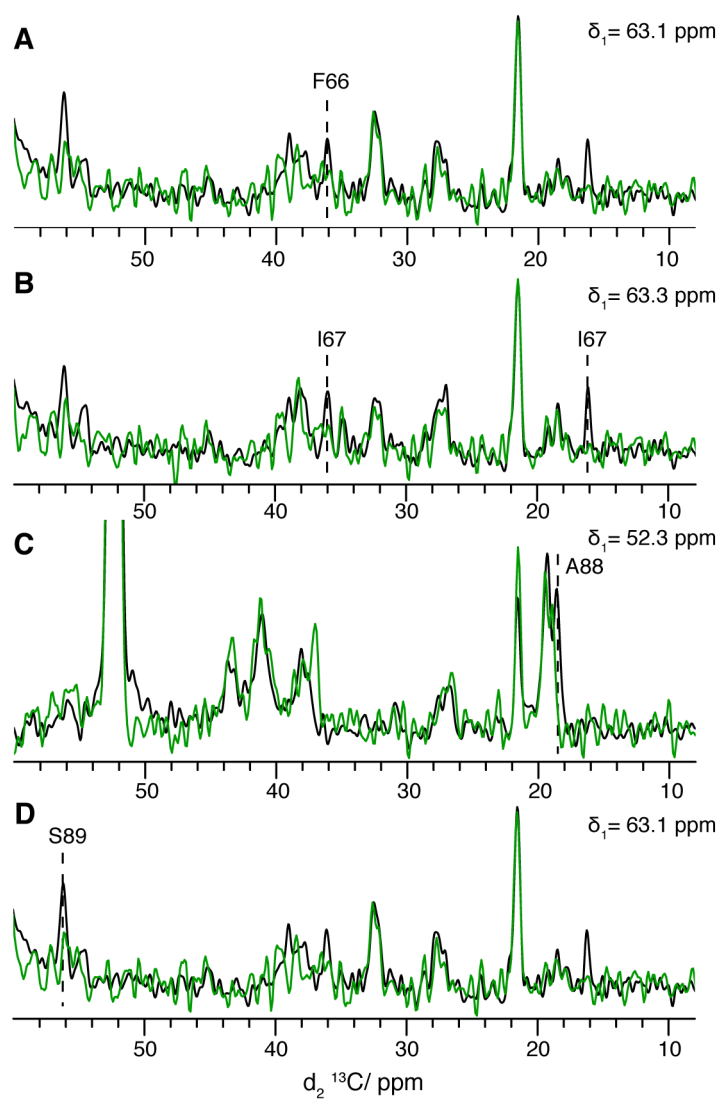


Figure S4: 1D traces corresponding to signals of extracts shown in Figure 2.

```

# Identity:      166/508 (32.7%)
# Similarity:   253/508 (49.8%)
# Gaps:         74/508 (14.6%)
# Score: 685.5

1  MDHLKHLQQL-----QNI--ERIVLSGIVLANHKIEEVHVSVLEPSDFY  HpDnaB
   :.:|      |:|  |:||...:|...:.....:|.|.|||
1  -----MSELFSERIPPQSIEAEQAVLGAFLDPAALVPASEILIPEDFY  BstDnaB

42  YPPNGLFFFEIALKLHEEDCPIDENFIRQKMPKDKQIKEEDLVAIF-----  HpDnaB
   .....|...|:.....|:|...:.....:|:|...|:..
45  RAAHQKIFHAMLRVADRGEVLDLVTVAELAASEQLEEIGGVSYLSELAD  BstDnaB

87  AASPIDNIEAYVEEIKNASIKRKLFLGLANTIREQALESAQKSSDILGAVE  HpDnaB
   |.....|:|.|...:..|:|:|...|.:|:.....:.....|...:
95  AVPTAANVEYYARIVEEKSVLRLRRLIRTATSIAQDGYTREDEIDVLLDEAD  BstDnaB

137 REVYALLNGSTIEGFRNIKEVLESAMDLITE-NQRKGSLEVTGIPTGFVQ  HpDnaB
   |:.....:.....|:|:|:|:|...|.|. .:|.|  |:|:|:|:|:|.
145 RKIMEVSQRKHSQAFKNIKDILVQTYDNIEMLHNRDG--EITGIPTGFTE  BstDnaB

186 LDNYTSGFNKGS�VIIGARPSMGKTSLMMNMVLS-ALNDDRGVAVFSLEM  HpDnaB
   ||.||||:..|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
193 LDRMTSGFQRSDLIIVAARPSVGKTAALNIAQNVATKTNENVAIFSLEM  BstDnaB

235 SAEQLALRALSDLTINMHDLESGRLDDQWENLAKCFDHLQKLLFFYD  HpDnaB
   |:|:|:|:|:|.|...:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
243 SAQQLVMRMLCAEGNINAQNLRTGKLTPEWGWKLTAMGSLSNAGIYIDD  BstDnaB

285 KSYVRIEQIRLQLRKLKSQHKELGIAFIDYLQMSGKATKE-RHEQIAE  HpDnaB
   ...:|:..|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
293 TPSIRVSDIRAKCRRLK-QESGLGMIVIDYDLQLIQGSGRSKENRQEVSE  BstDnaB

334 ISRELKTLARELEIPIIALVQLNRSLENRDDKRPILSDIKDSGGIEQDAD  HpDnaB
   |||.|||.|||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
342 ISRSLKALARELEVPIALSQLSRSVEQRQDKRPMSDIRESGSIEQDAD  BstDnaB

384 IVLFLYRGYIYQMRAEDNKIDKLKKEGKIEEAQELYLKVNEERRIHKQNG  HpDnaB
   ||.||||...|...:|:..|
392 IVAFLYRDDYYNKDSSENKNI-----  BstDnaB

434 SIEEAEIIVAKNRNGATGTVYTRFNAPFTRYEDMP---IDSHLEEGQETK  HpDnaB
   .|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
412 ----IEIIIAKQRNGPVGTVQLAFIKEYNKFVNLERRFDEAQIPPGA---  BstDnaB

481 VDYDIVTT  HpDnaB

455 -----  BstDnaB

```

Figure S5: Sequence alignment of *HpDnaB* and *BstDnaB* using EMBOSS (version 6.6.0) (Rice et al. 2000). Legend: identity (|), similarity with a small positive score (.) and similarity with a score larger 1.0 (:).

```

# Identity:      24/192 (12.5%)
# Similarity:   54/192 (28.1%)
# Gaps:         96/192 (50.0%)
# Score: 27.5

415 VSFQPFYPKTEKPNRPQRFQFAHVSSAPSLFLEKLVIRYLLEDRSLLDLAV HpDnaG (CTD)
      ...|:.....|:|:.....| |:|:
455 -----KLLPAFQNAERLLLAHMMRSR---DVAL BstDnaG (CTD)

465 -----GYIHSGVFLHKKQEFDALCQEKLDLDPKLVVAL HpDnaG (CTD)
      .||:: |.....|.|. .||
480 VVQERIGGRFNIEEHRALAAIYA--FYEEGHEADP-----GAL BstDnaG (CTD)

496 LLDANLPLKKGFEKELRLLILR--YFERQLKEIPKSSLPFSEKMICLKK HpDnaG (CTD)
      :  ::|:.....|:|:|:.. ..|:|:|:..... .|.:
517 I--SRIPGELQPLASELSLLLIADVSEQELEDYIRH-----VLNR BstDnaG (CTD)

544 ARQAIMKCLKQGELVAI----- HpDnaG (CTD)
      .....|:|:|.|.
556 PKWMLMLKVKEQEKTEAERRKDFLTAARIAKEMMKMLSSS BstDnaG (CTD)

```

Figure S6: Sequence alignment of *HpDnaG*(CTD) and *BstDnaG*(CTD) using EMBOSS (version 6.6.0) (Rice et al. 2000). Legend: identity (|), similarity with a small positive score (.) and similarity with a score larger 1.0 (:).

References

Rice P, Longden I, Bleasby A (2000) EMBOSS: the European molecular biology open software suite.

Table S1: Chemical shift differences between the signals disappearing in the *HpDnaB* sediment and their nearby appearing signals in the *HpDnaB/HpDnaG*-CTD co-sediment as shown in Figure 2.

Residue	Atom Name	δ DnaB (ppm)	δ DnaB/DnaG (ppm)	$\Delta\delta$ (ppm)
F66	C α	63.17	63.81	0.63
			65.12	1.95
F66	C β	39.00	39.30	0.30
			38.74	1.98
I67	C α	63.24	62.63	-0.61
			62.36	-0.88
I67	C β	36.12	35.72	-0.40
			36.39	0.27
A88	C α	52.24	51.27	-0.97
			51.28	-0.96
A88	C β	18.57	19.14	0.57
			19.80	1.23
S89	C α	56.17	56.91	0.74
			55.84	-0.33
S89	C β	63.16	63.61	0.45
			62.66	-0.50

Table S2 : Overview about experimental parameters of the performed solid-state NMR experiments.

Experiment/ Sample	DARR/ NTD <i>HpDnaB</i>	DARR/ CTD <i>HpDnaB</i>	DARR/ <i>HpDnaB</i>	DARR/ <i>HpDnaB:HpDna</i> G-CTD
MAS frequency/ kHz	17.0	17.5	17.0	17.5
Field/ T	20.0	18.8	20.0	18.8
Transfer I	HC-CP	HC-CP	HC-CP	HC-CP
¹H field/ kHz	59.4	65.0	58.6	64.0
X field/ kHz	40.9	50.0	41.6	50.0
Shape	Tangent ¹ H	Tangent ¹ H	Tangent ¹ H	Tangent ¹ H
¹³C carrier/ ppm	95	59	103	55
Time/ ms	0.5	0.7	0.5	0.7
Transfer II	DARR	DARR	DARR	DARR
¹H field/ kHz	17.0	17.5	17.0	17.5
Carrier/ ppm	95	59	103	55
Time/ ms	20	20	20	20
t1 increments	2000	1876	2560	2000
Sweep width (t₁)/ kHz	100	94	100	94
Acquisition time (t₁)/ ms	10	10	13	11
t2 increments	2988	2802	3072	2988
Sweep width (t₂)/ kHz	100	94	100	94
Acquisition time (t₂)/ ms	14.9	14.9	15.4	16.0
¹H Spinal64 decoupling power/ kHz	89	90	90	90
Interscan delay/ s	2.1	2.1	2.5	2.2
Number of scans	12	12	16	36
Measurement time/ h	14	13	28	45