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^1H , ^{13}C , and ^{15}N backbone resonance assignments of the full-length 40 kDa *S. acidocaldarius* Y-family DNA polymerase, *dinB* homolog

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Abstract The *dinB* homolog (Dbh) is a member of the Y-family of translesion DNA polymerases, which are specialized to accurately replicate DNA across from a wide variety of lesions in living cells. Lesioned bases block the progression of high-fidelity polymerases and cause detrimental replication fork stalling; Y-family polymerases can bypass these lesions. The active site of the translesion synthesis polymerase is more open than that of a replicative polymerase; consequently Dbh polymerizes with low fidelity. Bypass polymerases also have low processivity. Short extension past the lesion allows the high-fidelity polymerase to switch back onto the site of replication. Dbh and the other Y-family polymerases have been used as structural models to investigate the mechanisms of DNA polymerization and lesion bypass. Many high-resolution crystal structures of Y-family polymerases have been reported. NMR dynamics studies can complement these structures by providing a measure of protein motions. Here we report the ^{15}N , ^1H , and ^{13}C backbone resonance assignments at two temperatures (35 and 50 °C) for *Sulfolobus acidocaldarius* Dbh polymerase. Backbone resonance assignments have been obtained for 86 % of the residues. The polymerase active site is assigned as well as the majority of residues in each of the four domains.

Keywords Y-family polymerase · Bypass polymerase · Translesion synthesis · *Sulfolobus acidocaldarius* · Dbh (*dinB* homolog)

Biological context

Dbh (*dinB* homolog) is a 354 amino acid DNA polymerase from the thermophilic archaeon *Sulfolobus acidocaldarius* (Boudsocq et al. 2004). Dbh belongs to the Y-family of translesion (TLS) DNA polymerases that can accurately bypass a variety of damaged DNA templates (Yang and Woodgate 2007). Members of the Y-family include Dpo4 from *Sulfolobus solfataricus*, pol IV [*E. coli*, (Wagner et al. 1999)] mammalian pol κ (Ogi et al. 1999; Ohashi et al. 2000; Gerlach et al. 2001), pol η (Johnson et al. 1999a, b), pol ι (McDonald et al. 1999; Tissier et al. 2000), and Rev1 (Lin et al. 1999; Wiltout and Walker 2011). In particular, Dbh shares 54 % sequence identity to Dpo4. Members of the Y-family share canonical polymerase architecture consisting of a catalytic core composed of palm, fingers, and thumb domains, and have an additional unique C-terminal domain termed the “wrist”, “polymerase-associated domain” (PAD), or “little finger” (LF) domain. Notably, the LF domain is tethered to the catalytic core by a flexible linker and has been found to occupy a variety of conformations (Pata 2010). Y-family polymerases also have a significantly larger active site than that of replicative polymerases, which allows the accommodation of bulky DNA adducts, but increases the error-rate extending primers across from undamaged DNA (Sale et al. 2012). Hence, Y-family polymerases are usually highly regulated in eukaryotes, most often through monoubiquitination of PCNA, to prevent error-prone replication of cellular DNA (Yang et al. 2013). TLS polymerases can allow cancer cells

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to tolerate DNA damage caused by chemotherapeutics; for example, pol η can bypass cisplatin adducts (Alt et al. 2007). In this way, TLS polymerases that are upregulated can confer resistance to chemotherapy. Furthermore, loss of translesion synthesis can generate chromosomal abnormalities that can contribute to a cancerous phenotype (reviewed in Lange et al. 2011).

Different Y-family polymerases generate characteristic mutations; in particular, Dbh is highly prone to generating single-base deletions resulting in -1 frameshift mutations in vitro (Potapova et al. 2002). The first crystal structures of the apo form of Dbh were reported in 2001 (Silvian et al. 2001; Zhou et al. 2001), followed by preinsertion binary (Dbh, primer/template DNA), insertion ternary (Dbh, primer/template DNA, incoming dNTP) and postinsertion binary (Dbh, primer $+1$ /template DNA) crystal structures (Wilson and Pata 2008). The binary and ternary structures indicated that Dbh generates single-base deletions by a template-slippage mechanism (Wilson and Pata 2008). Dpo4 also generates single-base deletions at high rates (Kokoska et al. 2002), albeit with a somewhat altered mutational spectrum from Dbh. Intriguingly, the mutational and lesion bypass properties of Dbh can be made to resemble those of Dpo4 by generating chimeras of Dbh/Dpo4 containing the catalytic core of Dbh with the LF domain of Dpo4 (Boudsocq et al. 2004), Dbh core/LF with the linker region of Dpo4 (residues 231–245) (Wilson et al. 2013), or even merely three residues of the Dpo4 linker (R242, K243, S244) substituted for residues (K243, I244, P245) of Dbh (Mukherjee et al. 2014). The crystal structures of the Dbh chimeras (Boudsocq et al. 2004; Wilson et al. 2013; Mukherjee et al. 2014) reveal that the orientation of the LF domain with respect to the core resembles that of the LF in the Dpo4 crystal structure (Ling et al. 2001). This observation raises the question of how the position, flexibility, and dynamics of the LF domain affects substrate alignment in the active site of Dbh. Here we report the ^{15}N , ^1H , and ^{13}C backbone assignments of Dbh, which will be used to interpret residue specific T_1 , T_2 , and ^{15}N – ^1H NOE relaxation data, and CPMG relaxation dispersion data. The relaxation data will provide valuable information on the mechanism of polymerization of DNA and generation of single-base deletion errors by Dbh.

Methods and experiments

Protein expression and purification

The Dbh gene was incorporated into the vector pKKT7-H (a derivative of pKK233, Promega) containing an N-terminal His₆ tag (MHHHHHHLVPRGM). Quick-change mutagenesis (Stratagene) was used to change Cys31 to Ser (hereafter referred to as C31S-Dbh) to eliminate potential

formation of disulfide bonds. Transfected *E. coli* BL21 cells were grown in 1L Neidhart's minimal media (Neidhardt et al. 1974) at 37 °C containing 1 g ^{15}N ammonium chloride (^{15}N -labeled samples), or 1 g ^{15}N ammonium chloride, 3 g ^{13}C glucose, and 80 % D_2O (^2H , ^{15}N , ^{13}C -labeled samples) to ~ 1.0 OD; expression was induced by the addition of 1 mM IPTG. Protein was expressed for 5 h; subsequently, the cells were harvested by centrifugation and frozen at -80 °C. Dbh or C31S-Dbh were purified from cell lysate by Ni-NTA affinity chromatography under native conditions, and then dialyzed into buffer (20 mM HEPES, 100 mM NaCl, 50 μM EDTA, 50 μM NaN_3 , pH 7.5) at 4 °C, then one change of buffer without EDTA (20 mM HEPES, 100 mM NaCl, 50 μM NaN_3 , pH 7.5). To prepare the NMR samples, Dbh or C31S-Dbh protein was concentrated to at least 0.5 mM, and transferred into a Shigemi NMR tube. D_2O was added to the sample for a final concentration of 10 % v/v. Since polymerase enzymes use aspartic acid side chains to coordinate Mg^{2+} at the active site, we were particularly interested in assigning the Asp groups. ^{15}N -HSQC spectra of selectively ^{15}N labeled (Asp) and un-labeled (Asn, Arg, Gly, Lys, His, Met, Ser) samples were used to confirm amino acid identity within the sequence. Selectively labeled ^{15}N -Asp C31S-Dbh was prepared using the *E. coli* auxotroph strain EA1, which is unable to convert Asp to Asn (Muchmore et al. 1989), and by supplementing 1L of Neidhart's media with 100 mg ^{15}N -Asp. Selectively unlabeled Asn, Arg, Gly, Lys, His, Met, Ser C31S-Dbh samples were prepared using 1 g ^{15}N ammonium chloride per L of Neidhart's minimal media, BL21 cells, and by supplementing with 0.5 g of each ^{14}N amino acid separately. Since the HNCO experiment on C31S-Dbh was by far the most sensitive, an HNCO spectrum of a selective $^{13}\text{C}'$ -Leu, fully ^{15}N enriched sample was used to confirm amide resonances preceded by leucine residues. $^{13}\text{C}'$ -Leu was incorporated by expressing the protein in BL21 cells and adding 150 mg of $^{13}\text{C}'$ -labeled Leu to 1 L of Neidhart's medium containing 1 g ^{15}N ammonium chloride.

NMR experiments

NMR data were acquired at 35 and 50 °C on a Varian INOVA 800 MHz NMR spectrometer equipped with a 5-mm triple resonance XYZ-gradient probe. The chemical shifts were referenced using 2,2-dimethyl-2-silapentane-sulfonic acid (DSS). All spectra were processed using NMRPipe/NMRDraw (Delaglio et al. 1995) and analyzed using CcpNmr Analysis (Vranken et al. 2005). A set of 3D triple resonance experiments, including HNCO, HN(CA)CO, HN(CO)CA, and HNCACB were carried out using TROSY (Pervushin et al. 1997) for the sequential backbone resonance assignment (Kay 1997). In addition,

visible in the ^{15}N -HSQC spectrum and the 3D spectra; we were unable to find connectivities for these resonances. In addition, four resonances in the ^{15}N -HSQC did not have visible corresponding resonances in the 3D data or in the ^{15}N -edited NOESY spectrum. Figure 2 presents the C31S-Dbh amino acid sequence with assigned residues indicated. Residues 36–38 and the C-terminus (residues 345–354) were disordered in the crystal structures of Dbh [PDB entries 1K1S/1K1Q (Silvian et al. 2001) and 3BQ0 (Wilson and Pata 2008)]; unfortunately, we were unable to assign the backbone resonances of the regions that were not visible in the crystal structures. If we omit the 13 residues

ible in the crystal structures. If we omit the 13 residues

[illegible]

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1  MIVIFVDFDYFFAQVEEVLPQYKKGKPLVVSVMYSGRTKTSGAVATANYEA
51 RKLGVKAGMPIIKAMQIAPSAIYVPMRKPIIYEAFSNRIMNLLNKHADKIE
101 VASIDEAYLDVTNKVEGNFENGIELARKIKQEILEKEKITVTVGVPNKI
151 LAKIIADKSKPNGLGVIRPTEVQDFLNELDIDEIPCIGSVLARRLNELGI
201 QKLRDILSKNYNELEKITGKAKALYLLKLAQNKYSEPVENKSKIPHGRYL
251 TLPYNTRDVKVILPYLKKAINEAYNKVNGIPMRITVIAIMEDLDILSKGK
301 KFKHGISIDNAYKVAEDLLRELLVRDKRRNVRRIGVKLDNIIINKTNLSD
351 FFDI

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Fig. 2 C31S-Dbh amino acid sequence with assignments indicated. Grey shade is used to indicate residues with at least one backbone atom assigned. Black background with white lettering indicates residues where only the $^{13}\text{C}'$ or $^{13}\text{C}^{\alpha}$ were found but not the NH. White background represents residues that have not been assigned. In

blue lettering, amino acids 36–38 and 345–354 are too disordered in crystal structures to be detected [PDB entries 1K1S/1K1Q (Silvian et al. 2001) and 3BQ0 (Wilson and Pata 2008)]. These regions are also not detected by NMR

(36–38; 345–354) that are disordered in the crystal structures of Dbh, we can account for 89 % of residues with at least one assigned backbone resonance. Likewise, most of the linker region (residues 232–245) between the LF and thumb domains was not assigned. The linker and disordered regions from the crystal structure are flexible and solvent exposed. Hence, the signals may be missing due to intrinsic exchange with the solvent. In conclusion, for portions of C31S-Dbh where we would expect rigid structure to enable us to detect NMR signals, we have assigned 94 % of the protein.

A small part of the structure appears to have an alternate conformation; several residues have two corresponding peaks in the ^{15}N -HSQC spectrum and 3D data (97D, 98K, 100E, 101V, 102A, 103S, 108Y, 109L, and 110D), consistent with slow chemical exchange. These residues are located in the β -sheet structure of the active site palm domain, surrounding the metal ion coordinating residues 105D and 106E.

The mutation of residue 31 from Cys to Ser does not appear to affect the structure. For example, the crystal structures of apo C31S-Dbh [PDB entries 1K1S/1K1Q (Silvian et al. 2001)] superimpose very well with ligand-bound forms of WT Dbh [PDB entries 3BQ0, 3BQ1, and 3BQ2 (Wilson and Pata 2008)]. In addition, this mutation did not significantly affect the NMR spectrum, as the ^{15}N -HSQC of WT Dbh and C31S-Dbh overlay extremely well (data not shown). Only 11 peaks (9D, 10Y, 12F, 31S, 32V, 45T, 56K, 64A, 77R, 140T, and 301K) were observed to have shifted by any appreciable amount (>0.05 ppm for ^1H or >0.2 ppm for ^{15}N) upon mutation. Unsurprisingly, the peaks for residue 31 and adjacent 32V are shifted, and all but two (140T, 301K) of the remaining shifted peaks are located in the same domain (finger) as residue 31.

The backbone resonance assignments of Dpo4 catalytic core (Ma et al. 2010) and LF domain (Ma et al. 2011) at 50 °C have been published. Given the homology between the two proteins (54 % sequence identity) and similar tertiary structure, some of the resonances of the two proteins

would be expected to have similar chemical shifts. It should be noted that all of the assignments of C31S-Dbh were completed independently using only our own data; the Dpo4 assignments were compared to those of C31S-Dbh after we completed our assignments. The mutually assigned amide peak positions of identical residues do not correlate very well, with only 62 of 135 (46 %) available ^1H shifts within 0.2 ppm and 54 of 135 (40 %) available ^{15}N shifts within 0.8 ppm. However, the nearest neighbors of a particular residue can have a significant effect on the amide chemical shift, even if the residues are identical. For instance, a neighboring isoleucine residue on the C-terminal side of an amide could influence the ^1H shift downfield by ~ 0.2 ppm (Schwarzinger et al. 2001) and the ^{15}N shift downfield by almost 5 ppm (Braun et al. 1994; Schwarzinger et al. 2001). Eliminating identical residues from the comparison that do not also have identical neighbors, the correlation between the two sets of shifts is improved: 30 of 54 (56 %) ^1H shifts within 0.2 ppm and (57 %) 31 of 54 ^{15}N shifts within 0.8 ppm. All but one of the ^1H shifts and one of the ^{15}N shifts in the preceding comparison are found in the polymerase core: 29 of 45 (64 %) ^1H shifts within 0.2 ppm and 30 of 45 (67 %) ^{15}N shifts within 0.8 ppm. This is not surprising, since the polymerase core between the two proteins shares 59 % sequence identity, while the LF domains of the two proteins only have 41 % sequence identity. Even though the assignments of Dpo4 were completed on the polymerase core and the LF as separate constructs, the core and LF domains appear to fold independently into roughly the same native structure as found in the full-length protein based on chemical shifts.

The chemical shifts of C31S-Dbh polymerase at 308K and 323K (35 and 50 °C) have been deposited in the BioMagResBank database under accession number 26564 [<http://www.bmrb.wisc.edu>, (Ulrich et al. 2008)]. The backbone assignments will be used for relaxation dynamics studies of Dbh, and the published crystal structures of Dbh will be used to interpret NMR relaxation data through molecular modeling and molecular dynamics simulations.

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