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^1H , ^{13}C , ^{15}N backbone resonance assignments of the apo and holo forms of the ABC transporter solute binding protein PiuA from *Streptococcus pneumoniae*

Katherine A. Edmonds¹, Yifan Zhang^{1,2,3}, Daniel J. Raines⁴, Anne-K. Duhme-Klair⁴, David P. Giedroc^{1,3}

¹Department of Chemistry, Indiana University, Bloomington, Indiana, USA

²Graduate Program in Biochemistry, Indiana University, Bloomington, Indiana, USA

³Department of Molecular and Cellular Biochemistry, Indiana University, Bloomington, Indiana, USA

⁴Department of Chemistry, University of York, Heslington, York YO10 5DD, UK

Abstract

Streptococcus pneumoniae is a Gram-positive human pathogen that causes millions of infections worldwide with an increasing occurrence of antibiotic resistance. Iron acquisition is essential for its survival and virulence, especially under host-imposed nutritional immunity. *S. pneumoniae* expresses several ATP-binding cassette (ABC) transporters to facilitate acquisition under iron limitation, including PitABCD, PiaABCD, and PiuBCDA. The substrate specificity of PiuBCDA is not fully established. Herein, we report the backbone ^1H , ^{13}C and ^{15}N resonance assignments of the 31 kDa soluble, extracellular domain of the substrate binding protein PiuA in the apo form and in complex with Ga(III) and the catechol siderophore-mimic 4-LICAM. These studies provide valuable information for further functional studies of interactions with other proteins, metals, and small molecules.

Keywords

Iron acquisition; ABC transporter; Pathogen; Catechol; Siderophore; *Streptococcus pneumoniae*

Biological context

Streptococcus pneumoniae is an important Gram-positive human pathogen (Lanie et al. 2007; Lynch and Zhanel 2010). Iron (Fe) is essential for its survival and pathogenicity, as it is essential for most living organisms, due to its ability to perform redox reactions and function as a cofactor for many proteins (Andreini et al. 2018). During infection, the host often limits the availability of iron in a process known as nutritional immunity (Hood and Skaar 2012). To respond to iron limitation, various *S. pneumoniae* strains express three different ABC transporters. PitABCD, PiaABCD, and PiuBCDA each facilitate iron uptake

from different sources and as different Fe(III)-chelate complexes. Transporters such as these commonly recognize iron bound to heme or other small organic molecules, including various classes of siderophores (Zhang et al. 2020). *S. pneumoniae* is not known to endogenously biosynthesize siderophores and thus must rely on Fe piracy to meet nutritional Fe demands (Cheng et al. 2013). Hence, the regulated uptake of siderophore- or other small molecule-Fe(III) complexes produced by other, often competing, bacteria in polymicrobial communities or by the host is foundational to Fe piracy in *S. pneumoniae*.

PiuA is the solute-binding protein (SBP) of the PiuABC transporter. As an extracellular membrane-bound lipoprotein, its role is to recognize and bind a specific Fe(III)-complex and transfer it to the transmembrane subunit of the ABC complex embedded in the cytoplasmic membrane for import into the cell. Like most SBPs known to be involved in iron transport in bacteria, PiuA belongs to class III (cluster A), consisting of two α/β lobes connected by a rigid helical linker, with the ligand binding site in a cleft between the lobes (Delepleaire 2019). Members of this class typically display very little structural rearrangement upon ligand binding (de Boer et al. 2019). This feature leaves some questions as to how substrate binding promotes transport.

Previous nuclear magnetic resonance studies of a similar protein, the *E. coli* FepB, an SBP involved in import of Fe(III) in complex with the siderophore enterobactin, shows that Ga(III) is a suitable diamagnetic substitute for the paramagnetic Fe(III), enabling detailed structural and dynamic analysis without extensive line-broadening (Chu et al. 2014). Similarity to the SBP CeuE of *Campylobacter jejuni* predicts that PiuA might also bind coordinately unsaturated iron complexes of tetradentate siderophores (Raines et al. 2016) and their mimics, for example 4-LICAM, a *bis*-catecholate which contains a four-atom linker between two simple catecholamide units (Raines et al. 2013; Wilde et al. 2017).

Herein, we report the resonance assignments of the backbone ^1H , ^{13}C and ^{15}N atoms of the 31 kDa soluble, extracellular domain of *S. pneumoniae* PiuA in the apo form and in complex with Ga(III) and 4-LICAM, which provide valuable information for further studies of the dynamics and interactions with other proteins, metals, and small molecules.

Methods and experiments

Sample preparation

The region of the gene encoding the soluble, extracellular domain of PiuA, SPD_1652 residues 37–321 was PCR-amplified from the genomic DNA of *S. pneumoniae* D39 strain (Lanie et al. 2007), and inserted into the pSUMO expression vector (Peroutka et al. 2011). Uniformly ^{15}N , ^{13}C , ^2H protein was expressed in *E. coli* BL21 (DE3) cells in M9 minimal medium containing 1 kg D_2O , as well as 1.0 g of $^{15}\text{NH}_4\text{Cl}$ and 2 g $^{13}\text{C}_6,^2\text{H}$ -glucose as the sole nitrogen and carbon sources, respectively. Expression was induced by addition of 1 mM isopropyl β -D-1-thiogalactopyranoside (IPTG) at OD_{600} 0.7 and allowed to continue overnight at 18 °C. Cells were harvested by centrifugation, resuspended in buffer A (25 mM Tris pH 8.0, 500 mM NaCl, 10% glycerol, and 20 mM imidazole, pH 8.0), then lysed by sonication on ice. The crude lysate was clarified by centrifugation. 70% ammonium sulfate was added to the soluble fraction, and the precipitated protein was collected by

centrifugation, and then resuspended in buffer A. The tagged protein was isolated by Ni(II) affinity chromatography using a 5 mL HisTrap FF column (GE Healthcare Life Sciences) with a gradient from 100% buffer A to 100% buffer B (25 mM Tris pH 8.0, 500 mM NaCl, 10% glycerol, and 500 mM imidazole, pH 8.0). Eluted fractions were pooled and cleaved with SUMO protease while dialyzing overnight at room temperature into buffer A supplemented with 2 mM dithiothreitol (DTT). Uncleaved protein, protease, and the SUMO tag were removed by passage through the HisTrap FF column equilibrated in buffer A prior to further purification by size-exclusion chromatography in buffer C (25 mM Tris pH 8.0, 500 mM NaCl, 2 mM ethylenediaminetetraacetic acid [EDTA]), using a Superdex-75 column on an Akta Pure system (GE Healthcare Life Sciences).

To facilitate exchange of deuterated amides back to protons, the purified protein was incubated with 2.5 M guanidinium-HCl and 5 mM EDTA for 3 h, then dialyzed into buffer C and then into NMR buffer (25 mM MES pH 6.5, 150 mM NaCl, treated with Chelex (Biorad). ^{15}N TROSY spectra on samples labeled with only ^{15}N were used to confirm nearly complete back-exchange of the deuterated sample.

The holo sample was prepared as follows. 4-LICAM was synthesized according to a previously published procedure (Raines et al. 2013), and dissolved in neat DMSO to a concentration of 30 mM. GaCl_3 was prepared by dissolving Ga_2O_3 (Sigma Aldrich) in 0.3 M HCl to a concentration of 20 mM. Then GaCl_3 and 4-LICAM were mixed to achieve a final concentration of 15 mM 4-LICAM and 10 mM GaCl_3 . This mixture was then diluted to a final concentration of 1 mM in GaCl_3 in 50 mM HEPES, pH 7.2, 150 mM NaCl. This was used as a stock Ga(III)-4-LICAM complex and added to $\approx$ 3-fold molar excess over [PiuA] in the same buffer, incubated 4 h at RT, and buffer exchanged into NMR buffer using an Amicon ultra centrifugal filter unit (10 kDa cut-off) while removing free ligand.

NMR experiments

NMR samples for backbone assignment contained 0.8–0.9 mM protein, with 25 mM MES pH 6.5, 150 mM NaCl, and 10% v/v D_2O , with 0.3 mM 2,2-dimethyl-2-silapentanesulfonic acid (DSS) as an internal reference. NMR spectra were recorded at 35 °C on a 600 MHz Bruker Avance Neo spectrometer equipped with a cryogenic probe in the METACyt Biomolecular NMR Laboratory at Indiana University, Bloomington. Backbone chemical shifts were assigned for each state using TROSY versions of the following standard triple-resonance experiments: HNCACB, HNCOCACB, HNCA, HNCOCOA, HNCO, and HNCACO (Salzmann et al. 1999), using non-uniform sampling with Poisson gap schedules (Hyberts et al. 2010). Data were processed using NMRPipe (Delaglio et al. 1995) and istHMS (Hyberts et al. 2012), and analyzed using CARA (Keller 2004) and Sparky (Lee et al. 2015), all on NMRbox (Maciejewski et al. 2017).

Assignments and data deposition

The soluble, extracellular domain of PiuA consists of 285 residues. Nearly complete backbone assignments were obtained for the apo (ligand-free) and holo [Ga(III)-4-LICAM] forms of PiuA, as shown in the ^{15}N TROSY spectra in Figs. 1 and 2, respectively. Amide proton and nitrogen assignments are missing for R237 and H238 of the apo form, and for

R216 and R237 in the holo form. These crosspeaks are missing likely due to line-broadening caused by conformational exchange; the crystal structure of a close homolog (PDB 4JCC) from *S. pneumoniae* Canada MDR_19A shows that residues 237–238 are in the middle of a long loop, between β strands 10 and 11, with relatively high B-factors, and R216 is also in a loop between β strand 9 and helix 6 (Fig. 3a). With two missing amide ^1H and amide ^{15}N assignments and 12 prolines, 99.3% of all possible amide ^1H and ^{15}N atoms are assigned in both the apo and holo states. 99.6% of all possible $^{13}\text{C}'$, $^{13}\text{C}\alpha$, and $^{13}\text{C}\beta$ atoms are assigned in the apo form, and 100% of all possible $^{13}\text{C}'$, $^{13}\text{C}\alpha$, and $^{13}\text{C}\beta$ atoms are assigned in the holo form.

Chemical shifts were analyzed for indications of secondary structure using TALOS-N (Shen and Bax 2014). The results, shown in Fig. 3b, c, are in good agreement with the existing crystal structure of the apo form of PiuA (PDB 4JCC). As expected, the secondary structure regions for apo and holo PiuA are nearly identical. The main differences are a slightly increased likelihood of β -strand in residues 232–234 (magenta), as if Fe(III) coordination by H238 induces a β -hairpin in the relatively flexible loop formed by residues 231–243, and perhaps two residues unwinding from the N-terminus of α 3, as R126 is likely to interact electrostatically with the siderophore (Wilde et al. 2017). F217 also may open up from α 6 upon ligand binding. Although conformational changes are limited, the spectral changes are more than adequate to use NMR to monitor changes in metalation state. All chemical shifts have been deposited in the BioMagResBank (<http://www.bmrb.wisc.edu>) under accession number 28056 (apo) and 28057 (holo).

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Fig. 1.
 2D ^1H - ^{15}N TROSY spectrum of apo PiuA, annotated with the backbone assignments. The assignments are labeled with the one-letter amino acid code and residue number of full-length PiuA. The TROSY spectrum was collected at 600 MHz ^1H frequency. The crowded region at the center is shown in expanded view in the lower right, and the tryptophan sidechain region is shown in the inset in the upper left

Fig. 2.
 2D ^1H - ^{15}N TROSY spectrum of PiuA in complex with Ga(III) and 4-LICAM, annotated with the backbone assignments. The assignments are labeled with the one-letter amino acid code and residue number of full-length PiuA. The TROSY spectrum was collected at 600 MHz ^1H frequency. The crowded region at the center is shown in expanded view in the lower right, and the tryptophan sidechain region is shown in the inset in the upper left

Fig. 3.

a Sequence alignment of *S. pneumoniae* D39 PiuA studied here by NMR with the previously solved crystal structure (PDB 4JCC) of *S. pneumoniae* Canada MDR_19A in the apo form. The four residues that are different are highlighted in a red background. Secondary structures from the crystal structure are shown above, prepared with ESript (Robert and Gouet 2014). **b** Chemical-shift derived secondary structure assignments of apo and holo PiuA are painted onto the ribbon representation of the crystal structure. Where assignments are identical for apo and holo forms, helices are painted blue and beta strands

red. Magenta indicates residues in beta strand conformation only in the holo form, while cyan indicates residues in alpha helical conformation only in the apo form. Sidechains are depicted as spheres for those residues that differ between strains. **c** Chart of chemical-shift derived secondary structure predictions for apo (open bars) and holo (filled bars) PiuA, showing only subtle differences upon ligand binding