Hydration Changes Accompanying the Binding of Minor Groove Ligands with DNA

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ABSTRACT 4',6-Diamidino-2-phenylindole (DAPI), netropsin, and pentamidine are minor groove binders that have terminal –C(NH2)2+ groups. The hydration changes that accompany their binding to the minor groove of the (AATT)2 sequence have been studied using the osmotic stress technique with fluorescence spectroscopy. The affinity of DAPI for the binding site decreases with the increasing osmolality of the solution, resulting in acquisition of 35 ± 1 waters upon binding. A competition fluorescence assay was utilized to measure the binding constants and hydration changes of the other two ligands, using the DNA-DAPI complex as the fluorescence reporter. Upon their association to the (AATT)2 binding site, netropsin and pentamidine acquire 26 ± 3 and 34 ± 2 additional waters of hydration, respectively. The hydration changes are discussed in the context of the terminal functional groups of the ligands and conformational changes in the DNA.

INTRODUCTION

Water is an important structural component of DNA, and its effect on the function of DNA is also significant, especially in cellular environments where high concentrations of solutes limit the amount of water (1–3). This work considers hydration effects for small molecules that are used as drugs to target particular DNA sequences via the minor groove (4). To understand the role of water in these reactions, several approaches are used. For example, high-sensitivity calorimetry with structural data shows that the change in the heat capacity for reaction is correlated with the loss of structured water from a DNA-ligand interface (5). However, water is involved beyond the entropic contribution due to expulsion from solvent-exposed regions. To illustrate, crystallography studies demonstrate that waters make intermolecular contacts between ligands and DNA (6). Spectroscopic techniques highlight the complicated dynamics of macromolecules such as DNA, and special attention has been focused on bound water in the ligand-DNA complex (7,8). Thermodynamic studies in conjunction with theoretical calculations and x-ray crystallography demonstrate that water mediates contacts between ligands and the DNA binding site (9,10). Insight into the larger population of waters involved in the DNA-small molecule reaction is obtained from volumetric studies (11).

Another approach for measuring the total hydration changes is the osmotic stress method (12). In these experiments, water is treated as a ligand, and its involvement in the ligand-DNA association is analyzed using linkage thermodynamics (13). The activity of water in the bulk solution is affected via cosolutes that are excluded from the vicinity of the DNA (14). Via this method, water has been shown to be involved in the association of proteins with DNA, and the extent of the hydration changes is influenced by the base sequence and conformational changes (15,16). A significant result from studies of small molecules interactions with DNA is that water acquisition occurs with complex formation (17–19). To consider how the structure of the ligand influences the hydration changes, three small molecules with terminal –C(NH2)2+ groups——4',6-diamidino-2-phenylindole (DAPI), netropsin, and pentamidine——were reacted with the (AATT)2 binding site (Fig. 1). To measure how the binding affinities vary with osmotic pressure, the fluorescence reporter DAPI is used to measure relative equilibrium constants for the netropsin and pentamidine exchange reactions. The hydration changes are examined from the standpoint of the common terminal functional groups in the ligands and their conformational effects on DNA.

MATERIALS AND METHODS

All experiments were performed in 10 mM H2PO4−/HPO42− buffer with 50 mM NaCl and 1 mM Na2EDTA. Single strands of 5'-GGCGCGAATTGC-GCG-3' (Integrated DNA Technologies; Coralville, IA) were annealed by slow cooling of the solution from 95°C to 10°C over 48 h. The concentration of the single stranded DNA was measured spectrophotometrically using extinction coefficient derived from the nearest neighbor approximation (125, 800 M−1 cm−1 at 260 nm) or after the complete digestion using nuclease BAL-31 (New England Biolabs; Ipswich, MA) (1). The concentrations of the drugs were determined using the following extinction coefficients: 27,000 M−1 cm−1 at 342 nm for DAPI; 21,500 M−1 cm−1 at 296 nm for netropsin; 28,900 M−1 cm−1 at 262 nm for pentamidine (20). The buffer components, netropsin, pentamidine (Sigma-Aldrich; St. Louis, MO), and DAPI (Molecular Probes; Carlsbad, CA) were used without purification. The osmolytes acetamide (Sigma-Aldrich), betaine monohydrate, trimethylamine N-oxide dihydrate (Fluka; St. Louis, MO), and triethylene glycol (TCI, Portland, OR) were also used as received. The osmolalities of the solutions were measured on a Wescor 5520 (Wescor; Logan, UT). Absorbance measurements were carried out with a Cary 50 spectrometer (Varian; Palo Alto, CA). Fluorescence data were collected on a Fluorolog spectrometer (Jobin-Yvon Horiba; Edison, NJ) at room temperature. Isothermal titration calorimetry (ITC) experiments were performed in triplicate on Microcal VP-ITC.

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the relative equilibrium constant for the exchange reaction
with the competitors netropsin or pentamidine were analyzed to determine
length of 461 nm (19). Titrations of the preformed DNA-DAPI complex
tributions from the bound and unbound DAPI:
the same relative affinities, suggesting that small amounts of unbound oligo-
needed at each osmolality. Titrations with 99% bound oligonucleotide yielded
association reaction (Eq. 1) were used to calculate the DAPI concentrations
preformed complex with DAPI, the equilibrium constants for the DAPI
and the duplex oligonucleotide. To insure that 95% of the DNA was in the
two strands. The relative equilibrium constants were similar for the hairpin
TTGCTTTTGCAATTGC-3
experiments, netropsin was titrated to the preformed DAPI-DNA complex
DNA-ligand complexes did not dissociate during the time period of the
experiments. Netropsin was titrated to the preformed DAPI-DNA complex
using a hairpin-forming oligonucleotide. This oligonucleotide—5'-GCAA-
TTGCTTTTGCAATTGC-3'—contained the (AATT)2 binding site in the (AATT)2
binding site like the
960 Degtyareva et al.
Biophysical Journal 92(3) 959–965

\[
F = F_1[\text{DAPI}] + F_0[\text{DNA} - \text{DAPI}],
\]

where \([\text{DAPI}]\) is the unbound DAPI concentration during the DNA titration and \(F_1\) is the intrinsic fluorescence of the unbound DAPI (for a 1 M concentration). \([\text{DNA} \cdot \text{DAPI}]\) and \(F_0\) are the corresponding parameters for the bound DAPI. During the titration with the competitors, the observed emis-

\[
F = F \frac{[\text{DNA} \cdot \text{DAPI}]}{[\text{DAPI}]} + F \frac{[\text{DNA} - \text{DAPI}]}{[\text{DNA}]} = F_1[\text{DAPI}] + F_0[\text{DNA} - \text{DAPI}],
\]

\[
\frac{d\ln K}{d\text{Osm}} = \frac{-\Delta N_a}{55.6}.
\]

RESULTS

DAPI

For the study of the minor groove complexes in this study, DAPI was used as the fluorescent reporter in the competitive equilibria. Thus, the reference reaction of DAPI with the (AATT)2 binding site is discussed first. The fluorescence quantum yield of DAPI associated with DNA is significantly higher relative to the free DAPI, thus allowing a distinction of bound and unbound ligand for the measurement of the equilibrium constant. DAPI has a high affinity for the minor groove of A/T-rich binding sites (21). To ascertain the hydration changes for the association reaction, DAPI was titrated with the oligonucleotide in the presence of varying concentrations of the osmolytes acetamide, betaine, tri-
extered, with 35 ± 1 additional waters being measured using the four osmolytes (Fig. 2 B). The structures of the osmolytes differ significantly, but their in-
fluence on the equilibrium constants is similar. This observation suggests that the osmolytes are excluded from the vicinity of the DNA and thus primarily act to reduce the activity of water in the bulk solution. In further support of osmolyte exclusion, extrapolation to the buffer conditions in

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\left(1 - \frac{1}{K}\right)[\text{DNA} - \text{Competitor}]^2
\]

\[
- \left(\frac{[\text{DAPI}]}{K} + [\text{Competitor}] + [\text{DNA}]\right)
\]

\[
\times [\text{DNA} - \text{Competitor}] + [\text{DNA}]_i[\text{Competitor}]_i = 0,
\]

where \(K\) is the equilibrium constant for the exchange reactions (Eq. 2) and the total concentrations of the competitor and oligonucleotide are \([\text{Competi-
tor}]\) and \([\text{DNA}]\), respectively. The values of \(K\), \(F_1\), and \(F_0\) were obtained from Eqs. 3 and 4 using least squares fitting of the observed emission in-
tensity as a function of added competitor. The hydration changes (\(\Delta N_a\)) were measured using the change in the observed equilibrium constant (\(K\)) with the change in the osmolality (Osm) of the cosolute (12)

\[
\frac{d\ln K}{d\text{Osm}} = \frac{-\Delta N_a}{55.6}.
\]

\[
F = F_1[\text{DAPI}] + F_0[\text{DNA} - \text{DAPI}],
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\[
\frac{d\ln K}{d\text{Osm}} = \frac{-\Delta N_a}{55.6}.
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\[
\left(1 - \frac{1}{K}\right)[\text{DNA} - \text{Competitor}]^2
\]

\[
- \left(\frac{[\text{DAPI}]}{K} + [\text{Competitor}] + [\text{DNA}]\right)
\]

\[
\times [\text{DNA} - \text{Competitor}] + [\text{DNA}]_i[\text{Competitor}]_i = 0,
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tor}]\) and \([\text{DNA}]\), respectively. The values of \(K\), \(F_1\), and \(F_0\) were obtained from Eqs. 3 and 4 using least squares fitting of the observed emission in-
tensity as a function of added competitor. The hydration changes (\(\Delta N_a\)) were measured using the change in the observed equilibrium constant (\(K\)) with the change in the osmolality (Osm) of the cosolute (12)
with the DNA. Preferential interaction coefficients provide a direct measure of the extent of interaction of cosolutes with DNA, and the exclusion of betaine from the vicinity of the DNA has been demonstrated by vapor pressure osmometry (23). The changes in the dielectric constant due to the osmolytes are not a significant factor, as comparable results were obtained using betaine and acetamide that have opposite effects on the dielectric constant (24).

**Netropsin and pentamidine**

Having established the binding affinities and hydration changes that accompany the binding of DAPI with (AATT)$_2$, netropsin and pentamidine (Fig. 1) were studied. Like DAPI, these drugs also favor binding in the minor groove of A/T-rich sequences. To measure the relative affinities of netropsin and pentamidine for the oligonucleotide, competition titrations using DAPI-(AATT)$_2$ were used, and the progress of the titration was followed using the decrease in the fluorescence intensity (Figs. 3 A and 4 A). The netropsin and pentamidine complexes are hydrated to a comparable extent as the DAPI-(AATT)$_2$ complex, resulting in relative binding affinities that do not change significantly the osmolality of the solution (Figs. 3 B and 4 B). Using the hydration changes for the DAPI-(AATT)$_2$ complex, the differences in hydration between the DNA and DNA-ligand complex are 26 ± 3 waters for netropsin and 34 ± 2 waters for pentamidine.

The DAPI concentration was relatively low (<90 nM at the highest osmolyte concentrations) compared to the concentrations where additional binding sites are populated, so aggregation with the DNA should not perturb the results (19,25). For the weaker binding pentamidine, higher concentrations were needed to displace the DAPI. To discern possible additional binding for pentamidine at these higher concentrations, isothermal titration calorimetry studies were used. The derived binding site size is 0.94 ± 0.06 pentamidines/oligonucleotide, which is consistent with binding in the minor groove (Fig. 5). Importantly, titration to higher stoichiometries showed no additional binding modes. To support the exchange model in Eq. 2, the DAPI emission after adding saturating amounts of netropsin or pentamidine was comparable to the emission of the unbound DAPI (Figs. 3 A and 4 A). Furthermore, the fluorescence intensity changes upon addition of netropsin or pentamidine were well fit using the model provided by Eq. 2. Relative binding constants determined from extrapolation to the buffer conditions agree with the values measured in buffer (Figs. 3 B and 4 B). This agreement was also observed for the DAPI reaction with the oligonucleotide and supports osmolyte exclusion. Using the relative binding affinities with the affinity of DAPI for the oligonucleotide, the binding constants for netropsin and pentamidine with the oligonucleotide in the buffer were determined. For netropsin, the equilibrium constant of 3.4 (± 0.7) × 10$^9$ M$^{-1}$ is consistent with the high affinities measured using different experimental techniques (22,26). For pentamidine, the binding affinity of 1.3 (± 0.1) × 10$^6$ M$^{-1}$ is in good agreement with the value of 1.8 (± 0.3) × 10$^6$ M$^{-1}$ derived from the calorimetry studies (Fig. 5).

**DISCUSSION**

The hydration changes that accompany the interaction of three minor groove binding ligands—DAPI, netropsin, and pentamidine—with the (AATT)$_2$ sequence have been studied. Hydration is an important feature of this binding site, as
exemplified by its spine of hydration and cations (27,28).

The complementary fit of these ligands into the minor groove results in displacement of most of the interfacial waters, as demonstrated by x-ray crystallography and thermodynamic studies (5,29–33). However, waters remain an important feature of these complexes. For example, the crystallography studies show that waters bridge the ligand and DNA, particularly at the ends of the ligands (29–33). The significance of terminally bound water is highlighted by recent studies of ligands whose shapes are not complementary to the minor groove binding site. For example, a diamidine compound with a linear shape was reacted with an (AATT)2 binding site (9).

Based on x-ray crystallography and molecular dynamics simulations, waters mediate contacts between one of the amidine groups and the functional groups of the minor groove. The high affinity of this ligand for the binding site strongly suggests that hydration is critical for the formation of this complex. The thermodynamic significance of terminally bound water is highlighted by recent studies of ligands whose shapes are not complementary to the minor groove binding site.
bound waters was also demonstrated in a calorimetry study of the binding of netropsin with an (AATT)$_2$ binding site (10,22). Two binding modes were observed, of which one was attributed to the complementary fit of the crescent shaped netropsin with the binding site. The conformational flexibility of netropsin permits the formation of alternate structures, and the second binding mode was assigned to the interaction of a more linear netropsin with the binding site. For this complex, simulations supported water-mediated contacts between the minor groove and the amidino nitrogens at the terminus of the netropsin. In addition, the enthalpy and entropy changes for this second binding mode are consistent with previous studies that considered bridging waters in the complex (34).

The above studies show that small numbers of waters link small molecule ligands and the DNA via intermolecular bonding. The involvement of a larger population of waters in the complex is supported by theoretical, spectroscopic, and thermodynamic studies. The full range of bound waters is treated in recent theoretical treatments of macromolecular hydration (35,36). Spectroscopic studies of hydration suggest that bound waters extend away from the macromolecular surface (37). Thermodynamic methods measure the total hydration changes for reactions involving macromolecules (11,12). For example, DNA binding sites gain up to 30 additional waters when associated with small molecule intercalators (18). Importantly, the amount of solvent exposure of the DNA bound ligand dictates the amount of water uptake, a trend that is supported by studies of minor groove binders. For example, Hoechst 33258 binding with the (AATT)$_2$ minor groove enhances the hydration of the complex relative to the reactants by 60 ± 13 waters (19). Netropsin binding with mixed sequence DNA is coupled with the uptake of 50–60 waters (38). Consistent with the results from our study, higher affinity binding affinity sites such as the minor groove are associated with a smaller amount of additional waters. For both Hoechst and netropsin, volumetric techniques measure water expulsion for the binding of netropsin with A/T-rich oligonucleotides (39,40). These differences could be expected because the volumetric methods measure changes in the hydrogen bonding structure of water, whereas the osmotic stress method measures the stoichiometry of water exchange based on the exclusion of the osmolytes.

DNA conformational changes may also contribute to the water acquisition. For example, osmotic stress studies showed that oxygen binding to hemoglobin is accompanied by ~60 additional waters (41,42). Using structural data for the two forms of hemoglobin, the additional solvation was attributed to the surface area changes due to the reaction. Other studies have also related hydration changes to alterations in the solvent exposure (43,44). For the minor groove binders used in this study, the conformational changes are small but potentially significant (45–48). To further understand the significance of this effect, enhanced solvent exposure could be probed using osmolytes with varying sizes (43,49).

**CONCLUSION**

Understanding the role of water is important for efforts to target specific DNA sequences using drugs such as small molecules that bind in the minor groove (10,50). The results of this study show that the association of the minor groove binders DAPI, netropsin, and pentamidine with the (AATT)$_2$ binding site is accompanied by the acquisition of water. The ligands differ significantly with respect to their size and functional groups that bind with the DNA, but they have the common structural feature of terminal -C(NH$_2$)$_2$$^+$ groups. The similar amounts of water gained in the complex suggest that the termini of the ligands are important for the hydration changes. This conclusion is consistent with x-ray crystallography and thermodynamic studies which show that terminally located waters provide mediating interactions between the ligand and DNA. In addition, the ligands may produce similar conformational changes in the DNA that could result in enhanced hydration. The osmotic stress method offers a convenient method for assessing the involvement of water in DNA reactions, and our studies provide insight into structural factors that influence water acquisition.

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