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Solution Structure of the Complex between the Head-to-Tail Dimer of Calicheamicin γ_1^I Oligosaccharide and a DNA Duplex Containing d(ACCT) and d(TCCT) High-Affinity Binding Sites

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Abstract: The head-to-tail dimer of the calicheamicin oligosaccharide domain exhibits substantially higher DNA binding affinity and sequence selectivity and greater bioactivity than the monomer from which it is derived. To determine the structural basis for these functional properties, the solution structure of the 1:1 complex between the head-to-tail dimer of the calicheamicin oligosaccharide and the oligonucleotide duplex d(GCACCTTCCTGC)·d(GCAGGAAGGTGC) has been solved by restrained molecular dynamics calculations using NMR-derived distance and torsion angle constraints. The final input data consisted of 562 internuclear distance and 114 dihedral angle constraints, an average of 27 constraints per residue. In contrast to observations made for a complex between a DNA duplex and the head-to-head dimer of calicheamicin oligosaccharide, the head-to-tail dimer exhibits a unique binding mode in the DNA minor groove. A comparative analysis of the carbohydrate–DNA interactions at the two different binding sites explains at the atomic level how calicheamicin derivatives are able to effectively recognize both d(ACCT) and d(TCCT) sites. This study brings deeper insight into the factors governing DNA-binding affinity and the sequence preferences of calicheamicin and its derivatives.

Introduction

Despite the vast potential of new therapeutic modalities such as antisense and gene therapies, small molecule ligands remain as the mainstay of pharmaceutical agents in use today. Molecules that recognize and bind to DNA have been widely studied, and some have found clinical applications, for example, in cancer chemotherapy. One promising class of these agents are those that not only bind to DNA but also perform chemistry on this substrate. The enediyne, which include calicheamicins, esperamicins, dynemicin A, neocarzinostatin chromophore, kedarcin chromophore, and C1027 chromophore, form one family of extremely potent natural products of this type.

Calicheamicin γ_1^I (**1**)¹ (Scheme 1) has the strongest cytotoxicity among the natural enediynes, and it exhibits an intriguing mode of binding to DNA.^{2,3} Two chemical functionalities have been shown to be responsible for its high cytotoxic activity (Scheme 1): a molecular device, the enediyne

aglycon that upon activation forms a phenyl diradical, which leads to the cleavage of the DNA, and a binding moiety (the carbohydrate domain) that directs the reaction to certain sequences. Previous structural studies from this and other laboratories have shown that the oligosaccharide moiety binds in the DNA minor groove.⁴ The binding to duplex DNA actually occurs in a selective manner, with high affinity for four specific, primarily polypurine/polypyrimidine sites: d-TCCT·d-AGGA, d-TTTT·d-AAAA, d-TCTC·d-AGAG, d-ACCT·d-AGGT.⁵

The methyl glycoside of calicheamicin (**2**) has been shown to inhibit binding of transcription factors to sequences containing the TCCT high-affinity binding site and to interfere with the corresponding DNA transcription in vivo.⁶ As part of the effort to explore and build upon this remarkable observation, we have designed and synthesized oligomers of the calicheamicin oligosaccharide domain.⁷ The concept of linking two or more DNA binding agents to generate higher binding affinity, greater

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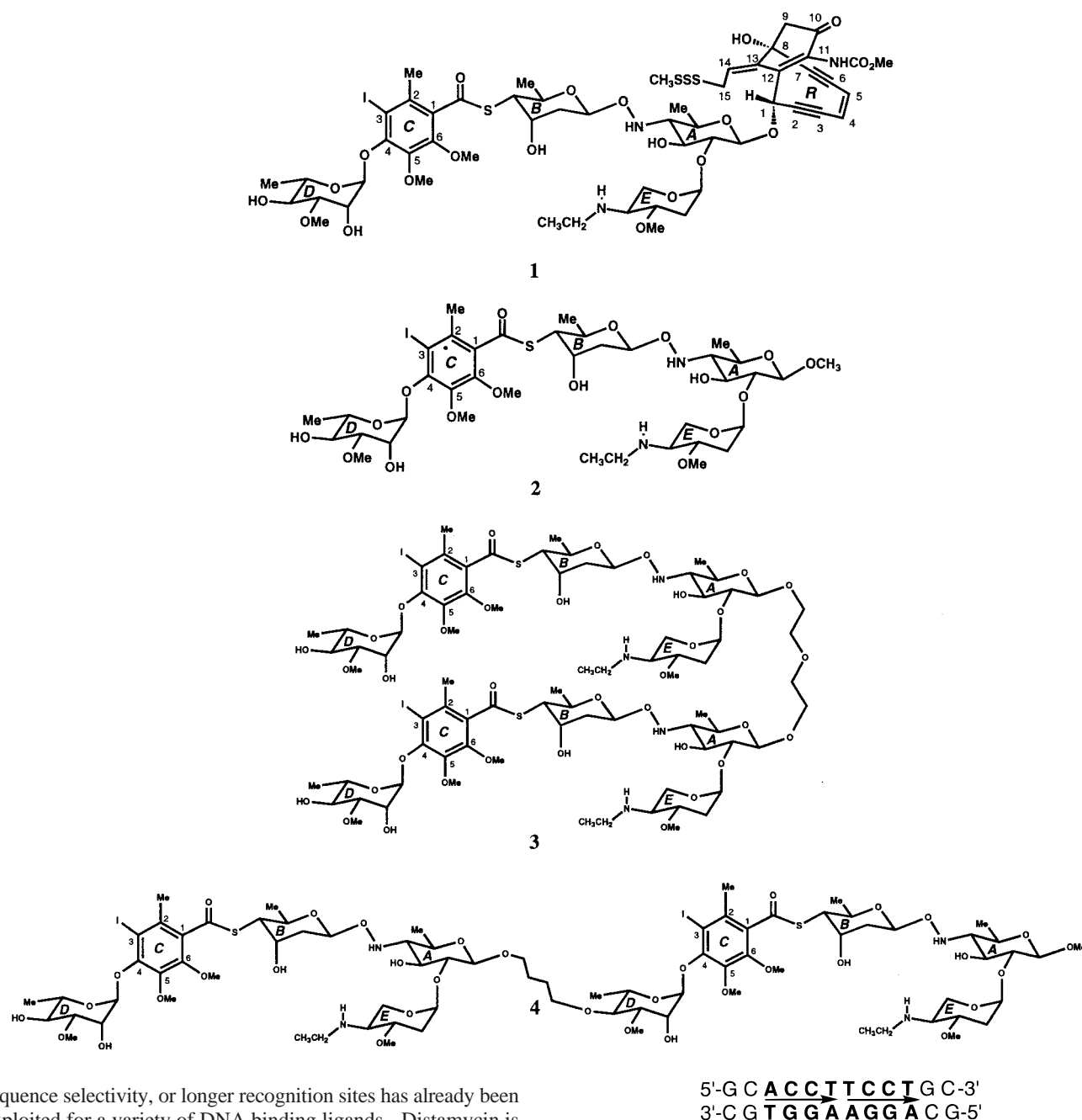
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Scheme 1. Structures of Calicheamicin (1), Calicheamicin Oligosaccharide Domain (2), Head-to-Head Calicheamicin Oligosaccharide Dimer (3), and Head-to-Tail Calicheamicin Oligosaccharide Dimer (4)



sequence selectivity, or longer recognition sites has already been exploited for a variety of DNA binding ligands. Distamycin is probably the most well-studied example; analogues were shown to have increased affinity many years ago, and ultimately, composite distamycin-based agents capable of high sequence specificity have been developed.⁸ The calicheamicin analogues examined in this study represent a unique effort to explore the DNA ligand oligomerization concept using a carbohydrate motif.

The head-to-head^{7a} (3) and head-to-tail^{7b} (4) dimers (Scheme 1) of calicheamicin oligosaccharide domain have been shown

Figure 1. DNA sequence with the dual head-to-tail binding site in bold and the two subsites identified by the arrows.

to possess superior binding properties and improved bioactivity relative to the monomer. The dimers exhibit increased binding affinity ($K_D \sim 10^{-6}$ M for the monomer vs $K_D \sim 10^{-9}$ for the dimers) and site specificity (100–1000-fold higher than the monomer vs the theoretical increase of 64-fold),⁹ as well as more than 10-fold greater inhibition of transcription factor binding and transcriptional activity.¹⁰

In this paper, we report the results of the NMR structural analysis of the complex between the head-to-tail dimer (HTD)

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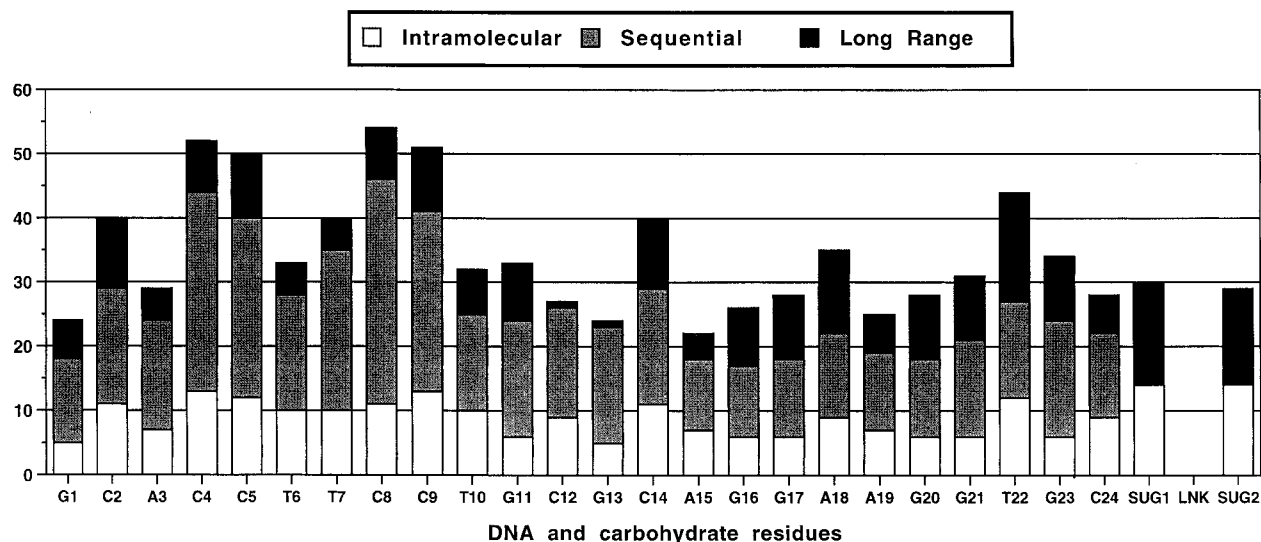


Figure 2. Plot of all the NMR-derived distance constraints used in the structure calculation of the HTD-DNA complex ordered by type.

(4) and duplex DNA (see Figure 1). This structure complements the previous study of a complex of the head-to-head dimer bound to a DNA duplex,¹¹ providing further insight into the molecular basis for the binding properties of these unique dimeric, carbohydrate-based DNA binding agents and into primary structure-function relationships for the calicheamicins. These results also contribute to a growing effort to understand the fundamental principles of biological recognition of carbohydrates.

Experimental Section

Synthesis of the Oligonucleotide. The two complementary oligonucleotides 5'-GCACCTTCCTGC-3' and 5'-GCAGGAAGGTGC-3' were synthesized on a Beckmann 200A automatic synthesizer using phosphoramidite chemistry. Both oligomers were purified by ion-exchange HPLC on a Partisil 10 SAX column, eluting with 20% aqueous CH₃CN and a linear gradient of KH₂PO₄ from 10 mM to 0.35 M at pH = 7.0. The oligos were then desalted by gel filtration on a Biogel P2 column.

Preparation of the Complex. The two complementary strands were mixed in water in equimolar amounts and annealed by heating at 90 °C for 5 min, followed by slow cooling to room temperature. The NMR sample was prepared by lyophilizing the duplex two times from 99.6% D₂O and then dissolving the lyophilized material in 500 μ L of 10 mM phosphate buffer containing 10 mM NaCl and 0.1 mM EDTA at pH 7.0 in 99.996 D₂O. For experiments to assign the labile protons, the sample was lyophilized and redissolved in 90% H₂O/10% D₂O. The final concentration of the duplex was 1.5 mM.

The ligand-DNA complex was prepared according to the following procedure. Known amounts of the ligand dissolved in MeOH were added to 1 mL of a 0.75 mM solution of the duplex, stirred at 25 °C for 10 min, and then lyophilized and dissolved in 99.96% D₂O. The extent of the titration was monitored by examining the DNA 6H/8H resonances in the ¹H NMR spectrum. The final solution was lyophilized and redissolved in 500 μ L of 99.996% D₂O. To examine the labile protons of the complex, the solution was lyophilized and redissolved in 500 μ L of a 90% H₂O/10% D₂O solution.

NMR Experiments. All NMR experiments were performed on a Bruker AMX2-500 spectrometer. The temperature was 27 °C for both the free DNA duplex and the ligand-DNA complex. All spectra were acquired in the phase-sensitive mode. The transmitter was placed on the solvent resonance, and the TPPI method was used to achieve frequency discrimination in the ω_1 dimension.¹² The standard pulse

sequence and phase cycling were used for 2Q¹³ and PE-COSY¹⁴ spectra. A total of 64 scans/ t_1 value were acquired for the 2Q ($t_{\text{mix}} = 30$ ms, $t_{1\text{max}} = 50$ ms) and PE-COSY ($t_{1\text{max}} = 80$ ms). A TOCSY spectrum was acquired using the DIPSI-2 sequence¹⁵ for spin locking with $t_{\text{mix}} = 70$ ms, 64 scans/ t_1 and $t_{1\text{max}} = 40$ ms. NOESY¹⁶ spectra from H₂O solutions were recorded with the last pulse replaced by a jump and return composite sequence.¹⁷ NOESY spectra from D₂O solution were recorded with saturation of the residual HOD resonance during the preparation and mixing periods and a Hahn-echo to improve the quality of the baseline.¹⁸ The NOESY spectra were acquired with mixing times of 50 and 200 ms, 64 scans/ t_1 , and $t_{1\text{max}} = 50$ ms.

The NMR data were processed on a SGI INDIGO² workstation using FELIX 95 software (Biosym-MSI, San Diego, CA).

Restrained Molecular Dynamics Calculations. The structures of the complex were generated by a protocol involving several iterations through cycles of simulated annealing docking and refinement (Scheme 2). All restrained energy minimizations (rEM) and restrained molecular dynamics calculations (rMD) were performed on a Convex C-240 meta-cluster using the SANDER module of the AMBER 4.1 software package.¹⁹ The potential functions for distance and dihedral constraints were flat between the given upper and lower bounds and rise parabolically outside of these bounds. These parabolic functions are turned smoothly into a linear function using a special feature of SANDER in order to avoid large violations. No explicit solvent molecules were included in these calculations, so a distance-dependent dielectric and reduced net charges on the phosphate oxygens (scaled by 0.2) were used to partially compensate for the absence of solvent. A distance cutoff of 9 Å was set for all nonbonded interactions. The force constant for both distance and dihedral constraints was set to 32 kcal/mol·Å².

The starting conformation of the calicheamicin oligosaccharide dimer was generated by linking two oligosaccharide molecules (derived from our structural model of the calicheamicin-DNA complex) through a saturated four carbon chain linker. Twenty different DNA starting conformations were constructed using Nucleic Acid Builder (NAB),²⁰ which allows generation of coordinates for nonstandard nucleic acid structures with a range of helical parameters. The initial geometries

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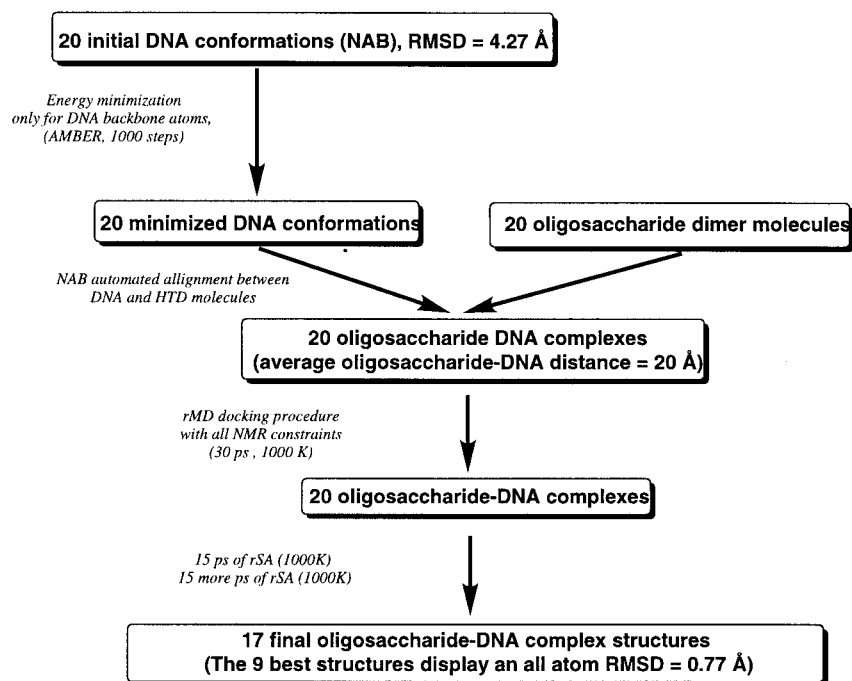
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Scheme 2. Protocol Used for the Calculation of the Solution Structure of the HTD–DNA Complex

were chosen with the stipulation of their being within the family of right-handed DNA helices. These structures sampled a range of conformations, exhibiting an RMSD over all atoms of 4.27 Å. The variation in their x -displacement, inclination, twist, and rise is listed in Table S1 (Supporting Information).

The docking procedure consisted of an “automated” alignment of the oligosaccharide and DNA molecules using NAB, followed by 30 ps of rMD docking, as described previously.^{4c} The alignment between the oligosaccharide molecule with each DNA conformation was achieved using the following procedure: (a) six pairs of DNA and ligand atoms were chosen among the residues that were more likely to be involved in the binding (based on intermolecular NOEs), and (b) the two molecules were positioned so that all ligand atoms are separated from DNA atoms by at least 20 Å. The distance between ligand and DNA atoms was selected assuming that a short distance is more likely to generate van der Waals (VDW) clashes, while a larger value would have compromised the rate of convergence of the subsequent steps of rSA. After the two molecules were positioned, the complex was obtained by performing 30 ps of rSA (1000 K) with a very slow ramping up of the force constant of the penalty function for the experimental constraints (0.1–32 kcal/mol-Å²). These docked structures were subjected to another 20 ps cycle of rSA (1000 K). These were then arranged in order of increasing experimental constraint violation energy, and the nine lowest violation energies were selected for detailed conformational analysis and display in Figure 3. The coordinates of these nine solution structures are being deposited with the nucleic acids section of the Brookhaven Protein Data Bank.

Results and Discussion

¹H NMR. The general strategies for sequence-specific assignment of the ¹H resonances of small DNA duplexes have been amply reviewed,²¹ and the specific protocols used here for the free DNA and ligand–DNA complex have been described.²² The shorthand notation described by Wüthrich^{21b}

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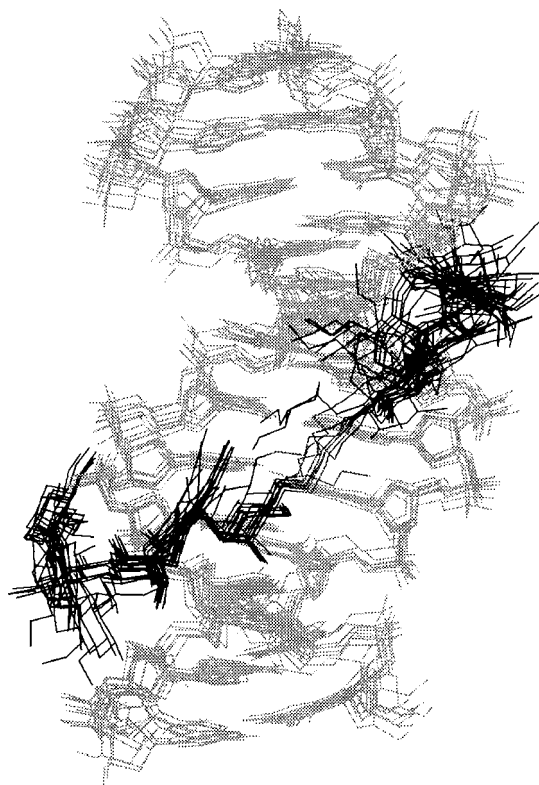


Figure 3. Ensemble of nine conformations representing the three-dimensional solution structure of the HTD–DNA complex. The DNA is displayed in light gray, the ligand subunits in black, and the linker between the two subunits of the ligand in dark gray.

is used to specify DNA interproton distances and corresponding NOEs. For the free duplex, all cytosine 5H–6H and thymine 5Me–6H resonances were readily identified by scalar connectivities in 2Q, PE-COSY, and TOCSY spectra. All 24 discrete 1′H–2′H–2′′H–3′H spin subsystems were identified in the 2Q, PE-COSY, and TOCSY spectra and were confirmed by inspection of the NOESY spectra. The sequential resonance assignments were obtained from the 50 ms mixing time NOESY

Table 1. ^1H Chemical Shifts in the $\text{HTD}-\text{d}(\text{GCACCTTCCTGC})\cdot\text{d}(\text{GCAGGAAGGTGC})$ Complex at $\text{pH} = 7.0$, $T = 300\text{ K}$

DNA Chemical Shifts ^a (ppm)										
residue	N1H, N3H	NH2	2H, 5H, 5-Me	6H, 8H	1'H	2'H	2''H	3'H	4'	5'H/5''H
G1				7.74	5.95	2.53	2.72	4.98	4.31	
C2		6.37, 8.46	5.40	7.51	5.41	2.23	2.46	4.91	4.20	
A3			7.92	8.30	6.52	2.75	3.12	5.12	4.52	
C4		6.26, 8.05	5.19	7.05	6.06	1.85	2.89	4.93	4.15	3.63, 3.97
C5		6.49, 7.81	5.46	7.35	5.66	1.85	2.76	4.65	4.55	4.23
T6	13.30		1.59	7.49	6.29	1.91	2.73	4.57	4.21	
T7	13.88		1.59	7.32	6.25	2.11	2.88	4.99	4.28	
C8		6.57, 8.19	5.56	7.40	6.16	1.94	2.94	4.94	4.21	
C9		6.55, 8.00	5.49	7.38	5.50	2.00	2.64	4.59	4.47	3.73, 4.30
T10	13.28		1.61	7.44	5.94	1.95	2.47	4.68	4.05	
G11	12.87			7.93	6.05	2.47	2.79	5.02	4.41	
C12			5.53	7.48	6.21	2.20	2.21	4.55	4.08	
G13				7.74	5.95	2.46	2.72	5.07	4.22	
C14		6.48, 8.54	5.49	7.52	5.61	2.18	2.48	4.90	4.22	
A15			7.68	8.14	6.28	2.71	3.10	5.08	4.46	
G16	13.33			7.49	5.44	2.51	2.56	4.99	4.41	
G17	12.67			7.96	5.99	2.63	2.79	4.87	4.24	
A18			7.37	7.99	6.04	2.65	2.83	4.87	4.27	3.75, 3.72
A19			7.71	7.93	6.25	2.48	3.05	4.86	4.42	
G20	13.28			7.59	5.48	2.46	2.53	4.96	4.36	
G21	12.92			7.94	6.13	2.47	2.74	5.10	4.38	
T22	14.09		1.22	7.00	6.06	2.08	2.54	4.83	3.46	3.09, 3.59
G23	13.00	8.05		7.96	5.98	2.59	2.79	4.96	4.15	
C24			5.49	7.51	6.27	2.22	2.22	4.55	4.13	
Ligand Chemical Shifts ^a (ppm)										
residue	H1	H2ax	H2eq	H3	H4	H5	Me	<i>N</i> -CH ₂ CH ₃	OMe	
A	4.31	3.32		4.41	2.46	3.97	1.54			
B	5.24	1.98	2.39	4.39	3.78	4.18	1.44			
C							2.43			3.87 (C5)–4.10 (C6)
D	5.19		4.68	4.09	3.78	4.39	1.50			3.64
E	5.36	1.69	2.80	3.96	3.30	4.25		3.18–1.39		3.63
A'	4.31	3.32		4.46	2.44	4.01	1.56			
B'	5.28	1.96	2.51	4.28	3.73	4.14	1.47			
C'							2.46			3.87 (C5)–4.10 (C6)
D'	5.35		4.65	4.04	3.67	4.26	1.44			3.64
E'	5.39	1.71	2.78	4.01	3.29	4.14		3.24–1.32		3.65

^a The chemical shifts are referenced to 3-(trimethylsilyl)propionic-2,2,3,3-*d*₄ acid, sodium salt, using the HOD resonance previously calibrated in stock buffer solution.

spectrum. The combination of $d_i(6,8;2')$, $d_s(2'';6,8)$, $d_i(5\text{Me};6)$, $d_s(6,8;5\text{Me})$ NOEs was sufficient to obtain complete sequence-specific resonance assignments. The relative intensities of characteristic NOE cross-peaks [e.g., $d_i(6,8;2')$, $d_s(2'';6,8) \gg d_i(6,8;2'')$, $d_s(2';6,8)$ and $d_i(5\text{Me};6) > d_s(6,8;5\text{Me})$] and the cross-peak patterns in the PE-COSY spectrum clearly indicated that the free duplex is a B-form duplex with C2'-endo-like sugar conformations.

Upon the binding of HTD (**4**) to the DNA, a small broadening of the line widths of all DNA and saccharide resonances was observed. The observation of one set of resonances over the full range of temperatures (10–37 °C) suggested a unique binding mode of the ligand to the DNA, in contrast to what was observed in the case of the HHD (**3**)–DNA complex, where two binding modes were detected.¹¹ The analysis of 2Q, TOCSY, and NOESY experiments enabled sequence-specific assignments to be made for the ligand and each DNA strand. The resonance assignments are provided in Table 1. The relative intensities of cross-peaks in the 50 ms NOESY indicated that in the presence of HTD (**4**) $\text{d}(\text{GCACCTTCCTGC})\cdot\text{d}(\text{GCAGGAAGGTGC})$ remains as a B-form duplex with C_{2'}-endo-like sugar conformations. However, a unique pattern of NOEs and scalar connectivities has been observed around the second cytosine of each recognition site (i.e., C₅ and C₉), suggesting that the deoxyribose ring conformation must be somewhat altered for these residues.

NMR-Derived Distance and Torsion Constraints. The chemical shift dispersion and resonance line widths of the HTD (**4**)–DNA complex were highly favorable relative to our experience with previously studied calicheamicin–DNA complexes. This facilitated both resonance assignment and accurate cross-peak volume measurements and ultimately led to a relatively large number of distance constraints. NOESY cross-peaks that could be unambiguously assigned for the first round of calculations were categorized into weak, medium, and strong on the basis of the cross-peak intensities and then converted into distance constraints with upper bounds of 5.0, 4.0, and 3.0 Å, respectively. The structures obtained using these constraints were used as input for hybrid relaxation matrix back-calculations (MARDIGRAS 5.1²³) to set target distances more precisely and assign additional constraints. Cross-peak intensities used for these calculations were taken from three NOESY spectra $\tau_{\text{mix}} = 50$ and 200 ms in D₂O and $\tau_{\text{mix}} = 200$ ms in H₂O/D₂O (9:1). The target distances obtained from this hybrid relaxation matrix approach were adjusted by adding 10% of the distance to set the upper bound, to account for experimental and systematic errors. Note that the newer RANDMARDI^{23b} protocol would obviate the need for this correction. Forty hydrogen-bonding restraints were also included on the basis of characteristic imino

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Table 3. Summary of the Molecular Energies, RMSDs, and Residual Violations of the Ensemble of the 9 Final Structures of the 1:1 Complex of HTD (**4**) and d(GCACCTTCCTGC)·d(GCAGGAAGGTGC)

molecular energies (kcal)	
E_{amber}	-926.1 ± 12.5
E_{viol}	8.9 ± 1.7
distance violations (Å)	
$0.01 < d < 0.20$	23.9 ± 2.1
$0.20 < d$	0.4 ± 0.7
dihedral angle violations (deg)	
$2 < \theta < 6$	3.56 ± 1.26
$6 < \theta < 10$	0.22 ± 0.44
average rmsd, all heavy atoms in the binding site (Å)	
HTD + DNA (pairwise)	1.16
HTD + DNA (from the mean)	0.77

Table 4. Intermolecular H Bonds^a Observed in the Solution Structure of the 1:1 Complex of HTD (**4**) and d(GCACCTTCCTGC)·d(GCAGGAAGGTGC)

donor	acceptor
B -3OH	T ₂₂ -O2
G ₂₁ -NH ₂	C -SC=O
D -2OH	T ₂₂ -O1P
B' -3OH	A ₁₈ -N3
G ₁₇ -NH ₂	C' -SC=O
D' -2OH	A ₁₈ -O1P

^a The H bonds are defined on the basis of a distance cutoff of 3.0 Å.

oligosaccharide residues E and E', which face out away from the DNA and are highly exposed to solvent.

To facilitate the discussion of the structure of the complex at the atomic level, a tube diagram of the representative structure (that closest to the mean) is shown in Figure 4 with color coding and labeling of the binding site. The binding of the HTD (**4**) molecule in the DNA minor groove is stabilized by the existence of many favorable intermolecular interactions. Among these are six hydrogen bonds (Table 4), two salt bridges between the positively charged nitrogen atoms of rings E and E' and the O1P oxygen atoms of residues C₄ and C₈, respectively, and a large buried intermolecular surface with extensive van der Waals contacts. In addition, each of the iodine atoms of HTD (**4**) exhibits a hydrogen-bonding-like interaction with a guanine NH₂ in the binding site as observed in several other studies on DNA complexes of calicheamicin-based ligands.^{4,11}

A detailed analysis of the DNA geometry was obtained using NEWHELIX software.²⁵ The helicoidal parameters, sugar pucker, and backbone torsion angles indicate that the oligonucleotide occupies a conformation in the B-DNA family. By way of example, the mean values and standard deviation of χ -displacement, inclination, twist, and rise are provided in Table 5. Some small local perturbations from typical B-DNA structure are observed, with the most significant being a shift in the deoxyribose ring conformation of the second cytosine (in direction 5' → 3') of each binding site (i.e., C₅ and C₉) toward O_{4'}-endo sugar pucker (PPA = 90°). This shift appears to be due to the high steric demand of the HTD iodine atom in ring C. A similar deoxyribose ring pucker for the corresponding DNA residue has been reported in other calicheamicin–DNA complexes.⁴ Although the experimental observables have been interpreted in terms of a perturbed sugar ring conformation, it should be noted that a similar structural result could be obtained if the sugar rings occupied an equilibrium distribution of C_{2'}-endo and C_{3'}-endo conformations. In fact, the currently

Table 5. Mean and Standard Deviation of a Selection of Representative Helical Parameters of the Ensemble of the 9 Final Structures of the 1:1 Complex of HTD (**4**) and d(GCACCTTCCTGC)·d(GCAGGAAGGTGC)

base pair	X displacement	inclination	twist	rise
1	-0.12 ± 0.53	12.54 ± 4.60	9.08 ± 5.70	7.43 ± 0.46
2	-0.70 ± 0.51	14.52 ± 3.10	38.77 ± 2.59	2.78 ± 0.19
3	-1.16 ± 0.33	15.13 ± 2.60	32.94 ± 2.12	2.95 ± 0.20
4	-1.19 ± 0.25	15.47 ± 2.69	39.04 ± 1.52	2.76 ± 0.20
5	-1.41 ± 0.21	14.42 ± 2.06	36.34 ± 2.31	3.08 ± 0.19
6	-1.57 ± 0.35	12.94 ± 1.55	33.81 ± 3.71	3.24 ± 0.17
7	-1.67 ± 0.33	13.23 ± 1.47	36.24 ± 3.30	3.21 ± 0.17
8	-1.47 ± 0.39	13.44 ± 1.23	39.30 ± 2.28	2.76 ± 0.20
9	-1.31 ± 0.53	12.34 ± 1.90	39.21 ± 4.54	2.77 ± 0.21
10	-1.53 ± 0.49	10.16 ± 2.86	34.64 ± 1.34	2.98 ± 0.18
11	-1.38 ± 0.38	7.49 ± 4.30	35.41 ± 2.20	3.16 ± 0.25
12	-0.50 ± 0.43	7.77 ± 5.07		
A-DNA	-3.5	19.0	33	2.6
B-DNA	0.0	2.0	36	3.4

available information is insufficient to distinguish unambiguously between the static and dynamic interpretations. However, the small difference in energy between O_{4'}-endo and C_{2'}-endo and C_{3'}-endo low-energy sugar ring conformations (2 kcal/mol or less)²⁶ suggests that the shift to the higher energy PPA could be readily accommodated.

Comparative Analysis of Calicheamicin Oligosaccharide Binding in the TCCT and ACCT Recognition Sites. The TCCT and ACCT sites to which the two subunits of the HTD (**4**) dimer are bound correspond to two of the four high affinity DNA sequences for calicheamicins; hence, it is clearly of interest to compare the structures of the two sites. We find that the intermolecular interactions within each of the dimer subunits are very similar. This similarity even extends to the position where the sequences are different: a hydrogen bond is formed from the carbohydrate B-3OH of each subunit, either to the thymine O2 in the ACCT site (5'-ACCT-3'/5'-AGGT₂₂-3') or the adenine N3 in the TCCT site (5'-TCCT-3'/5'-AGGA₁₆-3'). These homologous interactions are displayed in Figure 5, where the similarity in the position of the T₂₂–O2 and A₁₈–N3 is clearly evident. Thus, both AT and TA base pairs can be bound interchangeably at the 5' end of the binding site because both have a hydrogen-bond acceptor (the carbonyl O2 of thymine or the ring N3 of adenine) protruding into the center of the DNA minor groove.

The conformations of the two subunits of the HTD (**4**) dimer and their positioning within their DNA binding sites are also very similar, even though the sites are different (Figure 4). In the ACCT site, the HTD A and B rings are proximate to the A₃/T₂₂ and C₄/G₂₁ base pairs, and the amino sugar E is outside of the minor groove. The hexasubstituted aromatic ring C occupies a position between the two guanine residues G₂₁ and G₂₀ and appears to be closer to the purine than to the pyrimidine strand. Part of sugar residue D is actually outside the DNA minor groove with C-1, C-2 and the ring oxygen in close proximity to the phosphate backbone of the purine strand. The positioning of the ligand in the TCCT site, and the oligosaccharide–DNA interactions for the carbohydrate unit, are virtually identical to the ACCT site and are also similar to that observed in previous studies conducted on complexes between calicheamicin-based monomers and TCCT-containing oligonucleotides.⁴

Implications for Sequence Selectivity of the Calicheamicin Oligosaccharide. This work brings deeper insight into the

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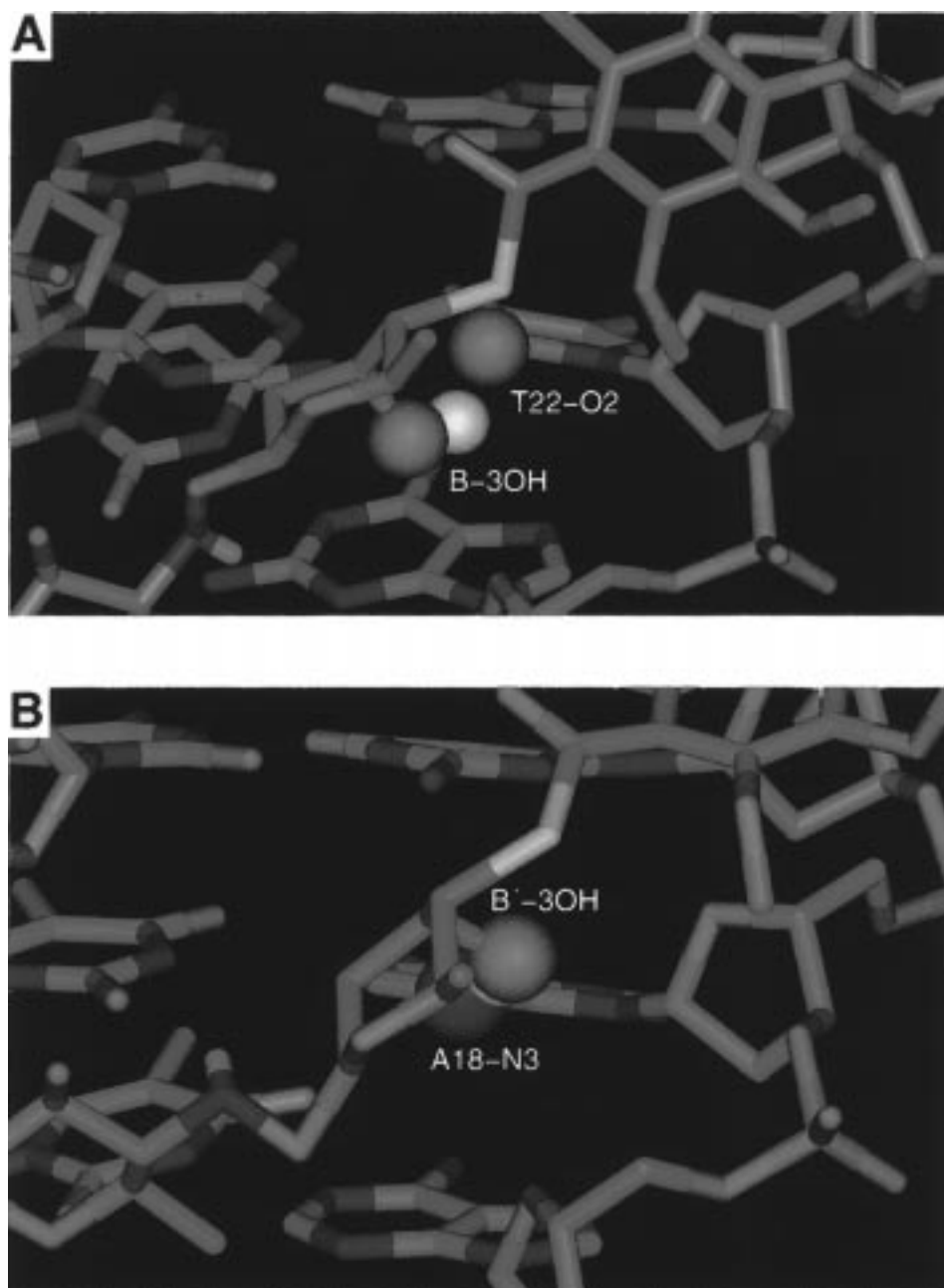


Figure 5. Close-up view of the homologous hydrogen-bonding interactions of HTD to different DNA functional groups in the HTD-DNA complex. The hydrogen bonds between (A) T₂₂-O₂ and the 3OH of ring B and (B) between A₁₈-N₃ and the 3OH of ring B' are highlighted with spheres colored red for oxygen, blue for nitrogen, and gray for hydrogen.

factors governing the DNA binding affinity of calicheamicin and its derivatives. The comparison of the structures and the carbohydrate-DNA interactions at the two different binding sites explains at the atomic level how calicheamicin derivatives are able to effectively recognize both d(ACCT) and d(TCCT) sites. A number of specific hydrogen-bonding and salt bridge interactions have been identified for both recognition sites. Three sequence-specific intermolecular interactions are present for each DNA site: (a) the H-bond between the carbonyl group connecting carbohydrate B and C rings and the G₁₇-NH₂ (and correspondingly, G₂₁-NH₂ in the other site); (b) the H-bond between B-3OH and A₁₈-N₃ (and T₂₂-O₂); and (c) the interaction between the C ring iodine atom and a guanine amino group G₁₆-NH₂ (and G₂₀-NH₂), whose importance in the binding affinity of these carbohydrates has been well estab-

lished.^{8,27} Thus, intermolecular interactions appear to play an important role in the sequence preferences of DNA binding observed for these compounds.

Structural distortions away from standard geometry are expected in the binding of any structurally complex ligand, as the two molecules optimize their structures to maximize interaction energy. The conformational adaptability of duplex DNA is clearly evident from the wide range of perturbations observed in structures of complexes with proteins and natural product ligands.²⁸ The structural perturbations of the d(GCACCTTCCTGC)·d(GCAGGAAGGTGC) duplex in the HTD (**4**) complex are quite modest overall, with the most extreme effect at C₅ and C₉, whose sugar ring pseudorotation angles shift on the order of 50°. The energy landscape of the

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sugar ring conformations found in duplex DNA is fairly shallow, with the O_{4'}-endo maximum only 1.5–2 kcal/mol less stable than the C_{2'}-endo and C_{3'}-endo minima.²⁶ Thus, the energetic penalty to pay for C₅ and C₉ occupying a less favorable conformation is small and is outweighed by the favorable hydrogen bond, salt bridge, and van der Waals interactions. These are important considerations for understanding the factors directing the propensity for calicheamicins to bind certain DNA sites.

The results reported here and in other structural studies have highlighted the importance of specific and nonspecific intermolecular interactions for the binding of the calicheamicin oligosaccharide in the DNA minor groove. In addition, some degree of induced fit of the DNA site to ligand (and possibly of the ligand to the DNA) is likely. Kahne and co-workers have proposed that DNA conformational adaptability is a primary factor determining the DNA sequence selectivity of calicheamicin.^{4a,c} In this view, binding to the recognition site will be facilitated in regions possessing greater intrinsic conformational flexibility. Unfortunately, there is little if any direct experimental evidence available on the sequence-dependence of conformational propensities and relative flexibility of duplex DNA. Furthermore, in the structure reported here, we observe no substantial distortions from B-DNA geometry, so the requirement for induced fit or indirect read-out appears to be minimal. This is true, even though the two ligands are constrained by their covalent linkage! Thus, we do not interpret the relatively small (50°) shift in sugar pucker as supportive of the “indirect readout” hypothesis. In contrast, the great similarity in the intermolecular interactions in the two binding sites of the HTD complex suggests that sequence selectivity *might* be due to the specific interactions not being available in the requisite locations in DNA duplexes with nonoptimal sequences. Solvation effects may also play an important role.²⁹ Further structural studies of the other high-affinity DNA binding sites and molecular modeling of lower affinity sites are in progress to test this hypothesis and ultimately establish the relative importance of the factors directing the propensity of the calicheamicin oligosaccharide to bind to duplex DNA.

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Conclusions

The NMR solution structure has been determined for the 1:1 complex between the head-to-tail dimer of the calicheamicin oligosaccharide and the duplex oligonucleotide d(G-CACCTTCCTGC)·d(GCAGGAAGGTGC) that contains two different DNA binding sites (ACCT and TCCT). The nanomolar affinity associated with the bidentate mode of binding of the HTD dimer to the DNA duplex combined with the snug fit of the ligand in the DNA minor groove strongly suggest that the HTD molecule possesses a very high degree of complementarity to the duplex containing the dual d(ACCTTCCT) binding site. The ability of each of the oligosaccharide subunits to assume correct positioning for effective binding to the DNA sequence is in turn attributed to the favorable attributes of the linker. Thus, the flexible, open-chain linkers of the HHD (**3**) and HTD (**4**) dimers appear to enable effective localization of each subunit into the respective binding site. On the basis of these results, an effort is currently underway to design corresponding ligands with rigid linkers that, in addition to ensuring correct positioning of each unit, bear specific structural elements for further stabilization of the ligand in the DNA minor groove.

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Supporting Information Available: Three figures showing expanded regions of NOESY spectra of the HTD–DNA complex (Figures S1–S3). Two tables listing helical parameters used to generate initial DNA conformations (Table S1) and all NMR derived distance constraints used in the structure calculation (Table S2) (16 pages, print/PDF). See any current masthead page for ordering information and Web access instructions.

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