Crystal structure of epidoxorubicin–formaldehyde virtual crosslink of DNA and evidence for its formation in human breast-cancer cells

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Epidoxorubicin and daunorubicin are proposed to be cytotoxic to tumor cells by catalyzing production of formaldehyde through redox cycling and using the formaldehyde for covalent attachment to DNA at G bases. The crystal structure of epidoxorubicin covalently bound to a d(CGCGCG) oligomer was determined to 1.6 Å resolution. The structure reveals slightly distorted B-form DNA bearing two molecules of epidoxorubicin symmetrically intercalated at the termini, with each covalently attached from its N3′ to N2 of a G base *via* a CH_2 group from the formaldehyde. The structure is analogous to daunorubicin covalently bound to d(CGCGCG) determined previously, except for additional hydrogen bonding from the epimeric O4′ to O2 of a C base. The role of drug–DNA covalent bonding in cells was investigated with synthetic epidoxorubicin–formaldehyde conjugate (Epidoxoform) and synthetic daunorubicin–formaldehyde conjugate (Daunoform). Uptake and location of drug fluorophore in doxorubicin-resistant human breast-cancer cells (MCF-7/ADR cells) was observed by fluorescence microscopy and flow cytometry. The fluorophore of Daunoform appeared more rapidly in cells and was released more rapidly from cells than the fluorophore of Epidoxoform over a 3 h exposure period. The fluorophore appeared predominantly in the nucleus of cells treated with both conjugates. The difference in uptake is explained in terms of the slower rate of hydrolysis of Epidoxoform to the species reactive with DNA and a proposed slower release from DNA based upon the crystal structures.

1. Introduction

Epidoxorubicin, the 4′-epimer of the clinically important antitumor drug doxorubicin, is a broad-spectrum anthracycline antitumor drug approved for human use in all countries of the world except the USA. A third relevant anthracycline, daunorubicin, is identical to doxorubicin except for the absence of the 14-hydroxyl group. Recent experiments by us and others indicate that all three of these anthracyclines derive at least some, if not most, of their antitumor activity through induction of oxidative stress leading to formaldehyde production. The drugs bind the formaldehyde at their 3′-amino substituents and use this formaldehyde moiety for covalent attachment to DNA at the 2-amino substituents of G bases (Cullinane, Cutts *et al.*, 1994; Cullinane, van Rosmalen *et al.*, 1994; Cutts *et al.*, 1996; Cutts & Phillips, 1995; Taatjes, Fenick, Gaudiano *et al.*, 1998; Taatjes, Gaudiano & Koch, 1997; Taatjes, Gaudiano, Resing *et al.*, 1997; Taatjes *et al.*, 1996; Wang *et al.*, 1991). The crystal structure of daunorubicin covalently attached to the self-complementary DNA oligomer

d(CGCGCG) *via* formaldehyde was solved even before drug-induced formaldehyde production was discovered (Wang *et al.*, 1991), and the structure was important in the elucidation of the mechanism. The drug–DNA covalent-binding mechanism then led to the synthesis of daunorubicin–, doxorubicin– and epidoxorubicin–formaldehyde conjugates, Daunoform (DAUF), Doxoform (DOXF) and Epidoxoform (EPIF), respectively, as improved antitumor drugs (Fenick *et al.*, 1997; Taatjes, Fenick & Koch, 1998). Of particular importance is that the conjugates are significantly more active against resistant cancer cells than the corresponding clinical drugs (Fenick *et al.*, 1997; Taatjes, Fenick & Koch, 1998). In each case, the formaldehyde conjugate is proposed to be a prodrug of the active metabolite of the clinical drug, with the metabolite released through partial hydrolysis of the conjugate. Structures for the clinical drugs, the formaldehyde conjugates and the active metabolites are shown in Fig. 1.

Of the three conjugates, Epidoxoform has emerged as the lead structure for further development because it is hydrolytically more stable with respect to release of formaldehyde (Fenick *et al.*, 1997; Taatjes, Fenick & Koch, 1998). Consequently, Epidoxoform or the product of its partial hydrolysis has a longer potential residence time in the vascular system. On this basis, the crystal structure of epidoxorubicin covalently attached to d(CGCGCG) *via* formaldehyde was solved and is reported here. The structure is compared with that of daunorubicin covalently attached to d(CGCGCG) (Wang *et al.*, 1991) and the structure of epidoxorubicin intercalated in d(CGATCG) (Williams *et al.*, 1990). Both structures are discussed in terms of uptake of Epidoxoform and Daunoform by resistant breast-cancer cells.

2. Materials and methods

2.1. Materials

All tissue-culture materials were obtained from Gibco Life Technologies (Grand Island, NY) unless otherwise stated. MCF-7/ADR breast-cancer cells were a gift from Dr William W. Wells (Michigan State University). Daunoform (DAUF) and Epidoxoform (EPIF) were synthesized from daunorubicin and epidoxorubicin, respectively, by reaction with formaldehyde, as described previously (Fenick *et al.*, 1997; Taatjes, Fenick & Koch, 1998). DAPI (4,6-diamidino-2-phenylindole), spermine and sodium cacodylate were from Sigma Chemical Co. DNA oligonucleotide was obtained from Integrated DNA Technologies (Coralville, IA) and purified by reverse-phase HPLC as described previously (Taatjes, Guadiano, Resing *et al.*, 1997). Formaldehyde was obtained from Mallinckrodt as a 37%(w/w) solution in water containing 10–15% methanol; barium chloride was obtained from Fisher Chemical Co.; 2-methyl-2,4-pentanediol (MPD) was from Aldrich Chemical Co. Water was distilled and purified with a Millipore Q-UF Plus purification system to 18 MΩ cm.

2.2. X-ray crystallography

Crystallization trials were based upon conditions described for daunorubicin–CH$_2$–d(CGCGCG) (Wang *et al.*, 1991). The crystallization mixture consisted of 1.2 mM d(CGCGCG), 1.2 mM epidoxorubicin and 9 mM formaldehyde in pH 6.0 buffer containing 30 mM sodium cacodylate, 16.1 mM spermine and 4.4 mM BaCl$_2$. Tetragonal crystals were grown in sterile 24-well plates (ICN Biomedicals, Aurora, OH) by the hanging-drop method on silanized cover slips. A 4 µl drop of
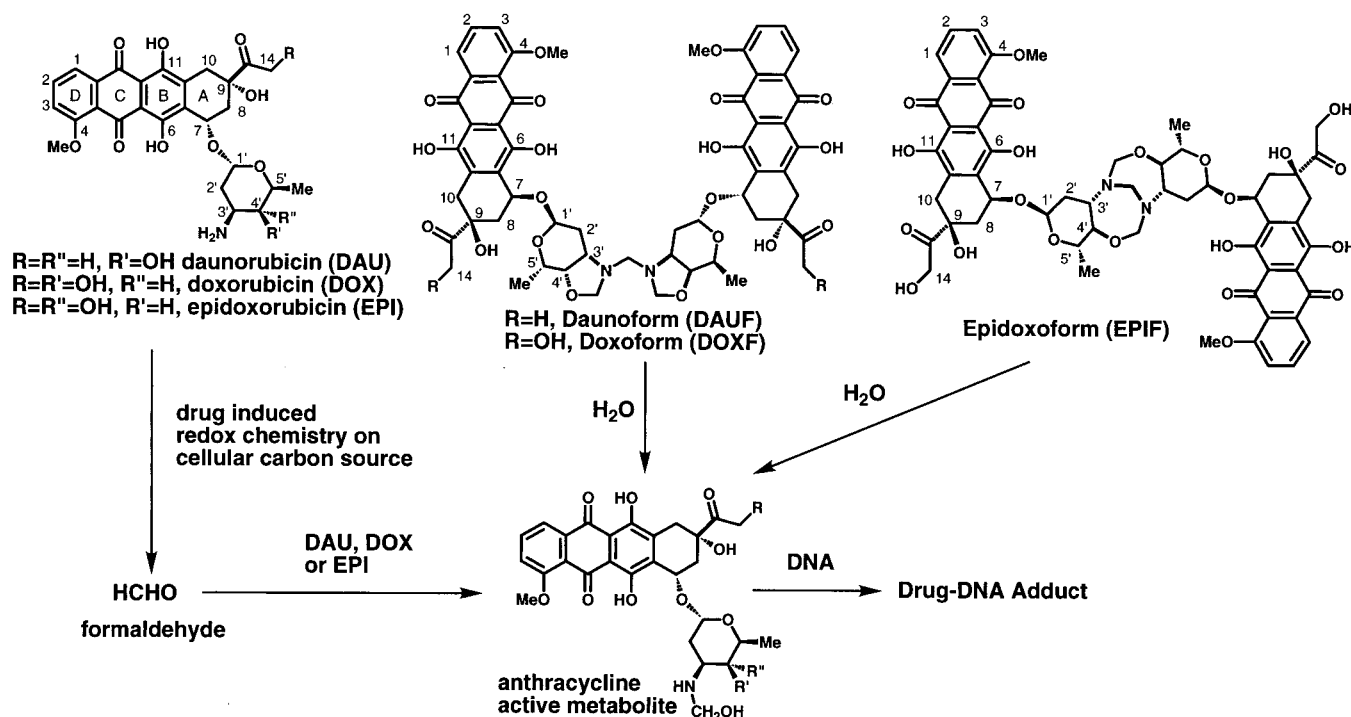

Figure 1
Structures of clinical anthracycline antitumor drugs, their metabolites and their conjugates with formaldehyde.

Table 1
Crystallographic parameters.

Data-collection and refinement statistics.

Unit-cell dimensions (Å, °)	$a = 28.11$, $b = 28.11$, $c = 52.54$; $\alpha = 90$, $\beta = 90$, $\gamma = 90$
Space group	$P4_12_12$
Temperature (K)	295
Number of collected reflections to 1.6 Å	47174
Number of unique reflections to 1.6 Å	3101
Number of reflections used for refinement [$F > 2\sigma(F)$]	2054†, 2612‡
R_{sym}§ (%)	9.9
R factor (%)	20.52†, 21.64‡
R_{free} (%)	26.78†
R.m.s. deviation of bonds from ideality (Å)	0.0197
R.m.s. deviation of angles from ideality (°)	2.087
Average B value (Å^2)	19.37
Number of observed solvent molecules	30
Overdeterminacy ratio	2.9†, 3.7‡

Number of reflections, R_{sym} and completeness by resolution.

Resolution range (Å)	No. of reflections	R_{sym} (%)	Completeness (%)
99.0–3.45	338	7.5	90.0
3.45–2.74	302	9.2	95.0
2.74–2.39	303	11.1	96.2
2.39–2.17	287	12.1	94.1
2.17–2.02	285	13.4	94.4
2.02–1.90	281	14.3	92.1
1.90–1.80	272	18.4	92.5
1.80–1.72	282	22.8	92.5
1.72–1.66	256	26.8	89.8
1.66–1.60	2482	30.9	82.7
All hkl	2854	9.9	92.0

† After sequestering 17.3% of the data. ‡ With all the data. § $R_{sym} = \sum[\text{Abs}(I - \langle I \rangle)] / \sum I$.

the crystallization mixture was placed on a cover slip and the drop was sealed over a 1 ml well solution containing 40% MPD in water. Crystals were allowed to grow at ambient temperature; crystal formation was complete in about 6 d. The crystal was mounted and sealed in a glass capillary tube, and diffraction data were collected on an R-AXIS II image-plate system using Cu $K\alpha$ radiation from a Rigaku RU-H2R rotating-anode X-ray generator (Molecular Structures Corp., Houston, TX). The data were reduced and scaled using the HKL package (Gewirth, 1997; Otwinowski & Minor, 1997) and are summarized in Table 1. Epidoxorubicin–CH$_2$–d(CGCGCG) crystals belong to the space group $P4_12_12$, like many anthracycline–hexanucleotide complexes, and have unit-cell parameters $a = b = 28.11$, $c = 52.54$ Å, $\alpha = \beta = \gamma = 90°$. Useful diffraction was seen to a minimum Bragg spacing of 1.6 Å.

Crystallographic refinement and σ_A-weighted electron-density map calculations were carried out using CNS (Brünger et al., 1998), with a new parameter dictionary for standard DNA nucleotides (Parkinson et al., 1996). The initial model was constructed from the coordinates of daunorubicin covalently linked to d(CGCGCG) with formaldehyde (PDB entry 1d33) modified to include the position of O14 from epidoxorubicin (PDB entry 1d54). All refinement was performed on

$|F_o| > 2\sigma|F_o|$. In order to obtain a statistically significant set of reflections for R_{free} calculations (Brünger & Rice, 1997), 529 reflections (17.3%) were sequestered from the data set. Initial refinement used the R_{free}, while the final series of conjugate-gradient minimizations utilized the entire data set in order to include a significant percentage (84.5%) of the theoretical data to 1.6 Å. The first set of refinement rounds consisted of torsion-angle simulated annealing with the maximum-likelihood target using amplitudes as implemented in CNS, followed by cycles of water picking based on positive peaks in $F_o - F_c$ difference Fourier maps. After each cycle of water picking, conjugate-gradient minimization and individual temperature-factor refinement was carried out and water positions were checked by examining $2F_o - F_c$ and $F_o - F_c$ difference Fourier maps calculated by CNS and displayed on a Silicon Graphics Onyx 2 workstation using O (Jones et al., 1991). After 29 waters were picked, the model was annealed again and the cycles continued with some waters being manually repositioned into density as necessary. In the last cycle, waters with temperature factors greater than 60 Å^2 were deleted. In the final series of refinement cycles, the model was refined against all reflections using the standard crystallographic residual target and a bulk-solvent correction. As in the first set, cycles of conjugate-gradient minimization and individual temperature-factor refinement were performed, followed by scrutinizing the $F_o - F_c$ and $2F_o - F_c$ maps. Additional waters were deleted or repositioned based on the same criteria of lower temperature factors and occupied density. No other ions (Na$^+$, spermine or Ba^{2+}) could be unambiguously identified. The final model contained 30 waters distributed throughout the epidoxorubicin–CH$_2$–d(CGCGCG) asymmetric unit. Crystallographic and refinement data are reported in Table 1.

2.3. Measurement of drug uptake by flow cytometry

MCF-7/ADR cells were maintained in vitro by serial culture in RPMI 1640 medium supplemented with 10% fetal bovine serum (Gemini Bio-Products, Calbasas, CA), L-glutamine (2 mM), HEPES buffer (10 mM), penicillin (100 units ml^{-1}), streptomycin (100 mg ml^{-1}) and 5 µM doxorubicin. Cells were maintained at 310 K in a humidified atmosphere of 5% CO$_2$ and 95% air. Cultured cells were dissociated with trypsin–EDTA and plated in 6-well plates (~500000 cells per well) and allowed to adhere overnight. The cells were treated with 0.25 µM DAUF or EPIF at 310 K for various amounts of time (5 min, 30 min, 1 h, 2 h, 3 h). For each time point, the medium was removed and the cells were trypsinized and suspended in 3 ml of RPMI 1640 medium (−) phenol red and (−) FBS. The cells were transferred to 15 ml conical vials and centrifuged at 1000 r min^{-1} for 5 min at 283 K. The supernatant was removed and replaced with 1 ml of RPMI 1640 medium (−) phenol red and (−) FBS. The cells were kept at 273 K until analysis (up to 3 h). Control experiments showed no significant loss of fluorescence in samples kept at 273 K for up to 4 h. The extent of drug uptake was determined by flow cytometry, as previously described (Durand & Olive, 1981). All flow-

cytometry measurements were made with a Becton Dickinson FACScan flow cytometer, using a Hewlett–Packard 9000 Series Model 340 computer for data storage and analysis. Drug-treated cells were analyzed with excitation at 488 nm (15 mW argon-ion laser), with emission monitored between 570 and 600 nm. Instrument settings were held constant for all experiments and 5000 cells were counted per measurement. The emission of drug-free cells was similarly measured in order to determine background fluorescence. The final data are plotted as mean fluorescence (as determined by computer data analysis) *versus* drug incubation time (or recovery time) for ease of data representation.

2.4. Analysis of intracellular drug distribution by fluorescence microscopy

Cells were plated in 6-well plates (∼300000 cells per well). Each well contained a sterile cover slip, and the cells were allowed to adhere to the cover slip overnight. Each well contained 3 ml of RPMI 1640 medium. The DAUF and EPIF were dissolved in DMSO to a concentration of 25 μM (100× solution). 30 µl of each DMSO solution was then added to the appropriate well, resulting in a 100× dilution (0.25 μM conjugate, 1% DMSO). The cells were incubated with the drug for 3 h. Following drug treatment, the medium was removed and the cells were washed with 2 ml PBS (phosphate-buffered saline). The cells were then fixed to the cover slips by submerging in 3 ml cold (253 K) methanol and storing on ice for 5 min. The methanol was then removed and the cells were washed with 3 ml PBS. The cover slips were then removed from the wells and inverted on a drop (30 µl) of DAPI (0.2 µg ml^{-1} in PBS) placed on parafilm. The cover slips were kept on the DAPI solution for 5 min at ambient temperature. The cover slips were then rinsed with PBS and mounted on a microscope slide using a drop (30 µl) of mowiol mounting medium. The slides were allowed to dry in the dark overnight. Microscopic images were observed at a magnification of 1000× and recorded with a Zeiss Axioplan fluorescence microscope equipped with a Photometrics Sensys digital CCD camera system. Images were manipulated using *IP-LAB Spectrum* software. Drug fluorescence in cells was observed at wavelengths longer than 590 nm with excitation at 546 ± 6 nm. DAPI fluorescence was observed at wavelengths longer than 420 nm with excitation at 355 ± 20 nm.

3. Results and discussion

3.1. Crystal structure of epidoxorubicin covalently bound to d(CGCGCG)

The crystallographic asymmetric unit consists of a single strand of d(CGCGCG) covalently attached to a 4′-epidoxorubicin molecule *via* a methylene group at one of the guanines and 30 ordered water molecules. Two asymmetric units form a slightly distorted B-type DNA duplex with six Watson–Crick base pairs. Fig. 2 shows [4′-epidoxorubicin–CH$_2$–d(CGCGCG)]$_2$ with the DNA displayed as a CPK structure and the drug as a framework structure. As observed in other anthracycline–DNA structures, the aglycon rings of epidoxorubicin are intercalated at CpG steps, with the D ring reaching into the major groove while the sugar moiety lies in the minor groove. The intercalation sites are C(1)G(2)/ G(12)C(11) and C(5)G(6)/G(8)C(7). [The nucleotides labeled C(1)–G(6) are on a strand in one asymmetric unit, with C(7)–G(12) located on the complementary strand generated by twofold crystallographic symmetry]. Two methylene groups link the 3′-amino substituents of two epidoxorubicins to the 2-amino substituents of G(10) and G(4). Fig. 3 shows a stereoscopic view of the $F_o - F_c$ electron-density map superimposed on the final model of 4′-epidoxorubicin–CH$_2$–d(CGCGCG).

Comparison of the structure with the covalent daunorubicin–CH$_2$–d(CGCGCG) (Wang *et al.*, 1991) and the non-covalent 4′-epidoxorubicin–d(CGATCG) structures (Williams *et al.*, 1990) is instructive and is summarized in Table 2 and Fig. 4. The general coordinate error based on the Luzzati plot for our model is 0.20 Å. Of particular significance are hydrogen-bonding interactions between the drug and the DNA. The O9 hydroxyl groups of the drugs in all three structures are hydrogen bonded to N2 and N3 of G(8) with approximately the same bond distances (3.12–3.15 and 2.89–2.59 Å, respectively). The
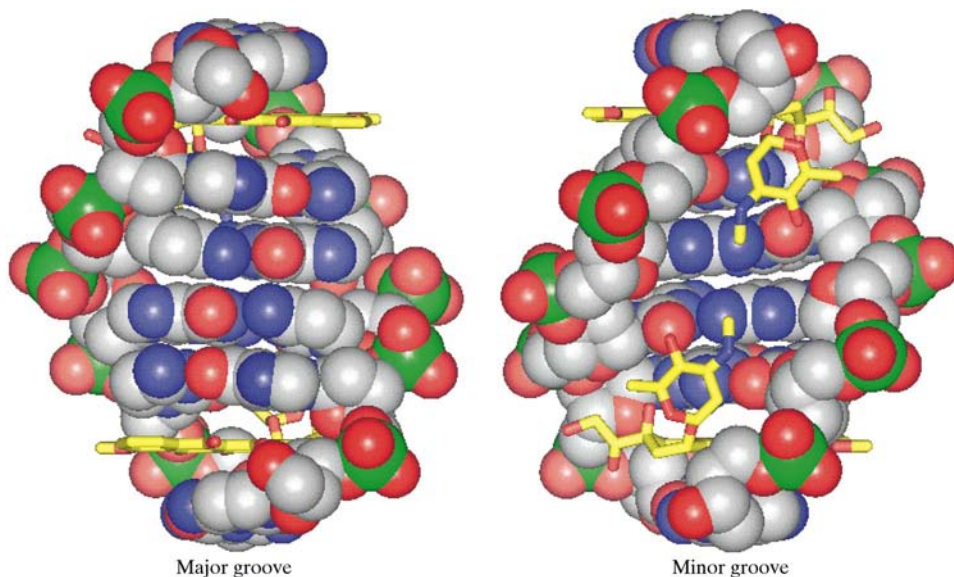

Figure 2
Epidoxorubicin–CH$_2$–d(CGCGCG), with DNA displayed as a CPK structure and epidoxorubicin–CH$_2$ displayed as a framework structure. This figure was created with *Insight*II (Biosym Technologies, San Diego, CA).

Table 2
Distances.

Drug atom	DNA atom	Water atom†	Distance (Å)		
			Epidox–CH₂–d(CG)₃‡	Dauno–CH₂–d(CG)₃	Epidox–d(CGATCG)
O9	G(8) N3		2.89	2.86	2.59
	G(8) N2		3.12	3.12	3.15
O7	G(8) N2		2.87	3.13	3.35
O14	C(9) C5′		3.70	—	—
	A(9) C5′		—	—	3.40
O4′	C(9) O2		2.70	>4.0	—
	A(9) N3		—	—	3.06
N3′	G(4) N2		2.50	2.48	—
	T(4) O2		—	—	3.37
	C(5) O2		3.33	3.24	3.27
	C(5) O4′		3.55	3.16	3.31
	CH₂		1.52	1.53	—
CH₂	G(4) N2		1.46	1.45	—
O4		W(1)	2.81	3.06	3.02
O5		W(1)	3.02	3.21	3.11
	G(6) N7	W(1)	2.69	2.72	2.83
O12		W(25)	2.90	3.81	>4.0
	G(8) N7	W(25)	3.07	3.01	2.61
O13		W(6)	2.89	3.11	3.34
	C(7) O2	W(6)	2.84	2.60	2.65

† Naming convention from epidoxorubicin–CH₂–d(CGCGCG) structure. ‡ Luzzati coordinate error = 0.20 Å.

distance between the anomeric O7 of the drug and the N2 of G(8) is shorter in epidoxorubicin–CH₂–d(CGCGCG) than in the other structures (2.87 *versus* 3.13 and 3.35 Å), suggesting that there is a stronger hydrogen bond between these atoms. The primary structural difference between epidoxorubicin and daunorubicin is the configuration of the hydroxyl at the 4′-position. This leads to a good hydrogen-bonding interaction (2.70 Å) with the O2 of C(9) on the complementary strand in the structure of epidoxorubicin–CH₂–d(CGCGCG). In epidoxorubicin–d(CGATCG), this hydrogen bonding is with the N3 of the adenine on the complementary strand. Perhaps the extra hydrogen bonding causes the aglycon A ring of the drug to lie closer to G(8) on the complementary strand in epi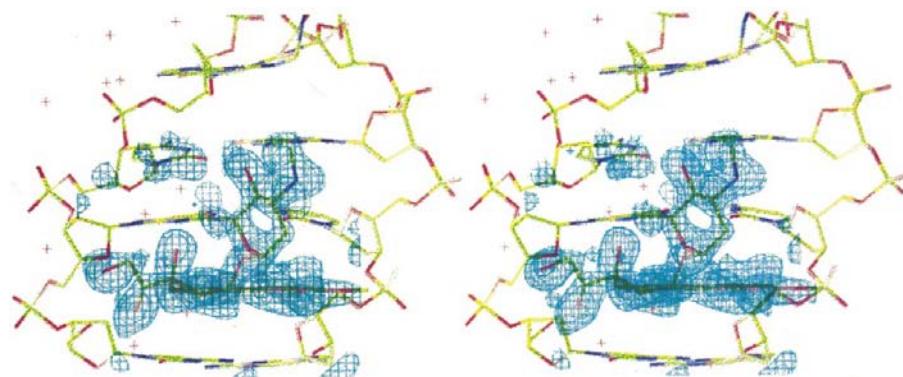doxorubicin–CH₂–d(CGCGCG). The 14-hydroxyl, present in the epidoxorubicin structures but not in the daunorubicin structure, lies farther away from the C5′ of C(9) on the complementary strand in the covalent structure than in the non-covalent structure (3.70 *versus* 3.40 Å). Differences in the A ring, O7, O14 and aminosugar of the drugs in the three structures are shown *via* an overlay diagram in Fig. 5. The alignment was performed using all the common atoms, including the atoms of the nucleic acid moieties, between epidoxorubicin–CH₂–d(CGCGCG) as the reference molecule and either daunorubicin–CH₂–d(CGCGCG) or epidoxorubicin–d(CGATCG). For clarity, the superimposed DNA from the structures was omitted from Fig. 5. The O12 is within hydrogen-bonding distance (2.90 Å) of a water molecule, W(25), which is associated with the N7 of guanine G(8) (3.07 Å). [Solvent water molecules are labeled W(1)–W(30).] While the other two structures also have a water molecule near N7 of G(8) (3.01 or 2.61 Å), it is too far away from O12 (3.81 or >4.01 Å) to mediate contact between the drug and the DNA. All of these drug–DNA and drug–water molecule contacts are summarized in Table 2 and Fig. 4.

The drug–DNA adducts are hydrolytically unstable, and the drug is released at two different first-order rates, one with a half-life of 5 h and the other of 40 h (Cullinane, van Rosmalen *et al.*, 1994; van Rosmalen *et al.*, 1995). Prior to an understanding of the molecular nature of the covalent bonding, the drug released more slowly was thought to be covalently attached to both strands of the DNA at the G bases of a GC site. The crystal structure reported here and the earlier structure of daunorubicin covalently bound to d(CGCGCG) (Wang *et al.*, 1991) provide a simple explanation for slower release of drug from a GC site, even though the drug has only one covalent attachment. In addition to the covalent attachment to a G on one strand, the drug is strongly hydrogen bonded between its 9-hydroxyl and the N atoms at the 2- and 3-positions of the guanine on the complementary strand. The [epidoxorubicin–CH₂–d(CGCGCG)]₂ structure also suggests hydrogen bonding between O7 of the drug and N2 of G(8) and between O4′ of the drug and O2 of C(9) on the complementary strand. Because the original proposal for the structure of more stable lesions was thought to be a crosslink of DNA by the drug, we proposed the term virtual crosslink for this bonding, which is unique to the GC dinucleotide. The less stable DNA–drug adducts are then proposed to have covalent attachment to isolated G bases *via* methylene groups, and to be missing the hydrogen-bonding interactions with the opposing strand.

The virtual crosslink of the self-complementary hexanucleotide d(ATGCAT) at the TGC site from reaction with a single doxorubicin and formaldehyde has recently been characterized using NMR spectroscopy (Zeman *et al.*, 1998). This study also reported the effect of the virtual cross-

Figure 3
Stereoscopic view of an $F_o - F_c$ electron-density map superimposed on the final model of epidoxorubicin–CH₂–d(CGCGCG). All atoms of the drug were omitted during the map calculation using only the coordinates in the final model of the DNA molecule for the phase information. The density is contoured at 2σ. Bonds and atoms are colored according to atom type.

link on the stability of duplex DNA from measurements of the rate of DNA-strand exchange. Strand exchange was 3.9 times slower with a doxorubicin merely intercalating in an oligonucleotide but 637 times slower with a doxorubicin virtually crosslinking the oligonucleotide.

3.2. Uptake and distribution of drug in MCF-7/ADR cells

MCF-7/ADR cells are mutants of the MCF-7 human breast-cancer cell line which are highly resistant to doxorubicin (Adriamycin). They are also resistant to daunorubicin and epidoxorubicin but not to Daunoform, Epidoxoform or Doxoform (Fenick *et al.*, 1997; Taatjes, Fenick & Koch, 1998). Important resistance mechanisms are increased activity of enzymes which neutralize the effects of drug-induced oxidative stress and overexpression of the drug efflux pump, P-170 glycoprotein (Pgp) (Mimnaugh *et al.*, 1991; Sinha & Mimnaugh, 1990). This cell line, therefore, is particularly useful for studying the effect of the conjugation of the drug with formaldehyde, because daunorubicin and epidoxorubicin are maintained at lower concentrations by drug efflux. The efflux pump recognizes the drugs partly because of the positive charge at the amino substitutent (Lampidis *et al.*, 1997). Formaldehyde conjugates are uncharged because of a change in pK_a resulting from the hydroxymethylene substituent and hence are predicted to be poorer substrates for Pgp. Further, formation of active metabolite by the clinical drug is enzymatically inhibited in MCF-7/ADR cells.

Uptake of Daunoform and Epidoxoform by MCF-7/ADR cells at 310 K as a function of time of exposure to 0.25 μ*M* drug was measured by flow cytometry (Fig. 6). Drug fluorophore in cells was detected by its fluorescence in the region of 590 nm. The fluorophore in cells treated with Daunoform appeared rapidly, with the maximum level occurring after approximately 7 min, followed by a steady decrease during the
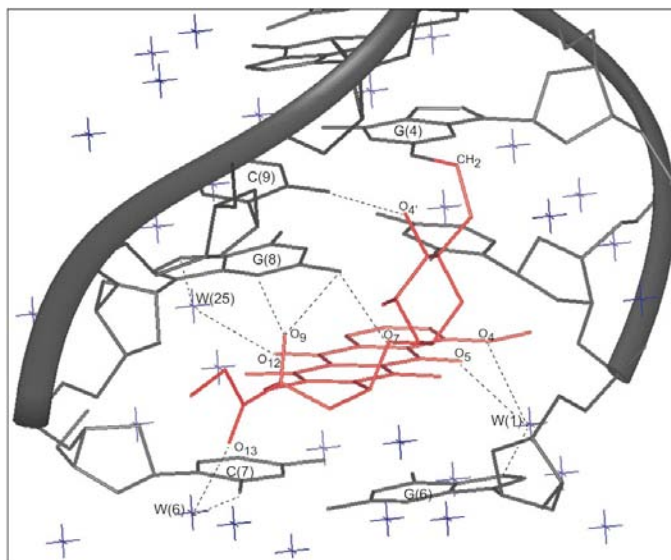
remaining 3 h exposure period. In contrast, fluorophore in cells treated with Epidoxoform steadily increased over the 3 h exposure period. The rates of uptake parallel the rates of hydrolysis of conjugates to the active metabolites of the respective clinical drugs (Fig. 1). Hydrolysis of Daunoform to the active metabolite occurs with a half-life of less than 10 min in cell-culture medium at 298 K (Fenick *et al.*, 1997), while hydrolysis of Epidoxoform to the active metabolite occurs with a half-life of 2 h at 310 K in cell-culture medium (Taatjes, Fenick & Koch, 1998). Further, the half-lives for hydrolysis of the active metabolites to daunorubicin and epidoxorubicin are less than 10 min and greater than 2 h, respectively. This parallel suggests that the slow step for drug uptake by MCF-7/ADR cells is hydrolysis to the active metabolite.

The location of drug fluorophore in the nucleus of MCF-7/ADR cells treated with either Daunoform or Epidoxoform was established by fluorescence microscopy, as shown in Fig. 6. Anthracycline fluorescence was primarily in the nucleus, as indicated by co-staining with the nuclear stain DAPI. The abundance of anthracycline in the nucleus is even higher than indicated by the micrographs because DNA partially quenches the fluorescence (Crooke & DuVernay, 1980; Roche *et al.*, 1994). Since the anthracycline is predominantly in the nucleus, the release of fluorophore from cells exposed to Daunoform after 7 min of drug treatment can be explained by hydrolysis of drug covalently bound to DNA occurring faster than new covalent binding of drug metabolite. In fact, after about 30 min most of the Daunoform in the cell-culture medium has hydrolyzed to daunorubicin, which is not significantly taken up by MCF-7/ADR cells because of the Pgp efflux pump. Further, formation of the active metabolite from daunorubicin is inhibited in these cells by the increased activity of enzymes which neutralize oxidative stress. The rate of fluorophore release over the 3 h time period is most consistent with predominant hydrolysis of drug–DNA lesions at isolated G bases. This hydrolysis releases daunorubicin which is pumped out of the cell. Steady increase of fluorophore in MCF-7/ADR cells upon treatment with Epidoxoform over the 3 h time period is a consequence of much slower
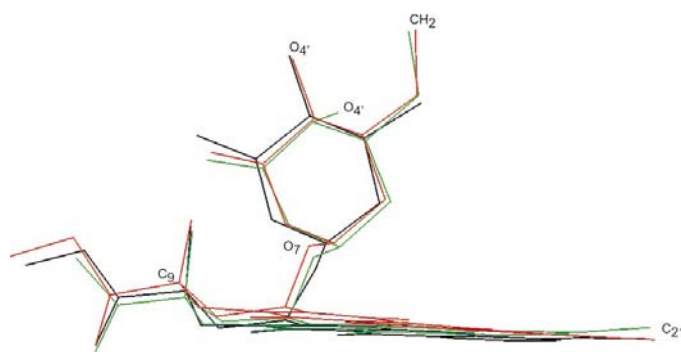

Figure 4
Hydrogen-bonding contacts between epidoxorubicin–CH$_2$ and d(CGCGCG). This figure was created with *Insight*II (Biosym Technologies, San Diego, CA).

Figure 5
Overlay of epidoxorubicin (in black) and daunorubicin–CH$_2$ (in green) on epidoxorubicin–CH$_2$ (in red). R.m.s.d. in distance of all atoms common to epidoxorubicin–CH$_2$–d(CGCGCG) and daunorubicin–CH$_2$–d(CGCGCG) is 0.923 Å and between epidoxorubicin–CH$_2$–d(CGCGCG) and epidoxorubicin–d(CGATCG) is 0.829 Å. For clarity, the overlaid DNA strands have been omitted from the figure.

hydrolysis to the active metabolite and increased stability of the metabolite with respect to further hydrolysis to epidoxorubicin. Hence, covalent bonding of drug to DNA is occurring faster than hydrolysis of drug–DNA lesions. Additional evidence for covalent binding of drug to DNA in cells comes from a comparison of release of drug from cells treated with
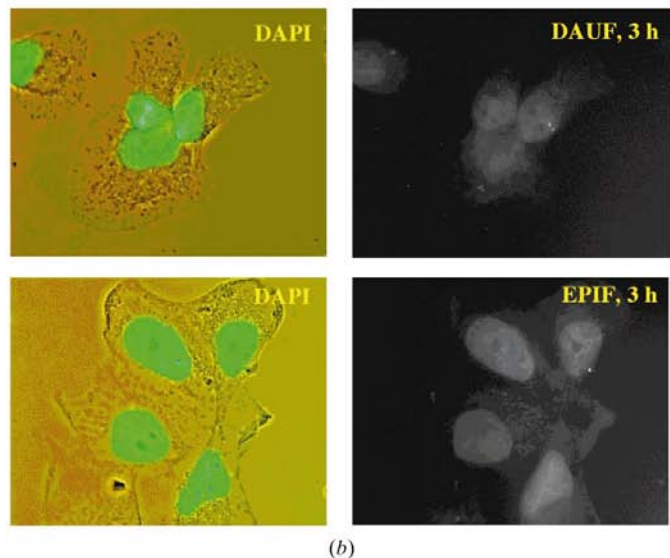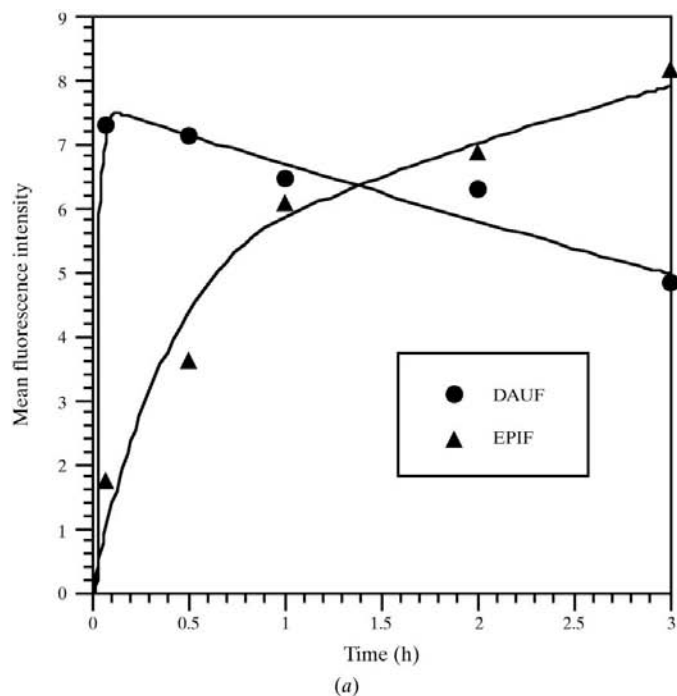

Figure 6

(*a*) Uptake of the fluorophore of Daunoform (DAUF) or Epidoxoform (EPIF) by MCF-7/ADR cells as a function of time of exposure to 0.25 μ*M* drug, as determined by flow cytometry measuring fluorescence intensity at 570–600 nm. (*b*) Fluorescence micrographs of MCF-7/ADR cells after treatment with either 0.25 μ*M* Daunoform or Epidoxoform for 3 h. After drug treatment, cells were fixed and stained with the nuclear stain DAPI. Each micrograph showing DAPI fluorescence in green superimposed on the whole cell image is located adjacent to the respective micrograph showing drug fluorophore emission.

either parent drug or formaldehyde conjugate and from analysis of cells treated with tritiated formaldehyde conjugate (Taatjes *et al.*, 1999).

The crystal structure provides some additional insight with respect to drug uptake by MCF-7/ADR cells. The additional hydrogen-bonding interaction between the epidoxorubicin's 4′-hydroxyl and the O2 of C(9) on the complementary strand should lead to slower hydrolysis of the virtual crosslink of DNA with epidoxorubicin than with daunorubicin. This slower release from DNA gives rise to continued uptake as Epidoxoform is slowly hydrolyzing to the active metabolite. Daunoform hydrolyzes to its active metabolite very rapidly, which causes a rapid buildup of drug in the nucleus of tumor cells. The intracellular levels then also decrease because of the predicted less stable daunorubicin virtual crosslink.

In summary, the 1.6 Å crystal structure of [epidoxorubicin–CH$_2$–d(CGCGCG)]$_2$ shows a slightly distorted B-type DNA duplex symmetrically bearing two intercalated epidoxorubicin molecules located at the termini which are covalently attached to the DNA *via* methylene groups. Covalent attachment is from the 3′-amino group of epidoxorubicin to the 2-amino group of G(4) or G(10). The structure is analogous to the structure of [daunorubicin–CH$_2$–d(CGCGCG)]$_2$, except for additional hydrogen bonding from the 4′-hydroxyl group of epidoxorubicin to the O2 of C(3) or C(9). This hydrogen bonding, together with hydrogen bonding between O7 and O9 of the drug and N2 and/or N3 of G(2) or G(8) and the covalent attachment, result in a virtual crosslinking of DNA by drug. Flow cytometry and fluorescence microscopy show that daunorubicin–formaldehyde conjugate (Daunoform) and epidoxorubicin–formaldehyde conjugate (Epidoxoform) are readily taken up by doxorubicin-resistant human breast-cancer cells and locate predominantly in the nucleus. Differences in the rates of uptake are consistent with differences in the rates of hydrolysis of the conjugates to the proposed DNA-reactive form of the drugs and differences in the respective structures of the drugs covalently bound to DNA.

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