

Cobalt(III), nickel(II) and ruthenium(II) complexes of 1,10-phenanthroline family of ligands: DNA binding and photocleavage studies

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Abstract. DNA binding and photocleavage characteristics of a series of mixed-ligand complexes of the type $[M(\text{phen})_2\text{LL}]^{n+}$ (where $M = \text{Co(III)}, \text{Ni(II)}$ or Ru(II) , $\text{LL} = 1,10\text{-phenanthroline (phen), phenanthroline-dione (phen-dione)}$ or $\text{dipyridophenazine (dppz)}$ and $n = 3$ or 2) have been investigated in detail. Various physico-chemical and biochemical techniques including UV/Visible, fluorescence and viscometric titration, thermal denaturation, and differential pulse voltammetry have been employed to probe the details of DNA binding by these complexes; intrinsic binding constants (K_b) have been estimated under a similar set of experimental conditions. Analysis of the results suggests that intercalative ability of the coordinated ligands varies as $\text{dppz} > \text{phen} > \text{phen-dione}$ in this series of complexes. While the Co(II) and Ru(II) complexes investigated in this study effect photocleavage of the supercoiled pBR 322 DNA, the corresponding Ni(II) complexes are found to be inactive under similar experimental conditions. Results of detailed investigations carried out inquiring into the mechanistic aspects of DNA photocleavage by $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$ have also been reported.

Keywords. Phenanthroline family of ligands; metal complexes; DNA binding and photocleavage.

1. Introduction

Current burgeoning interest in small molecules that are capable of binding and cleaving DNA is related to their utility in the design and development of synthetic restriction enzymes, new drugs, DNA footprinting agents etc. and also to their ability to probe the structure of DNA itself^{1,2}. In this regard, metal complexes have been found to be particularly useful because of their potential to bind DNA *via* a multitude of interactions and to cleave the duplex by virtue of their intrinsic chemical, electrochemical and photochemical reactivities^{3–10}. Prominent among the various metal complexes employed so far in studies with DNA are those metallo–intercalators which incorporate either 1,10-phenanthroline (phen) or a modified phenanthroline moiety as a ligand. A singular

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advantage in using these metallo-intercalators for such studies is that the ligands or the metal in them can be varied in an easily controlled manner to facilitate an individual application.

We have been interested to know the effect of variation of the metal ion and also the ligand on the ability to bind and photocleave DNA in mixed ligand complexes containing the phenanthroline family of ligands¹¹⁻¹³. During our studies, it occurred to us that complexes of the type $[M(\text{phen})_2(\text{LL})]^{n+}$ where M is a transition metal ion and LL is a modified phenanthroline ligand are well-suited for this purpose. Although DNA interactions of a number of $[M(\text{phen})_2(\text{LL})]^{n+}$ type complexes have previously appeared in the literature, relatively less attention seems to have been paid to systematic investigations inquiring into the effects brought about by changing M and LL in such complexes. In this paper, we compare the DNA binding and photocleavage characteristics of a family of $[M(\text{phen})_2(\text{LL})]^{n+}$ type complexes (M = Co(III), Ni(II) or Ru(II), LL = phen, phenanthroline-dione (phen-dione) or dipyridophenazine (dppz) and $n = 3$ or 2) (figure 1). Various physico-chemical and biochemical techniques including UV/Visible-, fluorescence-, and viscometric titration, thermal denaturation, differential pulse voltammetry and gel electrophoresis have been utilized to probe the nature of interaction of these complexes with the duplex. Also reported in this paper is a detailed mechanistic investigation on the DNA photocleavage by $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$.

2. Experimental

All common chemicals, solvents as well as cobalt(II), nickel(II) and ruthenium(III) salts, 1,10-phenanthroline monohydrate and 1,2-diaminobenzene were purchased either from BDH (Mumbai, India) or Merck (Mumbai, India). All the solvents were purified before use as per the standard procedures¹⁴. Deionized, triply distilled water was used for preparing various buffers. Calf thymus DNA (CT DNA), 1,4-diazabicyclo(2.2.2.)octane (DABCO), superoxide dismutase (SOD), phenyl-*t*-butylnitron (PBN), D₂O, tetrabutylammonium chloride (TBACl) and tetrabutylammonium hexafluorophosphate (TBAPF₆) were obtained from Sigma Chemicals, USA. The supercoiled pBR 322 DNA (CsCl purified, Bangalore Genie, Bangalore, India) was used as received. Agarose

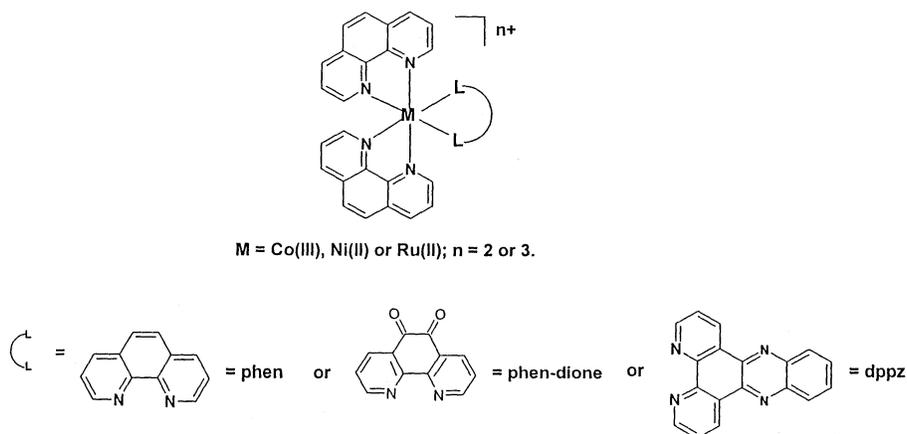


Figure 1. Structures of the $[M(\text{phen})_2(\text{LL})]^{n+}$ type complexes investigated in this study.

(molecular biology grade) and ethidium bromide (EtBr) were purchased from Bio-Rad, USA.

2.1 Synthesis

Ligands phen-dione and dppz were synthesized as per the reported procedures^{15,16} [Co(phen)₃]³⁺¹⁷, [Co(phen)₂Cl₂]¹⁸, [Co(phen)₂Cl₂]⁺¹⁹, [Co(phen)₂(phen-dione)]²⁺²⁰, [Co(phen)₂(dppz)]²⁺¹¹, [Ni(phen)₃]²⁺²¹, [Ni(phen)₂Cl₂]²², [Ni(phen)₂(dppz)]²⁺¹¹, [Ru(phen)₃]³⁺²³, [Ru(phen)₂Cl₂]²⁴, [Ru(phen)₂(phen-dione)]²⁺²⁵ and [Ru(phen)₂(dppz)]²⁺^{25,26} were prepared employing methods available in the literature. [Ni(phen)₂(phen-dione)]²⁺ was prepared following a procedure analogous to that adopted for the synthesis of [Ru(phen)₂(phen-dione)]²⁺²⁵. The hexafluorophosphate salts of the synthesized complexes have been recrystallized by dissolving them in the minimum volume of acetone and reprecipitating, after filtration, by the addition of ether. All these complexes gave correct elemental analysis consistent with their molecular formulae.

The hexafluorophosphate salts of the complexes have been converted to the water-soluble, chloride salts by treating the former salt solutions with excess TBACl in acetone. The chloride salts, being insoluble in acetone, were instantaneously precipitated out of the solution. They were filtered and vacuum dried before use.

2.2 Physical methods

UV/Visible and IR spectra were recorded with a Shimadzu model UV-160 A (coupled with a temperature controller Model TCC-240A) and a JASCO Model 5300 FT-IR spectrophotometer, respectively. Fluorescence spectra were recorded with a JASCO Model 7700 spectrofluorometer for solutions having an absorbance less than 0.2 at the excitation wavelength. [Ru(phen)₃]²⁺ was used as the standard for this purpose (ϕ , = 0.028 in CH₃CN)²⁷. ¹H NMR spectra were recorded with a Bruker NR-FT 200 spectrometer using DMSO-*d*₆/CDCl₃ as the solvent and tetramethylsilane (TMS) as an internal standard. ESR spectra were recorded with JEOL JM-FE3X spectrometer with diphenylpicrylhydrazide (DPPH) as an ESR standard. Magnetic susceptibility measurements for solid samples of the complexes were carried out using a Cahn Instruments (Model 6612) system. CuSO₄ and Hg[Co(CNS)₄] were employed as magnetic susceptibility standards. Diamagnetic corrections to the apparent magnetic susceptibility values have been incorporated, as specified²⁸. Cyclic voltammetric and differential pulse voltammetric experiments were performed on a Princeton Applied Research (PAR) 174A polarographic analyzer coupled with a PAR 175 universal programmer and a PAR RE 0074 X-Y recorder. A platinum-button working electrode, a platinum-wire counter electrode and a saturated calomel reference electrode (SCE) were employed for experiments involving the nonaqueous solvents (PF₆ salts of the complexes). The SCE was separated from the deaerated (N₂) bulk electrolytic solution by a fritted-glass-disc-junction containing the solvent (CH₃CN) and the supporting electrolyte (TBAPF₆). Ferrocene was used as an internal standard for these experiments.

2.3 Studies with DNA

Concentration of CT DNA was measured by using its known extinction coefficient at 260 nm (6600 M⁻¹ cm⁻¹)²⁹. BufferA (5 mM *tris*, pH 7.1, 50 mM NaCl) was used for

absorption titration experiments and luminescence measurements, bufferB (1 mM phosphate, pH 7.0, 2 mM NaCl) was used for thermal denaturation and differential-pulse voltammetric experiments. BufferC (1.5 mM Na₂HPO₄, 0.5 mM NaH₂PO₄, 0.25 mM Na₂EDTA, pH = 7.0) was used for the viscometric titrations. The chloride salts of the complexes were used in studies with DNA.

2.3a Absorption titration experiments: These experiments were performed by maintaining a constant concentration of the complex while varying the nucleic acid concentration. This was achieved by dissolving an appropriate amount of the metal complex in the DNA stock solution and by mixing various proportions of the metal complex and DNA stock solutions while maintaining the total volume constant (1 ml). This resulted in a series of solutions with varying concentrations of DNA but with a constant concentration of the complex. The absorbance (*A*) of the most red-shifted band of each investigated complex was recorded after successive additions of CT DNA. The intrinsic binding constant, K_b , was determined from the plot of $[\text{DNA}]/(\epsilon_a - \epsilon_f)$ vs $[\text{DNA}]$, where $[\text{DNA}]$ is the concentration of DNA in base pairs, ϵ_a , the apparent extinction coefficient is obtained by calculating $A_{\text{obsd}}/[\text{complex}]$ and ϵ_f corresponds to the extinction coefficient of the complex in its free form. The data were fitted to (1) where ϵ_b refers to the extinction coefficient of the complex in the fully bound form.

$$[\text{DNA}]/(\epsilon_a - \epsilon_f) = [\text{DNA}]/(\epsilon_b - \epsilon_f) + 1/K_b(\epsilon_b - \epsilon_f). \quad (1)$$

Each set of data, when fitted to the above equation, gave a straight line with a slope of $1/(\epsilon_b - \epsilon_f)$ and a y-intercept of $1/K_b(\epsilon_b - \epsilon_f)$. K_b was determined from the ratio of the slope to intercept. An in-house nonlinear least square analysis program or the MicroCal Origin software package run on an IBM-compatible Pentium 166 computer was used for curve-fitting the data.

2.3b Fluorescence titration experiments: A procedure analogous to that used for the absorption titration experiment was used in these experiments. The concentration of the Ru(II) complex employed, however, was between about 10^{-6} and 10^{-5} M and that of DNA was between about 10^{-6} and 10^{-3} M (base pairs).

2.3c Thermal denaturation studies: DNA melting experiments were carried out by monitoring the absorption (260 nm) of CT DNA (160 μM) at various temperatures, in both the absence and the presence (0–10 μM) of each investigated complex. The melting temperature (T_m) and the curve width σ_T (= temperature range between which 10% to 90% of the absorption increase occurred) were calculated as described³⁰. The shape of the melting curves, T_m and σ_T values for CT-DNA and for CT-DNA in the presence of $[\text{Ru}(\text{phen})_3]^{2+}$ were consistent with the literature data³⁰. Some of the metal complexes were seen to absorb at 260 nm, but control experiments suggested that this absorption is independent of temperature.

2.3d Viscometry: Viscometric titrations were performed with a Cannon-Ubbelohde viscometer at $25 \pm 1^\circ\text{C}$. Each compound (3–40 μM) was introduced into the degassed DNA solution (300 μM in base-pairs) present in the viscometer using a Hamilton syringe fitted with a glass extender. Mixing of the drug and DNA was done by bubbling with

nitrogen. Flow times were measured, using a digital stop-watch, at least thrice and were accepted if they agreed within 0.1s. Reduced specific viscosity was calculated according to Cohen and Eisenberg³¹. Plots of η/η_0 (η and η_0 are the reduced specific viscosities of DNA in the presence and absence of the drug) versus [drug]/[DNA] were constructed. Plot of η/η_0 versus [EtBr]/[DNA] was found to be similar to that reported in the literature³².

2.3e Differential pulse voltammetry: Differential pulse voltammetric experiments were performed for 0.1 mM chloride salts of the cobalt(III) complexes in the presence and absence of 0–3 mM (base-pairs) CT-DNA. The working electrode used was made up of glassy carbon and it was polished prior to each voltammetric run with 0.25 μm diamond paste on a nylon buffing pad and then subjected to ultrasonic cleaning for ≈ 5 min in 95% ethanol. Both, the peak currents and the peak potentials, were reproducible to better than 10% under our experimental conditions^{33,34}.

2.3f Gel electrophoresis: For the gel electrophoresis experiments, supercoiled pBR 322 DNA (100 μM) in *tris*-HCl buffer ($\text{pH} = 8.0$) was treated with an 10–100 μM of the metal complex and the mixture was incubated for 1 h in the dark. The samples were then analyzed by 0.8% agarose gel electrophoresis (*tris*-acetic acid-EDTA buffer, $\text{pH} = 8.0$) at 40 V for 5 h. The gel was stained with 1 $\mu\text{g}/\text{ml}$ ethidium bromide for 0.5 h after which it was analyzed using the UVP gel documentation system GDS 2000 and was also directly photographed and developed as described previously^{11,12,35,36}. Irradiation experiments were carried out by keeping the pre-incubated (dark, 1 h) samples inside the sample chamber of a JASCO Model FP-7700 spectrofluorometer. The excitation wavelength was either 313 ± 5 or 350 ± 5 nm for the Co(III) and Ni(II) salts and 450 ± 5 nm for the Ru(II) complexes.

2.3g Spin trapping studies: These experiments were carried out for irradiated (> 350 nm, 150 W Xenon arc lamp, 5–60 s irradiation time) solutions containing $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$ and PBN. ‘On-line’ ESR spectra of the samples were recorded during irradiation.

Unless otherwise specified, all the experiments were carried out at 293 ± 3 K.

3. Results and discussion

Although a few among the investigated complexes have previously been spectrally characterized to their structure^{17–26}, this study has provided an opportunity to compare the spectroscopic and other physical properties of all the complexes by using data obtained under the similar set of experimental conditions. In addition, during the course of this study it was found necessary to compile and compare the physico-chemical characteristics of each investigated complex in order to choose the appropriate technique for probing the DNA interaction. Thus, we compare the physical and spectroscopic characteristics of the investigated Co(III), Ni(II) and Ru(II) complexes before we discuss their DNA binding and photocleavage properties.

As far as the DNA interactions of these complexes are concerned, DNA binding by $[\text{M}(\text{phen})_3]^{n+}$ ($\text{M} = \text{Co(III)}$ or Ru(II) and $n = 2$ or 3) and $[\text{Ru}(\text{phen})_2(\text{dppz})]^{2+}$ have been investigated in great detail by several groups^{3,6,10,30,37–39} and that of $[\text{M}(\text{phen})_2(\text{dppz})]^{n+}$ ($\text{M} = \text{Co(III)}$ or Ni(II) and $n = 2$ or 3) by us in lesser detail¹¹. However, relatively few

studies seem to have been attempted to investigate the effect of variation of the metal ion and also the ligand on the ability to bind and photocleave DNA in mixed ligand complexes containing the phenanthroline family of ligands. In the present study, we have endeavoured to compare the DNA binding and photocleavage properties of a series of $[M(\text{phen})_2(\text{LL})]^{2+}$ type complexes under similar experimental conditions.

3.1 Spectroscopic characterization

The hexafluorophosphate salts of the complexes employed in this work have been fully characterized by UV/Visible, IR and ^1H NMR spectroscopic (for diamagnetic complexes) and magnetic susceptibility (for paramagnetic complexes) measurements. These data summarized in tables 1 and 2 agree well with the reported values, in cases where applicable^{11,17-26}.

Whereas Co(III) and Ru(II) complexes were found to be diamagnetic and hence were amenable for characterization by the ^1H NMR method, magnetic susceptibility data for nickel complexes clearly show that the metal ion is in the 2+ oxidation state in these complexes (table 2). The IR spectra of phen-dione clearly exhibits a band in the region of $1703 \pm 2 \text{ cm}^{-1}$ that is ascribable to a stretching frequency of the C=O bonds on the ligand. This band was seen to be not shifted much in the corresponding complexes (table 1), which is reasonable since the C=O moieties are far removed from the site of coordination of this ligand with the metal. The UV/Visible spectral data of all the nine complexes investigated in this study are summarized in table 1. Based on the literature data on the spectral properties of phen, phen-dione, dppz and the various other complexes containing these ligands^{15-27,40}, bands appearing in the spectra of the Co(III) and Ni(II) complexes can be assigned exclusively to the intraligand transitions. The ruthenium(II) complexes,

Table 1. UV/Visible and IR data.

Complex	UV-visible λ_{max} (nm) ($\log \epsilon$) ^a	IR $\bar{\nu}$ (cm^{-1}) ^b
$[\text{Co}(\text{phen})_3]^{3+}$	349 (4.28), 303 (4.17), 275 (4.60)	3659, 1609, 1524, 1433, 839, 557
$[\text{Co}(\text{phen})_2(\text{phen-dione})]^{3+}$	303 (4.31), 274 (4.79), 259 (4.74), 220 (5.05)	3647, 1703, 1524, 1433, 837, 557
$[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$	377 (4.23), 359 (4.29), 282 (5.11), 223 (5.10)	3653, 1608, 1524, 1433, 858, 557
$[\text{Ni}(\text{phen})_3]^{2+}$	293 (4.57), 269 (5.05), 227 (5.11)	3668, 1626, 1520, 1427, 835, 557
$[\text{Ni}(\text{phen})_2(\text{phen-dione})]^{2+}$	358 (4.13), 325 (4.14), 273 (5.03), 226 (4.92)	3652, 1701, 1520, 1428, 839, 557
$[\text{Ni}(\text{phen})_2(\text{dppz})]^{2+}$	376 (4.12), 358 (4.13), 325 (4.14), 273 (5.03), 223 (4.96)	3646, 1607, 1585, 1521, 1431, 839
$[\text{Ru}(\text{phen})_3]^{2+}$	446 (4.28), 422 (4.25), 263 (5.07), 223 (4.93)	3654, 1622, 1496, 1428, 840, 557
$[\text{Ru}(\text{phen})_2(\text{phen-dione})]^{2+}$	441 (4.29), 329 (4.51), 261 (5.11), 225 (4.92)	3659, 1701, 1498, 1429, 838, 558
$[\text{Ru}(\text{phen})_2(\text{dppz})]^{2+}$	443 (4.33), 369 (4.39), 360 (4.33), 277 (5.12), 225 (5.02)	3651, 1628, 1491, 1427, 837, 557

^aSpectra were measured in CH_3CN . Error limits: λ_{max} , $\pm 2 \text{ nm}$; $\log \epsilon$, $\pm 10\%$

^bSpectra were measured as KBr pellets

Table 2. ¹H NMR and magnetic susceptibility data ^a.

Complex	δ ppm ^a	μ_{eff} (BM) ^b
[Co(phen) ₃] ³⁺	9.19 (<i>d</i> , 6H), 8.58 (<i>s</i> , 6H), 7.98 (<i>dd</i> , 6H), 7.68 (<i>d</i> , 6H)	
[Co(phen) ₂ (phen-dione)] ³⁺	9.17 (<i>d</i> , 6H), 8.56 (<i>s</i> , <i>br</i> , 4H), 7.96 (<i>m</i> , 6H), 7.66 (<i>d</i> , 6H)	
[Co(phen) ₂ (dppz)] ³⁺	9.95 (<i>dd</i> , 2H), 9.21 (<i>d</i> , 2H), 8.18 (<i>d</i> , 2H), 8.59 (<i>m</i> , 8H), 8.32 (<i>d</i> , 4H), 8.00 (<i>m</i> , 4H), 7.68 (<i>d</i> , 4H)	
[Ru(phen) ₃] ²⁺	8.80 (<i>dd</i> , 6H), 8.39 (<i>s</i> , 6H), 8.09 (<i>dd</i> , 6H), 7.76 (<i>d</i> , 6H)	
[Ru(phen) ₂ (phen-dione)] ²⁺	9.26 (<i>d</i> , 6H), 8.71 (<i>s</i> , <i>br</i> , 4H), 8.12 (<i>m</i> , 6H), 7.82 (<i>d</i> , 6H)	
[Ru(phen) ₂ (dppz)] ²⁺	9.73 (<i>dd</i> , 2H), 8.82 (<i>dd</i> , 2H), 8.79 (<i>dd</i> , 2H), 8.60 (<i>dd</i> , 2H), 8.50 (<i>dd</i> , 2H), 8.48 (<i>m</i> , 2H), 8.42 (<i>s</i> , 4H), 8.39 (<i>dd</i> , 2H), 8.18 (<i>m</i> , 2H) 7.95 (<i>dd</i> , 2H), 7.83 (<i>dd</i> , 2H), 7.81 (<i>dd</i> , 2H)	
[Ni(phen) ₃] ²⁺		3.00
[Ni(phen) ₂ (phen-dione)] ²⁺		3.11
[Ni(phen) ₂ (dppz)] ²⁺		3.16

^aSpectra were measured in DMSO-*d*₆/CDCl₃ using TMS as an internal standard. Error limit: ± 0.1 ppm

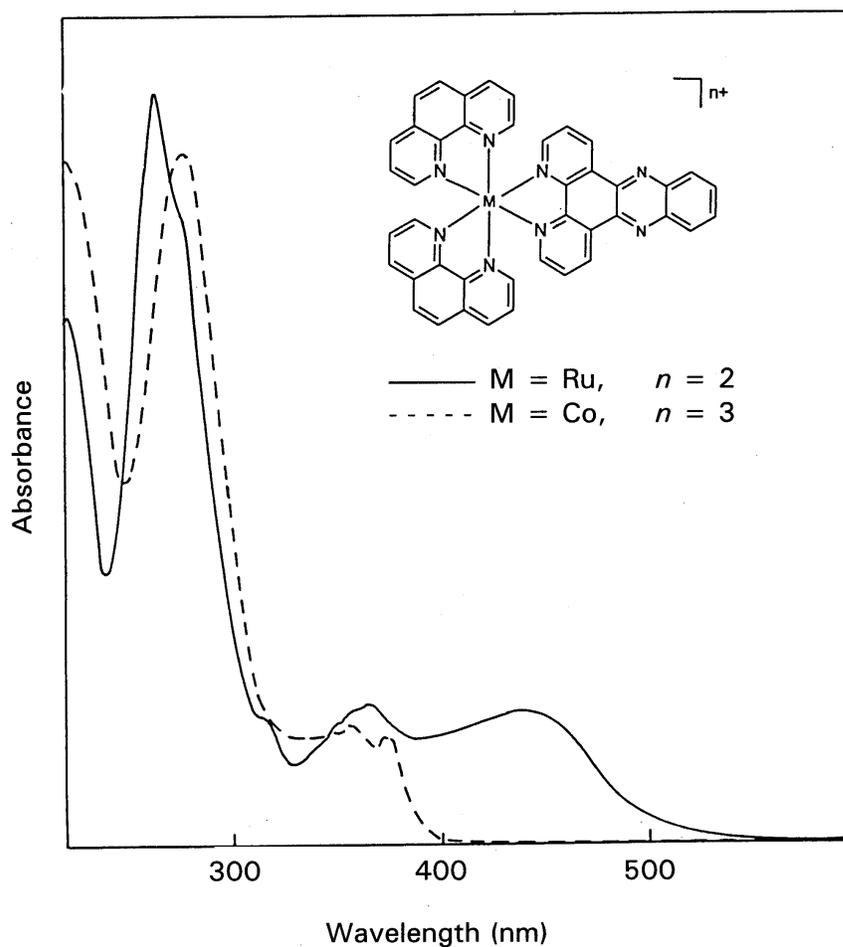
^bmeasured at 293 \pm 3 K. Error limit: $\pm 10\%$

on the other hand, showed additional MLCT charge transfer bands between 400–500 nm. This is illustrated in figure 2 which compares the UV/Visible spectra of [Ru(phen)₂(dppz)]²⁺ and [Co(phen)₂(dppz)]³⁺. Interestingly, each mixed ligand complex containing dppz absorbs at red-shifted wavelengths compared to the complexes containing phen-dione. While none of the Co(III) and Ni(II) complexes studied here were found to be fluorescent, each Ru(II) complex showed intense emission when excited into the MLCT band in CH₃CN. The emission quantum yield values (ϕ) of these complexes vary as [Ru(phen)₃]²⁺ (0.028) > [Ru(phen)₂(dppz)]²⁺ (0.008) > [Ru(phen)₂(phen-dione)]²⁺ (0.002).

Redox potentials of the ligands and the complexes, as obtained by the cyclic voltammetric method are summarized in table 3. Wave analysis suggested that most of the voltammetric peaks represent diffusion controlled (ip vs $v^{1/2}$ = constant where ip is the peak current and v is the scan rate), reversible (ip_a/ip_c = 0.9–1.0 where ip_a and ip_c refer to anodic and cathodic peak potentials, respectively), one-electron transfer (ΔE_p = 60–70 mV where E_p is the peak potential) reactions, while the others are either quasi-reversible (ip_a/ip_c = 0.7–0.9 and ΔE_p = 80–200 mV) or totally irreversible, as indicated in table 3. The peak assignments to the metal or the ligand based redox reactions are based on the reported electrochemical data of [M(phen)₃]²⁺ and [M(phen)₂(LL)]ⁿ⁺ systems ^{11,16,27,33}. A comparison of the reduction potential data of free and metal-bound dppz and phen-dione in the mixed-ligand complexes containing these π -acceptor ligands suggests that the potential displacement between the free and the coordinated species varies between 0.03 and 0.35 V. It is possible that the electronic coupling between these ligands and the metal ions is weak in their complexes. On the other hand, the potential displacement between the free and coordinated phen is > 1 V for the series of complexes investigated in this study. Finally, the redox wave corresponding to the Co^{III/II} couple for each Co(III) complex investigated here was found to be reversible and appear at potentials as low as 0.37–0.40 V in CH₃CN, 0.1 M TBAP.

Table 3. Redox potential data in CH₃CN, 0.1M TBAPF₆^a.

Complex	Metal	Ligand
Phen		-2.14*
Phen-dione		-0.45, -1.08
dppz		-1.22, -1.74*
[Co(phen) ₃] ³⁺	0.38	-0.98, -1.67*
[Co(phen) ₂ (phen-dione)] ³⁺	0.37	-0.42*, -0.90*
[Co(phen) ₂ (dppz)] ³⁺	0.40	-0.95, -1.17, -1.83*
[Ni(phen) ₃] ²⁺		-1.32, -2.02*
[Ni(phen) ₂ (phen-dione)] ²⁺		-0.17, -0.89
[Ni(phen) ₂ (dppz)] ²⁺		-0.99, -1.32*
[Ru(phen) ₃] ²⁺	1.26	-1.28, -1.44, -1.71*
[Ru(phen) ₂ (dppz)] ²⁺	1.33	-0.87, -1.29, -1.53

^aError limit: ± 0.03 V*Quasi-reversible/irreversible (electrochemical behaviour of [Ru(phen)₂(phen-dione)]²⁺ was found to be ill-defined)**Figure 2.** UV/Visible spectra of [Co(phen)₂(dppz)]³⁺ and [Ru(phen)₂(dppz)]²⁺ in CH₃CN.

Thus, the UV/Visible data suggest that DNA interaction by all the dppz-based complexes, having absorption peaks at wavelengths > 300 nm (up to which DNA itself absorbs), can be monitored by the absorption titration method. In addition, while the interaction of each Ru(II) complex with DNA can be specifically probed by emission titration, that by the Co(III) complexes can be probed by electrochemical methods.

3.2 DNA binding studies

Addition of increasing amounts of CT-DNA resulted in a decrease of absorbance for each investigated dppz complex and also that of $[\text{Ru}(\text{phen})_2(\text{phen-dione})]^{2+}$ and $[\text{Ru}(\text{phen})_3]^{2+}$. Representative spectra illustrating this hypochromicity and the presence of isobestic points observed for the interaction of $[\text{Ru}(\text{phen})_2(\text{phen-dione})]^{2+}$ with CT-DNA are given in figure 3. Change in absorbance at the peak maximum of the most red-shifted band of each complex with increasing concentration of DNA has been monitored for an evaluation of the intrinsic binding constant using (1) (see figure 3 inset, for the plot using (1)); the binding constants thus obtained are given in table 4. As reported earlier, dppz complexes of Co(III), Ni(II) and Ru(II) are found to be avid binders of CT DNA with the

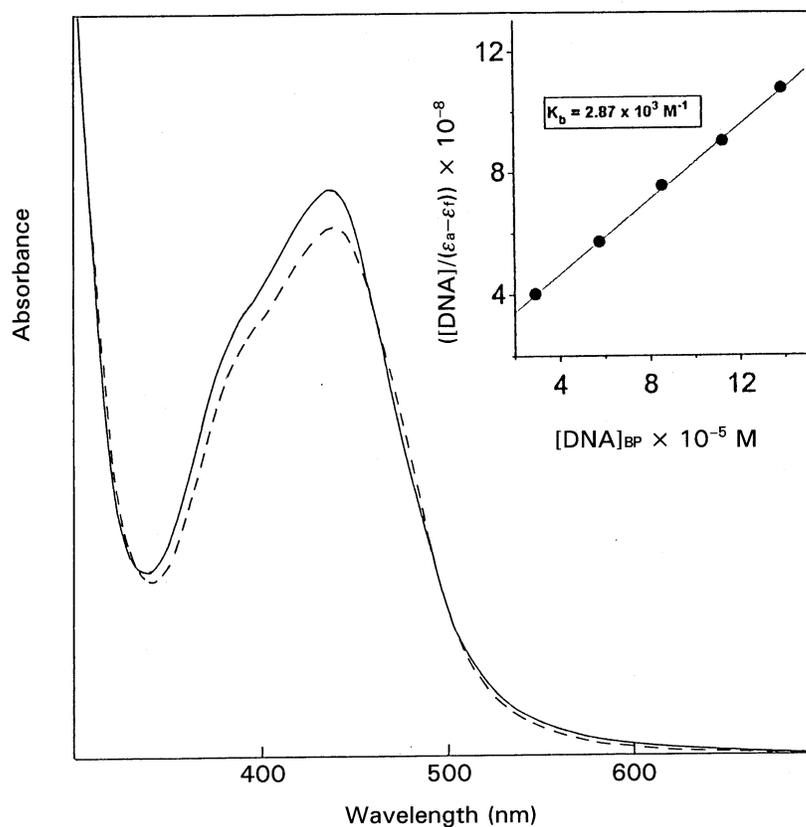


Figure 3. UV/Visible spectra of $[\text{Ru}(\text{phen})_2(\text{phen-dione})]^{2+}$ ($10 \mu\text{M}$) with (---) and without (—) CT-DNA ($100 \mu\text{M}$ in base-pairs) in buffer A. Inset: Plot of $[\text{DNA}]/(\epsilon_a - \epsilon_f)$ vs $[\text{DNA}]$ for this interaction.

Table 4. Results of absorption titration (K_b) and thermal denaturation (T_m) studies.

Complex	K_b , M ⁻¹ ^a	T_m °C ^b	σ_T ° ^c
[Co(phen) ₃] ³⁺	–	66	26
[Co(phen) ₂ (phen-dione)] ³⁺	–	65	24
[Co(phen) ₂ (dppz)] ³⁺	5.05×10^5	68	25
[Ni(phen) ₃] ²⁺	–	64	26
[Ni(phen) ₂ (phen-dione)] ²⁺	–	64	24
[Ni(phen) ₂ (dppz)]	1.51×10^5	66	27
[Ru(phen) ₃] ²⁺	7.88×10^3	66	27
[Ru(phen) ₂ (phen-dione)] ²⁺	2.87×10^3	66	25
[Ru(phen) ₂ (dppz)] ²⁺	$>10^6$	67	26

^aError limit: $\pm 10\%$ ^b[DNA nucleotide phosphate]/[drug] = 25; error limit $\pm 1^\circ\text{C}$ ^cError limit: $\pm 2^\circ$

binding constants being in the range of 1.5×10^4 – $>10^6$ M⁻¹ ^{11,37–39}. On the other hand, both [Ru(phen)₃]²⁺ and [Ru(phen)₂(phen-dione)]²⁺ bind less efficiently; the derived K_b values in these cases are \approx more than two orders of magnitude less than that for the corresponding dppz complex. In fact, K_b values as derived from the absorption titration method for these Ru(II) complexes vary as [Ru(phen)₂(dppz)]²⁺ $>$ [Ru(phen)₃]²⁺ $>$ [Ru(phen)₂(phen-dione)]²⁺.

A similar effect was noticed during the fluorescence titration experiments using these emissive Ru(II) complexes in the presence of DNA. Luminescence due to each Ru(II) complex was seen to increase steadily with increasing addition of CT-DNA, figure 4. As seen, the maximum enhancement factors (at saturation) noticed for [Ru(phen)₂(dppz)]²⁺, [Ru(phen)₃]²⁺ and [Ru(phen)₂(phen-dione)]²⁺ are $>10^4$, 2 and 1.8 respectively. It is pertinent to mention here that the 10^4 -fold emission enhancement observed for [Ru(phen)₂(dppz)]²⁺ has been ascribed earlier to protection of the imine nitrogens of the fused phenazine subunit on dppz from attack by water and consequent decrease in the non-radiative processes upon intercalation with DNA ^{37,38,41–44}. Such a process is obviously lacking in [Ru(phen)₃]²⁺ and [Ru(phen)₂(phen-dione)]²⁺, both of which do not possess the fused phenazine subunit in their architecture.

Owing to the lack of an absorption band above ≈ 300 nm, a wavelength up to which DNA itself absorbs, the binding affinities of the phen and phen-dione complexes of Co(III) and Ni(II) could not be investigated by the absorption titration method. Nor was it possible to monitor the binding of these complexes by the fluorescence titration method, as they were all found to be essentially non-emissive both in the absence and presence of DNA. However, application of the differential-pulse voltammetric method permitted an estimate of DNA binding affinities of the Co(III) complexes as described below.

Differential-pulse voltammetric experiments carried out in buffer B for [Co(phen)₃]³⁺, [Co(phen)₂(phen-dione)]³⁺ and [Co(phen)₂(dppz)]³⁺ both in the presence and absence of CT DNA have revealed that there is a decrease in the peak-current due to Co^{III}/Co^{II} redox couple in the presence of DNA. The decrease in the current value is more pronounced for the dppz containing complex compared to that for the phen and phen-dione complexes. The current values for [Co(phen)₂(dppz)]³⁺, [Co(phen)₃]³⁺ and [Co(phen)₂(phen-dione)]³⁺

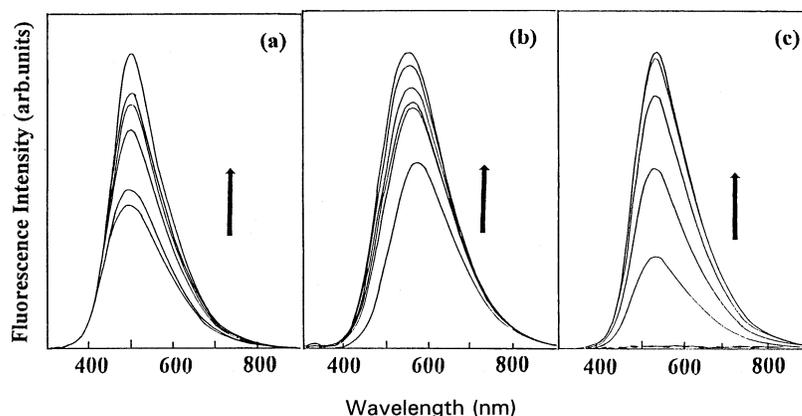


Figure 4. Emission enhancement observed for the three Ru(II) complexes ($10 \mu\text{M}$, buffer A) with increasing concentration of CT-DNA. (a) $[\text{Ru}(\text{phen})_3]^{2+}$, (b) $[\text{Ru}(\text{phen})_2(\text{phen-dione})]^{2+}$ and (c) $[\text{Ru}(\text{phen})_2(\text{dppz})]^{2+}$. Maximum DNA concentrations added (nucleotide phosphates) are 800 , 800 and $100 \mu\text{M}$ for (a), (b) and (c), respectively.

at a $[\text{DNA}]/[\text{Co}] = 40$ were respectively 30 , 50 and 80% of the corresponding values in the absence of DNA. This trend is consistent with that noticed above for the ruthenium(II) complexes during the absorption and fluorescence titration experiments.

A comparison of the available intrinsic binding characteristics by these complexes thus appears to suggest that, in general, DNA binding by the dppz containing complexes is far too strong than that by the complexes containing phen or phen-dione ligands, with the latter species being the weakest binding agents among all the systems investigated in this study. This supposition is further substantiated by the results of thermal denaturation, viscometric titration and agarose gel electrophoresis experiments, the results of which are described below.

Thermal denaturation curves for DNA in the presence and absence of a representative complex are given in figure 5 and the relevant data for all the complexes investigated in this study are summarized in table 4. It is clear from this figure and the data given in table 4 that while the dppz containing complexes shift the T_m values by up to 8° , the T_m values for DNA samples containing the *tris*-phen and *bis*(phen)(phen-dione) complexes, in general, are not so high compared to that of the pure DNA sample ($60 \pm 1^\circ\text{C}$)³⁰.

Intercalation of a ligand to DNA is known to cause a significant increase in the viscosity of a DNA solution due to an increase in the separation of the base pairs at the intercalation site and, hence, an increase in the overall DNA molecular length. In contrast, a ligand that binds in the DNA grooves causes either a less pronounced change (positive or negative) or no change in the viscosity of a DNA solution^{45,46}. The effect of each investigated complex on the viscosity of CT-DNA solution was studied in order to assess the binding mode and strength of these complexes with DNA. Representative plots of η/η_0 vs $[\text{drug}]/[\text{DNA}]$ are shown in figure 6 for the cobalt(III) complexes. As seen in this figure, while $[\text{Co}(\text{phen})_3]^{3+}$ does not affect the DNA viscosity as previously reported for this *tris*-phen complex³², positive and negative changes of viscosity with increasing addition of the complex are seen for $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$ and $[\text{Co}(\text{phen})_2(\text{phen-dione})]^{3+}$

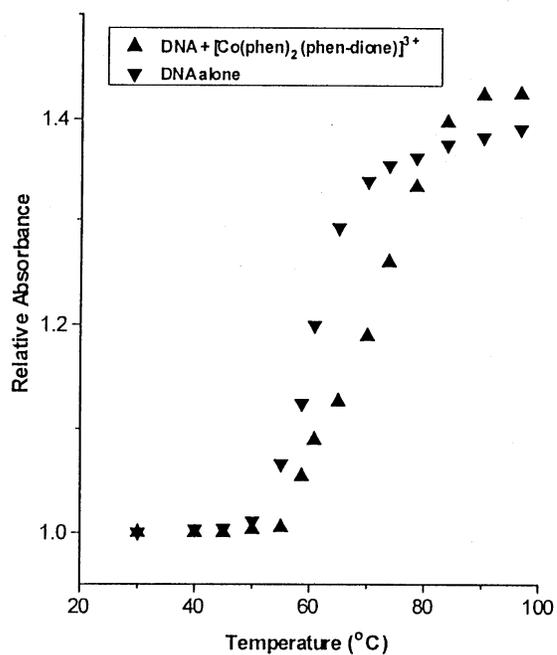


Figure 5. Melting curves (buffer B) for DNA alone (\blacktriangledown) and DNA + $[\text{Co}(\text{phen})_2(\text{phen-dione})]^{3+}$ (\blacktriangle). $[\text{DNA}] = 160 \mu\text{M}$. $[\text{Drug}] = 8 \mu\text{M}$.

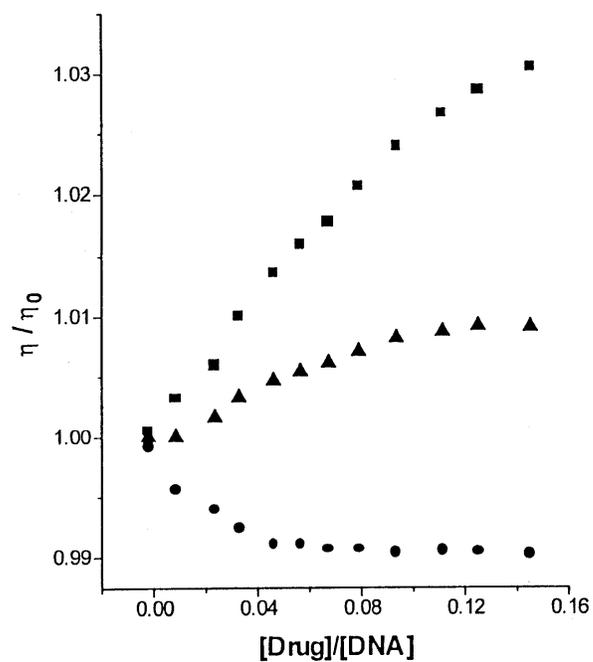


Figure 6. Plots of η/η_0 vs $[\text{drug}]/[\text{DNA}]$ (buffer B) for $[\text{Co}(\text{phen})_2(\text{phen-dione})]^{3+}$ (\blacksquare), $[\text{Co}(\text{phen})_3]^{2+}$ (\bullet) and $[\text{Co}(\text{phen})_2(\text{dppz})]^{2+}$ (\blacktriangle).

suggesting an intercalative mode of binding by the dppz-complex^{31,32}. A similar trend was noticed for the Ni(II) and Ru(II) analogues.

It has been suggested in the case of *tris*-phen complexes of various metal ions, that only one among the three phen ligands is involved in the intercalative mode of binding with DNA⁴⁷. On the other hand, results of various spectroscopic and biochemical studies have shown that the mixed-ligand complexes $[\text{Ru}(\text{bpy})_2(\text{dppz})]^{2+}$, $[\text{Ru}(\text{phen})_2(\text{dppz})]^{2+}$, $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$, $[\text{Ni}(\text{phen})_2(\text{dppz})]^{2+}$ and $[\text{Ru}(\text{O})(\text{dppz})(\text{tpy})]^{2+}$ (tpy = terpyridine) strongly bind to DNA and that it is the dppz ligand on them that binds with DNA^{11,37,38,41-44,48}. In particular, bathochromic shifts and hypsochromism in their UV/Visible spectrum, increased values of both the DNA melting temperature and the curve width in the thermal denaturation experiments and enhanced viscosity changes in the presence of DNA have all been argued in favour of an intercalative mode of binding by the coordinated dppz. Several factors which include the shape and hydrophobicity of the complex as well as extension of planarity and the presence of additional donor functionalities on the dppz ligand have been cited to be of importance in rendering these complexes to be such strong intercalative, major groove binding agents. It should be noted here that each dppz-based complex investigated in the present study also exhibited spectral, electrochemical and viscosity changes in the presence of DNA that are analogous to those exhibited by the complexes mentioned above. On the other hand, less pronounced spectral and viscosity changes have been observed for the phen and phen-dione complexes in the presence of DNA. Clearly, the intercalative ability of dppz is far too superior to that of phen and phen-dione.

Taking the above facts into consideration, we propose that only one among the three ligands on each complex is involved with the intercalative binding with the CT DNA and that the intercalative ability of the ligands varies as $\text{dppz} > \text{phen} > \text{phen-dione}$ in this series of complexes. The fact that coordinated dppz and phen bind to DNA is well-known. However, to our knowledge, DNA binding abilities of the phen-dione complexes have never been tested. The results obtained in this study indicate that coordinated phen-dione is a poor intercalator unlike dppz and phen. Indeed, the viscosity and thermal denaturation parameters of DNA have been noticed to be relatively insensitive to the presence of *tris*(phen-dione) complexes $[\text{M}(\text{phen-dione})_3]^{n+}$ (M = Co(III) or Ni(II) and $n = 2$ or 3) (data not shown). Notwithstanding this latter observation, it is unclear at present whether it is phen-dione or phen that is involved in the binding of $[\text{M}(\text{phen})_2(\text{phen-dione})]^{n+}$ systems investigated in this study. Nonetheless, it is interesting to note that extending the π -conjugation in the phenanthroline family of ligands, as is done in the case of dppz, results in strong intercalative agents. On the other hand, substitution of the hydrogen-bonding acceptors onto the phenanthroline ligands, such as for example the carbonyl groups of phen-dione, does not seem to provide any additional stabilization for binding by the ensuing complexes with DNA. Indeed, whereas complexes of modified phen ligands with fused aromatic groups are known to strongly bind to DNA^{49,50} $[\text{Ru}(\text{phen})_2(\text{flone})]^{2+}$, where flone is 4,5-diazafluorene-9-one – a potentially hydrogen-bonding ligand – shows poor affinity towards DNA as is the case with the phen-dione complexes investigated in this study⁵⁰.

3.3 Photocleavage of DNA

DNA photocleavage by the cobalt(III) complexes has been investigated in detail in this study. Control experiments have suggested that untreated DNA does not show any

cleavage in the dark and even upon irradiation by light. Control experiments have also suggested that phen, phen-dione and dppz (free-ligands, dissolved in 10% DMF) are not detectably active under the dark and light irradiated conditions. DNA nicking was not observed for pBR 322 treated with any of the complexes investigated in this study in the dark experiments. On the other hand, irradiation of DNA in the presence of the three Co(III) complexes caused the generation of relaxed circular DNA ($\lambda_{\text{exc}} = 313 \pm 5$ nm) with the nicking efficiency roughly following the order $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$ (1.0) > $[\text{Co}(\text{phen})_3]^{3+}$ (0.8) > $[\text{Co}(\text{phen})_2(\text{phen-dione})]^{3+}$ (0.7). It should be noted that, at this wavelength these complexes show more equal absorbance ($\log \epsilon$ values at 313 nm are 4.23, 4.16 and 4.10 for the three complexes in that order) than at 350 nm where only $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$ is capable of absorbing light. Accordingly, when the λ_{exc} was changed to 350 ± 5 nm from 313 ± 5 nm, only $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$ was found to effect the DNA cleavage. These observations highlight the importance of dppz in this class of cobalt complexes and suggest that it is essential to further probe the interaction of $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$ with pBR 322 DNA in order to gain insight into the mechanism of the photocleavage reaction.

Irradiation of pBR 322 samples containing $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$ was carried out in the presence of various 'inhibitors' (figure 7). Neither N_2 (used to purge O_2 from the sample, lane 3) nor DABCO (lane 5) – a $^1\text{O}_2$ 'quencher' and SOD – a O_2^- 'scavenger' (lane 6) inhibit the photocleavage by the complex. On the other hand, DMSO (and also mannitol or ethanol, data not shown) which scavenges OH^\bullet radical seems to inhibit the photocleavage (lane 4). Further support for the generation of OH^\bullet upon photolysis of the complex comes from the spin trapping experiments. In the presence of PBN as the spin trap, irradiated solutions of $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$ in deaerated aqueous buffer (phosphate buffer $\text{pH} = 7.0$) indeed exhibited an ESR spectrum consisting of three doublets ($g = 2.006$, $a^{\text{N}} = 15.2$ G and $a^{\text{H}} = 2.8$ G) typical of the OH^\bullet spin adduct of PBN⁵¹. In addition, a complex ESR spectrum was obtained when irradiation of the complex was carried out in deaerated acetonitrile solution containing $\approx 1\%$ water, figure 8. As seen in this figure, apart from a major three-line pattern due to PBN- OH^\bullet spin adduct, the

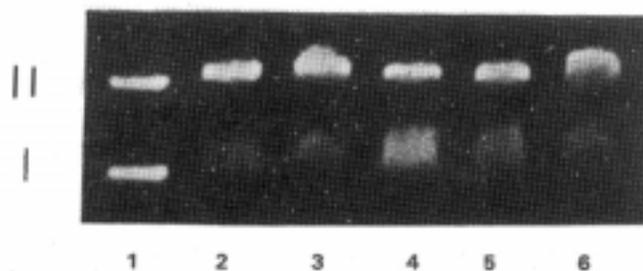


Figure 7. Photograph showing effects of 'inhibitors' on the light-induced nuclease activity of $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$: Lane 1: Untreated pBR 322 (100 μM nucleotide phosphate)⁴¹, Lane 2: pBR 322 + $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$ (100 μM)⁶⁵, Lanes 3–6: pBR 322 + $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$ in the presence of N_2 ⁷⁰, DMSO (0.2 M)⁴⁵, DABCO (10 mM)⁶² and SOD (20 $\mu\text{g}/\text{ml}$)⁷², respectively. Irradiation time = 45 min in each case and $\lambda_{\text{exc}} = 350 \pm 5$ nm. Numbers given in square-parentheses refer to percentages of form II DNA.

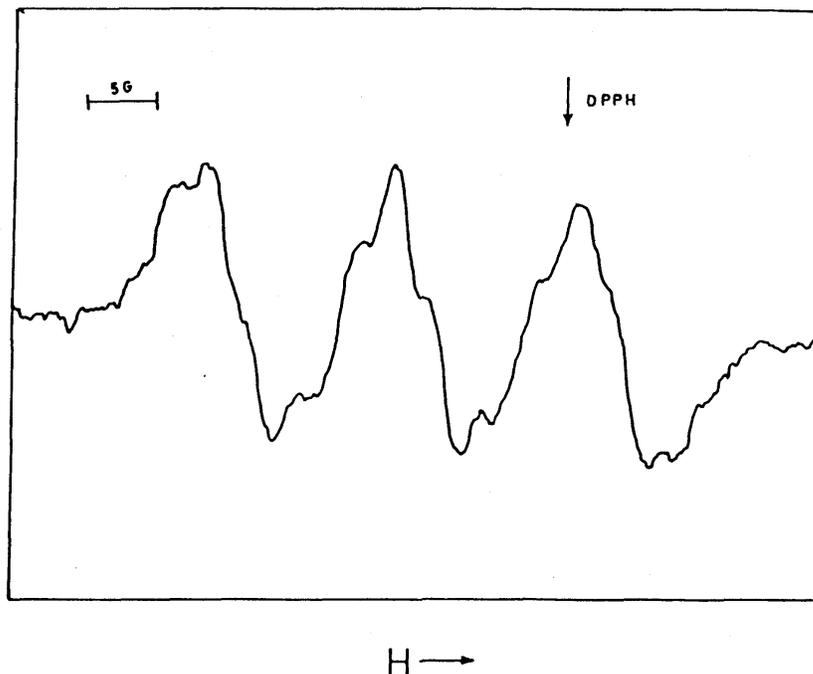


Figure 8. ESR spectrum obtained when 1 mM of $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$ was irradiated ($\lambda_{\text{exc}} = 350 \pm 5 \text{ nm}$) in acetonitrile solution containing 1% water.

spectrum also shows several splittings which we ascribe to the resonances of a ligand C/N-based radical⁵². Thus, it is possible that irradiation facilitates a process of ligand reduction in $[\text{Co}(\text{phen})_2(\text{dppz})]^{3+}$, a process that leads to the production of DNA-cleaving OH \cdot radical via the reaction of a C/N-based radical intermediate with the buffered water. Notwithstanding these evidences in favour of the presence of hydroxyl radicals, we believe that the participation of other reactive species cannot be altogether ruled out in the observed photocleavage reaction.

The three ruthenium(II) complexes also exhibited light induced nuclease activity when irradiated into their respective MLCT bands ($\lambda_{\text{exc}} = 450 \pm 5 \text{ nm}$). Detailed studies on the DNA photocleavage by $[\text{Ru}(\text{phen})_3]^{2+}$ have been carried out previously and it has been reported that $\text{O}_2^{\cdot-}$ and $^1\text{O}_2$ are the two important species responsible for the DNA damage^{53,30}. We expect that photocleavage reactions of $[\text{Ru}(\text{phen})_2(\text{dppz})]^{2+}$ and $[\text{Ru}(\text{phen})_2(\text{phen-dione})]^{2+}$ also involve these cytotoxic species. In addition, $[\text{Ru}(\text{phen})_2(\text{dppz})]^{2+}$ has recently been reported to induce oxidative damage to DNA by rapidly oxidizing the guanine moieties upon irradiation⁵⁴. Finally, none of the d^8 nickel(II) systems showed light-induced nuclease activity under our experimental conditions, probably because of the paramagnetic nature of these complexes that, in principle, would render the excited states of these molecules ineffective.

In summary, the results described in this study demonstrate that substitution by different ligands or metal ions in metallo-intercalators of the type $[\text{M}(\text{phen})_2\text{LL}]^{n+}$ can bring about subtle modulation in the properties of this class of mixed-ligand complexes and, consequently, in their interactions with DNA.

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