Synthesis, Characterization, DNA Binding and Cytotoxic Studies of Platinum(II) and Palladium(II) Complexes of the 2,2'-Bipyridine and an Anion of 1,1-Cyclobutanedicarboxylic Acid

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Two neutral complexes of formula [M(bpy)(cbdca)] [where M is palladium(II) (Pd(II)) or platinum(II) (Pt(II)), bpy is 2,2'-bipyridine and cbdc is anion of 1,1-cyclobutanedicarboxylic acid] have been synthesized. These water soluble complexes have been characterized by chemical analysis and conductivity measurements as well as 'H-NMR, ultraviolet-visible and infrared spectroscopy. In these complexes the ligand cbdc coordinates to Pt(II) or Pd(II) as bidentate with two oxygen atoms. They are nonelectrolyte in conductivity water. These complexes inhibit the growth of P388 lymphocytic leukemia cells and their targets are DNA. They invariably show ID50 values less than cisplatin. [Pt(bpy)(cbdca)] and [Pd(bpy)(cbdca)] have been interacted with calf thymus DNA and bind to DNA through coordinate covalent bond. In addition, the influence of binding of these complexes on the intensity of EtBr–DNA have been studied. They bind to DNA via a nonintercalating mode.

Key words platinum complex; palladium complex; DNA-binding study; 1,1-cyclobutanedicarboxylic acid

Since the discovery of the antineoplastic agent cisplatin (c-DDP) by Rosenberg et al., the drug has been in wide-spread use to date. However, because of its toxic side-effects, particularly renal damage, the search for safer platinum complexes and administration methods is still continuing. Several platinum and palladium containing compounds have subsequently been synthesized and tested. One of these, carboplatin, was approved for treatment of ovarian cancer. This has no kidney toxicity, there is less nausea, neurotoxicity, otoplotoxicity than cisplatin and the drug could be given without hydration treatment. Recent work on the reaction mode of carboplatin focused on the interaction with DNA building blocks. It was concluded that the reaction with carboplatin proceeds via a direct substitution mechanism, since the reaction with chloride was too slow to account for an aqua or chloro complex as the reactive species.

Tobe and Kokhar reported that change in the amine ligands have primary effect on antitumor properties of platinum(II)/palladium(II) [Pt(II)/Pd(II)] complexes. Recent kinetic work by van Eldik and his co-workers. on [Pd(bpy)(cbdca)] [where bpy is 2,2'-bipyridine and cbdc is anion of 1,1-cyclobutanedicarboxylic acid] indicates that a decrease in reactivity was induced by increasing the steric hindrance on the amine ligands, and attributed to the axial hindrance becoming blocked for the incoming ligands. On substitution the two monodentate amine groups of carboplatin with bpy, the resulting complexes have a UV-absorption band in a region where DNA does not show absorption. This enable us to carry out DNA-binding studies using A.M.Q. King method. Thus, in this paper we report the synthesis, characterization DNA-binding and cytotoxic studies of bpy Pt(II) and Pd(II) complexes containing cbdc which are analogous to carboplatin.

Experimental
Commercially pure chemicals: K2PtCl4, PdCl2, bpy, cbdc, highly polymerized calf thymus DNA sodium salt and Tris–HCl buffer were purchased from Aldrich U.S.A. and used as such. Solvents used were of laboratory reagent grade and they were purified before use by the standard methods.

Electronic absorption spectra of the titled metal complexes were measured on a Shimadzu UV-265 recording spectrophotometer. Infrared spectra of the metals complexes were recorded on a Nicolet 5-DSB FT-IR spectrophotometer in the range of 4000—400 cm-1 in KBr pellets. Conductivity measurements were carried out on a Systronics conductivity bridge, model 305, with a cell (cell constant 0.59) using conductivity water as a solvent. Micro chemical analysis of carbon, hydrogen and nitrogen for the complexes were carried out on an elemental analysis of CHNO-Rapid Heraeus. 'H-NMR spectra were recorded on a Brucker DRX-500 Avance spectrometer at 500 MHz in D2O using sodium 3-trimethylpropionate as internal reference. Fluorescence measurements were carried out on a SPEX-Flurolog spectrophotometer equipped with 1681, 0.22 mm double spectrometer and DMIB spectroscopy laboratory coordination using a 1 cm² spectrophotometer quartz cuvette.

Synthesis of Pd(II) and Pt(II) Compounds [Pt(bpy)Cl2] and [Pd(bpy)Cl2] were prepared by the procedures followed for making analogous bpy complexes. [Pt(bpy)(cbdca)] [Pt(bpy)(cbdca)] (420 mg, 1 mmol) was suspended in 100 ml double distilled water and 19.7 ml of 0.10 M AgNO3 (1.97 mmol) solution was added to it with constant stirring. This mixture was heated at 60°C with stirring for 12 h at 50°C and then filtered in hot. The clear solution was concentrated on water bath at 45°C to 5 ml. The yellow precipitate was filtered, washed with small amount of chilled double distilled water and finally recrystallized from double distilled water and dried in vacuum desiccator over anhydrous calcium chloride. The yield was 70%. Anal. Calcd for [Pt(bpy)(cbdca)]: C, 38.95; H, 2.84; N, 5.68; Pt, 39.57% . Found: C, 39.04; H, 2.93; N, 5.61; Pt, 39.49%. Absorption spectrum (H2O): λmax (εM), 318.6 (7630), 263.2 (7780), 232.5 (10820) nm.

[Pd(bpy)(cbdca)]: This complex was synthesized and recrystallized by following the preparative method of [Pt(bpy)(cbdca)] except [Pd(bpy)Cl2] was used in place of [Pt(bpy)Cl2]. The yield was 65%. Anal. Calcd for [Pd(bpy)(cbdca)]: C, 47.52; H, 3.47; N, 6.93; Pd, 26.33. Found: C, 47.46; H, 3.50; N, 6.93; Pd, 26.27%. Absorption spectrum (H2O): λmax (εM), 309.3 (11750), 245.7 (14880), 220.6 (31630).

Cytotoxic Studies The procedure followed was similar to that reported earlier except that 2×104 P388 lymphocytic leukemia cells per ml were used in place of 1×106 P388 cells per ml in Tris–HCl buffer solution.

Binding Studies The difference UV absorption and fluorescence methods were used to determine the binding of metal complexes to DNA as described earlier except that the metal complexes–DNA systems were incubated at 37°C for 10 h before spectral measurements.

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Results and Discussion

The mixed-ligand complexes of formula [M(bpy)(cbdca)] (where M is Pd(II) or Pt(II)), have been prepared by interaction of α,α-diiminedinitro Pt(II)/Pd(II) complexes with cbdca. These complexes (1 × 10⁻³ M) show molar conductance values in the range 5—6 cm⁻¹·mol⁻¹·mol⁻¹ and these values are unchanged after 2 d. Thus, they can be considered as neutral complexes.20)

Spectral Studies The selective infrared spectral data of the Pt(II) and Pd(II) complexes in the range 4000 to 400 cm⁻¹ are discussed. The infrared spectra of cbdca in KBr pellets showed a strong band at 1700 cm⁻¹ which is assigned to the COOH stretching band of this ligand.21) In the Pt(II) and Pd(II) complexes the COO stretching band is observed at 1655 and 1642 cm⁻¹, respectively. This suggest that the carboxyl groups of the above dicarboxylic acid is coordinated as bidentate through two oxygen atoms as shown in Fig. 1.

The electronic absorption maxima of the Pt(II) and Pd(II) complexes in distilled water with their extinction coefficients are given in the experimental section. The band at 318 nm for complexes in distilled water with their extinction coefficients.

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The fluorescence of ethidium bromide (EtBr) is greatly enhanced upon interaction of EtBr to DNA. The influence of binding of some \( \alpha \)-diimine Pt(II) and Pd(II) complexes of cbdca on the fluorescence intensity of EtBr–DNA complex has been studied. The number of EtBr molecules intercalated to DNA in the presence of [Pt(bpy)(cbdca)] and [Pd(bpy)(cbdca)] were calculated using Scatchard analysis. In these experiments the wavelength of excitation was set at 546 nm and the wavelength of emission at 605 nm. The saturation curves of fluorescence intensity for [Pt(bpy)(cbdca)]–DNA system at different \( r_f \) values (0.0—0.314) in the presence of increasing concentration of EtBr (2.4—24 \( \mu \)M) in 10 mM Tris–HCl buffer of pH 7.4 in the presence of 0.01 M sodium chloride were obtained. The fluorescence Scatchard plots were obtained for binding of EtBr to DNA in the absence (\( \circ \)) and in the presence (\( \bullet \)) of various concentrations of [Pt(bpy)(cbdca)] and are shown in Fig. 4. This Figure shows that the above complexes noncompetitively inhibit the EtBr binding to DNA (type-D behavior), in which the slope that is \( K \) (association constant) remain constant while the intercept on the abscissa that is \( n \) (the number of binding sites per nucleotide) decrease with increasing concentrations of metal complex. This suggests that the [Pt(bpy)(cbdca)] binds to DNA \textit{via} nonintercalating mode. The values of \( K \) and \( n \) are listed in Table 1. Similar Scatchard plots have been observed for [Pd(bpy)(cbdca)].

Other studies have also been carried out to determine the mode of the above interaction as followd. Each of the above metal complexes (48 mM) was incubated with calf thymus DNA (120 \( \mu \)M) for 10 h at 37°C in Tris–HCl buffer pH 7.4. Each DNA–metal complex was passed through a Sephadex G.25 column equilibrated with same buffer. The elution of the column fraction of 2.5 ml was monitored at 319 and 259 nm for DNA–Pt(II) complex systems and 314 and 259 nm for DNA–Pd(II) complex systems. These results are given in Figs. 5a and b, respectively. These plots show that the peak obtained for the two wavelengths were not clearly resolved. This indicates that the DNA is not separated from the metal complexes and their binding to DNA is covalent and irreversible. The irreversibility of metal complex to DNA is strictly true for kinetically inert Pd(II)–DNA complex and there is expected partial reversibility for kinetic labile Pd(II)–DNA complex as observed.
above.

The precipitation of DNA from DNA–metal complex with absolute alcohol was also attempted. The presence of DNA–metal complex in the precipitate suggests the coordinate covalent bond which is responsible for stabilizing the above DNA–metal complexes.

Studies with heat denatured DNA show virtually no reversal in the binding of metal complex to DNA as compared to the native DNA. This suggests that the metal complex is strongly binding to denatured single stranded DNA.

Conclusions

Two water soluble complexes of formula [M(bpy)(cbdca)] which are analogous to carboplatin have been prepared by interacting appropriate \( \alpha \)-diiminePt(II) and Pd(II) compounds with cbdca sodium salt. These complexes have been found to be better cytotoxic agents than cisplatin against P388 lymphocytic leukemia cells. [M(bpy)(cbdca)] complexes bind to calf thymus DNA through coordinate covalent bond as suggested earlier for the binding of cisplatin to DNA.

Acknowledgments

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