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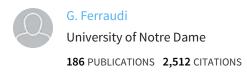
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Preparation and Reactivity of Pendent- -- CO₂Rh^{III}(phthalocyanine) Bound to a Poly(acrylate) Backbone. Effects of the Hypercoiled Backbone on the Association, Photochemical, and Thermal Redox Reactions of the Pendent Macrocycle

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Received June 7, 2006; Revised Manuscript Received July 19, 2006

ABSTRACT: Poly(acrylate) was derivatized with Rh^{III}(pc) (pc = phthalocyanine) and the morphology of the strands was established by AFM microscopy. The strands of the polymer, poly-Rh^{III}(pc), were seen as the strings of spherules expected for an hypercoiled poly(acrylate) backbone. Phthalocyanine pendants in a polymer, poly-Rh^{III}(pc), having a 1:8 stoichiometric relationship of - - -Rh^{III}(pc) to - --CH₂CH(CO₂-)- - - exhibited chemical properties different from those communicated for Rh^{III}(pc-tetrasulfonate)³⁻ in aqueous medium or XRh^{III}(pc), X = Cl, Br, I, in nonaqueous media. Reactions of poly-Rh^{III}(pc) with reducing radicals like a e^{-}_{aq} , (CH₃)₂C*OH, NiL⁺ (L = [14]aneN₄, Me₆[14]aneN₄), or oxidants, i.e., N₃*, Cl₂*-, Br₂*-, or NiL³⁺ were investigated with the pulse radiolysis technique. The phthalocyanine radicals formed in these redox reactions were characterized by means of the transient UV-vis spectrum, and the kinetics of the formation and decay of the radicals were followed at various wavelengths. The photophysical and photochemical processes of poly-Rh^{III}(pc) were studied by laser flash photolysis. The lifetime of the excited state, t = 133 ns, correlates well with the slow component of the decay of the luminescence detected in the NIR region. This transient has been assigned to a ligand field, LF, excited-state not observed in previous works when XRh^{III}(pc) complexes were irradiated in homogeneous solution. The thermal and photochemical properties of poly-Rh^{III}(pc) are discussed on the basis of environmental conditions affecting the pendent macrocycles in the strand of polymer.

Introduction

The study of the chemical properties of metalated and metalfree phthalocyanines remains a very active field of research. Application of the phthalocyanines to numerous fields of chemistry has motivated the interest in these compounds. 1-28 Among these many applications is the catalysis of various reactions in homogeneous solution and heterogeneous phase with transition metal phthalocyanines grafted to organic polymers. Aside the polymerization of pendent groups in a ligand of the complex,²⁹ the grafting of phthalocyanine and other transition metal complexes into polymers has been based in two general strategies.^{23,24,28} They respectively involve the coordination of transition metal complexes through the metal ion to pendent ligands of a polymer backbone or linking one ligand by covalent bonds to a polymer backbone. In the former strategy, metal ions are inert to ligand substitution, e.g., Co^{III}, allowing the complex to be irreversibly fastened to ligands in the polymer. The inertness toward ligand substitution suppresses the detachment of the transition metal complex from the strand of polymer and its migration to the bulk of the solution. On this basis, Rh^{III} was expected to be a convenient metal center for the coordination of the phthalocyanine to a polymer backbone.

Previous studies have shown that monomeric XRh^{III}(pc) complexes (pc = phthalocyanine; X = Cl, Br, I) are inert with regard to the thermal substitution of the ligand X⁻.^{25,26} In addition, the redox chemistry of the XRh^{III}(pc) could be used for the catalysis of processes in homogeneous solution.²⁵ The redox potential of the couples XRh^{III}(pc)/XRh^{II}(pc) and XRh^{IV}-(pc)/XRh^{III}(pc) greatly disfavors changes of the metal center's oxidation state.^{27,28} Thermal or photochemical redox reactions of XRh^{III}(pc) are therefore mediated by XRh^{III}(pc•[±]) ligand

Figure 1. Model showing the average distribution of isolated $Rh^{III}(pc)^+$ groups in a short segment of poly($Rh^{III}(pc)$). To simplify the presentation, no hydrogen atoms and dimers are shown in the structure. The total number of $-CO_2-$ groups and the ratio of $Rh^{III}(pc)+$ to $-CO_2-$ groups in the strand are indicated in the figure.

radical species. Nevertheless trace concentrations of XRh^{II}(pc)⁻, formed after a photoinduced hydrogen abstraction from an organic solvent, accelerate the substitution of the ligand X⁻.

When coordination complexes are hosted by a poly(anion), electrostatic interactions and environmental conditions created by the poly(anion) can change photochemical and thermal reactions of the complexes grafted in the polymer. In this work, the incorporation of ---Rh^{III}(pc) in poly(acrylate), Figure 1, has been investigated as a mean to place the phthalocyanine in aqueous solution and to change the thermal and photochemical reactions of the ---Rh^{III}(pc) group.

⁼Rh(III)
-CO₂-= 6.3 x 10³
Rh(pc) / -CO₂-= 1/8

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Experimental Section

The spectroscopy, photochemistry and redox reactions of the polymeric compounds described below were investigated with freshly prepared aqueous solutions of a pH 6 unless explicitly stated.

Flash Photochemical Procedures. Absorbance changes, ΔA , occurring in a time scale longer than 10 ns were investigated at room temperature with a flash photolysis apparatus described elsewhere. 30 In these experiments, 10 ns flashes of 351 nm light were generated with a Lambda Physik SLL-200 excimer laser. The energy of the laser flash was attenuated to values equal to or less than 20 mJ/pulse by absorbing some of the laser light in a filter solution of Ni(ClO₄)₂ having the desired optical transmittance, T = I_t/I_0 where I_0 and I_t are respectively the intensities of the light arriving to and transmitted from the filter solution. The transmittance, $T = 10^{-A}$, was routinely calculated by using the spectrophotometrically measured absorbance, A, of the filter solution. A right angle configuration was used for the pump and the probe beams. Concentrations of the chromophores in the polymer were adjusted to provide homogeneous concentrations of photogenerated intermediates over the optical path, l = 1 cm, of the probe beam. To satisfy this optical condition, solutions were made with an absorbance equal to or less than 0.4 over the 0.2 cm optical path of the pump. These solutions were deaerated with streams of N₂ before and during the photochemical experiments.

Pulse-Radiolytic Procedures. Pulse radiolysis experiments were carried out with a model TB-8/16-1S electron linear accelerator. The instrument and computerized data collection for time-resolved UV-vis spectroscopy and reaction kinetics have been described elsewhere in the literature. 26-28,31 Thiocyanate dosimetry was carried out at the beginning of each experimental session. The details of the dosimetry have been reported elsewhere.^{31,32} The procedure is based on the concentration of (SCN)₂⁻ radicals generated by the electron pulse in a N_2O saturated $10^{-2}\ M\ SCN^-$ solution. In the procedure, the calculations were made with G=6.13 and an extinction coefficient, $\epsilon = 7.58 \times 10^3 \, \mathrm{M}^{-1} \, \mathrm{cm}^{-1}$ at 472 nm, for the (SCN)₂⁻ radicals. ³²⁻³⁵ In general, the experiments were carried out with doses that in N2 saturated aqueous solutions resulted in $(2.0 \pm 0.1) \times 10^{-6} \text{ M}$ to $(6.0 \pm 0.3) \times 10^{-6} \text{ M}$ concentrations of e⁻_{aq}. In these experiments, solutions were prepared by the procedure indicated above for the photochemical experiments. The liquids were deaerated with streams of the O2-free gas, N2 or N2O, that was required for the experiment. Reactions of the radiolytically generated OH* radicals with N₃⁻, halides or 2-propanol were used for the preparation of reactive radicals, i.e., N₃, Cl₂, Br₂ and $(CH_3)_2C^{\bullet}OH$, eq 1.

radiolytic pulse
$$+ N_2O$$
, $+ H^+$
 $OH^- \dots$
 $X^ X^{\bullet}$
 $OH^ (1)$

 $(X^- = C1^-, Br^-, N_3^-, (CH_3)_2 CHOH \; ; \; X^- = C1^+, Br^-, N_3^-, (CH_3)_2 C^+OH + H^+)$

The oxidation of the Ni(II) complexes with $\text{Cl}_2^{\bullet-}$ radicals was used for the in situ preparation of Ni(III) complexes. The specific conditions used in the preparation of the radicals and Ni(III) complexes are given in the Results.

To radiolyze a fresh sample with each pulse, an appropriate flow of the solution through the reaction cell was maintained during the experiment. Other conditions used for the time-resolved spectroscopy of the reaction intermediates or in the investigation of the reaction kinetics are given in the Results.

The time-resolved changes in the absorption spectrum were modeled with a commercial package, Mathcad 2001i and VisSim V5. A series solution of the rate equations for the reactions described elsewhere in the text and known extinction coefficients of the reactants and reaction intermediates were used in the calculation. ^{25–27}

AFM Microscopy. AFM images were obtained using a Digital Instruments (DI) Multimode Nanoscope III A in the tapping mode. A sharp silicon nitride crystal tip attached to a cantilever was used

as the AFM probe for imaging the polymer on glass or mica slides. The glass slides were immersed in a 4:1 (v:v) H_2SO_4/H_2O_2 mixture for ~ 1 h and then rinsed with distilled water and acetone before the preparation of the samples. The mica slides were treated with 1 M Mg^{2+} before the preparation of the samples. Solutions containing 2 \times 10⁻¹¹ to 3 \times 10⁻⁸ M strands of poly-Rh(pc) in water at a pH between 4 and 9 were evaporated at room temperature on the microscope slides. The evaporation was carried out under a blanket of N_2 . Some of the preparations were also kept under vacuum for several hours to eliminate any trace of solvent.

Materials. The monomer phthalocyanines $XRh^{III}(pc)$ (X = Cl, Br, HSO₄), $Na_3[Rh^{III}(tspc)]$ (tspc = 4,4',4'',4'''-phthalocyaninetetrasulfonate), the Ni^{II} macrocyclic complexes $[Ni^{II}L](ClO_4)_2$ ($L = [14]aneN_4$ (a), $Me_6[14]aneN_4$ (b), and $[Fe^{III}py[14]dieneN_4](ClO_4)_3$ (c)) were available from previous works.

Rh^{III}(pc)-Derivatized Poly(acrylate). The polymeric compounds were prepared by a modification of the literature procedure for the synthesis of $XRh^{III}(pc)$ (X = Cl, Br, I) complexes from HSO₄Rh^{III}(pc).^{25,26} The procedure and quantities described next were used for the preparation of a polymer with a 1:8 stoichiometric relationship of - - -Rh^{III}(pc) to - --CH₂CH(CO₂⁻)- --, Figure 1. A solution containing 200 mg of poly(acrylic acid), Aldrich average $M_{\rm w} \sim 4.5 \times 10^5$, in 200 cm³ of 1:3 (v/v) water in methanol was warmed and neutralized with an equivalent amount of Na₂CO₃, \sim 300 mg. The resulting clear solution was added to a rapidly stirred solution containing 270 mg of HSO₄Rh^{III}(pc) in a minimum volume of a 1:3 (v/v) water in methanol and refluxed for 72 h. Dark solids, formed during the reaction, were removed by centrifugation. The near pure poly-RhIII(pc) was obtained as a dark blue powder after rotovaporation of the solvent. Two further purifications were carried out: first, a fractional recrystallization from aqueous solution followed, second, by dialysis through Spectra/Por membranes MWCO: 3500, and rotovaporation to dryness. The pure material was dried 72 h under vacuum at a pressure equal to or less than 1 Torr. Anal. Calcd for RhC₅₆H₃₀N₈O₁₆Na₇: H, 2.39; C, 49.70; N, 8.29. Found: H, 2.45; C, 49.79; N, 8.17. Other experiments made for the polymers characterization are described in the Results.

The procedure described above, with different molar relations of poly(acrylic acid) and HSO₄Rh^{III}(pc), was used for the preparation of Rh^{III}(pc)-derivatized poly(acrylate) with stoichiometric relationships of ---Rh^{III}(pc) to ---CH₂CH(CO₂⁻)--- between 1:3 and 1:30. Polymers containing stoichiometric relationships of ---Rh^{III}(pc) to ---CH₂CH(CO₂⁻)--- between 1:6 and 1:10 showed similar physical features, i.e., UV—vis spectra and solubilities. Polymers with loads of Rh^{III}(pc) larger than 1:8 proved to be unsuitable for the studies described here.

Other materials and solvents were reagent grade and used without further purification. Solutions for the experiments described herein were prepared with triple distilled water.

Results

Poly(acrylate) polymers containing various loads of ---Rh^{III}(pc) coordinated to pendent ---CO₂⁻ groups, Figure 1, were obtained by the general procedure described in the Experimental section. Polymers containing loads larger than 1:10 stoichiometric relationships of ---Rh^{III}(pc) to acrylate

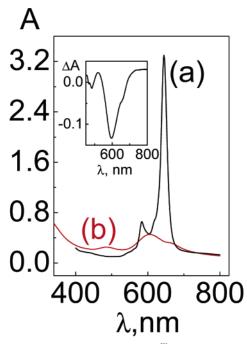


Figure 2. Spectra of monomeric HSO₄Rh^{III}(pc) (a) and poly-Rh^{III}(pc) (b) in solutions containing 5×10^{-5} M Rh(III) chromophores. The inset shows the difference spectrum of two solutions of poly- $Rh^{III}(pc),\, 5\times 10^{-5}\,M$ in - - -CO2Rh^{III}(pc) chromophores. The spectrum of one solution was subtracted from the spectrum of another containing 10^{-2} M UO₂²⁺ in addition to the poly-Rh^{III}(pc).

units, ---CH₂CH(CO₂-)---, were in general too insoluble in aqueous media and their properties were not investigated. In contrast to these compounds, those with loads equal to or smaller than 1:10 exhibited good solubilities in aqueous media. Phthalocyanine pendants in a polymer with a 1:8 stoichiometric relationship of - - -Rh^{III}(pc) to - - -CH₂CH(CO₂⁻)- - - exhibited chemical properties different of the Rh^{III}(tspc)³⁻ in aqueous medium or the $XRh^{III}(pc)$, X = Cl, Br, I, in nonaqueous media. The properties of this particular polymer, named poly-Rh^{III}(pc), are described next.

Stacking of - - - CO₂Rh^{III}(pc). A less intense and somehow broader Q-band than in the spectrum of HSO₄Rh^{III}(pc) is seen in the spectrum of poly-Rh^{III}(pc), Figure 2. The Rh^{III}(tspc)³⁻ spectrum in aqueous solution shows an intense absorption band at \sim 640 nm, i.e., the Q-band of the phthalocyanine ligand.²⁷ It is assigned to the 0-0 vibronic transition from the ¹A ground state to the first ${}^1\pi\pi^*$ excited state, Figure 2a. Other vibronic components of the band are seen at shorter wavelengths. In metallophthalocyanines, the broadening of the Q-band has been associated with the stacking of metallophthalocyanines in dimers containing two or more units of the macrocyclic complex. The broad Q-band observed in the spectrum of poly-Rh^{III}(pc) must also be associated with a stacking of metallophthalocyanine pendants. Given the reduced load of -Rh^{III}(pc) and limits to the flexibility of the backbone, only the dimers are probably formed by the π stacking of two phthalocyanine pendants. Further evidence of the - - - CO₂Rh^{III}(pc) stacking in poly-Rh^{III}-(pc) was provided by changes in the Q-band of poly-Rh^{III}(pc) UV-vis when aqueous solutions of the polymer contained metal ions, e.g., UO22+, capable of forming complexes with carboxylates. A decrease in the intensity of the Q-band, inset to Figure 2, shows that the stacking of - - -CO₂Rh^{III}(pc) pendants in the polymer strand increases when UO22+ is complexed by free carboxylate groups. Comparisons were made between the spectra of solutions having the same ionic strength, $I = 10^{-3}$ M, adjusted with NaClO₄. Changes in the UV-vis spectrum of

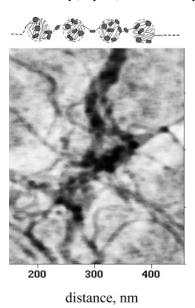


Figure 3. Typical AFM image of poly-Rh^{III}(pc) showing strings of spherules, bottom. The sample was prepared with a procedure described elsewhere in the text from a solution containing 2.5×10^{-8} M strands of poly-Rh $^{\rm III}$ (pc). The data scale is 4.0 nm. The approximate distribution of $-CO_2Rh^{III}(pc)$ pendants, blue circles, in the hypercoiled strand are shown in the top figure.

poly-Rh^{III}(pc) were also induced with 10^{-5} – 10^{-4} M Ni(II) macrocyclic complexes a, b, or c. Changes in the spectrum of poly-Rh^{III}(pc) caused by any of these Ni(II) complexes bear a strong resemblance to those observed in the presence of UO_2^{2+} .

In comparing this behavior to the behavior of the metallophthalocyanines in solution phase, the stacking of the - - - CO₂Rh^{III}(pc) pendants demands large concentrations of the pendant in the strand of poly-Rh^{III}(pc). Because of the low load of - - -CO₂Rh^{III}(pc) in poly-Rh^{III}(pc), a particular morphology of the strand is necessary to cause the high concentration of pendants.

Strand Morphology. The morphology of the polymer strands was investigated in an AFM microscope in the tapping mode. A string of spherules was observed in samples prepared with different procedures, Figure 3. These strings of spherules indicated the hypercoiling of the poly(acrylate) backbone in a manner resembling the hypercoiled poly(metaacrylate).^{36,37}

Reactivity of Poly-Rh^{III}(pc) with Radicals. The spectroscopy of the products formed when reductants, i.e., e-aq or $(CH_3)_2C^{\bullet}OH$, NiL⁺ (L = [14]aneN₄, Me₆[14]aneN₄), or oxidants, i.e., N₃•, Cl₂•-, Br₂•-, or NiL³⁺, react with poly-Rh^{III}(pc) and the kinetics of the products formation were investigated with the technique of pulse radiolysis. Transient absorption spectra resulting from the oxidation of poly-Rh^{III}(pc) by N₃*, Cl₂•-, or Br₂•- were recorded with solutions of poly-Rh^{III}(pc) having $\sim 10^{-4}$ M - - -CO₂Rh^{III}(pc) chromophores and 0.1 M Cl⁻, $0.1~M~N_3^-$, or $0.05~M~Br^-$. The reactions of these oxidants produce a bleach of the Q-band in the near-IR and a broad absorption band with a maximum at \sim 500 nm, Figure 4. This spectrum is similar to the literature spectrum of ClRh^{III}(pc•+) and it has been attributed to the formation of - - -CO₂Rh^{III}(pc•+) chromophores.²⁷

The kinetics of the poly-Rh^{III}(pc) oxidation by N₃• radicals was investigated at $l_{\rm ob} = 500$ nm where the difference spectrum of - - -CO₂Rh^{III}(pc•+) chromophores exhibits a maximum. The oscillographic traces used for the calculation of the rate constant were obtained by radiolyzing N2O saturated solutions of poly-Rh^{III}(pc) containing 2.9×10^{-5} M to 2.4×10^{-4} M

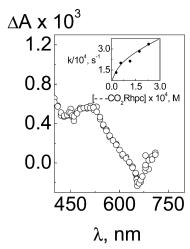


Figure 4. Transient absorption spectrum of - - - CO₂Rh^{III}(pc•+) radicals in the strand of polymer. The radicals were formed when poly-Rh^{III}(pc) was oxidized by pulse radiolytically generated N₃* radicals. In this experiment, N₂O-saturated solutions of poly-Rh^{III}(pc), 1.0 × 10⁻⁴ M in ---CO₂Rh^{III} (pc) chromophores, contained 0.1 M NaN3. The inset shows the dependence of the ---CO₂Rh^{III}(pc•+) radical formation rate constant on the - - -CO₂Rh^{III}(pc) concentration.

---CO₂Rh^{III}(pc) with an ionic strength, I = 0.1 M, adjusted with NaN₃, Figure 4. A nonlinear dependence of the rate constant with the concentration of chromophores was ascribed to poly-RhIII(pc) concentration-dependent changes in the - - - CO₂Rh^{III}(pc) stacking within the strand.

The oxidation of poly-Rh^{III}(pc) by Ni(III) macrocyclic complexes, is markedly different of the reactions with N₃°, Cl₂°-, or $\mathrm{Br_2}^{\bullet-}$ described above. Solutions containing 0.1 M Cl⁻, 10^{-2} M NiL²⁺, and 2.4 \times 10⁻⁴ M - - -CO₂Rh^{III}(pc) were saturated with N₂O and pulse radiolyzed. The rapid growth of the NiL³⁺ spectrum was observed on a time scale t < 400 ns, Figure 5a. The formation of the Ni(III) products occurs with a pseudo firstorder kinetics. Second-order rate constants, $k = 1.2 \times 10^9 \,\mathrm{M}^{-1}$ s^{-1} for L = Me₆[14]aneN₄ and $k = 2.4 \times 10^9 M^{-1} s^{-1}$ for L = [14]aneN₄, were calculated from the rate constants obtained under the pseudo first-order kinetics. Further spectral changes took place in a time scale t > 400 ns due to the reaction of the NiL³⁺ complexes with - - -CO₂Rh^{III}(pc) chromophores of the poly-Rh^{III}(pc), Figure 5b. A recovery of the solution's absorbance at wavelengths of the Q-band and an absorption band, $\lambda_{\text{max}} = 530 \text{ nm}$, were observed at the end of the reaction. The rate of the spectral changes exhibited a first-order kinetics and the rate constant of the process, $k = 4.4 \times 10^6 \text{ s}^{-1}$, was independent of λ_{ob} between 450 and 600 nm. Similar spectral changes were observed when solutions containing 0.1 M Cl⁻, 10^{-2} M NiL²⁺, and 2.4×10^{-4} M Rh^{III}(tspc)³⁻ were saturated with N₂O and pulse radiolyzed. However, the oxidation of NiL²⁺ was faster, $k \sim 10^{10} \text{ M}^{-1} \text{ s}^{-1}$, and the formation of the adduct was slower, $k = 10^5 \,\mathrm{s}^{-1}$, than that in the presence of poly-Rh^{III}(pc).

In contrast to the facile oxidation of - - - CO₂Rh^{III}(pc) chromophores by the above-mentioned radicals, poly-Rh^{III}(pc) proved to be more discriminative toward electron donors. Neither e-aq nor NiL+ macrocyclic complexes were observed to react with solutions of poly-Rh^{III}(pc) where the concentration of - - - CO₂Rh^{III}(pc) chromophores was equal to or less than 2.4 \times 10⁻⁴ M. However, the spectrum of the reduced chromophore ---CO₂Rh^{III}(pc•-), similar to the literature spectrum of ClRh^{III}(pc^{•-}), was observed when N₂O saturated solutions of poly-Rh^{III}(pc), 2.4×10^{-4} M in - - -CO₂Rh^{III}(pc) chromophores, reacted with pulse radiolytically generated (CH₃)₂C•OH radicals.

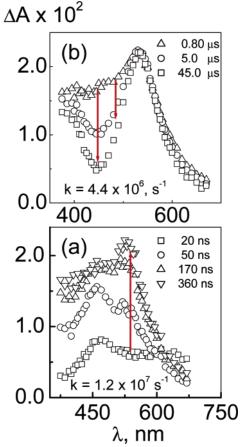


Figure 5. Transient spectra recorded when pulse radiolytically generated Ni^{III}(Me₆[14]dieneN₄)Cl²⁺ reacted with poly-Rh^{III}(pc). The N₂O-saturated solution was 0.1 M in Cl⁻ and it also contained 10⁻³ M $Ni^{III}(Me_6[14]dieneN_4)^{2+}$ and a concentration of poly-Rh^{III}(pc) equivalent to 1.0×10^{-4} M- - -CO₂Rh^{III}(pc) chromophores.

An overall rate constant, $k = 7.5 \times 10^9 \,\mathrm{M}^{-1} \,\mathrm{s}^{-1}$, was calculated for the reaction of (CH₃)₂COH radicals with poly-Rh^{III}(pc) monomeric pendants, - - - CO₂Rh^{III}(pc), and dimeric pendants, (---CO₂Rh^{III}(pc))₂, eqs 2 and 3.

$$---CO_{2}Rh^{III}(pc) + (CH_{3})_{2}C^{\bullet}OH \xrightarrow{-H^{+}}$$

$$---CO_{2}Rh^{III}(pc^{\bullet^{-}}) + (CH_{3})_{2}CO \quad (2)$$

$$(---CO_{2}Rh^{III}(pc))_{2} + (CH_{3})_{2}C^{\bullet}OH \xrightarrow{-H^{+}}$$

$$(CH_{3})_{2}CO + ---CO_{2}Rh^{III}(pc) + ---CO_{2}Rh^{III}(pc^{\bullet^{-}}) \quad (3)$$

Photophysical and Photochemical Processes. Deaerated solutions of poly-Rh^{III}(pc), 2.4×10^{-4} M in - - -CO₂Rh^{III}(pc) pendants, were flash irradiated at 351 nm. The irradiation produced a transient absorption spectrum with a broad band centered at ~500 nm, Figure 6. The decay of the spectrum was monitored at several wavelengths, $\lambda_{ob} = 450$, 500, and 550 nm. Oscillographic traces collected at these wavelengths showed the same rate of decay. A rate constant, $k = 7.5 \times 10^6 \text{ s}^{-1}$, was calculated from the lifetime, $\tau = 133$ ns, of the absorbance decay. To correlate these transients to excited-state processes, the poly-Rh^{III}(pc) luminescence was investigated with a solution of the polymer, 2.4×10^{-5} M in - - -CO₂Rh^{III}(pc) pendants, irradiated at 337 nm. The decay of the luminescence exhibited two components, one faster than the other. A slow component of the emission decay was detected in the NIR region. The lifetime of the NIR emission was approximately the same, $\tau =$ 140 ns, communicated above for the decay of the transient

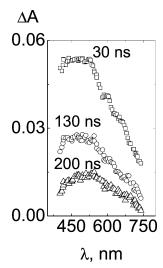


Figure 6. Photogenerated transient spectra recorded when deaerated solutions of poly-Rh^{III}(pc), 2.4 × 10⁻⁴M in - - -CO₂Rh^{III}(pc) pendants, were flash irradiated at 351 nm.

spectrum. This component of the luminescence was not previously observed with $XRh^{III}(pc)$ (X = Cl, Br, I). In addition, a fast decaying component of the emission, t < 20 ns, exhibited a maximum at ~440 nm. It was assigned to the radiative relaxation of the second $1\pi\pi^*$ excited state. This is the same anti-Kasha emission previously communicated for XRh^{III}(pc) $(X = Cl. Br. I)^{26}$

The spectra of the pendent ligand radicals, - - - CO₂Rh^{III}(pc•-) or --- CO₂Rh^{III}(pc•+), were recorded in flash photochemical experiments where poly-Rh^{III}(pc), were flash irradiated in solutions containing $2.4 \times 10^{-4} \text{ M}$ - - -CO₂Rh^{III}(pc) pendants and either an electron donor or an electron acceptor, eq 4 and 5.

$$--CO_{2}Rh^{III}(pc) + h\nu \rightleftharpoons ---(---CO_{2}Rh^{III}(pc))^{*} \stackrel{S}{\rightarrow}$$
$$---CO_{2}Rh^{III}(pc^{\bullet-}) + S^{\bullet+}$$
 (4)
$$S = TEA \text{ or } TEOA$$

$$--CO_{2}Rh^{III}(pc) + h\nu \rightleftharpoons --(--CO_{2}Rh^{III}(pc))^{*} \stackrel{S}{\rightarrow}$$
$$---CO_{2}Rh^{III}(pc^{\bullet+}) + S^{\bullet-} (5)$$
$$S = MV^{2+} \text{ or } Fe^{III}(pv[14]dieneN_{4})^{3+}$$

The various concentrations used in the reduction, eq 4, and oxidation, eq 5, of the excited state were 1 M TEA (TEA = triethylamine); 0.1, 0.4, and 1.0 M TEOA (TEOA = 2,2',2''triethanolamine); 10^{-2} M MV²⁺ (MV²⁺ = methyl viologen); and 10^{-4} M Fe^{III}(py[14]dieneN₄)³⁺.

In contrast to the reactions of the polymer with pulse radiolytically generated radicals, the formation of - - - CO₂Rh^{III}- $(pc^{\bullet-})$ and - - -CO₂Rh^{III} $(pc^{\bullet+})$ occurred in a time scale, $t = 10^2$ ns. These rates were fast by comparison to those of the reactions between poly-Rh^{III}(pc) and pulse radiolytically generated radicals. The difference was attributed to the presence of the electron donor or acceptor S near the electronically excited pendants, (---CO₂Rh^{III}(pc))*, in the photochemical processes. Back electron-transfer reactions were observed in a time scale, t >10² ns. These processes exhibited second-order kinetics. In Table 1, $k/\Delta\epsilon$ is the ratio of the second-order rate constant to the difference, $\Delta \epsilon$, between the extinction coefficients of the radicals and the polymer. It was calculated at wavelengths of the maximum in the difference spectrum of - - -CO₂Rh^{III}(pc•-) or - - - CO₂Rh^{III}(pc•+). Since the pulse radiolytic experiments yield

Table 1. Ratio of the Rate Constant to the Difference of Extinction Coefficient, $k/\Delta\epsilon$, of the Coordinated pc Radical at 470 nm^a

reaction	$k/\Delta\epsilon$, cm s ⁻¹
$\begin{split} &TEA^{\bullet+} +CO_2Rh^{III}(pc^{\bullet-}) \to TEA +CO_2Rh^{III}pc \\ &TEOA^{\bullet+} +CO_2Rh^{III}(pc^{\bullet-}) \to TEOA +CO_2Rh^{III}pc \\ &MV^{\bullet+} +CO_2Rh^{III}(pc^{\bullet+}) \to MV^{2+} +CO_2Rh^{III}pc \\ &Fe^{II}(py[14]dieneN_4)^{2+} +CO_2Rh^{III}(pc^{\bullet+}) \\ &\to Fe^{III}(py[14]dieneN_4)^{3+} +CO_2Rh^{III}pc \end{split}$	$\begin{array}{c} 1.3 \times 10^8 \\ 2.4 \times 10^8 \\ 4.5 \times 10^8 \\ 2.7 \times 10^8 \end{array}$

^a The rate constant is for the back-electron-transfer reaction of the photooxidized or photoreduced - -- CO₂Rh^{III}(pc) pendants.

an approximate value for the difference of extinction coefficients, $\Delta \epsilon \sim 10^3 \ \mathrm{M}^{-1} \ \mathrm{cm}^{-1}$, the second-order rate constants calculated with values of $k/\Delta\epsilon$ in Table 1 must have a $10^{11} \, M^{-1}$ s^{-1} order of magnitude.

Discussion

The UV-vis spectroscopy of poly-Rh^{III}(pc) shows a large fraction of --- CO₂Rh^{III}(pc) pendants forming stacks in the strand of poly(acrylate) polymer. A similar stacking has not been observed in homogeneous solutions of $XRh^{III}(pc)$ (X = Cl, Br, I), and it must be regarded as an intrinsic phenomenon of the poly-Rh^{III}(pc). The stacks of ---CO₂Rh^{III}(pc) pendants are likely to be located inside the spherules observed in the AFM microscope. This distribution of the pendants in the strand of polymer is favored by the hydrophobic character of - - -CO₂Rh^{III}(pc) pendants; i.e., it will isolate them inside the spherules from the aqueous media. Constrained to the small volume of the spherules, the - - - CO₂Rh^{III}(pc) pendants achieve local concentrations far larger than those of the XRh^{III}(pc) (X = Cl, Br, I) complexes in homogeneous solution, a condition also favoring the formation of stacks. Bonding between - - - CO₂Rh^{III}(pc) pendants in the stacks must also be responsible of the persistence for the hypercoiled morphology when carboxylate pendants, not coordinated to Rh^{III}(pc), are deprotonated at pH > 6. Indeed, the bonding between - - - CO₂Rh^{III}(pc) pendants in a stack is, to a large extent, pH independent. In acidic media, e.g., pH = 6, the H-bonds from the protonated carboxylates could make an additional contribution to the hypercoiled structure of poly-Rh^{III}(pc).

Hypercoiling and Redox Reactions. The hypercoiled structure of poly-Rh^{III}(pc) accounts for the diverse redox behavior of the polymer. Oxidation of the --- CO₂Rh^{III}(pc) pendants is most likely to be carried out in the spherules by neutral X^o radicals, eq 6

- - -CO₂Rh^{III}(pc) + X[•] - - -CO₂Rh^{III}(pc^{•+}) + X⁻ (6)

$$X^{\bullet} = Cl^{\bullet}, Br^{\bullet}, N_3^{\bullet}; X^{-} = Cl^{-}, Br^{-}, 3/2 N_2$$

that are hydrophobic in nature and exist in equilibrium with the spectroscopically observable $X_2^{\bullet-}$ anion radicals, eq 7.

$$X^- + X^{\bullet} \hookrightarrow X_2^{\bullet -}; X^{\bullet} = Cl^{\bullet}, Br^{\bullet}$$
 (7)

Reactions of the larger hydrophilic anion radicals with - -- CO₂Rh^{III}(pc) pendants are expected to be kinetically disfavored by electrostatic interactions and the hydrophobic environment hosting the stacks. In contrast to the - - - CO₂Rh^{III}-(pc•+) radical produced in eq 6, the reactions of poly-Rh^{III}(pc) or Rh^{III}(tspc)³⁻ with Ni(III) macrocyclic complexes gave products with a spectrum having a band at ~530 nm and no bleach of the \sim 675 nm pc's or tspc's Q-band. The spectrum is devoid, therefore, of the characteristic features of the radical's spectrum. On the basis of this spectroscopic features, the products must be assigned as adducts of the Ni(III) complex with $Rh^{III}(tspc)^{3-}$ or ---CO $_2Rh^{III}(pc)$ pendants in poly-Rh $^{III}(pc)$. A single equilibrium, eq 8

$$Rh^{III}(tspc)^{3-} + NiL^{3+} \rightleftharpoons [Rh(tspc)\cdots NiL]$$

$$L = [14]aneN_4, Me_6[14]aneN_4$$
(8)

accounts for the spectral changes observed in the reaction of Rh^{III}(tspc)³⁻ with NiL³⁺ to form the adduct [Rh^{III}(tspc)···NiL].

The mechanism of the reaction of NiL³⁺ with poly-Rh^{III}(pc) is more complex than eq 8. It must first involve the coordination of NiL²⁺ and NiL³⁺ to carboxylates not bonded to Rh^{III}(pc) by a process similar to the one observed with the unreactive UO₂²⁺, Figure 2. Coordination of the Ni macrocycles to the strand of polymer makes the formation of the adduct more than 1 order of magnitude faster, i.e., 10^6 s⁻¹ vs 10^5 s⁻¹, than eq 8. In a summarized manner, the processes initiated with Ni^{II}(Me₆[14]-aneN₄)²⁺ oxidation and formation of the adduct are described in Scheme 1, where radiolytically generated OH• radicals oxidize $X = Cl^-$ or Br^- to produce radicals X^\bullet . The radical X^\bullet oxidize Ni^{II}(Me₆[14]aneN₄)²⁺ which will form the corresponding adducts with pendent - - -CO₂Rh^{III}(pc).

Modeling of the overall reaction between (CH₃)₂C•OH radicals and - - - $CO_2Rh^{III}(pc)$ was made on the assumption of different contributions from the reaction of the radicals with the dimers to form - - - CO₂Rh^{III}(pc•-) radicals and monomeric - - - CO₂Rh^{III}(pc), eq 3. The model providing the best agreement with the time-resolved bleach of the Q-band at ~ 650 nm and the formation of the --- CO₂Rh^{III}(pc•-) radical absorption at \sim 500 nm, Figure 7a, suggests a significant participation of the dimers in the overall reaction, eqs 2 and 3. If the reaction of the dimers with the radical is diminished or ignored altogether, the bleach of the monomer's Q-band became too large in relationship to the absorption of the - - -CO₂Rh^{III}(pc•-) radical, Figure 7b. In the other limit, a contribution too large from the dimer's reaction, eq 3, gives an absorbance growth at \sim 650 nm, Figure 7c. This new absorption band results from an overestimated generation of - -- CO₂Rh^{III}(pc) monomer which has a much larger extinction coefficient at ~ 630 nm than the (- - -CO₂Rh^{III}(pc))₂ dimer. The best fit of the spectral changes yields rate constants $k = 5.0 \times 10^9 \,\mathrm{M}^{-1} \,\mathrm{s}^{-1}$ for eq 2 and $k = 10^{-1} \,\mathrm{m}^{-1}$ $8.0 \times 10^8 \text{ M}^{-1} \text{ s}^{-1} \text{ for eq 3.}$

Hypercoiling and Excited-State Processes. In aqueous solution, stacks of ---CO₂Rh^{III}(pc) pendants in hydrophobic pockets of the strand account for differences among the

photophysical processes of the pendants and those previously communicated for $XRh^{III}(pc)$ (X = Cl, Br, I). These differences are shown in Figure 8. The most marked difference is the photogeneration of an excited state with the spectrum and lifetime of a ligand field, LF, excited-state not observed when

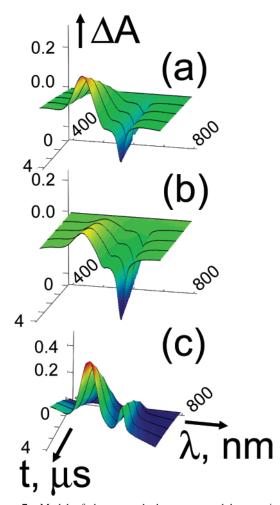


Figure 7. Model of the spectral changes caused by reactions of $(CH_3)_2C^{\bullet}OH$ radicals with monomeric and dimeric --- $CO_2Rh^{III}(pc)$ pendants. A rate constant, $k = 5 \times 109 \text{ s}^{-1}$, was used for the reaction of $(CH_3)_2C^{\bullet}OH$ radicals with monomeric --- $CO_2Rh^{III}(pc)$ pendants. The rate constant used for the reaction of $(CH_3)_2C^{\bullet}OH$ radicals with --- $(--CO_2Rh^{III}(pc))_2$ pendants is: $1 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$ (a), $8 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$ (b), and $3 \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$ (c).

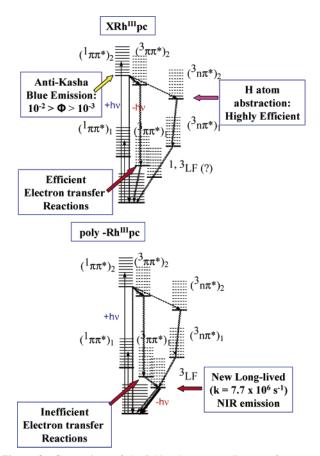


Figure 8. Comparison of the Jablonsky energy diagrams for monomeric XRh^{III}(pc), top, and ---CO₂Rh^{III}(pc) pendants in poly-RhIII(pc), bottom. The question mark in the top diagram shows where LF excited states of Rh(III) could be expected to be in the XRh^{III}(pc)

XRh^{III}(pc) complexes were irradiated in homogeneous solution. In addition the efficiency of the lowest phthalocyaninecentered excited state, $({}^{3}\pi\pi^{*})_{1}$, to undergo electron-transfer reactions is decreased in relation to similar reactions of the $(^{3}\pi\pi^{*})_{1}$ in XRh^{III}(pc). The decrease in the efficiency $(^{3}\pi\pi^{*})_{1}$ to undergo electron-transfer reactions can be related to the $(^{3}\pi\pi^{*})_{1}$ to ^{3}LF conversion in poly-Rh^{III}(pc), a process that was not previously observed in the photophysics of XRh^{III}(pc).

$$---Rh^{III}(pc^{\bullet\pm}) + S^{\bullet\mp} \rightarrow ---Rh^{III}(pc) + S$$
 (9)

Although the electron-transfer processes of the $(^3\pi\pi^*)_1$ with electron donors or acceptors is inefficient, the formation products, eqs 1 and 2, and the back-electron-transfer reactions of the phthalocyanine radicals, eq 9, were followed by means of the intense absorbance change at wavelengths of the Q-band. The reactions depicted in eq 9 occur with faster rates, rate constants $k = 10^{11} \text{ M}^{-1} \text{ s}^{-1}$, than those of related reactions between the XRh^{III}(pc $^{\bullet\pm}$) and S $^{\bullet-}$ radicals, rate constants k= 10^{7} or 8 M⁻¹ s⁻¹. The faster rates suggest that the radicals S^{•-} are not able to diffuse too far into the bulk of the solution before they react back with the - - - CO₂Rh^{III}(pc^{•±}) radicals in the strand of polymer.

Conclusions

In relation to poly(acrylate), the phthalocyanine pendants play an important stabilizing role in the strand's hypercoiled morphology. Environmental conditions created by the hypercoiling change the thermal and photochemical reactivity of pendent ---CO₂Rh^{III}(pc) groups. To a large extent, these changes in the reactivity of the - - - CO₂Rh^{III}(pc) pendants appear to be the

dual result of the medium and steric effects. They can be associated with a large number of pendants inside the spherules of the hypercoiled strand and placed, therefore, in a medium much different from the bulk of the solution.

Acknowledgment. The work described herein was supported by the Office of Basic Energy Sciences. of the US Department of Energy. This is Contribution No. NDRL-4668 from the Notre Dame Radiation Laboratory.

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MA061264U