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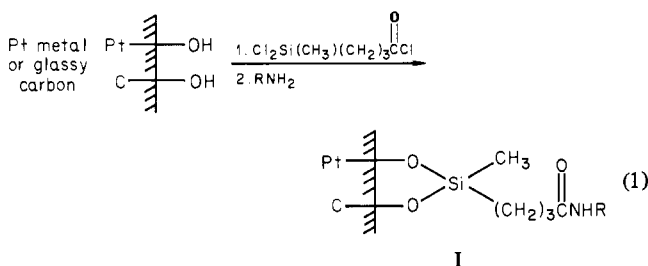
Electronic and Photoelectron Spectral Studies of Electroactive Species Attached to Silanized C and Pt Electrodes

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Abstract: Depending on silanization reaction conditions, aminophenylferrocene and tetrakis(*p*-aminophenyl)porphyrin can be covalently attached in monolayer or multimolecular layer quantities to glassy carbon and superficially oxidized Pt electrodes by using 4-(methyldichlorosilyl)butyryl chloride. This study applies various spectroscopies to surface structure of the attached porphyrin: XPS (number of surface amide bonds), fluorescence (quantity of porphyrin attached to a siloxane polymer film compared to quantity of electroactive porphyrin), and reflectance spectroscopies (spectroelectrochemical observation of oxidation state changes with electrode under potential control, including an electrochemically silent Co(III/II) porphyrin reaction). Electroactive films 500 monolayers thick can be prepared from the ferrocene by spin coating reactive silane; in aqueous sulfuric acid voltammetry of these films indicates a phaselike property.

This is a study of surface chemistry, electrochemistry, and spectroscopy of amine-substituted electron-transfer couples (RNH₂), attached to superficially oxidized glassy carbon and platinum electrodes, using 4-(methyldichlorosilyl)butyryl chloride (designated BCl silane). When RNH₂ is tetrakis(*p*-amino-



phenyl)porphyrin, I is designated Pt,C/ \sim (NH₂)₄TPP; when RNH₂ is *p*-aminophenylferrocene, I is Pt,C/ \sim NH₂PhCpFeCp. Amine derivatives constitute an important pathway to attaching redox molecules to electrode surfaces and have been used to immobilize chiral² and redox³⁻⁷ reagents on thionyl chloride ac-

tivated carbon and in chemisorption on Pt oxide⁸ and carbon⁹ surfaces. The acid chloride BCl silane is a general surface linking reagent for monolayers of amines on both metal oxides¹⁰ and carbon, and as shown here can also be used in a spin coated polymerized (siloxane) form to attach multilayers of amines to electrodes. Earlier reactions of organosilane reagents with glassy carbon surfaces¹¹ ultimately proved ineffectual for immobilizing redox materials.¹² Reaction 1 is the first example of immobilizing redox substances on glassy carbon with organosilane chemistry.

Spectroscopic data to further our understanding of electron-transfer reagents attached to electrodes is at present limited, especially with respect to correlations with electrochemical behavior and solution spectra. We apply here spectroscopic measurements, to Pt/ \sim (NH₂)₄TPP surfaces, not previously employed on electrochemically active modified electrodes. We report the first example of reflectance spectroscopy of a covalently assembled modified electrode, including spectroelectrochemical experiments in which the electrode is under active potential control so changes in surface oxidation state can be spectrally as well as electrochemically followed. Secondly, we have used a fluorescence technique to compare quantitatively the amount of (NH₂)₄TPP actually bound to Pt with that which is electrochemically active.

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Table I. Electrochemical Data for BCl Silane Attached Aminophenylferrocene and (NH₂)₄TPP

entry	electrode	solvent	E°_{surf} vs. SSCE ^a	ΔE_p , mV ^b	$\Gamma \times 10^{10}$, mol/cm ²	E°_{soln}
1	C/ \sim NH ₂ PhCpFeCp	CH ₃ CN	+0.46	10	4.5	+0.40 ^c
2	Pt/ \sim NH ₂ PhCpFeCp	CH ₃ CN	+0.43	23-176	60-1500	+0.40 ^c
3	Pt/ \sim (NH ₂) ₄ TPP	Me ₂ SO	-1.11, -1.49	20	5.7	-1.08, -1.45 ^d
4	C/ \sim (NH ₂) ₄ TPP	Me ₂ SO	-1.10, -1.48	25, 20	2.5	-1.08, -1.45 ^d
5	C/ \sim (NH ₂) ₄ TPP	CH ₃ CN	-1.24, -1.62	40	4.5	
6	C/ \sim (Co)(NH ₂) ₄ TPP	Me ₂ SO	-0.85	50	4.1	-0.82 ^c
7	C/ \sim (Ni)(NH ₂) ₄ TPP	Me ₂ SO	-1.19, -1.75	15	3.6	-1.18, -1.75 ^d
8	C/ \sim (Cu)(NH ₂) ₄ TPP	Me ₂ SO	-1.18, -1.64		3.8	-1.20, -1.68 ^d
9	C/ \sim (Zn)(NH ₂) ₄ TPP	Me ₂ SO	-1.37, -1.73		3.2	-1.31, -1.72 ^d

^a Average potential of cathodic and anodic peaks. ^b Difference in potential of cathodic and anodic peaks. ^c Solution analogue is 4-acetamidophenylferrocene. ^d Felton, R. H.; Linschitz, H. *J. Am. Chem. Soc.* 1966, 88, 1113.

Thirdly, we have applied an X-ray photoelectron spectroscopy (XPS) technique¹² to determine the average number of surface amide bonds formed by (NH₂)₄TPP in reaction 1. Besides yielding useful surface structural insights, the spectroscopic experiments reveal some interactions between the attached molecules and the electrode.

Experimental Section

Preparation of Modified Glassy Carbon Electrodes. Polished glassy carbon (Atomergic Grade V-10, 2.6 mm diameter) cylinders were oxidized in a low wattage 380 mtorr O₂ radiofrequency plasma for 30 min, reacted with a 5% solution of BCl silane in toluene at room temperature for 1 h, rinsed briefly with toluene, then reacted with ~4 mM aminophenylferrocene at 25 °C or (NH₂)₄TPP in refluxing toluene for ~2 h, thoroughly washed with toluene and methanol, and stored under CH₃CN. C/ \sim (NH₂)₄TPP surfaces were metalated by reaction with CoCl₂ or CuCl₂ solutions.³

Preparation of Modified Pt/PtO Electrodes. For reflectance spectroscopy and cyclic voltammetry, the surface of a superficially oxidized 0.29 cm² Pt disk, mirror polished (1 μm diamond compound) and Teflon shrouded, was covered with a drop of neat BCl silane and allowed to stand in laboratory air for ~1.5 min. Excess silane was rinsed off with a stream of toluene and a drop of solution of 1 mg of (NH₂)₄TPP in 5 mL of toluene was allowed to stand on the Pt surface for ~5 min. If the porphyrin became protonated (green), a fresh drop was added. The amidized surface was thoroughly washed with DMF which is effective at removal of adsorbed porphyrin. The surfaces were metalated by warming (40–50 °C) in a 0.5 M solution of CoCl₂ or CuCl₂ for 2 days or several hours, respectively, followed by washing in DMF.

XPS of dangling amine groups on Pt/ \sim (NH₂)₄TPP was conducted by reacting the surface with 0.1 M 3,5-dinitrobenzoyl chloride in toluene for 1 h, washing with toluene, and observing the N 1s spectra with a Du Pont Model 650B electron spectrometer.

Preparation of Pt/ \sim (NH₂)₄TPP surfaces for fluorescence experiments was as above, except Pt flag electrodes were immersed in neat BCl silane and then (NH₂)₄TPP–toluene solution.

Spin Coating of BCl Silane on Pt. A polished, Teflon-shrouded, 0.16 cm² disk, preoxidized with HNO₃, was spun at 28 000 rpm in a 6 N HCl-saturated N₂ atmosphere. A specific number of droplets of 10% BCl silane in benzene was slowly delivered to the center of the horizontally spinning electrode. This surface was exposed to a refluxing benzene solution of aminophenylferrocene for 30 min and washed with benzene and CH₃CN. Depending on the number of BCl silane droplets applied, the still highly reflecting Pt surface is either metallic or shiny golden in color.

Chemicals. Tetrakis(*p*-aminophenyl)porphyrin [(NH₂)₄TPP] was synthesized¹³ and purified as before.^{3b} Aminophenylferrocene was a gift of Professor W. F. Little, University of North Carolina, Chapel Hill. The BCl silane, 4-(methylchlorosilyl)butyryl chloride (Silar), used as received, was stored under N₂. Toluene and benzene were dried by Na^o; CH₃CN, DMF and Me₂SO were stored over molecular sieves.

Reflectance Spectroscopy. The unpolarized beam of a computer-interfaced Harrick rapid scanning spectrometer¹⁴ was reflected at near-normal incidence from a Pt/ \sim (NH₂)₄TPP or Pt/ \sim (M)(NH₂)₄TPP modified surface mounted in the cell of Figure 1. This cell has connections for auxiliary and reference (NaCl-SCE; SSCE) electrodes so the potential of the modified electrode surface can be controlled while acquiring reflectance spectra. About 500 spectra can be acquired, av-

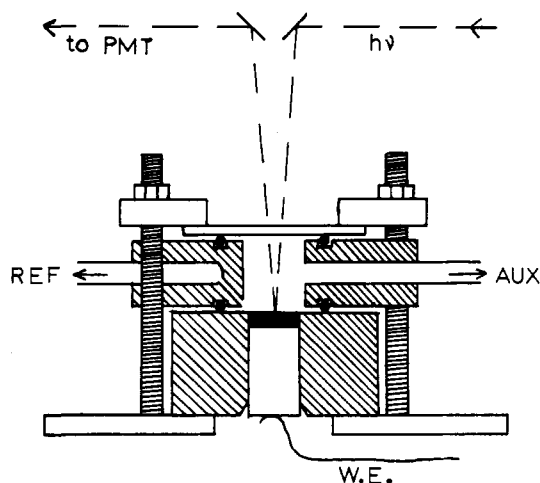


Figure 1. Cell for reflectance spectroelectrochemistry. Quartz window, 1 cm of deaerated supporting electrolyte/solvent. Cross-hatched: Teflon, solid: modified Pt electrode.

eraged, and stored in 15 s. A background spectrum was similarly acquired for subtraction from the modified electrode spectrum by removing, polishing, and carefully replacing the Pt disk in the cell. Comparison spectra of (NH₂)₄TPP and Co(NH₂)₄TPP solution in the desired oxidation states were obtained in an optically transparent thin layer (5 × 10⁻³ cm) electrode cell.¹⁵

Fluorescence Spectroscopy. Modified Pt/ \sim (NH₂)₄TPP flags were first scanned with cyclic voltammetry in 0.1 M Et₄NClO₄/Me₂SO, to determine the coverage of electrochemically reactive porphyrin, Γ_{elec} , and then hydrolytically stripped in a small, measured volume of refluxing HCl (concentrated) for 1 h. Exhaustive removal of the porphyrin was confirmed by reflectance and electrochemical inspection. The hydrolysate fluorescence spectrum was taken with a Hitachi MPF-2A spectrofluorimeter ($\lambda_{\text{exc}} = 439$ nm), calibrating the intensity axis with standard (NH₂)₄TPP solutions.

Results and Discussion

Cyclic Voltammetry of Modified Electrodes. The electrochemical behaviors of aminophenylferrocene and (NH₂)₄TPP immobilized as in reaction 1 are well behaved and similar to observations by other established attachment chemistries.^{3,5,7-10} Exemplary cyclic voltammetry of the attached molecules is shown in Figure 2 and data are given in Table I, entries 1, 3–9. Surface formal potentials E°_{surf} are similar to solution analogue values¹⁶ and the small ΔE_p values reflect fast electron-transfer rates. Coverage values from *anhydrous* silanization reactions (entries 1, 4–9) are within monomolecular expectations. The carbon-immobilized (NH₂)₄TPP is easily metalated, which is useful in “tuning” the porphyrin potential as an electrocatalytic electron-transfer mediator.¹⁷ The electrochemistry of Pt/ \sim (NH₂)₄TPP

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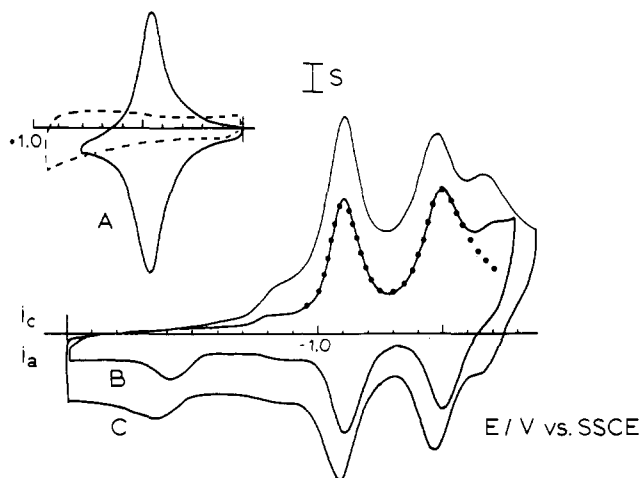


Figure 2. Cyclic voltammetry of BCl-silanized electrode. (A) $C/\infty\text{-NH}_2\text{PhCpFeCp}$ in 0.1 M $\text{Et}_4\text{NClO}_4/\text{CH}_3\text{CN}$ at 200 mV/s, $\Gamma = 4.5 \times 10^{-10}$ mol/cm², $S = 9.3 \mu\text{A}/\text{cm}^2$; dashed curve same as curve A except carbon electrode was not silanized, $S = 7 \mu\text{A}/\text{cm}^2$. (B) $\text{Pt}/\infty(\text{NH}_2)_4\text{TPP}$ in 0.1 M $\text{Et}_4\text{NClO}_4/\text{Me}_2\text{SO}$ at 100 mV/s, $\Gamma = 5.7 \times 10^{-10}$ mol/cm², $S = 14 \mu\text{A}/\text{cm}^2$, points are fit of surface activity theory¹⁰ to experiment, $r = r_0 = r_R = -4.04$ and -6.74×10^9 for the two waves. (C) $C/\infty(\text{NH}_2)_4\text{TPP}$ in 0.1 M $\text{Et}_4\text{NClO}_4/\text{Me}_2\text{SO}$ at 200 mV/s, $\Gamma = 2.5 \times 10^{-10}$ mol/cm², $S = 7 \mu\text{A}/\text{cm}^2$.

and $C/\infty(\text{NH}_2)_4\text{TPP}$ surfaces is virtually identical (Figure 2, curves B, C), except that Γ on the former are larger due to the nonanhydrous silanization procedure (Experimental Section). The broadening of surface waves of BCl-attached species can usually be accounted for by surface activity effects as illustrated by the fit¹⁰ of the theory of Brown and Anson¹⁸ to experiment in Figure 2, curve B. The derived activity parameters are the same on C and Pt, so this broadening is not specific to the electrode material.

That the BCl silane actually promotes aminophenylferrocene attachment to carbon is confirmed by omitting the silanization step, whereupon the ferrocene wave is much smaller or entirely absent¹⁹ (Figure 2A). The $C/\infty\text{-NH}_2\text{PhCpFeCp}$ electrode is very stable in the ferrocene state and has $t_{1/2} \sim 3$ h when electrochemically maintained in the ferricenium state²⁰ in CH_3CN . XPS of $C/\infty\text{-NH}_2\text{PhCpFeCp}$ surfaces shows a Si 2p peak at 102.4 eV and Fe 2p_{3/2} peak at 708.0 eV, characteristic of the silane and of ferrocene, respectively. Conversion of the 2.18 $I_{\text{Fe}}/I_{\text{Si}}$ intensity (area) ratio to an atom ratio²¹ indicates that about 50% of the attached silane undergoes amide coupling with the aminophenylferrocene. The rest of the BCl silane is hydrolyzed to carboxylic acid; XPS typically shows no residual chloride.

That BCl silane becomes attached to superficially oxidized Pt surfaces by Pt-O-Si bonds as represented in reaction 1 seems fairly well established.^{8,10,20,22} How BCl bonds stably to glassy carbon surfaces is less certain. Carbon oxide surface functionalities are important since the rf plasma preoxidation of the glassy carbon increases the intensity of the Si 2p silane band by about threefold over polished but not preoxidized carbon. We previously represented¹¹ carbon silanization as a surface silyl ether bond, but these surfaces are surprisingly stable if they are silyl ethers. Perhaps formation of two surface bonds as assumed in reaction 1 provokes this stability. Although it is possible that some siloxane polymer formation enhances stability, our experience with organosilanes suggests that polymers are unlikely under the anhydrous reaction

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(23) The slight λ_{max} difference may be due to unhydrolyzed amide bonds in the $(\text{NH}_2)_4\text{TPP}$ hydrolysate.

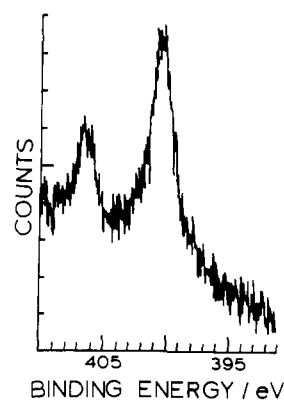


Figure 3. N 1s XPS of a BCl-silanized Pt/PtO surface after reaction with 3,5-dinitrobenzoyl chloride. Counts are expressed as 1×10^3 counts/division. Band at left is nitro (B.E. = 406.6 eV), band at right is porphyrin nitrogen plus amide nitrogen (overlap) (B.E. ~ 400 eV). Relative band areas are 1.9 after correction of 400 eV band for typical adsorbed level of reduced N 1s. No nitro correction necessary as indicated by control experiment.

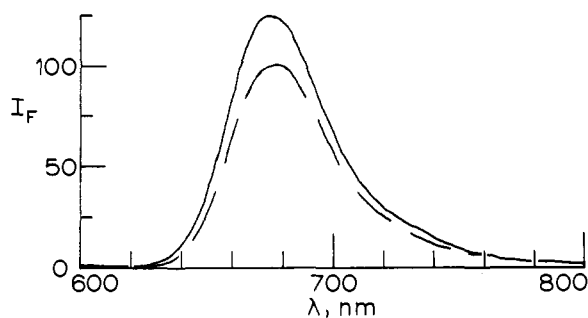


Figure 4. Fluorescence spectra ($\lambda_{\text{exc}} = 439$ nm) in HCl (concentrated) of $\text{Pt}/\infty(\text{NH}_2)_4\text{TPP}$ hydrolysate solution (—) and 2.0×10^{-7} M standard $(\text{NH}_2)_4\text{TPP}$ solution (---). I_F axis is % of intensity of the standard solution. For this example (determination 2 from Table II), $(\text{NH}_2)_4\text{TPP}$ hydrolysate concentration is 2.5×10^{-7} M.

conditions used for glassy carbon in this study. However the BCl silane bonds to glassy carbon, this acid chloride silane provides a bridge by which the same amine redox reagent can be attached to either carbon or a metal oxide in a similar chemical environment.

XPS Analysis of $(\text{NH}_2)_4\text{TPP}$ Bonding to BCl Silanized Pt. $(\text{NH}_2)_4\text{TPP}$ is a tetrafunctional amine, but all of the amine sites do not necessarily couple to BCl silane sites. This problem can be analyzed by derivatizing any dangling amine with 3,5-dinitrobenzoyl chloride followed by N 1s XPS of the nitro (406.6 eV) vs. amide plus porphyrin nitrogen bands (~ 400 eV) (Figure 3). The relative N 1s band areas indicate that the nitro tag reagent reacts on the average with 1.9 dangling amine groups per porphyrin, which leads us to surmise an average of two amide bonds between each porphyrin and BCl silane sites. We obtained⁷ the same result for $(\text{NH}_2)_4\text{TPP}$ attached to thionyl chloride activated carboxylic acid groups on glassy carbon. Given the added steric flexibility afforded by the carbon chain on the BCl silanized Pt as compared to acid chloride sites presumably affixed directly to a glassy carbon lattice, observation of the same average amide bond number to these surfaces with $(\text{NH}_2)_4\text{TPP}$ is surprising. Perhaps the formation of multiple surface bonds is more constrained by steric interactions between adjacently bonded $(\text{NH}_2)_4\text{TPP}$ molecules than by simple steric register of their amine groups with surface acid chloride sites.

Fluorescence of Hydrolytic Product of $\text{Pt}/\infty(\text{NH}_2)_4\text{TPP}$ Surface. The reaction conditions used in this paper for BCl silanization of Pt/PtO surfaces were deliberately "wet" (see Experimental Section), in contrast to the anhydrous procedures used above for glassy carbon and by us previously for metal oxides. In moist laboratory air, some polymerization of the BCl silane may occur, so that multimolecular layers of redox substances become bonded to the electrode surface. In such a multilayer film,

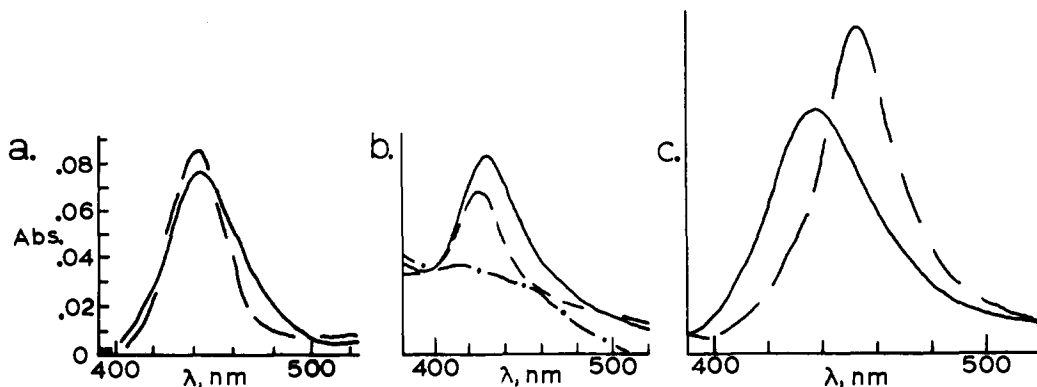


Figure 5. (A) Reflectance spectrum, $\Gamma_{\text{elec}} = 4.1 \times 10^{-10}$ mol/cm² of Pt/ \sim (NH₂)₄TPP surface in contact with 0.1 M Et₄NClO₄/Me₂SO (—); transmission spectrum of 1-cm cell containing 4.1×10^{-7} M solution of (NH₂)₄TPP in Me₂SO (---); (B) reflectance spectrum of Pt/ \sim (NH₂)₄TPP in contact with 0.1 M Et₄NClO₄/Me₂SO at 0 V vs. SSCE (—); same except -1.4 V vs. SSCE applied potential (---); transmission spectrum of Cu(NH₂)₄TPP in Me₂SO (---); (C) reflectance spectrum of Pt/ \sim (NH₂)₄TPP in contact with 0.1 M Et₄NClO₄/DMSO at 0 V vs. SSCE (—); same except after ~5 min at -1.3 V vs. SSCE (---). Absorbance axis for (B) and (C) uncalibrated.

Table II. Comparison of Electrochemical and Fluorimetric Coverages of (NH₂)₄TPP on Pt/ \sim (NH₂)₄TPP Electrodes

detmn	elec- trode area, cm ²	hydrolysate ^a concn, M	Γ_{fluor} , mol/ cm ^{2b}	Γ_{elec} , mol/ cm ^{2b}
1	0.84	1.8×10^{-7}	9.7×10^{-10}	10.5×10^{-10}
2	1.15	2.5	7.6	7.7
3	1.15	1.8	10.5	10.3

^a From hydrolysate solution of fluorescence intensity, solution volume is in 4–6-mL range. ^b Correction for Pt electrode microscopic roughness (~ 1.5)²⁰ has not been made to these values; it is probably inappropriate at multilayer coverages.

the arrangement of the polymeric structure may cause some of the immobilized redox molecules to be electrochemically silent. This is an important question with polymer-coated electrodes on which little quantitative data are available. Accordingly, we hydrolytically cleaved the porphyrin from Pt/ \sim (NH₂)₄TPP surfaces into a limited volume of acid and determined its fluorescence intensity. Figure 4 shows that the hydrolysate fluorescence spectrum is nearly identical with that of a standard (NH₂)₄TPP solution. Cyclic voltammetry of the Pt/ \sim (NH₂)₄TPP surface before hydrolysis affords an electrochemical coverage Γ_{elec} for comparison with that calculated from the fluorescence intensity data, Γ_{fluor} ; Table II shows that the two coverages agree quite well.

The geometrical area requirements for (NH₂)₄TPP estimated from models^{3b} are $\sim 1.2 \times 10^{-10}$ and 3.5×10^{-10} mol/cm² for coplanar and perpendicular orientations, respectively, of the porphyrin ring with respect to the Pt surface. On either account, the porphyrin coverages in Table II are more than monomolecular as expected for attachment to a polymeric BCl silane matrix. Between three and eight monolayer equivalents of porphyrin are attached to the BCl silane polymer and are quantitatively electroactive. Larger quantities have not been successfully attached by this particular silanization procedure. A technique for attaching thicker layers of amine redox substances to Pt, using spin-coating of BCl silane, is described.

If a Pt/ \sim (NH₂)₄TPP surface is viewed under fluorescent light, no visible emission from the surface is seen, although the amount of porphyrin suffices to give a visualized emission in the hydrolysate. We observed²⁴ (but did not report) a similar phenomenon during fluorescence studies of dansylsulfonamide attached to Pt with a different silane. We have reported²⁵ that the photostability characteristic of certain ruthenium complexes dissolved in solution vanishes upon their immobilization on Pt using silane chemistry; the surfaces are photostable. These three qualitative observations suggest that excited state lifetimes are generally shortened by the

silanization attachment to Pt. The excited-state quenching may occur via energy transfer through weak spectral overlap with the Pt electrode absorption. A far more interesting eventuality would be electron-transfer quenching, since in it an electron-transfer event akin to that operative in the electrochemical reactions of these molecules occurs at a rate competitive with excited-state emission lifetimes. We suggest that quantitation of the quenching effect might be an avenue to electrode kinetic studies.

Reflectance Spectra of Pt/ \sim (NH₂)₄TPP Electrodes. BCl silanized glassy carbon surfaces to which (NH₂)₄TPP has been attached exhibit a faint porphyrin coloration but are not very suitable for sensitive electronic spectroscopy. Reflectance spectra of Pt/ \sim (NH₂)₄TPP surfaces, which have a golden tint to the eye, are on the other hand readily obtained. A reflectance spectrum of Pt/ \sim (NH₂)₄TPP ($\Gamma_{\text{elec}} = 4.1 \times 10^{-10}$ mol/cm²) is compared in Figure 5a to a transmission spectrum of a solution containing the same number of porphyrin molecules per unit beam area. The Soret λ_{max} are virtually identical (see Table III), reflecting weak (or equal) interactions between the ground and excited porphyrin states and the Pt electrode, a result consistent with the general equality of $E^{\circ'}_{\text{surf}}$ and $E^{\circ'}_{\text{soln}}$ values (Table I). A weak porphyrin-electrode interaction does exist; the surface spectrum is broadened toward longer wavelengths. Similar results are obtained for Pt/ \sim (Cu)(NH₂)₄TPP (Figure 5b). One interpretation of this broadening effect would invoke charge-transfer interaction of the excited state with the metal conduction band.²⁶

For interactions between electrode and porphyrin to be weak, it is important that the Pt/ \sim (NH₂)₄TPP surface be properly solvated. A reflectance spectrum with the surface in contact with air is severely broadened and red shifted as compared to the Me₂SO solvated surface.

Quantitative Aspects of Pt/ \sim (NH₂)₄TPP Reflectance Spectra. The quantity of attached (NH₂)₄TPP is known unequivocally from the analysis of Table II, so the reflectance spectral intensity can be analyzed. In reflection from a surface coated with a thin absorbing film, of the available quantitative models,^{27,28} that by Hansen²⁸ is most readily applied to the present situation.

$$A = \frac{n_2 \alpha_2 d_2}{2.3 n_1 R_0 \cos \theta} \langle E^2 \rangle_0 \quad (2)$$

α_2 is the absorption coefficient of the porphyrin layer of thickness d_2 (i.e., $\ln(I_0/I) = \alpha_2 d_2 = 2.3 \times 10^3 \epsilon \Gamma$ where ϵ is molar absorptivity), and d_2 is taken as small compared to the thickness of the standing, reflecting wave²⁹ set up at the electrode-solution interface. The node of this wave is slightly below the surface plane;

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Table III. Reflectance Spectral Intensities for Pt/(NH₂)₄TPP Electrodes All in Me₂SO Solvent

solution ^a	electrode	Γ_{elec} , mol/cm ²	A at λ_{max}	λ_{max}	ϵ_{max} , L/(mol cm)	f^b
(NH ₂) ₄ TPP	Pt/(NH ₂) ₄ TPP	5.4×10^{-10}	0.100	442	3.2×10^5	3.2
	Pt/(NH ₂) ₄ TPP	4.1×10^{-10}	0.074	440	3.1×10^5	3.2
Cu(NH ₂) ₄ TPP	Pt/(Cu)(NH ₂) ₄ TPP	3.8×10^{-10}	0.074	432	3.4×10^5	2.3
				425	2.6×10^5	1.5
Co ^{III} (NH ₂) ₄ TPP ^c	Pt/(Co ^{III})(NH ₂) ₄ TPP	3.3×10^{-10}	0.070	450	3.6×10^5	2.9
				450	1.4×10^5	1.0
Co ^{II} (NH ₂) ₄ TPP ^c	Pt/(Co ^{II})(NH ₂) ₄ TPP	3.3×10^{-10}	0.056	435	2.9×10^5	3.2
				433	1.1×10^5	1.3
Co ^I (NH ₂) ₄ TPP ^c	Pt/(Co ^I)(NH ₂) ₄ TPP	3.3×10^{-10}	0.038	442	2.0×10^5	
				434	0.75×10^5	

^a Transmission spectra of standard solutions. ^b Calculated from $f = 4.32 \times 10^{-9} f \epsilon d \bar{\nu}$. ^c Due to solubility problems, Co(NH₂)₄TPP gave difficulties in preparing an adequate standard solution, so the Co(NH₂)₄TPP concentration was measured assuming that f for Co^{II}(NH₂)₄TPP is equal to f for Co^{II}TPP.

for visible light reflected normally from Pt the mean square intensity of the standing wave at the surface plane²⁷ $\langle E^2 \rangle_0 = 0.26$. Using literature parameters³⁰⁻³² and assuming that R_0 is unaffected by the porphyrin, eq 2 simplifies to

$$A = 0.58 \times 10^3 \epsilon \Gamma \quad (3)$$

Values of ϵ thus derived (Table III) are larger than those of their unattached analogues. A better comparison of attached and unattached intensities can be based on band areas expressed as oscillator strengths,³³ $f = 4.32 \times 10^{-9} \int \epsilon d\nu$, which (Table III) again indicates a higher absorption probability for the attached versions. Oscillator strengths for the unattached porphyrin Soret peak are very large, approaching the hypothetical limit of 2. The attached versions have $f_{\text{surf}} > 2$, which may indicate (i) we have seriously overestimated n_2 (unlikely), (ii) R_0 is substantially decreased by the porphyrin layers, (iii) we have underestimated reflectance intensity in the long-wavelength spectral tail, or (iv) the (NH₂)₄TPP ring has a preferred orientation with respect to the Pt surface. The last would not be surprising since the porphyrin is connected to the surface by two amine groups. The Soret transitions are polarized in the plane of the porphyrin ring,³⁴ so a larger f_{surf} implies that more porphyrins are oriented with absorption vector parallel to the electric vector of the (normal incidence) incoming light (i.e., with the porphyrin ring parallel to the Pt) than are oriented perpendicular to the surface.

Reflectance Spectroelectrochemistry. Virtually all of the spectroscopy to date on modified electrodes has been ex situ, i.e., XPS, IETS, fluorescence, etc. In a cyclic voltammogram such as in Figure 2 there is no real question that the voltammetry reflects reduction and oxidation of attached porphyrin; however, electrochemical potentials usually have limited structural information content. For more chemically involved electron-transfer reactions, supplementary in situ spectroscopy would often be of value. The reflectance spectral experiment is suited to such observations with attached, potent chromophores like the porphyrins.

The reflectance spectrum of a Pt/(NH₂)₄TPP surface in contact with Me₂SO solvent is the same whether the electrode is at open circuit or potentiostatted at 0 V vs. SSCE. Application of $E = -1.3$ V vs. SSCE (as observed by the cyclic voltammetry to be sufficient to reduce the porphyrin to its radical anion) for 5 min changes the reflectance spectrum to that in Figure 5c. The

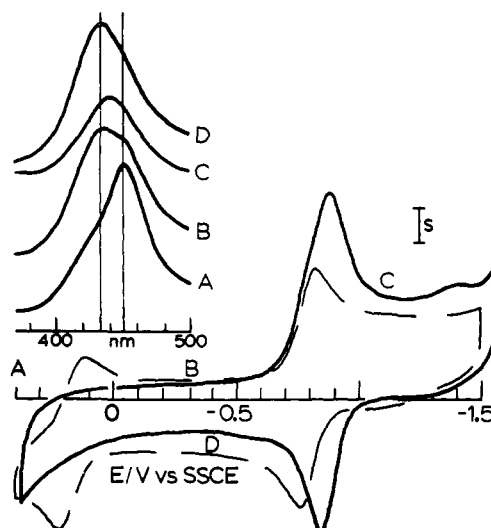


Figure 6. Reflectance spectroelectrochemistry of a Pt/(NH₂)₄TPP surface in 0.1 M Et₄NClO₄/Me₂SO. From the cyclic voltammetry (—), $\Gamma_{\text{elec}} = 4.1 \times 10^{-10}$ mol/cm² [from the Co(I → II) anodic wave], and $Q_{\text{cath}}/Q_{\text{anod}} = 1.8$. $S = 10 \mu\text{A}/\text{cm}^2$. The dashed curve (---) is a Co(TPP)-Me₂SO solution at a glassy carbon electrode, 100 mV/s. The reflectance spectra were taken at the potentials (A–D) indicated on the voltammogram of Pt/(NH₂)₄TPP, in that order. The line at 433 nm is λ_{max} for Co(II)(NH₂)₄TPP in Me₂SO solution, and the 450-nm line is λ_{max} for Co(III)(NH₂)₄TPP in Me₂SO solution, both from thin-layer spectroelectrochemistry.

peak at 453 nm is probably *not* the simple radical anion, but is instead a protonated form. Reduction of Pt/(Cu)(NH₂)₄TPP to its radical anion (Figure 5b) is not accompanied by protonation. In the radical anion, the Soret peak collapses and appears split.

The cobalt-metalated surfaces provide better illustrations of the power of in situ spectroelectrochemistry. Pt/(NH₂)₄TPP and C/(NH₂)₄TPP surfaces exhibit very unusual electrochemical behavior. Figure 6 compares typical cyclic voltammograms for Pt/(NH₂)₄TPP and for a CoTPP solution. The quasi-reversible dissolved Co(TPP) wave for Co(III/II) at $E^{\circ'}_{\text{soln}} = +0.1$ V vs. SSCE is vanishingly small on the Pt/(NH₂)₄TPP electrode; it can be observed by differential pulse voltammetry at the reversible potential (+0.12 V), but is only 1–5% as large as the Co(II/I) surface wave at -0.86 V vs. SSCE. The second unusual feature of the Pt/(NH₂)₄TPP voltammogram is that the charge underneath the cathodic surface wave at the Co(II/I) potential is *larger* than that of the anodic member. The exact value of $Q_{\text{cath}}/Q_{\text{anod}}$ depends on the electrode's potential scanning history; in Figure 6 it is 1.8. If only the potential region immediately around the Co(II/I) wave is scanned, $Q_{\text{cath}}/Q_{\text{anod}} = 1.0$.

We have observed⁷ qualitatively similar behavior for cobalt-metalated (NH₂)₄TPP attached to activated glassy carbon electrodes, and Jester has hypothesized³⁵ that the cobalt becomes

(30) Refractive indices $n_1 = 1.48$ for the incident medium Me₂SO and $n_2 \approx 2.0$ for the silane-porphyrin layer (estimated from ellipsometric data³¹), substrate reflectivity $R_0 = 0.6$ for Pt in the 400–500-nm range³², and angle of (normal) incidence $\cos \theta = 1.0$.

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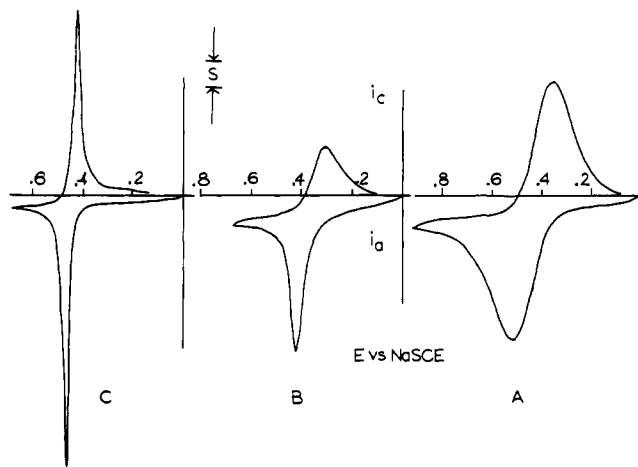


Figure 7. Cyclic voltammograms of aminophenylferrocene coupled to spin-coated, polymerized BCl silane film on Pt. (A) 0.1 M $\text{Et}_4\text{NClO}_4/\text{CH}_3\text{CN}$, 2 mV/s, $S = 6.5 \mu\text{A}/\text{cm}^2$; (B) 0.1 M $\text{LiClO}_4/\text{H}_2\text{O}$, 2 mV/s, $S = 7.0 \mu\text{A}/\text{cm}^2$; (C) 0.1 M $\text{H}_2\text{SO}_4/\text{H}_2\text{O}$, 1 mV/s, $S = 7.0 \mu\text{A}/\text{cm}^2$. Apparent coverage (measured by base line current extrapolation) of ferrocene is $\Gamma = 1.5 \times 10^{-7}$, 1.5×10^{-8} , and $3.0 \times 10^{-8} \text{ mol}/\text{cm}^2$, respectively. E_{fwhm} (anodic) = 200, 55, and 28 mV for curves A–C, respectively.

axially coordinated by carbon surface ligands, probably carboxylate, which centrally underlie the cobalt porphyrin site.

Reflectance spectroelectrochemistry of $\text{Pt}/\text{Co}(\text{NH}_2)_4\text{TPP}$ was useful in exploring this proposal. In this case Figure 6 shows that after potentiostating at +0.4 V vs. SSCE for a few minutes, the 452-nm λ_{max} reflectance spectrum agrees well with that (450 nm) for a solution of $\text{Co}(\text{III})\text{TPP}$ obtained in an optically transparent thin-layer cell at the same potential. At -0.3 V vs. SSCE the absorbance decreases at 452 nm and grows at 435 nm which is near λ_{max} for a solution of $\text{Co}(\text{II})\text{TPP}$ (433 nm). The change, however, is exceedingly slow; the spectrum shown (curve B) was taken *after several minutes of potentiostating* and a significant $\text{Co}(\text{III})$ porphyrin shoulder is still apparent. At a potential of -1.1 V vs. SSCE, the surface spectrum changes immediately to curve C, the $\text{Pt}/\text{Co}(\text{I})(\text{NH}_2)_4\text{TPP}$ state. Returning to -0.4 V vs. SSCE, an immediate restoration of the $\text{Co}(\text{II})$ spectrum (curve D) is obtained, now without any shoulder for $\text{Co}(\text{III})$ porphyrin. Finally, application of +0.4 V vs. SSCE restores the $\text{Co}(\text{III})$ spectrum as in curve A. This last reaction occurs slowly, but not as slow as the $\text{Co}(\text{III}) \rightarrow \text{Co}(\text{II})$ transformation.

The reflectance spectroelectrochemistry shows that the $\text{Co}(\text{III}/\text{II})$ reaction does occur on the $\text{Pt}/\text{Co}(\text{NH}_2)_4\text{TPP}$ surface, at a potential between +0.4 and -0.3 V vs. SSCE, but it is much too slow to contribute a noticeable current peak in a cyclic voltammogram. The reaction occurs rapidly at potentials sufficient to reduce the porphyrin to the $\text{Co}(\text{I})$ state; in Figure 6, Q_{cath} corresponds to a $\text{Co}(\text{III}/\text{I})$ reaction.

Our rationalization of the slow $\text{Co}(\text{III}/\text{II})$ reaction rests on the expectation³⁶ that cobalt is six-coordinate in the $\text{Pt}/\text{Co}(\text{III})-(\text{NH}_2)_4\text{TPP}$ state but five-coordinate as $\text{Pt}/\text{Co}(\text{II})(\text{NH}_2)_4\text{TPP}$, so dissociation of an axial ligand will accompany electron transfer. Such a change in coordination by pyridine was established on carbon by Jester.³⁵ If the axial ligand is carboxylate, coimmobilized on the Pt surface with the porphyrin, such axial dissociation would be very slow. Reassociation would be comparatively faster, and we spectrally observe that the $\text{Co}(\text{II}/\text{III})$ transformation is faster than $\text{Co}(\text{III}/\text{II})$.

Spin Coating of BCl Silane on Pt. Electrodes coated with polymeric redox substances^{21,37-43} offer possibilities of many in-

teresting applications, including electron-transfer mediation at semiconductor photoanodes, for which polymerized silylferrocenes^{44,45} have been used. To expand the scope of polyferrocene-coated electrodes, we have spin-coated films of BCl silane onto oxidized Pt surfaces (see Experimental Section). The relative reactivities of the chlorosilyl and acid chloride groups are such that PtOSi surface bonding and siloxane polymerization can occur without exhaustive hydrolysis of the acid chloride groups so that reagents such as aminophenylferrocene can amide bond to the preformed siloxane polymer matrix. The shiny brownish surface thus prepared shows an intense Fe $2p_{3/2}$ XPS peak at 708 eV (characteristic of ferrocene) and a Pt 4f band which is very weak or absent depending upon the thickness of the polymeric film.

Cyclic voltammetry of the film shows a charge for the oxidation of aminophenylferrocene which varies with the amount of spin-coated BCl silane from 10^{-9} to as large as $2 \times 10^{-7} \text{ mol}/\text{cm}^2$. In acetonitrile, symmetric voltammetric wave shapes observed at lower coverages grow tails (Figure 7A) at high coverages. In Figure 7A, electrochemical charge is being transported through a film equivalent to at least ~ 500 monolayers of aminophenylferrocene. Exchange of electrons between adjacent sites is the most likely mechanism for this, but the exchange itself is not necessarily rate controlling.^{39c}

In aqueous LiClO_4 (and HClO_4), the anodic current peak becomes quite sharp (Figure 7B), and in aqueous H_2SO_4 both anodic and cathodic peaks are sharp (Figure 7C). $E_{fwhm} = 28$ mV for Figure 7C whereas surface waves for redox polymers typically have $E_{fwhm} = 100\text{--}200$ mV. The sharp current peaks mean that the ratio of activities of the oxidized and reduced surface molecules changes relatively little over the course of exhaustive electrolysis of the surface layer. This is a *phaselike* property, as we have pointed out in connection with the behavior of plasma polymerized vinylferrocene films in aqueous LiClO_4 .⁴² The present results demonstrate that phaselike behavior can occur in quite diverse types of ferrocene polymer matrices and is moderated both by choice of solvent and electrolyte. Presumably, the important factor in this activity phenomenon is whether the polymer matrix becomes significantly swollen by solvent in its various oxidation state/electrolyte counterion forms. Finally, we note that the proper choice of solvent and electrolyte allows switching between ferrocene and ferricenium surface states over a rather narrow range of potentials. Considering the redox film as a molecular surface state (layer), a phaselike property yields a very narrow energy band, a factor of possible utility in applications to semiconductor electrodes.

Further details and applications of spin-coated BCl and other siloxane polymer films will be presented in another report.

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