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Kinetics of Simple Inner- and Outer-Sphere
Electrochemical Reactions at Rotating Silver
Electrodes as Examined using Surface-Enhanced
Raman Spectroscopy

by

S. Farquharson, D. Milner, M. A. Tadayyoni

and M. J. Weaver

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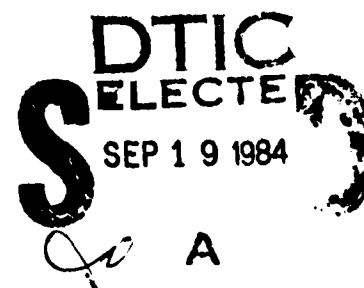
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An application of surface-enhanced Raman spectroscopy (SERS) to evaluating the kinetics of electrochemical reactions is outlined. This involves monitoring the potential-dependent SERS intensities of the adsorbed reactant at a rotating disk electrode in a potential region where the kinetics are under mixed mass transfer-heterogeneous electron transfer control. Comparisons are made between the kinetic behavior extracted from the SERS intensity-potential dependence and that obtained from the faradaic current flow. For $\text{Co}(\text{NH}_3)_6^{3+}$ reduction at silver, where the reactant is electrostatically attracted but not bound to the surface, the SERS and electrochemical rate behavior is closely compatible. However, for $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ and $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ reduction, where the reactants bind directly to the surface prior to electron transfer, the reactivities at the SERS-active sites are substantially higher than for the prevalent adsorbate as obtained electrochemically. These findings suggest that surface sites displaying efficient Raman scattering may also provide centers of catalytic activity.

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It is now well established that a variety of species when adsorbed on certain metal surfaces, most prominently silver, gold, and copper, can yield easily detectable Raman spectra as a result of surface-enhanced Raman scattering (SERS).¹ This *in situ* spectroscopic technique is especially applicable to electrochemical systems; Raman signals arising entirely from interfacial species can be obtained using conventional cell configurations even in the presence of substantial bulk concentrations (≤ 0.1 M) of the Raman scatterer. Although considerable application to the molecular characterization of metal-solution interfaces is envisaged, such developments are still in their infancy.

A key issue to be addressed in the application of SERS to surface electrochemistry is the correspondence between the properties of the adsorbate molecules that contribute predominantly to the spectral signals and those which are primarily responsible for the observed electrochemical behavior, such as faradaic current flow. This issue is somewhat clouded by the likelihood that the sites responsible for SERS constitute only a small minority of the overall surface. These sites may be associated with metal clusters or other surface morphologies.² Thus the viability of SERS as a surface electrochemical technique hinges on the correlation, if any, between the chemical properties of the SERS-active species and those of the overall adsorbate. However, surprisingly few such comparisons have been reported.³⁻⁷ The present communication describes some preliminary evaluations of the electroreduction kinetics of adsorbed species using SERS, and the comparison with the kinetics as determined from faradaic current flow. These experiments appear to be the first application of SERS to electron-transfer dynamics at metal surfaces.

We have recently been examining surface-enhanced Raman (SER) spectra for a variety of simple inorganic adsorbates at the silver-aqueous interface under conditions where the surface composition can be characterized by independent means.³⁻⁸ These include substitutionally inert transition-metal complexes, especially Co(III), Cr(III), Ru(III), and Os(III) ~~ammines~~ amines, whose surface concentrations can be obtained in favorable cases from the charge required to reduce the metal cation as measured using electrochemical perturbation techniques.^{9,10} Suitably intense SER spectra can be obtained for a number of such couples that are bound to the surface via bridging ligands, notably bromide, thiocyanate, or pyrazine. We have also observed that SER spectra can be obtained at silver for *unbound* molecules, most notably metal hexaamine cations, when they are electrostatically attracted into the diffuse layer⁷ (*vide infra*).

The occurrence of electron-transfer reactions involving such interfacial species can readily be detected using SERS since several features of the Raman spectra, including vibrational frequencies, are sensitive to the metal oxidation state. Redox thermodynamic data have been obtained from potential-dependent SER spectra for several adsorbed Ru(III)/(II) and Os(III)/(II) couples.^{6,7} These data are in good agreement (surface formal potentials within ca 20 mV) with the corresponding information obtained from conventional electrochemistry.^{6,7}

These recent findings regarding redox equilibria have led us to consider ways in which SERS can be utilized to probe the redox *reactivity* of molecules at metal surfaces. Two general approaches can be envisaged. The first involves monitoring the time dependence of the adsorbed reactant (or product) concentration in response to a rapid electrochemical perturbation. Such experiments are feasible using an array detector system as employed in our laboratory (PAR OMA-2).

However, the experiments reported here employ a simpler approach, involving monitoring the SER spectra at a rotating disk electrode as a function of potential. Cobalt(III) and chromium(III) reactants were chosen that exhibit totally irreversible electroreductions at silver electrodes under these conditions. In other words, the rotating disk voltammograms for these systems are controlled by the heterogeneous electron-transfer kinetics along with mass transport of the reactant to the electrode surface. This leads to the following analysis.

The reactant concentration close to the electrode surface, C_s , (i.e. the three-dimensional concentration just outside the double layer) will progressively decrease relative to the bulk concentration, C_b , on the rising part of the voltammetric wave according to^{11,12}

$$C_s = C_b [(i_l - i)/i_l] \quad (1)$$

where i is the current density at a given potential, and i_l is the mass transport-limited current density. The integrated intensity of a given Raman band for an adsorbed species, I_{SER} , is expected to be roughly proportional to its (two-dimensional) surface concentration, Γ_s , at the SERS-active surface sites. As the reaction rate at these sites increases, the steady-state value of Γ_s and hence I_{SER} will decrease. Provided that the reactivity of the SERS-active adsorbate does not differ from that associated with the remaining surface sites, these decreases in Γ_s will occur uniformly across the surface. If Γ_s is in turn proportional to C_s (i.e. Henry's Law applies) then

$$I_{SER} = I_{SER}^0 [(i_l - i)/i_l] \quad (2)$$

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The quantity I_{SER}^c is the value of I_{SER} that would be observed in the absence of an electrode reaction, (i.e. when $i = 0$). This should be approximated by the value of I_{SER} at the foot of the voltammetric wave, where $C_s \approx C_b$. Consequently, the comparison between the potential dependence of I_{SER} and the corresponding current-potential curves in terms of Eq. (2) should provide useful insight into the relative redox reactivities of molecules at SERS-active and -inactive sites.

Experimental

The rotating silver electrode had a disk radius of 0.2 cm, surrounded by a teflon sheath of radius 0.6 cm. The electrode was mechanically polished on a wheel successively with 5, 1, and 0.3 μm alumina immediately prior to use. A compact electrode rotator of in-house design was utilized, featuring a photodiode speed controller. The cell was of conventional design, with a window at the base through which the krypton laser beam (647.1 nm, 100-300 mW power) focussed on the electrode surface. The cell was tilted so that the scattered light could be collected at 90° to the incident beam and about 60° to the electrode surface using a SPEX Triplemate/PAR OMA-2 photodiode array detector system. This enabled sequential Raman spectra to be recorded rapidly, each being obtained by signal accumulation for 1-5 seconds so to yield suitably large signal-to-noise ratios ($> 5-10$) at the peak frequencies for the relevant SER bands. Simultaneous SERS- and rotating disk current-potential data could easily be obtained throughout the voltammetric wave within a time-scale (30-60 seconds) where the faradaic currents were reproducible and essentially no laser photolysis occurred.

Further electrochemical and chemical details for the systems studied here are given elsewhere.⁸⁻¹³ Representative spectra are given in refs. 7 and 8. All electrode potentials are quoted versus the saturated calomel electrode (s.c.e.).

Results and Discussion

Three model reactions were examined in this study. The first involves the one-electron reduction of hexaamminecobalt(III) at silver. This reaction is totally irreversible since the Co(II) product rapidly aquates to yield $\text{Co}(\text{OH}_2)_6^{2+}$ which cannot reoxidize except at extreme positive potentials.¹³ The electroreduction mechanism is expected to be outer-sphere since $\text{Co}(\text{NH}_3)_6^{3+}$ is substitutionally inert and lacks ligands capable of binding to the electrode surface. Indeed, after double-layer corrections its electroreduction kinetics are essentially independent of the electrode material.^{11,12} In perchlorate or hexafluorophosphate electrolytes at silver, well-defined rotating disk voltammograms are obtained with half-wave potentials, $E_{1/2}$, around -400 to -450 mV. Not surprisingly, no SER signals for $\text{Co}(\text{NH}_3)_6^{3+}$ were detected using these electrolytes.

However, the addition of even small concentrations of chloride to acidified 0.1 M NaClO_4 produced sizable positive shifts in $E_{1/2}$ for $\text{Co}(\text{NH}_3)_6^{3+}$ reduction. No further changes were seen for chloride concentrations above ca 0.5 mM; the $E_{1/2}$ shift (ca 300 mV) corresponding to rate increases at a given potential of around 10^4 .¹⁴ This catalytic influence of chloride can be accounted for quantitatively by the enhancement of the $\text{Co}(\text{NH}_3)_6^{3+}$ concentration at the outer Helmholtz plane (o.H.p.) caused by chloride specific adsorption.¹⁵ Most significantly for the present purpose, easily detectable SER spectra are obtained for $\text{Co}(\text{NH}_3)_6^{3+}$ under these conditions.⁷ Spectral bands observed include the symmetric N-H stretch at 3200 cm^{-1} , and the A_{1g} and E_g Co-N stretching modes at 515 and 450 cm^{-1} .⁷ (Some mild roughening of the silver via an oxidation-reduction cycle in chloride media⁵ is necessary to optimize the Raman

signal intensity. This has little effect on the electrochemical, including kinetic, properties of the surface beyond that resulting from a ca 1.5-2 fold increase in the microscopic electrode area.⁴⁾

Figure 1 contains plots (dashed curves) of the relative integrated intensity of the A_{1g} mode, I_{SER} , against the electrode potential for a pair of rotation speeds, 800 and 1600 r.p.m. The solution contained 1 mM $Co(NH_3)_6^{3+}$ in 0.1 M $NaClO_4$ + 0.01 M $NaCl$, with 2 mM $HClO_4$ to avoid hydrolysis of the Co(II) product. The values of I_{SER} at the least negative potentials (+50 to -100 mV) are normalized to the values of i_l at these two rotation speeds; the corresponding i -E curves are also shown (solid curves) in Fig. 1. (In actuality, the values of I_{SER} in the region +50 to -100 mV are independent of rotation speed.) Comparison of the corresponding I_{SER} -E and i -E curves reveals that somewhat different behavior is encountered than that predicted by Eq. (2) in that the decreases in I_{SER} only commence at potentials close to the top of the voltammetric wave, where $i \geq 0.9 i_l$ (Fig. 1). For example, according to Eq. (2), I_{SER} should equal $0.5 I_{SER}^\circ$ at $i = 0.5 i_l$ (i.e. at $E_{1/2}$), whereas in actuality $I_{SER} \approx I_{SER}^\circ$ at that point.

This apparent discrepancy can nonetheless be reconciled by noting that at potentials where $I_{SER} \approx I_{SER}^\circ$ (≥ -110 mV), the current is virtually independent of rotation speed, even though a sizable variation in C_s is expected on the basis of Eq. (1). Since it is expected that the current will be proportional to Γ_s at a given electrode potential, this result suggests that Γ_s is virtually independent of, rather than proportional to, C_s as is assumed in the derivation of Eq. (2). (In other words, the apparent reaction order, x , approaches zero, rather than equalling unity as expected under Henry's Law conditions.¹² Values

of x in the region ca 0.1-0.3 are obtained from an analysis of the dependence of the voltammograms upon rotation speed from 400-2,500 r.p.m.) This is consistent with values of Γ_s for $\text{Co}(\text{NH}_3)_6^{3+}$ that are sufficiently large so as to approach saturation ($\Gamma_s \sim 10^{-10} \text{ mol cm}^{-2}$), thereby accounting both for the large catalyses seen upon adding chloride and the appearance of the SERS signals. (Note that the reaction rate, and hence i , will be proportional to Γ_s .) Indeed, values of Γ_s approaching saturation (ca $2 \times 10^{-11} \text{ mol cm}^{-2}$) are determined directly from rapid scan cyclic voltammetry for $\text{Co}(\text{NH}_3)_6^{3+}$ and $\text{Ru}(\text{NH}_3)_6^{3+}$ at silver in chloride-containing media, even for very small values of C_s (0.05 mM).⁷ Consequently, Γ_s should remain virtually constant at values approaching saturation until $C_s \ll C_b$, corresponding to $i \geq 0.9 i_l$ [Eq. (1)]. The observation that I_{SER} only decreases around this point close to the top of the voltammetric wave is therefore wholly consistent with the electrochemical behavior. This therefore suggests that the reaction energetics for $\text{Co}(\text{NH}_3)_6^{3+}$ at the SERS-active sites do not differ noticeably from those at the sites predominantly responsible for the overall reaction rate.

The second reaction studied, $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ reduction, was chosen since, unlike $\text{Co}(\text{NH}_3)_6^{3+}$ reduction, it proceeds at silver via a well-defined inner-sphere pathway where the bromide ligand binds the reactant directly to the metal surface.⁹ Diagnostic criteria supporting this mechanism include the rate *decreases* observed for $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ reduction at silver upon the addition of chloride anions.⁹ This contrasts the large rate *increases* noted above for $\text{Co}(\text{NH}_3)_6^{3+}$ reduction and other outer-sphere reactions.^{9,12} A metal-amine stretching mode, ν_{MN} , is observed (450 cm^{-1}) in the SER spectra for $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ in perchlorate media, along with a surface-bromide stretching mode at 160 cm^{-1} , thereby confirming the mode of attachment to the silver surface.⁷ Similar to

$\text{Co}(\text{NH}_3)_6^{3+}$, the reduction of $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ is irreversible due to the lability of $\text{Cr}(\text{II})$, leading to a loss of the ν_{MN} band. Reoxidation of $\text{Cr}(\text{II})$ only occurs at potentials well positive of those for $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ reduction, and therefore does not influence the cathodic voltammogram.

Figure 2 shows a typical plot of I_{SER} of the ν_{MN} band for adsorbed $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ against electrode potential (dashed curve) at a rotation speed of 800 r.p.m., along with the simultaneously recorded voltammogram (solid curve). As in Fig. 1, the I_{SER} -E and i-E curves are normalized to one another. Both the SER spectra and the reduction currents on the rising part of the wave were somewhat time dependent; smaller values of i and I_{SER} were obtained when returning from more negative potentials. The data in Fig. 2 are an average of several runs. Nevertheless, the results stand in sharp contrast to those in Fig. 1 in that I_{SER} drops sharply to zero at potentials close to the foot of the wave, where $i/i_1 \leq 0.3$.

These results are clearly at variance with the predictions of Eq. (2), in the opposite direction to those for $\text{Co}(\text{NH}_3)_6^{3+}$ reduction. In contrast to the latter, the former cannot be rationalized on the basis of fractional electrochemical reaction orders since this only increases the discrepancies between the SERS and electrochemical data. Indeed apparent reaction orders somewhat below unity (ca 0.5-1) were obtained for $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ reduction. The I_{SER} -E data for $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ reduction therefore suggest that the adsorbate present at the SERS-active sites is reduced at noticeably smaller overpotentials, i.e. has a greater reactivity, than the prevalent adsorbate which is responsible for the i-E curves.

In order to examine this matter further, we undertook a similar comparison for $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ reduction. Like $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ reduction, this reaction proceeds via an irreversible inner-sphere mechanism since the remote thiocyanate sulfur binds strongly to the silver surface.⁹ The SERS ν_{CN} mode for $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ (450 cm^{-1}) is rather weak, making it unsuitable for monitoring the adsorbate concentration. Nevertheless, the C-N stretching mode, ν_{CN} , is suitably intense for this purpose. Although the thiocyanate ligand released upon formation of Cr(II) is also adsorbed and exhibits a SERS ν_{CN} band, its intensity is ca 5 to 10-fold weaker, and occurs at frequencies about 20 cm^{-1} lower than ν_{CN} for adsorbed $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ at the potentials (ca -600 mV) where $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ reduction commences.⁸ Consequently, the reduction of adsorbed $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ can be followed either from the intensity decreases, or from the decreases in the peak frequency of the ν_{CN} band.

Figure 3 contains a plot of the integrated intensity of the ν_{CN} band, I_{SER} , against electrode potential (dashed line) along with the corresponding voltammogram at 800 r.p.m. (solid curve) for $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ reduction (c.f. Figs. 1, 2). Similar to $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ reduction, the SER signal for adsorbed Cr(III) is almost entirely lost at the foot of the voltammetric wave, i.e. for $i/i_1 \leq 0.2$. It might be argued that this decrease in I_{SER} results from a potential dependence of the surface Raman scattering efficiency or from reactant desorption rather than from reduction of $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$. Indeed I_{SER} is somewhat potential-dependent in the region well positive of the voltammetric wave (Fig. 3) as in I_{SER} for $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ (Fig. 2). However, strong evidence against these possibilities is provided by the corresponding plot of the peak frequency, ν_{CN} , against electrode potential (the dotted-dashed curve in Fig. 3). Thus the sharp decrease in I_{SER} over the potential region -500 to -600 mV coincides with a marked change

in ν_{CN} from 2125 to 2090 cm^{-1} , consistent with complete reduction of adsorbed $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ to form NCS^- .⁸ This result is especially striking since apparent reaction orders approaching zero (≤ 0.3) are obtained from the rotation speed-dependent voltammograms for $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ reduction at silver.⁹ As for $\text{Co}(\text{NH}_3)_6^{3+}$ above, such fractional reaction orders are consistent with adsorbed reactant coverages that approach a monolayer, as is indeed found ($\Gamma_s \sim 2 \times 10^{-10} \text{ mol cm}^{-2}$) from rapid cyclic voltammetry.^{9,10} On the basis of the arguments presented above, the I_{SER} -E and i-E curves for $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ reduction also differ very markedly with the prediction of Eq. (2).

Rotating-disk voltammograms were recorded both in the absence and presence of laser illumination. Although virtually identical voltammograms were obtained, small additional cathodic currents ($\sim 20 \text{ } \mu\text{A cm}^{-2}$) were detected during laser irradiation. These photocurrents are only mildly dependent on the electrode potential, and were observed even at potentials well positive of the rising part of the voltammograms, i.e. where potential-independent SERS intensities are obtained and the thermal electroreduction currents are undetectably small. The photocurrents are only observed in the presence of high reactant surface concentrations. The possibility that the effect is due to enhancement of the thermal electrochemical kinetics by surface laser heating was considered. However, this was eliminated both on the basis of the above evidence and model calculations (as described in ref. 16) which indicate that the laser-induced temperature rise at the silver surface was negligible ($\leq 1^\circ\text{C}$) under the conditions employed. Further study of the effect has led us to identify it tentatively as photoemission involving photoreduction of the adsorbed reactant. Details will be given elsewhere.¹⁷ For the present purposes, it is important to note that the photocurrents do not appear to be connected with either the electroreduction pathways

as monitored by the rotating-disk voltammetry, or with the potential dependence of the SERS intensities.

The present results therefore indicate that both $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ and $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ adsorbed at SERS-active sites undergo noticeably more facile electroreduction than the majority of the adsorbate. This finding contrasts that obtained for the outer-sphere reduction of $\text{Co}(\text{NH}_3)_6^{3+}$ where the SERS-active species display reactivities more representative of the overall surface. Therefore such reactants bound directly to SERS-active sites are perturbed energetically in a manner which is not observed for unbound reactant. This is consistent with the observation that linear sweep voltammograms for reduction of adsorbed $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ and related complexes at silver yield unexpectedly broad reduction peaks.^{9,10,16} A typical such voltammogram is shown in Fig. 4 (solid curve); under the conditions chosen (sweep rate = 50 v sec^{-1} , reactant concentration = $50 \text{ }\mu\text{M}$) the current for $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ reduction arising from diffusing reactant can be neglected. Very similar voltammograms were obtained for smooth electropolished and electrochemically roughened (i.e. SERS-active) electrodes, save from the ca 1.5 to 2 fold higher currents for the latter associated with the larger effective electrode area. Although the cathodic reduction peak is somewhat ill-defined, it is substantially broader than that expected (half-peak width ca 125 mV .¹⁹) for a surface where all the adsorption sites are energetically equivalent.

This result, which is typically obtained for adsorbed $\text{Cr}(\text{III})$ and $\text{Co}(\text{III})$ reactants at polycrystalline surfaces^{9,10} but not at mercury,^{9,18} suggests that a polycrystalline metal provides a number of surface binding sites with significantly different electrocatalytic properties. Presumably the currents at the foot of the steady-state voltammetric waves in Figs. 2 and 3 are associated with reduction of adsorbate at the most catalytic sites

which also contribute predominantly to the SERS signals. In this regard it is interesting to note that the cyclic voltammetric reduction wave for adsorbed $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ (Fig. 4) also commences beyond -600 mV, although comparison with the steady-state voltammetric currents in Fig. 3 is difficult given the different solution conditions. The occurrence of such "active sites" for the present ligand-bridged reactions requires that the unimolecular rate constants for the adsorbed reactants are sensitive to the substrate structure. Such a dependence is indeed indicated from experimental data.⁹ Although it is difficult to estimate the fraction of the overall adsorbed reactant bound to such SERS-active sites, on the basis of the present evidence it is liable to be small, probably less than 5%.

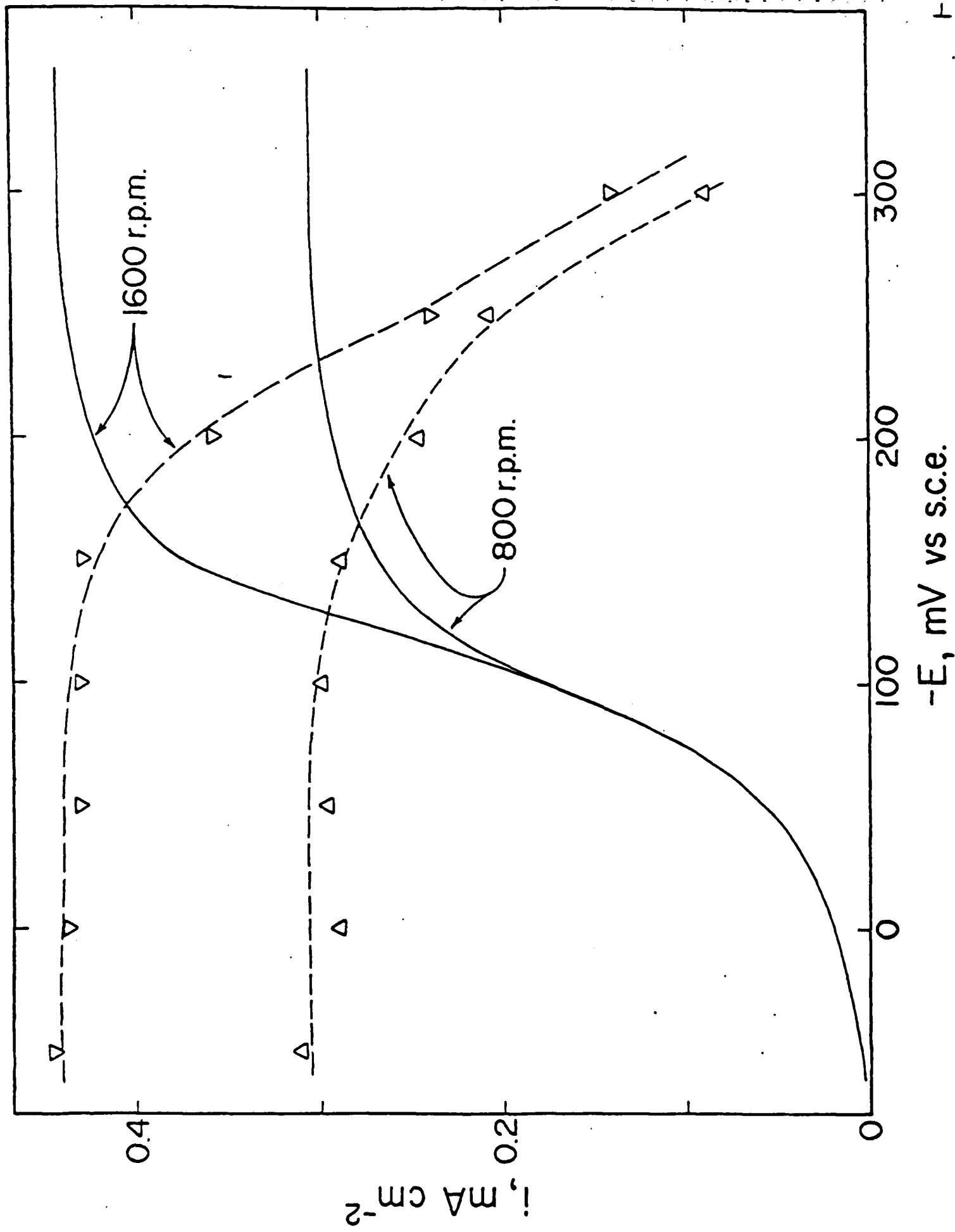
The observed electrocatalytic potency of the SERS-active sites is not entirely unexpected given the likelihood that particular surface morphologies, such as surface metal defects or clusters, contribute importantly to SERS.^{1,2} Such sites should bind strongly to incoming reactant, and are anticipated to yield especially large unimolecular rate constants and therefore enhanced overall reactivities for inner-sphere electron transfer,⁹ as observed. Whether or not these results portend well for the applicability of SERS to surface reaction dynamics is a somewhat subjective issue at this point. Regardless, the utilization of rapid photodiode array detectors to gather time-resolved SER spectra in conjunction with transient kinetic experiments should yield fresh perspectives into the molecular basis of heterogeneous catalysis.

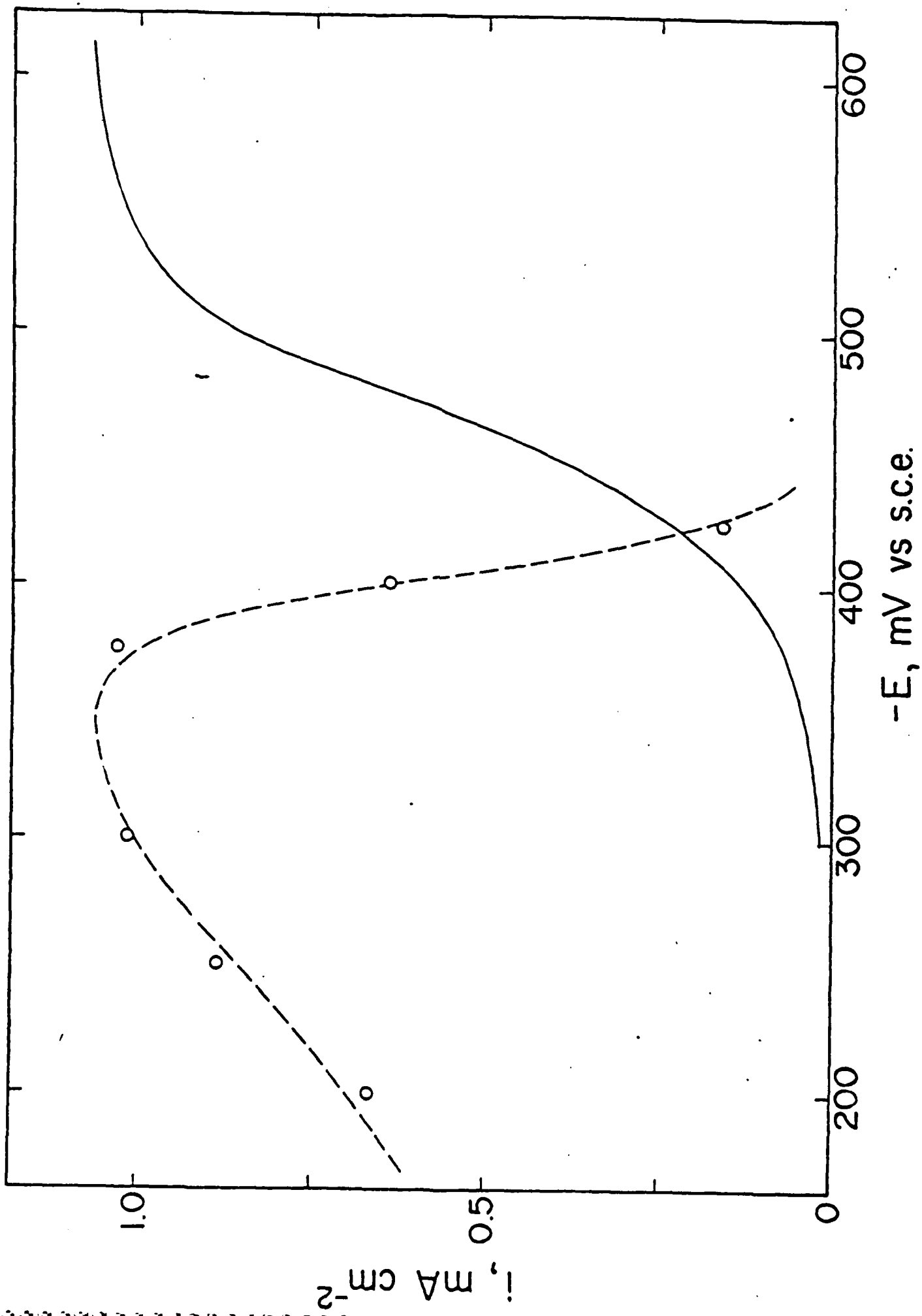
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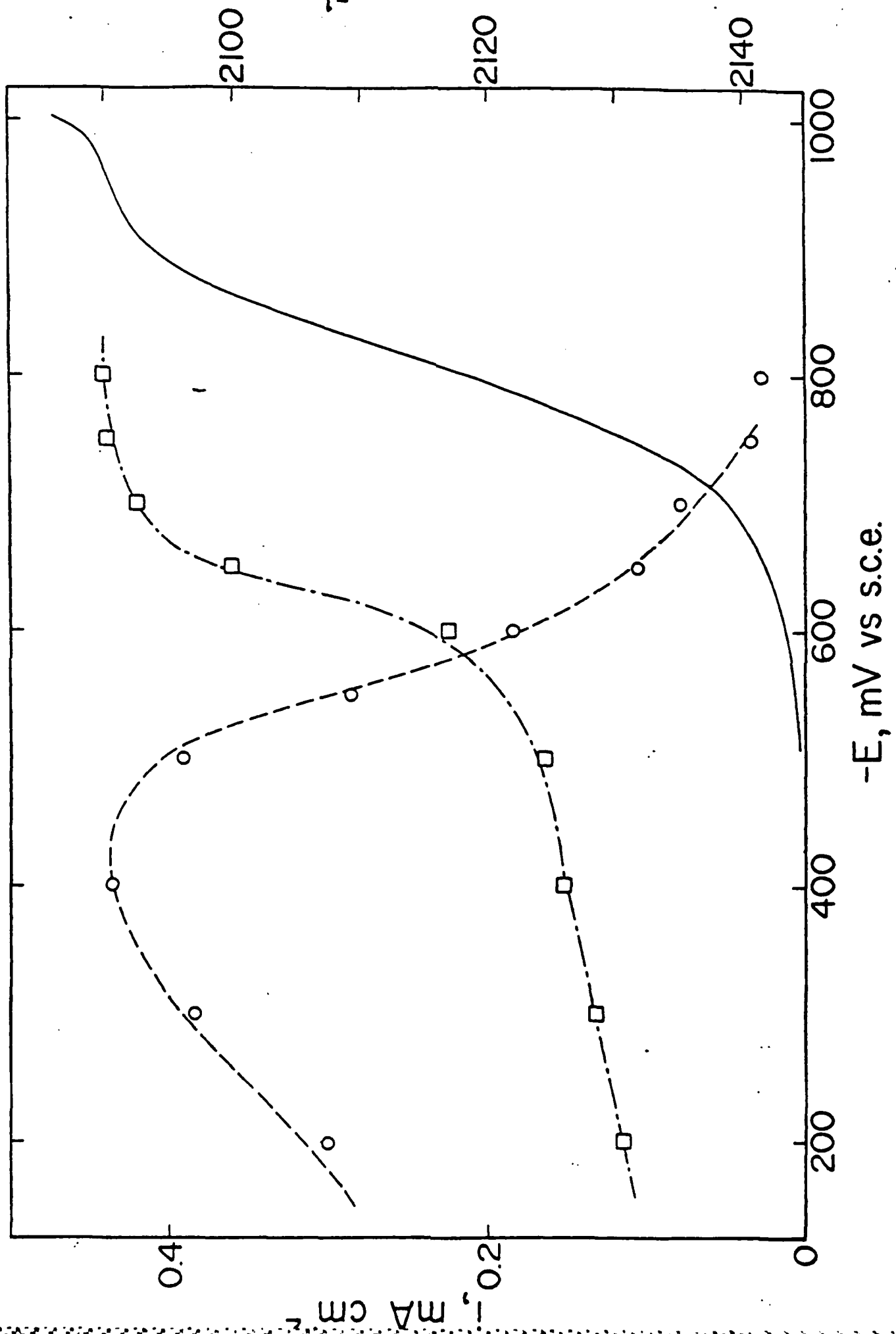
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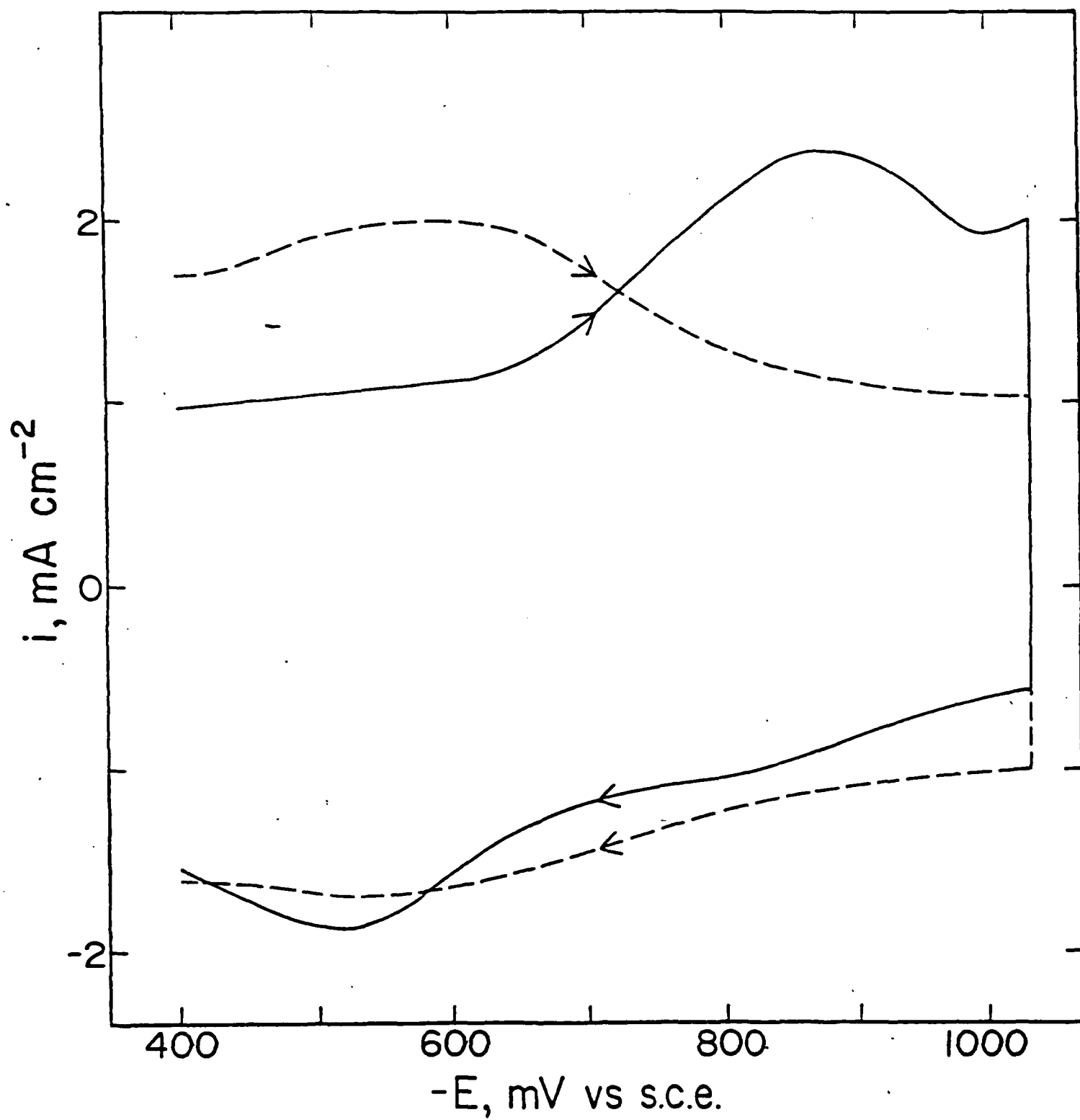


Figure Captions

Figure 1 Plots of current density, i , for $\text{Co}(\text{NH}_3)_6^{3+}$ reduction at a rotating silver electrode against electrode potential, E , (solid curves) compared with corresponding plots of integrated intensity, I_{SER} , of SERS $\text{Co}^{\text{III}}\text{-NH}_3$ stretching mode (515 cm^{-1}) against E (dashed curves). Solution contained $1\text{ mM Co}(\text{NH}_3)_6^{3+}$ in $0.1\text{ M NaClO}_4 + 0.01\text{ M NaCl} + 2\text{ mM HClO}_4$. Lower curves: 800 r.p.m. ; Upper curves: 1600 r.p.m. rotation speed. Maximum values of I_{SER} (arbitrary scale) set equal to maximum values of i at the same rotation speed. Uncertainties in I_{SER} values typically less than 5-10%.

Figure 2 Plot of current density, i , for $\text{Cr}(\text{NH}_3)_5\text{Br}^{2+}$ reduction against electrode potential, E (solid curve), in comparison with integrated intensity, I_{SER} , of SERS $\text{Cr}^{\text{III}}\text{-NH}_3$ stretching mode (450 cm^{-1}) against E (dashed curve). Solution contained $3\text{ mM Cr}(\text{NH}_3)_5\text{Br}^{2+}$ in $0.1\text{ M NaClO}_4 + 5\text{ mM NaCl} + 2\text{ mM HClO}_4$. Silver electrode was rotated at 800 r.p.m. Maximum value of I_{SER} set equal to maximum value of i .

Figure 3 Plot of current density, i , for $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ reduction against electrode potential, E (solid curve), in comparison with integrated intensity, I_{SER} , of SERS C-N stretching mode ($2090 - 2130\text{ cm}^{-1}$) against E (dashed curve). Solution contained $1.2\text{ mM Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ in $0.1\text{ M NaClO}_4 + 5\text{ mM NaCl} + 2\text{ mM HClO}_4$. Silver electrode was rotated at 800 r.p.m. Maximum value of I_{SER} set equal to maximum value of i . Dotted-dashed curve is corresponding plot of peak frequency of C-N stretching mode, ν_{CN} , (right-hand y axis) against electrode potential.

Figure 4 Cathodic-anodic cyclic voltammogram (solid curves) for reduction of adsorbed $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ at silver electrode. Electrolyte conditions as in Fig. 3 except that $\text{Cr}(\text{NH}_3)_5\text{NCS}^{2+}$ concentration was 50 μM ; electrode was electropolished as in ref. 15. Sweep rate was 50 V sec^{-1} . Dashed curve is corresponding voltammogram obtained in absence of adsorbed reactant.