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Hydrogen Oxidation and Evolution Reaction Kinetics on Platinum: Acid vs Alkaline Electrolytes

Wenchao Sheng,a,d Hubert A. Gasteiger,e,*,z and Yang Shao-Hornb,c,d,z

The kinetics of the hydrogen oxidation reaction (HOR) and hydrogen evolution reaction (HER) on polycrystalline platinum [Pt(pc)] and high surface area carbon-supported platinum nanoparticles (Pt/C) were studied in 0.1 M KOH using rotating disk electrode (RDE) measurements. After corrections of uncompensated solution resistance from ac impedance spectroscopy and of hydrogen mass transport in the HOR branch, the kinetic current densities were fitted to the Butler–Volmer equation using a transfer coefficient of \( \alpha = 0.5 \), from which HOR/HER exchange current densities on Pt(pc) and Pt/C were obtained, and the HOR/HER mechanisms in alkaline solution were discussed. Unlike the HOR/HER rates on Pt electrodes in alkaline solution, the HOR/HER rates on a Pt electrode in 0.1 M HClO4 were limited entirely by hydrogen diffusion, which renders the quantification of the HOR/HER kinetics impossible by conventional RDE measurements. The simulation of the hydrogen anode performance based on the specific exchange current densities of the HOR/HER at 80°C illustrates that in addition to the oxygen reduction reaction cell voltage loss on the cathode, the slow HOR kinetics are projected to cause significant anode potential losses in alkaline fuel cells for low platinum loadings (>130 mV at 0.05 mgPt/cm2anode and 1.5 A/cm2anode), contrary to what is reported for proton exchange membrane fuel cells.

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pensions of Pt/C of −0.15 mg/mL were obtained by dispersing the catalyst in deionized (DI) water (18.2 MΩ cm, Millipore) using ultrasonication of the suspension in an ice bath. 20 μL of the Pt/C catalyst water suspension was deposited on a glassy carbon electrode (GCE) disk (5 mm diameter, Pine Instruments), which was polished with 0.05 μm alumina (Buehler), and dried in air at room temperature, resulting in a Pt loading of ~7.0 μgPt/cm² on GCE. The Pt(pc) disk electrode (5 mm in diameter, Pine Instruments) was also polished with 0.05 μm alumina and was cleaned by ultrasonication in DI water.

Modification of the reference electrode.— Glass corrodes in an alkaline solution and the arising contaminants (lead and silica) can significantly influence the ORR and the HOR/HER activity of Pt electrodes.25,26 To minimize contamination from the gradual dissolution of glass, a fresh alkaline electrolyte was prepared for each electrochemical measurement at each temperature, and all electrochemical data were collected within 30 min. More importantly, we also modified the Luggin capillary, replacing the highly KOH-soluble Vycor glass tip of the Luggin capillary with a 50 μm thick Nafion film, analogous to what was shown previously.27 The short exposure time and modification of the Luggin capillary tip eliminated contamination; consequently, as shown later, the HOR/HER and ORR activities obtained in our study are equal to those reported for measurements in a Teflon cell.26

Electrochemical measurements.— The electrochemical measurements were carried out in a jacketed glass cell (Pine Instruments) connected to a circulating temperature-controlled water bath (Thermo Electron Corp.). As-prepared Pt/C thin-film electrodes or a Pt(pc) disk was mounted onto a rotator (Pine Instruments) and was immersed into 0.1 M potassium hydroxide (99.99% purity, Sigma- Aldrich) solution. A Pt wire served as the counter electrode and a saturated calomel electrode (Analytical Sensor, Inc.), immersed into the Nafion film-modified Luggin capillary filled with 0.1 M KOH, was used as the reference electrode. However, all potentials reported in this paper are referenced to the reversible hydrogen electrode (RHE) potential, calibrated in the same electrolyte by measuring HOR/HER currents on the Pt RDE, whereby the potential at zero current corresponds to 0 V vs RHE. After the electrolyte was saturated with argon, the cyclic voltammogram (CV) at 294 K was recorded between −0.08 and −1.0 V vs RHE at a scan rate of 50 mV/s after it reached steady state.

The HOR/HER rates were measured using RDE measurements. After the electrolyte was saturated with pure hydrogen, polarization curves were recorded between −0.08 and −1.0 V vs RHE at a sweep rate of 10 mV/s and rotation rates of 400, 900, 1600, and 2500 rpm. A freshly prepared 0.1 M KOH electrolyte was used for each measurement at each temperature, as described above. Exchange current densities (i0) of the HOR/HER were obtained by fitting the experimental data to the Butler–Volmer equation. The HOR/HER was also measured on Pt(pc) in H2-saturated 0.1 M HClO4 (70%, GFS Chemicals Inc.) using the same method and experimental conditions.

After measuring the HOR/HER at 294 K, the ORR kinetics were also measured on Pt(pc) and Pt/C in O2-saturated 0.1 M KOH at 1600 rpm and 10 mV/s. The kinetic current at 0.9 V vs RHE was calculated based on the Koutecky–Levich equation

\[
i = \frac{i_0 + i_d}{1 + \frac{i_0}{i_d}} = \frac{1}{B_{\text{RHE}}/i_d^{1/2}} \]

where i is the measured current, i0 is the kinetic current, id is the diffusion limited current, s is the rotation rate, and \(B_{\text{RHE}}\) is a constant related to the number of electrons transferred in a reaction, the gas concentration and diffusivity, as well as the electrolyte kinematic viscosity.28 The kinetic currents were then normalized to the electrochemical surface area (ESA) to obtain the specific activity and, in Pt/C, also to the Pt mass to obtain the mass activity.

Figure 1. (Color online) CVs of Pt(pc) at 294 ± 1.5 K in Ar-saturated 0.1 M KOH at 50 mV/s with two different positive potential limits. The current density is referenced to the actual Pt surface area.

Impedance measurements.— The cell resistance was measured immediately after RDE measurements using VoltaMaster (Voltlab), taking the ac impedance spectra from 32 to 0.1 kHz and a voltage perturbation of 10 mV. The real part of the resistance at 1 kHz was taken as the cell resistance and was used to obtain the iR-free potential of the working electrode. The solution resistances measured at 275 and 314 K were 65 and 35 Ω, respectively. These rather high uncompensated solution resistances in the relatively conductive 0.1 M KOH were due to the highly skewed potential distribution near the RDE surface,29 with half of the potential drop in the electrolyte phase occurring at a distance of only a disk radius from the disk surface.

Results

Measurements of HOR/HER kinetics of Pt(pc).— The CV of a Pt(pc) disk in 0.1 M KOH (Fig. 1) shows the typical Pt–H underpotential deposition region (0.08–0.5 V), double-layer region (0.5–0.6 V), and Pt–oxide region (above 0.6 V), similar to that reported previously.30 The Pt–H peaks at ~0.3 and ~0.4 V can be attributed to the Pt–H interaction on Pt(110) and Pt(100) planes, respectively, based on the CVs of Pt single-crystal surfaces in 0.1 M KOH.25–27 To determine the Pt surface area, the positive potential window was restricted to 0.6 V to eliminate the interference from Pt oxide reduction in quantifying the double-layer contribution to the H adsorption/desorption region. The Pt surface area was then obtained by integrating the H adsorption/desorption region, subtracting the double-layer charging currents, and dividing the resulting coulombic charge by 2 and 210 μC/cm².23,30 The thus obtained roughness factor of the Pt(pc) disk from four independent experiments, which was defined as ESA/area_sdisk, was ~1.6 ± 0.2 cm²/cm².

Figure 2a shows the HOR/HER polarization curves on Pt(pc) in 0.1 M KOH at 294 K as a function of rotation rate, with well-defined hydrogen mass transport controlled (limiting) current densities occurring at potentials above 0.2 V. They scale linearly with the inverse of the square root of the rotation rate, as shown by the Koutecky–Levich plot constructed at 0.5 V (see inset of Fig. 2a), yielding a zero y-axis intercept and a \(B_{\text{C}}\) value (Eq. 1) of 0.0664 mA/(cm²·cmH₂·rpm). The limiting current density of 3.31 mA/cm² at 2500 rpm agrees excellently with the value obtained in a Teflon cell,26 where it was shown from KOH-induced glass corrosion would lead to substantially lower limiting current densities. In addition, our measured HOR half-wave potential of 0.065 V on Pt(pc) at 2500 rpm and 294 K (Fig. 2a) is actually slightly higher than that of 0.08 V obtained in a Teflon cell26 at essentially the same conditions. The identical diffusion limited current densities and the slightly higher HOR activity in our study compared to the data reported for measurements in a Teflon cell26.
Figure 2. (Color online) (a) HOR/HER polarization curves (positive-going scans) on Pt(pc) at different rotation rates at 294 ± 1.5 K. The inset shows a Koutecky–Levich plot at 0.5 V/RHE. (b) HOR/HER polarization curves on Pt(pc) at 1600 rpm at different temperatures (positive-going scans). The data were collected in H2-saturated 0.1 M KOH at 10 mV/s.

The HOR/HER kinetic current densities \( i_0 \) on Pt(pc) are shown as gray solid lines in Fig. 3a and b (at 275 and 314 K, respectively), which were obtained from correcting the polarization curves by the measured IR and the hydrogen mass transport in the HOR branch (Eq. 1). The voltage corrections associated with the IRs can be quite substantial at the higher end of the current density scale, amounting to, for example, 25 mV (275 K) and 12 mV (314 K) at 2 mA/cm\(^2\)\(_{\text{disk}}\), as shown in the insets of Fig. 3a and b. Following the IR correction, the HOR/HER kinetic current densities were obtained by hydrogen mass transport resistance correction for the HOR branch using the Koutecky–Levich equation (Eq. 1). Only data below 80% of the HOR diffusion limited current density were used to minimize errors in the analysis, which can become substantial as the measured current approaches the mass transport limited current. Because hydrogen oversaturation and bubble formation can introduce mass transport resistances for the HER at high current densities, only data with HER current densities equal to or lower than \(-2 \text{ mA/cm}^2_{\text{disk}}\) were used, where no bubble formation was observed and the measured current was rotation rate independent (between 900 and 2500 rpm).

The thus obtained HOR/HER kinetic current densities \( i_0 \) on Pt(pc) were then plotted in Fig. 3 (gray solid lines) at (a) 275 and (b) 314 K. The HOR/HER exchange current densities \( i_0 \) on Pt(pc) were extracted subsequently by fitting the HOR/HER kinetic current densities to the Butler–Volmer equation (dashed black lines in Fig. 3a and b) as a function of the HOR/HER overpotential \( \eta \)

\[
i_k = i_0 e^{(\alpha F \eta)/R T} \left[ 1 - e^{-(1-\alpha) F \eta/R T} \right]
\]

where \( \alpha \) represents the transfer coefficient, \( F \) is Faraday’s constant (96,485 As/mol), \( R \) is the universal gas constant (8.314 J/mol/K), and \( T \) is the temperature in Kelvin. In fitting all the HOR/HER data in this study, \( \alpha \) and \( i_0 \) were first set as variables, where the best fit was obtained for \( \alpha = 0.5 \). Therefore, the reported \( i_0 \) values herein were obtained by setting \( \alpha = 0.5 \). Fig. 3 shows that the simple Butler–Volmer equation with \( \alpha = 0.5 \) perfectly fits the experimental HOR/HER kinetics on Pt(pc) at both 275 and 314 K, but equally good fits were obtained at intermediate temperatures, corresponding to Tafel slopes (TS) defined as \( \text{TS} = 2.303 R T / (\alpha F) \) with values of 109 mV/dec at 275 K and 125 mV/dec at 314 K. The symmetry of the anodic (HOR) and cathodic (HER) current densities with respect to the reversible potential suggests that the HOR and HER have the same reaction mechanism and intermediates, which is discussed in more detail later. The above determined potential-independent TSs of \(-120 \text{ mV/dec} \) disagree with the variable TSs ranging from \(-50 \text{ to } -150 \text{ mV/dec} \) at low and high overpotentials, respectively, reported for the HOR/HER on Pt(hkl) in 0.1 M KOH.\(^{20,21}\) However, the origin of this discrepancy is simply a misinterpretation of the definition of the TS because the latter can only be obtained from a...
Table I. Summary of average HOR/HER exchange current densities ($i_0$), mass activity ($i_{0,m}$) at 294 ± 1.5 K, and activation energy for Pt(pc) and Pt/C in 0.1 M KOH.

<table>
<thead>
<tr>
<th></th>
<th>$i_0$ at 294 K (mA/cm$^2$)</th>
<th>$i_{0,m}$ at 294 K (A/mg$_{Pt}$)</th>
<th>$E_a$ (kJ/mol)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pt(pc)</td>
<td>0.69 ± 0.03 (4')</td>
<td>–</td>
<td>28.9 ± 4.3 (4')</td>
</tr>
<tr>
<td>Pt/C</td>
<td>0.57 ± 0.07 (2')</td>
<td>0.35 ± 0.05 (2')</td>
<td>29.5 ± 4.0 (2')</td>
</tr>
</tbody>
</table>

The asterisk indicates the number of independent repeat experiments. Activation energies ($E_a$) were obtained by fitting the data between 275 ± 1.5 and 314 ± 1.5 K. Data were iR- and mass-transport corrected as described in the text.

The plot of log($i_0$) vs $\eta$ only if $\eta$ is significantly larger in value than roughly one-half of the TS, so that the fitting of a TS to the low overpotential region in these studies is incorrect. Consequently, only their reported value of −150 mV/dec at high overpotentials is determined correctly and is reasonably close to the TS obtained in our work. Based on fitting at least three independent data sets at each temperature, specific HOR/HER exchange current densities of Pt(pc), normalized to the Pt ESA, are 0.25 ± 0.01, 0.69 ± 0.01, and 1.26 ± 0.30 mA/cm$^2$ at 275, 294, and 314 K, respectively. At 294 K, our value of 0.69 mA/cm$^2$ (listed in Table I) is larger than the ~0.1 mA/cm$^2$ reported for Pt(pc) in 0.5 M NaOH 4 and for Pt(110), which was the most active surface for the HOR/HER among the three low index Pt single-crystal faces reported in an early study. The origin of this discrepancy is likely caused by both the lack of iR correction in these studies and electrolyte contamination effects, which is supported by the fact that the more recently measured HOR/HER specific exchange current density on Pt(110) at 293 K of 0.6 mA/cm$^2$ 20 is much larger and comparable to our value on Pt(pc) (the exchange current density given in Ref. 20 must be multiplied by a factor of 2 to be consistent with the definition used in Eq. 2 here). In addition, the activation energy of the HOR/HER on Pt(pc) was obtained by plotting $i_0$ vs 1/$T$; where a linear relationship was found between 275 and 314 K. A representative data set is shown in Fig. 4. Based on four independent sets of repeat measurements, the HOR/HER activation energy ($E_a$) on Pt(pc) was determined to be 28.9 ± 4.3 kJ/mol, which is listed in Table I. This value agrees reasonably with 23 kJ/mol obtained for Pt(110) in the same electrolyte. 20

Measurements of HOR/HER kinetics of Pt/C.—The same HOR/HER electrochemical measurements as on Pt(pc) were made on Pt/C. Fig. 5 depicts the CV of Pt/C in 0.1 M KOH at 294 K and 50 mV/s, where the current was normalized to the true Pt surface area. The observed features resemble those of Pt(pc) except that it has higher currents in the double-layer region due to the capacitive contribution from the high surface area carbon support. Using the double-layer-corrected Pt–H underpotential deposition region, the specific surface area of Pt/C was determined to be 62 m$^2$/g$_{Pt}$, corresponding to a roughness factor of 4.3 cm$^2$/cm$^2$ for thin-film RDE electrodes of Pt/C.

HOR/HER kinetic current densities of Pt/C RDE electrodes are shown in Fig. 6 and were extracted from HOR/HER polarization curves following processes analogous to what was used for Pt(pc) in Fig. 3. The hydrogen transport limited current densities on Pt/C RDE electrodes (not shown) were within 10% of those shown for Pt(pc) in Fig. 2. The fitting of the HOR/HER kinetic current densities using the Butler–Volmer equation with $\alpha = 0.5$ was excellent, from which the exchange current density $i_0$ was obtained. The experimentally extracted and fitted kinetic current densities at 275 and 304 K are shown in Fig. 6 as examples, whereas fits at intermediate temperatures (data not shown) are equally good. The specific exchange current density of the HOR/HER on Pt/C at 294 K is 0.57 ± 0.07 mA/cm$^2$ which is comparable to that obtained on Pt(pc) (Table I). Normalized to Pt mass, the mass exchange current density of Pt/C is 0.35 ± 0.05 A/mg$_{Pt}$. A representative Arrhenius plot of $i_0$ on Pt/C vs 1/$T$ ranging from 275 to 304 K is shown in Fig. 4. Based on two independent repeat measurements, Pt/C has an HOR/HER activation energy of 29.5 ± 4.0 kJ/mol, which is essentially identical to that on Pt(pc) (Table I), considering experimental uncertainty. There are essentially identical specific exchange current densities, activation energies, and transfer coefficients for the HOR/HER on Pt(pc) thus on Pt/C, and it is hypothesized that there is no significant Pt particle-size effect for the HOR/HER in 0.1 M KOH. Considering the roughly 10-fold lower specific exchange current densities reported for Pt(111) compared to Pt(110), future studies on Pt NPs of different shapes are needed to examine this hypothesis further. To our knowledge, the HOR/HER kinetics on Pt/C in alkaline electrolytes have never been reported before and this is the first comparison of the HOR/HER kinetics of Pt(pc) and Pt NPs.

Discussion

Comparing HOR/HER kinetic data with proposed mechanisms in alkaline electrolyte.—Due to the experimental difficulties of measuring the kinetic reaction rates of very fast reactions, the HOR/HER mechanism on Pt remains inconclusive and under active inves-
For the reactions occurring in an alkaline solution\textsuperscript{32}, although the Heyrovsky and Volmer reactions as written above appear in some textbooks\textsuperscript{27} as the Heyrovsky–Volmer mechanism,\textsuperscript{32,34} the three elementary steps \textit{Heyrovsky}–\textit{Volmer} and \textit{Tafel}–\textit{Volmer} will be assumed and successfully described the HER activity trends of a wide range of metal catalysts in an acid electrolyte\textsuperscript{35}. This and later DFT\textsuperscript{36} calculations, nominally evaluated in an acid electrolyte, show that the following elementary steps: the dissociative adsorption of H\textsubscript{2} without electron transfer (Tafel reaction, Eq.\textsuperscript{5}) or with simultaneous electron transfer (Heyrovsky reaction, Eq.\textsuperscript{4}) and the discharge of adsorbed hydrogen (H\textsubscript{ad}) (Volmer reaction, Eq.\textsuperscript{5}), whereby the asterisk represents an adsorption site on Pt\textsuperscript{32,34}. The simplest kinetic HOR/HER models assume that one of the reactions in each of the two reaction sequences is rate-determining (rds),\textsuperscript{32,34} which leads to four limiting cases: Tafel–Volmer (rds), Heyrovsky–Volmer (rds), Heyrovsky (rds)–Volmer, and Tafel (rds)–Volmer.

In the following, we review and discuss the kinetic HOR/HER expressions for the first two cases (Volmer reaction being rds) in an alkaline solution, which was first derived by Vetter.\textsuperscript{25} The anodic reaction rate of Eq.\textsuperscript{7} is proportional to the surface coverage of H\textsubscript{ad}, \theta_{H\textsubscript{ad}} and the activity of OH\textsuperscript{−}, whereas the cathodic reaction rate is proportional to the coverage of unoccupied surface sites (1 − \theta_{H\textsubscript{ad}}) as well as the activity of H\textsubscript{2}O. Using the Butler–Volmer equation (Eq.\textsuperscript{2}) to describe this elementary reaction step, the kinetic expression can be written

\[ i_k = k_0 \theta_{H\textsubscript{ad}} \alpha F \exp \left( \frac{\alpha F}{RT} \right) - k_1 \]

\[ \text{where } e_V \text{ represents the electrode potential across the solid/liquid interface measured against an arbitrary reference electrode. At the reversible potential of the Volmer reaction \( e_{0,V} \), Eq.\textsuperscript{8} can be rewritten as} \]

\[ i_{0,V} = k_0 \theta_{H\textsubscript{ad}} \alpha F \exp \left( \frac{\alpha F}{RT} e_{0,V} \right) = k_0 \]

\[ \text{where } i_{0,V} \text{ is the exchange current density of the Volmer reaction. Dividing Eq.\textsuperscript{8} by Eq.\textsuperscript{9} yields} \]

\[ i_k = i_{0,V} \left\{ \frac{\theta_{H\textsubscript{ad}}}{\theta_{H\textsubscript{ad}}} \exp \left( \frac{\alpha F}{RT} (e_V - e_{0,V}) \right) \right\} \]

\[ \text{in the assumed case that the Volmer reaction is the rds and } (e_V - e_{0,V}) \text{ corresponds to the overpotential (}\eta\text{) of the overall HOR/HER. Furthermore, if the HOR/HER kinetics are measured in a reasonably narrow potential range (e.g., from −0.075 to +0.15 V in our study), the variation in \theta_{H\textsubscript{ad}} is small (i.e., } \theta_{H\textsubscript{ad}} \approx \theta_{H\textsubscript{ad}}^0). \text{To a first approximation, the terms } \theta_{H\textsubscript{ad}}^0, \theta_{H\textsubscript{ad}}^0 \text{ and } (1 - \theta_{H\textsubscript{ad}})/(1 - \theta_{H\textsubscript{ad}}^0) \text{ can be approximated to be one, as the potential dependence of the exponential terms influences kinetic current densities much more strongly. Thus, Eq.\textsuperscript{10} can be approximated as} \]

\[ i_k = i_{0,V} \left\{ \exp \left( \frac{\alpha F}{RT} \eta \right) - \exp \left( - \frac{(1 - \alpha) F}{RT} \eta \right) \right\} \]

\[ \text{Formally, Eq.\textsuperscript{11} is equivalent to the Butler–Volmer equation (Eq.\textsuperscript{2}), which was used to fit the HOR/HER kinetic current densities in our study, providing an excellent fit with all the experimental data (Fig.\textsuperscript{3} and \textsuperscript{6}).} \]

An analogous derivation assuming the Heyrovsky reaction to be the rds is given in Ref.\textsuperscript{32}, yielding the same expression as Eq.\textsuperscript{11} in the limit that \theta_{H\textsubscript{ad}} = \theta_{H\textsubscript{ad}}^0. Therefore, the Butler–Volmer equation given by Eq.\textsuperscript{11} describes the HOR/HER kinetics in the vicinity of the HOR/HER equilibrium potential for three of the four limiting cases, i.e., for the Tafel–Volmer(rds), Heyrovsky–Volmer(rds), and Heyrovsky(rds)–Volmer mechanisms.\textsuperscript{32,34} A very different kinetic expression is obtained only for the Tafel(rds)–Volmer mechanism,\textsuperscript{32,34} so that it can be excluded as a possible HOR/HER mechanism in alkaline electrolytes on the basis of our measurements.

In recent density functional theory (DFT) calculations, a Tafel–Volmer mechanism was assumed and successfully described the HER activity trends of a wide range of metal catalysts in an acid electrolyte.\textsuperscript{35} This and later DFT\textsuperscript{36} calculations, nominally evaluated...
for pH 0. should apply equally to alkaline pH, as long as the potential scale is referenced to RHE. In the later DFT study by the same group on the HER mechanism on Pt(111),35 however, it was concluded that the activation barrier at zero overpotential of the Volmer reaction (−0.15 eV) is substantially lower than that of the Tafel (−0.8 eV) and Heyrovsky (−0.6 eV) reactions, suggesting that either one or both of the latter reactions would be rds. The calculated transfer coefficients for the Tafel and Heyrovsky reactions were $\alpha = 0.64$ and $\alpha = 0.45$, respectively.13 One major discrepancy between experimental data and their DFT calculations noted by the authors is the fact that the calculated activation barriers (0.6–0.8 eV) are so much larger than the experimentally reported activation energies on Pt(hkl) in acid electrolytes (<0.2 eV14) measured via RDE. As argued below, meaningful HOR/HER kinetics in an acid electrolyte cannot be obtained by RDE measurements, so that these experimental activation energies are not meaningful. However, experimental RDE-based activation energies in 0.1 M KOH, where the HOR/HER kinetics are slow, should be reliable and, indeed, the value of $\sim 0.5$ eV on Pt(111) in 0.1 M KOH reported by Schmidt et al.24 is quite comparable with the DFT-based values, considering the expected uncertainties of the latter. If one were to assume that the basic learnings from the DFT model33 are correct, i.e., that the Volmer reaction would have the lowest activation barrier and that the Heyrovsky reaction would have a transfer coefficient of $\alpha = 0.45$, our experimental observation that the HOR/HER on Pt in 0.1 M KOH can be described by a simple Butler–Volmer equation with $\alpha = 0.5$ would favor the Heyrovsky(rds)–Volmer mechanism. However, this hypothesis is very tentative. Further experimental studies on the effect of pH and the partial pressure of H$_2$ on the HOR/HER kinetics should be carried out to explore its mechanism in more detail.

Comparing HOR/HER kinetics in acid with alkaline electrolyte.— Although HOR/HER kinetics on Pt in alkaline can be measured accurately from conventional RDE measurements with or without iR correction and the HOR/HER kinetics on Pt in acid cannot be separated from the hydrogen diffusion overpotential in RDE measurements due to the very fast reaction rate. To illustrate this point, we conducted RDE measurements of the HOR/HER on Pt(pc) in 0.1 M HClO$_4$, including appropriate correction of the uncompensated solution resistance measured by ac impedance. Fig. 7a shows the polarization curve of the HOR/HER on the Pt(pc) electrode at 1600 rpm and 294 K before (solid black lines) and after (dotted red lines) iR correction; the dashed black lines represent the calculated Nernstian diffusion overpotential ($\eta_{\text{diffusion}}$), assuming ideal reversibility of the HOR/HER, i.e., infinitely fast reaction kinetics$^{12,32}$

$$\eta_{\text{diffusion}} = \frac{RT}{2F} \ln \left(1 - \frac{i}{i_c}\right)$$  \[12\]

where $i_c$ is the hydrogen diffusion limited current density. Clearly, after iR correction, the experimental current densities of the HOR/HER are indistinguishable from the Nernstian diffusion overpotential and therefore the latter assumes infinitely fast reaction kinetics, it is not reasonable to attempt to extract kinetic rates for the HOR/HER on Pt in acid using the RDE method, consistent with previous conclusions.13 Consequently, the several orders of magnitude lower HOR/HER exchange current density values from RDE measurements (0.1 to 1 mA/cm$^2$) compared to those obtained from microelectrode$^{16,17}$ or gas-diffusion electrode$^{2,12,14,31}$ measurements are caused by both the rather low hydrogen diffusion limited current densities in the RDE configuration (on the order of several mA/cm$^2$) as well as the lack of iR correction. Kinetic currents cannot be measured with reasonable precision if the exchange current density is one or two orders of magnitude larger than the diffusion limited current density. Consequently, many RDE-based studies on the HOR/HER on Pt in an acid electrolyte report TS values in the vicinity of TS $= 2.303RT/(2F)$,14,36 which simply reflects the apparent TS of the Nernstian diffusion overpotential (Eq. 12). Finally, based on RDE and microelectrode data for the HOR on Pt in acid electrolytes, Wang et al.17 recently developed a complex kinetic model, which suggests that the HOR proceeds through parallel Heyrovsky–Volmer and Tafel–Volmer mechanisms. However, considering the difficulty of extracting HOR kinetics on Pt in an acid electrolyte from RDE measurements, the complexity of the model is perhaps not warranted by the uncertainties associated with available electrochemical data.

However, the iR-corrected polarization curve of the HOR/HER on Pt(pc) in 0.1 M KOH under the same experimental conditions deviates substantially from the Nernstian diffusion overpotential, as demonstrated in Fig. 7b. This is caused by the much slower HOR/HER kinetics on Pt in KOH, so that the diffusion limited current density is only reached at $\sim 0.25$ V (the diffusion limited current densities are essentially the same in both 0.1 M KOH and 0.1 M HClO$_4$). Because the difference between the Nernstian diffusion overpotential relationship and the experimental data in KOH is very large, kinetic parameters for the HOR/HER can be extracted easily without interference from mass transport resistances. As was shown, this is not the case in an acidic electrolyte (see Fig. 7a), which is the reason why reliable HOR/HER kinetics on Pt with the RDE method can be obtained in alkaline but not in acid electrolytes. The fact that the HOR/HER kinetics on Pt are slower in alkaline compared to an acid electrolyte was noted before,15 but its cause remains unclear.

Figure 7. (Color online) (a) HOR/HER polarization curve on Pt(pc) in 0.1 M HClO$_4$ at 1600 rpm before ($E_{\text{RDE}}$, solid black lines) and after ($E_{\text{RDE, iR-free}}$, dotted red lines) iR correction. The inset is the Koutecky–Levich plot obtained at $E = 0.1$ V/RHE for the HOR at different rotation rates. (b) HOR/HER polarization curve on Pt(pc) in 0.1 M KOH at 1600 rpm before ($E_{\text{RDE, iR-free}}$, solid black lines) and after ($E_{\text{RDE, iR-free}}$, dotted red lines) iR correction; the dashed black lines are the Nernstian diffusion overpotential calculated according to Eq. 12. The data are shown for the positive-going scans at 10 mVs$^{-1}$ at 294 ± 1.5 K. The $x$-axis potential range in (b) is double of that in (a).
activities measured on 5 wt % Pt/C in a PEMFC at 353 K. Pt/C catalyst used in Ref.12 extrapolated to 353 K compared to that obtained in 0.1 M KOH in this study. The resulting polarization curves in KOH are compared to that measured in PEMFCs, which is shown in Fig. 8b. Considering an envisaged anode catalyst loading of 0.05 mg/cm$^2$ and an operating current density of 1.5 A/cm$^2$, corresponding to a required mass activity of 30 A/mgPt. Fig. 8b illustrates that one would expect a hydrogen anode overpotential in an AFC/AMFC of 130–150 mV depending on the specific Pt surface area. This is in stark contrast to the negligibly small anode overpotential under the same conditions measured in a PEMFC. This projection clearly emphasizes that ultralow Pt anode loadings are not feasible in AFCs/AMFCs (in contrast to PEMFCs) and that novel anode catalysts must be developed for these applications, which either have a much higher mass activity for platinum group metals (PGM)-based anodes or are PGM-free.

Lastly, we compare the ORR kinetics of Pt/C in alkaline with that in acid and discuss the implications in the performance of AFCs/AMFCs relative to PEMFCs. As Pt NPs are shown to exhibit a much lower ORR activity than Pt bulk surfaces in acid, we examined here the ORR specific activity of both Pt(pc) and Pt/C in 0.1 M KOH at 294 K and 100 kPa$_{\text{O}_2}$. The activities at 0.9 V are summarized in Table II. The ORR activity of Pt(pc) is 0.95 ± 0.25 mA/cm$^2$, which is consistent with the value of 0.43 mA/cm$^2$ reported in a Teflon cell (confirming our measurements of ORR activity are free of glass-cell related contamination effects). Unlike HOR/HER kinetics on Pt, the ORR specific activity of Pt/C is roughly two times lower than that of Pt(pc) (Table II), indicating a Pt particle-size effect on the ORR kinetics in alkaline. The ORR specific activities of Pt/C and Pt(pc), and the mass activities of Pt/C in 0.1 M KOH are identical to those in 0.1 M HClO$_4$ within the experimental error (Table II). This is perhaps not surprising because the ORR rate-controlling formation of adsorbed oxygen-containing species (OH$_{ad}$) occurs at essentially the same potential vs RHE in both electrolytes. For practical applications, the identical ORR activities in acid and base predict that the ORR overpotential on a Pt cathode in AFCs/AMFCs would be the same as that of PEMFCs under the same conditions (i.e., under the same mass specific current density (A/g$_{Pt}$), temperature, and oxygen partial pressure).

### Conclusion

Our RDE measurements show that the HOR/HER on Pt(pc) and Pt/C in an alkaline electrolyte have very comparable specific exchange current densities (showing no size effect of HOR/HER on Pt) of −0.6 mA/cm$^2$ at 294 K, and activation energies of −29 kJ/mol. In addition, based on the fact that the kinetics of the

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<th>Table II. Summary of ORR specific activity ($i_{m,0.9}, \text{V}<em>{294}, \text{K}$) and mass activity ($i</em>{m,0.9}, \text{V}<em>{294}, \text{K}$) at 0.9 V vs RHE, 100 kPa$</em>{\text{O}_2}$, and 294 ± 1.5 K in 0.1 M KOH and 0.1 M HClO$_4$ for Pt(pc) and Pt/C.</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Pt(pc)</strong></td>
</tr>
<tr>
<td>$i_{m,0.9}, \text{V}_{294}, \text{K}$ (mA/cm$^2$)</td>
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<tr>
<td>$i_{m,0.9}, \text{V}<em>{294}, \text{K}$ (A/mg$</em>{Pt}$) in 0.1 M KOH</td>
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<tr>
<td>$i_{m,0.9}, \text{V}_{294}, \text{K}$ (mA/cm$^2$) in 0.1 M HClO$_4$</td>
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<tr>
<td>$i_{m,0.9}, \text{V}<em>{294}, \text{K}$ (A/mg$</em>{Pt}$) in 0.1 M HClO$_4$</td>
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The asterisk indicates the number of independent repeat experiments. Data were obtained from the positive-going scans at 10 mV/s and were iR- and mass-transport corrected as described in the text.
HOR/HER in our study are very well described by the Butler–Volmer equation and considering recent DFT findings, it is hypothesized that the mechanism of the HOR/HER in KOH may go though the Heyrovsky–Volmer mechanism with the Heyrovsky step as the rds.

Analogous RDE measurements of the HOR/HER on Pt(pc) in 0.1 M HClO4 clearly demonstrate that the HOR/HER current densities in an acid solution simply follow the Nernstian hydrogen diffusion overpotential relationship, which is based on assuming reversible, i.e., infinitely fast HOR/HER kinetics. This implies that the HOR/HER exchange current densities in an acid electrolyte are one or more orders of magnitude larger than the typical hydrogen diffusion limited current density and can therefore not be quantified in RDE measurements. This finding is consistent with the much higher HOR/HER exchange current densities in measurements based on microelectrode or gas-diffusion electrodes.

Our measurements and analysis suggest that the HOR/HER kinetics on Pt(pc) and Pt/C are several orders of magnitude slower in alkaline electrolyte and can therefore not be quantified in RDE measurements. This work was supported in part by the DOE Hydrogen Initiative program under award no. DE-FG02-05ER15728, an Air Products Faculty Excellence grant, and the Toyota Motor Co. This research made use of the Shared Experimental Facilities supported by the MRSEC Program of the National Science Foundation under award no. DMR 08-19762.

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