HYDROGEN OVERVOLTAGE ON PLATINUM IN AQUEOUS SODIUM HYDROXIDE

Part I. Rate of The Discharge Step

By

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Abstract

Hydrogen overvoltage on Pt in aqueous NaOH and Na2SO4 was studied by means of the galvanostatic transient methods to distinguish the electrostatic part of the overvoltage from that due to the electronic work function of the electrode. The differential capacity of the electric double layer at the metal-solution interface and the rates of the electron transfer step were determined from the overvoltage-time curves at various values of pH and Na concentration as a function of overvoltage. It was concluded that: 1) the electron transfer step is the discharge of Na, but not of a water molecule, and its rate obeys FRUMKIN's equation with the Tafel constant \( \alpha = 1/2 \), 2) the exchange rate of the electron transfer step is proportional to the square root of Na concentration and 10\( \sim \)100 times greater than that of the overall reaction, 3) the greater part of the overvoltage in steady states is attributed to the change in the work function of the electrode due to the formation of intermetallic compounds or solid solutions between Pt and Na produced by the discharge step.

Introduction

The hydrogen overvoltage is defined as the difference in the chemical potential of metal electron \( \Delta \mu^e \) between the reversible and the irreversible states of the hydrogen electrode:

\[
\eta = -\Delta \mu^e / F \tag{1}
\]

where \( \mu^e \) is the chemical potential of metal electron and \( F \) the Faraday. The quantity \( \eta \) can be measured as the difference in the electrostatic potential between the lead wires of the same metal connected respectively to the reversible and the irreversible electrodes in the same environment. The chemical potential of metal electron is composed of the electrostatic potential \( \varphi \) and the

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165
electronic work function \( W \) of the metal as**)

\[ \mu' = -F(\varphi + W). \]  

(2)

The hydrogen overvoltage is thus given by the difference in the electrostatic potential \( \varphi \) and that in the electronic work function \( W \) between the irreversible and the reversible states of the hydrogen electrode:

\[ \eta = \Delta \varphi + \Delta W, \]  

(3)

Fig. 1. Schematic diagrams of the distribution of the electrostatic potential and the FERMI level of an electrode when the surface is negatively charged. Full and dotted lines correspond respectively to the reversible and the irreversible states of the electrode reaction.

(a): the overvoltage is caused by the change in the potential difference across the metal-solution boundary, \( \eta = \Delta \varphi \);

(b): the overvoltage is caused by the change in the work function of the electrode, \( \eta = \Delta W \).

\( \eta \) overvoltage, \( \varphi \) cavity potential of the electrode referred to that inside the solution, \( A \) electrode surface, \( E \) electrode, \( L \) lead wire, \( H_0 \) and \( H_0 \) inner and outer HELMHOLTZ planes, \( O \) inside the solution, \( F \) or \( F' \) FERMI level of electrons.

** The term electrostatic potential in the present paper means the cavity potential introduced by GRAHAME\(^1\) which is defined as the potential of a point within a macroscopic cavity in a conductor. The work function is thus defined as the work necessary to take out an electron from interior of a conductor into a macroscopic cavity in the conductor.
The $\Delta \varphi$ is caused by charge up of the electric double layer at the metal-solution interface, while $\Delta W$ causes the change in the contact potential between the electrode and the lead wire, each of which alters readings of potentiometer, as illustrated in Fig. 1. The former depends on the structure of the double layer at the metal-solution interface and the latter on the electronic states of the electrode.

When the discharge step is rate-determining, the reaction rate has been derived as a function of $\Delta \varphi$ by FRUMKIN in connection with the structure of the double layer at the metal-solution interface. In general case, however, neither the rate nor the species which discharges across the double layer has been separately determined because of experimental difficulties and in consequence no informations have been given on the intermediate species produced by the discharge step.

When the structural changes occur under the polarization due to intermediates produced by the discharge step or other elementary steps which compose the overall reaction, it is expected that the contact potential between the lead wire and the electrode or between the interior and the surface layers of the electrode metal itself plays an important role of determining the hydrogen overvoltage, as pointed out by LANGMUIR, since the work function of the electrode may be changed by the intermediates.

PAST and JOPA have shown on the basis of the galvanostatic transient studies that the build up and decay of the hydrogen overvoltage on mercury can be attributed to $\Delta \varphi$ in the initial stages of the build up and the decay of the overvoltage. The galvanostatic transient methods also are applicable in solid hydrogen electrodes for determining $\Delta \varphi$ separately and further the rate of the discharge step as a function of $\Delta \varphi$ on the basis of informations on the differential capacity of the double layer, identifying the latter as the leak current of the double layer.

The aim of the present work is, on one hand, to determine the differential capacity of the double layer at the metal-solution interface and to find the rate of the discharge step as a function of $\Delta \varphi$ by the galvanostatic transient methods on the platinum hydrogen electrode in alkaline solutions containing sodium ions, and on the other hand, to elucidate the contribution of the change in the contact potential between the electrode and the lead wire to the hydrogen overvoltage due to the intermediates produced by the discharge step or other elementary steps in these solutions.

1. Theoretical treatments of the overvoltage-time curves

It will be discussed here how to find the differential capacity of the double
layer at the metal-solution interface and the rate of the discharge step from the overvoltage-time curves, which enable the change in the potential difference across the double layer to be distinguished.

When a constant polarizing current is applied to a hydrogen electrode, the overvoltage increases with time until it reaches a steady value and when the current is switched off, the overvoltage decreases first rapidly approaching zero asymptotically. We first consider the build up of overvoltage after switching on a current. On switching the current, the electrode surface is charged by the polarizing current to increase the electrostatic potential of the electrode. The discharge of the hydrogen electrode now proceeds to form intermediates on the electrode surface. The hydrogen overvoltage is thus caused first by charge up of the double layer at the metal-solution interface and then by the increase of intermediate as well. From the initial stage of the overvoltage-time curve where the hydrogen overvoltage can be regarded as due solely to charge up of the double layer, it is possible to determine the differential capacity of the double layer and the time constant of the discharge step on the basis of the theory of the galvanostatic transients developed by Frumkin et al. A little different formula from that of Frumkin et al will be derived here for the analysis of the overvoltage-time curves in the region where the overvoltage increases due to charge up of the double layer at the metal-solution interface.

Electrons supplied by the polarizing current to the electrode are partly charged on the metal surface and partly consumed by the discharge reaction, i.e., the leakage of the double layer. From the material balance of electrons within time interval $\Delta t$, we have

$$\delta q = \delta q' - \delta q'', \quad (4)$$

where $\delta q$, $\delta q'$ and $\delta q''$ are respectively an increment of electricity charged on the electrode surface, supplied by the polarizing current to the electrode and used for the discharge reaction. These quantities are respectively expressed as

$$\delta q = C_n \delta \varphi, \quad (5a)$$
$$\delta q' = -i \Delta t, \quad (5b)$$
$$\delta q'' = -i_1 \Delta t, \quad (5c)$$

where $\varphi$ is the electrostatic potential of the electrode referred to that in the bulk of solution, $C_n$ the differential capacity of the electric double layer at the metal-solution interface, $i$ the polarizing current and $i_1$ the net rate of the discharge step expressed in terms of the electric current, the cathodic current being taken positive. Eliminating $\delta q$, $\delta q'$ and $\delta q''$ from Eqs. (4) and (5), we have
Hydrogen Overvoltage on Platinum in Aqueous Sodium Hydroxide Part I.

\[ C_D = -(i - i_i)/\hat{\psi}, \quad (6a) \]

or

\[ i_i = i + C_D \eta, \quad (6b) \]

which enables \( i_i \) to be evaluated when \( C_D \) is known in case where the variation of \( \eta \) consists practically in that of \( \phi \).

In the overvoltage region where \( |\eta| \ll RT/F \), the rate of the discharge step is proportional to the overvoltage on the basis of the general theory of Horiuti\(^6\) admitted that the latter is the only irreversible step at the moment in question,

\[ \eta = -r_1 i_i, \quad (7) \]

where

\[ r_1 = RT/F_i_0, \quad (8) \]

is a constant in terms of the exchange current \( i_0 \) of the discharge step. Eliminating \( i_i \) from Eqs. (6b) and (7) and integrating, we have

\[ \eta = r_1 i \{ \exp(-t/r_1) - 1 \}, \quad (9) \]

where

\[ \tau_1 = C_D r_1. \quad (10) \]

As seen from Eq. (9), \( \tau_1 \) gives approximately a measure of time in which \( \eta \) attains \(-r_1 i\) independent of the surface area of the electrode and is called the time constant of the discharge step. Differentiating Eq. (9) with time, we have

\[ \ln(-i/\hat{\eta}) = t/\tau_1 + \ln C_D. \quad (11) \]

which states a linear relation between \( \ln(-i/\hat{\eta}) \) and \( t \). The \( C_D \) and \( \tau_1 \) can be experimentally evaluated from the latter linear relation, if observed at all at various current densities.

We consider now the decay process of the hydrogen overvoltage starting from a steady state by switching off the polarizing current. At the moment when the current is switched off, the overvoltage decays by decrease of electrons charged on the electrode surface through the discharge step with the rate equal to the polarizing current, since the concentration of the intermediate is kept constant at the moment. It follows that the differential capacity \( C_D \) of the double layer can be determined from the initial rate of the decrease of the overvoltage as

\[ C_D = i/\eta t = 0. \quad (12) \]

When the overvoltage thus decays exclusively by discharge of the double layer, the rate of the discharge step \( i_i \) can be given by the following equation
Especially when the electrostatic component of the overvoltage in the steady state $\Delta \phi$ is small as compared with $RT/F$, putting $\Delta \phi = \eta_1$, we have similarly to the case of the build up

$$\eta_1 = -r_1 \dot{i}_1. \tag{14}$$

Eliminating $i_1$ from Eqs. (13) and (14) and putting $\dot{i} = \dot{\eta}_1$, we have

$$\eta_1 = -\tau \dot{\eta}_1, \tag{15}$$

where $\tau_1 = C_\rho \tau_1$ as in the case of build up. Integrating Eq. (15) with time and putting the initial condition $\eta_1 = -r_1 i$ at $t=0$, we have

$$\eta_1 = -r_1 i \exp \left( -t/\tau_1 \right). \tag{16}$$

Differentiating Eq. (16) with time and putting $\dot{\eta}_1 = \dot{\eta}$, we have similarly to the case of build up

$$\ln \left( i/\dot{\eta} \right) = t/\tau_1 + \ln C_\rho, \tag{17}$$

$C_\rho$ and $\tau_1$ are determined by fitting Eq. (17) to the initial stage as functions of the overvoltage from which the decay curve starts.

The experimental results in the galvanostatic transient studies will be described in the following section on the platinum hydrogen electrode in alkaline solutions including sodium ions.

2. Experimental procedures

Special precaution was taken of the preparation of solutions and electrodes and of the purification of glass material used for the construction of cells against impurities, which are known to affect sensitively the hydrogen electrode reaction.

2.1. The reaction apparatus and preparation of solutions

Glass materials used for the reaction apparatus were preliminarily purified by steaming in a hydrogen stream for several hours after ordinary treatments of boiling in a mixture of concentrated nitric and sulfuric acids and washing with hot equilibrium water. Fig. 2 shows the reaction apparatus made of Halio glass (SHIBATA Co. Tokyo). Vessels 1~11 show the hydrogen purification train which is composed of cylinder hydrogen (1), trap (2), silica gel (3), hop- kalite (4), soda lime (5), platinized asbestos (6), silica gel (7), trap (8), molecular sieve immersed in liquid nitrogen (9), liquid nitrogen traps (10) and (11). Silica gel and molecular sieve were preliminarily evacuated through stop cocks (A)
and (B) at temperatures 120 and 450°C respectively. Deionized water of specific resistance $10^7 \Omega$ was introduced into vessel (a), and distilled with addition of potassium permanganate in a nitrogen stream into vessel (b) containing barium hydroxide, and then the water was distilled into flask (c), where the water thus distilled twice was boiled for several hours in a stream of purified hydrogen to remove traces of organic substances. The water thus purified was now ready for distillation into vessel (d) in which the solution of desired concentration was prepared as follows. Guaranteed reagent of sodium hydroxide or sodium sulfate (Koso Chemical Co. Tokyo) was introduced into vessel (d) through the intake (d$_1$) covered with a ground cap in a hydrogen stream and 150 cc of water purified in (c) was distilled into (d) in a stream of hydrogen. Then the solution thus prepared was transferred into pre-electrolysis cell (f) by hydrogen pressed in through greaseless stop cock ($q_1$) purging hydrogen in (f) through another greaseless stop cock ($p_2$). The solution was finally purified by pre-electrolysis in (f) between two electrodes of platinum foil with 20~30 mA/cm$^2$ for 30~50 hours. The solution thus purified was transferred into

![Diagram of reaction apparatus](image-url)

**Fig. 2.** The reaction apparatus.

(1)-(11) hydrogen purification train, (a)-(c) distillation apparatus of water, (d) solution vessel, (f) pre-electrolysis cell, (g) cell including the test electrode, (h) solution storage vessel, (i) glass electrode for pH measurement, (d$_1$) ground glass cap, (e) purified SO$_3$ sealed in glass tube with breakable joints, ($p_1$)~($p_4$), ($q_0$)~($q_4$) greaseless valves for hydrogen path.
cell (g) containing the test electrode by hydrogen pressure. After each series of experiments in cell (g) the solution was transferred into vessel (h), a part of which being transferred into vessel (i) provided with a calomel and a glass electrode for the determination of pH of the solution.

2.2. Preparation of platinum film electrodes

The structure of the cell containing the test electrode is shown in Fig. 3(a), where (T), (R) and (A) are respectively the test, the reference and the counter electrodes. All electrodes are made of platinum films of ca. 1 μ thick which were prepared under vacuum of $10^{-6}$ mmHg by evaporating a platinum disk of purity 99.99% (JOHNSON MATTHEY) on a ground glass surface, heating the disk by means of an induction furnace at temperatures just below the melting point.

**Fig. 3.** Electrolytic cell (a), evaporation apparatus (b) and geometrical configuration of evaporated platinum film electrodes (c). (T), (R) and (A) test, reference and counter electrodes, (E) ground glass surface on which platinum is evaporated, (P) piece of platinum, (S₁) and (S₂) sample holder and cover made of quartz, (H) coil of induction furnace.
point of platinum, as shown in Fig. 3(b). Fig. 3(c) shows an example of the geometrical configuration of (T) and (R). Counter electrode (A) is 25 mm in diameter. The solution is stirred and circulated by hydrogen bubbles through glass filters (D).

Although platinum hydrogen electrodes thus prepared have high activities and rapid response, well reproducible results have been obtained in the galvanostatic transient measurements after repeated anodic polarizations for a few seconds at 1.8 v referred to the reversible hydrogen electrode.

2.3. The electric circuit in the galvanostatic transient methods

The electric circuit is shown in Fig. 4. A polarizing current is supplied to cell (g) from 24 v battery through high resistance (R₁) and its strength is determined by measuring the potential drop across fixed resistance (R₂).

![Fig. 4.](image)

Fig. 4. The electric circuit for galvanostatic transient methods. (R₁), (R₂) and (R₃) resistances for current control, current measurement and compensation of ohmic overvoltage, (E) storage battery, (S) proximity switch, (D) high gain differential preamplifier, and (M) memoscope oscilloscope.

For the galvanostatic experiments it is desirable to switch a polarizing current in a shortest possible time, e.g., within 1 μsec. This can be achieved by the use of a proximity switch (TATEISH DENKI Co. Kyoto). The current is switched on by contact a piece of iron (P) with proximity switch (S) or off by separating (P) from (S). An example of the current-time relations is shown
in Plate I. Test and reference electrodes (T) and (R) are connected respectively to terminals of a high gain differential preamplifier (D) attached to a oscilloscope (M) (HUGH's Aircraft). The potential difference between (T) and (R) is observed on the oscilloscope screen. The ohmic potential drop between (T) and (R) caused by the resistance of solution can be separately determined from the oscillograms as the initial potential jump on switching on or off. This ohmic drop was compensated for expedient observations with a potential drop of reverse sign through resistance (R₃) imposed on terminals of (D) on switching.

Plate 1. The current-time variation after switching on. The ordinate shows potential drop across R₂ (100 Ω), 2 mV/Division; the abscissa time after switching on, 4 μsec/Div.; solution 0.1 N Na₂SO₄, pH 8.3.

3. Experimental results

The overvoltage-time curves after switching on or off the polarizing current have been observed in aqueous sodium hydroxide and sodium sulfate of various concentrations. Examples of the oscillograms are shown in Plates 2~11. As seen from the Plates, the potential difference due to the ohmic resistance of solutions can be clearly discriminated from the change in the true hydrogen overvoltage. From these oscillograms the rate of change in the overvoltage \( \dot{\gamma} \) is calculated at a given time after switching on or off the polarizing current, and the values of \( \log (\mp i/\dot{\gamma}) \) are plotted against time in order to
Plate 2. The potential-time variation after switching on a cathodic polarizing current on Pt in 0.1 N NaOH\textsubscript{aq}, $i=6.1 \times 10^{-4} \text{ amp., } 1 \text{ mV/Div, } 20 \mu \text{s/Div.}$

Plate 3. The potential-time variation after switching on a cathodic polarizing current on Pt in 0.1 N NaOH\textsubscript{aq}, $i=4.6 \times 10^{-3} \text{ amp, } 2 \text{ mV/Div, } 10 \mu \text{s/Div.}$
Plate 4. The potential-time variation after switching on a cathodic polarizing current on Pt in 0.1 N NaOHaq,
\[ i = 4.6 \times 10^{-3} \text{ amp.}, 10 \text{ mv/Div, 100 \mu s/Div}. \]

Plate 5. The potential-time variation after switching on a cathodic polarizing current on Pt in 0.1 N NaOHaq,
\[ i = 8.4 \times 10^{-3} \text{ amp.}, 5 \text{ mv/Div, 10 \mu s/Div}. \]
Plate 6. The potential-time variation after switching on a cathodic polarizing current on Pt in 0.1 N NaOH\(_{aq}\), 
\(i = 8.4 \times 10^{-3}\) amp., 10 mv/Div, 100 \(\mu\)s/Div.

Plate 7. The potential-time variation after switching on an anodic polarizing current on Pt in 0.1 N NaOH\(_{aq}\), 
\(i = 6.1 \times 10^{-4}\) amp., 1 mv/Div, 20 \(\mu\)s/Div.
Plate 8. The potential-time variation after switching on an anodic polarizing current on Pt in 0.1 N NaOH$_{aq}$, $i=6.1 \times 10^{-4}$ amp., 1 mv/Div, 50 $\mu$s/Div.

Plate 9. The potential-time variation after switching off a cathodic polarizing current on Pt in 0.1 N NaOH$_{aq}$, $i=1.4 \times 10^{-3}$ amp., 2 mv/Div, 20 $\mu$s/Div.
Plate 10. The potential-time variation after switching off a cathodic polarizing current on Pt in 0.1 N NaOH$_{aq}$, $i=1.4 \times 10^{-3}$ 10 amp., 10 mv/Div., 100, 200, 500 $\mu$s/Div.

Plate 11. The potential-time variation after switching on a polarizing current on Pt in aqueous 0.1 N Na$_2$SO$_4$, pH 8.3, $i=1 \times 10^{-3}$ mp., 1 mv/Div, 10 $\mu$sec/Div.
Fig. 5. The relation between $\log\left(-i/\dot{i}\right)$ and time after switching on the cathodic polarizing current on an evaporated platinum film in NaOH$_{aq}$, pH 13.0; the apparent area 0.13 cm$^2$; the polarizing currents $i$ (amp): (○) 6.1·10$^{-4}$, (●) 1.19·10$^{-3}$, (△) 2.7·10$^{-3}$, (▲) 4.9·10$^{-3}$, (□) 8.4·10$^{-3}$; $C_D = 17.5 \mu$F, $\tau_1 = 92.4 \mu$sec.

Fig. 6. The relation between $\log(i/\dot{i})$ and time after switching off the cathodic polarizing current, the electrode and the solution are the same as in Fig. 6; the current $i$ and the starting overvoltage $\eta_s$: (○) 1.4·10$^{-3}$ amp. and $\eta_s = -50$ mv, (△) 8.9·10$^{-3}$ amp. and $\eta_s = -106$ mv; $C_D = 20.4 \mu$F and $\tau_1 = 87 \mu$sec.
Fig. 7. The relation between log \((i/\dot{i})\) and time after switching on the anodic polarizing current, the electrode and the solution are the same as in Fig. 6; the currents \(i\); (\(\bigcirc\) 1.19 x 10^{-3} \text{ amp.}, (\(\triangle\) 2.73 x 10^{-4} \text{ amp}; \(C_D=18.1 \mu \text{ F and } \tau_1=91.4 \mu \text{ sec.}

put Eqs. (7) and (13) to the test. Examples of the relation between log \((\mp i/\dot{i})\) and time are illustrated in Figs. 5~11. These Figures show the linear relation between log \((\mp i/\dot{i})\) and time every magnitude of polarizing current and every solution used in the present experiment valid at the initial stages of the build up and decay curves throughout. \(C_D\) and \(\tau_1\) evaluated from these Figures according to Eqs. (7) and (13) are now examined whether they satisfy the necessary conditions for being the differential capacities and the time constants of the discharge step of the hydrogen electrode reaction respectively; the necessary condition for \(C_D\) is its constancy for an electrode of constant surface area, and that for \(\tau_1\) is its independence of the electrode area, which are checked in the following sections in a) and b) respectively, and on which basis the rate of the discharge step is formulated in c).

a) The differential capacity of the electric double layer

The relations between log \((\mp i/\dot{i})\) and time are shown in Figs. 5, 6, 7 and 8(b) as derived from the overvoltage-time curves observed with an evaporated film of apparent area 0.13 cm^2 in NaOH \text{aq,} pH 13.0 and in 0.1 N Na_2SO_4, pH 11.0. The value of \(C_D\) is found to be constant at 18 \mu \text{ F independent of pH within experimental errors or of the direction of polarization, as seen from comparison of Fig. 5 and 7. The same value of } C_D \text{ is also obtained from}
A. MATSUDA and R. NOTOYA

decay curves starting from \(-50\) and \(-106\) mv, as seen from Fig. 6. Comparing Figs. 5 and 8(b), it is found that \(C_D\) remains constant as the electrolyte varies from NaOH to \(\text{Na}_2\text{SO}_4\). The constant value of \(C_D\) is thus secured from the log \((\mp i/\eta)\sim t\) relations throughout every solution and polarization examined on one and the same electrode.

Figs. 8(a) and 9 show the relation between log \((-i/\eta)\) and time in sodium sulfate observed with an evaporated film of apparent area \(0.07\,\text{cm}^2\). The value of \(C_D\) is found \(11\,\mu\text{F}\).

![Graph](image)

**Fig. 8.** The relation between log \((-i/\eta)\) and time in the build up processes on platinum in 0.1 N \(\text{Na}_2\text{SO}_4\). (a): apparent area \(0.07\,\text{cm}^2\), pH 8.3 and the currents \((\bigcirc)\ 4.7\times10^{-4}\,\text{amp}, (\triangle)\ 1.0\times10^{-3}\,\text{amp}; C_D=11\,\mu\text{F} \text{ and } \tau_1=90\,\mu\text{sec.}\) (b): apparent area \(0.13\,\text{cm}^2\), pH 11.0 and the currents \((\bullet)\ 5.25\times10^{-4}\,\text{amp}, (\bigtriangleup)\ 2.25\times10^{-4}\,\text{amp}; C_D=18\,\mu\text{F} \text{ and } \tau_1=84\,\mu\text{sec.}\)

The overvoltage-time curves were observed further on two platinum films of different area evaporated at the same time on different parts of one and the same ground glass surface, using a quartz cap with two openings in the evaporation apparatus in Fig. 3(b). The log \((-i/\eta)\sim t\) relations on these electrodes in \(\text{NaOH}_{aq}\) of pH 13.45 and 11.9 are shown respectively in Figs. 10 and 11. As seen from these Figures, a constant definite value of \(C_D\) is obtained on each electrode, while the values of \(C_D\) of the two electrodes are proportional to their apparent area. The proportionality of \(C_D\) to the apparent area for platinum film electrodes prepared under the same conditions of evaporation has been generally established, as shown in Fig. 12. It is concluded from these
Fig. 9. The relation between log \((-i/\gamma)\) and time in the build up processes on platinum in 0.01 N Na₂SO₄, pH 8.2, apparent area 0.07 cm², the currents (O) 2.3 \times 10^{-4} \text{ amp}, (△) 3.4 \times 10^{-4} \text{ amp}; C_D = 10 \mu \text{F} and \(\tau_1 = 246 \mu \text{sec}\).

Fig. 10. The relation between log \((-i/\gamma)\) and time in the build up processes on evaporated platinum films, apparent area 0.1 and 0.3 cm², in NaOHaq pH 13.45; the currents (O) 1.15 \times 10^{-3} \text{ amp}, (△) 2.25 \times 10^{-3} \text{ amp}, [□] 3.9 \times 10^{-5} \text{ amp}, (▲) 2.9 \times 10^{-3} \text{ amp}; C_D = 15 and 46 \mu \text{F} respectively and \(\tau_1 = 73 \mu \text{sec}\).
experimental results that $C_D$ determined from the linear relations between log ($\mp \dot{i}/\dot{\eta}$) and time satisfies the necessary condition mentioned in the foregoing section, of being the differential capacities of the electric double layers at the metal solution interfaces.

![Graph showing log ($\mp \dot{i}/\dot{\eta}$) vs. time](image)

Fig. 11. The relation between log ($\mp \dot{i}/\dot{\eta}$) and time in the build up processes in NaOH$_{aq}$, pH 11.9, the electrodes are the same as in Fig. 10; the currents (□) $2.23 \times 10^{-4}$ amp, (△) $4.06 \times 10^{-4}$ amp; [●] $C_D = 13$ and [▲] $44 \mu$F respectively and $\tau_1 = 326 \mu$sec.

b) The time constant of the discharge step

It is assumed that the overvoltage is that due to charge up of the double layer at the metal-solution interface in the overvoltage region where log ($\mp \dot{i}/\dot{\eta}$) varies linearly with time, inasmuch as $C_D$ obtained from the extrapolation of this linear part satisfies the condition of being the differential capacity of the double layer as described in a). The $\tau_1$'s obtained from these linear relations are then the time constants of the discharge step in accordance with Eq. (11) or (17). The values of $\tau_1$ thus determined are now commented upon in what follows.

First, Figs. 10 and 11 show that $\tau_1$ is independent of the electrode area, satisfying the condition that $\tau_1$ is the time constant of the discharge step. Figs. 5, 10 and 11 indicate on the other hand that $\tau_1$ obtained from the build up
curves in NaOH increases with decrease of pH. Comparison of Figs. 5 and 6 reveals that $\tau_1$ obtained from the decay curves coincides with that obtained from the build up curves in the same solution, i.e., $\tau_1$ is constant independent of the overvoltage. Figs. 8 and 9 show, on the other hand, that $\tau_1$ in Na$_2$SO$_4$ does not depend on pH, but on the concentration of sodium ion $C^{Na^+}$.

The $\tau_1$'s obtained in various solutions are tabulated in Table I and log $\tau_1$ is plotted against pH or log $C^{Na^+}$ in Fig. 13. As readily seen from the Figure, the log $\tau_1$ varies linearly with log $C^{Na^+}$ with gradient $-1/2$ independent of pH and irrespective as to whether $\tau_1$ is derived from the build up or the decay curves. It follows that the exchange current of the discharge step $i_{10}$ is proportional to the square root of $C^{Na^+}$,

$$i_{10} = k(C^{Na^+})^{1/2},$$

(18)

since $i_{10}$ is related to $\tau_1$ by Eqs. (8) and (10) as
Fig. 13. The relation between log $\tau_1$ and pH or log $C_{Na^+}$. (○) from build up curves in NaOH$_{aq}$; (△) or (●) from build up curves in aqueous Na$_2$SO$_4$ plotted against pH or log $C_{Na^+}$; (△) from decay curves in NaOH$_{aq}$.

**TABLE I.** Kinetic parameters of the discharge step of platinum hydrogen electrode reaction in alkaline solutions containing sodium ions

<table>
<thead>
<tr>
<th>Electrolyte</th>
<th>pH</th>
<th>$C_{Na^+}$ (N)</th>
<th>Build up</th>
<th>Decay</th>
<th>$-\eta_s$ (mv)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>$\tau$ (µ sec)</td>
<td>$i_{10}$ (amp/cm$^2$)</td>
<td>$\tau_1$ (µ sec)</td>
</tr>
<tr>
<td>NaOH</td>
<td>13.45</td>
<td>72.5</td>
<td>6.20 \times 10^{-3}</td>
<td>70</td>
<td>6.42 \times 10^{-3}</td>
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<td></td>
<td>13.00</td>
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<td>12.30</td>
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Note: the exchange current of the overall reaction $i_0=3.1 \times 10^{-4}$ amp/cm$^2$.

$$i_{10} = \frac{C_pRT}{F\tau_1}.$$  (19)

It may be concluded from these results that the discharge step of the hydrogen
Hydrogen Overvoltage on Platinum in Aqueous Sodium Hydroxide Part I.

electrode reaction in these solutions is not the discharge of a water molecule as usually accepted\textsuperscript{2}), but the discharge of a sodium ion. On the other hand, the exponent of $C_{Na}^{+}$ in Eq. (18) can be identified with the TAFEL constant of the discharge reaction on the basis of the FRUMKIN\textquoteright s theory\textsuperscript{5}, so that it may be valid to assume that the rate of the discharge step can be expressed as

$$i_{i} = k(C_{Na}^{+})^{1/2} \left\{ \exp \left( -F\eta_{i}/2RT \right) - \exp \left( F\eta_{i}/2RT \right) \right\},$$

where $\eta_{i}$ denotes $\Delta \varphi$ or the overvoltage of the discharge step.

c) The rate of the discharge step

The rate of the discharge step $i_{i}$ can be calculated by Eq. (6b) from $i$, $C_{D}$ and $\dot{\eta}$ relevant to the overvoltage curve. The relations between $i_{i}$ and $\eta$ thus calculated from the build up curves in aqueous NaOH of pH 13.0 and 13.3

Fig. 14. The relation between the current density of the discharge step and the overvoltage in NaOH\textsubscript{aq}, pH 13.0, in the build up processes; the polarizing current (○) $6.1 \times 10^{-4}$ amp/cm\textsuperscript{2} (△) $1.19 \times 10^{-3}$ amp/cm\textsuperscript{2} (□) $2.73 \times 10^{-3}$ amp/cm\textsuperscript{2} (●) $8.4 \times 10^{-3}$ amp/cm\textsuperscript{2}.
A. MATSUDA and R. NOTOYA

Fig. 15. The relation between log $i_1$ vs. $\eta$ in the build up processes, and log $i$ vs. $\eta$ in steady states in NaOH$_{aq}$, pH 13.0; the polarizing current (○) $1.2 \times 10^{-3}$ amp/cm$^2$, (△) $2.7 \times 10^{-3}$ amp/cm$^2$, (□) $4.8 \times 10^{-3}$ amp/cm$^2$, (●) $8.3 \times 10^{-3}$ amp/cm$^2$; (⊗) denotes log $i$ vs. $\eta$ in steady states.

The broken line shows log $i_1$ vs. $\eta$ calculated by the FRUMKIN's equation using $i_{10}$ obtained from $\tau_1$.

are respectively shown in Figs. 14, 15 and 16.

It is seen from Fig. 14 that $i_1$ is proportional to $\eta$ in the overvoltage region where $|\eta| \ll RT/F$. The proportionality constant $r_1 = -\eta/i_1$ is found $5\Omega \cdot$cm$^2$ which coincides with that calculated from $C_0$ and $\tau_1$ determined from log $(-i/\eta) \sim t$ relation in Figs. 5*). Fig. 15 and 16 show the log $i_1 \sim \eta$ relations in the whole region of the overvoltage in the build up processes. We see from these Figures that the discharge step nearly attains its steady state with very small value of $\eta$ or at the initial stage of the build up process, and that the log $i_1 \sim \eta$ curves coincide with the theoretical one calculated by Eq. (20)

*1 P. DOLIN, B. ERSHELR and A. FRUMKIN* have early obtained the value of $r_1 5\Omega \cdot$cm$^2$ in N NaOH$_{aq}$ by means of the conductivity measurements using alternating current at a frequency 375 c.p.s.*
using $i_{10}$ derived from $\tau_1$ and $C_B$ in the initial stages of the build up processes. These facts show that the increase of the overvoltage cannot be exclusively attributed to the increase of $\Delta \varphi$.

4. Discussions

As described in the introduction, the overvoltage is composed of $\Delta \varphi$ and $\Delta W$, i.e., the change in the potential difference across the double layer at the metal-solution interface and that in the contact potential between the electrode and the lead wire.

It has been experimentally found that there exists a linear relation between $\log (\mp i/\eta)$ and time in the initial stage of the build up and decay curves and $C_B$ determined from this linear relation on the basis of Eq. (11) or (17) is identifiable with the differential capacity of the electric double layer at the metal-solution interface. It can be concluded from these results that the increase of the overvoltage can be attributed exclusively to $\Delta \varphi$ in this initial stage. It has further been found that $C_B$ thus determined is constant independent of $\eta$ and the rate $i_i$ of the discharge step calculated by equation $i_i = i + C_B \eta$ on the basis of constancy of $C_B$ obeys the following equation in the initial stage within
time interval $\tau_1$ for every build up curve observed

$$i_i = i_{0} \left\{ \exp \left( -\Delta \phi / 2RT \right) - \exp \left( \Delta \phi / 2RT \right) \right\},$$

(20)

where $i_{0}$ is calculated from $\tau_1$ and $C_{D}$ by Eq. (19). This equation corresponds to FIRMKIN's equation simplified neglecting $\psi_{1}$-potential on the HELMHOLTZ plane in the double layer.

If the log $i_i$-$\eta$ curves coincide with the log $i$-$\eta$ in steady states throughout the build up of the overvoltage, the discharge step may be regarded as rate-determining. It has been found, however, that the log $i_i$-$\eta$ relation deviates definitely from log $i$-$\eta$ relation, as illustrated in Figs. 15 and 16, practically log $i_i$ being kept constant after the lapse of time interval $\tau_1$ and consequently the discharge step is excluded from being rate-determining.

We discuss here the increase of the overvoltage in the later stages of the build up curves after the lapse of time interval $\tau_1$. In this later stage the increase of the overvoltage cannot exclusively be attributed to $\Delta \phi$, since if it were so, $i_i$ should increase far beyond $i$ with increase of the overvoltage according to Eq. (20) contrary to the experimental results, as seen from Figs. 15 and 16. The discharge step can be regarded as nearly in steady state in this later stage, since $\tau_1$ measures the time in which the discharge step attains its steady state after switching on the polarizing current, as seen from Eq. (9). It is thus suggested that the charge density on the electrode surface is nearly kept constant and in consequence $\Delta \phi$ remains constant with increase of $\eta$ in this later stage, inasmuch as $C_{D}$ is constant independent of $\eta$ as stressed in the foregoing section. It is thus assumed that the increase of $\eta$ is attributed to $\Delta W$ in this later stage of the build up curves, which can be expected by the structural change of the electrode due to the formation of an intermetallic compound or a solid solution between platinum and sodium atom produced by the discharge step, as mentioned in the foregoing section.

The rate-determining step and the intermediates of the hydrogen electrode reaction will be discussed in the following papers.

**Conclusive remarks**

The platinum hydrogen electrode in aqueous NaOH and Na$_2$SO$_4$ has been studied by means of the galvanostatic transient methods.

It has been found that the discharge step is the discharge of sodium ion and its rate obeys FIRMKIN's equation simplified neglecting the $\psi_{1}$-potential on the HELMHOLTZ plane in the double layer and further suggested that the work function of the electrode is changed by the intermediate sodium atom produced by the discharge step.
Hydrogen Overvoltage on Platinum in Aqueous Sodium Hydroxide Part I.

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