2-step reaction kinetics for hydrogen absorption into bulk material via dissociative adsorption on the surface

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We have demonstrated that the process of hydrogen absorption into a solid experimentally follows a Langmuir-type (hyperbolic) function instead of Sieverts law. This can be explained by independent two theories. One is the well-known solubility theory which is the basis of Sieverts law. It explains that the amount of hydrogen absorption can be expressed as a Langmuir-type (hyperbolic) function of the square root of the hydrogen pressure. We have succeeded in drawing the same conclusion from the other theory. It is a 2-step reaction kinetics (2sRK) model that expresses absorption into the bulk via adsorption on the surface. The 2sRK model has an advantage to the solubility theory: Since it can describe the dynamic process, it can be used to discuss both the amount of hydrogen absorption and the absorption rate. Some phenomena with absorption via adsorption can be understood in a unified manner by the 2sRK model.

Hydrogen is the smallest element and a hydrogen ion H+ can be thought of as a point charge. If the size of neutral H0 is regarded as the Bohr radius, the size is about 0.05 nm. It is also known that hydrogen molecules are adsorbed dissociatively onto the surfaces of some kinds of materials and the hydrogen atoms then are absorbed into the bulk of them. The positive aspect of this phenomenon is that these materials can be used as storage materials1, hydrogen filters2, or sensors3,4; the negative aspect is that it may cause embrittlement of some materials and lead to their destruction. Sieverts’ law is the most well-known theory of hydrogen absorption into solids. It is obtained from solubility theory in thermodynamics, and is expressed as

\[ \theta = \frac{P_{H_2}}{P(T)} \]

where \( \theta \) is the ratio of the number of absorbed hydrogen atoms per the number of sites capable of absorption, \( P_{H_2} \) is the hydrogen gas pressure, and \( P(T) \) is a constant with a pressure dimension that depends on temperature \( T \). When the hydrogen concentration in the solid is low enough (\( \theta < 1 \)), the left side of Eq. (1) becomes \( \theta \), which is Sieverts’ law. Solving it for \( \theta \) without the approximation yields a Langmuir-type (hyperbolic) function. The Langmuir-type function appears in an ideal molecular adsorption on surface.

In our previous work, we demonstrated the detection of hydrogen by using a simple Pd film with a membrane-type stress sensor (MSS). The MSS is an optimized nanomechanical sensor which can measure mechanical deformation to nanometre precision and be used for observing various target analytes by selecting a sensitive film. We confirmed that the sensing signals of hydrogen absorption decreased with the number of absorption and desorption cycles because of the hysteresis behaviour of the cycles of them in the Pd–H system. By focusing on the hydrogen absorption rate (the time derivative of the absorption amount), we clarified experimentally that the absorption rate was fitted by a Langmuir-type function of the square root of the hydrogen concentration. Furthermore, the relation was proved theoretically as well with a 2-step reaction kinetics (2sRK) model.

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Considering the application as a sensor, short reaction and recovery times and reproducibility are important for hydrogen sensor. In particular, it is desirable that the reaction time is one second to a few ten seconds\(^3\). In the current study, we employed an amorphous Pd alloy, which has been studied as the sensing material in other types of hydrogen sensors owing to its low hysteresis and fast response\(^1\)\(^1\). With this alloy, we succeeded in demonstrating hydrogen sensing with low hysteresis and fast response. Our experimental results showed that the amount of the absorbed hydrogen was fitted by a Langmuir-type (hyperbolic) function of the square root of the hydrogen concentration instead of Sieverts’ law. We show in this paper that this relationship can be also explained by the 2sRK model we developed.

**Experimental results and discussion**

The principle of hydrogen detection using a hydrogen storage material and a stress sensor is that when the material stores hydrogen, the interatomic distance spreads and stress is generated, which is transmitted to the stress sensor and output. In this study, we used Pd\(_{75}\)Cu\(_{10}\)Si\(_{15}\) as a sensitive film for hydrogen sensing. The Pd\(_{75}\)Cu\(_{10}\)Si\(_{15}\) film was made by co-sputtering of the elements at room temperature onto the MSS. The thicknesses of the sensitive films on the MSS were set to 30 nm and 50 nm. For structural analysis of the film, we made a 50 nm film on a Si(100) substrate. The structure of the Pd\(_{75}\)Cu\(_{10}\)Si\(_{15}\) film was analysed by X-ray diffraction (XRD; Rigaku Smart Lab) at room temperature (Fig. 1). There were two peaks: a broad peak around 40 degrees from the amorphous structure and a sharp peak around 45 degrees from partial crystallization of Pd(200) or γ-PdCu\(^1\). In our hydrogen measurement system\(^7\), the hydrogen concentration is adjusted by mixing the hydrogen source gas with pure nitrogen gas (99.999%). We used two kinds of hydrogen source gas: 40 000 ppm (4.00%) ± 100 ppm (0.01%) and 100 ppm ± 10 ppm. Two mass-flow controllers (SEC-N112MGM; Horiba STEC) were used to adjust the hydrogen concentration flowing into the measurement chamber in which the MSS coated with the Pd\(_{75}\)Cu\(_{10}\)Si\(_{15}\) film was placed. Since the flowing gas was discharged without pumping, the total pressure of the mixture gas in the vicinity of the MSS was considered to be atmospheric pressure. The temperatures of the MSS measurement chamber and gas line were maintained at 298.0 ± 0.1 K. The stress signals were read as electrical voltages of the MSS with sampling times of 1 s (i.e., 1-Hz measurement).

Figure 2 shows the raw MSS signals for hydrogen concentrations of 2000, 4800, 10,000, 20,000, and 40,000 ppm. When hydrogen atoms are absorbed into the sensing film on the MSS, the induced mechanical stress of the film is transduced into an electrical signal via piezoresistors embedded in the narrow bridges that suspend the centre membrane\(^6\). Baselines reflect the internal stress of the Pd films and depend on the film thickness. Three cycles of 5 min hydrogen injection and 5 min nitrogen purge were executed for each concentration. The reaction time to saturation of 40,000 ppm was approximately 10 s, and the peak heights of the signals had good reproducibility. In the case of a MSS coated with a simple Pd film, it took 1 h or more to reach saturation and reproducibility was low\(^7\). It takes a little longer to saturate in the first absorption process for 2000 ppm in Fig. 2. The structure of the sensitive film should change slightly due to the first absorption and release process. Each release process is slower than each absorption one because the absorption process is an exothermic reaction and the release one is an endothermic reaction\(^1\). These results demonstrate the much faster hydrogen reaction in Pd\(_{75}\)Cu\(_{10}\)Si\(_{15}\) than in simple Pd and the good reproducibility for hydrogen absorption and release. We consider that the hydrogen diffusion was faster because hydrogen atoms can move easily along extended paths in the amorphous structure of the film.

Figure 3 shows the raw MSS signals for lower hydrogen concentrations of 12, 25, 50, and 100 ppm. Two cycles of 30 min hydrogen injection and 90 min nitrogen purge were executed for each concentration because

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**Figure 1.** XRD profile of Pd\(_{75}\)Cu\(_{10}\)Si\(_{15}\) thin film on Si(100). There are two peaks: a broad peak (■) around 40 degrees from the amorphous structure and a sharp peak (▼) around 45 degrees from partial crystallization of Pd(200) or γ-PdCu.
the hydrogen absorption and release required more time at low concentrations than at high concentrations. The peak heights of signals also showed good reproducibility at these concentrations.

From Figs. 2 and 3, we calculated a signal peak jump height (ΔV) for each injection-purge cycle and plotted ΔV against the square root of hydrogen concentration (C1/2) as shown in Fig. 4. The peak jumps are fitted by a Langmuir type (hyperbolic) function. In the inset of Fig. 4, ΔV exhibits a linear relationship to C1/2 at low concentrations with crossing about C1/2 = 2 (C = 4 ppm). It means that the baselines in Fig. 3 contains slight residual hydrogen, which is difficult to make the residual hydrogen completely 0 in cycles of hydrogen measurements, and the proportional to C1/2 means Sieverts' law. There is no film thickness dependence at low concentrations, due to the difficulty to eliminate residual hydrogen. The theoretical interpretation is given in the next section.

Theoretical model
This relationship can be explained by either of two different theories. One is the solubility theory (Eq. (1)); the other is the 2sRK model. With the 2sRK model, it is considered that there is dissociative adsorption of hydrogen onto surface site S and absorption into bulk site B. The reaction kinetics can be written as follows:

\[ S + \frac{1}{2} H_2 \xrightleftharpoons[k_{1-}]{k_{1+}} S-H \xrightleftharpoons[k_{2-}]{k_{2+}} B-H \]  (2)
where $S$–$H$ corresponds to the status of an adsorbed hydrogen atom on the surface site, $B$–$H$ corresponds to the status of an absorbed hydrogen atom in the bulk site, and $k_i$ ($i = +1, -1, +2, -2$) corresponds to the reaction-rate constants. The reaction kinetics are expressed as follows:

$$\frac{d}{dt}[S–H] = k_1 + [S][H_2]^\frac{1}{2} - k_1–[S–H] - k_2+[S–H][B] + k_2–[S][B–H]$$  \hspace{1cm} (3)$$

and

$$\frac{d}{dt}[B–H] = k_2+[S–H][B] - k_2–[S][B–H]$$  \hspace{1cm} (4)$$

where $[S–H]$ and $[S]$ are the numbers of occupied and empty sites on the surface, and $[B–H]$ and $[B]$ are the numbers of occupied and empty sites in the bulk, respectively. We can define a dimensionless $[H_2] = \frac{P_{H_2}}{P_a}$, where $P_{H_2}$ is the partial pressure of hydrogen and $P_a$ is atmospheric pressure. From this definition, $[H_2] = 1$ corresponds to a concentration of hydrogen in a gas mixture of 100%. Let the maximum number of the adsorbable sites on the surface be $S_0$ and the maximum number of the absorbable sites in the bulk be $B_0$. Then,

$$[S] + [S–H] = S_0.$$  \hspace{1cm} (5)$$

and

$$[B] + [B–H] = B_0.$$  \hspace{1cm} (6)$$

When we assume an equilibrium state in Eqs. (3) and (4), the following equation holds:

$$\frac{d}{dt}[S–H] = \frac{d}{dt}[B–H] = 0.$$  \hspace{1cm} (7)$$

From Eqs. (3), (4), and (7),

$$[S–H] = \frac{S_0K_1[H_2]^{1/2}}{1 + K_1[H_2]^{1/2}}$$  \hspace{1cm} (8)$$

and

$$[B–H] = \frac{B_0K_1K_2[H_2]^{1/2}}{1 + K_1K_2[H_2]^{1/2}}$$  \hspace{1cm} (9)$$

where $K_1 = \frac{k_1+}{k_1–}$ and $K_2 = \frac{k_2+}{k_2–}$. The surface adsorption $[S–H]$ obeys the well-known Langmuir’s law. Furthermore, the bulk absorption $[B–H]$ is also expressed as a Langmuir-type function with $K_1$ and $K_2$. Thus, we have simply and clearly explained that the hydrogen behaviour into bulk follows a Langmuir-type function, which is same conclusion of the solubility theory (Eq. (1)).

The MSS output signal voltage corresponding to the induced stress of hydrogen absorption, it can be expressed by the following formula:

![Graph](image-url)

**Figure 4.** Peak jump heights of MSS output voltage ($\Delta V$) plotted against square root of hydrogen concentration ($C^{1/2}$). Inset: Expansion at low concentrations.
where $V_v$ is an initial voltage depending on the residual stress of the sensitive film. The Langmuir-type dependence of hydrogen absorption shown in Fig. 4 is clarified.

We can also discuss Sieverts’ law in terms of this 2sRK model, as has already been proved by Davenport, Dienes and coworkers. In the case of $[B−H] < < B_0$, Eq. (6) can be approximated as:

$$[B] \approx B_0.$$  \hfill (11)

In this case, from Eqs. (3), (4), and (7), the following is obtained instead of Eq. (9),

$$[B−H] \approx B_0 K_1 K_2 [H_2]^{1/2}.$$  \hfill (12)

It can be also obtained directly from Eq. (9) in the case where the concentration of hydrogen is low enough (i.e., $[H_2] < < 1$).

A case that cannot be described by the solubility theory can be described by the 2sRK model. In previous work, we were able to evaluate the initial absorption rate as a function of concentration with a simple Pd film as a sensitive film. The theoretical essence is outlined below. In the case that Reaction 1 is much faster than Reaction 2 (i.e., $k_1, k_1 > > k_2, k_2$), the $k_1$ and $k_2$ in Eq. (3) can be ignored, thereby being simplified without $k_2, k_2$ as

$$\frac{d}{dt}[S−H] \approx k_1+[S][H_2]^{1/2}−k_1−[S−H].$$  \hfill (13)

Then the differential Eq. (13) can be solved analytically with Eq. (5) and a boundary condition $[S−H] = 0$ at $t = 0$:

$$[S−H] \approx \frac{S_0 K_1 [H_2]}{1 + K_1 [H_2]} \left[ 1 − \exp\left(−(k_1+[H_2]^{1/2}+k_1−)t \right) \right].$$  \hfill (14)

Since $k_1$ and $k_1$ are large enough and the exponential function in Eq. (14) rapidly approaches 0, $[S−H]$ can be regarded as Langmuir adsorption. If we consider the initial process of absorption into the bulk (i.e., $[B−H] \approx 0$), then Eq. (4) becomes,

$$\frac{d}{dt}[B−H] \approx k_2+[S−H][B] = \frac{S_0 B_0 K_1 K_2 [H_2]^{1/2}}{1 + K_1 [H_2]}.$$  \hfill (15)

This equation indicates that when Langmuir adsorption takes place quickly on the surface, the absorption rate at the initial process is proportional to the hydrogen density on the surface. As a result, a Langmuir-type function appears in the absorption rate.

**Summary and conclusion**

We showed experimentally that the process of hydrogen absorption into an amorphous Pd alloy film reaches saturation very quickly with little hysteresis. It was explained by each of the solubility theory and the 2sRK theory that the amount of absorbed hydrogen is a Langmuir-type (hyperbolic) function of the square root of the hydrogen partial pressure (or hydrogen concentration) instead of Sieverts’ law. The reason why the two theories draw the same conclusion is that they do not suppose the common assumption that the number of absorbable hydrogen partial pressure (or hydrogen concentration) instead of Sieverts' law. The reason why the two theories can be regarded as Langmuir adsorption. If we consider the initial process of absorption into the bulk (i.e., $[B−H] \approx 0$), then Eq. (4) becomes,

$$\frac{d}{dt}[B−H] \approx k_2+[S−H][B] = \frac{S_0 B_0 K_1 K_2 [H_2]^{1/2}}{1 + K_1 [H_2]}.$$  \hfill (15)

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Author contributions
T.Y. wrote the original manuscript and constructed the theoretical model. T.Y. and G.I. conducted the experiments. G.I. and G.Y. constructed the MSS measurement system and the software. T.Y., N.M., M.K., and A.N.I discussed mainly for hydrogen science. T.Y., G.I., and G.Y. discussed mainly for MSS detection. All authors discussed the results, provided good feedback and contributed to the final version of the manuscript.

Competing interests
The authors declare no competing interests.

Additional information
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