Theory of the Kinetics of Chemical Potentials in Heterogeneous Catalysis


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Catalysis is of paramount importance in our daily life. In particular, heterogeneous catalysts which speed up reactions on their solid surfaces play a vital role in a wide range of industries, such as petroleum, energy, and environment-related industries. There is no doubt that rational design of new catalysts is a major endeavor in chemistry. A key to achieve this goal is reaction kinetics, bridging the gap between microscopic elementary chemical reactions and macroscopic performance of catalysts. Although great progress in understanding reaction kinetics has been made, rational design of new catalysts remains one of the profound challenges. Here, we present a new formulism of reaction kinetics at surfaces in terms of the involved chemical potentials, which simplifies the reaction kinetics significantly. Furthermore, within this formulism we propose a new approach of searching for new catalysts. The effectiveness and universality of this theory are discussed.

There have been several major related developments in this field. First, density functional theory (DFT) approaches have been developed to such a level that the barriers of elementary steps are determined routinely. Many total-energy profiles of reaction systems from DFT calculations were reported. Second, with the energy profiles from first-principles calculations in hand, kinetic information, such as reaction rates and coverages of surface intermediates, are obtained by kinetic Monte Carlo simulations[1, 2] and microkinetic calculations.[3] Third, the linear relationship between the adsorption paradigm [Eq. (1); see the Supporting Information for the derivation][14–16]

$$\mu_i(T, \theta_i) = \mu_{i}^{0}(T) + RT \ln \frac{\theta_i}{\theta_{i}^{0}} = E^{*i} + \Delta H_i(T) + RT \ln \frac{\theta_i}{\theta_{i}^{0}}$$

(1)

where \(\theta_i\) and \(\theta_{i}^{0}\) are coverages of adsorbed species \(i\) on a surface, respectively, and \(\mu_{i}^{0}(T)\) is the standard chemical potential of species \(i\) at temperature \(T\) and is readily obtained from the total energy \(E^{*i}\); routinely computed using DFT, at 0 K with a small thermal correction term \(\Delta H_i(T)\).

Based on Equation (1), many expressions for the microkinetics, for example, the reaction rate and reversibility, can be reformulated by using only chemical potentials. The formulations for some typical elementary surface reactions are given in the Supporting Information. To illustrate our method, we apply a simple two-step kinetic model consisting of adsorption [Eq. (2)] and desorption [Eq. (3)] processes, which captures the essence of many heterogeneous catalytic reactions[9] where \(R\) and \(P\) are reactant and product in the gas phase, and \(I\) and \(*\) are surface intermediate and free site, respectively.

<table>
<thead>
<tr>
<th>Reaction</th>
<th>Equations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Adsorption</td>
<td>(R(g) + * \rightleftharpoons I^*)</td>
</tr>
<tr>
<td>Desorption</td>
<td>(I^* \rightleftharpoons P(g) + *)</td>
</tr>
</tbody>
</table>

[1] Dr. J. Cheng, Prof. Dr. P. Hu
School of Chemistry and Chemical Engineering
The Queen’s University of Belfast, Belfast BT9 5AG (UK)
E-mail: jcs90@cam.ac.uk
p.hu@qub.ac.uk
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[4] The ease of adsorption from the total-energy profiles obtained consequently is not straightforward, let alone rational design of new catalysts. As a result, new catalysts are traditionally developed using experimental trial-and-error methods. Therefore, to have better approaches, in particular better kinetic theories directed towards the design of catalysts, is essential to further develop the subject. Here, we introduce chemical potentials, which are widely used in electrochemistry under the name of electrochemical potential, to surface-catalytic reactions to reformulate the reaction kinetics. It will be manifested below that the chemical potentials of surface intermediates implicitly take into account surface coverages and temperature effects, and therefore reveal more chemical meanings. More importantly, we show that chemical potentials of surface intermediates can be used as a guide in searching for new catalysts without detailed kinetic analyses, and the interesting observation of the universal energy window can be understood using our approach.

The basis of our approach is the expression of the chemical potential of a surface species in the Langmuir adsorption paradigm [Eq. (1); see the Supporting Information for the derivation][14–16]
Following our formulation, we can rewrite the kinetic equations of reactions (2) and (3) as Equations (4)–(7), where $r_{\text{ads}}$ and $r_{\text{des}}$ are the reaction rates of the adsorption and desorption processes, respectively, $z_{\text{ads}}$ and $z_{\text{des}}$ are the reversibilities of the adsorption and desorption processes,[17] respectively, $\mu_R$, $\mu_I$, and $\mu_P$ are the chemical potentials of the reactant, the surface intermediate, and the product, respectively, and $\mu_R^{\circ\circ}$ and $\mu_I^{\circ\circ}$ are the standard chemical potentials of the transition states (TS) of adsorption and desorption, respectively. At the steady state, $r = r_{\text{ads}} = r_{\text{des}}$ where $r$ is the overall reaction rate.

\[
r_{\text{ads}} = \frac{k_B T}{h} \exp\left(\frac{\mu_R - \mu_R^{\circ\circ}}{RT}\right) \theta_s (1 - z_{\text{ads}}) \\
z_{\text{ads}} = \exp\left(\frac{\mu_R - \mu_R^{\circ\circ}}{RT}\right) \\
r_{\text{des}} = \frac{k_B T}{h} \exp\left(\frac{\mu_I - \mu_I^{\circ\circ}}{RT}\right) \theta_s (1 - z_{\text{des}}) \\
z_{\text{des}} = \exp\left(\frac{\mu_I - \mu_I^{\circ\circ}}{RT}\right)
\]

From Equations (4)–(7), we obtain Equation (8), where $\mu_R^{\circ\circ}$ and $\mu_I^{\circ\circ}$ are, according to the BEP relation, linearly related to $\mu_1^*$, which is an intrinsic property of the catalyst surface[8–10] and independent of reaction conditions. Thus, solving Equation (8) with the condition of conservation of the surface site $\theta_1 + \theta_2 = 1$, we calculate $\theta_1$ and $\theta_2$, as well as the overall reaction rate. If treating $\mu_1^*$ (equivalent to the adsorption energy) as a variable, the overall reaction rate will be a function of adsorption energy, giving rise to a typical volcano curve.[8–10]

\[
r = \frac{k_B T}{h} \exp\left(\frac{\mu_R - \mu_R^{\circ\circ}}{RT}\right) \theta_s \left[1 - \exp\left(\frac{\mu_I - \mu_I^{\circ\circ}}{RT}\right)\right] \\
= \frac{k_B T}{h} \exp\left(\frac{\mu_I - \mu_I^{\circ\circ}}{RT}\right) \theta_s \left[1 - \exp\left(\frac{\mu_P - \mu_P^{\circ\circ}}{RT}\right)\right]
\]

What can we learn from the kinetics of chemical potentials compared to traditional kinetic equations for catalytic reactions? First, the equations shown above contain some useful chemical insights. By plotting the reaction profiles of the chemical potentials (the gray curve in Figure 1), we obtain a deeper understanding of surface reactions compared to the traditional reaction profiles currently used, in which only total energies are considered without thermal correction and concentration terms (the black curve in Figure 1). Useful kinetic information can hardly be obtained from the conventional reaction profiles of total energies by DFT calculations, whereas in the profile of chemical potentials the reactants, intermediates, and products along the reaction coordinate must decrease step by step [Eq. (9)].

\[\mu_R > \mu_I > \mu_P\]  

This equation can be generalized to any sequential multistep reaction system, namely $\mu_{i+1} > \mu_i > \mu_{i+2} > \ldots > \mu_P$. Its significance will be revealed later. Second, the magnitude of the decrease of each step, which is related to the reversibility according to Equations (5) and (7), indicates the thermodynamic driving force for the step. Third, the heights of the TSs with respect to the reactant states (the barriers of the chemical potentials) are direct measures of the reaction rates, unlike the barriers of total energies in which the entropic effects and coverages are not taken into account. This is best manifested by the fact that for adsorption processes the sole use of the barriers of total energies as measures will significantly overestimate the reaction rates because of the lack of large negative entropic effects.

There are some significant implications in the above results for understanding heterogeneous catalysis. First, if the standard chemical potential of a TS of a prior step is smaller than the later step, that is, $\mu_{i+1} < \mu_{i+2}$, then $\mu_R \approx \mu_I$ is approximately satisfied (see the Supporting Information), indicating that the former step reaches a quasi-equilibrium at steady states. This result can be readily extended to other sequential elementary reactions, such as the multistep hydrogenation reaction $\text{C}_4 + 4\text{H} \rightarrow \text{CH}_4$ in the CO hydrogenation on metal surfaces.[18, 19] For a series of sequential reactions if the last step has the highest chemical potential at the TS, the previous steps can be approximated treated as being in quasi-equilibrium, and the last step is the rate-determining. Second, for a given reaction condition and a catalyst surface, the levels
of the chemical potentials of reactant and product ($\mu_R$ and $\mu_P$) and TSs ($\mu_R^0$ and $\mu_P^0$) are fixed in a reaction profile of chemical potentials. However, the chemical potential of the surface intermediate ($\mu_I$) consists of two terms, $\mu_I^0$ and $R \ln(\theta_1/\theta_0)$, where $\mu_I^0$ is invariant for a given catalyst and $R \ln(\theta_1/\theta_0)$ is the coverage-dependent term. Namely, $\mu_I$ is varied around the $\mu_I^0$ by $\theta_1$, considering that $\theta_1$ is a variable whereas $\theta_0$ is not an independent variable because of $\theta_1 + \theta_0 = 1$ in our two-step model (but generally, $\theta_1 + \theta_0 \approx 1$ if there are more than one intermediate and intermediate I is the main one). Upon approaching a steady state, however, the level of $\mu_I$ must reside between $\mu_R$ and $\mu_P$ to satisfy Equation (8).

Now we are in a position to illustrate an important application of our chemical potential kinetics theory for searching for new catalysts. In the Supporting Information, using a simple kinetic model we show that the coverage of free sites $\theta$ on the surface can be derived to be around $10^{-1}$ monolayer (ML) for the optimal catalysts, which is supported by experimental results: When catalytic reactions take place on good catalysts, $\theta$ is usually in the order of magnitude of $10^{-1}$ to $10^{-2}$ ML at steady states. For example, kinetic analyses showed that $\theta$ is around 0.08 ML for hydrogenation of isobutene on Pt,[17] and around 0.01 ML for the synthesis of ammonia on Fe and Ru catalysts.[7,17] This is also consistent with a general consensus in the field: 1) If $\theta$ is low, it is usually a sign of blockage of surface sites, leading to low activities. This often happens when the surface–adsorbate bonding interaction is too strong. 2) If $\theta$ is high and approaches 1 ML, it is difficult for molecules to adsorb on the surface. Namely, the surface is too inert to catalyse the reaction.

Since $\theta$ is about $10^{-1}$ to $10^{-2}$ ML on the best catalysts and $\theta_1$ and $\theta_0$ are related to each other because of the conservation of surface sites, the magnitude of the coverage-dependent term $R \ln(\theta_1/\theta_0)$ in Equation (5) has to lie in a small range, about 0.1–0.2 eV at 500 K, and is defined as $\epsilon$. In other words, for good catalysts [Eq. (10)].

$$\mu_I^0 + \epsilon > \mu_I > \mu_I^0 - \epsilon \text{ or } \mu_I \approx \mu_I^0$$

Combining Equations (9) and (10), we reach the key relation [Eq. (11)] for searching for good catalysts.

$$\mu_R + \epsilon > \mu_I > \mu_P - \epsilon \text{ or approximately } \mu_R > \mu_I^0 > \mu_P$$

In principle the coverage-dependent term $R \ln(\theta_1/\theta_0)$ varies in the range from $-\infty$ to $+\infty$ and approaches the infinity limits when the surface coverage is extremely low (no adsorption) and high (1 ML). Thus, $\mu_I$ can be changed by $\theta_1$ to locate anywhere in the diagrams of chemical potentials no matter where $\mu_I^0$ is. However, upon approaching the two limits the total reaction rates will be reduced dramatically because of the inertness of the surface or blockage of surface sites. For good catalysts $R \ln(\theta_1/\theta_0)$ should be small, and hence $\mu_I$ is mainly determined by $\mu_I^0$. Equation (11) may partially justify the assumption of a downhill requirement for the free-energy diagrams in the model for electrocatalytic reactions reported by Nørskov, Rossmeisl, and co-workers.[20]

Then, the searching procedure of catalysts, as illustrated in Figure 2, can be devised: First, based on the gas-phase energetics of an overall reaction under a certain reaction condition (i.e., temperature and pressure) which can be very easily obtained, determine the positions of the chemical potentials of the reactants and products in the diagram of chemical potentials, which approximately establishes the upper and lower boundaries of the standard chemical potentials of the key intermediates on the surface (zone 1 in Figure 2). Second, according to Equation (11), we slightly relax the boundaries (e.g., by 0.2 eV) and search for the appropriate catalyst surfaces which are able to offer the standard chemical potentials of the surface intermediates lying between the relaxed boundaries (zone 2 in Figure 2). As can be seen from Figure 2, $\mu_I^0$ and $\mu_P^0$ are very likely to be good catalysts, whereas surfaces related to $\mu_R^0$ and $\mu_P^0$ cannot be good catalysts.

![Figure 2](image-url)
Here, we use the ammonia synthesis as an example to elaborate our method. According to Equation (9), we can obtain the boundaries for the chemical potential of adsorbed N [Eq. (12), see the Supporting Information for details].

$$\frac{1}{2} \mu_{N_2} > \mu_N > \mu_{N_2} + \frac{3}{2} \mu_H$$  (12)

Substituting Equation (10) and applying experimental gas-phase energetic data under typical reaction conditions (H2 75 bar, N2 25 bar, NH3 1 bar, 673 K), we can rewrite Equation (12) into Equation (13), where $\Delta E_{N_2}$ is the disso- 

$$-1.3 + \epsilon > \Delta E_{N_2} + 2 \mu_{N_2} > -1.8 - \epsilon$$  (13)

ciative adsorption energy of N2.

Correcting the small term $\Delta T \mu_{N_2}$ as well as the zero-point energy (ZPE),[23] we have Equation (14).

$$-1 + \epsilon > \Delta E_{N_2} > -1.5 - \epsilon$$  (14)

Taking $\epsilon$ as 0.2 eV as suggested above, we will have adsorption energies of N2 in the range of around $-1.7$ to $-0.8$ eV for optimal catalysts, which is in good agreement with the observed energy window (around $-2$ to $-1$ eV).[9,13] This agreement is extraordinary: In predicting the optimal range of the adsorption energy, we only use the data from gas-phase reaction energetics, without recourse to detailed DFT calculations on surfaces except the small corrections of the adsorption energies of N2 in the range of around $0.6$ to $1.0$ eV at a typical temperature range of 300–500 K.

In summary, here we have applied chemical potentials to catalytic reactions on surfaces; reaction kinetics of surface processes, for example, the reaction rate and reversibility, has been reformulated in terms of the involved chemical potentials. The total energy profiles of surface reactions usually computed from DFT simulations can be readily converted to profiles of chemical potentials. The new formulation is simple but powerful to understand surface reactions both thermodynamically and kinetically. We have estimated with approximations that for many catalytic reactions the coverage of free site on surfaces of the best catalysts is usually in a medium range. Combining this simple, but important result and our formulation, a procedure in searching for good catalyst has been proposed. Our formulation has also provided the explanation of the universality of the adsorption-energy window in heterogeneous catalysis. We have further shown our method by using the ammonia synthesis as an example. Our method is not only able to predict the range of optimal adsorption energies, in agreement with reported values, but it also is very simple without the need of extensive calculations of reaction barriers and detailed kinetic analyses.

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[21] We use the vibration frequencies of N adsorbed on Ru(0001): 581, 581, and 556 cm$^{-1}$ (S. P. Liu, C. Hao, S. M. Li, Z. X. Wang, *Appl. Surf. Sci.* **2009**, *255*, 4232). Then, the entropy of adsorbed N is calculated to be 21.4 J mol$^{-1}$ K$^{-1}$ using the harmonic approximation, and the ZPE is 0.11 eV. The gaseous N$_2$ has a stretched vibrational mode of 2744 cm$^{-1}$, leading to a ZPE of 0.17 eV.