Elementary kinetic modeling and experimental validation of electrochemical CO oxidation on Ni/YSZ pattern anodes

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1. Introduction

Due to their high energy conversion efficiency and low pollutant emission level, solid oxide fuel cells (SOFC) are a promising electrical power source for a variety of applications, ranging from mobile technology to stationary power plants [1]. At the high operating temperatures of SOFCs (typically around 1100 K) a variety of fuels, such as H2, CO, H2/CO-rich reformate gases, and hydrocarbons, can be utilized [2]. However, although being widely investigated, the microscopic details of the electrochemical reaction mechanism of the SOFC anode, occurring at the three-phase boundary (TPB) of gas-phase, electrode, and the yttria-stabilized zirconia (YSZ) electrolyte, are not yet fully resolved [3]. This is true even for apparently simple chemical systems such as the electrochemical H2 or CO oxidation reactions. Yet, the knowledge of elementary kinetics of SOFC electrochemistry is important because the understanding at the fundamental level yields better predictive capability, which allows optimal design and operation of new fuel cell systems.

There have been only a few results published on Ni/YSZ SOFC anodes operating on CO/CO2 mixtures. Moreover, these studies so far did not lead to a consistent picture of the electrochemical CO oxidation mechanism. There are both qualitative (based on electrochemical measurements) and quantitative (based on kinetic modeling) mechanistic interpretations, both of which will be briefly reviewed in the following.

Holtappels et al. [4] investigated the CO electrochemical oxidation on Ni/YSZ cermet anodes. They showed that the rate of CO oxidation is lower than the rate of H2 oxidation by about one order of magnitude, which was attributed to a difference in both, mass-transfer and charge-transfer resistance. They also observed that CO oxidation is highly dynamical. These observations were attributed to a fundamental difference in the mechanisms of H2 and CO oxidation. Boulenouar et al. [5] studied CO oxidation on a Ni grid electrode using electrochemical impedance spectroscopy. The experiments showed at least two relaxation times for high pCO2/pCO ratios and a single relaxation time at low pCO2/pCO ratios. Based upon equivalent circuit analysis it was concluded that CO, O and CO2 participate in the reaction mechanism, where two species are adsorbed on a surface. Sukeshini et al. [6] carried out a series of CO oxidation measurements for Ni/YSZ pattern anodes. It was concluded that at low overpotentials, the rate-controlling process may likely not depend on temperature and may be CO...
adsorption/desorption. However, at high overpotentials a similar mechanism as for H₂ oxidation may become rate-controlling.

Although the results discussed so far are of great importance as qualitative observations, they do not provide sufficient insight for a quantitative analysis. Detailed quantitative studies were performed by Lauvstad et al. [7,8] who conducted electrochemical steady-state polarization and impedance measurements for point-shaped Ni electrodes. In a theoretical analysis, different mechanisms of CO oxidation were compared. Based upon impedance analysis, a mechanism was identified in which CO adsorbed on Ni reacts with oxygen atoms adsorbed on Ni to produce gaseous CO₂. This mechanism is very similar to the one proposed in a pioneer study by Ytset and Flengas [9] for the Pt/YSZ system. Ytset et al. concluded that either gaseous or adsorbed CO and gaseous CO₂ are directly involved in the electrochemical step of the reaction. Moyer et al. [10] performed combined experimental and modeling studies for reversible solid-oxide cells with Ni/YSZ cermet anodes operated on H₂/CO/H₂O/CO₂ gas mixtures. Using elementary kinetic models, they investigated the simultaneous presence of hydrogen and oxygen spillover mechanisms. The mechanism of CO oxidation included CO and CO₂ adsorption/desorption on Ni, as well as CO dissociation and oxidation via a Langmuir–Hinshelwood type surface reaction on Ni. An interaction of CO with the YSZ surface was not included in this mechanism.

In the case of the electrochemical H₂ oxidation, we have in the past carried out combined experimental and theoretical studies using micro-patterned model anodes, which have considerably increased our understanding of this system [3,11,12]. Applying a similar methodology to CO oxidation, we have recently developed an elementary kinetic reaction mechanism for the electrochemical CO oxidation [13]. The mechanism was validated against experimental electrochemical data (steady-state polarization curves and electrochemical impedance spectra) measured by Lauvstad et al. [7] for a Ni-point/YSZ model anode under different CO/CO₂ atmospheres. In Ref. [13], a thermodynamically consistent kinetics data set could be derived for the first time, which allows for a quantitative description of CO adsorption/desorption, and heterogeneous CO oxidation on both the Ni and YSZ surfaces, as well as the charge-transfer reaction kinetics taking place at the TPB. However, as the electrochemical characterization experiments of Lauvstad et al. [7] were performed for a single operating temperature (T = 1149 K) only, no information about the temperature dependence of the rate constants of the charge-transfer reactions could be obtained so far.

In the present work, our previously developed elementary reaction mechanism [13] is applied to analyze recent experiments of Utz et al. [14], which were conducted using well-defined and thoroughly characterized Ni/YSZ patterned model anodes. These experiments stand for the most complete data set available at present and cover an extended range of operating temperatures (973 K ≤ T ≤ 1073 K) and CO/CO₂ gas compositions (4.0 × 10⁻³ Pa ≤ pCO ≤ 5.1 × 10⁻² Pa and 9.5 × 10⁻³ Pa ≤ pCO₂ ≤ 9.2 × 10⁻² Pa). The experimentally observed electrochemical behavior is interpreted in terms of microscopic heterogeneous reaction pathways on both Ni and YSZ and charge transfer steps taking place at the TPB.

2. Theoretical and experimental methodology

2.1. Modeling and simulation

Numerical simulations of the CO electrochemical oxidation were performed using the software package DENIS [12,15,16], which is based on (i) an elementary kinetic approach for the thermodynamically consistent description of electrochemistry and heterogeneous surface chemistry, (ii) charge transfer reactions described as surface spillover reactions at the TPB, and (iii) surface diffusion on the Ni electrode and YSZ electrolyte surfaces described in one dimensions (1D) parallel and perpendicular to the TPB line. The model reproduces one repeat unit of the pattern anode, as shown schematically in Fig. 1. The model equations are given in detail in Ref. [12]. Experimental electrochemical impedance spectra were simulated using a potential step and current relaxation technique [15]. The impedance is obtained in the frequency domain by a Fourier transformation of the resulting time-domain traces of current and potential. The differential-algebraic equation (DAE) system solver LIMEX [17] is used for the numerical integration of the reaction–diffusion equations. Chemical source terms are calculated using the DETCHM software package [18].

The modeling framework has been used in our previous theoretical study of CO oxidation at a Ni/YSZ point model anode [13]. We use the reaction kinetics and thermodynamic datasets of that study (Tables 1 and 2 of Ref. [13]) as parameter base for the present work.

2.2. Experimental

The experimental setup and methodology has been described in detail elsewhere [14]. In the experiments, Ni/YSZ pattern anodes with an area of 10.25 × 10.25 mm² were used which were fabricated on 8.5 mol% polycrystalline Y₂O₃-stabilized ZrO₂ (YSZ) substrates. The anode consists of parallel Ni stripes with 25 μm width, a spacing of 160 μm and a thickness of 800 nm. An additional Ni frame with 500 μm width ensures contacting of all stripes resulting in an actual TPB length of 1.3 cm⁻² in the experiments, as determined by scanning electron microscopy (SEM) measurements. The geometric features of the Ni/YSZ pattern anode used in the electrochemical characterization studies are summarized in Table 1. As counter electrode, a screen-printed Ni/YSZ cermet anode was employed, the preparation of which has been described previously [19]. The counter electrode was applied to the substrate prior to Ni anode patterning.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Symbol</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thickness of electrolyte</td>
<td>d_{YSZ}</td>
<td>200 μm</td>
</tr>
<tr>
<td>Thickness of Ni stripe</td>
<td>d_{Ni}</td>
<td>0.8 μm</td>
</tr>
<tr>
<td>Width of YSZ area</td>
<td>w_{YSZ}</td>
<td>160 μm</td>
</tr>
<tr>
<td>Width of Ni stripe</td>
<td>w_{Ni}</td>
<td>25 μm</td>
</tr>
<tr>
<td>Length of TPB</td>
<td>h_{TPB}</td>
<td>1.3 cm⁻²</td>
</tr>
<tr>
<td>Electrode area</td>
<td>A</td>
<td>1 cm²</td>
</tr>
</tbody>
</table>

Electrochemical impedance spectra were recorded for different CO/CO\textsubscript{2}/N\textsubscript{2} gas mixtures at open circuit over a frequency range of 100 mHz–1 MHz with a voltage stimulus of 10 mV to derive the line-specific charge transfer resistance, which will be denoted as LSR\textsubscript{CT} in the following (for further experimental details see [14] and references therein). Values of LSR\textsubscript{CT} were calculated by multiplication of the charge transfer contribution with the polarization resistance (denoted as R\textsubscript{CT}), the actual TPB length per electrode area (h\textsubscript{TPB}) and the electrode area (A) via:

\[
LSR_{CT} = R_{CT} \times h_{TPB} \times A
\]  

(1)

The partial pressures of CO and CO\textsubscript{2} were varied in the range of \(4.0 \times 10^2\) Pa \(\leq p_{\text{CO}} \leq 5.1 \times 10^4\) Pa and \(9.5 \times 10^2\) Pa \(\leq p_{\text{CO}_2} \leq 9.2 \times 10^4\) Pa, respectively. The investigated operating temperature range was \(973\) K \(\leq T \leq 1073\) K. These conditions were chosen in order to avoid Ni oxidation and carbon formation due to the Boudouard reaction. The stability of the Ni patterned anodes used in the experiments was carefully controlled by pre- and post-test SEM analysis [14].

3. Results and discussion

3.1. Elementary kinetic reaction mechanism and model parameterization

In the elementary kinetic modeling approach, the global electrochemical CO oxidation reaction,

\[
\text{CO} + O_{\text{YSZ}}^\text{\textsubscript{O,YSZ}} = \text{CO}_2 + V_{\text{O,YSZ}}^\ast + 2e_{\text{Ni}}^-
\]  

(2)

is resolved into elementary steps taking place on the involved surfaces. The reaction mechanism is listed in Table 3 and schematically depicted in Fig. 2. The thermodynamic data of all involved species, which form the basis for a thermodynamically consistent kinetic model, are given in Table 2. The mechanism contains two gas-phase species (CO, CO\textsubscript{2}), four Ni surface species (\(\text{Ni}_{\text{Ni}}, \text{Ni}_{\text{Ni}}, \text{Ni}_{\text{Ni}}, \text{Ni}_{\text{Ni}}\)), five YSZ surface species (\(\text{O}_{\text{YSZ}}, \text{CO}_{\text{YSZ}}, \text{O}_{\text{YSZ}}, \text{O}_{\text{YSZ}}, \text{O}_{\text{YSZ}}\)) and two YSZ bulk species (\(V^\ast_{\text{O,YSZ}}, V^\ast_{\text{O,YSZ}}\)), which in Kröger–Vink notation [20] stand for bulk YSZ oxygen ions and vacancies, respectively. The species denoted as \(\text{O}_{\text{YSZ}}^\ast\) and \(\text{O}_{\text{Ni}}^\ast\) represent free surface adsorption sites on YSZ and Ni.

The thermal heterogeneous reaction mechanism for the Ni surface includes CO and CO\textsubscript{2} adsorption/desorption (Table 3, (R1) and (R2)) as well as CO oxidation steps via an Eley–Rideal (ER) reaction step (R3) and via a Langmuir–Hinshelwood (LH) reaction step (R4). In our previous work [13] we observed that for a consistent simulation of the high and low CO partial-pressure regimes, the addition of the ER pathway (R3) to the LH pathway (R4) was necessary. This can be rationalized because under low CO partial pressure the actual CO surface coverage is very low, which leads to a substantial decrease in the LH reaction rate. Under these conditions only an ER reaction,

\[\text{CO} + \text{Ni} \rightarrow \text{CO}_2 + \text{Ni} \]

Table 3

<table>
<thead>
<tr>
<th>Reaction</th>
<th>(k^0) (or (s^0))</th>
<th>(E^\ast) (kJ mol\textsuperscript{-1})</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ni surface reactions</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(\text{CO} + \text{Ni} \rightarrow \text{CO}_2 + \text{Ni}) (R1)</td>
<td>(s^0 = 5.0 \times 10^{-1})</td>
<td>0</td>
<td>[13]</td>
</tr>
<tr>
<td>(\text{CO}_{\text{Ni}} + \text{Ni} \rightarrow \text{CO}_2 + \text{Ni}) (R2)</td>
<td>(7.0 \times 10^{-2})</td>
<td>41</td>
<td>[13]</td>
</tr>
<tr>
<td>(\text{CO} + \text{Ni} \rightarrow \text{CO}_2 + \text{Ni}) (R3)</td>
<td>(1.0 \times 10^{-2}) cm\textsuperscript{2} mol\textsuperscript{-1} s\textsuperscript{-1}</td>
<td>181.8</td>
<td>[12,13]</td>
</tr>
<tr>
<td>(\text{CO} + \text{Ni} \rightarrow \text{CO}_2 + \text{Ni}) (R4)</td>
<td>(2.0 \times 10^{-2}) cm\textsuperscript{2} mol\textsuperscript{-1} s\textsuperscript{-1}</td>
<td>138.3</td>
<td>[12,13]</td>
</tr>
<tr>
<td>YSZ surface reactions</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(\text{CO} + \text{O}_{\text{YSZ}}^\ast \rightarrow \text{CO}<em>2 + \text{O}</em>{\text{YSZ}}^\ast) (R5)</td>
<td>(s^0 = 4.0 \times 10^{-2})</td>
<td>0</td>
<td>[13]</td>
</tr>
<tr>
<td>(\text{CO} + \text{O}_{\text{YSZ}}^\ast \rightarrow \text{CO}<em>2 + \text{O}</em>{\text{YSZ}}^\ast) (R6)</td>
<td>(1.0 \times 10^{-2}) cm\textsuperscript{2} mol\textsuperscript{-1} s\textsuperscript{-1}</td>
<td>115.1</td>
<td>[12,13]</td>
</tr>
<tr>
<td>(\text{CO} + \text{O}_{\text{YSZ}}^\ast \rightarrow \text{CO}<em>2 + \text{O}</em>{\text{YSZ}}^\ast) (R7)</td>
<td>(1.6 \times 10^{-2}) cm\textsuperscript{2} mol\textsuperscript{-1} s\textsuperscript{-1}</td>
<td>90.9</td>
<td>[12,13]</td>
</tr>
<tr>
<td>Charge-transfer reactions</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>(\text{O}<em>{\text{YSZ}}^\ast \rightarrow \text{O}</em>{\text{YSZ}}^\ast + e_{\text{Ni}}) (O1)</td>
<td>(8.4 \times 10^{-2}) mol cm\textsuperscript{-1} s\textsuperscript{-1}</td>
<td>75</td>
<td>Fit</td>
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<tr>
<td>(\text{O}<em>{\text{YSZ}}^\ast \rightarrow \text{O}</em>{\text{YSZ}}^\ast + e_{\text{Ni}}) (O2)</td>
<td>(3.5 \times 10^{-2}) mol cm\textsuperscript{-1} s\textsuperscript{-1}</td>
<td>135</td>
<td>Fit</td>
</tr>
<tr>
<td>(\text{O}<em>{\text{YSZ}}^\ast \rightarrow \text{O}</em>{\text{YSZ}}^\ast + e_{\text{Ni}}) (O3)</td>
<td>(1.4 \times 10^{-2}) mol cm\textsuperscript{-1} s\textsuperscript{-1}</td>
<td>82</td>
<td>Fit</td>
</tr>
</tbody>
</table>

where gaseous CO directly reacts with the more strongly adsorbed oxygen atom species, can maintain a sufficiently high CO$_2$ production rate. Following our previous work on Ni point electrodes, we have investigated CO oxidation on Ni via the ER pathway employing Density Functional Theory (DFT) [21]. In these calculations, an activation energy of 181.8 kJ mol$^{-1}$ was obtained for the ER reaction step (R4).

The reaction mechanism for the YSZ surface includes CO adsorption/desorption (R5) and a heterogeneous CO oxidation step (R6). According to temperature-programmed desorption (TPD) experiments [21], the CO oxidation proceeds predominately via an ER mechanism, where gas-phase CO directly reacts with YSZ surface-adsorbed oxygen atoms to yield gas-phase CO$_2$. This reaction is taken from Ref. [13] with modified kinetics according to the DFT results given in Ref. [21]. The transport of bulk oxygen species to the YSZ surface occurs by a bulk-surface migration process via electrolyte vacancies (Table 2, (R7)).

The model includes diffusion of the CO, CO$_2$ and O surface species on Ni and YSZ perpendicular to the TPB. We use values for the diffusion coefficients from our previous work [13] of $D(\text{CO}) = D(\text{CO}_2) = 4.7 \times 10^{-7}$ cm$^2$ s$^{-1}$, $D(\text{O}_2) = 1.3 \times 10^{-5}$ cm$^2$ s$^{-1}$, and $D(\text{YSZ}) = 2.3 \times 10^{-11}$ cm$^2$ s$^{-1}$, all values at 1073 K. It should be noted that at this temperature the nickel atoms themselves are mobile with a common pre-exponential factor of $1.4 \times 10^{-3}$ cm$^2$ s$^{-1}$ and an activation energy of 93.2 kJ mol$^{-1}$ [22]. The pre-exponential factor for bulk ion conductivity of the YSZ electrolyte was set to $3.6 \times 10^{-5}$ S cm$^{-1}$ with an activation energy of 90 kJ mol$^{-1}$ [13].

The charge-transfer (CT) mechanism derived in the present study comprises three CT reaction steps (denoted as (O1), (O2) and (O6) in Fig. 2 and Table 3), all of which take place at the TPB line. The CT reaction nomenclature follows that of our previous work on electrochemical H$_2$ and CO oxidation at Ni/YSZ anodes, where five different (O1–O5) CT reaction steps were investigated [12,13]. As described in detail in Ref. [13], only the reaction sequence (O1) followed by (O2) (denoted as (O1) and (O2)) resulted in a quantitative agreement with the point electrode experimental data of Lauvstad et al. [7]. In this reaction sequence, the first reaction (O1) represents a single electron oxidation step of an O$_{\text{YSZ}}$ species, followed by a second CT reaction (O2), which represent a spillover of an O$_{\text{YSZ}}$ onto the Ni electrode surface with a simultaneous electron transfer to the Ni. The additional CT reaction step (O6), introduced in the present simulation study describes the discharge of an YSZ oxygen ion, O$_{\text{YSZ}}^-$, which proceeds by an electron transfer to the Ni electrode with the resulting YSZ oxygen atom, O$_{\text{YSZ}}$, remaining on the YSZ electrolyte surface. The necessity for the implementation of the additional CT reaction (O6) will be rationalized below. The charge-transfer symmetry factor ($\beta$) is assumed to be 0.5 for all CT reactions. Although the modeling results are sensitive to the values of the symmetry factors, we decided to fix them to the “symmetric” value of 0.5 in order to reduce the number of fitting parameters. However, if further molecular-level details about the CT mechanism become available (such as the actual location of the transition state – closer to the reagent or closer to the product side), the refinement of the symmetry factors values might certainly be valuable.

A complex space-charge layer (electrical double layer) is present at the two-phase interface between the metal (Ni) and ion-conducting oxide (YSZ) [23], leading to additional capacitive effects upon transient electrochemical characterization (e.g., impedance spectroscopy). In the present model we assume that the double layer behavior can be described as ideal parallel-plate capacitor, where the capacitance is temperature-dependent via

$$C_d = C_{d,0} + C_{r,d} \times T$$  

By fitting to experimental impedance data, we obtained values of $C_{d,0}$ = 0.25 F m$^{-2}$ and $C_{r,d}$ = 8 × 10$^{-4}$ F K$^{-1}$ m$^{-2}$. At $T = 1073$ K, the obtained double layer capacitance has a value of 1.1 F m$^{-2}$. This value is somewhat lower than the value of 3.46 F m$^{-2}$ observed for pattern anodes in H$_2$/H$_2$O atmospheres [3]. The difference between the capacitances for these two reaction systems might be due to the high H$_2$O solubility in YSZ bulk [24–26], which can result in additional chemical capacitances in case of the H$_2$ system as compared to the CO system. For an ideal capacitor, the thickness $d$ of the interfacial double layer can be estimated via

$$d = \frac{\varepsilon \varepsilon_0}{\varepsilon_d}$$  

where $\varepsilon_0$ is the vacuum permittivity (8.85 × 10$^{-12}$ F m$^{-1}$), $\varepsilon$ is a dielectric constant ($\varepsilon = 27$ as obtained for bulk YSZ measured by Fork et al. [27]). Applying the values given above, we obtain $d = 0.21$ nm for CO/CO$_2$ and $d = 0.068$ nm for H$_2$/H$_2$O. Both values are smaller than the YSZ lattice parameters ($a = 0.36$ nm, $c = 0.518$ nm).

As already pointed out by Vels Hansen et al., such low values have no direct physical meaning and additional capacitances in parallel need to be assumed in order to interpret the measured capacitance values [28].

3.2. Electrochemical impedance spectra

Fig. 3 shows the comparison between experimental and simulated impedance spectra for four different conditions (varying pCO and pCO$_2$) at $T = 1073$ K. For each condition, both Nyquist plots (upper panels – a1, b1, c1, d1) and Bode plots (lower panels – a2, b2, c2, d2) are shown. We observe a qualitative agreement between model and experimental over the complete investigated range of experimental conditions. The quantitative discrepancy (rel. error <5%) needs to be considered as minor given the physical basis of the present model. Note that all simulations are based on one single parameter set. In contrast, equivalent circuit models usually show an excellent agreement with experimental data, however they are based upon fitting several parameters for each individual condition.

The polarization resistance obtained from the impedance spectra shown in Fig. 3 (R$_{\text{CT}}$ in Eq. (1)) was used to calculate LSR$_{\text{CT}}$, which will be discussed in the following sections.

3.3. Influence of CO/CO$_2$ gas-phase composition

Previous studies on model anodes have shown that gas-phase composition has a strong influence on the electrochemical performance, for both H$_2$ [3,29] and CO [13] oxidation. Fig. 4 shows a comparison between experimental and simulated values for LSR$_{\text{CT}}$ (open symbols) as a function of CO and CO$_2$ partial pressures for an operating temperature of 1073 K. The LSR$_{\text{CT}}$ shows a pronounced dependence on both CO and CO$_2$ partial pressures. Generally, increasing either CO or CO$_2$ partial pressure decreases LSR$_{\text{CT}}$ and therefore enhances the electrode kinetics. A particular interesting case is the pCO variations at low pCO$_2$ shown in Fig. 4a. Here, LSR$_{\text{CT}}$ shows different slopes at high and low pCO values and a broad minimum at pCO ~ 10$^3$ Pa. The minimum is also present for higher pCO$_2$, as seen in Fig. 4b, however, shifted towards higher pCO.

The observation that both reactant and product concentrations positively influence the electrode kinetics is non-intuitive. A similar behavior is known from the H$_2$/H$_2$O/Ni/YSZ model anode system, where both increasing hydrogen and water partial pressure increases the electrode kinetics [3,30]. We have showed before that this effect is due to the combined changes of concentration and electrode potential [3].
Simulation results for the full mechanism ((O1), (O2) and (O6), cf. Fig. 2), parameterized as given in Table 3, are shown as solid lines in Fig. 4. The simulations agree quantitatively with the experiments over the complete investigated range of CO and CO₂ partial pressures. This agreement is the basis for the model-based mechanistic interpretation below (Section 3.5). It should be noted that simulations based on the previously derived (O1) and (O2) consecutive CT reaction mechanism only (cf. Fig. 2), the experimentally observed minimum in LSR₉ could not be reproduced; only the high CO partial pressure region could be quantitatively described. This finding motivated the introduction of the additional CT reaction (O6).

3.4. Influence of temperature

The temperature dependence of LSR₉ was measured and simulated for two different CO partial pressures of pCO = 3 × 10⁴ Pa.
and $p_{CO} = 9.2 \times 10^2$ Pa at a constant $CO_2$ partial pressure of $p_{CO_2} = 2 \times 10^4$ Pa. These measurements form the basis of quantifying the activation energies of the charge transfer reactions (Table 3), which was not possible in previous work due to lacking temperature-dependent data [13]. The comparison between simulation (lines) and experimental results (open symbols) is shown in Fig. 5. The data show an Arrhenius-type temperature dependence with an activation energy that depends on $p_{CO}$. Both the absolute values and the slope of the simulated LSR$CT$ curves are in excellent agreement with the experimental data. This demonstrate the ability of (O1), (O2) and (O6) CT mechanism to reproduce the electrochemical data over the entire CO/CO$_2$ gas composition and operating temperature range investigated experimentally.

Fig. 5 also depicts the global activation energies for two cases at high and at low $p_{CO}$, determined from the slope of the experimental data. They have values of 141 kJ mol$^{-1}$ and 84 kJ mol$^{-1}$, respectively. The activation energies of the elementary charge-transfer steps (cf. Table 3) are very similar to these global activation energies. This finding confirms the change of rate-determining step for different $p_{CO}$ (see following section). Note that the globally observed activation energies are a convolution of the thermal activation of the different chemical and physical processes contributing to the polarization behavior.

3.5. Sensitivity analysis and mechanistic interpretation

In order to determine the influence of different physicochemical processes on the electrode performance, a sensitivity analysis was performed. In a sensitivity analysis, model parameters $P$ are individually varied by 10%, and the impact on the LSR$CT$ is quantified as dimensionless relative sensitivity, $s = (\Delta \text{LSR}_{CT}/\text{LSR}_{CT})/(\Delta P/P)$. A sensitivity of unity means that LSR$CT$ is directly proportional to the parameter. Negative sensitivities mean that a parameter increase leads to a decrease of LSR$CT$. Sensitivities close to zero mean that the process represented by that parameter is not rate-determining.

This analysis was carried out for two different gas compositions representing conditions left and right of the LST$CT$ minimum shown in Fig. 4a. Results are shown in Fig. 6. The parameters are separated into three blocks. The upper block describes kinetic parameters of all chemical and electrochemical reactions, the middle part includes the description of transport parameters of all surface species (diffusion coefficients) and the lower block describes the influence of physical parameters (TPB length, pressure and temperature). The change of heterogeneous reactions kinetics on both Ni and YSZ surfaces does not significantly influence LSR$CT$.  

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indicating that the interaction of the gaseous components with the Ni and YSZ surfaces is fast and not rate-determining. The analysis reveals that the charge-transfer reactions are rate-limiting. This is not surprising as these reactions are confined to the (one-dimensional) TPB line, whereas heterogeneous reactions take place at the (two-dimensional) surfaces. At low CO partial pressure ($p_{\text{CO}} = 9.2 \times 10^2$ Pa), reaction (O6) dominates the behavior with co-limitation of reaction (O2). Note that for this case oxygen diffusion on the YSZ surface also becomes rate co-limiting. This is a clear indication that, at low CO partial pressures, the dominant part of the conversion proceeds via (O1) and (O6) mechanism with CO oxidation on the YSZ surface. When the CO partial pressure increases ($p_{\text{CO}} = 3 \times 10^4$ Pa), the only rate-limiting step is the spillover reaction (O2), indicating that the conversion proceeds via (O1) and (O2) mechanism with CO oxidation on the Ni surface.

Fig. 6 further shows that the LSR$_{CT}$ is strongly and nonlinearly sensitive on temperature, as is already evident from Fig. 5. The sensitivity analysis indicates that different CO partial pressures result in different activation energy regimes.

For further insight into the mechanistic details, calculated coverages of all species on both Ni and YSZ surfaces are shown in Fig. 7 as a function of CO partial pressure. These results indicate that the oxygen coverage on the Ni adjacent to the TPB ($\theta_{\text{O,Ni}-\text{TPB}}$) is the key parameter inducing the change in the experimentally observed LSR$_{CT}$ slope. The value of $\theta_{\text{O,Ni}-\text{TPB}}$ is determined by the competition between oxygen formation at the Ni TPB region (due to the spillover CT reaction step (O2)) and the heterogeneous O removal reactions on Ni (via (R3) and (R4)), both of which depend on the CO/CO$_2$ gas composition. The present results show that at sufficiently high CO partial pressures the overall charge transfer occurs via (O1) and (O2) reactions, while at low CO partial pressures the overall charge transfer occurs via the (O1) and (O6), as the actual rate of (O2) spillover step strongly decreases with increasing $\theta_{\text{O,Ni}-\text{TPB}}$ due to surface site blocking. In the framework of the extended CT reaction model, the minimum in the experimental LSR$_{CT}$ values at $T = 1073$ K, $p_{\text{CO}_2} = 2 \times 10^4$ Pa and $p_{\text{CO}} = 1 \times 10^4$ Pa could be quantitatively reproduced in numerical simulations (Fig. 4). The simulations predict a value of $\theta_{\text{O,Ni}-\text{TPB}} = 0.15$ for these conditions (Fig. 7b).

4. Conclusions

We have presented a combined modeling and experimental study of electrochemical CO oxidation at SOFC Ni/YSZ anodes. An elementary kinetic model was developed and validated...
using well-defined micropatterned model anodes. By solving reaction–diffusion equations on both the Ni electrode and the YSZ electrolyte surfaces, experimental data were successfully reproduced over a wide range of CO/CO2/N2 gas compositions and operating temperatures. In contrast to commonly used Butler–Volmer equations, which are based on global reaction kinetics, the representation of all chemical processes in terms of elementary reactions allows for predictive simulations of the complex behavior of Ni/YSZ patterned anodes. Furthermore, the elementary kinetic description opens up the possibility of a direct mechanistic interpretation of the experimentally observed electrochemical characteristics via sensitivity analyses and surface coverage quantifications.

The main results of this study are summarized as follows: CO and CO2 concentrations have a strong and nonlinear influence on the electrode kinetics. Over a wide range of gas compositions, both an increase of CO or CO2 partial pressure enhances the electrode kinetics. This counter-intuitive effect is due to a simultaneous change of electric potential and coverages of species participating in the charge-transfer reaction. The gas composition also influences the macroscopically observed activation energy. It was shown that charge transfer can proceed via two different mechanisms: At high CO/CO2 ratios, oxygen spillover from the YSZ to the Ni surface takes place, and CO is oxidized in a Langmuir–Hinshelwood type heterogeneous reaction on the Ni surface. At low CO/CO2 ratios, the largest part of oxygen ions is fully reduced on the YSZ surface without undergoing spillover, and CO is oxidized in an Eley–Rideal type heterogeneous reaction on the YSZ surface.

The level of understanding achieved in the present study will be the basis for further investigations of the behavior of more complex SOFC fuels, in particular reformate gases or hydrocarbons.

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