Multicomponent Gas Diffusion in Porous Electrodes

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Multicomponent gas transport is investigated with unprecedented precision by AC impedance analysis of porous YSZ anode-supported solid oxide fuel cells. A fuel gas mixture of H2-H2O-N2 is fed to the anode, and impedance data are measured across the range of hydrogen partial pressure (10–100%) for open circuit conditions at three temperatures (800°C, 850°C and 900°C) and for 300 mA applied current at 800°C. For the first time, analytical formulae for the diffusion resistance (Rd) of three standard models of multicomponent gas transport (Fick, Stefan-Maxwell, and Dusty Gas) are derived and tested against the impedance data. The tortuosity is the only fitting parameter since all the diffusion coefficients are known. Only the Dusty Gas Model leads to a remarkable data collapse for over twenty experimental conditions, using a constant tortuosity consistent with permeability measurements and the Bruggeman relation. These results establish the accuracy of the Dusty Gas Model for multicomponent gas diffusion in porous media and confirm the efficacy of electrochemical impedance analysis to precisely determine transport mechanisms.

There is an extensive literature on modeling gas diffusion in porous media using Fick’s law, Stefan-Maxwell or Dusty Gas Model. It is thought that the Dusty Gas Model should be the most accurate, although it is also the most complicated and difficult to validate. Almost no analytical results are available, but the Dusty Gas Model has been used in a number of numerical simulations, albeit with constant pressure approximation which is inconsistent (see below). Moreover, no theoretical framework exists to analytically derive the diffusion resistance values from impedance data using these more complex diffusion models for porous media.

Instead, it is common practice for researchers to use limiting current values from the current-voltage or I-V curves to study gas diffusion in SOFC. Limiting current is usually obtained when the reactant is nearly or completely depleted at the reaction site. Therefore, it has often been used to derive properties of the porous electrode that would account for slow diffusion or sluggish mass transport. However, high tortuosity is commonly invoked to explain the limiting current values. Many previous attempts to fit models to I-V data for SOFC have been inconclusive with widely varying tortuosity values from 2 to 19 for the same system. Yet, most direct measurements conducted on anode materials and reconstruction of 3D microstructure indicate tortuosity values should be in the range of 1.5 to 3. At the same time, according to the theories about tortuosity, we should expect tortuosity of porous electrode with nice and open microstructures to be not too high. Actually, limiting current can have the signature of not only gas diffusion, but also dissociative adsorption, surface diffusion, catalytic redox reaction, or even gas transport in free channels outside the electrode. Therefore, we studied gas diffusion in porous electrodes using AC impedance, which better separates processes of different time scales and therefore provides better assignment of arcs in data to different processes.

The SOFC button cell we studied uses hydrogen as fuel, carried by nitrogen together with 1.7% of water vapor, at the anode. Oxygen in air is used as oxidant for cathode. Therefore, the electrochemical reaction
goes as follows. The oxygen molecule diffuses through the cathode bulk layer and reaches the functional layer, where it accepts electrons and is oxidized to oxygen ion, which is further conducted through the electrolyte layer. When it arrives at the anode functional layer, it reacts with hydrogen fuel, forming water and releasing electrons to the external circuit.

In this paper, we present a new theoretical approach to predict concentration profiles and diffusion resistance using Fick’s law (Fick), Stefan-Maxwell formulation (SM), and Dusty Gas Model (DGM) and compare with experimental data for SOFC. By using this approach in conjunction with AC impedance, we are able to show that DGM provides a very accurate description of multicomponent gas diffusion and can be used to subtract gas diffusion response from overall data for analyzing contributions from other physical processes.

**Theory**

Gas diffusion models.— Transport of gaseous components through porous media has been extensively studied over the years. In general, models including Fick’s model (FM), the Stefan–Maxwell model (SM) and the Dusty Gas model (DGM) are widely used to predict the concentration overpotential. Many researchers have concluded that among these three, the DGM is the most accurate and appropriate model to simulate gas transport phenomena inside a porous electrode such as SOFC electrodes. However, due to its complexity, this model has no analytical solutions, and the corresponding analysis requires complicated numerical simulation. In this work, we derive analytical formulae for the diffusion impedance of multicomponent systems because it considers the molecular collisions and can be carried out following Eq. 2. Where $D_i$ is the theoretical diffusion coefficient of species $i$, $\varepsilon_p$ and $\tau_p$ are the porosity and tortuosity of the porous electrode, respectively. Note that our standard definition of “tortuosity” is also sometimes called the “tortuosity factor”.

\[ D_{i,eff} = D_i \varepsilon_p \varepsilon_p \tau_p \]  

The Stefan-Maxwell model is more commonly used in multicomponent systems because it considers the molecular collisions among the different types of gas species by using a more complicated left hand side term (Eq. 3). However, it is more typically used in multicomponent systems because it considers the molecular collisions and can be used to subtract gas diffusion response from overall data for analyzing contributions from other physical processes.

In theory, compared to Stefan-Maxwell model, the Dusty Gas Model further takes into account the interactions between the gas molecules and the solid pore surface leading to Knudsen diffusion. This model assumes the pore walls consist of large molecules that are uniformly distributed in space. These pseudo ‘dust’ molecules also collide with real gas molecules, which leads to the Knudsen diffusion effect. Moreover, the fluxes due to pressure gradient are also taken into consideration. The general form of the DGM is as follows:

\[ \frac{N_i}{D_{K,i,j}^{eff}} + \sum_{j \neq i} X_j N_i - X_i N_j = -P \frac{dX_i}{RT} \frac{dx}{dx} \left( 1 + \frac{1}{D_{K,i,j}^{eff}} \frac{P}{\mu} \right) \frac{dP}{dx} \]  

where $B_0$ is the permeability of the porous medium and $\mu$ is the viscosity of the gas mixture. The first term in Eq. 4 describes Knudsen diffusion.

In both SM and DGM, the binary diffusion coefficients $D_{i,j}$ can be calculated using the Chapman-Enskog equation (Eq. 5), where $T$ is temperature in K, $P$ is pressure in Pa, $\Omega$ is the collision integral, $\sigma_{ij}$ is the collision diameter, and $M_i$ is the molecular weight of species $i$.

\[ D_{i,j} = 1.86 \times 10^{-3} \cdot T^{3/2} \cdot \left( \frac{1}{\mu_i} + \frac{1}{\mu_j} \right)^{1/2} \]  

\[ D_{K,i} = \frac{2}{3} \left( \frac{8RT}{\pi M_i} \right)^{1/2} T \]  

Note that this expression was derived from cylindrical pore geometry that having the mean radius $T$, but in reality, pore geometry can deviate from cylinders, therefore, this expression has some uncertainties in predicting Knudsen diffusivity. In this study, the mean pore size $T = 0.98 \mu m$ obtained by mercury porosimetry (which also assumes cylindrical pores) leads to relatively small Knudsen diffusion effects. Nevertheless, we shall see that the differences between the DGM and SM are noticeable and lead to improved accuracy in fitting the experimental data, and these differences would be amplified with smaller pore sizes.

The effective binary diffusivity and Knudsen diffusivity ($D_{i,j}^{eff}$ and $D_{K,i}^{eff}$) are defined as their theoretical counterparts ($D_{i,j}$ and $D_{K,i}$) times a geometric factor, which is porosity ($\varepsilon_p$) divided by tortuosity ($\tau_p$).

\[ D_{i,j}^{eff} = D_{i,j} \varepsilon_p \varepsilon_p \tau_p \]  

\[ D_{K,i}^{eff} = D_{K,i} \varepsilon_p \varepsilon_p \tau_p \]  

Numerous studies on transport through porous media in the absence of a chemical reaction reveal that the DGM is superior to the Fick’s law in its ability to predict the fluxes, and in porous catalyst, the Fick’s law is still frequently used because its simplicity allows explicit and analytical expressions to be derived. If nonuniform pressure is present in a porous media due to reactions involving a change in the number of molecules, additional permeation term has to be taken into account, and therefore the DGM should be adopted. Many works showed that the DGM can successfully predict the fluxes for these reactions in various reacting systems. For example, Davies used it for the SO2 oxidation reaction, Bli€ck applied it to the coal gasification where a large pressure gradient is present. However, the pressure gradient term requires additional computational time and cost. Therefore researchers started to use DGM without the permeation term if pressure gradient can be approximately neglected. And a comparison among different diffusion models, including Fick’s Law, Stefan-Maxwell model and Dusty Gas Model, to predict concentration polarization is presented in in Suwanwarangkul’s work.

**Graham’s law and the Dusty Gas Model.**— Strictly speaking, the Dusty Gas Model with the constant pressure assumption leads to a
paradox that has not been widely recognized in the SOFC community. Eq. 4 shows the general Dusty Gas Model with an extra permeation flux term due to the pressure variation, if we sum over all the gas species, the pressure gradient can be calculated as shown in Eq. 9.30 By taking a look at the numerator, we can find that the pressure gradient comes from the different effective Knudsen diffusivity $D_{ij}^{\text{eff}}$ of two active species in equi-molar counter-diffusion mode.

$$\sum_i \frac{N_i}{D_{ij}^{\text{eff}}} = -\frac{1}{RT} \left( 1 + \frac{R_i P}{\mu} \sum_i X_i \frac{D_{ij}^{\text{eff}}}{D_{ij}^{\text{eff}}_i} \right) \frac{dP}{dx}$$ \hspace{1cm} [8]

$$\frac{dP}{dx} = \left( 1 + \frac{R_i P}{\sum_i p x_i \frac{D_{ij}^{\text{eff}}}{D_{ij}^{\text{eff}}_i}} \right) \frac{dX_i}{dx}$$ \hspace{1cm} [9]

In the case of hydrogen molecules reacting to produce water vapor, the molar flux of all species should add up to zero. In this equi-molar counter diffusion mode, if the effective Knudsen diffusivity $D_{ij}^{\text{eff}}$ of hydrogen and water are the same, which means if the force exerted on the pore walls by $H_2$ and $H_2O$ are exactly the same but in the opposite direction, they will cancel each other and no pressure will build up. However, the molecular weight and size of the molecules vary among different species, therefore, Knudsen diffusivity must be different, which means total pressure has to change throughout the electrode. From another point of view, in the constant pressure assumption, the summation over all gas components will lead to Graham’s law,45 which states the sum of molar flux ($N_i$) times the square root of the molecular weight ($M_i$) should be zero (Eq. 10). Indeed, Graham’s law is valid in the absence of chemical reactions. However, when chemical reactions occur, the component fluxes are related through the reaction stoichiometry, and only isomerization reactions will be consistent with Graham’s law.

$$\sum_i N_i \sqrt{M_i} = 0$$ \hspace{1cm} [10]

In our case, moles of $H_2$ react to form equivalent number of moles of $H_2O$ and this is obviously contradictory to the flux relations imposed by the reaction. Since the algebraic derivation from Dusty Gas Model to Graham’s law is strict, this conflict indicates the Dusty Gas Model is intrinsically inconsistent with the constant pressure assumption. Actually, Graham’s law is only valid in the case of gas diffusion without reaction in general. In the case of SOFC, the gas diffusion in porous electrode has a boundary condition of surface reaction at the functional layer/electrolyte interface; therefore, the flux of active species ($H_2$ and $H_2O$) cannot be captured by Graham’s law. However, some current researches still use it to study gas transport in porous SOFC electrodes.44,46 In fact, the full DGM with permeation flux term due to pressure variation has no problem, and is accurate enough to satisfy chemical reaction boundary conditions. Yet with the permeation term, DGM is too complicated for deriving analytical results, therefore restrict its acceptability in some theoretical studies. But we will provide a proof, in the Validation of the isobaric approximation in the Dusty Gas Model section that in porous electrodes of SOFC, the pressure gradient effects on the gas transport is not significant and can be safely neglected.

**Steady-state concentration profiles.**— From the governing equations, we can analytically derive the concentration profiles throughout the porous electrode with the constant pressure assumption when the bulk concentrations of different gas species are taken to be known. In Eqs. 11–13, $X_i^0$ is the molar fraction of species $i$ in the bulk gas mixture outside the porous electrode, $X_i(x)$ is the molar fraction of species $i$ at position $x$. $I$ is the total current with the sign convention of $I > 0$ for hydrogen consumption, $F$ is the Faraday constant. $R$ is universal gas constant and $P$ is the total gas pressure.

$$X_{N_2}(x) = X_{N_2}^0 \exp \left( \frac{RT}{2FP} \left( \frac{1}{D_{N_2H_2O}^{\text{eff}}} - \frac{1}{D_{N_2H_2O}^{\text{eff}}} \right) x \right)$$ \hspace{1cm} [11]

$$X_{H_2} = X_{H_2}^0 - \frac{RT}{2FP} \left( \frac{1}{D_{N_2H_2O}^{\text{eff}}} + \frac{1}{D_{N_2H_2O}^{\text{eff}}} \right) x$$

$$- \left( \frac{D_{N_2H_2O}^{\text{eff}} - D_{N_2H_2O}^{\text{eff}}} {D_{N_2H_2O}^{\text{eff}} + D_{N_2H_2O}^{\text{eff}}} \right) X_{N_2}^0$$

$$\times \left[ \exp \left( \frac{RT}{2FP} \left( \frac{1}{D_{N_2H_2O}^{\text{eff}}} + \frac{1}{D_{N_2H_2O}^{\text{eff}}} \right) x \right) - 1 \right]$$ \hspace{1cm} [12]

$$X_{H_2O} = X_{H_2O}^0 + \frac{RT}{2FP} \left( \frac{1}{D_{N_2H_2O}^{\text{eff}}} - \frac{1}{D_{N_2H_2O}^{\text{eff}}} \right) x$$

$$+ \left( \frac{D_{N_2H_2O}^{\text{eff}} - D_{N_2H_2O}^{\text{eff}}} {D_{N_2H_2O}^{\text{eff}} + D_{N_2H_2O}^{\text{eff}}} \right) X_{N_2}^0$$

$$\times \left[ \exp \left( \frac{RT}{2FP} \left( \frac{1}{D_{N_2H_2O}^{\text{eff}}} - \frac{1}{D_{N_2H_2O}^{\text{eff}}} \right) x \right) - 1 \right]$$ \hspace{1cm} [13]

**Concentration polarization.**— From the concentration profile calculation, we know the gas concentration at the reaction surface and then concentration overpotential can be calculated using Nernst equation (Eq. 14)

$$\eta_{\text{node}, \text{conc}} = -\frac{RT}{2F} \ln \left( \frac{X_{H_2} X_{H_2O}^0}{X_{H_2}^0 X_{H_2O}} \right)$$ \hspace{1cm} [14]

In the case of impedance under current, the concentration outside the porous electrode is very close to the bulk concentration (the concentration in the feed gas). However, under a non-zero current, some of the reactants need to react electrochemically to support the current, therefore, there must be some concentration gradient resulting from the consumption of the reactants. We use a continuously stirred tank reactor (CSTR) assumption to approximately calculate the gas concentration outside the cell in the feed tube,16,47 as described in Eq. 21 and Eq. 22.

**Bulk diffusion impedance ($R_b$).**— The above mentioned gas diffusion models, including Fick’s law, Stefan Maxwell and Dusty Gas Model, are not new, and are widely used to predict I-V curves and fit the limiting current values as mentioned earlier.12,26 But the SM and DGM models have rarely been used before to analytically analyze impedance spectra of SOFC, although they have been used to describe gas diffusion in porous electrodes. By taking the derivative of the concentration overpotential with respect to current and evaluating it at a specified current, diffusion resistance ($R_b$) is obtained for all three models. In the Fick’s law derivation, the Bosanquet formula was adopted for the theoretical diffusivity derivation. By taking a look at zero current $R_b$ in Eqs. 16, 17, and 18, we notice that compared to $R_b$ value from Fick’s law, the $R_b$ of SM has a complicated extra term resulting from the consideration of interactions among different gas species. Also, the $R_b$ value derived from DGM further incorporated the Knudsen effect, which accounts for the collision of gas molecules with the pore wall. The multicomponent gas diffusion inside the porous electrodes was then studied by comparing these three different diffusion models. In Eqs. 15 to 18, $R_{\text{node}}(t)$ is the gas diffusion resistance at current $I$, $\eta_{\text{node}}$ is the anode concentration overpotential due to gas diffusion, $P_{\text{an}}$ and $X_{\text{an}}^0$ are the partial pressure and molar fraction of species $i$ in the bulk gas mixture outside the porous electrode. All other parameters are defined the same way as in the general Dusty Gas Model.

$$R_{\text{node}}(t=0) = \frac{d\eta_{\text{node}}}{dt} \mid_{(t=0)} \text{ or } R_{\text{node}}(t) = \frac{d\eta_{\text{node}}}{dt} \mid_{(t)}$$ \hspace{1cm} [15]
Note that in the $R_b$ expressions, there are not too many quantities that need to be fitted to data. Almost all the variables and parameters are determined from experimental inputs or estimations from kinetic gas theory, except for a microstructure factor (porosity divided by tortuosity), which links effective diffusivity inside porous electrode with its theoretical value. When the porosity is known, the only quantity need to be determined from fitting is the tortuosity value of the electrodes.

Similarly, cathode diffusion resistance can also be estimated by deriving from a specified diffusion model, e.g., Dusty Gas Model derivation was shown in Eq. 19:

$$R_{b,DGM\text{(cathode)}} = \left(\frac{RT}{2F}\right)^2 L_a \left\{ \frac{1}{P_{O_2}} \left( \frac{1}{D_{O_2}^{eff}} + \frac{1}{D_{N_2}^{eff}} \right) \right\}$$  \[19\]

$$R_{b,DGM\text{(anode)}} = \left(\frac{RT}{2F}\right)^2 L_a \left\{ \frac{1}{P_{H_2}} \left( \frac{1}{D_{H_2}^{eff}} + \frac{1}{D_{N_2}^{eff}} \right) + \frac{X_{N_2}^0}{D_{N_2}^{eff} - D_{H_2}^{eff}} \right\}$$  \[18\]

Comparing the theoretical $R_b$ at anode and cathode, Figure 1 shows the ratio between cathode $R_b$ and anode $R_b$ multiplied by 100%. It clearly shows that theoretical $R_b$ of cathode is less than 0.5% of that of the anode in anode supported cells. Though the porosity and tortuosity can be slightly different in two porous electrodes, we can still safely conclude that the diffusion resistance from anode side dominates. Therefore, in all the following discussion, we assume that the dominant contribution to the total gas diffusion resistance comes from the anode gas diffusion resistance.

Besides the successful fitting of the data below, this assumption can also be further justified theoretically. In particular, we estimate the gas conversion resistance due to the stagnant diffusion layer outside the porous electrode in the open circuit condition, studied by Prindah and Mogensen.\textsuperscript{48} With the relatively high total gas flow rate of 300 ml/min (at 25°C), operating conditions of 800 C and electrode area of 1 cm², the theory (Eq. 6) gives a gas flow rate of 2.045 mol/m²/s, and the predicted gas conversion resistance ranges from 0.0403 to 0.0511 ohm cm², which only accounts for 9.3% to 16.1% of the low frequency resistance in our data. Since the gas conversion resistance is relatively small, we proceed to consider only gas diffusion within the porous electrode.

The low frequency arc in the impedance data was fitted with a finite-length Warburg element in a Randles circuit (Fig. 2) to extract the anode diffusion resistance ($R_{b}$), which was then compared to analytical predictions from the three diffusion models (Eqs. 16–18). It has been shown by distribution of relaxation times (DRT) analysis that anode supported SOFC can exhibit multiple processes with similar time scales and overlapping impedance arcs at low frequency,\textsuperscript{51} which cannot be described by a Randles circuit. However, here we use this simple circuit model (which fits the low frequency impedance arc fairly well) only to extract the diffusion resistance for comparison with steady-state gas diffusion analyses, and not to model any frequency
dependence. Note that low frequency arc in SOFC impedance data could also possibly come from oxygen reduction and electrocatalysis at the cathode, but our study with some other in-house made cells found that, in the anode-supported cells studied in this work, contribution from oxygen reduction at cathode is very small. In other cells with thinner anodes, however, we are able to use the accurate gas model of this paper to show that electrocatalysis at the cathode can dominate the low-frequency impedance.

Nonlinear least-squares fitting.— A fitting procedure called complex nonlinear least-squares fitting (CNLS), was implemented, where data sets of \( Z_{\text{real}}, Z_{\text{imaginary}} \) versus frequency, or \((|Z|, \text{phase angle})\) versus frequency \( \omega \) are used. The aim of the least squares fitting procedure is to find a set of parameters which will minimize the sum of weighted deviations (Eq. 20).

\[
\sum_{k=1}^{n} w_k \cdot \left[ (Z_{r,k} - Z_{r,k}'(\omega))^2 + (Z_{i,k} - Z_{i,k}'(\omega))^2 \right]^{\frac{1}{2}}
\]

Where subscript \( k \) denotes the \( k \)th data point in impedance plot, \( Z_{r,k} \) is the real part of the experimental impedance data, while its counterpart \( Z_{r,k}' \) is the theoretical prediction of the real part of the impedance response. Similarly, \( Z_{i,k} \) and \( Z_{i,k}' \) are the imaginary parts of the impedance, experimentally and theoretically, respectively. Note that the theoretical prediction of the impedance is a function of frequency \( \omega \), which makes fitting of the Nyquist plot to be a three-dimensional curve fitting. Curves should not only match the correspondence of real and imaginary parts, but also need to match their frequency dependence as well. \( w_k \) is the weighting factor, for which we use reciprocal of the magnitude of the \( k \)th data point in this study. By minimizing the sum using the least square logarithm, a set of optimized parameters will be obtained.

We chose a Levenberg–Marquardt nonlinear least-squares fitting algorithm because of its straightforward implementation. Any parameter entering the model can in principle be used as a free fitting parameter; however, care must be taken to stay within limits of physical sense. It should also be noted that the Levenberg–Marquardt algorithm does not necessarily iterate to a global optimum of the fitting parameters, nor does it give any indication for the uniqueness of the optimized parameters. Therefore it is important to start from realistic initial guesses for the free parameters, and to exclude fitted results by analyzing its orders of magnitude and looking at the fitted graphs. Or upper and lower bounds can be set in the process of nonlinear least square fitting.

Validation of the isobaric approximation in the Dusty Gas Model.— As mentioned before, the Dusty Gas Model is intrinsically inconsistent with isobaric or constant pressure assumption. However, by comparing the gas composition profiles of \( \text{H}_2, \text{N}_2 \) and \( \text{H}_2\text{O} \), we can see pressure variation only leads to very small deviations of the gas composition profiles (Fig. 3). At the same time, theoretical prediction of anode gas diffusion resistance \( R_b \) values derived from the full Dusty Gas Model is practically the same as that derived from the isobaric Dusty Gas Model (Fig. 4). Therefore, our analysis proves that it is still safe to neglect total pressure variation inside the porous electrode we used when using the Dusty Gas Model, although theoretically there is some inconsistency between the model itself and the isobaric assumption. Therefore, all the analysis and results we show in this paper are based on isobaric assumption, assuming total pressure inside porous electrode does not vary in depth.

Before proceeding, we emphasize that validity of the isobaric assumption relies on the relatively large mean pore radius of our porous electrode (0.98 \( \mu m \), obtained by mercury porosimetry). At such scales, Knudsen diffusion is relatively insignificant, since gas molecules collide more often with each other than with the pore walls. For nanoporous solids or lower gas pressures, the isobaric assumption could break down, especially at high currents with strong concentration polarization.
Experimental

Anode-supported single cells were fabricated based on technology developed by Forschungszentrum Jülich. The anode was approximately 1.5 mm thick and 1 inch in diameter, while the cathode was 0.1 mm thick, and 0.5 inch in diameter. The anode side consists of an anode support layer and an anode functional layer with a thickness of 15~30 μm, both of which are composed of Ni/YSZ composites with different loadings and microstructures. The cathode side consists of a cathode current collection layer made from pure LSM and a cathode functional layer with a thickness of 15~30 μm, which is composed of YSZ/LSM composite.

A new cell was sealed at the circumference using LP-1071 glass from Applied technologies and dried in an oven at 120°C for 20 min. Then, it was placed into a spring loaded single cell testing fixture (Fig. 5). The fixture was put into a furnace with N2 (150 sccm) on the anode side and Air (150 sccm) on the cathode side. The furnace was then heated to 800°C at 5°C/min. The cell was reduced the next morning for 3 hours by gradually switching the anode gas from N2 to H2 flowing at 300 sccm. During testing, a tertiary gas mixture of hydrogen, nitrogen and 1.7% by volume water vapor was provided to the anode from a top feeding tube and air was fed from the bottom, carrying oxygen to the cathode. Pt lead wires were connected from the current collector layers (Pt mesh on cathode and nickel on the anode) to the data collecting equipment. A 1470 E Solartron Analytical current collector layers (Pt mesh on cathode and nickel on the anode) to the data collecting equipment. A 1470 E Solartron Analytical mounts and use it to further calculate concentration profiles of other active species, such as H2 and H2O.

Figure 7 shows the experimental Rb values we extracted from experimental data (black curve), and the theoretical predictions of anode Rb without taking into account the microstructure factor from three different models (colored curves). From the fitting, it is noteworthy that the Dusty Gas Model gives a constant structural factor (porosity/tortuosity), independent of hydrogen partial pressure over, with the anode porosity known to be 46%, the tortuosity fitted to the data is 2.30, which matches both theoretical expectations and independent experimental measurements by mercury porosimetry (in the range of 2~4). After taking into account the fitted tortuosity, the Dusty Gas Model best describes the gas diffusion, while the Stefan-Maxwell model shows some deviations, and Fick's law cannot capture the performance at all (Fig. 9). It is also interesting to note that Dusty Gas Model with constant N2 composition (green curve) does not give good enough results as well, which confirmed the necessity to calculate nitrogen concentration without any assumption, and use it to further calculate concentration profiles of other active species, such as H2 and H2O.

We also derived the diffusion resistance Rb from the impedance at a non-zero current that represents typical operating conditions for SOFC. The impedance was measured at the current of 300 mA/cm² (Fig. 10), and the corresponding Rb values for all three models were numerically evaluated using Maple software. It is worth noting that at a non-zero current, the bulk gas concentration (concentration at the interface between porous electrode and gas feeding tube) can deviate from the feeding concentration due to the concentration polarization resulting from consumption of reactants by electrochemical reactions. And, the concentration gradients in the gas feeding tube can be approximated using a continuous stirred tank reactor (CSTR) model. Therefore a CSTR correction (Eq. 21, 22) is introduced for calculating the concentration boundary condition of the bulk gas concentration. P0i is the ideal bulk concentration of species i, and Pbi is the corrected bulk concentration of species i after the CSTR formulation. In these equations, Nj is the molecular flux of species i in mol/(m²·s), A is

Results

Figure 7 shows the experimental Rb values we extracted from experimental data (black curve), and the theoretical predictions of anode Rb without taking into account the microstructure factor from three different models (colored curves). From the fitting, it is noteworthy that the Dusty Gas Model gives a constant structural factor (porosity divided by tortuosity), independent of hydrogen partial pressure (Fig. 8). This is consistent with real physics, where the microstructure of the porous media does not change with testing conditions. Moreover, with the anode porosity known to be 46%, the tortuosity fitted from the Dusty Gas Model is 2.30, which matches both theoretical expectations and independent experimental measurements by mercury porosimetry (in the range of 2~4). After taking into account the fitted tortuosity, the Dusty Gas Model best describes the gas diffusion, while the Stefan-Maxwell model shows some deviations, and Fick's

Figure 7. Experimentally extracted anode Rb and theoretical predictions of anode Rb of anode supported cell under different hydrogen partial pressures at OCV.

Figure 8. Structural factor (porosity/tortuosity) values fitted from three diffusion models under OCV.
Figure 9. Comparison between diffusion resistances (Rb) derived from models and the values extracted from experimental data after taking into account the fitted structural factor (porosity/tortuosity).

\[ p_{H_2}^{0*} = p_{H_2}^0 - \frac{N_{H_2} A}{m_T} p \]  

\[ p_{H_2O}^{0*} = p_{H_2O}^0 + \frac{N_{H_2O} A}{m_T} p \]

With the CSTR correction, the fitted porosity/tortuosity is almost independent of feeding gas composition and is practically the same as the zero current impedance analysis, which is another validation of the method of analyzing Rb to extract tortuosity. However, it is important to note that the CSTR approximation can lose accuracy in the limits of low gas flow rate, high operating temperature or very large current density (near the limiting current), outside the parameter regime considered here.

We further applied this gas diffusion analysis to zero-current impedance measured at different temperatures (800°C, 850°C and 900°C) (Fig. 12), and the tortuosity fitted at these temperatures only varies slightly, from 3.1 to 3.3 (Fig. 13). This also shows the applicability of the proposed anode gas diffusion resistance to AC impedance measured at different temperatures. Note that the data in Figures 9 and 11 come from two different cells prepared in exactly the same
Gas Model was identified, but numerical simulation confirmed that the total pressure variation only has very small effects on gas composition profiles and the predicted gas diffusion resistance ($R_b$) for the case of 0.98 μm mean pore radius. Therefore, we can still safely use the isobaric assumption with the Dusty Gas Model, which could otherwise break down at high currents in electrodes with much smaller pores in the Knudsen regime. By incorporating interactions between different gas molecules and collisions between gas molecules and pore walls (Knudsen effect), the Dusty Gas Model works best and gives a more or less constant tortuosity value over a wide range of operating conditions (10% to ∼100% of hydrogen partial pressure, zero and non-zero currents, and three different temperatures), and the fitted tortuosity value matches well with direct experimental measurements using mercury porosimetry.

In summary, we have developed a new theoretical approach to utilize AC impedance data and various analytical models to investigate multicomponent gas diffusion in porous media. The remarkable data collapse of the measured gas diffusion resistance for a wide range of hydrogen partial pressures, currents and temperatures with a single, reasonable tortuosity establishes DGM as the best model for gas diffusion in porous media (at least under these conditions). Therefore, this approach can be used to estimate tortuosity for porous media or to estimate gas diffusion resistance for further investigating other physical processes occurring inside the porous electrodes. For example, we show in a companion paper that the same model can be used to successfully distinguish the impedance signatures of gas diffusion from those of electrocatalysis in SOFC cathodes. This study further shows that electrochemical impedance analysis is a much more reliable method to measure gas diffusion in porous media than other methods based on permeability or limiting-current measurements.

Conclusions

In this work, we investigated the multi-component gas transport in porous electrodes using anode-supported solid oxide fuel cells, and proposed a new theoretical approach to predict gas diffusion resistance ($R_b$). Explicit analytical expressions for gas diffusion resistance ($R_b$) were derived at zero current conditions, and values of gas diffusion resistance ($R_b$) were evaluated numerically at non-zero current conditions. Comparison of cathode and anode gas diffusion resistance shows that in anode supported cells, the anode is the major contributor to gas diffusion resistance.

Experimental $R_b$ values were determined by fitting the low frequency arc of the anode supported cell to the finite length Warburg impedance in a Randles circuit. Then, they were compared to predictions from three analytical models, including Fick’s Law, Stefan-Maxwell model, and Dusty Gas Model, to determine the structure factor (porosity divided by tortuosity) or tortuosity when porosity is known. An inconsistency between isobaric assumption and the Dusty

References