Ultra large-scale simulation of polymer electrolyte fuel cells

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Abstract

This paper reports on the computational performance and detailed results of ultra large-scale simulations of a 200 cm² polymer electrolyte fuel cell (PEFC) using a 23.5 million gridpoint mesh. The computer code is based on a comprehensive single-phase PEFC model that features a detailed membrane-electrode assembly (MEA) model, electron transport, thermal and species transport, coolant heat transfer, in addition to other standard functionalities. Two cases under dry operation are simulated and compared. One case concerns an infinitely large coolant flowrate and consequently a constant temperature of bipolar plates. The other case involves a finite flowrate and a lower inlet coolant temperature designed to avoid membrane dryout in the inlet region while alleviating electrode flooding in the outlet region.

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1. Introduction

Computer simulation of PEFCs has been an active research area: see the most recent comprehensive review of Wang [1]. Significant capabilities exist using the single-phase modeling approach. Notable examples include a detailed model of membrane-electrode assembly (MEA) [2–4], electron transport and direct prescription of constant current boundary condition [5,6], non-isothermal modeling and hence coupled consideration of water and thermal management [7], variable gas flow [8,9], transient analysis [10], and experimental validation against detailed current distribution data [11,12].

For single-phase calculations, Meng and Wang [13] first proposed a parallel computing methodology to handle large-scale simulations involving millions of gridpoints. They simulated a variety of wet-to-dry operating conditions with a five-channel serpentine flow-field and benchmarked the parallel computing performance to be greater than 7× speed-up with 10 processors. The largest calculations reported so far were by Wang [1] (2.7 million cells); Shimpalee et al. [14] (5 million cells), and Wang and Wang [15] (2.7 million cells). However, the industry has a need for simulations using an order-of-magnitude larger mesh with reasonable computing time. This paper presents 23.5 million gridpoint calculations for the first time, herein termed ultra large-scale simulation to distinguish from prior studies of large-scale simulations. In addition, we will show a capability to perform co-simulation of electrochemical/transport phenomena in a PEFC and heat transfer in coolant channels. Such an integrated heat and water management tool is highly desirable in design and engineering of commercial-size PEFCs.

2. Numerical model

Fig. 1 shows the computational domain of the 200 cm² PEFC featuring a 24-channel, 3-pass serpentine flowfield, which includes the flow plates, gas and coolant channels, gas diffusion layers (GDLs), and catalyst layers on both the anode and cathode, as well as the membrane. The geometrical dimensions of various components in the cell are listed in Table 1.
2.1. Governing equations

The single-phase PEFC model employed in this work consists of nonlinear, coupled partial differential equations governing the conservation of mass, momentum, species, energy, and charge with electrochemical reactions. The equations can be written in the vector form, as [3–7]:

Continuity equation:
\[ \nabla \cdot (\rho \vec{u}) = 0 \quad (1) \]

Momentum conservation:
\[ \frac{1}{\varepsilon} \nabla \cdot (\rho \vec{u} \vec{u}) = -\nabla p + \nabla \cdot \tau + S_t \quad (2) \]

Species conservation:
\[ \nabla \cdot (\vec{u} \mathbf{C}_k) = \nabla \cdot (D^{\text{eff}} \nabla C_k) + S_k \]

Energy conservation:
\[ \nabla \cdot (\rho C_p \vec{u} T) = \nabla \cdot (k^{\text{eff}} \nabla T) + S_T \quad (4) \]

Proton transport equation:
\[ \nabla \cdot (\kappa^{\text{eff}} \nabla \phi_e) + S_{\phi_e} = 0 \quad (5) \]

Electron transport equation:
\[ \nabla \cdot (\sigma^{\text{eff}} \nabla \phi_s) + S_{\phi_s} = 0 \quad (6) \]

where \( \rho, \vec{u}, p, C_k, T, \phi_e, \phi_s \) denote the gas density, superficial fluid velocity vector, pressure, molar concentration of species \( k \), temperature, electrolyte and electronic phase potentials, respectively. The species considered here are hydrogen, oxygen, and water. The various source terms, electrochemical properties, and thermophysical properties identified for various regions of a PEFC, as well as necessary boundary conditions, have been detailed in [3–7] and thus are not repeated here. Select physical properties most relevant to the present simulation results are listed in Table 1.

2.2. Numerical procedure

The conservation equations, Eqs. (1)–(6), are solved by the commercial package, Fluent®, with the SIMPLEC algorithm [16], using a parallel computational methodology for a Linux PC cluster. The SIMPLEC algorithm is to update the pressure and velocity fields from the solution of the pressure corrections equation, solved by algebraic multi-grid (AMG) method [17]. Following the solution of the flow field, the energy, species, proton, and electron equations are solved. Approximately 23.5 million computational elements \((46 \times 900 \times 600)\) are employed to capture details of flow, thermal, electrochemical and mass transport phenomena in the PEFC. The MEA, where the important electrochemical
reactions and water back-diffusion process occur, is accurately captured by placing five and eight gridpoints in the catalyst layer and the membrane, respectively, along the through-plane direction. Details of the computational mesh are shown in Fig. 1. A constant average current density of 1.0A cm$^{-2}$ is specified as an input parameter, allowing the local current density and electronic phase potential to be predicted to vary spatially according to local operating conditions [6]. A simulation using 23.5 million cells typically requires about 600 iterations with the global mass balance less than 1% and residuals of species equations less than 10$^{-6}$, and takes nearly 20 h on 32 nodes of 2.8 GHz Intel Pentium 4 CPU and 1.0 GB DDR RAM.

### 3. Results and discussion

A 200 cm$^2$ PEFC is considered with the inlet gases having relative humidity (RH) of 50% relative to 80 $^\circ$C on both anode and cathode sides. Other common operating conditions are detailed in Table 1. Two simulation cases are presented in this work. Case 1 involves an infinitely large flow rate of the coolant, thus effectively keeping the flow plate temperature constant at 80 $^\circ$C due to infinitely large coolant flowrate, there is ∼3 K temperature rise in the dryout of the membrane in the inlet region, thereby increasing the local ionic resistance. As gases travel through the fuel cell, membrane hydration improves by product water and hence its ionic resistance decreases downstream and the current density increases.

Fig. 2 shows the current density distribution in Case 1. It can be seen that the current density initially increases along the cathode flow, as can be explained by the ohmic control of cell performance. The dry anode and cathode gases result in 26 mV gain in cell voltage in Case 2 as compared with Case 1. In addition, detailed contours of the current density, membrane water content, and membrane temperature, membrane water content, and membrane water crossover coefficient are presented below to illustrate the predictive capabilities of such an ultra large-scale simulation tool.

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### Table 1

<table>
<thead>
<tr>
<th>Quantity</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gas channel depth</td>
<td>1.0 mm</td>
</tr>
<tr>
<td>Gas channel width</td>
<td>1.0 mm</td>
</tr>
<tr>
<td>Land width</td>
<td>1.0 mm</td>
</tr>
<tr>
<td>Diffusion layer thickness</td>
<td>0.2 mm</td>
</tr>
<tr>
<td>Catalyst layer thickness</td>
<td>0.01 mm</td>
</tr>
<tr>
<td>Membrane thickness</td>
<td>0.045 mm</td>
</tr>
<tr>
<td>Fuel cell footprint area</td>
<td>143 mm × 143 mm</td>
</tr>
<tr>
<td>Anode/cathode/catalyst layer</td>
<td>Co-flow</td>
</tr>
<tr>
<td>Anode/cathode pressure, $P$</td>
<td>2.072 atm</td>
</tr>
<tr>
<td>Average current density, $I_{av}$</td>
<td>1.0 A cm$^{-2}$</td>
</tr>
<tr>
<td>Stoichiometric ratio ($\xi$)</td>
<td>2.0/2.0</td>
</tr>
<tr>
<td>RH of anode/cathode inlet @ 80 $^\circ$C</td>
<td>50%/50%</td>
</tr>
<tr>
<td>GDL porosity, $\varepsilon$</td>
<td>0.6</td>
</tr>
<tr>
<td>Cataly layer porosity, $\varepsilon$</td>
<td>0.4</td>
</tr>
<tr>
<td>Volume fraction of ionomer in catalyst layer, $\varepsilon$</td>
<td>0.26</td>
</tr>
<tr>
<td>GDL permeability, $K$</td>
<td>10$^{-11}$ m$^2$</td>
</tr>
<tr>
<td>Specific heat, $c_p$, of coolant (de-ionized water)</td>
<td>4182 J kg$^{-1}$ K$^{-1}$</td>
</tr>
<tr>
<td>Water content in membrane, $\lambda$</td>
<td>[9,19]</td>
</tr>
</tbody>
</table>

# Table 2

<table>
<thead>
<tr>
<th>Quantity</th>
<th>Case 1</th>
<th>Case 2</th>
</tr>
</thead>
<tbody>
<tr>
<td>Coolant flow rate</td>
<td>Infinite</td>
<td>2.9 × 10$^{-4}$ m s$^{-1}$</td>
</tr>
<tr>
<td>Coolant inlet temperature</td>
<td>353.15 K (80 $^\circ$C)</td>
<td>345.15 K (72 $^\circ$C)</td>
</tr>
<tr>
<td>Predicted cell voltage</td>
<td>0.590 V</td>
<td>0.616 V</td>
</tr>
</tbody>
</table>
Fig. 4 presents the distribution of membrane water content at the interface between the membrane and cathode catalyst layer for Case 1. It can be seen that the cathode is dry in the inlet region in spite of water production. In addition, water content reaches the fully hydrated level, i.e. $\lambda = 14$, in the outlet region, resulting in likely occurrence of cathode flooding.

Fig. 4 points out two problems inherent in the design of Case 1. First, the dry gas feed, while preferred for the automotive application, causes membrane dryout in the inlet region. Secondly, the outlet gas stream is over-saturated. These two issues can be resolved by proper thermal control through the coolant flow design, such as in Case 2. Fig. 5 presents the temperature distribution in the cathode flow plate for Case 2. It is seen that this coolant design, using a lower inlet temperature and a small flow rate, creates a thermal gradient (345–355 K) from the inlet to the outlet regions. A significant benefit of this thermal gradient along the flow direction can be seen in Fig. 6, where membrane water content is displayed. As compared to Fig. 3 with isothermal cooling, Case 2 increases the membrane water content in the inlet region under lower local temperatures and at the same time maintains fully saturated gas in the outlet region under higher temperatures. The net result of this thermally graded fuel cell is that the membrane water content becomes more uniform.

As a result of the more uniform degree of membrane hydration, the current density near the inlet becomes higher, as shown in Fig. 7. In addition, more uniform current distribution results.

It is worth noting that the coolant condition in Case 2, however, results in greater temperature gradient in the membrane, as shown in Fig. 8. There is $\sim 12$ K temperature difference between the inlet and outlet areas. This temperature non-uniformity could potentially cause membrane degradation due to thermal stresses [18].

Fig. 9 displays the net water transfer coefficient through the membrane, $\alpha$, for Case 2. It can be seen that this coefficient, indicating water crossover through the membrane, varies spatially. In the inlet region of both anode and...
cathode, the anode loses water to the cathode. Once the cathode gas gains moisture from product water, the net water flow is directed from the cathode to anode due to dominance of water back-diffusion.

4. Conclusions

Ultra large-scale simulations of a commercial-size PEFC with 24-channel 3-pass flowfield have been carried out to demonstrate the computational performance of a parallelized code for PEFCs, and to illustrate the internal phenomena and distributions of key parameters in a representative industrial cell as well as the profound coolant effects. Using a 32-processor PC cluster, we have shown a computing time of roughly 20 h for a simulation using a 23.5 million gridpoint mesh that considers comprehensive single-phase phenomena in a PEFC. In addition, the computed contours of current and water distributions indicate that low-humidity operation commonly used in automotive applications leads to membrane dryout in the inlet region and cathode flooding in the outlet region. These two seemingly conflicting issues can be simultaneously resolved by using a cooling scheme where a lower inlet coolant temperature and a small coolant flowrate are employed to create a thermal gradient along the cathode gas flow. The resulting “cold” inlet region maintains high relative humidity despite the dry gas supply, and the “hot” outlet region alleviates susceptibility of cathode flooding.

While the present paper exemplifies only one application of the ultra large-scale simulation tool, numerous other applications of the same computational intensity have been carried out routinely at this research laboratory as well as several automotive companies.
Acknowledgements

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References