

## Review Article

# Investigation of Fuel Cells under Transient (Dynamic) Conditions to Improve the Efficiency of Polymer Electrolyte Fuel Cells in Dead-Ended Anode Mode: Review Article

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## Abstract

Polymer Electrolyte Membrane Fuel Cells (PEMFCs) operating in Dead-Ended Anode (DEA) mode present a promising alternative to traditional flow-through systems by simplifying design and reducing costs. However, their efficiency and durability are challenged by transient phenomena such as water accumulation, nitrogen buildup, and carbon corrosion throughout operation. This review investigates the dynamic behavior of DEA PEMFCs under dynamic operating states, aiming to improve their efficiency. By analyzing purge cycle optimization and transient response characteristics, we identify strategies to mitigate hydrogen loss, maintain voltage stability, and extend stack lifetime. The key findings indicate that precise purge scheduling and effective water management are critical for optimizing performance, with dynamic models providing insights into time-dependent processes. This study underscores the potential of DEA PEMFCs for high-efficiency applications provided transient effects are effectively managed.

## Introduction

Proton Exchange Membrane Fuel Cells (PEMFCs) are increasingly valued for their high efficiency and low environmental impact [1,2], making them viable for use in automotive, stationary, and portable power systems. In Dead-Ended Anode (DEA) mode, hydrogen is supplied at a regulated pressure without recirculation, eliminating reliance on complex balance-of-plant (BOP) components like ejectors and humidifiers found in Flow-Through Anode (FTA) systems. This simplification enhances practicality by reducing system weight, volume, and cost. However, DEA operation introduces dynamic challenges, including the accumulation of nitrogen and liquid water in the anode channel, which leads to hydrogen starvation, voltage drops, and cathode carbon corrosion over time.

Dynamic operating states—such as startup, load changes, and purge events—significantly influence DEA PEMFC performance. Understanding these dynamics is essential for optimizing efficiency\*\* (*defined as the ratio of energy*

*output to hydrogen fuel input (Foot note)*), while minimizing degradation. Research by Chen et al. [3] explores purge cycle optimization to balance hydrogen utilization and durability, while Moçotéguy, et al. [4] examine transient responses to current steps, highlighting water management's role in performance decline. This review synthesizes these studies to investigate DEA PEMFCs under dynamic operating states, aiming to enhance efficiency through improved purge strategies and dynamic modeling. The objectives are to: (1) evaluate the impact of transient phenomena on efficiency [5], (2) assess methods for optimizing purge cycles, and (3) propose approaches to mitigate degradation in DEA mode.

## Methodology

This review draws on two seminal studies from the provided document: Chen, et al. [3] and Moçotéguy, et al. [4], and supplemented with additional peer-reviewed sources.

The methodology encompasses:

### 1. Source selection

## More Information

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The chosen articles focus on DEA PEMFC dynamics and efficiency. Chen, et al. provide a model-based approach to purge optimization, while Moçotéguy, et al. offer experimental and simulation insights into transient behavior [6].

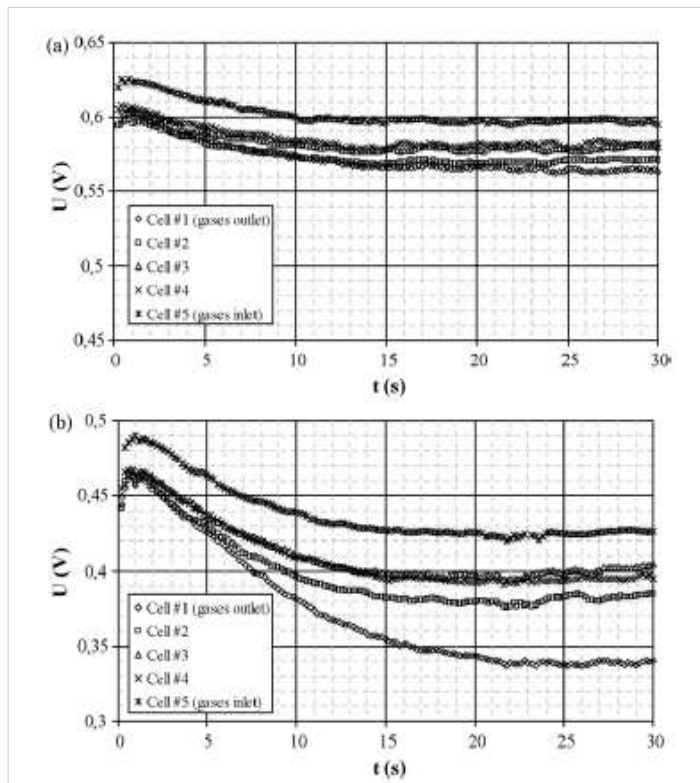
## 2. Modeling techniques

- Chen, et al. [3] developed a two-phase, 1+1D model simulates anode channel dynamics, including nitrogen and water accumulation, carbon corrosion, and purge flow (e.g., Eq. 2). Efficiency ( $\eta$ ) is calculated over multiple cycles (Eq. 1) [7], with purge intervals scanned (20-900 ms) to identify optima. The model was validated with a single-cell experiment (50 cm<sup>2</sup> MEA, 0.6 A/cm<sup>2</sup>, 50 °C).

$$\eta = \frac{\int_0^{t_{tot}} E_{cell} I A dt}{\sum_1^m \Delta h_f (Q_{i,ext} H_2 + Q_{i,loss} H_2)} \quad (1)$$

$$W_{total} = A_0 \left( C_{turb} \left[ \frac{2\Delta p}{\rho} + \left( \frac{\nu R_t}{2C_{turb} D_h} \right)^2 \right]^{0.5} \nu - \frac{R_t}{2D_h} \right) \quad (2)$$

- Moçotéguy, et al. [2] used a one-dimensional dynamic model to predict to a current step (0 to 0.5 A/cm<sup>2</sup>), incorporating double-layer capacitance and water-induced porosity reduction (Eq. 3). Parameters like initial water content ( $s_0$ ) and equilibrium current density ( $j_{eq}$ ) were fitted to data from a five-cell stack (300cm<sup>2</sup>, 288K) (Figure 1).



**Figure 1:** Evolution of each cell voltage after a current step from OCV: (a) fresh stack and (b) aged stack.

$$\varepsilon = \varepsilon_{dry} (1 - s) \quad (3)$$

## Experimental setup

- Chen, et al. measured the evolution of cell voltage and liquid water mass in a single cell with periodic purges triggered at 0.4 V, using high current (0.6 A/cm<sup>2</sup>) and 100% cathode RH to induce water accumulation.
- Moçotéguy, et al. applied a 150 A current step to a stack, recording voltages at 2000 Hz (first second) and 5Hz (1-30s), comparing fresh and aged states.

## 3. Analysis approach

Transient effects on efficiency, hydrogen loss, and degradation were quantified. Time constants, voltage stability, and water management strategies were compared to identify efficiency-enhancing measures.

## Results

The results are organized into two main themes: transient effects on purge optimization and dynamic responses under load changes.

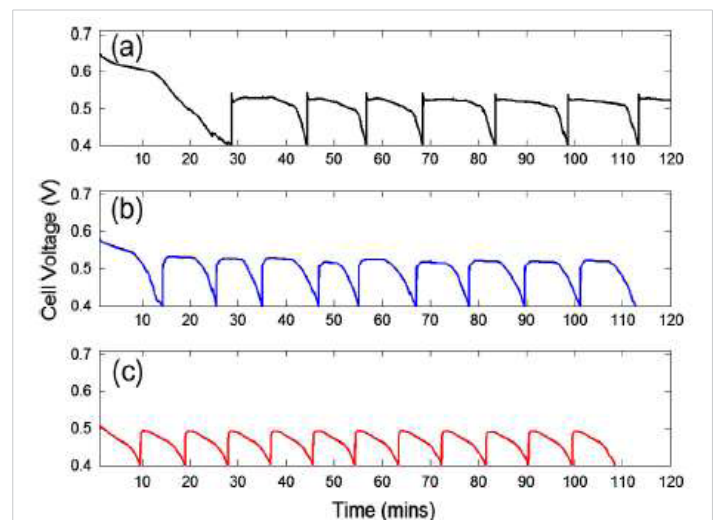
### Transient Effects on Purge Optimization [8].

#### Purge cycle dynamics

Voltage exhibited cyclic behavior with purges, dropping due to nitrogen and water buildup and recovering post-purge. Efficiency peaked within a target purge interval range (e.g., 20-54 ms for 900 s cycles) (Figure 2), declining beyond due to excessive hydrogen loss.

#### Cycle duration impact

Short cycles (< 600 s) reduced carbon corrosion but



**Figure 2:** Cell voltage variations of the single cell with a dead-enabled anode operated at different constant current densities: (a) 0.4 A cm<sup>-2</sup>; (b) 0.6 A cm<sup>-2</sup>; and (c) 0.8 A cm<sup>-2</sup>

increased hydrogen waste [10], while longer cycles ( $> 900$  s) lowered corrosion rates after a peak cathode potential ( $\sim 650$  s) but reduced voltage output (Figure 3).

#### - Hydrogen loss

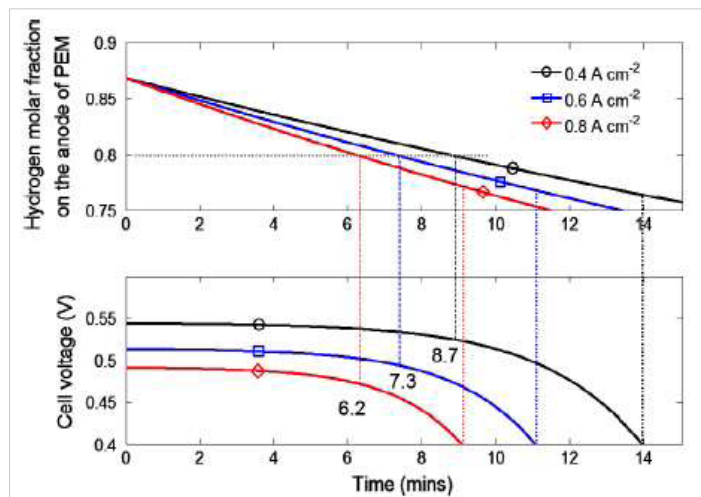
Purge flow, modeled as plug flow [12] (Eq.2), showed minimal hydrogen loss at the minimum interval ( $\delta t_1$ ) and higher loss at the maximum ( $\delta t_2$ ), affecting efficiency (Eq. 2).

#### - Degradation

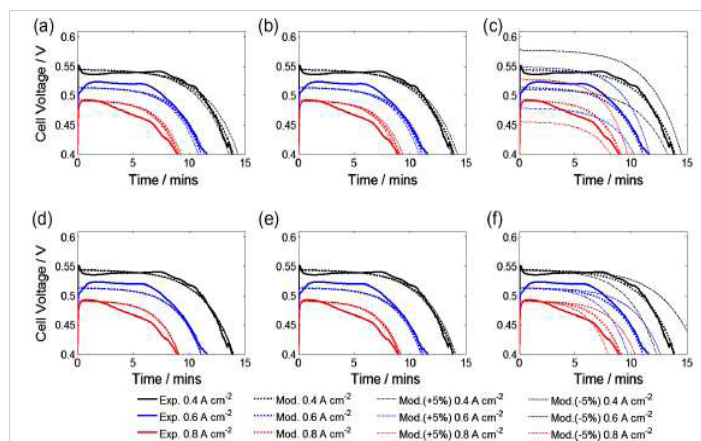
After 150 cycles (900 s each), an irreversible 10 mV voltage drop occurred [13], linked to  $\sim 25\%$  carbon loss at the channel end (Figure 4), driven by transient hydrogen starvation.

### Dynamic responses under load changes [2]

- **Time constants:** Two responses emerged: a rapid 40 ms phase (double-layer charging) and a slower 15-20 s phase (mass transport stabilization) (Figure 5). The 40 ms response was consistent across fresh and aged stacks, while the 15-20 s phase extended with ageing.



**Figure 3:** Simulation of hydrogen molar fractions on the anode side of the membrane and corresponding cell voltages at selected current densities over a defined time frame. The results highlight the impact of current density on hydrogen distribution and cell performance, providing insights into optimizing operational conditions for enhanced system efficiency.

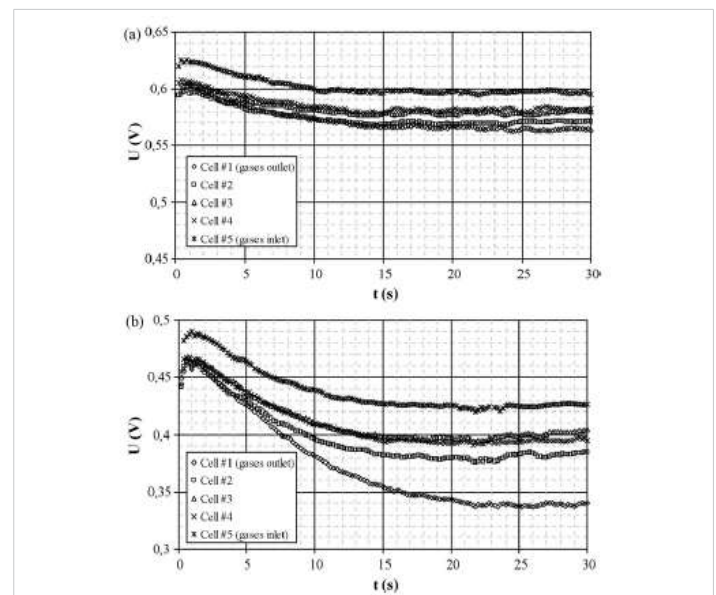


**Figure 4:** Sensitivity analysis of modeling parameters.

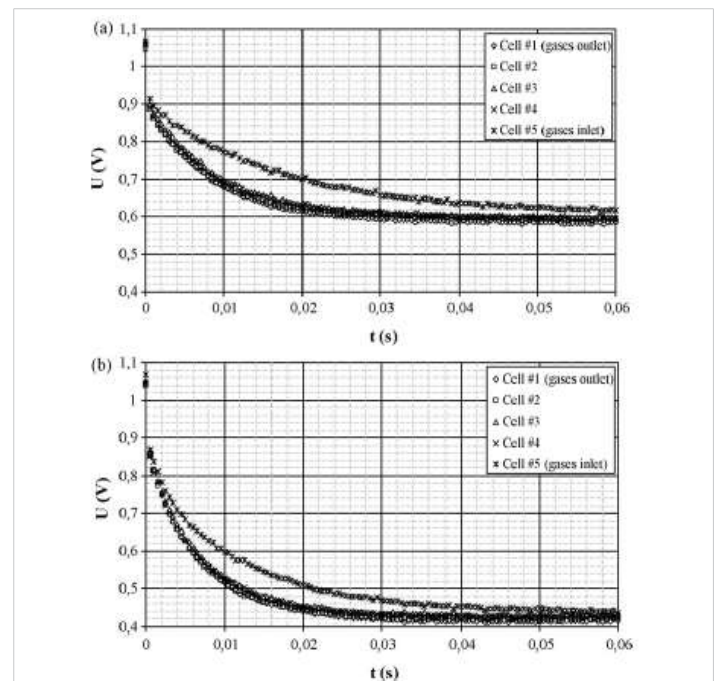
- **Voltage evolution:** Fresh stacks showed a 10 mV overshoot and 0.03 V cell variance, increasing to 24 mV and 0.08 V in aged stacks (Figure 6). Steady-state voltage stabilized after 15-20 s, with outlet cells lagging due to water buildup.

- **Water effects:** High initial water content ( $s_0 = 82\% - 85\%$ ) rose with ageing, reducing GDL porosity and oxygen pressure (Figure 3), dropping  $j_{eq}$  from  $110 \text{ A/m}^2$  (fresh) to  $10 \text{ A/m}^2$  (aged).

- **Oxygen limitation:** Interfacial oxygen pressure fell rapidly (e.g., to  $5 \times 10^4 \text{ Pa}$  in 7-19 s), indicating mass transfer constraints exacerbated by transient water accumulation.



**Figure 5:** Evolution of each cell voltage after a current step from OCV: (a) Fresh stack; and (b) Aged stack.



**Figure 6:** Evolution of each cell voltage after a current step from OCV: (a) Fresh stack and (b) Aged stack.

## Discussion

The transient behavior of DEA PEMFCs reveals critical insights into efficiency optimization:

### 1. Purge cycle dynamics

- Chen, et al. highlight that transient accumulations of nitrogen and water drive voltage instability, necessitating purges. Optimal purge intervals (e.g., 20-54 ms) balance hydrogen retention and channel clearing, enhancing efficiency ( $\eta$ ). However, frequent purges increase hydrogen loss, while infrequent ones risk starvation and corrosion, aligning with automotive lifetime targets ( $\sim 5000$  h).

- The non-monotonic efficiency trend (Figure 7) suggests dynamic control of purge timing—adjusting intervals based on real-time voltage or gas composition—could further improve performance.

### 2. Transient load responses

- Moçotéguy, et al. demonstrate that rapid electrochemical responses (40 ms) are robust, but slower mass transport processes (15-20 s) are sensitive to water and ageing. This dual timescale reflects the interplay of capacitance and diffusion limitations. These factors are especially critical under transient loads, such as during startup or power surges.

- Water's dominance in DEA mode contrasts with FTA systems, where membrane hydration is a primary concern. The high  $s_0$  (82% - 85%) indicates flooding as the key transient challenge, reducing oxygen access and efficiency.

### 3. Efficiency enhancement strategies

- Chen, et al.'s purge optimization, combined with Moçotéguy, et al.'s water management insights, suggests a hybrid approach. This includes short, precise purges to clear

anode accumulations, paired with cathode-side strategies like increased stoichiometric ratios for effective water removal. This could stabilize voltage and reduce corrosion.

- Ageing amplifies transient effects, increasing water content and resistance [14]. Preemptive purge adjustments in aged stacks could mitigate these losses, extending operational life.

### 4. Practical implications

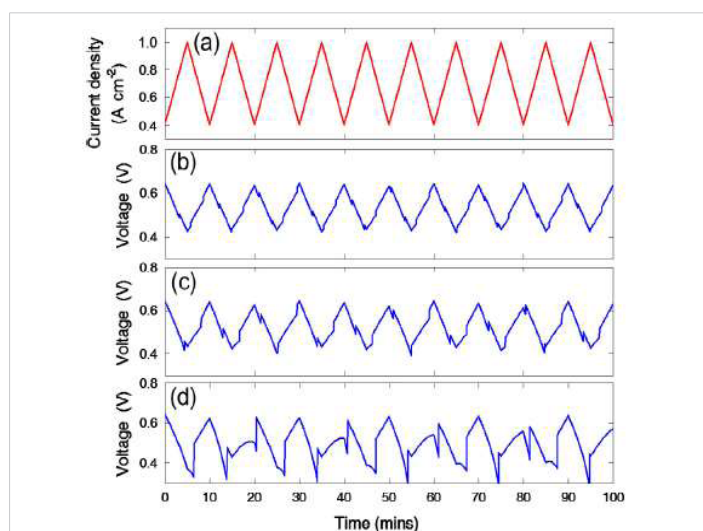
- For DEA PEMFCs to achieve high efficiency, transient phenomena must be managed dynamically. Real-time monitoring of cell voltage or gas pressure variations, coupled with adaptive purge algorithms, could optimize efficiency under varying loads.

- The reduced need for humidification in DEA mode, owing to inherent water retention, offers a cost advantage. However, controlling flooding is a key consideration for portable and automotive systems.

## Future work and research gaps

Despite the advancements reviewed in optimizing purge strategies and understanding transient responses in DEA-mode PEMFCs, several critical gaps remain that warrant focused future research:

- Real-time control algorithms:** Most current studies rely on pre-defined purge intervals and response models. There is a need for adaptive, sensor-based control algorithms that dynamically adjust purge timing and water management based on live voltage, gas concentration, and humidity data [15].
- Integration with Balance-of-Plant (BOP) components:** The interaction between DEA operation and ancillary systems such as humidifiers, cooling loops, and pressure regulators remains underexplored. Future studies should assess how BOP simplification in DEA mode affects long-term system robustness and control complexity.
- Long-term aging and durability studies:** While short-term degradation trends are established, experimental validation over thousands of hours is limited. Research is needed to evaluate degradation mechanisms under cyclic purge conditions and repeated dynamic loads over full stack lifetimes.
- Stack-level modeling and validation:** Much of the modeling work to date has been at the single-cell level. Scaling these models to multi-cell stacks, including non-uniform flow, temperature, and aging effects, is essential for realistic system integration.
- Transient response under real-world load cycles:** Current testing primarily uses step changes or



**Figure 7:** (a) Periodically varying current density applied to model; simulated voltage response to the varying current when the anode is purged at different nitrogen molar fractions: (b) 0.07; (c) 0.1; and (d) 0.16



simple current profiles. There is a gap in evaluating DEA PEMFCs under actual automotive or portable application load cycles (e.g., New European Driving Cycle or WLTP).

- f) **Water distribution imaging and mapping:** Limited studies visualize or map liquid water saturation dynamically during transient events. Advanced diagnostics like neutron imaging or X-ray tomography could help optimize purge strategies and GDL designs.
- g) **Hybrid strategies for cathode-anode coordination:** Coordinating anode purging with cathode-side water removal or oxygen stoichiometry adjustments remains a largely theoretical concept. Experimental studies on hybrid dynamic strategies could improve system efficiency without increasing purge frequency.
- h) **Hydrogen utilization efficiency vs. performance trade-offs:** Research should explore the balance between hydrogen conservation and voltage stability in more detail. Identifying optimal operating points for different applications (e.g., portable vs. vehicular) can guide purge design.
- i) **Effect of environmental conditions:** Studies rarely address how temperature, ambient humidity, or elevation affect purge dynamics and efficiency. These are critical for deploying PEMFCs across varied geographic and climatic zones.
- j) **Material-specific behavior under transient loads:** GDL, catalyst, and membrane material responses under rapid transients and long-duration purging cycles are not fully understood. Tailoring materials for DEA-specific stresses is a potential area for innovation.

## Conclusion

Numerous nations are exploring future transport strategies based on the idea that electric vehicles are the future of transportation. Moreover, according to researchers, electric vehicle sales are expected to reach 45 million vehicles per year by 2040, bringing the total global electric vehicle stock to 323 million. However, there are also challenges to the future of electric vehicles. For example, battery technology has come a long way in recent years, but for larger vehicles, the power output is still not enough.

Technically, such a future is conceivable with fuel cell vehicles capable of that uses hydrogen to produce electricity. A fuel cell vehicle could act as a generator and distributor [16], and Homes could function as energy producers and refueling stations [17]. If the hydrogen and electricity produced come from renewable sources and technologies, the amount of emissions from energy consumption for your home and car could be considered zero.

Research into advanced fuel-battery technologies that uses

hydrogen to power vehicles, without producing pollution, will be the achievement of competitive vehicles with higher efficiency, lower maintenance costs, and less pollution.

One of the subsets analyzed is the battery and its thermal efficiency. Different batteries tend to perform best in their own specific temperature range, so controlling temperature changes can affect the life and efficiency of the vehicle. Other important components include those that affect passenger comfort, such as heating, ventilation, cooling, and air purification, which can consume a lot of energy. NREL highlights the importance of optimizing these components to enhance passenger comfort with the least fuel consumption and pollution. This systems-level approach may reduce annual fuel consumption significantly, thereby reducing dependence on imported fuel and reducing pollution.

This investigation of DEA PEMFCs under non-steady-state conditions, reveals that efficiency can be significantly enhanced by addressing dynamic challenges. Chen, et al. [1] show that optimizing purge cycles—balancing intervals (e.g., 20-54 ms) and durations (e.g., 600-900 s)—mitigates hydrogen loss and carbon corrosion, achieving peak efficiency. Moçotéguy, et al. [2] indicate that transient responses (40 ms and 15-20 s) are governed by water accumulation, with ageing exacerbating mass transfer limitations. Effective strategies include precise purge scheduling and water management to prevent flooding, ensuring stable voltage and prolonged stack life. These findings support the viability of DEA PEMFCs, efficient alternatives to FTA systems, with future research needed to develop adaptive control systems for real-world transient operation [18].

One of the key issues of proton exchange membrane fuel cells is water management. This issue is not possible without considering the transport of liquid water within the cell. The conservation equations of mass, momentum, energy, ions, oxygen species and water vapor and the transport equation of liquid water in all layers of the cell including cathode and anode electrodes have been solved. A general study of the processes occurring in the cell has been carried out by providing contours of velocity, temperature, concentration of different species and saturation of liquid water, and the effect of liquid water on the efficiency of the cell has been considered. Also, the effect of the passage of reacting gases through the membrane on the efficiency of the cell has been investigated. The results show that by considering the effect of liquid water in the proton exchange membrane fuel cell, the polarization curve and the corresponding efficiency curve are lowered. The effects of several key variables such as coefficient of liquid water diffusion, current density, membrane thickness, and the oxygen-to-nitrogen ratio on the conditions and performance of the proton exchange membrane fuel cell, such as liquid water saturation, oxygen molar concentration, power density, cell temperature, transient efficiency, overall efficiency, potential, and ohmic potential drop, were investigated [6,19,20]. Liquid



water saturation at the anode was found to be significant [21], highlighting the need for further investigation into liquid water behavior at this electrode. Liquid water saturation increased at both the cathode and anode with rising current density with increasing [22] current density.

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