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Published in:
Journal of the Electrochemical Society

Link to article, DOI:
10.1149/2.0741802jes

Publication date:
2018

Document Version
Publisher's PDF, also known as Version of record

Link back to DTU Orbit

Citation (APA):

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Towards High Power Density Metal Supported Solid Oxide Fuel Cell for Mobile Applications

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For use of metal supported solid oxide fuel cell (MS-SOFC) in mobile applications it is important to reduce the thermal mass to enable fast startup, increase stack power density in terms of weight and volume and reduce costs. In the present study, we report on the effect of reducing the Technical University of Denmark (DTU) SoA MS-SOFCs support layer thickness from 313 µm gradually to 108 µm. The support layer thickness decrease in the DTU co-sintering MS-SOFC fabrication route results in an increased densification of the support layer and a slight decrease in performance. To mitigate the performance loss, two different routes for increasing the porosity of the support layer and thus performance were explored. The first route is the introduction of gas channels by puncturing of the green tape casted support layer. The second route is modification of the co-sintering profile. In summary, the cell thickness and thus weight and volume was reduced and the cell power density at 0.7 V at 700°C was increased by 46% to 1.01 W cm⁻² at a fuel utilization of 48%. All modifications were performed on a stack technological relevant cell size of 12 cm × 12 cm.

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Manuscript submitted November 29, 2017; revised manuscript received January 3, 2018. Published January 30, 2018.

Solid oxide fuel cell (SOFC) is attractive due to the excellent power generation efficiency and fuel flexibility. The conventional ceramic electrolyte and anode supported SOFC (AS-SOFC) is limited to stationary power generation applications as a result of the difficulty of quick startup and the high operating temperatures of 700 to 1000°C. If quick startup of a SOFC is realized, mobile applications may be considered. To enable this, it is important to shorten the startup time and reduce the heat cycle performance degradation and increase stack power density both in terms of weight and volume. As an approach to solving these problems, metal supported SOFCs (MS-SOFCs) have attracted attention. The development of this new generation of SOFC is currently in progress. DTU Energy’s MS-SOFC technology is based on co-sintering of laminated tape casted electrolyte, anode and support layers in a reducing atmosphere. This implies that the sintering shrinkage of the different layers must be matched sufficiently so that the mechanical stesses originating from any mismatch in sintering shrinkage of the individual layers can be absorbed by the cell structure. If the cell structure is unable to absorb the mechanical stresses, the cell will crack. Perfect matching of sintering shrinkage of the layers is practically very difficult as e.g. the electrolyte layer needs to be completely dense and thus gas tight, while the anode and support layers in contrast needs to be highly porous to allow sufficient gas transport.

An overview and recent progress of MS-SOFCs have been reviewed in Ref. 1. The company Ceres Power founded in 2001 is at present the organization, which most effectively has demonstrated up scaled large cell sized >80 cm² MS-SOFC stack technology. This is followed by consortia involving the company Plansee. At last MS-SOFC stacking has been demonstrated within consortia consisting of Topsoe fuel cell (TOFC no longer in operation), Plansee, and others. Apart from the mentioned players other MS-SOFC R&D activities are predominantly performed on button cell level by various university groups. 2-11 The present MS-SOFC designs are to our knowledge not optimized for mobile applications in terms of rapid startup and volumetric and gravimetric power densities. For example, the Plansee design consist of very thick substrates of 1 mm,12,13 whereas exact specification of the Ceres Power cell and stack design is shrouded in some uncertainty. However, from early publication within Ceres Power’s relatively long history it is possible to get a substrate thickness of 300 µm from a modelling study in 2005.14 The performance of the different MS-SOFC designs are for various reasons difficult to compare for among other reasons very different aimed operation temperatures e.g. Plansee 800°C and DTU 650–700°C. Ceres Power is aimed at an even lower temperature of 500–600°C due to the use of a doped ceria based electrolyte layer. Doped ceria based electrolyte is a mixed ionic and electronic conductor (MIEC). The MIEC behavior increases with increasing temperature and results in a partial short circuit. This is observed as a lowered open circuit voltage (OCV) compared to the theoretical value. It furthermore complicates performance evaluation and comparison as the degree of electronic conduction of the doped ceria electrolyte is polarization dependent. Nonetheless, to overcome the issues of electrolyte MIEC behavior, Ceres Power has recently introduced a so-called electronic blocking layer.15 The result in a 3 layered electrolyte configuration. How such further complications to the cell design affects the overall stack cost is unclear.

It is the aim of the present study to explore the feasibility of reducing the metal support (MS) layer thickness and for improved performance explore the feasibility of opening up the support layer. For opening up the support layer two routes are explored. The first route is introduction of gas channels via puncturing of the green tape casted support layer. The second route is modification of the sintering profile. Realizing this may serve as a means of enabling faster startup and increase the stack power density in terms of weight and volume.

Experimental

Cell fabrication.—The half-cell (i.e. metal support, anode, electrolyte) processing and infiltration route, described in Ref. 16,17, was also used in this study. The processes involved tape casting of the layers: metal-support (a ferritic stainless Fe22Cr steel alloy), cermet backbone (with 0–50 vol.% Y-doped doped ZrO2 with respect to metal), and ScYSZ electrolyte, followed by a subsequent co-sintering of these layers in a reducing atmosphere (H2/Ar). The electrocatalytically active phase, comprising a precursor solution of Ce0.8Gd0.2O1.9 and Ni (hereafter referred to as Ni:CGO), was infiltrated as a next step. The infiltration of electrocatalysts followed the same procedure as described elsewhere.15 Cells being evaluated for electrochemical performance were deposited with a Ce0.9Gd0.1O1.95 inter-diffusion barrier layer on the electrolyte, using the physical vapor deposition technique as described in Ref. 16. The electrocatalytically active phase, comprising of (La0.8Sr0.2)0.95Co0.8Fe0.2O3-δ (LSC) was applied by screen printing as the last component. For introduction of gas channels into the MS-SOFC support layer, the green tape casted support layer was rolled to various degrees with a spiked roller. The increasing porosity of the support layer was achieved by a 5 fold reduction of the dwell time at the sintering temperature T max.

Electrochemical characterization.—The single cells were cut out of large 12 cm × 12 cm cells and had a 54 mm × 54 mm cell area with
an active area of 16 cm² (defined by the screen-printed cathode layer). The cells were tested in an alumina housing used for conventional anode-supported cells, and the test house and positions of voltage probes and current pick-up points were as described in Ref. 19. A flat Ni net was used as a current collector on the anode side, and Au net was used on the cathode side. The cell rested directly on the flat Ni net with the edges being sealed with glass. The fuel gas was distributed from one side to the other via milled gas trenches in a 1 mm thick Ni block, which was embedded in the alumina test housing. On top of this arrangement, on the cathode side, an alumina block with the flat current collecting Au nets was put and a weight of 4 kg was applied. No sealing was used on the cathode side since the gas was distributed from the middle of the cathode and outwards via milled gas trenches in the alumina top block. The cells were heated to 800 °C for 5 h in order to seal the cells and sinter the cathode before the performance and durability tests were started. During startup air was supplied to the cathode and dry 9% H₂ in N₂ to the anode side (to prevent corrosion of the metal support at the elevated temperature).

Polarization curves and impedance data were collected in the temperature range 650–750 °C with 4% and 20% humidified H₂ on the anode side, and air or oxygen on the cathode side. The fuel and air flows were maximum 1.5 and 8.75 Nl h⁻¹ cm⁻², respectively. Specified fuel utilization (FU) corrected Area Specific Resistance (ASRcorr) within the present paper was calculated as outlined in Ref. 19.

The electrochemical impedance (EIS) data was recorded using a Solartron 1260 Gain-Phase Analyser and a Solartron 1255B frequency response analyser (Solartron Instruments, Houston, Texas) with a perturbation amplitude of 60 mA. The impedance was recorded in the frequency range 96.850 kHz–0.096850 Hz with 12 points per decade. For visualization at which frequencies changes occur in the impedance spectra during testing, the method “Analysis of Difference in Impedance Spectra” (ADIS) was used along with the method “Distribution of Relaxation Times” (DRT). In the ADIS method the difference between the derivative of the real part of the impedance with respect to frequency is plotted as a function of log(frequency).

Microstructural characterization.—The microstructure of the various cell samples was investigated using polished cross-sections. The polished cross-sections were prepared by vacuum embedding the samples in Struers epoxy resin (Epofix); ground using SiC paper; polished using 6, 3 and 1 μm diamond paste, and then carbon coated to eliminate surface charging. Scanning electron microscopy (SEM) imaging with backscattered electrons was performed using a Hitachi TM1000 tabletop SEM.

**Results and Discussion**

**Effect of reducing the metal support thickness.**—The effect of reducing the half-cell metal support layer thickness is shown in Figure 1. As can be seen, the effect of reducing the support layer thickness from the standard DTU state-of-the-art (SoA) MS-SOFC thickness of ∼313 μm gradually to 108 μm is predominantly an increase in support layer densification. The cause for the densification is the sintering mismatch between the layers. The ScYSZ layer sintering starts at a lower temperature than the sintering of the anode and support layer. Thus, at reduced support layer thicknesses the electrolyte layer has a more dominating role in the overall shrinkage of the co-sintered multilayered half-cell structure. A lower support layer thickness in itself will lower the anode gas diffusion length. However, the increase in support layer densification results in a reduced porosity ε, reduced
Increased support porosity and channel connectivity will also have an effect on the electrocatalyst loading and distribution when deposited by infiltration. A higher porosity will accumulate a larger amount of liquid and thus electrocatalyst upon an infiltration cycle. For these reasons it is expected that higher loadings and thus thicker electrocatalyst coatings will be the result for highly porous microstructures with large pores. Furthermore, in the case of dense microstructures it may be difficult to deposit sufficient anode electrocatalyst by infiltration without blocking or severely reducing the gas transport properties of the support layer. For a broad pore size distribution, there seems to be a tendency that there is an accumulation of electrocatalyst associated with the smaller pores. Thus, the electrocatalyst is not evenly distributed within the microstructure. A possible explanation could be that the capillary forces has a more predominant effect for the smaller pores. Therefore upon water evaporation when the sample is heated after infiltration, the water within the smallest pores is the last to evaporate. Since the concentration increases as water is evaporated a larger electrocatalyst deposition will take place in association with the smallest pores. For a more even electrocatalyst deposition it therefore seems desirable with a narrow pore size distribution. However, for other purposes such as a given desirable sintering characteristics this may not be feasible. It should be noted that the effect of infiltration procedures and microstructure on electrocatalyst distribution has been considered before for example in Ref. 27.

### Table I. Gas channel density of studied cells.

<table>
<thead>
<tr>
<th>Cell</th>
<th>Density of gas channels per mm²</th>
<th>Average distance between gas channels / μm</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>2</td>
<td>710</td>
</tr>
<tr>
<td>B</td>
<td>18</td>
<td>235</td>
</tr>
</tbody>
</table>

Figure 2. Sketch of envisioned consequences of increased support layer densification.
Figure 4. Electrochemical characteristics at 650°C and 700°C with air as oxidant and 20% humidified H2 as fuel. A) IV characteristics at 650°C and 700°C. The fuel utilization corrected area specific resistance (ASRcorr) of the cells at 650°C are as follows: ASRcorr (DTU SoA 313 μm MS) = 0.453 Ω cm², ASRcorr (700°C 175 μm MS) = 0.626 Ω cm², ASRcorr (700°C 175 μm MS with gas channels) = 0.492 Ω cm². B) The associated impedance characteristics at open circuit voltage at 650°C. The solid points indicate the frequencies 96.85 kHz, 9.685 kHz, 968.5 Hz, 96.85 Hz, 9.685 Hz, 968.5 mHz, 96.85 mHz.

Figure 5. Distribution of Relaxation Times (DRT) and Analysis of Difference in Impedance Spectra (ADIS) plots of the impedance data in Figure 4B. For the ADIS plots the impedance spectrum of the cell with 175 μm support layer has been used as reference.

Table II. Summary of impedance spectroscopy identified anode related processes.28 The given impedance arc summit frequencies are for a temperature of 650°C.

<table>
<thead>
<tr>
<th>Freq.</th>
<th>Impedance resolved processes in MS-SOFC Ni:CGO impregnated anode</th>
</tr>
</thead>
<tbody>
<tr>
<td>~500 kHz</td>
<td>Possible charge transfer of O2 at anode/electrolyte interface</td>
</tr>
<tr>
<td>~100 Hz</td>
<td>Electrochemical response of mixed ionic and electronic conducting composite anode</td>
</tr>
<tr>
<td>~30 Hz</td>
<td>Gas diffusion</td>
</tr>
<tr>
<td>~1 Hz</td>
<td>Gas conversion</td>
</tr>
</tbody>
</table>
The process is presumably somewhat overlapped with the response of the electrolyte grain boundaries. The two processes affected are both dependent on the Ni/CGO electrocatalyst coating thickness. A thicker electrocatalyst coating will improve the oxide ion conduction of the anode and thus the anode transmission line response at $\sim 100$ Hz. Furthermore, a thicker coating will make it more favorable for the oxide ion charge transfer processes at the anode/electrolyte interface to spread out and utilize a larger area. This leads to a lowering of the resistance. However, despite that the anode functional layer fabrication procedure is unchanged, it cannot be ruled out that the anode microstructure is affected by the changes in the support layer during co-sintering. A denser anode microstructure is expected to have a similar effect as a thinner electrocatalyst coating as discussed previously. The porous structure of the anode can as in porous electrode theory be modelled as an array of columns each representing a transition line. A denser anode microstructure would in this interpretation correspond to a lower density of columns/transmission lines per area. Thus, it is a simple active anode/electrolyte area affect, where both the processes at $100$ Hz and $> 50$ kHz are affected.

For comparison of performances we will from here on use 0.7 V at 700 $^\circ$C as bench marking condition for the performance of MS-SOFCs. In Figure 4 it is possible to see that the performance at bench marking conditions of DTU SoA MS-SOFC is 0.69 Wcm$^{-2}$ at a FU of 33%.

Increased support porosity via 5 fold reduction in the dwell time of the maximum sintering temperature.—In Figure 6A a cross section of a MS-SOFC half cell with a $\sim 175$ μm thick support is seen, where the porosity of the support layer has successfully been increased in comparison to the DTU SoA MS-SOFCs in Figure 1. The associated performance is shown in Figure 6B, where it is possible to see an improved performance of 0.89 Wcm$^{-2}$ at bench marking conditions @700 $^\circ$C, 0.7 V with a FU of 43%. In addition, the performance characteristics can be seen of using pure oxygen as oxidant, which results in significant performance improvements. This illustrates that there also is a clear scope for further improvement of the cathode in future cell performance development. For historical reasons and as a standard polarization curves are stopped at 0.6 V at DTU. However, there are no indications or guideline within the literature that SOFCs cannot be operated at lower potentials and thereby higher power densities. Nevertheless, it is commonly accepted that degradation rates are usually dependent on the degree of polarization/current load, temperature and used fuel utilization. To illustrate the potential of the studied MS-SOFCs the polarization curves are linearly extrapolated, which furthermore provide potential peak power densities (PPD). In Figure 7A a cross section of a MS-SOFC half cell with a $\sim 240$ μm thick support can be seen, with the most open support microstructure (highest porosity) of the studied MS-SOFCs. From Figure 7B the associated performance characteristics can be seen, which shows a significantly improved performance of 1.01 Wcm$^{-2}$ at a FU of 48%.

For comparison of performances we will from here on use 0.7 V at 700 $^\circ$C as bench marking condition for the performance of MS-SOFCs. In Figure 4 it is possible to see that the performance at bench marking conditions of DTU SoA MS-SOFC is 0.69 Wcm$^{-2}$ at a FU of 33%.

Figure 6. A) Cross section image of MS-SOFC with support layer thickness of 175 μm with a more open microstructure than DTU SoA in Figure 1. B) Electrochemical characteristics of MS-SOFC in A) with air and oxygen as oxidant and 20% humidified H$_2$ as fuel.

Figure 7. A) Cross section image of MS-SOFC with support layer thickness of 240 μm with a significantly more open microstructure than DTU SoA in Figure 1. B) Electrochemical characteristics of MS-SOFC in A) with air and oxygen as oxidant and 20% humidified H$_2$ as fuel.
The driving force for the flux $J(H_2)$ will be the concentration gradient. Figure 7 will be somewhat lower than the 1.56 W cm$^{-2}$ governed by diffusion (Fick's first law):

It was demonstrated that it was feasible to decrease the support layer thickness substantially. From a processing perspective MS-SOFCs with a 108 μm thick support is handling wise somewhat challenging, whereas MS-SOFC with a support thickness of at least 150 μm seems okay. However, an increasing densification of the support layer was observed as the thickness was lowered. To avoid this, a better sintering shrinkage match of the individual co-sintered layers need to be achieved or alternative routes for opening up of the support layer needs to be explored. In this study, it was demonstrated that it is possible to introduce gas channels in the MS-SOFCs via puncturing of the green support layer tape. From the results, it is clear that a certain density of gas channels is needed before it is possible to get an effect on the cell performance. Even with a relatively high density of gas channels with a mutual average spacing of 235 μm, gastight electrolyte layers on a large cell size of 12 cm × 12 cm were achieved. However, it is expected that when continuously increasing the gas channel density by puncturing of the green support layer tape, the sintering characteristics will be affected at some point. This change in sintering characteristics might ultimately affect the co-sintering process and thus the quality of the electrolyte layer. A further implication of the introduced gas channels is that it represents a significant broadening of the pore size distribution. This is expected to have some impact on the distribution of Ni:CGO electrocatalyst deposition via infiltration. For instance it seems reasonable to expect a somewhat higher electrocatalyst loading at or near the large pores (e.g. gas channels). The impedance and the change hereof in Figures 4B and 5 were found to be dependent on the support layer densification. This can be understood in terms of the deposited electrocatalyst coating thickness as a function support layer densification. However, a denser support microstructure is also expected to affect and limit gas transport properties as illustrated in Figure 2. The tested condition with 20% humidified $H_2$ as fuel minimizes both the gas diffusion and gas conversion resistances and are at this condition relatively insensitive to small variations. This is unlike the situations of gas mixtures with either low $H_2$ or $H_2O$. This may explain why these processes seem relatively unaffected upon the microstructural changes. However, if we for simplicity consider the $H_2$ gas flux $J(H_2)$ to be governed by diffusion (Ficks first law):

$$J(H_2) = -D_{H_2} \frac{\partial C_{H_2}}{\partial x}$$

The driving force for the flux $J(H_2)$ will be the concentration gradient through the anode and support layers, which still is high with 20% humidified $H_2$ as fuel at OCV conditions. This is unlike the SOFC technology relevant condition of high fuel utilization, where the $H_2$ concentration gradient through the anode and support layer at the fuel outlet is small and hence the $H_2$ flux may be insufficient resulting in an apparent fuel starvation. Indications of such unwanted and limiting situations have been observed for cells with a dense metal support layer. Thus, from a performance and technological perspective a more open microstructure is needed. It should be noted that for a more accurate modelling of the microstructural gas transport properties more advanced models such as e.g. the dusty gas model is needed. Since single cell testing is used for performance evaluation, the fuel utilization is not insignificant as Figures 4, 6 and 7 illustrate. This is unlike button cell testing where the cells are flushed with fuel and the effect of fuel utilization is suppressed. This is practical feasible as the active area is very small (typically ≤ 0.5 cm$^2$). In addition many variations of button cell setups exist with different gas flow configurations. In the used single cell testing, the flow configuration is similar to what will be present in a stack that is plug flow. Thus, the condition in single cell testing mimics stack conditions and the measured performance of the cell will therefore be very close to what can be expected in a stack environment. The importance and impact of fuel utilization has been considered before in Ref. 19. In here, a methodology for compensating for the fuel utilization has been suggested. An exact compensation of fuel utilization is not feasible, but estimation of a conservative first approximation is feasible. Basically, the methodology attempts to compensate for the change in Nernst potential as the fuel is utilized. Area specific resistances (ASRs) corrected by this methodology is specified as ASR$_{corr}$. In Figure 8 the effect of fuel utilization is illustrated on the performance characteristics with air as oxidant of the cell in Figure 7. Assumining linear polarization relationship the calculated ASR$_{corr}$ from the measured data has been used to forecast the effect of fuel utilization. As expected it is possible to see that the higher performance at the highest temperature of 750°C is affected the most. From the plot it is possible to see a fuel utilization corrected performance of 1.20 W cm$^{-2}$ at bench marking conditions @0.7 V, 700°C and a PPD of 1.4 W cm$^{-2}$ at 700°C. Key performance characteristics of the studied cells are for convenience summarized in Table III. Thus, MS-SOFCs with higher porosity and therefore improved gas transport properties and higher electrocatalyst loadings does lead to improved performance of the anode and therefore also the cell. Recently, M. Tucker has also shown for a MS-SOFC design with comparable electrode architectures and electrocatalysts that boosting infiltrated electrocatalyst loadings lead to significant performance improvements. The result was an impressive boosting of the corrected performance of 1.20 W cm$^{-2}$ at 700°C and 0.7 V for 1200 h showed severe degradation with a peak power density loss of 73%. A summary of reported key performances in laboratory scale, using different manufacturing process and materials is provided in Table 13 of the recent MS-SOFC review from 2017 by Krishnan. From the summary it is clear that the performances reported within the present study are superior compared to the reviewed studies in Ref. 1 in particular if temperature is taken into consideration. From the present data it is uncertain how much the anode can be further improved. In any case, there is for even further performance improvements a clear potential for the cathode to be further optimized considering the effect on performance of varying the oxidant gas between air and pure oxygen in both Figures 6B and 7B. At last the electrolyte layer can be further densified and made thinner and thereby additionally boost the performance.

Figure 8. Fuel utilization corrected polarization and power density curves as outlined in Ref. 4.
It was demonstrated possible in the DTU MS-SOFC design and fabrication route to reduce the metal support considerably from standard 313 μm to 108 μm. However, upon the reduction of the metal support layer thickness an increased densification of the layer and a supported loss was observed. To mitigate the performance loss, two routes for increasing the porosity of the support layer was demonstrated possible. The first route was introduction of gas channels via puncturing of the green tape casted support layer. In the case of an average gas channel spacing of 235 μm on MS-SOFCs with a support thickness of 175 μm a significant performance improvement was observed. The second route was a 5 fold reduction of the sintering temperature dwell time, which lead to significant increase of the support porosity. This resulted in significant performance improvements with a maximum increase of 46% in performance to 1.01 Wcm⁻² at 0.7 V, 700 °C with a FU of 48%. This corresponds to fuel utilization corrected peak power densities of 0.84 Wcm⁻² at 650 °C. 1.40 Wcm⁻² at 700 °C and 1.90 Wcm⁻² at 750 °C. All results were obtained on gastight stack technology relevant sized cells of 12 cm × 12 cm.

Acknowledgment

Financial support by NISSAN MOTOR CO., LTD. is gratefully acknowledged.

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