

PAPER

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Metal nanoparticles support materials play a crucial role in many fields, including energy conversion/storage, catalysis and photochemistry. Here, the exsolution is reported as an *in situ* method to fabricate metal nanoparticles supported on perovskite ($\text{La}_{0.52}\text{Ca}_{0.28}\text{Ni}_{0.06}\text{Ti}_{0.94}\text{O}_3$) powder and fiber materials. Significantly decreased polarisation resistance can be achieved by applying electrochemical switching within 3 min on the fiber electrode fuel cell to facilitate the exsolution. The fuel cell activated by electrochemical switching under wet hydrogen shows a promising performance with a maximum output power density of about 380 mW cm^{-2} at 900°C in hydrogen. The phase-field model shows that the exsolution under extreme low oxygen partial pressure induced by electrochemical switching performs faster nucleation than the chemical-reduced case. This work provides a further understanding of electrochemically driven exsolution with fiber structure platform and simulation with phase-field models.

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10th Anniversary Statement

I have been a strong supporter of the *Journal of Materials Chemistry* as a strong vehicle for advancing *Materials Chemistry* as a core *Chemistry Discipline* since its inception by my colleague and mentor Tony West in Aberdeen. I served as an editorial board member before the journal's subdivision and am honoured when my articles are published in this key journal.

1 Introduction

Supported nanoparticle materials received much attention over the last decades due to their prominent roles in many fields of modern industry.^{1–3} A recently studied cost- and time-efficient method named exsolution was widely investigated to fabricate fine and embedded metal nanoparticle decorated materials.^{4–6} The exsolution materials have been utilized in catalysis,^{7,8} solid oxide fuel/electrolysis cells,^{9,10} and membrane reactors.^{11,12} The number of applications based on exsolved functional materials greatly increases across multiple areas due to their compositional, structural and functional tunability.^{13–16}

The exsolution process can be simplified into three steps: diffusion, nucleation, and growth.^{4,6} The diffusion of cation is typically determined by the concentration and mobility of charges, thus could be affected by the crystal structure, geometry (mainly surface) and environment.^{6,17} The nucleation and growth are tuned by the free energy,¹⁸ temperature¹⁹ and other parameters.²⁰ In general, the exsolution could be driven by the reduction,²¹ cation deficiency,²² strain²³ and electrochemical switching.²⁴ Among different driving forces, electrochemical switching by applying bias has been proven as an effective way to facilitate exsolution, enabling fast exsolution (within minutes) with homogeneous particle distribution compared with chemical treatment by hydrogen reduction.^{9,25,26} However, the process of electrochemical switching has been employed on few system and not been fully understood.²⁷

Moreover, systems with high surface may afford larger interactive surfaces while guaranteeing the supply of cations, such as ordered microporous frame, thin film and fibers.^{19,28,29} The solid-state reaction was widely used to synthesis perovskite oxides by calcination at high temperature, for examples, above 1400°C . The bulk or coarse powder displays a pure phase but low surface area due to the sintering under such a high calcination temperature. Differentiating from bulk samples and nano-powder, nanofibers offer a continuous structure to

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provide a unique pathway for the diffusion of ions.³⁰ The hierarchical, porous and less tortuous pathway in the fiber electrode may contribute to a fast mass and charge transport.³¹ Besides, the nanofiber provides higher surface area which affords more active sites and resulting in better catalytic activity.³² The one-dimensional structure may provide a way to understand the diffusion of cations/vacancies in the support materials contributing to their unique structure and morphology, for example the surface effect and the continuity of support materials.³³

Herein, the exsolution process of Ni particles was studied on both powder and fiber structure with the same composition, $\text{La}_{0.52}\text{Ca}_{0.28}\text{Ni}_{0.06}\text{Ti}_{0.94}\text{O}_3$ (LCNT), to illustrate the surface and structure effects. The 20% A-site defect was designed to achieve oxygen stoichiometry to understand the influence of exsolution on the formation of oxygen deficiency.²² The results demonstrate that the fiber performed a faster exsolution process than the powder with hydrogen reduction. Furthermore, the electrochemical switching possesses an efficient exsolution process on the fiber electrode in wet H_2 solid oxide fuel cell (SOFC) with the polarization resistance decreased from 6.02 to 0.69 $\Omega \text{ cm}^2$. The phase-field model (thermodynamic function) shows that the extremely low $p\text{O}_2$ achieved through electrochemical switching provokes instant nucleation, thus facilitating particle exsolution.

2 Results and discussion

2.1. Characterization of Ni exsolution

The crystal structure evolution for perovskite samples before and after hydrogen reduction was examined during the exsolution. The LCNT powder samples were obtained by grinding the pellet prepared by solid-state reaction (SSR), while the fiber samples were prepared by electrospinning to achieve a continuous nano structure. Fig. 1(a) displays the XRD patterns of the powder and fiber samples, showing a similar perovskite

structure. The crystal structure of the perovskite was investigated based on Rietveld refinement with GSAS II,³⁴ revealing quite similar unit cell parameters (Fig. S1 and Table S2†). Indexing on a double cubic cell reveals that both (400) and (404) are single peaks without broadening. Weak reflections at (211) and (301) indicative of out-of-phase tilting are visualized from the crystal structure in Fig. 1(a). Thus, the most likely space group chosen based on the observation is tetragonal $I4/mcm$ ($a = b = 5.4769(0) \text{ \AA}$, $c = 7.7543(8) \text{ \AA}$). Besides, there are small peaks (mark with diamond) indicate the presence of NiTiO_3 (PDF No. 33-0960, rhombohedral, $R\bar{3}H$ space group).³⁵ The formation of NiTiO_3 phase can be avoided by reducing the doping amount to 0.03 (Fig. S2†).

To trigger the exsolution of Ni nanoparticles, the samples were reduced in 5% H_2/N_2 at 800 °C for 4 h. There is no obvious peak shift shown in the XRD pattern before and after reduction indicates the perovskite structure was well retained. Obvious reflection around 44.5° show up for the fiber sample reveals the emergence of metallic Ni species, as shows in Fig. 1(b). However, the metallic Ni reflection cannot be observed for powder samples from the XRD pattern. The absence of metallic Ni reflection from the XRD may contribute to small nanoparticles and low exsolved Ni amount. The powder pieces exhibit a dense surface prepared by solid-state synthesis, as shown in Fig. 2(a and c). Sparse nanoparticles with an average size of about 15.9 nm exsolved from the powder surface reduced at 800 °C. A more significant number of nanoparticles decorated on the surface can be seen from the sample reduced at 900 °C (Fig. S3†).

As the exsolved particles exhibit nano-size, thus, the micro-structure of the samples was important to evaluate the degree of exsolution. The fiber calcined at 1100 °C consists of dense grains of about 200 nm (Fig. S4†). The smooth surface of fiber from high-resolution SEM figures indicates no particles on the surface before reduction. The fiber morphology was maintained

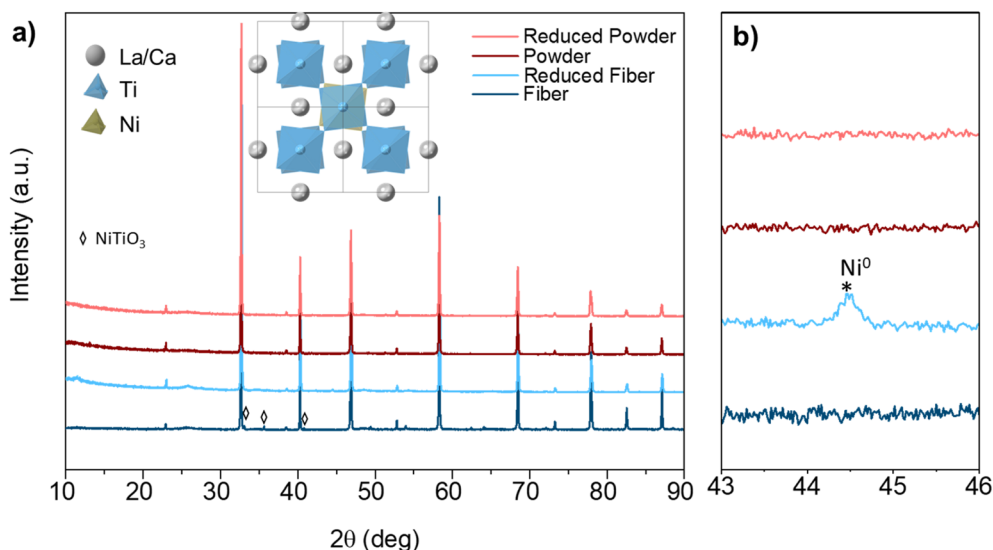


Fig. 1 Exsolution of Ni nanoparticles from LCNT perovskite materials. (a) XRD pattern of powder and fibers samples before and after reduced in 5% H_2/N_2 at 800 °C for 4 h. (b) The magnified 2θ range between 43 to 46° to illustrate the presence of Ni phase.



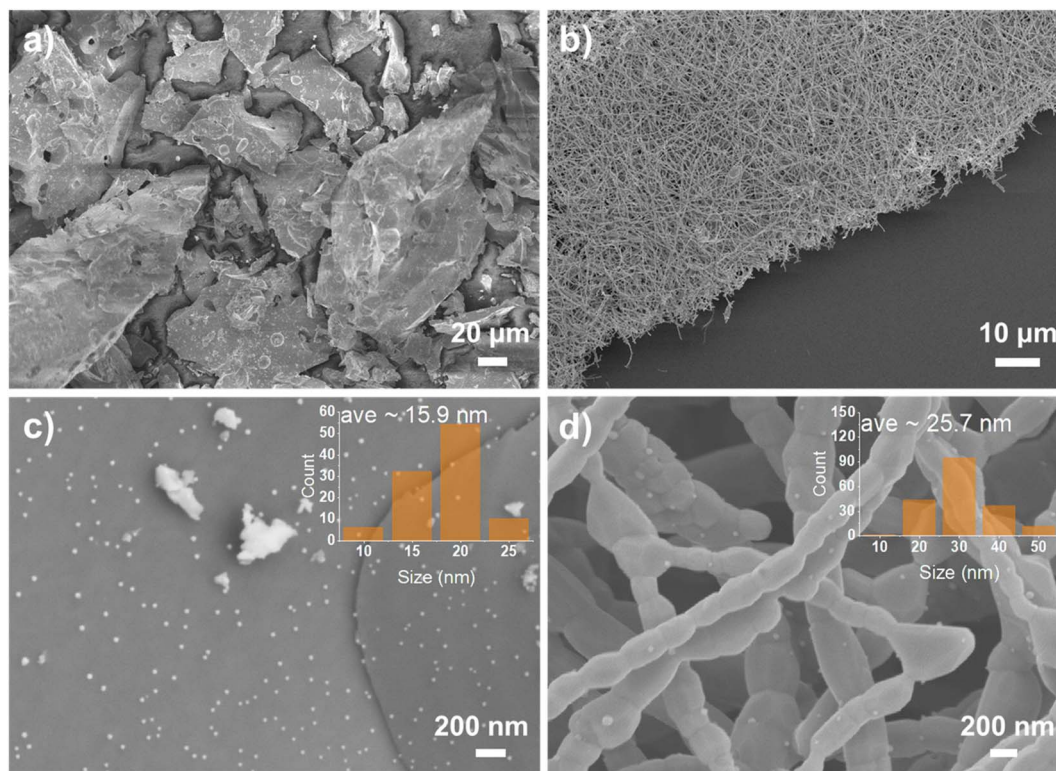


Fig. 2 Morphology of exsolution of Ni nanoparticles from LCNT perovskite materials. SEM figures of (a and c) powder and (b and d) fiber reduced in 5% H_2/N_2 at 800 °C for 4 h. The insets show the size distribution of the exsolved nanoparticles.

after reduction, as shown in Fig. 2(b). Many nanoparticles formed on the surface of the nanofibers with an average size of about 25.7 nm, as shown in Fig. 2(d). The nanoparticles on fiber can be categorized into two types based on the location at the boundaries and on the facets of grains. Most nanoparticles are located on the grain with some particles on the grain boundary in which reduced surface energy was achieved.³⁶ The nanoparticles size is larger on fibers than on pellets probably due to the lower surface energy and the short diffusion distance for the cations in lattice. The growth of exsolved nanoparticles may also benefit from the accumulation of cation diffusion through periphery grains.³⁷

The Ni exsolution process of the fiber can be demonstrated from X-ray absorption spectra by the changes in oxidation state. Fig. 3(a) shows the X-ray absorption near-edge spectrum (XANES) at Ni K-edge for the reduced fiber samples. Significant absorption intensity suggests that Ni present in the high-oxidation state in fibers reduced at 400 and 600 °C, similar to Ni^{2+} in $\text{Ni}(\text{NO}_3)_2$ standard. XANES of the fiber sample under reduced below 600 °C almost stayed unchanged, while the sample reduced at 800 °C shows a decreased intensity for the absorption edge which reveals the formation of metallic species. However, the intensity of XANES of fiber reduced at 800 °C is still higher than the Ni foil, indicating mixed oxidation states with metallic and Ni^{2+} .

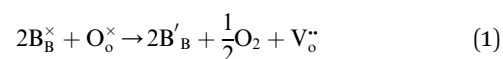
Extended X-ray absorption fine spectra (EXAFS) at Ni K-edge were used to confirm the chemical states for Ni in the reduced fiber samples, as shown in Fig. 3(b). The Fourier transforms

spectra in the R space from the k^3 -weighted EXAFS of fiber before reduction exhibits one predominant peak in the region $\sim 1.0\text{--}2.0$ Å, assigned to the Ni–O bond indicating that Ni is in octahedra environment.³⁸ The monotonous feature of samples before reduction and reduced at 400 and 600 °C consisted with $\text{Ni}(\text{NO}_3)_2$, indicating samples possessed a single coordination environment surrounding the Ni atoms. Regarding the sample reduced at 800 °C, the peak shift to a longer distance (about 2.1 Å) contributes to the arising from the Ni–Ni coordination.³⁹ A small shift distance (about 0.2 Å) to the Ni–O bond in $\text{Ni}(\text{NO}_3)_2$ revealing some nickel presents in a higher oxidation state.

The nanoparticles embedded in the fiber can be seen from the TEM image in Fig. 3(c–f). The exsolved particles are spherical, with about 20 nm in diameter, and socketed into the fiber with various depths as shown in Fig. 3(d–f) and S5.† The lattice space for the support is about 0.27 nm corresponding to the spacings of (200) planes of perovskite. Fig. 3(g) shows the selected area electron diffraction pattern of areas shown in Fig. 3(e), indicating the coexistence of the exsolved metallic Ni and the perovskite support.

2.2. Surface effect on exsolution

In reduction conditions, the oxygen in the perovskite oxide will be stripped away from the lattice followed with the exsolution of cations form perovskite:



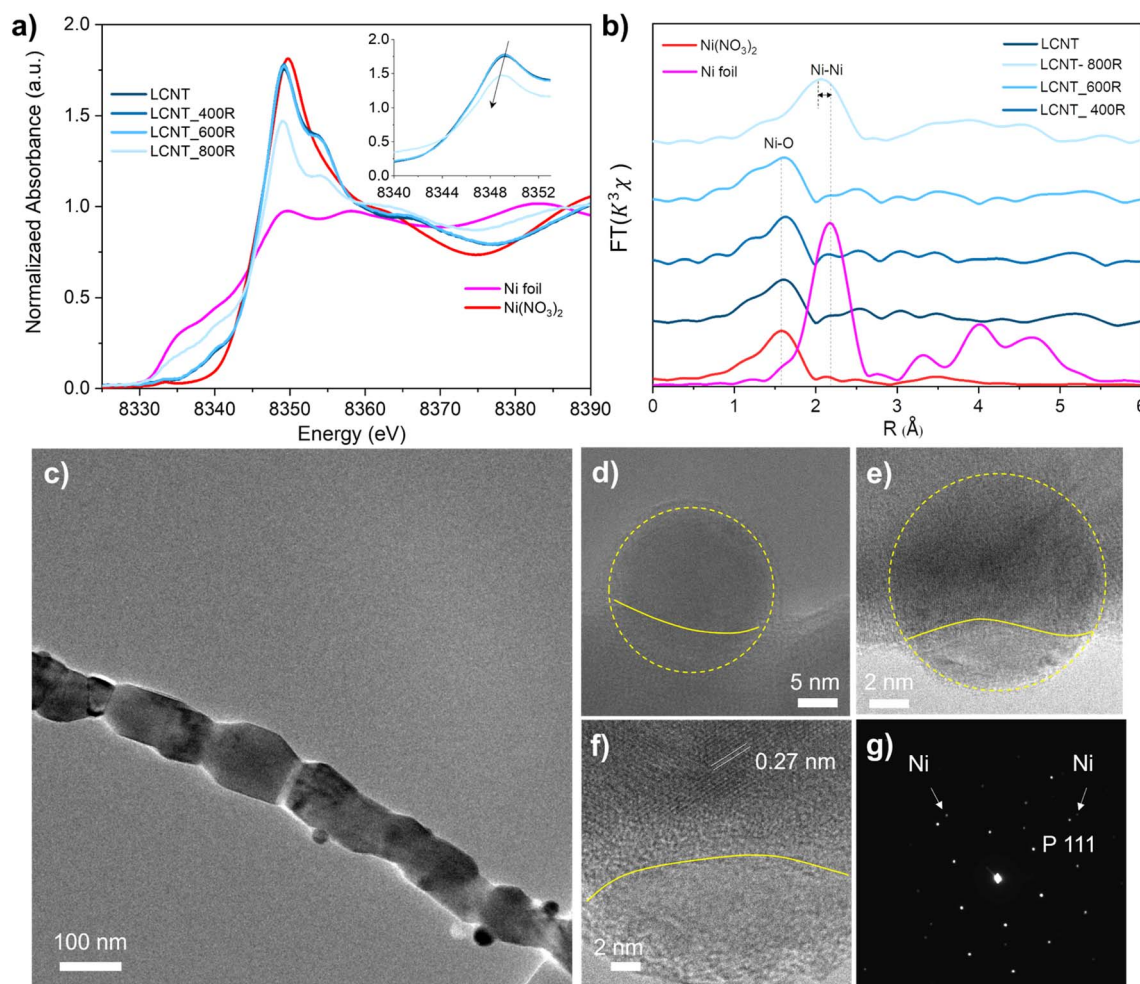


Fig. 3 Characterization and microstructure of reduced LCNT perovskite fibers. (a) Ni K-edge XANES spectra and (b) Fourier-transformed $k^3\chi$ data for fibers reduced at 400, 600, and 800 °C with standard samples of Ni foil and $\text{Ni}(\text{NO}_3)_2$. (c) TEM images of reduced fiber. (d–f) TEM image of exsolved particle, (g) the corresponding SAED pattern of (c).

where B_B^\times , O_O^\times and V_O^\bullet denote lattice cation, lattice oxygen and oxygen deficiency sites, respectively. The exsolution studied in this work can be simplified as cations diffusion in the solid followed by nucleation and growth.³⁶ The diffusion of substitutional cation corresponds to the movement of the oxygen vacancies in another direction. By studying the formation of oxygen vacancies, the proceeding of exsolution can be elucidated to some extent. To further understand the influence of the structure of perovskite samples, the thermogravimetric analysis (TGA) was performed for samples in 5% H_2/N_2 , as shown in Fig. 4(a) and S6.† The TGA with temperature curves shows the onset of weight loss at 670 and 800 °C for fibers and powder, respectively. The corresponding weight loss before dwelling at 800 °C for fibers is 0.31%, about 4.4 times higher than the powder with 0.07%. The fast weight loss for the fiber sample benefits from the presence of NiTiO_3 and also the higher specific surface due to the small grain size and connection of the individual grains. It seems that the higher surface area mainly contribute to the fast weight loss as the perovskite fiber without NiTiO_3 phase displays a fast weight loss under the same

reduction process, as shown in Fig. S7.† Moreover, it seems that the NiTiO_3 endures a quick loss weight under 5% H_2/N_2 around 800 °C, as shown in Fig. S8.†

The oxygen deficiency formed in each sample can be calculated based on the weight loss. When increase the temperature, marginally oxygen was expelled from the bulk of the sample. Oxygen deficiency for the reduced samples can be calculated for the samples according to eqn (2):

$$\delta = \frac{M_{\text{ABO}_3}}{M_{\text{O}}} \times \frac{m_{\text{ABO}_3} - m_{\text{ABO}_{3-\delta}}}{m_{\text{ABO}_3}} \quad (2)$$

After being reduced at 800 °C for 10 h, the weight change reaches a plateau indicating a sufficient reduction (Fig. S5†). The weight loss is 0.45% and 0.27% for fiber and powder samples corresponding to oxygen deficiencies of 0.05 and 0.03.

The TGA results can be used to fit the particle growth model to understand the determining factors for the exsolution on materials. The fitting result is depicted in Fig. S9† and suggests that the reactant concentration limits the rate of reduction of



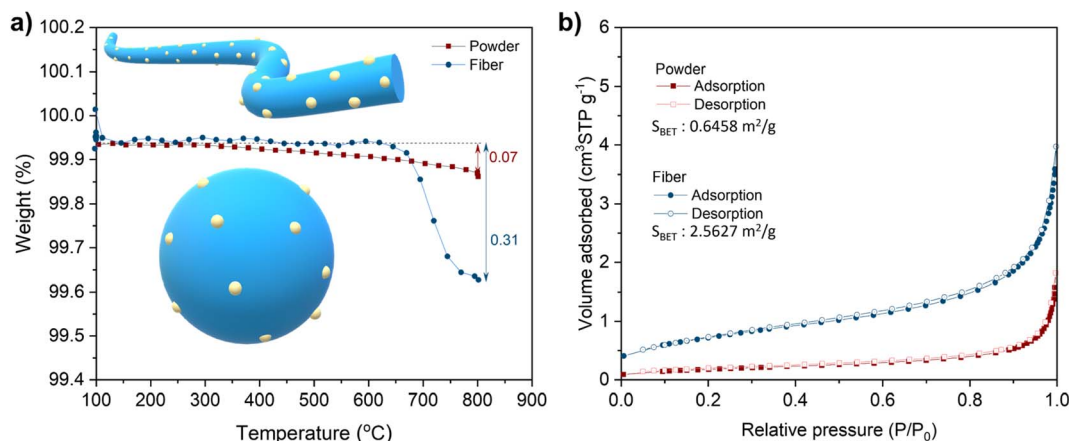


Fig. 4 Exsolution process of LCNT perovskite materials under chemical reduction. (a) Weight loss of LCNT samples reduced in 5% H_2/N_2 from 100 to 800 °C, (b) N_2 adsorption-desorption isotherms and BET surface area of samples.

the fibers. The powder reduction is too slow during the ramping step that none of the factors seems to fit the corresponding results well. When the reduction dwelled at 800 °C (shown in the range between 250–800 min), the reduction matched the diffusion-limited model. The reduction of fibers is well matched with the reactant limited model revealing that the exsolution in fibers was determined mainly by the concentration of Ni.

The difference in rate-limiting factors may due to the morphology variation of samples, such as surface area, grain size and continuity. The N_2 adsorption-desorption isotherms data for LCNT powder and fibers show that specific surface areas (S_{BET}) for samples are $0.6458 \text{ m}^2 \text{ g}^{-1}$ and $2.5627 \text{ m}^2 \text{ g}^{-1}$, respectively, as shown in Fig. 4b. The higher S_{BET} comes from the nano-grain size of fiber indicating more exposed surface areas, as shown in the SEM figures of the powder and fibers (Fig. 2). Besides, the reduction of NiTiO_3 may also contribute to this difference.

2.3. Electrochemically driven exsolution on fiber cells

Above discussions show that the fiber sample exhibits a fast exsolution, thus fiber fuel cells was built to further explored their electrochemical properties. The fiber was prepared in slurry and painted on both sides of the YSZ electrolyte to fabricate symmetric cells (LCNT fiber/YSZ/LCNT fiber). The symmetric cells were then loaded in a sealed quartz tube filled with 5% H_2/N_2 to simulate *in situ* reduction. The electrodes bond well with the electrolyte after calcined at 1100 °C on the YSZ electrolyte with an optimized thickness of about 20 μm . The woven structure was maintained and built up a rough and porous surface indicating the perovskite fibers were resistant to sintering with weaker contact between each other (Fig. S10†). A well-distributed interface was formed based on intimate contact between the electrolyte and electrode.

The impedance spectra of the symmetric cell are shown in Fig. 5. The arc at high frequency is associated with the charge-transfer process and the interface. The low-frequency arc can be assigned to the gas adsorption and dissociation at the surface of the electrode. The offset resistance (R_s) represents the

ohmic resistance that is contributed to the ionic and electronic conduction. The R_s is related to the electrode ohmic resistance, the contact resistance between electrode and electrolyte. The electronic resistance of both the YSZ electrolyte (which is generally stable in a wide range of hydrogen partial pressures) and electrodes. R_s for fiber-based symmetric cells decreased from 0.57 to $0.32 \Omega \text{ cm}^2$ after being reduced for 2 h indicating an improved electronic conductivity due to the increased electron carriers.³³

Furthermore, the R_s of the fiber-based symmetric cell can be further decreased by reducing in pure hydrogen (Fig. S11†). The spectra arc at low frequency reflects the total polarization resistance (R_p) of the electrode, including fuel adsorption and decomposition. The total polarization resistance R_p (at 0 h) decreased from $300 \Omega \text{ cm}^2$ to $150 \Omega \text{ cm}^2$ at 800 °C after a reduction in 5% H_2/N_2 (R_p after 2 h) indicating the improvement of the electrochemical kinetics of the electrode surface.⁴⁰ The summit frequency increased from 0.6 Hz to 1.6 Hz indicates the dominated process still at low-frequency which assign to the gas conversion and/or diffusion at the surface of the electrode.⁴¹ The resistance for powder based symmetric cell is comparable with fiber based electrode but slow reduction, as shown in Fig. S12.†

The activity of the electrode can be further improved by anodic polarization (refer as ‘switching’ later in this work) which facilitates exsolution in the steam.^{9,42} Electrochemically switching significantly decreases polarization resistance when applied potential from 1.8 V to 2.1 V, as shown in Fig. 5(b) and Table S3†. The R_p decreased from 6.02 to $1.10 \Omega \text{ cm}^2$ after switching at 1.8 V for 3 min which was further reduced to $0.69 \Omega \text{ cm}^2$ after switching at 2.1 V. As measured in an LSM half-cell measurement at OCV in air, it can be seen that the total polarisation resistance negligible contribution from the air electrode, as shown in Fig. S13.† The dramatic enhancement in cell performance was mainly due to the activation of fuel electrode. Moreover, the summit frequency for LCNT fiber based cell gradually increased from 200 Hz to 1262 Hz indicates the dominated reaction are mainly charge transfer process after switching.



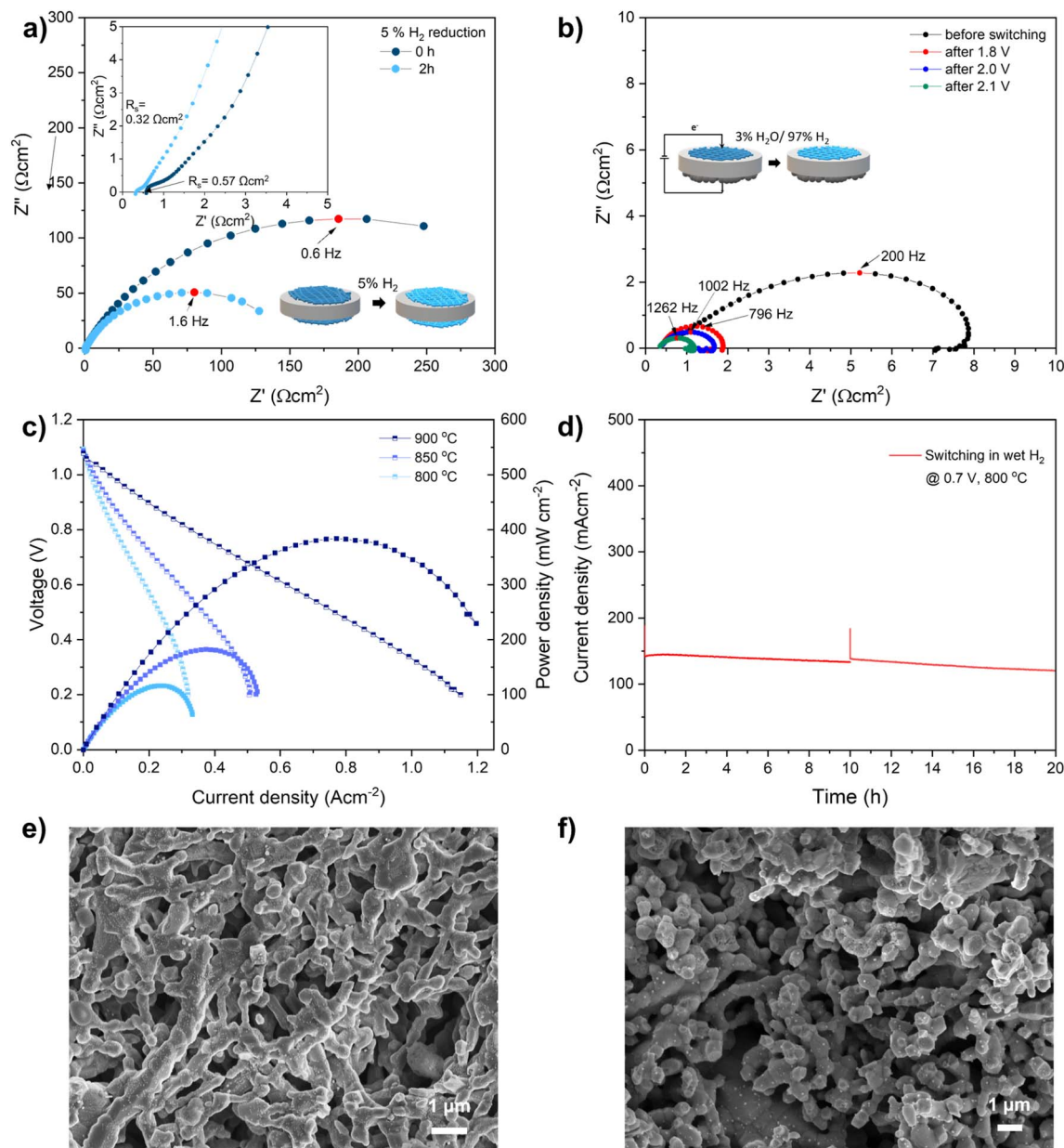


Fig. 5 Performance and short-term stability of fiber electrode-based fuel cells. (a) Impedance spectra of the symmetric cells reduced at 800 °C in 5% H₂/N₂ for 0 and 2 h. (b) Impedance spectra after switching for 180 s at 900 °C at OCV in ~3% H₂O/97% H₂. (c) current–voltage–power curves of the cell at different temperatures after switching in wet H₂. (d) Short term stability in wet H₂ for 20 h. SEM figure of (e) fiber electrode surface in symmetric cell after reduction, (f) fiber electrode in cell after stability test.

The cell reaches a maximum output power density of about 380 mW cm⁻², as shown in the IV curve in Fig. 5(c), which is compatible with the cell based on superior Zr_{0.89}Sc_{0.1}Ce_{0.01}O₂ (SSZ) electrolyte supported cell.⁴⁰ The switched fiber cell demonstrates good short-term stability by operating in wet hydrogen at 800 °C for 20 h, as shown in Fig. 5(d). The current density decreased from 142 mA cm⁻² to 120 mA cm⁻² after the stability test. The surface and cross-section morphology of the symmetric cell and whole fiber cell were examined after measurement. A number of nanoparticles emerged on the surface and the cross-section of the electrodes, seen from the SEM images of cells switching in wet H₂ (Fig. S14[†]). A careful

inspection reveals a uniform distribution of nanoparticles decorating on the surface of the fiber electrode across the whole electrode, as fine nanoparticles are readily seen on the interface area and the surface. The small and uniform nanoparticle can be seen from both the surface of the electrode and the interface close to the electrolyte after reduction, as shown in Fig. 5(e) and S14(a).[†] The exsolved particles are uniform with slightly increased size after stability test, as shown in Fig. 5(f). A careful inspection reveals a uniform distribution of nanoparticles decorating on the surface of the fiber electrode across the whole electrode area, as shown in Fig. S14(b).[†] The emergence of nanoparticles



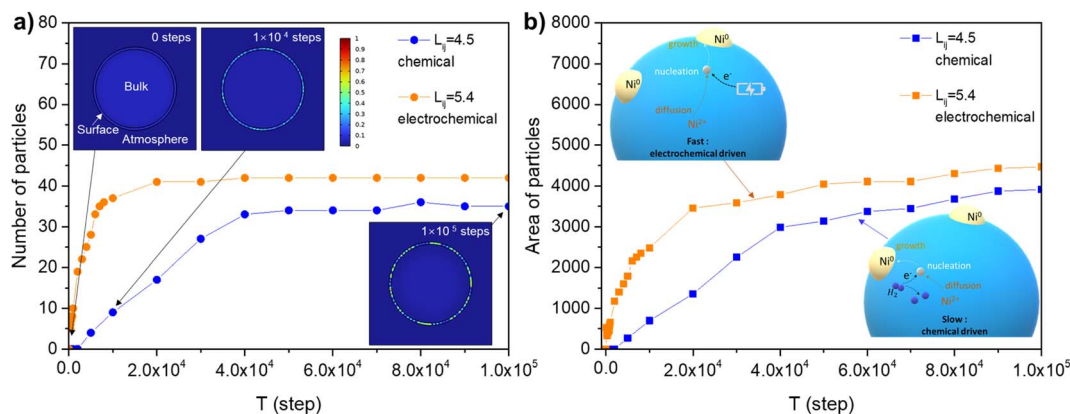


Fig. 6 Phase-field simulation results for exsolution. (a) The number of particles and the exsolution particles profile vs. steps from the model, the insets show the contour plots of exsolved particles. (b) The area of particles and schematic of bias-driven exsolution of nanoparticles from perovskite fiber cells, the insets corresponding to the exsolution process under electrochemical and chemical driven, respectively.

may contribute to the significantly reduced polarization resistance.

2.4. The phase-field model on the exsolution under different driven cases

The electrochemical reduction creates a low oxygen partial pressure (estimated to be about 10^{-35} atm from the Nernst equation) which is much higher than chemical reduction-induced exsolution. The electrochemical switching provides an swift driving force for the formation of oxygen deficiency which significantly improves the exsolution, and simultaneously promoting the electrode performance compared with hydrogen reduction.⁹ However, it is not clear why the electrode display fast exsolution under electrochemical switching. Thus, we proposed that the applied bias provides an additional driving force for the creation of defective sites which serve as nucleation sites.⁴³ To understand the exsolution process under different treatment condition, the phase-field model was used to simulate the emergence of nanoparticles under both chemical reduction and electrochemical switching (see ESI Section 1.4† for model details). The model results consist with classical development of phase transformation stages, as shown in the contour plots of x_B (molar ratio of exsolved metal) under the reference condition in the inset pictures in Fig. 6(a), which describe the microstructure evolution at 0, 1×10^4 steps and 1×10^5 steps. The implicit relationship between $p\text{O}_2$ and interaction energies between components was represented by $L_{ij}(p\text{O}_2)$. The implicit relationship between interaction parameter and $p\text{O}_2$ of $L_{ij}(p\text{O}_2)$ simulates the thermodynamic basis of the exsolution. Noted that, herein the speculated constant values were employed in this work to simplify the model. More experimental and theoretical works are needed to provide the L_{ij} for perovskite oxides.

The value of L_{ij} was set as 4.5 to achieve an oxygen partial pressure in chemical reduction situation (about 10^{-18} atm the 5% H_2/N_2). The nucleation rate reaches a plateau at 5×10^4 steps, as shown in Fig. 6(a), which consistent to our experimental observation (Fig. 4(a)). The systems experienced

different incubation states in which a metastable phase was initially reached and eventually stable (not revert to matrix) nuclei appeared. This quasi-steady state is followed by the continuous production of stable nuclei (state II in Fig. S15†).

The case with high L_{ij} (5.4) represents an electrochemical switching system in which the number of stable particles rapidly increased. The high L_{ij} was assigned to a fast transformation of metastable regions corresponding to fast oxygen extracting from the host by electrochemical switching. The number of particles for the electrochemical switching ($L_{ij} = 5.4$) system is more than chemical reduced system ($L_{ij} = 4.5$). Similar tendency can be seen from the area of emergent particles, as shown in Fig. 6(b). It is not surprising that the first incubation period is long for the system with a small interaction factor $L_{ij} = 3.4$, which indicates a higher barrier under oxidation conditions (Fig. S16†). The model results indicate a rapid nucleation of metal occurred under electrochemical switching which offers a quickly activated exsolution process. The fast nucleation was consistent with the observation at high temperatures and under high vacuum.²¹ Besides, the nucleation in the high specific surface area fiber may be more important than ion diffusion in the bulk since the ions travel in short distances and faster in the continuous pathway.^{4,44}

In the exsolution in perovskite, the formation of oxygen vacancies induces the subsequent ion diffusion and precipitating as well as the particle growth in the exsolution process. Our experimental and model results summarize that bias-driven exsolution was induced by the large $p\text{O}_2$ gradient between the host lattice and the external environment. The oxygen stoichiometry variation *via* bias potential may also induce the generation of chemical strain and interfacial stress, followed by facilitation on nanoparticle exsolution. The bias to control the oxygen stoichiometry and induce the exsolution should be further investigated with quantitative techniques.^{45,46}

3 Conclusion

This work developed a perovskite fiber catalyst as the fuel electrode of SOFC. The high surface fibers provide a special



platform for investigating the surface/morphology effect on the exsolution of nanoparticles. The weight loss of the perovskite fiber under hydrogen reduction exhibits faster exsolution than powder, an indication corresponding to the formation of more oxygen deficiencies. Moreover, the exsolution on the fiber electrode was quick triggered by electrochemically switching. The electrolyte-supported solid oxide fuel cell based on fiber electrodes shows promising performance with the maximum power density reaching 380 mW cm^{-2} at 900°C in hydrogen fuel after the electrochemically switching. Combined with the phase-field model simulation, we proposed that fast nucleation during electrochemical switching may contribute to extraordinary exsolution performance.

Author contributions

M. X.: designed and carried out the experiments and wrote the manuscript; R. C: performed the modelling calculation; S. T. W.: TEM; D. C: funding acquisition and wrote the manuscript; J. T. S. I.: funding acquisition and resources, and wrote the manuscript. All authors discussed the results and commented on the manuscript.

Conflicts of interest

There are no conflicts to declare.

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