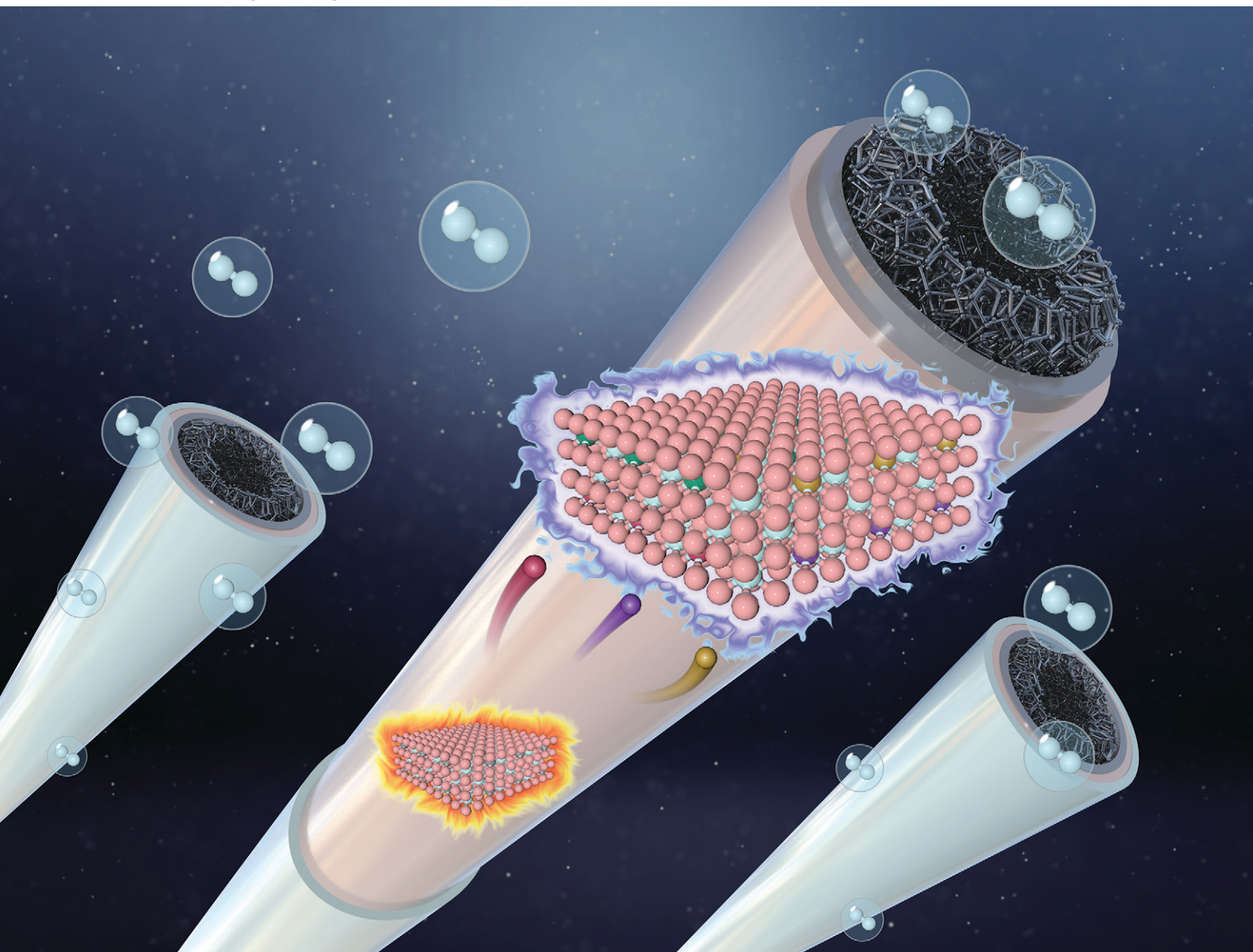


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Multi-doped ceria-based composite as a promising low-temperature electrolyte with enhanced ionic conductivity for steam electrolysis†

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Steam electrolysis is one of the most efficient approaches for producing green hydrogen. This method is based on the application of solid oxide electrolysis cells (SOECs) fabricated from functional ceramic composites for water splitting at high temperatures. Gadolinium doped ceria (GDC) is a promising electrolyte material for the fabrication of SOECs. However, the effective sintering temperature for GDC composites is usually above 1250 °C, which makes it impossible to use conventional supporting materials like ferritic steel for stack fabrication. In this work, for the first time, we have developed a lithium–bismuth–copper co-doped GDC composite capable of sintering at ~750 °C. The physicochemical and electrochemical characteristics of the co-doped GDC electrolyte were systematically analysed using thermogravimetric analysis (TG/DTA), Raman spectroscopy, SEM/EDX, XRD, EIS, XPS and dilatometry analysis. The fabricated electrolyte pellets sintered at 750 °C for 6 hours in an inert atmosphere (argon) showed high densification, obtaining 96.70% relative density. Also, the electrical conductivity obtained for the synthesised composite $\text{Ce}_{0.712}\text{Gd}_{0.178}\text{Li}_{0.05}\text{Bi}_{0.05}\text{Cu}_{0.01}\text{O}_{1.801}$ (sintered at 950 °C for 6 h) was 29.6 mS cm^{-1} at 750 °C with activation energy as low as 0.13 eV. The result of this study helps to understand better the properties of co-doped electrolyte materials for the fabrication of more efficient steam electrolyzers for environmentally-friendly hydrogen generation.

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Design, System, Application

Gadolinium-doped ceria (GDC) is one of the most common types of polycrystalline solid-state electrolytes for the fabrication of solid-oxide electrolyzers and fuel cells operating at intermediate temperatures (500–700 °C). In this work, for the first time, we have developed a lithium–bismuth–copper co-doped GDC composite ($\text{Ce}_{0.712}\text{Gd}_{0.178}\text{Li}_{0.05}\text{Bi}_{0.05}\text{Cu}_{0.01}\text{O}_{1.801}$) capable of sintering at ~750 °C. Such a super-low sintering temperature with high ionic conductivity (29.6 mS cm^{-1} measured at 750 °C) makes it possible to apply conventional ferritic stainless steel supporting materials for the fabrication of metal-supported steam electrolyzers. The physicochemical and electrochemical characteristics of the co-doped GDC electrolyte were systematically analysed using thermogravimetric analysis (TG/DTA), Raman spectroscopy, SEM/EDX, XRD, EIS, XPS and dilatometry analysis. The fabricated electrolyte pellets sintered at 750 °C for 6 hours in an inert atmosphere (argon) showed high densification properties (96.70% relative density) and low activation energy (0.13 eV). The result of this study helps to understand better the properties of co-doped electrolyte materials for the fabrication of more efficient steam electrolyzers for environmentally friendly hydrogen generation. Steam electrolysis is one of the most efficient approaches for producing green hydrogen.

Introduction

Solid oxide electrolysis cells (SOECs) are eco-friendly hydrogen energy generation equipment that directly utilises electrical energy to produce hydrogen and other chemical products with high energy efficiency.¹ Compared with proton-exchange membrane electrolyzers (PEM) and alkaline electrolyzers

(AEM), the SOEC system shows the highest hydrogen production rate and system exergy efficiency.^{2,3} Due to environmental concerns and supply limits associated with fossil fuels, SOECs have recently gained considerable attention on a global scale.⁴ Over the past 10 to 15 years, the SOEC technology has generated significant research and advancement.⁵ According to steam electrolysis current–voltage curves, the initial performance of SOECs has improved by more than a factor of 2.5 during the last 15 years.^{6,7} The durability and degradation of SOECs also showed obvious progress.^{8,9} According to the estimation of Hauch's group, the SOEC technology is now mature enough for industrial scale-up, and this scale-up is currently proceeding quickly.¹

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SOECs are fabricated in various structural designs (electrolyte-supported, ceramic electrode-supported and metal-supported). Ceramic supports are applied in electrolyte-supported cells and ceramic electrode-supported cells. Metal-supported cells employ low-cost supporting metals and alloys.¹⁰ Compared with other structural designs, metal-supported cells show several advantages. The basic idea of metal-supported cells is to support the smallest ceramic electrode/electrolyte layers with durable, low-cost metal layers made of stainless steel.¹¹ The expensive ceramic electrochemically active layers are only as thick as required for effective electrochemical operation, lowering the cell fabrication cost and maintaining robustness. Conventional cells face changes in the physical and mechanical properties of the metal–ceramic composite electrode caused by redox cycling, which highly decreases the performance of cells.¹² In contrast, the metal-supported design offers more robust cell polarisation behaviour and improves electrochemical performance.

However, the major challenge of metal-supported cells is the sintering temperature of the electrolyte. Ceramic electrolytes require a high sintering temperature (>1000 °C) to achieve approximately full density (≥95%) to provide high ionic conductivity with no gas crossover between the electrodes. While yttria-stabilised zirconia (YSZ) and gadolinium-doped ceria (GDC) served as the most common types of solid-state electrolytes for SOECs, GDC offers a significantly better ionic conductivity at lower operating temperatures.^{13,14} GDC is considered an ideal intermediate-temperature (500–700 °C) solid electrolyte for prospective commercial applications.^{15–17} However, the average sintering temperature for GDC composites is usually above 1250 °C, which makes it impossible to use conventional ferritic stainless steel supporting materials for stack fabrication. The high operating temperature applied during the fabrication steps (e.g., sintering of the electrolyte layer) can damage the metal support structure and decrease its mechanical strength.¹⁸

Reducing the sintering temperature of electrolytes is a crucial requirement for designing metal-supported SOECs with improved thermochemical stability at high temperatures. The reduced thermal energy associated with such a low sintering temperature can also effectively contribute toward reducing the overall cost of cell manufacturing.

It is well-known that the initial connections between the GDC particles cause the creation of particle necks, followed by densification and grain growth, during the sintering process.¹⁹ Mass transport processes mainly occur at the surface and grain boundaries at lower temperatures, i.e., during the initial stage of the sintering process. Thus, the number of interactions between the particles, which strongly relies on the initial particle and pore size of the GDC powder, is crucial. However, during both the intermediate and final stages of sintering, the densification can be significantly altered by the pinning effects of porosity.²⁰ According to Herring's study, nanopowders reduce temperature sintering,

as smaller particle size allows densification to occur primarily *via* grain-boundary diffusion instead of lattice diffusion.²¹ The grain-boundary diffusion can be described *via* the flux of atoms along a grain boundary; it can be given as:²²

$$J = MC\nabla\mu$$

where J is the flux of atoms; M is the atomic mobility along the grain boundary; C is the vacancy concentration; $\nabla\mu$ is the driving force for sintering. In theory, it is possible to reduce the sintering temperature by using a sintering aid that increases one or more of these parameters. Also, if the sintering aid works, it must segregate to the grain boundaries rather than dissolve in bulk. The solubility of the sintering aid is highly affected by the material of the electrolyte.

Various methods for applying sintering aids have been reported, in which the whole sintering progress is processed in an air atmosphere.²³ In the prediction and evaluation of Nicholas's group, doping Cu, Co, Fe, Mn, Li, and Zn into GDC can decrease sintering temperature, and 3 mol% lithium-doped GDC can be sintered at 800 °C to 99% density.²² They indicated that doping with Li (lithium) as a single dopant could decrease the sintering temperature, gaining a final density of 95% at ~950 °C. Chen *et al.* applied Li₂O as a sintering aid for lab-synthesised GDC using LiNO₃ as a Li source.²⁴ According to their results, 5 mol% lithium-doped GDC with a relative density of 99.3% was achieved at a sintering temperature of 900 °C and showed a maximum shrinkage rate (−0.18) at 800 °C. Kim found that the sintering temperature of GDC was decreased from 1400 °C to 1100 °C when CuO was added as a sintering aid at levels exceeding 0.25 mol%.²⁵ Yoon's group synthesised Bi nano-doped GDC by direct sol-gel combustion, and the sintering temperature was reduced to about 1200 °C.²⁶ The composed sintering aid containing Li, Cu and Zn for GDC was investigated by Nicolle's group.²⁷ According to their results, the sintering temperature went down to 930 °C for Zn, Cu, and Li addition. Lithium (Li)-, bismuth (Bi)-, and copper (Cu)-based composites have already been reported in the literature to reduce the sintering temperature of solid oxide electrolytes.^{22,24–27} These research studies have effectively prepared highly dense oxide electrolytes with satisfactory ionic conductivity results at relatively low sintering temperatures.

However, most of the electrolytes produced from previous studies were sintered in an air atmosphere.^{28–32} Moreover, very few studies have illustrated the results of sintering behaviour and electrochemical performance for doping ceria-based composites sintered in an inert atmosphere.³³ Co-doping the common solid oxide composites with aliovalent transition metal cations is a new approach to improve the densification properties and ionic conductivity of the resulting polycrystalline electrolytes.³⁴ Considering the protection for the metal support in the fabrication process, the study of sintering behaviour in an inert atmosphere for doping ceria-based composites is necessary. Also, the physicochemical and electrochemical properties for multi-doped ceria-based



composites were rarely reported. Synergistic effects and mutual effects between multi-doping elements still lack detailed explanations and understanding of the associated mechanism, which this study aims to address.

In this work, for the first time, we have developed a lithium–bismuth–copper co-doped GDC composite capable of sintering at 750 °C in an inert atmosphere. The sintering temperature of commercial GDC was optimised by the doping of Li, Bi, and Cu. Li, Bi, and Cu-doped GDC sintered at 750 °C for 6 h achieved 96.70% relative density. A novel preparation method instead of ball milling was designed and applied for commercial GDC. The homogeneous powders of Li, Bi, Cu-doped GDC and Li, Bi-doped GDC were successfully prepared. The effects of high ratio (10%) Li and Bi doping in commercial GDC were observed. Synergistic effects and mutual effects of Li, Bi, and Cu-doping were investigated. The physicochemical characteristics of the multi-doped GDC electrolyte were analysed using thermogravimetric analysis (TG/DTA), Raman spectroscopy, field emission scanning electron microscopy (FESEM)/energy dispersive spectroscopy (EDX), and X-ray diffraction (XRD). The sintering behaviours of the calcined powders were evaluated using thermodilatometry measurements, and the electrochemical performance of multi-doped ceria-based composite sintered pellets was studied using electrochemical impedance spectroscopy (EIS) tests.

Experimental section

Reagents

GDC powder (nanopowder, containing 20 mol% gadolinium as a dopant), lithium nitrate (LiNO_3 , 99.99%), copper nitrate ($\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$, 99.99%), bismuth nitrate ($\text{Bi}(\text{NO}_3)_3 \cdot 5\text{H}_2\text{O}$, 99.99%), polyvinylpyrrolidone (PVP, average mol wt 40 000), and 2-propanol ($\geq 99.5\%$) were purchased from Sigma Aldrich (Merck) and used as precursors without further purification. All the chemicals and solvents, including 2-propanol used in this study, were of analytical grade purchased from Sigma Aldrich and used without further purification.

Synthesis method

A fabrication method without ball-milling was applied to fabricate doped GDC ceramic pellets, shown in Fig. 1. The methodologies involved in preparation for low-temperature-sintered doped GDC include drying, calcination, pressing and sintering.

Preparation of mixed ceramic powder

Commercial ceramic powder, nitrate and PVP were firstly dispersed in IPA to form the inks. For the Li, Bi-doped GDC (5LB), the proportions of GDC, lithium nitrate and bismuth nitrate were 90 mol%, 5 mol% and 5 mol%. For the Li, Bi, and Cu-doped GDC, the proportions of GDC, lithium nitrate, bismuth nitrate and copper nitrate were 90 mol%, 5 mol%, 5 mol% and 1 mol%. The electrolyte compositions

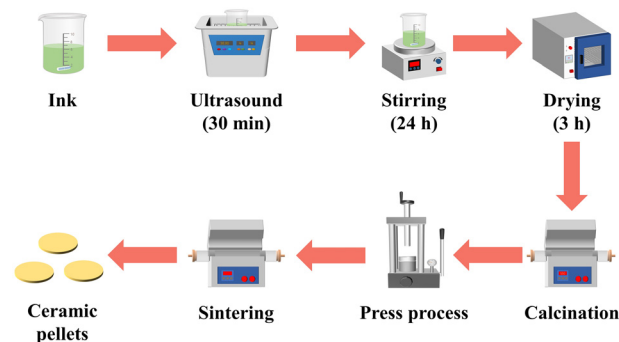


Fig. 1 Fabrication process of the doped GDC pellets.

are presented in Table 1. 3 g GDC, metal nitrates, 0.0361 g PVP and 7 mL IPA were used to prepare stable homogeneous inks *via* 30 min ultrasound and 24 h stirring at room temperature. The prepared ink was dried at 80 °C for 3 h to obtain ceramic powder.

Preparation of calcined ceramic powder

The prepared ink was dried at 80 °C for 3 h to obtain ceramic powder. The ceramic powder was calcined at 550 °C for 2 h in an air atmosphere to remove the carbon and organic components.

Preparation of calcined ceramic pellets

As shown in Table 2, several button-shape disks with a thickness of 0.3–0.5 mm were fabricated by pressing a given amount of the calcined electrolyte powders (Table 1) using a manual hydraulic press (2–10 MPa pressure) with uniaxial dies ($\phi = 3\text{--}15$ mm). The fabricated electrolyte substrates were sintered at 750, 850 and 950 °C for 6 h under a flowing argon atmosphere in a tube furnace.

Characterisation

Physicochemical characterisation

The crystallite information of the samples was studied using an X-ray powder diffractometer (PANalytical X'Pert3; Cu-K α radiation, $\lambda = 1.5406$ Å, 40 kV and 30 mA) in the range $20^\circ \leq 2\theta \leq 90^\circ$ with a scanning rate of $1.3^\circ \text{ min}^{-1}$.

The morphologies of the samples were observed *via* a field-emission scanning electron microscope (FESEM, JEOL 7100F; 15 kV, 8 A) equipped with an EDX detector (Oxford Instruments). Before the FESEM testing, the samples were attached to aluminium stubs using conductive carbon glue (TED PELLA, INC-16035 SDS), followed by sputter coating with a 10 nm layer of gold. TGA-DSC was carried out using a TA Instruments SDT-Q600 instrument (room temperature to 1000 °C at $5^\circ \text{ C min}^{-1}$ in atmospheric air). The synthesised samples were analysed by Raman spectroscopy in the range of 200–3000 cm^{-1} *via* a Thermo DXR2 spectrometer (excitation wavelength 532 nm, 8 mm optical objective 50 $\times$). An Agilent Cary 640 FTIR spectrometer was applied to detect



Table 1 Composition of the prepared electrolyte powders

Sample	Additive	Composition
5LB1C	5 mol%Li, 5 mol%Bi and 1 mol%Cu	Ce _{0.712} Gd _{0.178} Li _{0.05} Bi _{0.05} Cu _{0.01} O _{1.801}
5LB	5 mol%Li, 5 mol%Bi	Ce _{0.76} Gd _{0.19} Li _{0.03} Bi _{0.02} O _{1.85}

the infrared spectra of the samples in the range of 500–4000 cm⁻¹ at 4 cm⁻¹ resolution (32 scans). A push-rod vertical dilatometer (NETZSCH, DIL 402C) was also used to measure the linear shrinkage behaviour (room temperature to 1000 °C, with a heating rate of 2 °C min⁻¹ in atmospheric air). The apparent densities of the sintered ceramic pellets were measured using a gas pycnometer (AccuPyc II – 1345, Micromeritics).

Electrochemical characterisation

A potentiostat–galvanostat electrochemical workstation (Interface 1010E, Gamry, USA) was used to characterise the electrochemical impedance spectroscopy of the sintered ceramic pellets in the frequency range from 0.1 Hz to 2 MHz, at an AC voltage amplitude of 10 mV, and at temperatures ranging from 450 °C to 750 °C in an air atmosphere.

Results and discussion

Thermogravimetric analysis

Fig. 2 shows the thermogravimetric analysis (TGA-DSC) profiles of the 5LB1C and 5LB samples in air. As shown in Fig. 2(a), the complete decomposition of the 5LB1C powder takes place in three steps. The first step, starting from room temperature up to 130 °C, could be attributed to the dehydration of the sample with loss of the adsorbed water and other volatile compounds trapped in the solid polycrystalline structure, with a weight-loss of ~3.97%. The second weight-loss quantity (~8.25%) is observed in the range of 130–310 °C, which could be due to the thermal decomposition of the associated organic constituents of the sample and the partial decomposition of nitrate salt traces (Li, Bi, Cu).^{35–37} It is denoted that this step is associated with exothermic peaks appearing in the range of 130–310 °C, due to the combustion of the nitrate salts and the PVP residue. As can be seen in Fig. 2(a), extra peaks (140–160 °C) appear in the DSC curve for the 5LB1C powder, which can be explained by the decomposition of an additional phase

included in this sample, *i.e.*, the copper nitrate. This stage is followed by a more intense decrease of weight loss (~2.67%) within the temperature range between 310 and 600 °C that could correspond to the further decomposition of the residual organic materials and continuing decomposition and oxidation of the nitrate salt traces (*i.e.*, decomposition of Li, Bi, and Cu salts and the formation of metal oxides Li₂O, Bi₂O₃, and CuO, respectively). The pure PVP sample shows complete decomposition between 310 and 600 °C, according to the previous literature.^{38–40} The weight-loss process is stopped at ~600 °C, which corresponds to the full decomposition of lithium nitrate (500–600 °C),³⁶ bismuth nitrate (500–550 °C)³⁵ and copper nitrate (500–550 °C), respectively.³⁷ A similar trend of weight-loss behaviour is observed in Fig. 2(b) for the 5LB sample, which has been synthesised by the same method. The weight loss associated with the first step (20–160 °C) was ~5.63% for this sample, while it experienced ~6.67% of weight loss in the second step (at 160 to 310 °C), which likewise could be due to the thermal decomposition of sample's organic components (Li and Bi nitrates). In the next decomposition step (310 and 600 °C), a weight loss of 4.04% was observed. The total weight loss associated with the individual 5LB1C and 5LB samples was ~14.89% and ~16.34%, respectively. The TGA result was used to determine the calcination temperature of the electrolyte composite powders that was selected to be 550 °C in order to minimise the effects of the decomposition of nitrate salts and organic components.

Thermal expansion analysis

The linear shrinkage behaviour of the calcined powders was studied in the range from room temperature to 1200 °C. Fig. 3 shows the linear sintering shrinkage and the sintering rate of calcined powder, respectively. The calcined 5LB1C powder and calcined 5LB powder started to shrink at ~550 °C and ~500 °C, respectively. Both the GDC-based composites showed a single-step densification behaviour

Table 2 Sintering conditions for 5LB1C pellets and 5LB pellets

Sample	Conditions of sintering (°C)
5LB-1	950 (6 h)
5LB-2	850 (6 h)
5LB-3	750 (6 h)
5LB1C-1	950 (6 h)
5LB1C-2	850 (6 h)
5LB1C-3	750 (6 h)

The sintering temperature average error: ±1 °C.

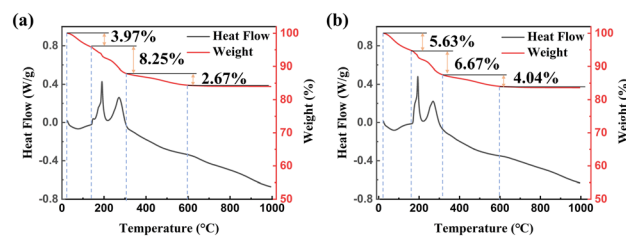


Fig. 2 TGA-DSC profiles of (a) 5LB1C powder without calcination and (b) 5LB powder without calcination.



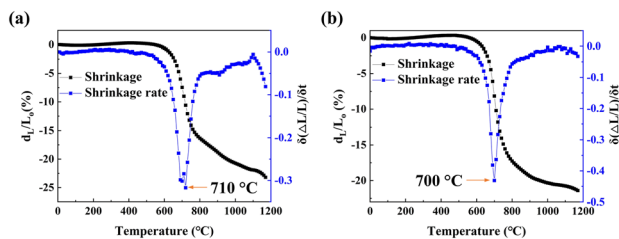


Fig. 3 The linear shrinkage and the shrinkage rate of calcined 5LB1C powder (a) and calcined 5LB powder (b).

with the maximum shrinkage rate at 710 °C and 700 °C, according to the shrinkage rate curves. The linear shrinkage profile in Fig. 3(a) shows that there was a small shrinking peak from 800 °C to 1100 °C after the main shrinking peak (550–800 °C). This could be due to the tiny evaporation of Li, Bi and Cu species from the polycrystalline solid composite.^{41–46} Fig. 3(b) shows a similar small shrinking peak from 800 °C to 1080 °C that could be similarly associated with the evaporation of Li and Bi species. A decrease in the shrinkage rate exhibited at ~1080 °C for the calcined 5LB sample could also be due to the increased rate of evaporation for Li and Bi at the elevated temperatures.⁴⁷ Also, at the end of the shrinkage process for the calcined 5LB1C sample, an obvious drop is observed (~1100 °C). The larger drop in the shrinkage rate curve from 1100 °C to 1200 °C for the calcined 5LB1C powder could be explained by the more substantial evaporation effect of Cu beyond 1050 °C compared with Li and Bi.²⁷

The results showed that multi-element substitution has a synergistic impact on Ce sites. As shown in Fig. 3, adding balanced loading fractions of Li and Bi (*i.e.*, at 5 mol%) can significantly decrease the maximum shrinkage rate of the resulting GDC composite to 700 °C, with the zero-shrinkage rate being reached at ~1080 °C. The addition of copper (sample 5LB), however, slightly increased the maximum shrinkage rate of 5LB1C to 710 °C, with a zero-shrinkage rate obtained at ~1100 °C. Therefore, the temperature range selected for sintering the electrolyte composite powders was set from 750 °C to 900 °C based on the dilatometer shrinkage analysis results. It is also evident that increasing the Li and Bi doping ratio can decrease the sintering temperature and improve the densification behaviour of the resulting electrolytes, which is in good agreement with the results obtained in the literature.⁴⁷

Structural characteristics

The prepared ceramic pellets are shown in Fig. 4. As can be seen, the colour of both samples became darker by increasing the sintering temperature. The change of colour in the samples could be due to the change of constituents. Increasing the sintering temperature enhances the evaporation rate of the compositional elements such as Li, Bi and Cu (please see the XRD elemental analysis below). The typical colours of Li₂O and Bi₂O₃ samples are white and

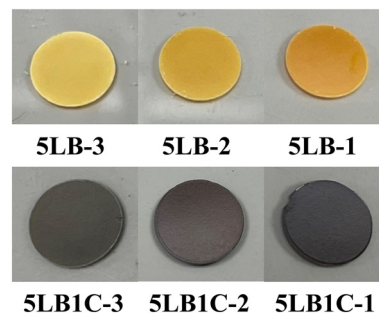


Fig. 4 Prepared ceramic pellets.

yellow, respectively. Therefore the change of colour in the 5LB sample could be attributed to the decrease of Li at the surface, which corresponds to the results of XPS characterisation. The standard colours of Cu and CuO are red-orange and black, respectively. Considering the colour change of the 5LB1C samples, this could be due to the decrease of Li and Cu and the increase of CuO at the surface of the samples, which is in accordance with the XPS characterisation results.

The XRD patterns of the sintered ceramic pellets are presented in Fig. 5. The crystal structure of GDC is calcium fluoride (CaF₂), with the space group *Fm* $\bar{3}$ *m* (no. 225). According to the reference card (ICDD data, reference code: 01-075-0162), the standard XRD patterns illustrated the presence of the (111), (200), (220), (311), (222), (400) and (331) planes of the face-centred cubic (FCC) fluorite type structure.⁴⁸ As can be seen, all the sintered ceramic pellets showed good purity, and the peaks obtained fitted well onto the reference card. In addition, the profile indicated the formation of a single crystalline phase with a cubic fluorite structure (like that of pure CeO₂), with no heterogeneous or non-soluble secondary phases in the electrolyte polycrystalline structure. The absence of the additional or shifted peaks in the XRD patterns can confirm obtaining a GDC electrolyte composite with excellent structural homogeneity with no (or insignificant) impurities. According to Fig. 5(a), the commercial GDC powder showed broad peaks, which could be due to its smaller particle size (nano-size grains).⁴⁹ The lower crystallinity of the commercial GDC powder with broader peaks indicated smaller crystal sizes.

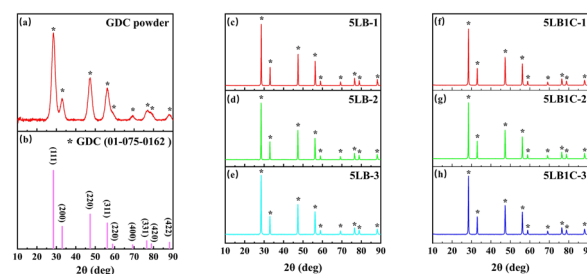


Fig. 5 XRD patterns of the standard GDC reference card (a), commercial GDC powder (b), 5LB-1 (c), 5LB-2 (d), 5LB-3 (e), 5LB1C-1 (f), 5LB1C-2 (g) and 5LB1C-3 (h).



Table S1† lists the doped GDC samples' lattice constants estimated using Bragg's equation (eqn (S1)†). Compared with pure GDC, all doped GDC samples showed distortions. Single Bi-doped GDC showed decreased lattice parameters with the promotion of the doping ratio.²⁶ The decrease in the lattice parameters with the promotion of the doping ratio also happened in single Li-doped GDC samples.⁵⁰ For the 5LB samples, increasing the sintering temperature leads to a decrease in the lattice parameter. This can be explained by applying a higher sintering temperature, which could result in more Li ions and Bi ions being doped into GDC rather than remaining in the grain boundary.^{26,50}

The tertiary doping of the Ce^{4+} ions (0.097 nm) with the Li^+ ions (0.092 nm), Bi^{3+} ions (0.117 nm), and Cu^{2+} ions (0.073 nm) in an 8-fold coordination leads to distortion, which could have different lattice parameters. The lattice parameters for 5LB1C-3, 5LB1C-2, and 5LB1C-1 were 5.430 Å, 5.431 Å and 5.423 Å. The lattice parameters of 5LB1C showed similar values at 750 °C and 850 °C. As shown in Table S1,† the samples sintered at the highest temperature (*i.e.*, 950 °C), resulted in a lower lattice parameter, confirming that more Li ions, Bi ions and Cu ions were doped into GDC leading to the decrease of lattice parameters.^{26,50,51}

According to the Scherrer equation (eqn (S2)†), the particle sizes of the samples were obtained, as shown in Table S1.† More literature data illustrated that higher sintering temperatures can increase the particle size of GDC.^{25,52} This tendency of increasing particle size matched the 5LB samples. For the 5LB samples, the particle size increased from 29.04 nm to 35.87 nm when the sintering temperature was raised from 750 °C to 950 °C. Also, the 5LB1C samples showed a continuous increase in particle size when the sintering temperature was raised from 750 °C to 950 °C.

Morphological characteristics

Fig. 6 shows the morphologies of the 5LB1C and 5LB samples at different sintering temperatures. Fig. 6(a) shows that the GDC grains are surrounded by small crystalline GDC at 750 °C for 5LB1C. Also, the dense crystalline GDC filled the gaps and pores in the grain boundaries, which

meant that the densification of GDC was enhanced even at 750 °C by doping Li, Bi and Cu. According to particle size analysis (analysed using a Nano measurer), the average particle sizes of the 5LB1C-1, 5LB1C-2 and 5LB1C-3 samples were 54.69, 74.71, and 110.54 nm. The average particle sizes on the surface of the samples increased with the increase of the sintering temperature. For the 5LB samples, the average particle sizes were 53.84, 69.15, and 98.37 nm after sintering at 750 °C, 850 °C, and 950 °C. The 5LB1C samples had larger grain sizes than the 5LB samples at the same sintering temperature. This can confirm that adding copper could enhance the grain growth in the sintering process. The results are aligned with the increasing tendency of particle size as shown in the XRD results. The smaller particle sizes estimated by the Scherrer equation (eqn (S2)†) in the XRD results could be due to the errors of the peak broadening induced by stress and amorphous forms.^{53,54} By increasing the sintering temperature, both the 5LB1C and 5LB samples showed improved crystallinity behaviour, particularly the 5LB1C-1 sample with a more apparent crystalline morphology.

Fig. 7 presents the EDX mapping results illustrating the elemental distribution at the cross-section of the 5LB1C-1 sample. As can be seen, the Bi and Cu elements are uniformly distributed into the GDC crystalline structure, which can cross-confirm the XRD analysis results. Also, a good dispersion of the Gd element is observed after the doping of Li, Bi and Cu in the resulting electrolyte composite. There were no aggregations of Li, Bi, Cu or Gd.

Table 3 shows the ratio of the elements for the 5LB1C and 5LB samples excluding Li. The Au in the samples was sourced from the Au layer by sputter coating. The carbon was mainly from the surrounding contamination. The little amount of carbon detected could come from the carbonisation of IPA and PVP. It could be seen that the ratio of Bi was kept in similar values in the 5LB1C samples when the sintering temperature increased. In the commercial powder, the proportion of Ce and Gd should be 4:1. As can be seen, the proportions of Ce and Gd in the 5LB1C and 5LB samples are similar to those of the commercial GDC powder.

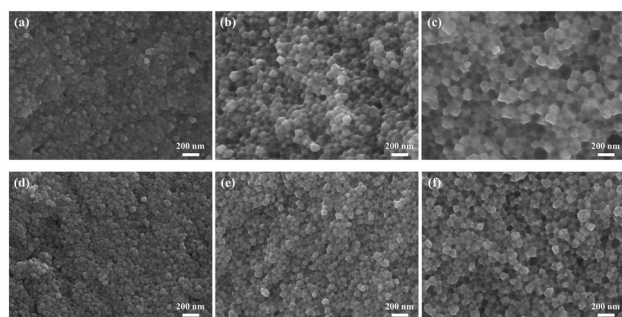


Fig. 6 Cross-sectional SEM images of 5LB1C-3 (a), 5LB1C-2 (b), 5LB1C-1 (c), 5LB-3 (d), 5LB-2 (e), 5LB-1 (f).

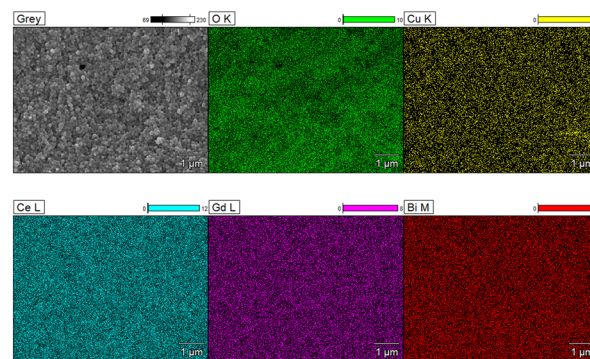


Fig. 7 EDX elemental mapping of the cross-section for the 5LB1C-1 sample.



Table 3 Ratio (atom%) of the elements from the EDX elemental mapping

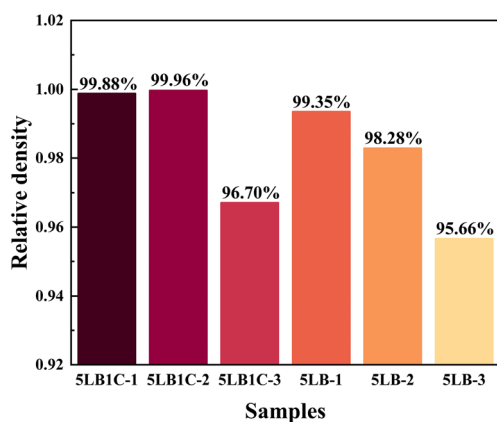
Samples	C-K	O-K	Cu-K	Ce-L	Gd-L	Au-M	Bi-M
5LB1C-3	3.8 ± 0.2	27.9 ± 0.3	0.8 ± 0.5	42.2 ± 0.4	12.7 ± 0.6	10.3 ± 0.1	2.3 ± 0.1
5LB1C-2	2.1 ± 0.1	36.0 ± 0.3	0.5 ± 0.4	41.4 ± 0.4	11.9 ± 0.5	5.8 ± 0.1	2.3 ± 0.1
5LB1C-1	2.8 ± 0.1	49.3 ± 0.4	0.4 ± 0.3	31.5 ± 0.3	9.3 ± 0.4	4.6 ± 0.1	2.2 ± 0.1
5LB-3	2.7 ± 0.1	30.7 ± 0.4	0 ± 0	42.2 ± 0.4	12.4 ± 0.6	9.6 ± 0.1	2.5 ± 0.1
5LB-2	2.3 ± 0.1	30.6 ± 0.3	0 ± 0	43.5 ± 0.4	12.7 ± 0.6	8.7 ± 0.1	2.2 ± 0.1
5LB-1	2.7 ± 0.2	28.1 ± 0.5	0 ± 0	44.9 ± 0.4	12.8 ± 0.6	9.3 ± 0.1	2.2 ± 0.1

Relative density

The densities, theoretical densities, and relative densities of the sintered pellets are shown in Fig. 8. The 5LB1C samples show 99.88%, 99.96% and 96.70% relative density at 950, 850 and 750 °C. The 5LB samples show 99.35%, 98.28% and 95.66% relative density at 950, 850 and 750 °C. The 5LB1C samples obtained the highest relative density at 850 °C, and the 5LB samples obtained the highest relative density at 950 °C. Both the 5LB1C and 5LB samples achieved full density (>95%) at 750 °C. It is also evident that the 5LB1C samples comparably achieved a higher relative density than the 5LB samples. This indicates that the addition of copper can potentially accelerate the process of densification at the same temperature.

Elemental and composition analysis

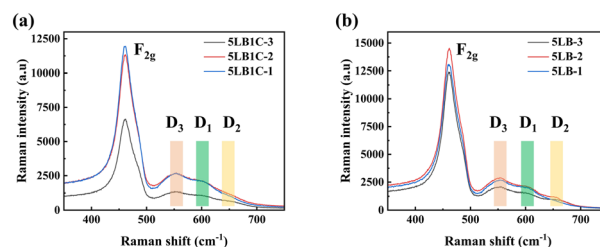
The influence of Li, Bi-doping, and Li, Bi, Cu-doping on the GDC structure and the creation of new defective sites were studied by Raman spectroscopy analysis (Fig. 9). Referring to this figure, the multiple peaks around two areas of 400–500 and 500–700 cm⁻¹ correspond to bulk ceria vibrational modes and surface defects. The most obvious sharp peak is observed at about 460 cm⁻¹ which corresponds to the F_{2g} mode of the fluorite phase in all spectra.^{55,56} This is a common characteristic of the fluorite lattice structure typical of ceria-based oxides.^{57–59} The 5LB1C-3 sample shows a relatively weaker F_{2g} peak compared to that of 5LB1C-2 and 5LB1C-1, indicating a lower level of crystallinity obtained with this sample during sintering at 750 °C.

**Fig. 8** Relative density of sintered pellets.

In a shifting and broadening range from 500 to 700 cm⁻¹, new defect-induced bands can be identified. The signal at ~550 cm⁻¹ is assigned to defect spaces formed by extrinsic oxygen vacancies.⁶⁰ The extrinsic oxygen vacancies usually appear with the presence of Ce³⁺ or other aliovalent cations.^{61,62} This vibrational feature has been labelled “D3” in Fig. 9. Another signal referred to as “D1” could be observed at ~600 cm⁻¹, which is attributed to the Frenkel defects, in which an oxygen atom has moved into an octahedral interstitial position generating a vacancy.^{63,64} This signal can also be related to the defect spaces including a dopant cation without any O²⁻ vacancy.^{65,66} A relatively weaker peak was detected between 630 and 650 cm⁻¹, labelled “D2”, which can be formed due to the extrinsic defect band, generated by the dopant addition.⁵⁶ The extrinsic defects are believed to be linked to the presence of MO₈ units without oxygen vacancies, where M is a foreign positive ion in its structure.^{56,67}

For the 5LB1C and 5LB samples, it could be seen that the strengths of the D1, D2, and D3 peaks have increased with increasing sintering temperature from 750 to 850 °C. This can confirm the enhanced level of oxygen vacancies and extrinsic defects in the samples after sintering.

The sintered samples were analysed further with XPS analysis to understand their oxidation states and electrical properties better (Fig. 10). As shown in Fig. 10a, the binding energies of Li 1s for pure Li and pure Li₂O are 55.35 and 55.60 eV, respectively, which are in good agreement with the literature data.^{68,69} The peak observed at ~55.38 eV for 5LB-3 indicates the presence of a metallic Li phase on the surface of the ceramic pellet after sintering the 5LB sample at 750 °C.^{70–72} This can be explained by the unique vaporisation mechanism of Li during the sintering process at a relatively low sintering temperature in an inert atmosphere (argon flow), which has been frequently reported in the preparation

**Fig. 9** Raman spectra of 5LB1C and 5LB samples.

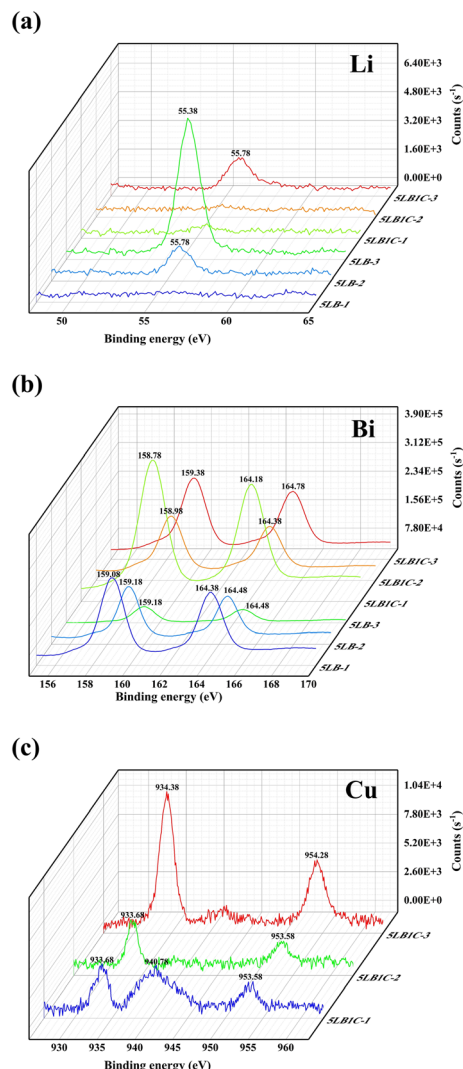


Fig. 10 Li (a) and Bi (b) of XPS elemental spectra for 5LB1C and 5LB samples; Cu (c) of XPS elemental spectra for 5LB1C samples.

of ceramic materials.^{24,73} For the Li-doped GDC sample, the generation of the Li_2O phase can also be cross-confirmed referring to its relatively higher sintering temperature ($>900^\circ\text{C}$) as shown in Fig. 3.²⁴ In our study, all samples were sintered at inert gas, and there was no oxygen source to oxidise the Li vapour from the Li_2O at the surface. The sintering temperature applied for the 5LB-3 sample was not high enough (850°C) to trigger the thermal reduction of the doped Li_2O phase into its metallic phase. In the case of the second sample (5LB2), with a sintering temperature of 850°C , no obvious peak was observed at ~ 55.38 eV. The peak at ~ 55.78 eV belongs to Li-O (Li_2O). The increased binding energy here can be due to the strong interaction with other cations.^{74,75} This implies that the metallic Li phase could be fully evaporated, leaving a small amount of Li-O (Li_2O) phase remaining at the surface. Considering the increased binding energy, Li-O (Li_2O) may also exist in the grain boundaries, and thus it cannot be readily evaporated due to its strong interaction at the interface. For the case with a sintering

temperature of 950°C , no Li 1s signal was detected. It is also noticeable that the Li 1s became weak and finally disappeared as the sintering temperature increased from 750 to 950°C . This can be due to the reduced amount of the Li phase during the sintering process for the 5LB samples. For the 5LB1C samples, only one peak at ~ 55.78 eV is detected with no obvious peaks observed for the 5LB1C-2 and 5LB1C-1 samples. This can be explained by the presence of copper in its structure which can boost the evaporation of Li. The tiny signals of Li detected in the 5LB1C-2 and 5LB1C-1 samples could be due to the diminutive residual Li-O (Li_2O).^{50,76,77}

Fig. 10b presents the XPS elemental spectra of Bi 4f. The binding energies of Bi $4f_{7/2}$ and Bi $4f_{5/2}$ are 157.1 and ~ 162.4 eV, respectively, for the elemental Bi, and ~ 159.1 and ~ 164.4 eV, respectively, in the case of Bi-O in Bi_2O_3 .^{78–81} Comparing these numbers with the locations of the peak observed for the 5LB1C and 5LB samples (Fig. 10b), the corresponding peaks could be attributed to Bi-O. Furthermore, the Bi 4f peaks associated with this oxide state have well-separated spin-orbit components (6 eV) detected at ~ 159 eV, with the symmetrical peaks indicating the absence of a reduced metal state.²⁶ For 5LB1C and 5LB, the positions of Bi peaks decreased with increasing sintering temperature. All samples showed clear Bi-O peaks, which evidenced the stable doping of the Bi phase at the surface of the electrolyte composites.

The XPS spectra of the samples containing Cu are shown in Fig. 10(c). The peaks with binding energies of 933 and 953 eV, respectively, are attributed to the $2p_{3/2}$ and $2p_{5/2}$ excitations of metallic Cu^0 , while those detected at ~ 940 eV belong to Cu^{2+} in CuO .^{82–84} Referring to Fig. 10c, only the Cu^0 peaks can be observed in 5LB1C-3 and 5LB1C-2, while these peaks became even weaker by increasing the sintering temperature from 750 to 850°C . This can be explained by the promotion of the evaporation effect for the Cu-containing samples by increasing the sintering temperature. In the case of sample 5LB1C-1 sintered at 950°C , the Cu^{2+} peak belongs to the CuO phase doped into the GDC crystal structure. Compared to the Li phase, the evaporation rate of Cu is slower, so the CuO phase would still be detectable after sintering the 5LB1C-1 sample at 950°C . The absence of the Cu^{2+} peak in 5LB1C-2 and 5LB1C-3 could be due to the thick layer of GDC formed by evaporated Cu which can cover the surface of CuO in the composite structure.

Electrochemical impedance analysis

Fig. 11(a) and (b) show the electrochemical characteristics of the 5LB1C and 5LB samples. 5LB-3 showed a relatively high total ionic resistance at 450°C , but its resistance decreased sharply at 750°C , which could be due to the high loading fraction of Li in this sample. According to the XPS results, 5LB-3 showed also a relatively high amount of Li compared with 5LB-2 and 5LB-1 (Fig. 10). At lower sintering temperatures, Li_2O may exist as Li-Gd-Ce-O precipitation phases at the grain boundary region. The ionic radii with approximately similar ionic sizes for Li, Gd, and Ce (*i.e.*,



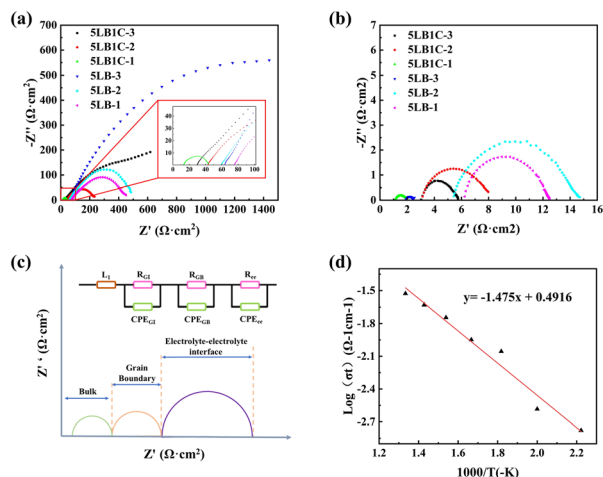


Fig. 11 (a) Nyquist plots of 5LB1C and 5LB samples at 450 °C (The inset shows the enlarged view of high frequency part), (b) Nyquist plots of 5LB1C and 5LB samples at 750 °C, (c) illustration of a typical impedance plot for a polycrystalline sample with equivalent circuits, and (d) Arrhenius plot of the 5LB1C-1 sample.

0.092, 0.1053, and 0.097 nm, respectively) could result in the doping of Li^+ into CeO_2 or Gd_2O_3 with the formation of negative charges as Li_{Ce}''' and Li_{Gd}'' . Both of these negative charges could potentially neutralise the positive charges in the grain boundary core (e.g., by substituting the excess oxygen vacancies), hence decreasing the overall conductivity of the resulting solid state electrolyte.⁸⁵

Compared with 5LB-3, the grain boundary resistance significantly decreases in the Li, Bi and Cu-doped GDC electrolytes in 5LB1C-1, indicating an improvement in the electrochemical properties. Through a naked-eye inspection of the Nyquist plots shown in Fig. 11, it can be seen that the 5LB1C-1 sample provides the highest total conductivity. This indicates that the total resistance of the electrolyte composite could be reduced by adding copper when it is sintered at 950 °C. The addition of Cu could also contribute to increasing the defects in the GDC backbone and hence help to improve the overall conductivity of the resulting electrolyte composite.²⁵

The 5LB1C-1 Nyquist plots were modelled using an equivalent electrical circuit to fit the impedance data. The Nyquist plots and their associated fitted models for 5LB1C-1 at 450–750 °C are presented in Fig. S1 (see the ESI†). The applied equivalent circuit was composed of three-sub circuits in series, each including a resistor (R) and constant phase elements (CPE) connected in parallel as shown in Fig. 11c. In the equivalent circuit, L_1 is the apparent inductor; R_{GI} stands for the resistance of the grain bulk/interior; R_{GB} is the resistance of

the grain boundary; R_{ee} is the resistance of the electrolyte-electrode interface (current collector). In general, the three contributions cannot always be distinguished during the experiments due to the relaxation characteristics of the various materials and the measured operating temperatures.

Table 4 summarises the corresponding resistance and electrical conductivity values calculated by fitting the equivalent circuit impedance data. The EIS curve fitting results showed that the R_{GI} and R_{GB} could be obtained directly. The electrical conductivity equation (eqn (S4), in the ESI†) was used to quantify the electrical conductivity of the samples. Based on the Arrhenius conductivity model (eqn (S5)†), the least squares curve fitting method was applied to obtain the total conductivity vs. operating temperature, as shown in Fig. 11(d).

As shown in Table 4, the total ionic conductivity values increased with increasing operating temperature, which is consistent with the polycrystalline solid electrolytes' temperature-dependent conductivity behaviour.⁸⁶ The ionic transport inside the lattice is usually correlated with the bulk conductivity values.²³ The value of the grain-boundary conductivity indicates the effect of microstructural characteristics on the ionic transport in the electrolyte.⁸⁷ For 5LB1C-1, the presence of bulk and grain boundary curves is hard to identify by the naked eye, but the entire cell resistance could still be calculated from the high-frequency range of the Nyquist plot. At low temperatures (below 550 °C), a significant decrease in overall conductivity is apparent. It can be found that the bulk conductivity and grain-boundary conductivity for 5LB1C-1 are promoted with increasing temperature. The total conductivity values of pure GDC sintered at 950 °C were 1.31×10^{-4} (450 °C), 8.68×10^{-4} (550 °C), 2.67×10^{-3} (650 °C), and 6.49×10^{-3} (750 °C).⁴⁷ Compared with pure GDC, 5LB1C-1 showed an obvious promotion in total conductivity.

Fig. 11(d) shows the Arrhenius plot of the 5LB1C-1 sample, where the straight lines are the least squares fits of the total conductivity data with the linearized form of the Arrhenius equation (eqn (S5)†). For the curve, data were collected from the 5LB1C pellets sintered at 950 °C for 6 h. The bulk and the grain-boundary conductivity dominate the total conductivity in all temperature range. Results obtained from equivalent circuits show a quasi-linear dependence on temperature. This is because the existence of the grain boundary arc in the intermediate frequency area of the Nyquist plot causes the conduction mechanism to change for all samples at a transition temperature of 450 °C. According to the slope and Arrhenius equation (eqn (S5)†), the activation energy is

Table 4 Resistance and ionic conductivity at different temperatures for 5LB1C-1

Temperature (°C)	R_{GI} ($\Omega \text{ cm}^2$)	σ_{GI} (S cm^{-1})	R_{GB} ($\Omega \text{ cm}^2$)	σ_{GB} (S cm^{-1})	σ_{t} (S cm^{-1})
450	22.35	1.70×10^{-3}	17.14	2.21×10^{-3}	1.66×10^{-3}
550	6.40	5.93×10^{-3}	1.08	3.51×10^{-2}	8.80×10^{-3}
650	2.93	1.29×10^{-2}	0.73	5.21×10^{-2}	1.79×10^{-2}
750	1.81	2.09×10^{-2}	0.41	9.24×10^{-2}	2.96×10^{-2}



calculated. The activation energy is the energy which is essential to begin the reaction and thus determines the overall reaction state.⁸⁸ The maximum electrical conductivity obtained for the 5LB1C-1 sample was 2.96×10^{-2} at 750 °C with an activation energy of 0.13 eV. The activation energy of pure GDC sintered at 950 °C was 0.83 eV.⁴⁷ The high activation energy could be due to the additional grain boundary resistance, which has been reported in other ceria-based materials.^{89,90}

Considering the low total conductivities of pure GDC (sintered at 950 °C), the results of activation energies fit well with the values of total conductivity. Compared with pure GDC (sintered at 950 °C), it showed a considerable improvement in activation energy for 5LB1C-1.

Conclusions

A multi-doped ceria-based electrolyte composite with a sintering temperature as low as 750 °C was successfully prepared in this work. The preparation method was based on adding controlled amounts of Li, Bi, and Cu dopants to modify the electrical and sintering properties of the commercial GDC solid-oxide electrolytes. The physicochemical and electrochemical characteristics of the resulting co-doped GDC electrolyte were systematically analysed. The impact of copper as an addition to co-doped GDC electrolytes was explored, and the influences of sintering temperature on ionic conductivity were reported. After sintering, the surface analysis of ceramic pellets for doping elements was investigated. The modified Li-Bi-Cu-doped GDC electrolyte sintered at 750 °C for 6 h achieved 96.70% relative density, which is one of the lowest sintering temperatures reported for the modified GDC composite. The maximum electrical conductivity obtained for the $\text{Ce}_{0.712}\text{Gd}_{0.178}\text{Li}_{0.05}\text{Bi}_{0.05}\text{Cu}_{0.01}\text{O}_{1.801}$ sample (sintered at 950 °C for 6 h) was 29.6 mS cm^{-1} at 750 °C with an activation energy of 0.13 eV. The results of this study help understand better the properties of co-doped electrolyte materials for the fabrication of more efficient steam electrolyzers for environmentally friendly hydrogen generation.

Author contributions

Yuheng Liu: methodology, investigation, data curation, formal analysis, visualisation, writing – original draft. Ming Xu: investigation, formal analysis. Yunlong Zhao: supervision, writing – review & editing. Bahman Amini Horri: conceptualisation, investigation, methodology, project administration, resources, supervision, writing – review & editing.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 A. Hauch, R. Kungas, P. Blennow, A. B. Hansen, J. B. Hansen, B. V. Mathiesen and M. B. Mogensen, *Science*, 2020, **370**, 6513.
- 2 M. M. Nejadian, P. Ahmadi and E. Houshfar, *Fuel*, 2023, **336**, 126835.
- 3 M. Fallah Vostakola, H. Ozcan, R. S. El-Emam and B. Amini Horri, *Energies*, 2023, **16**, 3327.
- 4 Y. Zheng, J. Wang, B. Yu, W. Zhang, J. Chen, J. Qiao and J. Zhang, *Chem. Soc. Rev.*, 2017, **46**, 1427–1463.
- 5 J. T. S. Irvine, D. Neagu, M. C. Verbraeken, C. Chatzichristodoulou, C. Graves and M. B. Mogensen, *Nat. Energy*, 2016, **1**, 15014.
- 6 X. Tong, S. Ovtar, K. Brodersen, P. V. Hendriksen and M. Chen, *J. Power Sources*, 2020, **451**, 227742.
- 7 A. Hauch, S. H. Jensen, S. Ramousse and M. Mogensen, *J. Electrochem. Soc.*, 2006, **153**, A1741.
- 8 A. Hauch, K. Brodersen, M. Chen, C. Graves, S. H. Jensen, P. S. Jørgensen, P. V. Hendriksen, M. B. Mogensen, S. Ovtar and X. Sun, *ECS Trans.*, 2017, **75**, 3.
- 9 J. Schefold, A. Brisse and H. Poepke, *Int. J. Hydrogen Energy*, 2017, **42**, 13415–13426.
- 10 M. Fallah Vostakola and B. Amini Horri, *Energies*, 2021, **14**, 1280.
- 11 H. Kurokawa, G. Y. Lau, C. P. Jacobson, L. C. De Jonghe and S. J. Visco, *J. Mater. Process. Technol.*, 2007, **182**, 469–476.
- 12 D. Sarantaridis, R. Chater and A. Atkinson, *J. Electrochem. Soc.*, 2008, **155**, B467.
- 13 Y. Zheng, Z. Chen and J. Zhang, *Electrochem. Energy Rev.*, 2021, **4**, 508–517.
- 14 M. Ni, M. Leung and D. Leung, *Int. J. Hydrogen Energy*, 2008, **33**, 2337–2354.
- 15 K. Neuhaus, R. Dolle and H.-D. Wiemhöfer, *J. Electrochem. Soc.*, 2018, **165**, F533.
- 16 X. D. Zhou, W. Huebner, I. Kosacki and H. U. Anderson, *J. Am. Ceram. Soc.*, 2002, **85**, 1757–1762.
- 17 M. Choolaei, E. Jakubczyk and B. Amini Horri, *Electrochim. Acta*, 2023, **445**, 142057.
- 18 D. J. Young, *High temperature oxidation and corrosion of metals*, Elsevier, 2008.
- 19 K. Lu, N. J. Manjooran, R.-i. Murakam and G. Pickrell, *Advances in Synthesis, Processing, and Applications of Nanostructures*, John Wiley & Sons, 2012.
- 20 V. Esposito, S. P. V. Foghmoes and S. Ramousse, *J. Eur. Ceram. Soc.*, 2014, **34**, 2371–2379.
- 21 C. Herring, *J. Appl. Phys.*, 1950, **21**, 301–303.
- 22 J. Nicholas and L. Dejonghe, *Solid State Ionics*, 2007, **178**, 1187–1194.



- 23 N. Ghaemi, R. C. T. Slade and B. Amini Horri, *Ceram. Int.*, 2021, **47**, 20009–20018.
- 24 T. Zhu, Y. Lin, Z. Yang, D. Su, S. Ma, M. Han and F. Chen, *J. Power Sources*, 2014, **261**, 255–263.
- 25 H.-J. Choi, Y.-H. Na, M. Kwak, T. W. Kim, D.-W. Seo, S.-K. Woo and S.-D. Kim, *Ceram. Int.*, 2017, **43**, 13653–13660.
- 26 G. Accardo, D. Frattini, H. C. Ham, J. H. Han and S. P. Yoon, *Ceram. Int.*, 2018, **44**, 3800–3809.
- 27 C. Nicollet, J. Waxin, T. Dupeyron, A. Flura, J.-M. Heintz, J. P. Ouweltjes, P. Piccardo, A. Rougier, J.-C. Grenier and J.-M. Bassat, *J. Power Sources*, 2017, **372**, 157–165.
- 28 A. Azzolini, J. Downs and V. M. Sglavo, *J. Eur. Ceram. Soc.*, 2015, **35**, 2119–2127.
- 29 J. P. F. Grilo, D. A. Macedo, R. M. Nascimento and F. M. B. Marques, *Electrochim. Acta*, 2019, **318**, 977–988.
- 30 H.-C. Lee, J.-A. Lee, J.-H. Lee, Y.-W. Heo and J.-J. Kim, *Ceram. Int.*, 2017, **43**, 11792–11798.
- 31 L. Spiridigliozzi, G. Dell'Agli, G. Accardo, S. P. Yoon and D. Frattini, *Ceram. Int.*, 2019, **45**, 4570–4580.
- 32 G. Accardo, G. Dell'Agli, L. Spiridigliozzi, S. P. Yoon and D. Frattini, *Int. J. Hydrogen Energy*, 2020, **45**, 19707–19719.
- 33 A. Alemayehu, M. Biesuz, K. Y. Javan, A. Tkach, P. M. Vilarinho, V. M. Sglavo and V. Tyrpekl, *J. Eur. Ceram. Soc.*, 2023, 4837–4843.
- 34 J. Li, Q. Cai and B. A. Horri, *Chem. – Eur. J.*, 2023, e202300021.
- 35 I. Van Driessche, R. Mouton and S. Hoste, *Mater. Res. Bull.*, 1996, **31**, 979–992.
- 36 M. L. Ruiz, I. D. Lick, M. I. Ponzi, E. R. Castellón, A. Jiménez-López and E. N. Ponzi, *Thermochim. Acta*, 2010, **499**, 21–26.
- 37 A. Kumar, E. Wolf and A. Mukasyan, *AIChE J.*, 2011, **57**, 3473–3479.
- 38 Y. K. Du, P. Yang, Z. G. Mou, N. P. Hua and L. Jiang, *J. Appl. Polym. Sci.*, 2006, **99**, 23–26.
- 39 I. Savva, A. S. Kalogirou, M. Achilleos, E. Vasile, P. A. Koutentis and T. Krasia-Christoforou, *Molecules*, 2016, **21**, 1218.
- 40 K. Zhu, G. Wang, S. Zhang, Y. Du, Y. Lu, R. Na, Y. Mu and Y. Zhang, *RSC Adv.*, 2017, **7**, 30564–30572.
- 41 J.-A. Lee, Y.-E. Lee, H.-C. Lee, Y.-W. Heo, J.-H. Lee and J.-J. Kim, *Ceram. Int.*, 2016, **42**, 11170–11176.
- 42 Z. Yang, Y. Yang, Y. Chen, Y. Liu, T. Zhu, M. Han and F. Chen, *J. Power Sources*, 2015, **297**, 271–275.
- 43 A. A. Solovyev, I. V. Ionov, V. A. Semenov, A. V. Shipilova and S. V. Rabotkin, *J. Phys.: Conf. Ser.*, 2019, **1393**, 012140.
- 44 A. Sanson, P. Mangifesta, E. Roncari, S. Fiameni, S. Barison and M. Fabrizio, *Proceedings 8th European Fuel Cell Forum*, 2008.
- 45 V. Gil, C. Moure, P. Duran and J. Tartaj, *Solid State Ionics*, 2007, **178**, 359–365.
- 46 V. Gil, J. Tartaj, C. Moure and P. Duran, *J. Eur. Ceram. Soc.*, 2007, **27**, 801–805.
- 47 G. Accardo, E. Audasso and S. P. Yoon, *J. Alloys Compd.*, 2022, **898**, 162880.
- 48 B. Arndt, H. Noei, T. F. Keller, P. Müller, V. Vonk, A. Nenning, A. K. Opitz, J. Fleig, U. Rütt and A. Stierle, *Thin Solid Films*, 2016, **603**, 56–61.
- 49 M. Choolaei, Q. Cai, R. C. T. Slade and B. Amini Horri, *Ceram. Int.*, 2018, **44**, 13286–13292.
- 50 G. Accardo, G. S. Kim, H. C. Ham and S. P. Yoon, *Int. J. Hydrogen Energy*, 2019, **44**, 12138–12150.
- 51 C. G. M. Lima, T. H. Santos, J. P. F. Grilo, R. P. S. Dutra, R. M. Nascimento, S. Rajesh, F. C. Fonseca and D. A. Macedo, *Ceram. Int.*, 2015, **41**, 4161–4168.
- 52 V. Gil, J. Tartaj, C. Moure and P. Duran, *Ceram. Int.*, 2007, **33**, 471–475.
- 53 A. W. Burton, K. Ong, T. Rea and I. Y. Chan, *Microporous Mesoporous Mater.*, 2009, **117**, 75–90.
- 54 J. S. J. Hargreaves, *Catal., Struct. React.*, 2016, **2**, 33–37.
- 55 S. Agarwal, X. Zhu, E. Hensen, L. Lefferts and B. Mojet, *J. Phys. Chem. C*, 2014, **118**, 4131–4142.
- 56 M. Dosa, M. Piumetti, S. Bensaid, T. Andana, C. Novara, F. Giorgis, D. Fino and N. Russo, *Catal. Lett.*, 2018, **148**, 298–311.
- 57 C. Andriopoulou, A. Trimpalis, K. C. Petallidou, A. Sgoura, A. M. Efstathiou and S. Boghosian, *J. Phys. Chem. C*, 2017, **121**, 7931–7943.
- 58 S. Chen, L. Li, W. Hu, X. Huang, Q. Li, Y. Xu, Y. Zuo and G. Li, *ACS Appl. Mater. Interfaces*, 2015, **7**, 22999–23007.
- 59 Z.-Y. Pu, X.-S. Liu, A.-P. Jia, Y.-L. Xie, J.-Q. Lu and M.-F. Luo, *J. Phys. Chem. C*, 2008, **112**, 15045–15051.
- 60 S. Acharya, V. Gaikwad, V. Sathe and S. Kulkarni, *Appl. Phys. Lett.*, 2014, **104**, 113508.
- 61 S. Acharya, V. Gaikwad, S. D'Souza and S. Barman, *Solid State Ionics*, 2014, **260**, 21–29.
- 62 T. Vinodkumar, B. G. Rao and B. M. Reddy, *Catal. Today*, 2015, **253**, 57–64.
- 63 Y. Lee, G. He, A. J. Akey, R. Si, M. Flytzani-Stephanopoulos and I. P. Herman, *J. Am. Chem. Soc.*, 2011, **133**, 12952–12955.
- 64 Z. Wu, M. Li, J. Howe, H. M. Meyer III and S. H. Overbury, *Langmuir*, 2010, **26**, 16595–16606.
- 65 T. Taniguchi, T. Watanabe, N. Sugiyama, A. Subramani, H. Wagata, N. Matsushita and M. Yoshimura, *J. Phys. Chem. C*, 2009, **113**, 19789–19793.
- 66 A. Nakajima, A. Yoshihara and M. Ishigame, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **50**, 13297.
- 67 L. Li, F. Chen, J.-Q. Lu and M.-F. Luo, *J. Phys. Chem. A*, 2011, **115**, 7972–7977.
- 68 J. Contour, A. Salesse, M. Froment, M. Garreau, J. Thevenin and D. Warin, *J. Microsc. Spectrosc. Electron.*, 1979, **4**, 483–491.
- 69 M. Shek, J. Hrbek, T. Sham and G.-Q. Xu, *Surf. Sci.*, 1990, **234**, 324–334.
- 70 S. Oswald, *Appl. Surf. Sci.*, 2015, **351**, 492–503.
- 71 S. Xia, X. Zhang, C. Liang, Y. Yu and W. Liu, *Energy Stor. Mater.*, 2020, **24**, 329–335.
- 72 K. Rana, G. Kucukayan-Dogu, H. S. Sen, C. Boothroyd, O. Gulseren and E. Bengu, *J. Phys. Chem. C*, 2012, **116**, 11364–11369.
- 73 K. Waetzig, A. Rost, U. Langklotz, B. Matthey and J. Schilm, *J. Eur. Ceram. Soc.*, 2016, **36**, 1995–2001.
- 74 T. Šalkus, E. Kazakevičius, A. Kežionis, V. Kazlauskienė, J. Miškinis, A. Dindune, Z. Kanepe, J. Ronis, M. Dudek, M.



- Bučko, J. R. Dygas, W. Bogusz and A. F. Orliukas, *Ionics*, 2010, **16**, 631–637.
- 75 J. Cao, L. Li and Z. Gui, *J. Mater. Chem.*, 2001, **11**, 1198–1200.
- 76 H. M. Wang, M. C. Simmonds and J. M. Rodenburg, *Mater. Chem. Phys.*, 2003, **77**, 802–807.
- 77 Q.-H. Wu, A. Thißen and W. Jaegermann, *Surf. Sci.*, 2005, **578**, 203–212.
- 78 V. S. Dharmadhikari, S. Sainkar, S. Badrinarayan and A. Goswami, *J. Electron Spectrosc. Relat. Phenom.*, 1982, **25**, 181–189.
- 79 P. Ding, Y. Du and Z. Xu, *Chem. Res. Chin. Univ.*, 2004, **20**, 717–721.
- 80 W. E. Morgan, W. J. Stec and J. R. Van Wazer, *Inorg. Chem.*, 1973, **12**, 953–955.
- 81 S.-P. Photoemission, *Springer Series in Solid-State Sciences*, 1995, vol. 82.
- 82 G. V. Bazuev, A. P. Tyutyunnik, M. V. Kuznetsov and R. P. Samigullina, *Eur. J. Inorg. Chem.*, 2016, **2016**, 5340–5346.
- 83 A. Losev, K. Rostov and G. Tyuliev, *Surf. Sci.*, 1989, **213**, 564–579.
- 84 M. Al-Kuhaili, *Vacuum*, 2008, **82**, 623–629.
- 85 X. Guo and R. Waser, *Prog. Mater. Sci.*, 2006, **51**, 151–210.
- 86 J. Fergus, R. Hui, X. Li, D. P. Wilkinson and J. Zhang, *Solid oxide fuel cells: materials properties and performance*, CRC press, 2016.
- 87 V. T. D. S., P. R. Padala and R. U. A., *RSC Adv.*, 2015, **5**, 88675–88685.
- 88 X. Zhang, L. Wang, M. Espinoza, T. Li and M. Andersson, *Int. J. Energy Res.*, 2021, **45**, 12980–12995.
- 89 D. Ding, B. Liu, M. Gong, X. Liu and C. Xia, *Electrochim. Acta*, 2010, **55**, 4529–4535.
- 90 C. Tian and S.-W. Chan, *Solid State Ionics*, 2000, **134**, 89–102.

