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Engineering thermoelectric and mechanical properties by nanoporosity in calcium cobaltate films from reactions of Ca(OH)₂/Co₃O₄ multilayers†

Binbin Xin, (1)** Erik Ekström, (1)** Yueh-Ting Shih, (1)* Liping Huang, (1)** Jun Lu, (2)** Anna Elsukova, (3)** Yun Zhang, (2)** Wenkai Zhu, (2)** Theodorian Borca-Tasciuc, (1)** Ganpati Ramanath, (3)** Arnaud Le Febvrier, (1)** Biplab Paul (1)** and Per Eklund (1)** Arnaud Le Febvrier, (1)** Biplab Paul (1)** Arnaud Le Febvrier, (1)** Biplab Paul (1)** Arnaud Le Febvrier, (1)** Biplab Paul (1)** Biplab

Controlling nanoporosity to favorably alter multiple properties in layered crystalline inorganic thin films is a challenge. Here, we demonstrate that the thermoelectric and mechanical properties of $Ca_3Co_4O_9$ films can be engineered through nanoporosity control by annealing multiple $Ca(OH)_2/Co_3O_4$ reactant bilayers with characteristic bilayer thicknesses (b_t). Our results show that doubling b_t, e.g., from 12 to 26 nm, more than triples the average pore size from \sim 120 nm to \sim 400 nm and increases the pore fraction from 3% to 17.1%. The higher porosity film exhibits not only a 50% higher electrical conductivity of $\sigma \sim$ 90 S cm⁻¹ and a high Seebeck coefficient of $\alpha \sim$ 135 μ V K⁻¹, but also a thermal conductivity as low as $\kappa \sim$ 0.87 W m⁻¹ K⁻¹. The nanoporous $Ca_3Co_4O_9$ films exhibit greater mechanical compliance and resilience to bending than the bulk. These results indicate that annealing reactant multilayers with controlled thicknesses is an attractive way to engineer nanoporosity and realize mechanically flexible oxide-based thermoelectric materials.

1. Introduction

The rapid development of autonomous, portable and wearable devices and sensors has sparked a great deal of interest in self-sustaining energy sources to replace batteries that are typically limited by shape constraints, periodic recharging and replacement. Harvesting electricity from heat using devices made from mechanically flexible thermoelectric materials is promising for such applications. He seebeck a high thermoelectric figure of merit $ZT = \alpha^2 \sigma T/\kappa$ (α is the Seebeck coefficient, σ and κ the electrical and thermal conductivities, respectively, and T the absolute temperature for efficient energy conversion, properties such as high mechanical flexibility and toughness are key requirements.

Conducting polymer-based organic thermoelectric materials are mechanically flexible and exhibit ZT values as high as 0.42, 10-17 but are unsuitable for higher than near-room-temperature applications. Purely inorganic thermoelectrics 18 are usually brittle. Integrating nanograined films or nanocrystal

assemblies of inorganic thermoelectrics, e.g., $Bi_{2-x}Sb_xTe_3$, on flexible substrates that can withstand moderately high temperatures addresses these challenges to some extent, and yield $\alpha^2\sigma$ values up to 0.2 mW m⁻¹ K⁻² at \sim 200 °C. ^{19,20} Semiconductors with extraordinary metal-like ductility, e.g., Ag_2S , and Ag_2Se also hold promise as free-standing thermoelectric materials that obviate flexible substrates. ²¹⁻²³

Oxide-based thermoelectric films on layered substrates such as mica offer greater stability at even higher temperatures and can become viable alternatives if high α and σ , are achieved together with high mechanical flexibility.24,25 Calcium cobaltate Ca₃Co₄O₉ is an attractive p-type thermoelectric that exhibits inherently high α and high σ due to its layered crystal structure.26-30 The thermoelectric properties of layered cobaltates can be improved by nanostructuring, such as fabricating a highsingle-phase,31,32 nanostructuring approaches,33 elemental doping,34 growing textured films with c-axis orientation.35 However, the thermal conductivity was normally increased with the improving electrical conductivity.36,37 Introducing nanoscale with dimensions mainly in the ranges of phonon mean free paths is a possible approach to reduce thermal conductivity without inhibiting electrical conductivity due to phonon mean free paths are typically significantly higher than electronic mean free paths. The textured nanograins and faceted nanopores not only offer additional means to lower the κ , 38,39 but also alleviate the brittleness. 40,41 Flexible porous nanograined Ca₃Co₄O₉ films obtained by annealing CaO/CoO reactant multilayers exhibit power factors as high a 0.23 mW

[&]quot;Thin Film Physics Division, Department of Physics, Chemistry and Biology (IFM), Linköping University, SE-58183 Linköping, Sweden. E-mail: binbin.xin@liu.se; per. eklund@liu.se

^bDepartment of Materials Science and Engineering, Rensselaer Polytechnic Institute, Troy, New York 12180, USA

^cRensselaer Polytechnic Institute, Department of Mechanical, Aerospace, and Nuclear Engineering, Troy, NY 12180, USA

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m⁻¹ K⁻² at room temperature.⁴⁰ To further increase the porosity of nanoporous Ca₃Co₄O₉ films retaining α and σ , it might be an effective way to further reduce κ and improve flexibility.

Here, we demonstrate that nanoporosity characteristics in $Ca_3Co_4O_9$ films can be controlled by adjusting the reactant bilayer thickness in multilayer stacks. The porosities change from 11.2 to 17.1%. The nanoporous films show high σ with a narrow range from 80 to 95 S cm⁻¹ and high α of \sim 130 to \sim 135 μ V K⁻¹, which are close to the values.⁴²⁻⁴⁴ The film with the highest porosity 17.1% has the lowest κ (\sim 0.87 W m⁻¹ K⁻¹). The nanoporous $Ca_3Co_4O_9$ films exhibit greater mechanical compliance and resilience to bending than the bulk. Our findings showing pore engineering of layered-ceramics is attractive for realizing mechanically flexible thermoelectrics.

2. Experimental section

Nanoporous Ca₃Co₄O₉ thin films were synthesized by annealing Ca(OH)₂/Co₃O₄ multilayers on muscovite mica(00*l*) substrates. CaO/Co₃O₄ multilayers were deposited at 600 °C by reactive RFmagnetron sputtering from Ca and Co targets with a 2 mTorr plasma of a 0.5% O₂/99.5% Ar gas mixture⁴⁰. Ambient airexposure of the CaO/Co₃O₄ multilayers for one month resulted in Ca(OH)₂/Co₃O₄ multilayers via CaO hydration into Ca(OH)2.45 Subsequent annealing Ca(OH)2/Co3O4 multilayers at 700 °C for 2 hours in air led to nanoporous Ca₃Co₄O₉ formation. We prepared five sets of multilayers with different Ca(OH)₂/ Co₃O₄ bilayer thicknesses, b_t. The deposition times for CaO and Co₃O₄ were varied while keeping the total nominal multilayer thickness constant at ~140 nm. The individual thicknesses of Ca(OH)₂ and Co₃O₄ layers in each multilayer film were identical, i.e., $b_t/2$. Thus, altering the bilayer thickness in the $12 \le b_t \le$ 50 nm range is equivalent to varying the number of bilayers b_n in the $21 \ge b_n \ge 5$ range.

Phase identification was carried out by X-ray diffractometry (XRD) using a PANalytical X'Pert PRO instrument with monochromatic Cu K α radiation ($\lambda=1.5406$ Å) and a Ni filter. X-ray reflectivity (XRR) measurements were carried out in a PANalytical Empyrean diffractometer equipped with a copper Cu K α source with a hybrid mirror on the incidence beam path, a triple-axis Ge 220 analyzer on the diffracted beam path, and a PIXcel3D detector operated in open detection mode. The XRR data were fitted using the X'Pert reflectivity program.

Scanning electron microscopy (SEM) and energy dispersive X-ray (EDX) spectroscopy were carried out in a LEO Gemini 1550 Zeiss instrument operated at 10 kV to characterize film morphology and composition. The surface porosity fraction was determined by analyses of SEM micrographs using the Java version of image J software^{46,47}. Most of the nanopores were hexagonal in shape. The average nanopore sizes were estimated from the nanopore area with an uncertainty of 20%, by assuming all the pores to be regular hexagons. The average nanopore size is equal to $L + 2L \cos 60^\circ$ where L is hexagon side length. Transmission electron microscopy (TEM) was carried out in a FEI Tecnai G2 TF20 UT instrument operated at 200 kV on multilayer cross-sections prepared by face-to-face gluing of two sample pieces and mounting them on a Ti grid. The

samples were mechanically polished down to 50 μ m and ion-milled in a Gatan system with 2–5 kV Ar $^+$ beams incident at 5°.

Electrical conductivity σ was determined from the sheet resistance measured with a four-point probe Jandel RM3000 station and the film thickness determined from cross-sectional TEM images. Seebeck coefficient α was determined from the slope of the temperature gradient–voltage characteristics measured in a homemade Seebeck measurement setup system equipped with two K-type thermocouples placed at the same position as two Cu electrodes, a Keithley 2001 multimeter, two Peltier elements acting as temperature controller, and two thermometers 45,48.

Thermal conductivity of the annealed $Ca_3Co_4O_9$ film-on-mica samples was determined by non-contact scanning thermal microscopy (SThM) described elsewhere^{49–52}. This technique utilizes a Joule-heated 5 μ m-diameter Wollaston wire probe, whose thermal resistance was measured in air at 100 nm above the sample surface based on temperature-induced changes in the electrical resistance of the probe and the dissipated Joule heating power. For each film, we measured thermal resistance at three different locations on the sample surface. Thermal conductivity was determined by fitting the thermal resistance data with a 3D finite element model (3DFEM) of the probe-to-sample surface heat transfer assuming isotropic thermal properties for the films and the substrate.

Mechanical flexibility was evaluated by estimating the elastic moduli of the films by surface Brillouin scattering (SBS) spectroscopy and measuring bending-induced relative changes in electrical resistance. SBS spectra were collected by using a JSR Scientific Instruments six-pass high-contrast Fabry-Perot interferometer equipped with a 532.18 nm Verdi V2 DPSS green laser probe. Surface Rayleigh and Sezawa wave velocities were obtained by using $V = (\lambda_0 \Delta f)/(2 \sin \theta_s)$, where λ_0 is the laser wavelength, Δf the Brillouin frequency shift, and θ_s the scattering angle.⁵³ For the bending tests, we measured the normalized electrical resistance change $\Delta R/R_0$ for different bending radii and cycles, where the initial resistance R_0 includes contact resistances. A constant resistance during bending corresponds to $\Delta R/R_0 = 0$ and indicates good mechanical flexibility and retention of electrical properties.

3. Results and discussion

X-ray diffractograms acquired immediately after deposition for all b_t values investigated show 111 and 222 Bragg reflections from CaO besides those from the mica substrate (Fig. 1a). The 400 diffraction peaks reflected from $\mathrm{Co_3O_4}$ is overwhelmed by a stronger 00l reflection from the mica substrate (Fig. 1a) but is observable in diffractograms from as-deposited films on sapphire (Fig. 1b). Films on sapphire also show a weak 111 peak from CoO. The CaO and $\mathrm{Co_3O_4}$ peaks intensities, normalized to that of the 004 mica substrate reflection and sapphire reflection, increase with increasing b_t (Fig. 1c).

Air-exposure of the as-deposited multilayers growing on mica and sapphire leads to the diminution and eventual disappearance of the CaO 111 and 222 peaks (see Fig. 2a and b). Multilayers on sapphire also show a similar behaviour besides

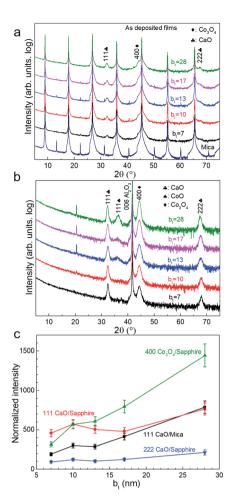


Fig. 1 X-ray diffractograms from as-deposited CaO/Co $_3$ O $_4$ multilayers with different bilayer thicknesses b $_t$, on (a) mica (00t), and (b) sapphire (006) substrates. (c) CaO and Co $_3$ O $_4$ peak intensities normalized with the intensity of the corresponding substrate peak, plotted as a function of b $_t$.

revealing a concomitant emergence of the $Ca(OH)_2$ 001 peak. This peak is not observable in diffractograms from films on mica due to substrate peak overlap. These results are indicative of the conversion of CaO to $Ca(OH)_2$ through ambient moisture uptake.⁴⁵ We note that the intensities of the $Ca(OH)_2$ and Co_3O_4 peaks increase with increasing b_t , the intensity of $Ca(OH)_2$ is not the highest when the b_t is 50 nm (Fig. 2c).

Air-exposed CaO/Co₃O₄ multilayers specified by low b_t (*e.g.*, ~12 nm) consist of equiaxed grains of Ca(OH)₂ and Co₃O₄ without any discernible interfaces (Fig. 3a). EDX spectral maps of Ca and Co indicate that Ca(OH)₂ and Co₃O₄ are interspersed across individual layers (Fig. 3a inset). This result is consistent with Co₃O₄ layers comprised of discontinuous grains that facilitate moisture intake and transport, as suggested in our recent work. Air-exposed multilayers with $b_t = 16$ nm exhibit more distinct Ca(OH)₂/Co₃O₄ interfaces (Fig. 3b). The Cacontaining layers that appear brighter due to a higher Z-contrast are about 81% thicker than the Co₃O₄ layers, consistent with a 95.2% unit cell volume expansion caused by the CaO \rightarrow Ca(OH)₂ conversion during air-exposure. For $b_t > 16$ nm,

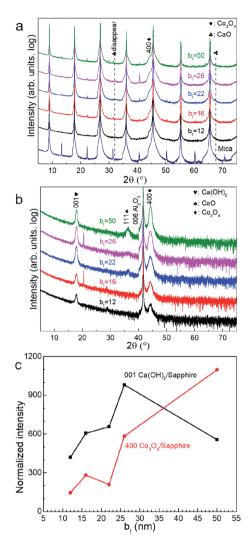


Fig. 2 X-ray diffractograms from Ca(OH) $_2$ /Co $_3$ O $_4$ multilayers formed by air-exposure of CaO/Co $_3$ O $_4$ multilayers with different bilayer thicknesses between $12 \le b_t \le 50$ nm on (a) mica (00t) and (b) sapphire (006) substrates. (c) Ca(OH) $_2$ and Co $_3$ O $_4$ peak intensities normalized to that of the 006 peak from the sapphire substrate, plotted as a function of b_t .

TEM images show distinct Co- and Ca-containing layers, but with a greater interface roughness that increases with b_t . Such interface roughening is attributable to the higher volume Ca(OH)₂ grains encroaching into the nearly unchanged adjacent Co_3O_4 layers. For $b_t = 50$ nm, the thickness of $Ca(OH)_2$ layer in top larger than that in bottle near mica, which may be due to the low hydration reactions by preventing of thicker Co₃O₄ layers and lead to the low intensity of Ca(OH)₂ 001 in Fig. 2c. SAED patterns indicate increased in-plane grain texturing with increasing b_t (see Fig. 3c and d). EDX spectral maps from air-exposed films with higher b_t show distinct Cocontaining layers, in contrast to the uniform distribution of Ca, indicating Ca diffusion (see ESI Fig. S1†). Our results indicate that the bilayer thickness needs to be greater than a critical value of $b_t > 12$ nm to form layers structure with sharp interfaces.

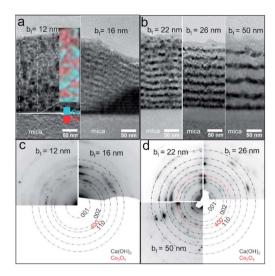


Fig. 3 (a and b) Cross-sectional bright-field TEM images and (c and d) selected-area electron diffraction (SAED) patterns from $Ca(OH)_2/Co_3O_4$ multilayers with different bilayer thicknesses b_f .

Otherwise, EDS maps from multilayers with $b_t = 16$ nm show clear Co layers and almost uniform distribution Ca elements in multilayers (see ESI Fig. S1a†).

X-ray diffractograms show that annealing the Ca(OH)₂/Co₃O₄ multilayers leads to Ca₃Co₄O₉ formation through a reaction between the Co₃O₄ and Ca(OH)₂ layers (Fig. 4a). The exclusive presence of multiple 00*l* peak reflections of Ca₃Co₄O₉ indicate textured basal planes oriented parallel to the substrate surface. The Ca₃Co₄O₉ grain size is largest for films specified by $b_t = 22$ nm, as indicated by the narrowest width of the Ca₃Co₄O₉ 002 reflection.

Surfaces of the annealed films exhibit faceted intergranular nanopores (see Fig. 5). Films with the smallest b_t in our studies reveal a relatively rough surface, probably due to the edge-on orientation of some platelet-shaped grains on the surface (see Fig. 5a). In contrast, annealed films with $b_t \ge 16$ nm exhibit smoother surfaces, with faceted nanopores between hexagonal terraces and plate-shaped grains (Fig. 5b–e). The average pore

size increases monotonically from around 120 to 400 nm with increasing b_t from 12 nm to 50 nm (see Fig. 5f). The porosity fraction increases with b_t from 3.7% to 17.1% for $12 \le b_t \le$ 26 nm but drops to 13.4% for $b_t = 50$ nm (Fig. 5f).

Cross-sectional TEM micrographs of films obtained by annealing Ca(OH)₂/Co₃O₄ multilayers with $12 \le b_t \le 50$ nm created from CaO/Co₃O₄ multilayers (Fig. 6a-e) reveal a polycrystalline Ca₃Co₄O₉ layer separated from the substrate by an amorphous glass layer. 40 Lattice images for the Ca3Co4O9 layer and SAED patterns (Fig. 6f-j) confirm that the (001) basal planes are oriented parallel to the film surface, corroborating our XRD and SEM results. The in-plane/out-of-plane aspect ratio of the pores is about four for the film with $b_t = 50 \text{ nm}$ (Fig. 6e). This observation is consistent with the SEM results and our recent work45 indicating that oriented nanopore formation arises from basal plane removal driven by local densification of textured $Ca_3Co_4O_9$. For all b_t values except $b_t =$ 50 nm, we observe nanopores spanning across the Ca₃Co₄O₉ layers and microporous gaps at the Ca₃Co₄O₉ - amorphous layer interface and interlayer nanoporosity (see Fig. 6a-e). No microporous gaps or interlayer porosity are discernible in the amorphous layers.

TEM micrographs reveal that for all our films, the Ca₃Co₄O₉ layer thickness is around twice the amorphous layer thickness (Fig. 7). Increasing the b_t from 12 nm to 26 nm increased the $Ca_3Co_4O_9$ layer thickness from 170 \pm 10 nm to 193 \pm 10 nm but decreased the amorphous layer thickness from 103 nm to 73 nm. EDX analyses of the amorphous layer revealed O, Al, Si and Ca (ESI Fig. S2†), but no traces of Co above the EDX detection limit. These results suggest that the amorphous layer is formed due to preferential Ca diffusion and incorporation into the mica substrate. This inference is supported by the inverse correlation between amorphous layer thickness and b_t and the higher Ca/Co ratio of 55: 45 in multilayer than that in the Ca₃Co₄O₉ layer. The anomalously low Ca₃Co₄O₉ layer thickness of 159 \pm 10 nm for $b_t = 50$ nm is likely an outlier due to a very low surface porosity fraction of 13.4% and no microporous gaps at the Ca₃Co₄O₉ - amorphous layer interface and needs further study.

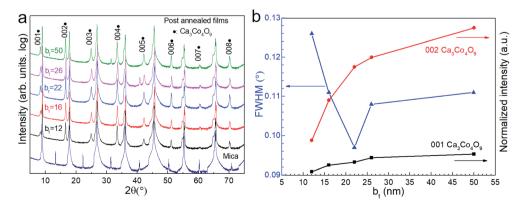


Fig. 4 (a) X-ray diffractograms showing $Ca_3Co_4O_9$ formation on mica (00*l*) substrate by annealing $Ca(OH)_2/Co_3O_4$ multilayers with different bilayer thicknesses in the $12 \le b_t \le 50$ nm range. (b) $Ca_3Co_4O_9$ 001 and 002 peak intensities normalized with the intensity of the corresponding substrate peaks, plotted as a function of b_t .

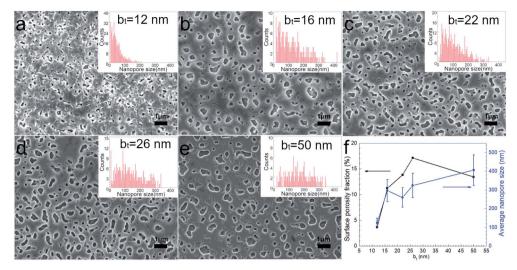


Fig. 5 (a–e) SEM images from annealed $Ca(OH)_2/Co_3O_4$ multilayers with $12 \le b_t \le 50$ nm grown on mica (001) substrates. (f) Surface porosity fraction determined from SEM image analyses.

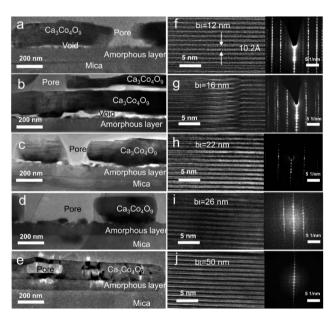


Fig. 6 (a–e) Cross-sectional bright-field TEM images from $Ca_3Co_4O_9$ films obtained by annealing $Ca(OH)_2/Co_3O_4$ multilayers with $12 \le b_t \le 50$ nm created from CaO/Co_3O_4 multilayers. (f–j) The corresponding high-resolution TEM images and SAED patterns capturing the layered atomic structure of $Ca_3Co_4O_9$.

Thermoelectric properties

Considering that amorphous materials normally have lower electrical conductivity σ than corresponding crystalline materials and the amorphous layer has a low thermal conductivity κ , the amorphous oxide layer is assumed as an insulator. The electrical conductivity σ of Ca₃Co₄O₉ films slightly increases with b_t (Fig. 8a) and peaks around 13% porosity (Fig. 8b). For example, $\sigma \sim 90$ S cm⁻¹ for the film with higher porosity (13.8%) is 50% higher than for the film with lowest porosity (3.7%). But σ for the film with different porosities from 11.2 to 17.1% shows

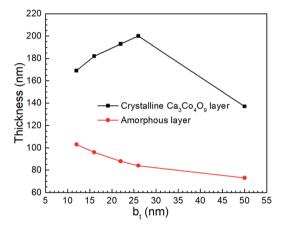


Fig. 7 The thicknesses of crystalline Ca₃Co₄O₉ and amorphous layers in porous films obtained by annealing Ca(OH)₂/Co₃O₄ multilayers created from CaO/Co₃O₄ multilayers with 12 \leq b_t \leq 50 nm.

a narrow range from 80 to 95 S cm $^{-1}$. The reason may be that the rough morphology and lowest quality layered structure of annealed film with $b_t = 12$ nm cause lower carrier mobility and the high-quality layered structure of the annealed films with $b_t \ge 16$ nm led to high carrier mobility as shown in Fig. 4 and the size range of nanopores mainly from 120 nm to 400 nm has a slight effect for the electrical conductivity.

The Seebeck coefficient α increases sightly with nanoporosity fraction, *e.g.*, from $\alpha=128~\mu V~K^{-1}$ observed for porosity = 3.7% to $\alpha=136~\mu V~K^{-1}$ for porosity = 17.1%. The power factor increases from 0.96 to 1.53 $\mu W~cm^{-1}~K^{-2}$ with increasing porosity (Fig. 8b). The highest power factor is $\alpha^2\sigma=1.63~\mu W~cm^{-1}~K^{-2}$ for the film with $b_t=50~nm$ with highest σ .

We used 3DFEM model fitting⁴⁹ to determine the effective thermal conductivity $\kappa_{\rm film}$ of the Ca₃Co₄O₉ films (see Fig. 9) by using a mica substrate thermal conductivity value of $\kappa_{\rm sub} = 0.421~{\rm W~m}^{-1}~{\rm K}^{-1}$, determined by SThM probe measurements.

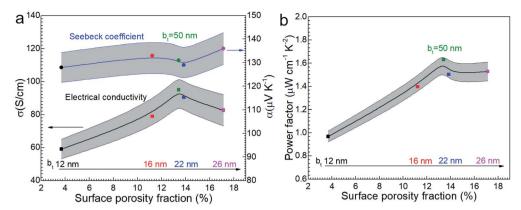


Fig. 8 (a) The electrical conductivity σ and the Seebeck coefficient α and (b) power factor $\alpha^2 \sigma$ of nanoporous Ca₃Co₄O₉ films as function of surface porosity fraction at room temperature.

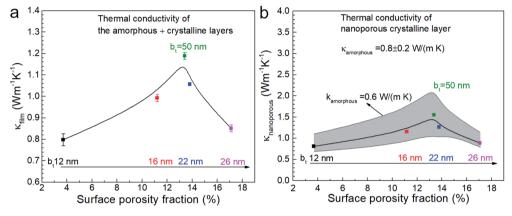


Fig. 9 (a) Effective thermal conductivity κ film of Ca₃Co₄O₉ films as a function of surface porosity fraction, and (b) thermal conductivity of the nanoporous crystalline Ca₃Co₄O₉ layer κ _{nanoporous}, assuming κ _{amorphous} = 0.8 W m⁻¹ K⁻¹ with a \pm 0.2 W m⁻¹ K⁻¹ uncertainty captured by error bars.

phous $(\kappa_{\rm amorphous})$ and nanoporous crystalline $(\kappa_{\rm nanoporous})$ layers, both of which are unknown, we computed $\kappa_{\rm nanoporous}$ by using a one-dimensional cross-plane thermal resistance network model for the crystalline-amorphous bilayer described by $\frac{t_{\rm nanoporous}}{\kappa_{\rm nanoporous}} = \frac{t_{\rm total}}{\kappa_{\rm film}} - \frac{t_{\rm amorphous}}{\kappa_{\rm amorphous}}$. We selected three possible $\kappa_{\rm amorphous}$ values of 0.6, 0.8, and 1 $(0.8 \pm 0.2)~{\rm W~m^{-1}~K^{-1}}$, which are comparable to the thermal conductivity of amorphous silica. We assumed $\kappa_{\rm amorphous}$ to be identical in all the five samples with $50 \ge b_t \ge 12~{\rm nm}$ and used the measured amorphous layer thickness for each sample, *i.e.*, 73 nm $\le t_{\rm amorphous}$ $\le 103~{\rm nm}$. This model is apt because both the amorphous and the nanoporous regions are much thinner than the $\sim 5~{\rm \mu m}$

Since $\kappa_{\rm film}$ depends on the thermal conductivities of the amor-

Plotting the effective $\kappa_{\rm film}$ and the values extracted for $\kappa_{\rm nanoporous}$ of the nanoporous layer as a function of porosity (see Fig. 9a and b) reveal peaks in both $\kappa_{\rm film}$ and $\kappa_{\rm nanoporous}$ around 13% porosity, at which σ also peaks, as shown earlier. We note that decreasing assumed $\kappa_{\rm amorphous}$ yields a higher $\kappa_{\rm nanoporous}$. Both the $\kappa_{\rm film}$ and σ are low for the lowest porosity film due to

probe-sample heat transfer radius and have low values of fitted

the relatively higher surface roughness compared to the other films. The film with the highest porosity 17.1% (*i.e.*, with $b_t = 26$ nm) has the lowest κ (\sim 0.87 W m⁻¹ K⁻¹) in annealed films with $b_t \geq 16$ nm. The annealed film with $b_t = 12$ nm shows low κ (\sim 0.8 W m⁻¹ K⁻¹) due to the rough morphology.

Mechanical flexibility

Surface Brillouin scattering (SBS) spectra from $Ca_3Co_4O_9$ films (Fig. 10a) reveal stiffness values that are significantly lower than that of bulk $Ca_3Co_4O_9$. Rayleigh peaks are seen around \pm 8.5 GHz at scattering angles $\theta_s \geq 70^\circ$. At lower scattering angles, the Rayleigh peaks merge into the central elastic peak, and we observe Sezawa peaks around \pm 11.5 GHz. Sezawa peaks are typical of soft films on hard substrates, se as is our case for $Ca_3Co_4O_9$ on sapphire. The invariance of the surface Rayleigh velocity with k||d (Fig. 10b) within experimental uncertainties is indicative of the elastic properties of the $Ca_3Co_4O_9$ film with negligible influence of the sapphire substrate. In contrast, the Sezawa velocity increases substantially with decreasing k||d due to the hard sapphire substrate. We calculated the shear modulus G of the nanoporous $Ca_3Co_4O_9$ film from

thermal conductivities.

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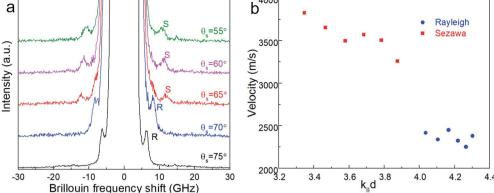


Fig. 10 (a) Representative Brillouin spectra from a \sim 185 nm-thick nanoporous Ca₃Co₄O₉ film (corresponding to $b_t = 26$ nm) on sapphire for different scattering angles θ_s showing Rayleigh (R) and Sezawa (S) peaks. (b) Surface Rayleigh and Sezawa velocities plotted a of k||d, where k|| $(4\pi \sin \theta_s)/\lambda_0$ is the surface acoustic wave vector, and d the film thickness.

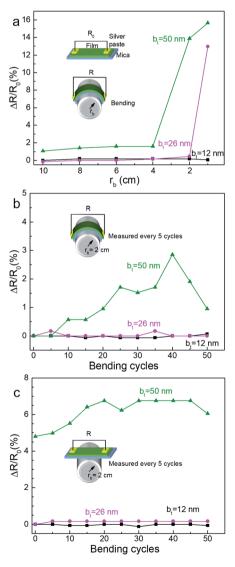


Fig. 11 The normalized resistance changing $\Delta R/R_0$ of the Ca₃Co₄O₉ films on mica (001) substrates obtained by reactions of multiple stacks of Ca(OH)₂/Co₃O₄ bilayers with $12 \le b_t \le 50$ nm as a function of (a) bending radius r_b and (b) the number of bending cycles at a bending radius of 2 cm, and (c) after restoration to non-bent geometry.

$$V_{\rm Rayleigh} = \beta \sqrt{\frac{G}{\rho}}$$
, where ρ is the density and $\beta = 0.94.^{56,57}$

Assuming isotropy, invoking $E = 2G(1 + \nu)$, using the bulk $Ca_3Co_4O_9$ Poisson's ratio $\nu = 0.31$, and a density $\rho \sim 3.0$ g cm⁻³ for the nanoporous Ca₃Co₄O₉ film determined from X-ray reflectivity measurements (ESI Fig. S3†), we get Young's and shear moduli values of $G=18.92\pm1.14$ GP, and $E=49.62\pm1.14$ 2.99 GPa, respectively. Both these values for nanoporous $Ca_3Co_4O_9$ film are 52.7% lower than the predicted values of G=39.98 GPa and E = 104.86 for bulk $Ca_3Co_4O_9$.58 The 21% lower density of the nanoporous Ca₃Co₄O₉ film compared to bulk Ca₃Co₄O₉ (3.8 g cm⁻³) partially explains the lower moduli. We propose that the low atomic bond density near the nanopore walls in the film also contribute to the increased mechanical compliance.

Mechanical bending of the film-substrate composite showed a high retention of the electrical properties, consistent with high mechanical compliance indicated by the SBS results. Twopoint resistance of the film measured as a function of bending radius r_b (see Fig. 11a) showed negligible changes in the normalized resistance ($\Delta R/R_0 \sim 0$) for $r_b \ge 4$ cm, where R_0 is the initial resistance for each film, indicating good mechanical flexibility and low electromechanical coupling. Films synthesized from multilayers specified by $b_t = 12$ nm and $b_t = 26$ nm show a greater resilience to bending and bend cycling than the film corresponding to $b_t = 50$ nm (Fig. 11b and c). While further studies are needed to understand correlations between nanoporosity, bend cycling, and mechanical compliance, our results clearly indicate that the nanoporous films are mechanically flexible and have a low electromechanical coupling for a large range of b_t and bending.

Conclusions

We have synthesized $Ca_3Co_4O_9$ films with different porosities by annealing multilayers of calcium and cobalt oxide bilayers specified by $Ca(OH)_2/Co_3O_4$ bilayer thicknesses b_t . Increasing b_t increases the nanoporosity fraction and average nanopore size in the Ca₃Co₄O₉ film formed during annealing. The higher Nanoscale Advances

porosity films exhibit a 50% higher electrical conductivity as well as a high Seebeck coefficient, together with a low thermal conductivity. The nanoporous Ca₃Co₄O₉ films show a higher mechanical compliance than the bulk Ca₃Co₄O₉ and are resilient to mechanical bending and bend cycling. These results indicate that engineering nanoporosity in layered oxides through reactions of multilayer stacks of component oxides could be attractive for achieving mechanically-flexible highfigure-of-merit thermoelectric nanomaterials for emergent applications.

Author contributions

Binbin Xin: performed experiments, data collection and analvsis, writing – original draft, and writing – review & editing; Erik Ekström: magnetron sputtering; Jun Lu and Anna Elsukova: TEM data; Yueh-Ting Shih and Liping Huang: surface Brillouin scattering (SBS) spectroscopy; Yun Zhang, Wenkai Zhu, and Theodorian Borca-Tasciuc: thermal conductivity; Biplab Paul and Per Eklund: experiment design and supervision; Erik Ekström, Yueh-Ting Shih, Liping Huang, Jun Lu, Anna Elsukova, Yun Zhang, Wenkai Zhu, Theodorian Borca-Tasciuc, Ganpati Ramanath, Arnaud Le Febvrier, Biplab Paul and Per Eklund: writing - review & editing.

Conflicts of interest

The authors declare no conflicts of interest.

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