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## Synthesis, structure, and superconductivity of $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$ <sup>†</sup>

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Sodium and potassium have been doped into  $\text{La}_7\text{SrCuO}_{16-\delta}$  to form  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  solid solutions by a solid-state reaction, which crystallizes in the  $I4/mmm$  space group when  $0 \leq x < 0.16$  and  $Fmmm$  when  $0.16 < x \leq 0.25$ . With the equivalent replacement of four  $\text{Sr}^{2+}$  by one  $\text{Na}^+$ , one  $\text{K}^+$  and two  $\text{La}^{3+}$ , it is very interesting to find that  $T_c^{\text{zero}}$  (the definition is presented in the introduction part) of  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  does not remain constant with an increase in  $x$  but first increases from 14.8 K for  $x = 0.00$  to 33.9 K for  $x = 0.10$  and then decreases to 18.5 K for  $x = 0.175$ .

## 1. Introduction

Since J. G. Bednorz *et al.*<sup>1</sup> first discovered superconductivity in the La–Ba–Cu–O system with  $T_c^{\text{onset}}$  (a transition temperature at which the resistivity begins to drop rapidly to “zero” with a decrease in temperature, indicating the onset of superconductivity) at about 30 K and  $T_c^{\text{zero}}$  (the highest temperature at which the resistivity of the sample reaches “zero”, marking the full superconducting state) at about 20 K, the mechanism of high  $T_c$  (a general term for the phase transition temperature, which can be one of  $T_c^{\text{onset}}$ ,  $T_c^{\text{zero}}$ , and  $T_c^{\text{mag}}$  (the temperature at which the magnetization of the sample starts to decrease to negative values with a decrease of temperature, indicating the start of the Meissner effect) for a superconductor) has been widely investigated, and new materials with higher  $T_c$  have been explored.<sup>2,3</sup>  $\text{La}_2\text{CuO}_{4-\delta}$  materials are called “model materials” for the study of superconductivity due to their simple formulation and structure,<sup>4</sup> which provides a unique opportunity to investigate the relationship between the structure and superconducting properties.<sup>5</sup> The widely studied Sr-substituted compound  $\text{La}_{2-x}\text{Sr}_x\text{CuO}_{4-\delta}$  is the copper oxide compound that can be varied over a wide enough range to obtain a full spectrum of electronic properties.<sup>6</sup> As the Sr content increases,

it goes from antiferromagnetic insulator ( $0 \leq x \leq 0.03$ ) to superconductor ( $0.07 \leq x \leq 0.27$ ) to non-superconducting metal ( $x \geq 0.27$ ).<sup>7</sup> Many attempts have been made to investigate the relationship with  $T_c$  in  $\text{La}_{2-x}\text{Sr}_x\text{CuO}_{4-\delta}$  systems by adding new doping elements and changing the doping amount. J. P. Attfield *et al.*<sup>8</sup> have shown that the  $T_c$  of  $(\text{Ln}_{1-x}\text{M}_x)_2\text{CuO}_{4-\delta}$  ( $\text{Ln} = \text{La}$  and other lanthanides,  $\text{M} = \text{Ca, Sr, and Ba}$ ) superconductors depends on the doping level  $x$  and the average A-site cation radius.<sup>9,10</sup> More complex doping compounds, such as  $(\text{La}_{0.925}\text{Sr}_{0.075-x}\text{Ba}_x)_2\text{CuO}_{4-\delta}$ ,<sup>11</sup>  $(\text{La}_{0.925-x}\text{Ln}_x\text{Sr}_{0.075})_2\text{CuO}_{4-\delta}$  ( $\text{Ln} = \text{Pr, Nd, Sm, Eu}$ ),<sup>10,12</sup> have also been studied, which indicates that  $T_c$  is also affected by the change in the ionic radius at the A-site. M. A. Subramanian *et al.*<sup>13</sup> have reported the superconductivity of  $\text{La}_{2-x}\text{A}_x^{1+}\text{CuO}_{4-\delta}$  ( $\text{A} = \text{Na, K}$ ) with  $T_c^{\text{onset}} \sim 40$  K, which confirms the feasibility of A-site doping with alkali metal cations. The elements Na and K can influence the crystal structure and electronic properties of copper oxide superconductors due to their unique electronic configurations.<sup>14,15</sup> They also help to lower the sintering temperature, promote sintering, and inhibit grain growth.<sup>16</sup> Therefore, combining Na and K into  $\text{La}_{2-x}\text{Sr}_x\text{CuO}_{4-\delta}$  matrices provides an interesting avenue to tailor the electronic structure and superconducting properties of the materials. In the  $\text{La}_{2-x}\text{Sr}_x\text{CuO}_{4-\delta}$  system, superconductivity is usually reported when  $0.05 < x < 0.27$ ,  $x = 0.15$ , which is identified as the optimal doping concentration.<sup>17</sup> Many studies relating to the optimal doping of  $\text{La}_{1.85}\text{Sr}_{0.15}\text{CuO}_{4-\delta}$  have been performed.<sup>18–22</sup> The research on the over-doped range ( $0.15 \leq x \leq 0.27$ ) should be enhanced. Therefore,  $\text{La}_{1.75}\text{Sr}_{0.25}\text{CuO}_{4-\delta/4}$  (which can also be noted as  $\text{La}_7\text{SrCu}_4\text{O}_{16-\delta}$ ) is chosen as the solid solution matrix, serving as a starting point for further investigation. To reduce the change in the average A-site cation radius to limit the size effect on  $T_c$  suggested by J. P. Attfield *et al.*,<sup>8</sup> one  $\text{K}^+$ , one  $\text{Na}^+$ , and two  $\text{La}^{3+}$  were chosen to replace four  $\text{Sr}^{2+}$  (the details are shown in section 3.1). Then, the nominal compounds  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  were tried. In this case,

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it is found that the above doping is just an equivalent doping, which introduces no additional electron or hole into the synthesized compound if  $\delta$  is not changed. This means that the  $T_c$  of  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  may remain constant if we believe that the  $T_c$  of doped  $\text{La}_2\text{CuO}_{4-\delta}$  is strongly related to the hole content. However, the  $T_c^{\text{zero}}$  of  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  is raised to 33.9 K for  $x = 0.10$  from 14.8 K for  $x = 0.00$ , then finally lowered to 18.5 K for  $x = 0.175$ . The details for such an interesting phenomenon and the possible explanation about the influence of K and Na ions on the superconductivity are presented below.

## 2. Experimental

The samples with the nominal formula  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  ( $x = 0.0, 0.025, 0.050, 0.075, 0.100, 0.125, 0.150, 0.175, 0.200, 0.225$  and  $0.250$ , denoted as LSKNCO1, LSKNCO2, ..., and LSKNCO11, respectively) were synthesized.  $\text{La}_2\text{O}_3$  (99.95%),  $\text{SrCO}_3$  (A.R.),  $\text{K}_2\text{CO}_3$  (A.R.),  $\text{Na}_2\text{CO}_3$  (A.R.), and  $\text{CuO}$  (A.R.) were used as raw materials. The oven-dried raw materials were weighed according to the molar ratios of the nominal formula  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  and homogenized by about 30 min of grinding for total 10 g of mixtures with an agate mortar and a pestle. These mixtures were sintered first at 900 °C for 12 h. Then, the reacted powders were pressed into pellets under 30 MPa and sintered at 920 °C for 12 h. The sintered mass was again crushed, pulverized, and pressed into cylindrical pellets to undertake five 12 h heat treatments at 1100 °C (LSKNCO1, LSKNCO2, LSKNCO3), 1080 °C (LSKNCO4, LSKNCO5), 1060 °C (LSKNCO6), 1020 °C (LSKNCO7), and 960 °C (LSKNCO8, LSKNCO9, LSKNCO10, LSKNCO11). All the treatments were performed in air.

Powder X-ray diffraction (PXRD) data of  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  were collected on a PANalytical X'Pert PRO (The Netherlands) with  $\text{Cu K}\alpha$  ( $\lambda_1 = 1.5405 \text{ \AA}$  and  $\lambda_2 = 1.5443 \text{ \AA}$ ) radiation over a wide  $2\theta$  range (5°–120°) at a scan rate of 1° per minute. Time-of-flight (TOF) neutron powder diffraction (NPD) data were collected for the samples LSKNCO2, LSKNCO5, and LSKNCO9 using the General-Purpose Powder Diffractometer (GPPD) at the China Scattering Neutron Source (CSNS, Dongguan, China). Rietveld refinements were performed for the XRD patterns and neutron diffraction patterns using TOPAS-Academic software.<sup>23</sup> Selected area electron diffraction (SAED) was performed on a JEM2100 (accelerating voltage: 200 kV). X-ray photoelectron spectroscopy (XPS) was carried out using a Kratos Axis Ultra spectrometer (UK) at a voltage of 15 kV and a current of 15 mA. The vacuum of the instrument during the test was about  $10^{-10}$  mbar, and the chamber pressure was not greater than  $5.0 \times 10^{-9}$  torr. All the acquired energy spectral data were calibrated with the C 1s emission at  $E_b = 284.8 \text{ eV}$ . Direct-current (dc) magnetic susceptibility was recorded in a magnetic field of 10 Oe while heating the sample from 2 K to 300 K after zero-field cooling (ZFC) and field cooling (FC) using an MPMS-3 Superconducting Quantum Interference Device (SQUID, Quantum Design) magnetometer. The resistivity was investigated with a cryogenic liquid helium-free physical property measurement system

(PPMS, supplied by East Changing, China) from 2 to 120 K. Scanning electron microscopy (SEM) imaging and elemental analysis through energy-dispersive X-ray spectroscopy (EDS) were performed using an SU8600 (Hitachi, Japan) field-emission scanning electron microscope equipped with an XFlash 760 (Bruker, Germany) EDS spectrometer.

## 3. Results and discussion

### 3.1 Solid solution and structure of $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$

The powder XRD patterns of samples LSKNCO1 to LSKNCO11 are depicted in Fig. 1. The powder XRD patterns of LSKNCO2 to LSKNCO7 closely resemble that of LSKNCO1 ( $\text{La}_{1.75}\text{Sr}_{0.25}\text{CuO}_{4-\delta/4}$ ), suggesting a similarity in their structures to  $\text{La}_{1.75}\text{Sr}_{0.25}\text{CuO}_{4-\delta/4}$ , which was reported to crystallize in a tetragonal space group.<sup>17</sup> When more Na and K are doped into the compound, some reflection peaks are split, as shown at the right hand of Fig. 1, at 2 theta of about 33.5° for the samples LSKNCO8 to LSKNCO11, which may mean that these samples crystallize in the orthogonal space group, similar as that of  $\text{La}_2\text{CuO}_{4-\delta}$ .<sup>3</sup> Therefore, the structures of  $\text{La}_{1.75}\text{Sr}_{0.25}\text{CuO}_{4-\delta/4}$  and  $\text{La}_2\text{CuO}_{4-\delta}$  were used as the starting structure in the Rietveld refinement of the X-ray diffraction patterns of LSKNCO1 to LSKNCO7 and LSKNCO8 to LSKNCO11, respectively. Reasonable refinement results were obtained with  $R_p^x \leq 0.032$ ,  $R_{wp}^x \leq 0.045$ . Neutron diffraction data of the selected samples LSKNCO2 (tetragonal phase), LSKNCO5 (tetragonal phase), and LSKNCO9 (orthogonal phase) were obtained and are refined well using the same lattice parameters and atomic coordination as those used in the refinement of the X-ray diffraction patterns for the corresponding samples with  $R_p^n \leq 0.048$ ,  $R_{wp}^n \leq 0.062$ , which confirms that the results obtained from the X-ray diffraction data are plausible. Typical Rietveld plots of the X-

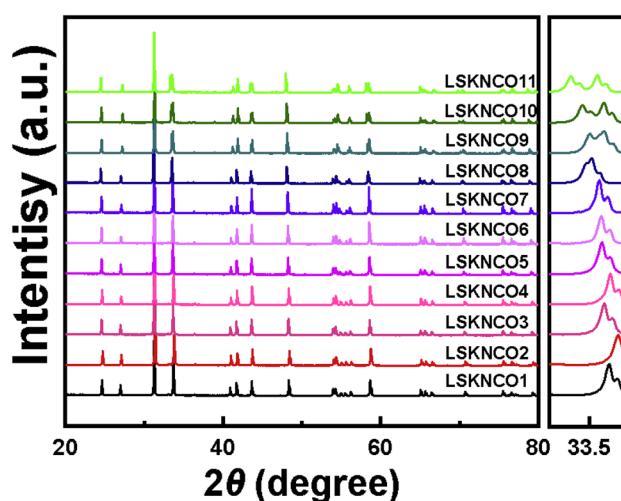


Fig. 1 X-ray diffraction data of  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$ . ( $x = 0.000$  (LSKNCO1), 0.025 (LSKNCO2), 0.050 (LSKNCO3), 0.075 (LSKNCO4), 0.100 (LSKNCO5), 0.125 (LSKNCO6), 0.150 (LSKNCO7), 0.175 (LSKNCO8), 0.200 (LSKNCO9), 0.225 (LSKNCO10) and 0.250 (LSKNCO11)).



ray and neutron diffraction data for  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  with  $x = 0.1$  and  $0.2$  are shown in Fig. 2 and the corresponding refinement details are listed in Table 1 (the others are shown in Fig. S1 to S9 and Tables S1 to S4 in ESI<sup>†</sup>).

Through the Rietveld refinement, the lattice parameters and unit cell volume of  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  were obtained and are presented in Fig. 3, where the data corresponding to the  $\sqrt{2}a_T$  and  $2V_T$  of the tetragonal phase were used to simplify the comparison. The lattice parameter  $a$  of the tetragonal  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  phase (noted as  $a_T$ ) increases with an increase in the Na and K content until  $x \approx 0.16$ . When  $x > 0.16$ , the orthogonal  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  phase appears. The lattice parameters  $a$  and  $b$  of the  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  phase (noted as  $a_o$  and  $b_o$ , respectively) also increase with an increase in the Na and K content. The lattice parameter  $c$  of the tetragonal  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  ( $c_T$ ) and the orthogonal  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  ( $c_o$ ) are similar to each other, as shown in Fig. 3b. They decrease in a different slope with the increase in the Na and K content in the samples with  $x = 0.16$  as the boundary. Simultaneously, the unit cell volume of the tetragonal  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  and the orthogonal  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  increases in a different slope before

and after  $x = 0.16$ . Therefore, the tetragonal to orthogonal phase transition occurs at about  $x = 0.16$  for the  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  system.

It is known that La, Sr, Na, K in  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  are situated at the same crystallographic site (noted as the La site here), which is nine-coordinated to oxygen atoms in the  $I4/mmm$  or  $Fm\bar{m}m$  space group. Therefore, the radius of  $\text{La}^{3+}$ ,  $\text{Sr}^{2+}$ ,  $\text{Na}^+$ , and  $\text{K}^+$  under nine-coordination is used to consider the change in the lattice parameters, which are  $1.212 \text{ \AA}$ ,  $1.31 \text{ \AA}$ ,  $1.24 \text{ \AA}$ , and  $1.55 \text{ \AA}$ , respectively. The change  $\Delta$  in the average radius of La site is denoted as follows.

$$\Delta = (2r_{\text{La}^{3+}}(\text{IX}) + r_{\text{K}^+}(\text{IX}) + r_{\text{Na}^+}(\text{IX}) - 4r_{\text{Sr}^{2+}}(\text{IX}))x/4 \\ = -0.0065x \text{ (\AA)} \quad (1)$$

Eqn (1) indicates that the average radius of the La site should decrease with an increase in the Na and K contents in the samples, which will reduce the volume of the unit cell of the sample. However, the data shown in Fig. 3b tell us that the volume of the unit cell of  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  increases with an increase in the Na and K contents in the sample, which means that the radius data for  $\text{La}^{3+}$ ,  $\text{Sr}^{2+}$ ,  $\text{Na}^+$ , and  $\text{K}^+$  under

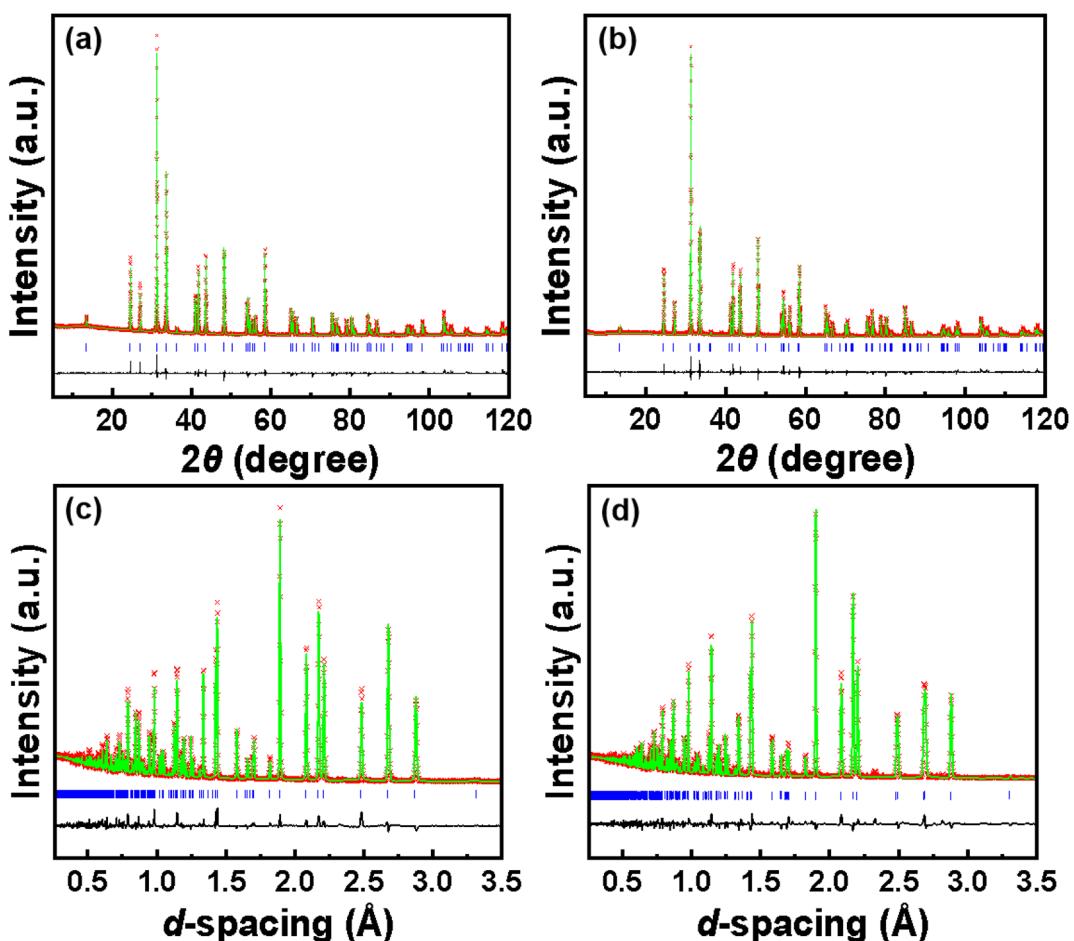


Fig. 2 Rietveld plot of the X-ray (a) LSKNCO5; (b) LSKNCO9 and neutron diffraction data (c) LSKNCO5; (d) LSKNCO9. The plus symbol represents the observed value, the solid line presents the calculated value, the marks below the diffraction patterns are the calculated reflection positions, and the difference curve is shown at the bottom in the figure.



Table 1 Rietveld refinement details of  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  with  $x = 0.100$  and  $0.200$ 

| Sample  | LSKNCO5, $x = 0.100$  | LSKNCO9, $x = 0.200$  |
|---|---|---|
| Lattice parameters ( $\text{\AA}$ )   | $a = b = 3.7781(2)$ , $c = 13.2411(5)$  | $a = 5.3540(2)$ , $b = 5.3769(1)$ , $c = 13.1962(5)$                                      |
| Space group   | $I4/mmm$  | $Fm\bar{m}m$  |
| Atom  | $(x, y, z)$   | $(x, y, z)$   |
| $\text{La}/\text{Sr}/\text{K}/\text{Na}^a$  | $0, 0, 0.3609(3)$   | $0, 0, 0.3611(3)$   |
| $\text{Cu1}$  | $0, 0, 0$   | $0, 0, 0$   |
| $\text{O1}$   | $0, 1/2, 0$   | $1/4, 1/4, 0$   |
| $\text{O2}$   | $0, 0, 0.1814(6)$   | $0, 0, 0.1867(8)$   |
| $B_{\text{eq}}(\text{La}/\text{Sr}/\text{K}/\text{Na})$ ( $\text{\AA}^2$ ) <sup>b</sup> | $1.02(1)$   | $0.65(1)$   |
| $B_{\text{eq}}(\text{Cu})$ ( $\text{\AA}^2$ )   | $1.29(1)$   | $0.99(1)$   |
| $B_{\text{eq}}(\text{O1})$ ( $\text{\AA}^2$ )   | $1.17(1)$   | $0.55(1)$   |
| $B_{\text{eq}}(\text{O2})$ ( $\text{\AA}^2$ )   | $2.13(1)$   | $2.42(2)$   |
| $R$ factor <sup>c</sup>   | $R_{\text{wp}}^x = 0.040$ , $R_{\text{wp}}^n = 0.062$ , $R_p^x = 0.020$ , $R_p^n = 0.044$ | $R_{\text{wp}}^x = 0.044$ , $R_{\text{wp}}^n = 0.054$ , $R_p^x = 0.032$ , $R_p^n = 0.048$ |

<sup>a</sup> The occupancy of  $\text{La}/\text{Sr}/\text{K}/\text{Na}$  is  $0.9000 : 0.0750 : 0.0125 : 0.0125$  for  $x = 0.100$ , and  $0.925 : 0.025 : 0.025 : 0.025$  for  $x = 0.200$ . <sup>b</sup>  $B_{\text{eq}}$  is the thermal displacement parameter. <sup>c</sup>  $R_{\text{wp}}^x$  and  $R_{\text{wp}}^n$  are the weighted  $R$  factors calculated by  $(\text{sum}(w(I_0 - I_c)^2)/\text{sum}(w(I_0)^2))^{0.5}$  for powder X-ray and neutron diffraction data, respectively;  $R_p^x$  and  $R_p^n$  are the  $R$  factors calculated by  $\text{sum}(|I_0 - I_c|)/\text{sum}(I_0)$  for powder X-ray and neutron diffraction data, respectively.

nine-coordination are not suitable for the present compound  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$ . For the  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  system, the average radius of the La site should increase with four  $\text{Sr}^{2+}$  replaced by two  $\text{La}^{3+}$ , one  $\text{Na}^+$  and one  $\text{K}^+$ . Then, the increase in the lattice parameters  $a_{\text{T}}$ ,  $a_{\text{o}}$ , and  $b_{\text{o}}$  can be attributed to this increase in the average radius of the La site. The decrease in the lattice parameters  $c_{\text{T}}$  and  $c_{\text{o}}$  may be explained by the “shrinkage effect”: expansion of the  $a_{\text{T}}$ ,  $a_{\text{o}}$ , and  $b_{\text{o}}$  axis allows the layers to be closer together to shrink the  $c_{\text{T}}$  and  $c_{\text{o}}$  axis. This phenomenon is similar to the findings in  $\text{ZrSi}_{1-x}\text{Ge}_x\text{Te}$  solid solutions by C. Wang *et al.*<sup>24</sup> A. Kimura *et al.* also found that on adding larger Cr and Ti atoms into the AlN lattice, the hexagonal lattice spacings expanded in the  $a$ -direction and shrank in the  $c$ -direction.<sup>25</sup>

It was found that the lattice parameter  $a$  of  $\text{La}_{2-x}\text{Sr}_x\text{CuO}_{4-\delta}$  ( $0.1 \leq x \leq 0.25$ ) reported by J. M. Tarascon<sup>26</sup> also decreases with an increase in Sr, which is very similar to that of our  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  samples (shown in Fig. S10†). However, the lattice parameter  $c$  and the unit cell volume of  $\text{La}_{2-x}\text{Sr}_x\text{CuO}_{4-\delta}$  are smaller than those of  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  with the same amount of Sr in the

sample (shown in Fig. S10†), which supports that  $\text{K}^+$  and  $\text{Na}^+$  should occupy the La sites as these larger ions expand the lattice (see ESI† for the details).

The SAED (selected-area electron diffraction) patterns are very useful to check the choice of the space group for the studied materials.<sup>27,28</sup> Therefore, the SAED patterns of the selected samples in the  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  system have been obtained with the typical data shown in Fig. 4 for LSKNCO5. The tetragonal space group  $I4/mmm$  with lattice parameters  $a \approx 3.778 \text{ \AA}$ ,  $b \approx 3.778 \text{ \AA}$ ,  $c \approx 13.241 \text{ \AA}$ ,  $\alpha = 90^\circ$ ,  $\beta = 90^\circ$ ,  $\gamma = 90^\circ$  can adequately index the corresponding points, indicating that the space group  $I4/mmm$  can be used to describe the structures of LSKNCO5. As the X-ray diffraction patterns of LSKNCO1 to LSKNCO7 are almost the same as that of LSKNCO5, the space group  $I4/mmm$  can also be used to describe the structures of LSKNCO1 to LSKNCO7.

In order to access the contents of Na and K in  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$ , EDX analysis was performed on LSKNCO9 and LSKNCO11 (see Fig. S11 and S12†). It was found that the La : Sr : K : Na : Cu ratio of LSKNCO9 was 27.88 : 0.68 : 0.76 : 0.82 : 15.76 (corresponding to 7.34 : 0.18 : 0.20 : 0.22 :

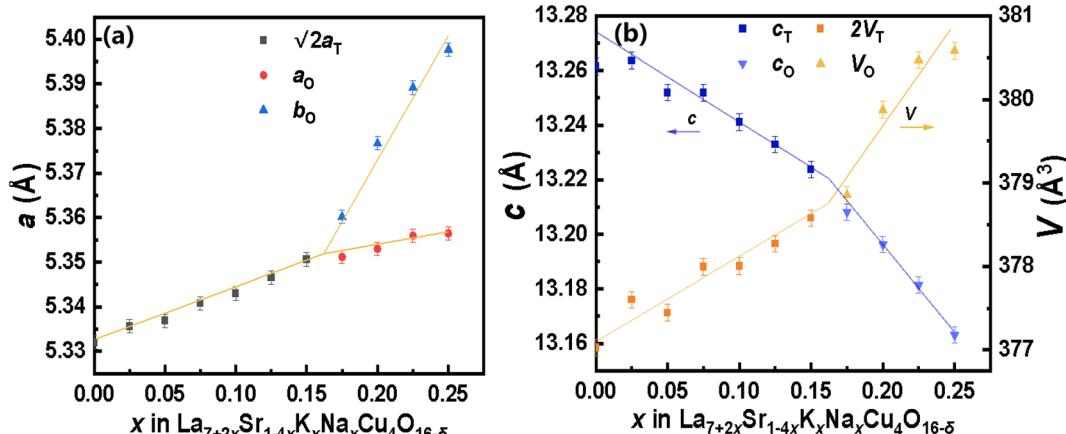


Fig. 3 Lattice parameters of  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  ( $0 \leq x \leq 0.25$ ).  $a_{\text{T}}$ ,  $c_{\text{T}}$ ,  $V_{\text{T}}$ , the lattice parameter  $a$ ,  $c$ , and the volume of unit cell for the tetragonal phase;  $a_{\text{o}}$ ,  $b_{\text{o}}$ ,  $c_{\text{o}}$ ,  $V_0$ , the lattice parameter  $a$ ,  $b$ ,  $c$ , and the volume of the unit cell for the orthorhombic phase.



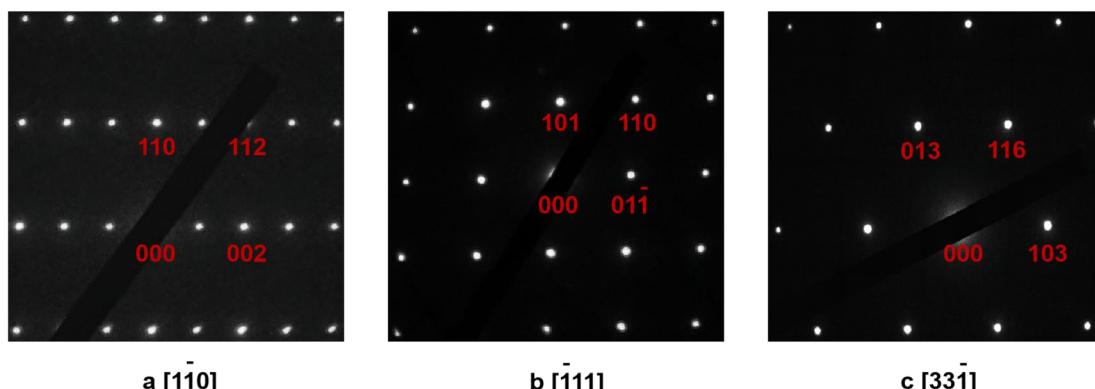


Fig. 4 The electron diffraction patterns of LSKNCO5 ( $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  with  $x = 0.1$ ) along [331] (a), [111] (b) and [110] (c) indexed by space group  $I\bar{4}/m\bar{m}m$  with  $a \approx 3.778 \text{ \AA}$ ,  $b \approx 3.778 \text{ \AA}$ ,  $c \approx 13.241 \text{ \AA}$ ,  $\alpha = 90^\circ$ ,  $\beta = 90^\circ$  and  $\gamma = 90^\circ$ .

4.15), and the La : Na : Cu ratio of LSKNCO11 was 28.23 : 0.82 : 0.94 : 14.70 (corresponding to 8.02 : 0.23 : 0.27 : 4.18), which are essentially identical to their ideal values. Therefore, the chemical composition of the samples aligns with the planned formula.

### 3.2 Valence of Na and Cu in $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$

As widely recognized, copper in most copper oxides exists in the oxidation states of +1, or +2. However, the presence of  $\text{Sr}^{2+}$ ,  $\text{Na}^+$ , and  $\text{K}^+$  in  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  can induce the partial oxidation of  $\text{Cu}^{2+}$  to  $\text{Cu}^{3+}$ . In order to accurately understand the changes in the oxidation state of copper in the samples, the X-ray photoelectron spectroscopy (XPS) data of the samples LSKNCO1, LSKNCO3, LSKNCO5, LSKNCO7, LSKNCO9, and LSKNCO11 was obtained. The full-scan XPS measurement spectrum of the LSKNCO9 sample is presented in Fig. 5a, which

indicates the presence of La, Sr, K, Na, Cu, and O elements in the sample (with C being introduced by the XPS instrument itself). High-resolution element-specific fine scans of Na 1s, Cu 2p, and O 1s were recorded, which will be discussed in detail later.

As shown in Fig. 5b, Na 1s XPS have no significant signals for the samples LSKNCO1 and LSKNCO3. This is probably due to the low doping concentration of  $\text{Na}^+$ . As the Na content in the samples increases, prominent peaks are observed at an energy position of about 1071.58 eV in LSKNCO5, LSKNCO7, LSKNCO9, and LSKNCO11, corresponding to the Na 1s orbital. The  $\text{Na}^+$  state (binding energy  $\sim$ 1071.0–1071.5 eV) can be confirmed based on the commonly used line position diagrams.<sup>29,30</sup> Due to the limited sensitivity of the spectrometer to K elements, the signal intensity for K is not significant, and the spectra for K elements were not measured.

Fig. 5c shows the Cu 2p XPS spectra of all the studied components containing  $\text{Cu} 2p_{3/2}$  and  $\text{Cu} 2p_{1/2}$  as well as satellite peaks. The main peaks with binding energies at  $\sim$ 933.14 eV and  $\sim$ 952.62 eV correspond to  $\text{Cu}^{2+}$  at  $\text{Cu} 2p_{3/2}$  and  $\text{Cu} 2p_{1/2}$ , respectively.<sup>31–33</sup> However, the peaks at higher energies of the binding energy at  $\sim$ 934.98 eV and  $\sim$ 954.63 eV correspond to the  $\text{Cu}^{3+}$  of  $\text{Cu} 2p_{3/2}$  and  $\text{Cu} 2p_{1/2}$ , respectively.<sup>31</sup> Rocking satellite peaks at binding energies 940.79 eV and 943.31 eV were observed, which can be assigned to  $\text{Cu}^{2+}$  and  $\text{Cu}^{3+}$ , respectively. The  $\text{Cu}^{2+} : \text{Cu}^{3+}$  ratios in these samples were obtained by fitting the XPS spectra of Cu 2p, as shown in Fig. 5d, which are listed in Table 2 and presented in Fig. 6. The  $\text{Cu}^{2+} : \text{Cu}^{3+}$  ratio of the

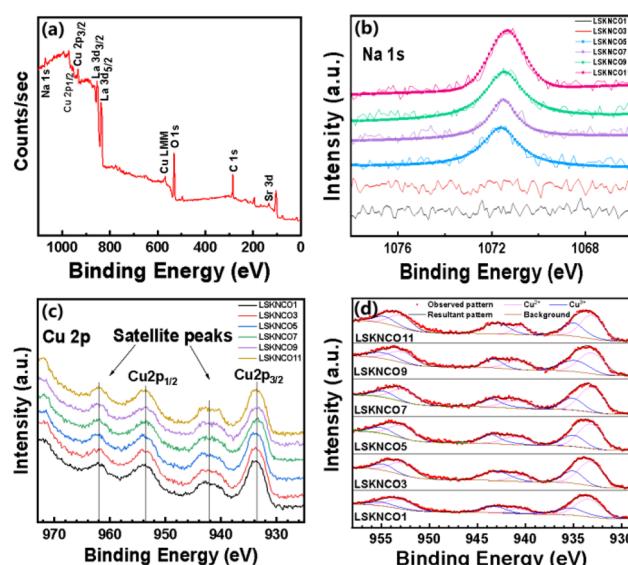


Fig. 5 Full scan spectra of LSKNCO9 (a), and the XPS spectra of Na 1s (b),  $\text{Cu} 2p_{3/2}$  and  $\text{Cu} 2p_{1/2}$  (c), the  $\text{Cu} 2p_{3/2}$  and  $\text{Cu} 2p_{1/2}$  XPS peak fitted with three different Gaussian peaks (d) for LSKNCO1, LSKNCO3, LSKNCO5, LSKNCO7, LSKNCO9 and LSKNCO11.

Table 2 Binding energies (eV) of  $\text{Cu} 2p_{3/2}$ ,  $\text{Cu} 2p_{1/2}$ , and the  $\text{Cu}^{2+} : \text{Cu}^{3+}$  ratio for  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$

| Sample   | $\text{Cu}^{2+}$  | $\text{Cu}^{3+}$  | $\text{Cu}^{2+}$ | $\text{Cu}^{3+}$ | $\text{Cu}^{2+} : \text{Cu}^{3+}$ |
|----------|-------------------|-------------------|------------------|------------------|-----------------------------------|
|          | 2p <sub>3/2</sub> | 2p <sub>1/2</sub> |                  |                  |                                   |
| LSKNCO1  | 933.40            | 935.19            | 953.00           | 954.90           | 0.47 : 0.53                       |
| LSKNCO3  | 933.20            | 935.04            | 952.79           | 954.76           | 0.70 : 0.30                       |
| LSKNCO5  | 933.22            | 935.00            | 952.81           | 954.96           | 0.83 : 0.17                       |
| LSKNCO7  | 933.14            | 935.10            | 952.62           | 954.76           | 0.72 : 0.28                       |
| LSKNCO9  | 933.11            | 934.86            | 952.85           | 954.76           | 0.62 : 0.38                       |
| LSKNCO11 | 933.10            | 934.98            | 952.98           | 954.63           | 0.54 : 0.46                       |



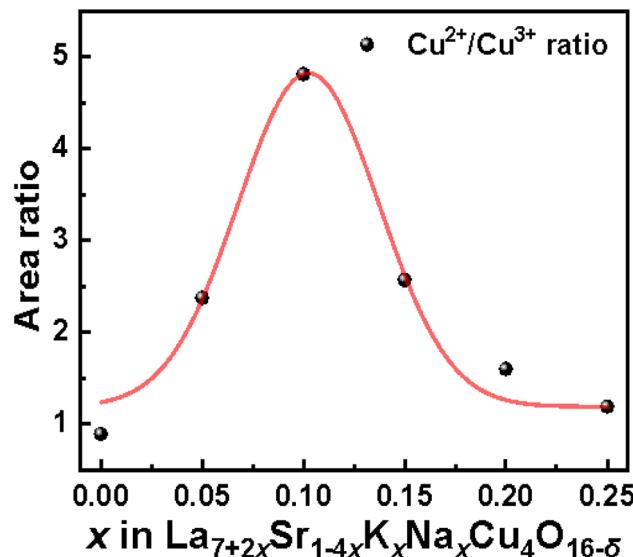


Fig. 6 Area ratio of  $\text{Cu}^{2+}/\text{Cu}^{3+}$  in  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$ .

substituted samples gradually increased in the doped samples to a maximum value of 4.882 at  $x = 0.10$ , followed by a gradual decrease, which indicates that the valence state of Cu firstly shifts from  $\text{Cu}^{3+}$  to  $\text{Cu}^{2+}$  and then moves back to  $\text{Cu}^{3+}$ . R. A. M. Ram *et al.* mentioned for the  $(\text{La}, \text{Ln})_{2-x}(\text{Ba}, \text{Sr})_x\text{CuO}_{4-\delta}$  system that the substitution of  $\text{La}^{3+}$  with  $\text{Sr}^{2+}$  usually oxidizes  $\text{Cu}^{2+}$  to  $\text{Cu}^{3+}$ .<sup>34</sup> However, in the  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  system, the oxidation tendency caused by the replacement of two  $\text{Sr}^{2+}$  by one  $\text{Na}^+$  and one  $\text{K}^+$  is cancelled by the co-replacement of two  $\text{Sr}^{2+}$  by two  $\text{La}^{3+}$ . The change in the valence of Cu in the  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  system may be due to other reasons.

### 3.3 Oxygen vacancy in $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$

The O 1s XPS spectra for all the investigated compositions are shown in Fig. 7a with two peaks at binding energies of 528.7 eV and 531.4 eV, denoted as O1 and O2, respectively. In the study on  $\text{Ga}^{3+}$ -doped  $\text{BiMn}_{1-x}\text{Ga}_x\text{O}_3\text{-BaTiO}_3$  ceramics, K. K. Rahangdale<sup>35</sup> found that the two peaks at binding energies of 527.75 eV and 529.0–532.0 eV in the O 1s XPS spectra are related to the lattice oxygen without oxygen vacancies in the neighboring vicinity and vacancy-related oxygen with oxygen vacancies in the neighboring vicinity, respectively. Additionally, D. Kumar *et al.*<sup>36</sup> in  $\text{Ti}^{4+}$ -doped  $\text{La}_{0.6}\text{Ba}_{0.4}\text{Mn}_{1-x}\text{Ti}_x\text{O}_3$  perovskites found the binding energies of lattice oxygen and vacancy related oxygen to be approximately 529.0–530.2 eV and 531.3–531.7 eV, respectively. Therefore, the binding energies O1 and O2 in our study may correspond to lattice oxygen without oxygen vacancies in the neighboring vicinity and vacancy related oxygen with oxygen vacancies in the neighboring vicinity, respectively.

Fig. 7b shows the fitted spectra of the two peaks of the selected samples. When  $x \leq 0.10$ , the intensity of the O2 peak ( $I_{\text{O}2}^{1s}$ ) increases with increasing doping concentration of K and Na. However, there is a decreasing trend in the intensity of the O2 peak when  $x > 0.10$ . The  $I_{\text{O}2}^{1s} : I_{\text{O}1}^{1s}$  ratio denoted by  $R_I$  for each sample is listed in Table 3 and shown in Fig. 7c. Compared to

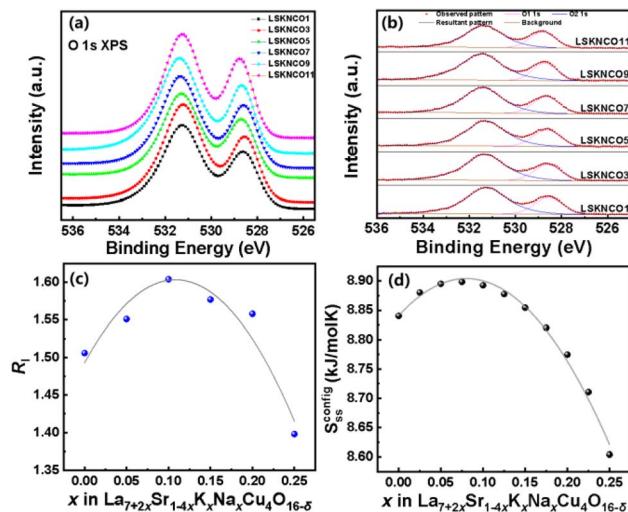


Fig. 7 The O 1s XPS spectra (a), peak fitting of O 1s XPS spectra (b), ratio  $I_{\text{O}2}^{1s} : I_{\text{O}1}^{1s}$  (c) of LSKNCO1, LSKNCO3, LSKNCO5, LSKNCO7, LSKNCO9 and LSKNCO11, and configurational entropy (d) of all the samples (LSKNCO1 to LSKNCO11).

Table 3 Binding energies (eV) of O1 and O2, the ratio  $I_{\text{O}2}^{1s} : I_{\text{O}1}^{1s}$  ( $R_I$ ) for  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$

| Sample   | O1 1s  | O2 1s  | $R_I$ |
|----------|--------|--------|-------|
| LSKNCO1  | 528.56 | 531.24 | 1.506 |
| LSKNCO3  | 528.61 | 531.32 | 1.551 |
| LSKNCO5  | 528.65 | 531.29 | 1.604 |
| LSKNCO7  | 528.66 | 531.43 | 1.577 |
| LSKNCO9  | 528.74 | 531.47 | 1.558 |
| LSKNCO11 | 528.73 | 531.35 | 1.398 |

Fig. 6, the  $I_{\text{O}2}^{1s} : I_{\text{O}1}^{1s}$  ratio can correspond well with the  $\text{Cu}^{2+} : \text{Cu}^{3+}$  ratio, which means that the change in the valence of Cu in the  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  system is related to the change in the content of oxygen vacancies in the samples. In fact, as  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  is an equivalent doping system, it is natural that the  $\text{Cu}^{2+} : \text{Cu}^{3+}$  ratio is directly related to the change in the oxygen vacancies. Therefore, the origin for the change in the  $\text{Cu}^{2+} : \text{Cu}^{3+}$  ratio may be the same as the origin of the change in the  $I_{\text{O}2}^{1s} : I_{\text{O}1}^{1s}$  ratio.

As shown in Fig. 3, the change in the lattice parameters of  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  is monotonous, which could not induce a parabolic change in the oxygen vacancies shown in Fig. 7c or a parabolic change in the  $\text{Cu}^{2+} : \text{Cu}^{3+}$  ratio shown in Fig. 6 with the increase in the  $\text{Na}^+$  and  $\text{K}^+$  contents in the samples. Therefore, it is estimated that the change in the content of oxygen vacancies in  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  may be related to the configurational entropy change in this system with different amounts of Na and K doping. The configurational entropy can be calculated by the following equation.<sup>37</sup>

$$S_{ss}^{\text{config}} = -R \sum_i X_i \ln(X_i) \quad (2)$$

where  $R$  is the gas constant, and  $X_i$  is the mole fraction of each element. The calculated results for  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$



are in the range of  $1.03\text{--}1.07R$  (the details are listed in ESI†), which is consistent with the  $A_2BO_4$ -type medium entropy material ( $1\text{--}1.5R$ ) discussed by M. Brahlek *et al.*<sup>38</sup> As shown in Fig. 7d, the entropy has a maximum at about  $x \approx 0.10$  with Na and K doping variations, which is similar to the change in the  $I_{O2}^{1s} : I_{O1}^{1s}$  ratio shown in Fig. 7c and the change in the  $Cu^{2+} : Cu^{3+}$  ratio shown in Fig. 6. The change in the content of oxygen vacancies in  $La_{7+2x}Sr_{1-4x}K_xNa_xCu_4O_{16-\delta}$  and the change in the  $Cu^{2+} : Cu^{3+}$  ratio may be due to the change in the configurational entropy caused by the Na and K doping.

### 3.4 Superconducting properties of $La_{7+2x}Sr_{1-4x}K_xNa_xCu_4O_{16-\delta}$

Temperature-dependent resistivity measurements were conducted on all the samples from LSKNCO1 to LSKNCO11 in the temperature range from 2 to 120 K. The typical data for the samples LSKNCO1, LSKNCO5, LSKNCO8, and LSKNCO9 are shown in Fig. 8a-d (the others are presented in the ESI†).

For the samples LSKNCO1 to LSKNCO6, the resistivity decreases with decreasing temperature to show a metallic behavior until the temperature reaches  $T_c^{\text{onset}}$ . After  $T_c^{\text{onset}}$ , the resistivity drops sharply with a decrease in temperature. Subsequently, as the temperature continues to decrease, the resistivity reaches "zero" at  $T_c^{\text{zero}}$ .  $T_c^{\text{onset}}$  is not changed significantly for these six samples, which is about 38 K, as shown in Fig. 8e. However, as shown in Fig. 8f, the  $T_c^{\text{zero}}$  of  $La_{7+2x}Sr_{1-4x}K_xNa_xCu_4O_{16-\delta}$  firstly increases and then decreases, which is a surprise at first glance. The main reason is that with an increase in the contents of Na and K in the sample, the average valence at the La site remains unchanged because the decrease tendency caused by the replacement of two  $Sr^{2+}$  with one  $Na^+$  and one  $K^+$  is cancelled by the replacement of two  $Sr^{2+}$

with two  $La^{3+}$ . In this case, the valence of Cu should not change. Therefore, the  $T_c^{\text{zero}}$  of  $La_{7+2x}Sr_{1-4x}K_xNa_xCu_4O_{16-\delta}$  should remain constant at about 16.5 K, as reported by L. Weckhuysen,<sup>39</sup> because most of us believe that the  $T_c^{\text{zero}}$  of doped  $La_2CuO_{4-\delta}$  is strongly related to the valence of Cu, as indicated by P. G. Radaelli *et al.*<sup>6</sup> However, the XPS data shown in Table 2 in section 3.2 indicates that the  $Cu^{2+} : Cu^{3+}$  ratio in  $La_{7+2x}Sr_{1-4x}K_xNa_xCu_4O_{16-\delta}$  first increases and then decreases. This means that the amount of  $Cu^{3+}$  in  $La_{7+2x}Sr_{1-4x}K_xNa_xCu_4O_{16-\delta}$  first decreases with an increase in  $x$  when  $x < 0.100$ , then increases when  $0.100 < x < 0.25$ . As mentioned by P. G. Radaelli *et al.*,<sup>6</sup>  $La_{1.75}Sr_{0.25}CuO_{4-\delta/4}$  (which is corresponding to  $La_{7+2x}Sr_{1-4x}K_xNa_xCu_4O_{16-\delta}$  with  $x = 0.000$ ) is a hole over-doped superconductor. Therefore, the decrease in the holes (corresponding to the decrease in the  $Cu^{3+}$  content) in  $La_{7+2x}Sr_{1-4x}K_xNa_xCu_4O_{16-\delta}$  with an increase in  $x$  when  $x < 0.100$  will help to move to the optimal doped state. Then, the  $T_c^{\text{zero}}$  of  $La_{7+2x}Sr_{1-4x}K_xNa_xCu_4O_{16-\delta}$  increases with an increase in  $x$  when  $x < 0.100$ . The  $T_c^{\text{zero}}$  of  $La_{7+2x}Sr_{1-4x}K_xNa_xCu_4O_{16-\delta}$  decreases from 33.9 K for  $x = 0.100$  to 26.7 K for  $x = 0.125$ , which can be due to the increase in the holes to move the sample back to the over-doped state.

With more Na and K doped into the sample, the metallic resistivity above  $T_c^{\text{onset}}$  for LSKNCO1 to LSKNCO6 changes to semiconductive resistivity for LSKNCO7 and LSKNCO8. This may be because the doping level directly affects the carrier concentration and metallic nature of the material. For samples LSKNCO1 ( $x = 0$ ) and LSKNCO5 ( $x = 0.1$ ), the doping level is low and the carrier concentration is sufficiently high to form effective conduction channels, which enhances the metallic properties. As the temperature decreases, superconductivity emerges, leading to increased carrier mobility and reduced

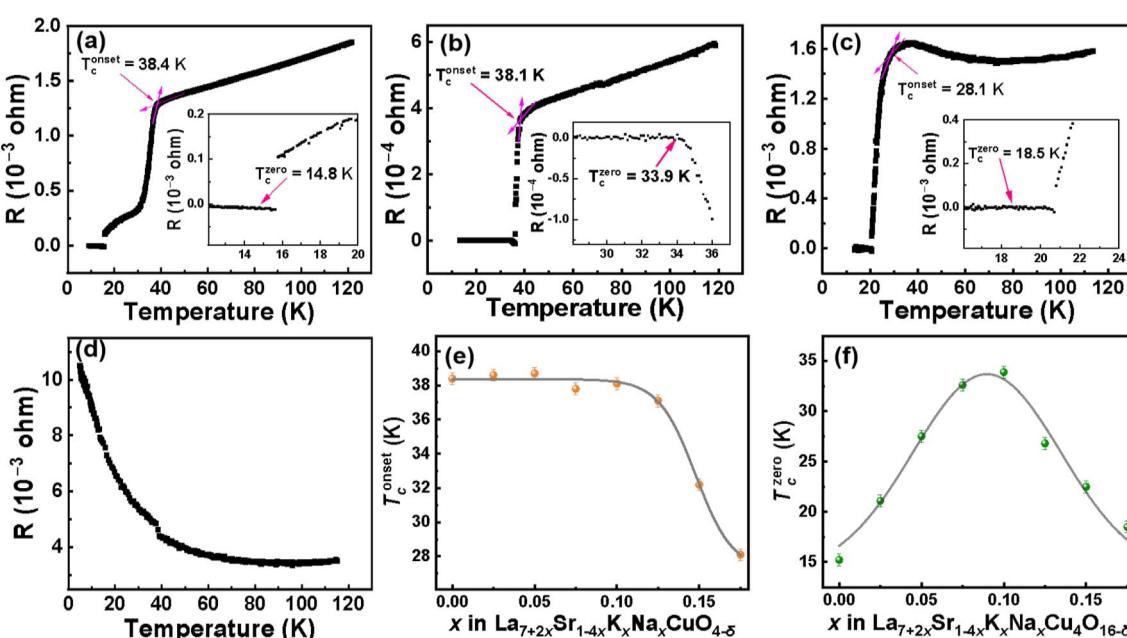


Fig. 8 Temperature-dependent resistance of LSKNCO1 (a), LSKNCO5 (b), LSKNCO8 (c), LSKNCO9 (d),  $T_c^{\text{onset}}$  ( $0 \leq x \leq 0.175$ ) (e) and  $T_c^{\text{zero}}$  of the samples  $La_{7+2x}Sr_{1-4x}K_xNa_xCu_4O_{16-\delta}$  ( $0 \leq x \leq 0.175$ ) (f).



electron scattering, which results in a decrease in resistivity.<sup>41,40</sup> In contrast, for LSKNCO7 ( $x = 0.15$ ) and LSKNCO8 ( $x = 0.175$ ) (Fig. S18† and 8(c)), the higher doping levels introduce increased disorder, leading to the presence of defects. This disorder enhances electron scattering, thereby hindering the decrease in resistivity, even though the materials exhibit superconductivity at low temperatures.<sup>41</sup> Additionally, the increased disorder due to higher doping levels may localize some carriers, preventing them from effectively contributing to conductivity at higher temperatures.<sup>42</sup> As the temperature decreases, these localized carriers contribute less to the metallic behavior, resulting in different resistivity characteristics compared to the lower-doped samples.  $T_c^{\text{onset}}$  decreases from 32.2 K for LSKNCO7 to 28.1 K for LSKNCO8, as shown in Fig. 8e, and  $T_c^{\text{zero}}$  continues to decrease from 22.5 K for LSKNCO7 to 18.5 K for LSKNCO8. The reason is similar to that for LSKNCO6, as mentioned above. It is noted that LSKNCO7 is tetragonal and LSKNCO8 is orthogonal, which means that the superconductivity can appear in both the tetragonal and orthogonal phases for the  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  system.

With a further increase in the Na and K contents in the sample, “zero” resistivity is not measured for the samples LSKNCO9 to LSKNCO11 with a semiconductive behavior for the temperature-dependent resistivity. The data for the temperature-dependent resistance of LSKNCO10 and LSKNCO11 is shown in Fig. S20–S21.†

In order to confirm the superconductivity of the  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  system, the magnetization of the samples was measured under a magnetic field of 10 Oe in the temperature range of 2–300 K. Typical data are shown in Fig. 9 and the data for other samples are presented in ESI (Fig. S13–S21†). As the temperature decreases, the magnetization of the samples starts to decrease to negative values at  $T_c^{\text{mag}}$ , which are 37.5 K and 28.0 K for LSKNCO5 and LSKNCO8, respectively. They both exhibit strong diamagnetic signals, confirming that superconductivity exists in these samples as “zero” resistivity is already observed above. Weak diamagnetic signals (about 2% to 0.1% of that for LSKNCO5 or LSKNCO8) are found in the magnetization of the samples for LSKNCO9 to LSKNCO11. Since the sample LSKNCO9 no longer exhibits “zero” resistivity, the temperature-dependent DC magnetization data (Fig. S19†) reveals that the zero-field-cooling (ZFC) and field-cooling (FC)

curves do not show any splitting, which may be attributed to weak flux pinning in the material.<sup>43</sup> This observation further suggests the absence of superconductivity properties in the sample. Additionally, the incomplete diamagnetic signals observed in the magnetic susceptibility measurements of LSKNCO10 and LSKNCO11 further corroborate the loss of superconductivity in these materials.

As discussed in section 3.3, the change in the  $\text{Cu}^{2+}:\text{Cu}^{3+}$  ratio is due to the change in the configurational entropy caused by the Na and K doping in the present equivalent doping  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  system. Therefore, the  $T_c^{\text{zero}}$  of  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  is increased to 33.9 K (for  $x = 0.10$ ) from 14.8 K (for  $x = 0.00$ ) and then lowered to 18.5 K (for  $x = 0.175$ ), which is related to the change in the configurational entropy caused by Na and K doping.

## 4. Conclusions

In summary, we demonstrated the synthesis of new series of solid solutions  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  ( $0.00 \leq x \leq 0.25$ ) by the traditional solid-state method, which crystallizes in the *I4/mmm* space group when  $0.00 \leq x < 0.16$  and *Fmmm* when  $0.16 < x \leq 0.25$ .

All the tetragonal phase  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  ( $0.00 \leq x < 0.16$ ) are superconductors with  $T_c^{\text{zero}}$  increasing from 14.8 K for  $x = 0.00$  to 33.9 K for  $x = 0.10$  and then decreasing to 22.5 K for  $x = 0.15$ . The change in the  $T_c^{\text{zero}}$  for the equivalent doping system  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  is related to the change in the  $\text{Cu}^{2+}:\text{Cu}^{3+}$  ratio in the samples confirmed by the XPS data, which may be caused by the change in the configurational entropy.

$T_c^{\text{zero}}$  for the orthogonal phase  $\text{La}_{7+2x}\text{Sr}_{1-4x}\text{K}_x\text{Na}_x\text{Cu}_4\text{O}_{16-\delta}$  with  $x = 0.175$  is about 18.5 K. The other samples with  $x = 0.200$ , 0.225, and 0.250 do not show superconductivity in the present study.

## Data availability

The data supporting this article have been included as part of the ESI.† All relevant data are available from the corresponding author upon reasonable request.

## Author contributions

Deyang Xu: methodology, data curation, writing—original draft preparation; Guohong Cai: conceptualization; Peiliang Huang: formal analysis; Xi Wu: formal analysis; Yan Wang: data curation; Jinling Geng: conceptualization; Jing Ju: methodology, data curation; Xiaoge Wang: investigation, formal analysis; Congling Yin: conceptualization, supervision, project administration, writing—review and editing; Guobao Li: conceptualization, formal analysis, supervision, writing—review and editing. All authors have read and agreed to the published version of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

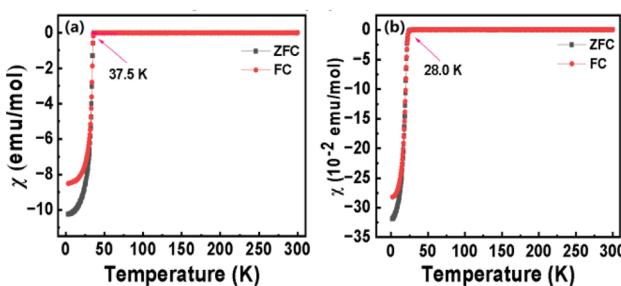


Fig. 9 Temperature dependence of DC magnetic susceptibility of LSKNCO5 (a) and LSKNCO8 (b) in an applied field of 10 Oe after zero field cooling.



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