One-Step Synthesis of LiCo$_{1-1.5x}$Y$_x$PO$_4$@C Cathode Material for High-Energy Lithium-ion Batteries

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Abstract: Intrinsically low ion conductivity and unstable cathode electrolyte interface are two important factors affecting the performances of LiCoPO$_4$ cathode material. Herein, a series of LiCo$_{1-1.5x}$Y$_x$PO$_4$@C ($x = 0, 0.01, 0.02, 0.03$) cathode material is synthesized by a one-step method. The influence of Y substitution amount is optimized and discussed. The structure and morphology of LiCo$_{1-1.5x}$Y$_x$PO$_4$@C cathode material does not lead to obvious changes with Y substitution. However, the Li/Co antisite defect is minimized and the ionic and electronic conductivities of LiCo$_{1-1.5x}$Y$_x$PO$_4$@C cathode material are enhanced by Y substitution. The LiCo$_{0.97}$Y$_{0.03}$PO$_4$@C cathode delivers a discharge capacity of 148 mAh g$^{-1}$ at 0.1 C and 96 mAh g$^{-1}$ at 1 C, with a capacity retention of 75% after 80 cycles at 0.1 C. Its good electrochemical performances are attributed to the following factors. (1) The uniform 5 nm carbon layer stabilizes the interface and suppresses the side reactions with the electrolyte. (2) With Y substitution, the Li/Co antisite defect is decreased and the electronic and ionic conductivity are also improved. In conclusion, our work reveals the effects of aliovalent substitution and carbon coating in LiCo$_{1-1.5x}$Y$_x$PO$_4$@C electrodes to improve their electrochemical performances, and provides a method for the further development of high voltage cathode material for high-energy lithium-ion batteries.

Keywords: lithium-ion battery; high energy; cathode; LiCo$_{1-1.5x}$Y$_x$PO$_4$@C

1. Introduction

With the rapid development of portable devices and electric vehicles, the demand for high energy batteries is increasing. The development of high-energy cathode [1–6] and anode [7] materials is imperative. Olivine LiCoPO$_4$ with a theoretical energy density of about 800 Wh kg$^{-1}$ is a good candidate cathode material for high-energy lithium-ion batteries [8–10]. However, the continuous oxidative decomposition of electrolyte [11,12] and the unstable cathode electrolyte interface [13,14] under 5 V high voltage caused by Co$^{2+}$/Co$^{3+}$, resulting in rapid capacity degradation during cycling, severely hindering the application of LiCoPO$_4$ cathode material. In addition, the Li/Co antisite exchange during the cycling process [15–17] and low intrinsically ionic and electronic conductivity [18–20] must also be overcome.

Lots of work have been undertaken to solve these problems, including decreasing the cathode particle size and controlling the morphology to shorten the Li-ion migration distance [21–24]; coating the cathode particle with stable materials [25–28] or conductive materials [29–33] to stabilize the interface and reduce the side reaction; partial substitution at Co site [20,34–37] to improve the intrinsic ionic and electronic conductivity [34,35,38–40]; and adding electrolyte additives to suppress the electrolyte decomposition [14,41,42]. Every method has some effect in improving LiCoPO$_4$ cathode performance. Generally, for LiCoPO$_4$ cathode material, surface coating is the most effective way to enhance the stability of the interface [25,31,43], while cation substitution can significantly improve the material ionic conductivity [16,36,44,45].
In this work, the LiCo$_{1-1.5x}$Y$_x$PO$_4$@C cathode material, the substitution of Y, and carbon coating are synthesized in one step. The amount of Y substitution is optimized and its influence discussed.

2. Materials and Methods

2.1. Synthesis of LiCo$_{1-1.5x}$Y$_x$PO$_4$@C Cathode Material

LiCo$_{1-1.5x}$Y$_x$PO$_4$@C (x = 0, 0.01, 0.02, 0.03) cathode material was synthesized by a one-step method. First citric acid (CA, Sinopharm Chemical Reagent, ≥ 99.5%), Y(NO$_3$)$_3$·6H$_2$O (Sinopharm Chemical Reagent, ≥ 99.0%), Co(NO$_3$)$_2$·6H$_2$O (Sinopharm Chemical Reagent, ≥ 98.5%), LiNO$_3$ (Sinopharm Chemical Reagent, ≥ 99.9%) and NH$_4$H$_2$PO$_4$ (Sinopharm Chemical Reagent, ≥ 99.0%) were dissolved at stoichiometric amounts (n$_{Li}$:n$_{Co}$:n$_{Y}$:n$_{P}$:n$_{CA}$ = 1:0.95:1-1.5x:x:1:2) in deionized water. Then the gel was formed by heating the solution at 80 °C. Subsequently, the wet gel was dried at 120 °C 24 h to obtain dry gel. Finally, the dry gel was calcined at 400 °C 3 h in rotary furnace at air atmosphere, and then changed to Ar atmosphere and calcined at 700 °C 2 h [39].

2.2. Material Characterization

The crystal information of the materials was detected by XRD (Smart Lab) with Cu Kα radiation, and TOPAS 5.0 software (Bruker AXS, America) was used to Rietveld refinements. The morphology of the materials was observed with SEM, EDS (Hitachi, BCPCAS-4800), and TEM (Tecnai, F20). FTIR spectra were obtained by IR spectrometer (PerkinElmer, Spectrum One).

2.3. Electrochemical Performance

The electrochemical tests were carried out with using the 2025-type coin cell. Coin cells were assembled with the dried LiCo$_{1-1.5x}$Y$_x$PO$_4$@C as cathode, Li metal as anode, Celgard2400 polyethylene as separator, and 1M LiPF$_6$ in a mixture of DMC/EC (v/v, 1/1) with 1 wt.% TMSB additive as electrolyte in an argon-filled glovebox. The LiCo$_{1-1.5x}$Y$_x$PO$_4$@C electrodes were dried at 120 °C 12 h in a vacuum oven.

The cycling and rate performances of LiCo$_{1-1.5x}$Y$_x$PO$_4$@C electrode were measured by the LAND CT2001A. The CV curve of the LiCo$_{1-1.5x}$Y$_x$PO$_4$@C electrode was collected by CHI660D within the voltage range 3.0~5.3 V, with 0.05 mV s$^{-1}$. EIS curve was conducted on Solartron SI 1260 and SI 1287 with a frequency range from 0.1 MHz to 10 MHz.

3. Results and Discussion

3.1. Composition and Morphology of LiCo$_{1-1.5x}$Y$_x$PO$_4$@C Cathode Material

Figure 1 shows the XRD patterns of the four cathode materials. All four cathode materials were well indexed to the olivine structure (JCPDS: 89-6192) with an orthorhombic Pnma space group, indicating the carbon layer and Y substitution did not change the main crystal structure of LiCoPO$_4$. An obvious impurity peak YPO$_4$ appeared when the Y substitution amount was 0.03. This means that at the substitution amount of 0.03, it was difficult for Y to be completely incorporated into the LiCoPO$_4$ lattice, as the Y atomic radius was larger than that of Co. Figure 2 presents the Rietveld refinement results of the four cathode materials. The crystal structural parameter is listed in Table 1, showing the details of the structural differences. Significantly, with the increase in the Y doping amount, the a and b parameters increased obviously, leading to an increase in unit cell volume (283.84, 284.09, 284.30 and 284.42 for x = 0, 0.01, 0.02 and 0.03). This change indicates that Y (III) replaced Co (II) and was incorporated into the LiCoPO$_4$ lattice. Since the ionic radius of Y (III) (90 pm) was larger than that of Co (II) (74.5 pm), it led to an increase in the unit cell volume. The increase in the unit cell volume could facilitate the migration of Li-ions, thus improving the electrochemical performance of LiCo$_{1-1.5x}$Y$_x$PO$_4$@C cathodes. According to the XRD and Rietveld refinement results, Y was successfully incorporated into the LiCoPO$_4$ lattice without altering the olivine structure, although it caused an increase in the unit cell volume as Y doping amount increased.
the XRD and Rietveld refinement results, Y was successfully incorporated into the LiCoPO₄ lattice without altering the olivine structure, although it caused an increase in the unit cell volume as Y doping amount increased.

Figure 1. XRD patterns of the four cathode materials.

![XRD patterns](image)

Figure 2. Refinement results of the four cathode materials.

Table 1. Crystal structural parameters of the four cathode materials.

| Samples                        | a  (Å)  | b  (Å)  | c  (Å)  | V  (Å³) | Rwp | Rp  | GOF |
|-------------------------------|---------|---------|---------|---------|-----|-----|-----|
| LiCoPO₄ (89-6192)             | 10.2021 | 5.9227  | 4.7003  | 284.01  |     |     |     |
| LiCoPO₄@C                     | 10.2007 | 5.9220  | 4.6987  | 283.84  | 3.85 | 2.71 | 1.73 |
| LiCo₀.⁹₈₅Y₀.₀₁PO₄@C           | 10.2027 | 5.9245  | 4.6999  | 284.09  | 3.96 | 2.74 | 1.78 |
| LiCo₀.⁹₇Y₀.₀₂PO₄@C            | 10.2058 | 5.9275  | 4.6995  | 284.30  | 4.02 | 2.82 | 1.84 |
| LiCo₀.₉₅₅Y₀.₀₃PO₄@C           | 10.2077 | 5.9288  | 4.6996  | 284.42  | 4.07 | 2.92 | 1.89 |

SEM images of the four cathode materials are displayed in Figure 3a–d. All the four cathode materials had similar morphology, which was composed of agglomerated nanoparticles of about 200 nm. This suggests that the basic morphology of the four cathode materials was unaffected by Y substitution. EDS of LiCo₀.⁹₇Y₀.₀₂PO₄/C cathode material
was performed to determine the Y distribution, and the result is shown in Figure 3e. It is evident that there was no region poorer or richer in Y, and the Y element was homogeneously dispersed in the LiCo$_{0.99}$Y$_{0.02}$PO$_4$/C cathode material. The same results were observed for the O, Co, and P elements. TEM images of the four cathode materials are presented in Figure 4. All the four samples had only one diffraction fringe and had a uniform 5 nm carbon layer on the LiCoPO$_4$ particle surface. From our previous studies [14,25,39], we found that a uniform carbon coating layer firstly refined particle size and improved material conductivity; secondly prevented direct contact between electrolyte and LiCoPO$_4$ particle, and inhibited the continuous oxidative decomposition of electrolyte under 5 V high voltage caused by Co$^{2+}$/Co$^{3+}$, and thirdly, stabilized the interface between the cathode and electrolyte and suppressed the continuous generation of CEI on the LiCoPO$_4$ particle surface, thus improving the electrochemical performance of the LiCoPO$_4$ material. SEM and TEM images suggest that the basic morphology of the four cathode materials was unaffected by Y substitution; that is, it consisted of clustered nanoparticles with a 5 nm uniform carbon film on the Li LiCo$_{1-1.5x}$Y$_x$PO$_4$@C particle surface.

![Figure 3. (a-d) SEM images of the four cathode materials and (e) EDS images of LiCo$_{0.97}$Y$_{0.02}$PO$_4$/C cathode materials.](image)

![Figure 4. TEM images of the four cathode materials.](image)
Figure 5 exhibits the FTIR spectra of the four cathode materials. It is obvious that with an increase in the Y substitution, the symmetric stretching at 987 cm\(^{-1}\) position shifted to 979 cm\(^{-1}\) position when x = 0.02, which indicated the decrease in the Li/Co antisite defect [8,15,16]. However, when the Y substitution amount increased to 0.03, the symmetric stretching position shifted to 984 cm\(^{-1}\) position, meaning the Li/Co antisite defect increase decreased the electrochemical performance of the LiCo\(_{0.955}\)Y\(_{0.03}\)PO\(_4\) material.

![FTIR spectra](image)

Figure 5. FTIR spectra of the four cathode materials.

The XPS spectrum of LiCoPO\(_4\)/C and LiCo\(_{0.97}\)Y\(_{0.02}\)PO\(_4\)/C cathode material is shown Figure 6. The spectra of the two cathode materials were similar, except for the LiCo\(_{0.97}\)Y\(_{0.02}\)PO\(_4\)/C sample with the Y characteristic peak. Figure 6b shows the Co2p spectra of the LiCoPO\(_4\)/C and LiCo\(_{0.97}\)Y\(_{0.02}\)PO\(_4\)/C cathode material, which were consistent with the reported binding energy of the Co [39,46]. The Y3d spectra for LiCo\(_{0.97}\)Y\(_{0.02}\)PO\(_4\)/C cathode material is displayed in Figure 6c. The Y3d spectra suggests that Y is present in the LiCo\(_{0.97}\)Y\(_{0.02}\)PO\(_4\)/C cathode material and that the oxidation state of Y is +3.

![XPS spectrum](image)

Figure 6. XPS spectrum of LiCoPO\(_4\)/C and LiCo\(_{0.97}\)Y\(_{0.02}\)PO\(_4\)/C cathode material.

3.2. Electrochemical Performances of LiCo\(_{1-1.5x}\)Y\(_{x}\)PO\(_4\)/C Electrodes

Electrochemical performances of the four electrodes were tested in half-cell. Figure 7 shows the first cycle CV curves of the four electrodes. The CV curves of the four electrodes had two similarities: first, all had an oxidation peak around 4.3 V that can be ascribed to the electrolyte oxidation reaction; then, all had two oxidation peaks in the range of 4.8 V~4.9 V and one reduction peak around 4.7 V (the reduction potential of LiCoPO\(_4\) was closer and the two overlapped as a larger reduction peak at the CV test sweep rate of 0.05 mV s\(^{-1}\), which is consistent with the reports in the reference) that corresponded to the
two steps of Li-ion extraction/intercalation [25,39]. The differences between the oxidation reaction potential, the reduction reaction potential, and the polarization potential of the CV curves are listed in Table 2. With increasing Y substitution amounts, the oxidation reaction potential decreased to 4.796 V and 4.892 V and the reduction potential increased to 4.71 V (x = 0.02); meanwhile, the polarization potential reduced to 0.182 V (x = 0.02). The changes in potential suggest that it is easier for Li-ion to migrate with Y substitution. These CV results reveal that the polarization of the four electrodes was reduced and that the Li-ion conductivity was enhanced with Y substitution. The improvement in Li-ion conductivity can be ascribed to the enlargement in unit cell volume of LiCo$_{1-1.5x}$Y$_x$PO$_4$/C cathode material, the decrement of the Li/Co antisite defect, and the increment of the Co-site vacancy with Y substitution, which offers additional channels for Li-ion migration.

![CV curves of the four electrodes.](image)

**Figure 7.** CV curves of the four electrodes.

**Table 2.** The differences of the CV curves of the four electrodes.

| Samples                  | Oxidation Potential (V) | Reduction Potential (V) | Polarization Potential (V) |
|--------------------------|-------------------------|-------------------------|-----------------------------|
| LiCoPO$_4$/C             | 4.829                   | 4.924                   | 4.695                       | 0.229                       |
| LiCo$_{0.985}$Y$_{0.01}$PO$_4$/C | 4.82                   | 4.903                   | 4.704                       | 0.199                       |
| LiCo$_{0.97}$Y$_{0.02}$PO$_4$/C | 4.796                  | 4.892                   | 4.71                        | 0.182                       |
| LiCo$_{0.955}$Y$_{0.03}$PO$_4$/C | 4.827                  | 4.908                   | 4.712                       | 0.196                       |

Figure 8 presents the first, second, third, tenth, twentieth, fortieth and hundredth cycle charge/discharge profiles of the four electrodes at 0.1C. The four electrodes displayed two oxidation plateaus at approximately 4.8 to 4.9 V, and two reduction plateaus at approximately 4.6 to 4.8 V, which agrees with the CV results. Noticeably, all four electrodes had one side reaction at about 4.3 V and a high overcharge capacity, which is ascribed to the electrolyte oxidation reaction. The overcharge capacity during the first charge is a major factor in the low initial coulombic efficiency [15,39]. However, at the second charge process, the overcharge capacity was reduced, which means the electrolyte oxidation reaction was restrained. This phenomenon confirms carbon film plays an important role in inhibiting the continuous oxidative decomposition of electrolyte under 5 V high voltage, and in stabilizing the interface between the cathode and electrolyte.

Figure 9a displays cycling stability and rate performances of the four electrodes. Figure 9a presents the cycling stability performance of the four electrodes at 0.1 C. The first discharge capacities of the four electrodes are 142.6, 144.5, 148, and 145.8 mAh g$^{-1}$, respectively. After 80 cycles, the discharge capacities are 76.8, 106.6, 111, and 97.6 mAh g$^{-1}$, with capacity retention of 53.8%, 73.7%, 75%, and 66.9%, respectively. The rapid capacity fading can be ascribed to the continuous generation of CEI on the LiCoPO$_4$ particle surface
and the Li/Co antisite defects during the cycling process [43,47]. Figure 9b presents the rate performances of the four electrodes. As expected, the LiCo$_{0.97}$Y$_{0.02}$PO$_4$/C electrode displayed the best performance, with the discharge capacities of 145 mAh g$^{-1}$ (0.1 C), 130 mAh g$^{-1}$ (0.2 C), 113 mAh g$^{-1}$ (0.5 C), and 96 mAh g$^{-1}$ (1 C); the discharge capacity returned to 130 mAh g$^{-1}$ when the discharge rate returned to 0.1 C, exhibiting good electrochemical performance stability. In comparison, the corresponding discharge capacity of the LiCoPO$_4$/C electrode was about 125 mAh g$^{-1}$ (0.1 C), 110 mAh g$^{-1}$ (0.2 C), 82 mAh g$^{-1}$ (0.5 C), and 66 mAh g$^{-1}$ (1 C). The improvement of the cycling stability and rate performances for LiCo$_{0.97}$Y$_{0.02}$PO$_4$/C electrode was ascribed to the decrease in the Li/Co antisite defect and the increase in ionic conductivity due to Y doping. Similar improvements in cycling stability and rate performance were reported for Cr-doped LiCoPO$_4$ [39,48] and V-doped LiCoPO$_4$ [16,37], which the authors, due to the facilitation of ion migration, caused by Cr or V substitution. Table 3 shows the cycling performance of the LiCo$_{0.97}$Y$_{0.02}$PO$_4$/C electrode in comparison with others reported.

### Table 3

| Method | Capacity (mAh g$^{-1}$) | Rate (C) | Capacity Retention (%) | Cycles |
|--------|------------------------|----------|-----------------------|--------|
| Carbon coating | 124 | 0.1 | 53.8 | 80 |
| LiCoPO$_4$/C | 120 | 0.1 | 73.7 | 80 |
| lithium metal coated | 16 | 0.1 | 75 | 80 |
| LiCo$_{0.97}$Y$_{0.02}$PO$_4$/C | 148 | 0.1 | 80 | 80 |

Figure 8. The charge/discharge profiles of the four electrodes.

Figure 9. The cycling stability (a) and rate (b) performances of the four electrodes.
Figure 10 shows the EIS spectra of the four electrodes and the corresponding equivalent circuits. The simulation results are listed in Table 4. It is clear that all four cathodes had similar ohmic resistance Re (1.6 Ω, 1.33 Ω, 1.41 Ω, and 1.53 Ω, respectively) due to their similar basic forms. However, the transfer resistance Rct (59.15 Ω, 25.41 Ω, 23.66 Ω and 21.03 Ω, respectively) decreased significantly, which implies that the electronic conductivity of the LiCo\(_{1-1.5x}\)Y\(_x\)PO\(_4\)@C cathode material was improved with the Y doping. The \(D_{Li^+}\) results are presented in Table 4. By increasing the Y doping amounts, the \(D_{Li^+}\) was improved, and the LiCo\(_{0.955}Y_{0.045}\)PO\(_4\)@C electrode showed the best values of \(6.16 \times 10^{-14}\) cm\(^2\) s\(^{-1}\), whereas the LiCoPO\(_4\)@C electrode only attained \(7.11 \times 10^{-15}\) cm\(^2\) s\(^{-1}\). The EIS results reveal that the intrinsic performance of ionic and electronic conductivities for LiCo\(_{1-1.5x}\)Y\(_x\)PO\(_4\)@C material was improved by Y substitution. This improvement can be ascribed to the enlargement in unit cell volume and the increment of the Co-site vacancy caused by aliovalent Y substitution that provided a convenient pathway for Li ion migration.

![EIS spectra](image)

**Figure 10.** The EIS spectra of (a) the four electrodes and (b) the linear relationship between \(Z'\) and \(\omega^{-1/2}\) in four electrodes.

**Table 3.** Cycling performance of the LiCo\(_{0.97}\)Y\(_{0.02}\)PO\(_4\)@C electrode in comparison with others reported.

| Samples                  | Rate | Initial Discharge Capacity (mAh g\(^{-1}\)) | Capacity Retention (%) | Cycles | Method                        |
|-------------------------|------|---------------------------------------------|------------------------|--------|-------------------------------|
| Our work                | 0.1C | 148                                         | 75                     | 80     | Y-Substituted and carbon coating |
| Ref. [33]               | 0.1C | 135                                         | 52                     | 30     | Carbon coating                |
| Ref. [49]               | 0.1C | 147                                         | 69                     | 40     | Carbon coating                |
| Ref. [31]               | 0.1C | 120                                         | 75                     | 20     | Carbon coating                |
| Ref. [32]               | 0.1C | 124                                         | 56                     | 100    | Carbon coating                |
| Ref. [16]               | 0.1C | 97                                          | 85                     | 20     | V-Substituted                 |
| Ref. [37]               | 0.1C | 145                                         | 52                     | 20     | V-Substituted                 |
| Ref. [44]               | 0.1C | 153                                         | 21                     | 30     | Y-Substituted                 |
| Ref. [15]               | 0.1C | 124                                         | 80                     | 20     | Fe-Substituted                |
| Ref. [36]               | 0.1C | 88                                          | 22                     | 20     | Mg-Substituted                |

**Table 4.** Impedance parameters of the four electrodes.

| Samples                  | Re (Ω) | Rct (Ω) | σ   | \(D_{Li^+}\) (cm\(^2\) s\(^{-1}\)) |
|-------------------------|--------|---------|-----|-------------------------------|
| LiCoPO\(_4\)@C          | 1.6    | 59.15   | 391 | \(7.11 \times 10^{-16}\)     |
| LiCo\(_{0.985}Y_{0.01}\)PO\(_4\)@C | 1.33   | 25.41   | 103 | \(1.02 \times 10^{-14}\)     |
| LiCo\(_{0.97}Y_{0.02}\)PO\(_4\)@C | 1.41   | 23.66   | 60  | \(3.02 \times 10^{-14}\)     |
| LiCo\(_{0.955}Y_{0.045}\)PO\(_4\)@C | 1.53   | 21.03   | 42  | \(6.16 \times 10^{-14}\)     |
4. Conclusions

In this work, the LiCo_{1-1.5x}Y_{x}PO_{4}@C (x = 0, 0.01, 0.02 and 0.03) cathode material was synthesized in one step. The uniform carbon layer stabilized the interface between the cathode and electrolyte, inhibiting the continuous side reaction on the LiCoPO_{4} particle surface; meanwhile, the Y substitution decreased the antisite defect, increasing the ionic and electronic conductivities of LiCo_{1-1.5x}Y_{x}PO_{4}@C sample. Thus, the LiCo_{0.97}Y_{0.02}PO_{4}@C cathode exhibited the best electrochemical performance, for instance, delivering an initial discharge capacity of 148 mAh g^{-1}, with a capacity retention of 75% after 80 cycles at 0.1 C, and delivered a capacity of 96 mAh g^{-1} at 1 C. The low Li/Co antisite defect, the enhancement of electronic and Li-ion conductivity caused by Y substitution, and the uniform carbon layer, worked together to improve the performance of LiCo_{1-1.5x}Y_{x}PO_{4}@C cathode.

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