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Graphene composite 3,4,9,10-perylenetetracarboxylic sodium salts with a honeycomb structure as a high performance anode material for lithium ion batteries

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In order to address the issues of high solubility in electrolytes, poor conductivity and low active site utilization of organic carbonyl electrode materials, in this work, the 3,4,9,10-perylenetetracarboxylic sodium salt (PTCDA-Na) and its graphene composite PTCDA-Na-G are prepared by the hydrolysis of 3,4,9,10-perylenetetracarboxylic dianhydride and the strategy of antisolvent precipitation. The obtained PTCDA-Na active substance has a porous honeycomb structure, showing a large specific surface area. Moreover, after recombination with graphene, the dispersion and specific surface area of PTCDA-Na are further enhanced, and more active sites are exposed and conductivity is improved. As a result, the PTCDA-Na-G composite electrode materials exhibit superior electrochemical energy storage behaviors. The initial charge capacity of the PTCDA-Na-G electrode is 890.5 mA h g⁻¹, and after 200 cycles, the capacity can still remain at 840.0 mA h g⁻¹ with a high retention rate of 94.3%, which is much larger than those of the PTCDA-Na electrode. In addition, at different current densities, the PTCDA-Na-G electrode also presents higher capacities and better cycle stability than the PTCDA-Na electrode. Compared with PTCDA-Na with a porous honeycomb structure and previously reported sodium carboxylic acid salts with a large size bulk structure, the PTCDA-Na-G composite material prepared in this work shows superior electrochemical energy storage properties due to its large specific surface area, high dispersion, more exposed active sites and large electrical conductivity, which would provide new ideas for the development of high performance organic electrode materials for lithium-ion batteries.

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1 Introduction

With the increasing demand of high power equipment for energy storage devices, lithium ion batteries with large capacity, long cycle and high rate performance are attracting extensive interest from researchers.¹⁻³ In lithium ion batteries, the negative electrode material is one of the key materials that affect the performance of the battery.⁴⁻⁶ At present, the traditional negative electrode material which has been commercialized is the graphite carbon negative electrode; however, there are still many problems such as a limited theoretical capacity (372 mA h g⁻¹), unstable layered structure, poor cycling stability and so on, which limit its application in high power energy storage batteries.⁷⁻⁹ Therefore, it is urgent to develop novel high performance anode materials to meet the increasing market demand. Subsequently, many other negative electrode

materials have been studied, such as hard carbon materials, transition metal oxides, silicon negative electrode materials, and so on.¹⁰⁻¹⁵ Some of the designed materials show superior properties and provide new ideas for the subsequent development of high performance negative electrode systems, but they have their own advantages and disadvantages.

Organic compounds¹⁶⁻²¹ as a new class of electrode materials are of great interest to researchers because of their advantages of high theoretical specific capacity, abundant raw materials, environmental friendliness, strong structural designability and so on. Among them, conjugated organic molecules containing unsaturated double bond structures have the advantages of fast redox reaction kinetics, high capacity and structural diversity due to their conjugated systems and multiple unsaturated groups, which is of broader concern in their roles as emerging electrochemical energy-storage materials.²²⁻²⁹ However, organic electrode materials with small molecules still have the problems of high solubility in electrolytes, poor conductivity and limited exposed active sites, which greatly limit the electrochemical performance of such materials.³⁰⁻³⁴ To address these issues, researchers have carried out many modification

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studies,^{35–41} such as grafting active groups onto macrocyclic conjugated structures, polymerization, doping with inorganic systems, and so on. These methods have greatly improved the performance of organic electrode and provided many new ideas for subsequent modification. However, many modification measures can lead to the decrease of the proportion and specific capacity of the active components, or cannot fundamentally solve the problem of poor conductivity.^{41–43} Therefore, it is still necessary to design new modification measures to solve the above problems of organic electrode materials at the same time.

In this work, 3,4,9,10-perylenetetracarboxylic dianhydride (PTCDA) was added to NaOH aqueous solution to hydrolyze, and ethanol anti-solvent is added to obtain a yellowish-brown precipitate of perylenetetracarboxylic sodium salts (PTCDA-Na). As a result, lower solubility in the electrolyte is observed, and at the same time, the composite material of perylenetetracarboxylic sodium salts with graphene (PTCDA-Na-G) is used as an anode material for lithium ion batteries, showing improved capacity, cycling stability and rate capability. The strategy of sodium salinization of an organic compound and its graphene recombination technology will provide new ideas for the design of novel organic electrode materials with high performance for lithium-ion batteries.

2 Experimental

2.1 Materials

The raw materials used in this work, such as 3,4,9,10-perylenetetracarboxylic dianhydride (PTCDA), NaOH, graphene, Super P, anhydrous ethanol, poly(vinylidene fluoride) (PVDF), *N*-methylpyrrolidone (NMP), *etc.*, are purchased from Chemical Reagent Co., Ltd, and are used directly without any purification.

2.2 Synthesis

2.2.1 Perylenetetracarboxylic sodium salts (PTCDA-Na). 9.8 g of 3,4,9,10-perylenetetracarboxylic dianhydride (PTCDA) was added to the aqueous solution of NaOH (8 g), and then the mixed solution was transferred to a round-bottom flask and stirred in a water bath at 80 °C until PTCDA was completely hydrolyzed to a clear, transparent brown solution. After cooling the hydrolyzed solution to room temperature, adding it to a large amount of anhydrous ethanol, a turbid suspension was gradually produced, and then the suspension was centrifuged, and the precipitate was washed several times with anhydrous ethanol. The collected precipitates were dried in an 80 °C vacuum oven to obtain an orange precipitated powder of perylenetetracarboxylic sodium salts (PTCDA-Na).

2.2.2 Graphene/PTCDA-Na composite material (PTCDA-Na-G). Few-layer graphene (0.05 g) prepared by a physical method was added into the aqueous solution of PTCDA-Na (0.258 g) until the solution became dark green. Slowly adding ethanol into the solution, the dissolved PTCDA-Na sodium salt gradually precipitates, and the resulting PTCDA-Na is attached to the surface of the thin layer of graphene. After dripping, the solution was filtered, washed several times with anhydrous ethanol, a gray-green precipitate was collected, and then dried

in a vacuum oven to obtain the graphene/PTCDA-Na composite material (PTCDA-Na-G).

2.3 Characterization and electrochemical tests

UV-Vis spectroscopy and dissolution experiments were conducted on a Hitachi U-3010 spectrophotometer. The infrared spectrum was obtained with a Fourier transform infrared (FTIR) spectrophotometer Bruker IFS66/S. X-ray diffraction spectroscopy (XRD) was performed with an X-ray diffractometer under Cu K α radiation. Scanning electron microscopy (SEM) images were obtained on a ZEISS Crossbeam 340 scanning electron microscopy analyzer. A transmission electron microscope (FEI Tecnai G2F20) was used to capture the TEM images. X-ray photoelectron spectroscopy (XPS) was carried out on a Thermo Fisher K-Alpha with an Al K α X-ray source.

Before the electrochemical performance test, all cells were pretreated by discharging to 0.001 V from open-circuit voltage (OCV) at a 100 mA g⁻¹ current density and eventually charged to 3.0 V at the same current density. The cyclic voltammetry (CV) curves were recorded using an Ivium-Stat multichannel electrochemical workstation (Ivium-n-Stat.Xri, the Netherlands), with the voltage range of 3 to 0.001 V and a scan rate of 0.1 mV s⁻¹. Electrochemical impedance spectroscopy (EIS) was also performed using an Ivium-Stat multichannel electrochemical workstation (Ivium-n-Stat.Xri, the Netherlands) with a frequency of 10⁻²–10⁵ Hz and a voltage amplitude of 5 mV. The charge–discharge test including cycle and rate performance analysis was performed on a Neware battery test system (BTS-5V10 mA) in a voltage range of 3 to 0.01 V.

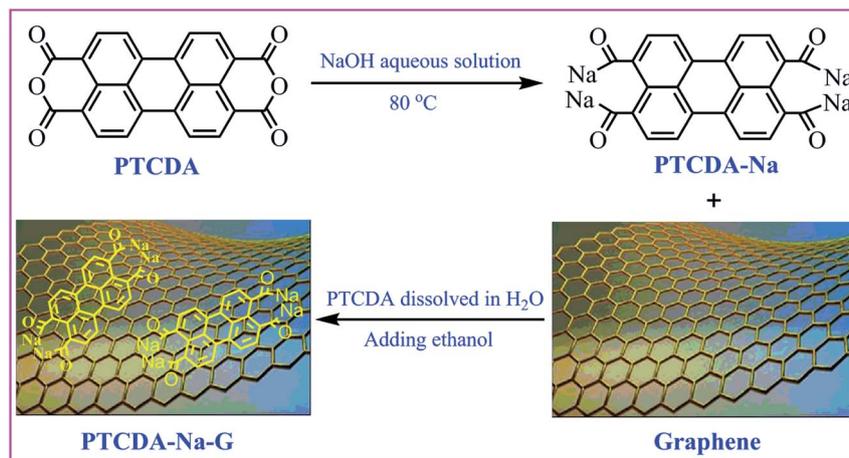
2.4 Preparation of electrode sheets and batteries

Batteries were assembled according to the following processes: PTCDA-Na powder or PTCDA-Na-G composite powder and Super P and poly(vinylidene fluoride) (PVDF) binder were mixed at a mass ratio of 6 : 3 : 1, and then 2 g of this mixture was added to 10 mL of *N*-methylpyrrolidone (NMP) and ball-milled until the slurry was uniform. The uniform slurry was coated on copper foil uniformly, and dried at 120 °C for an hour, and finally vacuum dried at 60 °C for 24 h. The film was cut into a circular piece of 1.2 cm diameter, and the actual loading amount of the active material was 0.81 mg and 1.04 mg for the PTCDA-Na and PTCDA-Na-G electrodes, respectively, with an areal mass loading of 0.72 and 0.92 mg cm⁻². Then PTCDA-Na/Li and PTCDA-Na-G/Li half batteries, and PTCDA-Na/NCM-811 and PTCDA-Na-G/NCM-811 full batteries are assembled with lithium metal sheets as counter electrodes, and 1 M LiPF₆ in a mixture of ethylene carbonate/diethyl carbonate/ethyl methyl carbonate (EC/DEC/EMC, 1 : 1 : 1 by volume) as the electrolyte (purchased from Capchem Technology (Shenzhen) Co., Ltd.).

3 Results and discussion

The PTCDA-Na and PTCDA-Na-G composites were prepared as shown in Scheme 1. A yellowish green transparent solution was obtained after PTCDA was fully hydrolyzed in NaOH aqueous solution, and the precipitate was produced after adding





Scheme 1 Preparation of PTCDA-Na and PTCDA-Na-G composite samples.

antisolvent ethanol to the solution, and then the yellow **PTCDA-Na** powder sample was obtained. Moreover, a certain amount of graphene was added to the saturated aqueous solution dissolved **PTCDA-Na**, and after vigorous stirring and ultrasonic dispersion, the antisolvent ethanol was added to the mixed solution. At this time, **PTCDA-Na** began to precipitate and adhere to the flake graphene surface, and the **PTCDA-Na-G** composite of graphene was obtained.

For **PTCDA** compounds, there is a common problem with most small molecular organic compounds, that is, they will dissolve gradually in the electrolyte solution, which will cause the loss of active substances and eventually lead to the loss of capacity. After sodium salinization, the solubility of the **PTCDA-Na** sample in the electrolyte will be effectively inhibited. In order to investigate and verify this effect, the solubility of **PTCDA** and **PTCDA-Na** samples was compared. They were added to the electrolyte solvent (DMC : EMC : EC = 1 : 1 : 1) and expected to be prepared into a solution of 1×10^{-5} M, and their UV-Vis absorption spectra were tested, as shown in Fig. 1. As can be seen, even the newly prepared **PTCDA** solution shows

obvious characteristic absorption peaks at 450–550 nm. However, for **PTCDA-Na** samples, even after 2 days of placement, the characteristic absorption peak was not observed, which fully demonstrates that sodium salinization effectively inhibits the dissolution of organic electrode materials in the electrolyte, which will be very beneficial to the improvement of **PTCDA-Na** electrode properties.

The morphology of the active material has a crucial effect on the performance of the electrode. To this end, the microstructure of the **PTCDA-Na** and **PTCDA-Na-G** samples was investigated, as shown in Fig. 2. Obviously, the **PTCDA-Na** sample is composed of flake particles with a thickness of about 50 nm, and the flake particles accumulate together to form a honeycomb porous microstructure (Fig. 2(a and b)), which would have a large specific surface area, and more active sites will be exposed and more capacity contributions will be produced. After compositing with graphene, for the **PTCDA-Na-G** sample, the flake **PTCDA-Na** active material is coated with a thin layer of graphene, and the **PTCDA-Na** particles are attached to the surface of graphene (Fig. 2(c and d)), which can effectively

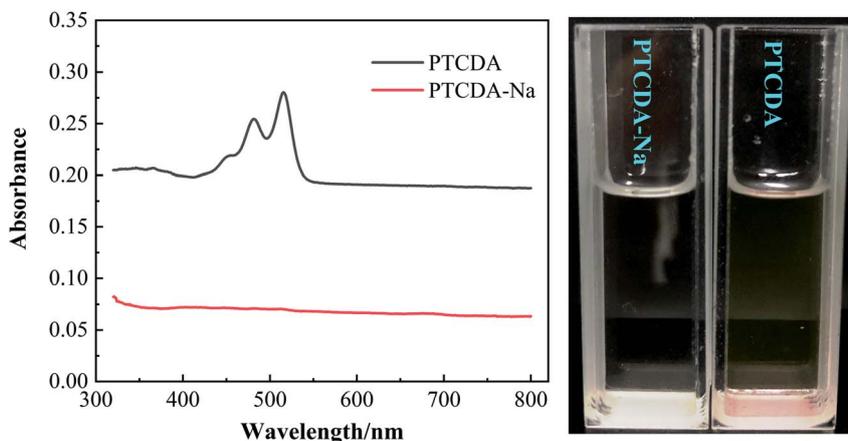


Fig. 1 UV-Vis spectra of PTCDA and PTCDA-Na composite samples (left) and their solution in electrolyte solvent (DMC : EMC : EC = 1 : 1 : 1) with an expected concentration of 1×10^{-5} M.



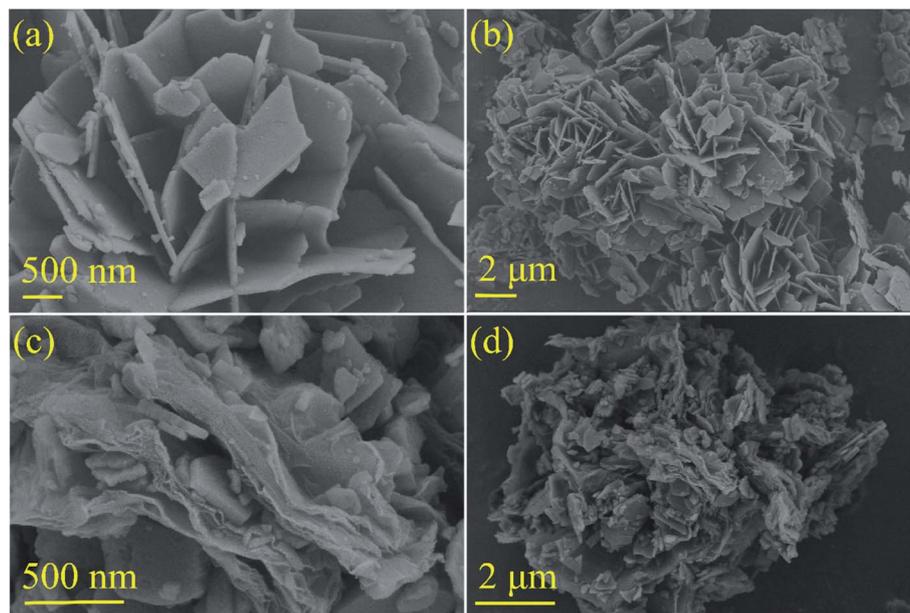


Fig. 2 SEM images of PTCDA-Na and PTCDA-Na-G composite samples (a and b) PTCDA-Na; (c and d) PTCDA-Na-G.

improve the dispersion degree and further enhance the exposure degree of the active point of the electrode material, and would be more conducive to improving the electrochemical performance.

The EDS spectra were also investigated to further characterize the elemental composition of electrode materials. For the **PTCDA-Na** sample, the C, O and Na elements were clearly observed (Fig. 3(a–d)), and their distribution is relatively uniform. Meanwhile after compositing **PTCDA-Na** with graphene, with the increase of the composition of the C element,

the blue color representing the C element in the total spectrum and the C element spectrum also deepens, while in the O and Na element spectra, the green color representing the O element and the purple color representing the Na element are slightly lighter (Fig. 3(e–h)), which fully confirms the recombination process of graphene.

A BET surface adsorption experiment was carried out, as shown in Fig. 4 and Table 1. For the **PTCDA-Na** sample, because of its porous and loose honeycomb structure (Fig. 3(a–d)), a large pore size of 53.17 nm, total pore volume of 0.5617 cm³

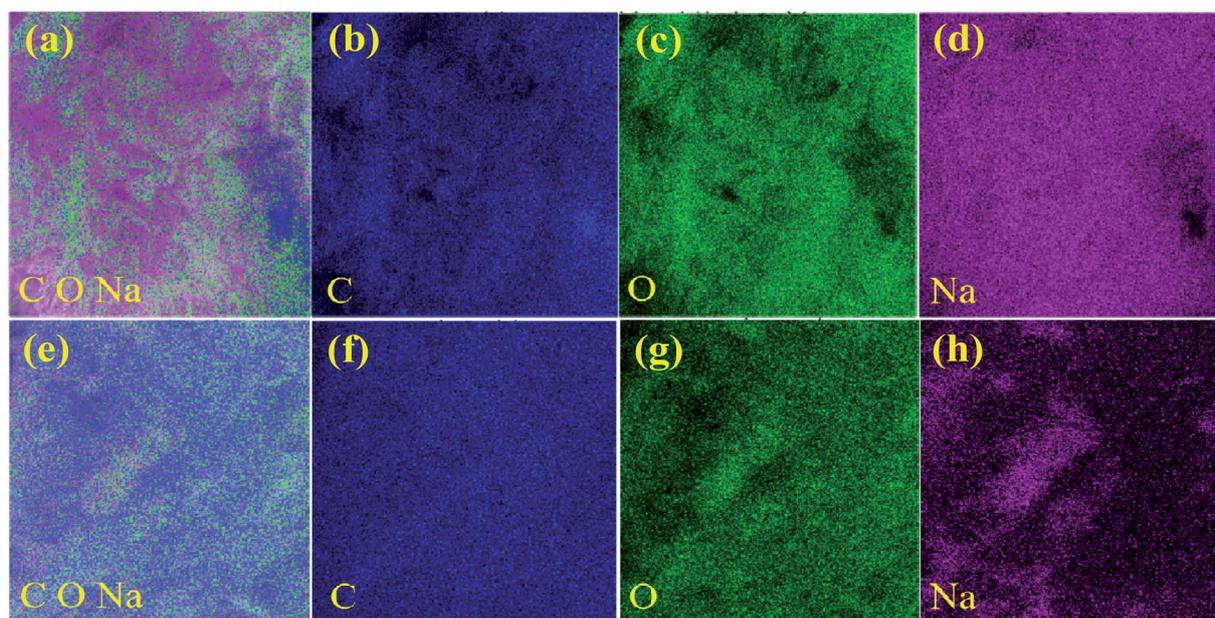


Fig. 3 EDS images of PTCDA-Na and PTCDA-Na-G composite samples (a–d) PTCDA-Na; (e–h) PTCDA-Na-G.



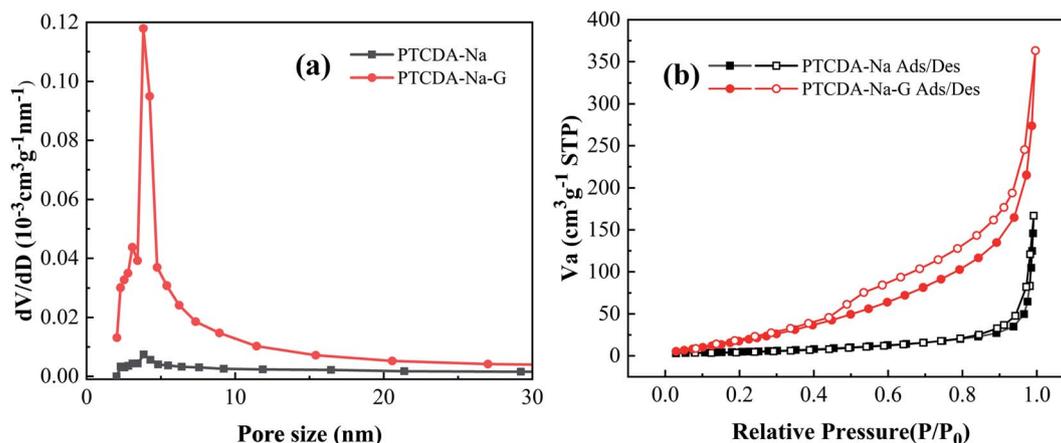


Fig. 4 (a) Pore size distribution curves of PTCDA-Na and PTCDA-Na-G samples; (b) adsorption curves of PTCDA-Na and PTCDA-Na-G samples.

Table 1 BET adsorption parameters of PTCDA-Na and PTCDA-Na-G samples

Samples	Specific surface area ($\text{m}^2 \text{g}^{-1}$)	Total pore volume ($\text{cm}^3 \text{g}^{-1}$)	Average pore diameter (nm)
PTCDA-Na	19.4	0.5617	53.17
PTCDA-Na-G	112.4	0.2583	19.99

g^{-1} and specific surface area of $19.4 \text{ m}^2 \text{g}^{-1}$ are observed. However, after graphene compositing with PTCDA-Na, the PTCDA-Na-G sample presents a larger specific surface area of $112.4 \text{ m}^2 \text{g}^{-1}$ with a lower total pore volume of $0.2583 \text{ cm}^3 \text{g}^{-1}$ and small size of 19.99 nm , respectively. After graphene recombination, the pore size of the PTCDA-Na-G sample is decreased, and the specific surface area of the sample is significantly enhanced, which clearly indicates that the recombination of graphene can further increase the dispersion of PTCDA-Na active substances and expose more active sites. This will further improve the electrochemical performance of the electrode.

The symmetric and antisymmetric peaks of $-\text{COO}-$ bonds in PTCDA-Na compounds can be clearly observed in the IR

spectrum (Fig. 5(a)). After graphene recombination, there is no change of new organic functional groups and bonds, and the IR spectrum of PTCDA-Na-G composites has almost no change. XRD diffraction patterns show that the PTCDA-Na sample has good crystallinity (Fig. 5(a)), which is mainly determined by its regular morphology and ordered lamellar structure as shown in Fig. 2(a and b). Due to the disorder of graphene itself, in the process of recombination with graphene, the PTCDA-Na compound is attached to the graphene surface after precipitation, which naturally reduces the disorder degree of the PTCDA-Na-G sample. Therefore, the XRD diffraction peaks in Fig. 5(b) also indicate that the crystallization degree of the sample is obviously decreased and the disorder degree is increased, which can also be clearly observed from Fig. 2(c and d). Moreover, for the PTCDA-Na-G sample, in its XRD diffraction pattern, it can be obviously observed that the wide drum-wrapped peaks around 26° belongs to the characteristic diffraction peak of graphene, which indicates the existence of graphene.

Changes in the morphology and structure before and after graphene recombination are also reflected in TEM images. As shown in Fig. 6(a and b), the lamellar PTCDA-Na particles piled together, and obvious lattice fringes were observed on the lamellar particle surface with a fringe spacing of 1.265 nm .

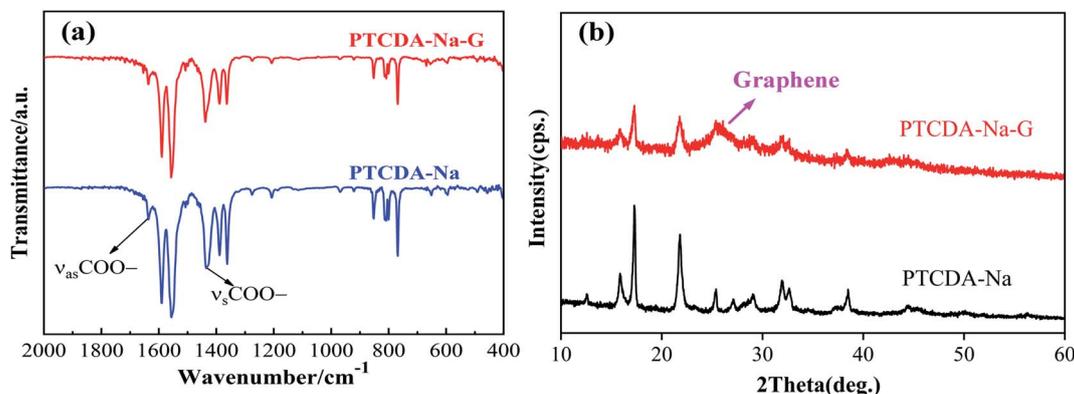


Fig. 5 (a) IR spectra and (b) XRD diffraction patterns of PTCDA-Na and PTCDA-Na-G composite samples.



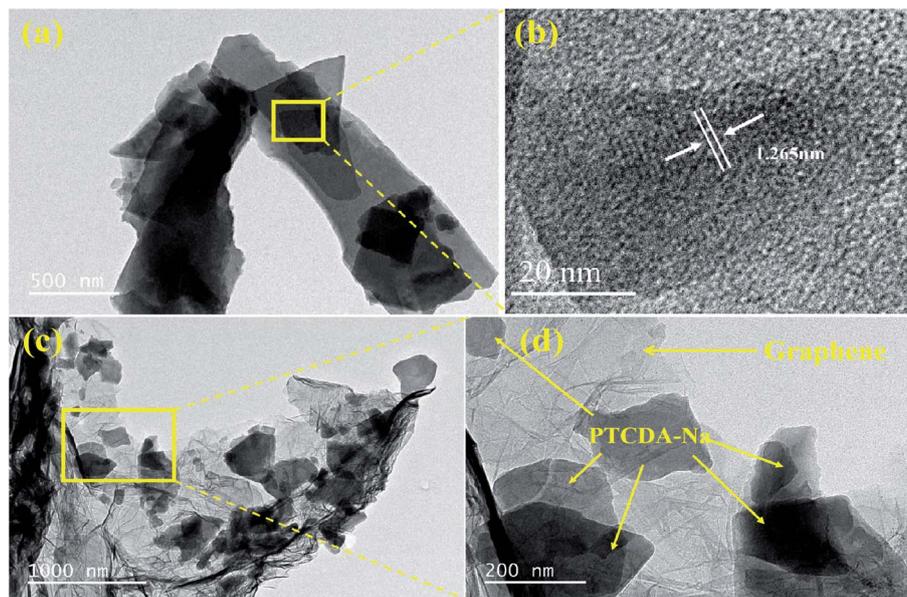


Fig. 6 TEM images of (a and b) PTCDA-Na and (c and d) PTCDA-Na-G composite samples.

Meanwhile in the **PTCDA-Na-G** sample, **PTCDA-Na** consists of mainly bulk particles, which are loosely bound on the surface of the graphene layer and no obvious lattice fringes are observed (Fig. 6(c and d)). The TEM image further indicates that **PTCDA-Na** samples have better order and crystallinity, while **PTCDA-Na-G** samples are more disordered.

PTCDA-Na and **PTCDA-Na-G** electrodes were prepared and assembled into a Li-ion battery, and cyclic voltammetry (CV) was carried out from 0.001 to 3.0 V with a scan rate of 0.1 mV s^{-1} . As shown in Fig. 7(a), **PTCDA-Na** shows three pairs of redox peaks at 2.2/1.9 V, 1.5/1.1 V and 0.2/0.01 V, which correspond to the lithium intercalation reaction of C=O and C=C functional groups in the **PTCDA** compound as well as interlayer spacing, respectively. Fig. 7(b) clearly shows that there is a weak ground peak at 1.9 V through differential analysis. The second CV curve has a good coincidence with the third, and their area decreases slightly compared with the first CV curve, which indicates that the **PTCDA-Na** electrode has smaller irreversible capacity and better cycling reversibility. After compositing with graphene, the **PTCDA-Na-G** sample mainly has a disordered amorphous structure, and the redox peak caused by the lithium intercalation process between layers becomes very weak; only two pairs of obvious redox peaks at 1.5/1.1 V and 0.2/0.01 V correspond to the lithium intercalation reaction of C=O, C=C functional groups as well as a disordered amorphous structure was observed for the **PTCDA-Na-G** electrode (Fig. 7(c)). Similarly, the second CV curve has a good coincidence degree with the third one, and compared with the first CV curve, the area decreases slightly, indicating that the **PTCDA-Na-G** electrode also has a small irreversible capacity loss and better cycling reversibility. From the TEM image in Fig. 6, it can be seen that the **PTCDA-Na** sample is mainly mixed with an ordered layered structure and a disordered unstructured structure, which determines that the oxidation reduction peak in its CV curve is derived from the

ordered layered structure and the disordered unstructured structure, respectively. As a result, as shown in Fig. 7(a), the **PTCDA-Na** electrode shows three pairs of redox peaks which correspond to the lithium intercalation reaction of C=O and C=C functional groups in the **PTCDA** compound, interlayer spacing and disordered amorphous structure, respectively. However, for **PTCDA-Na-G** samples, no orderly layered structures were found, mainly consisting of rambling amorphous structures. As a result, the CV curve mainly has two main groups of oxidation reduction peaks, and no redox peaks at 2.2/1.9 V resulting from ordered layered structures were found. Fig. 7(d) shows that the area of the CV curve for the **PTCDA-Na-G** electrode is slightly larger than that of the CV curve for the **PTCDA-Na** electrode, indicating that graphene recombination will improve electrode capacity to some extent. This is mainly because the *in situ* recombination of graphene will increase the dispersion of **PTCDA-Na** active particles, improve the conductivity of the electrode, and finally expose more active sites.

This conclusion can also be confirmed by the analysis of the EIS impedance spectra of the two electrodes, as shown in Fig. 8. It's obvious that after graphene recombination, the semicircle diameter in the high frequency region of the **PTCDA-Na-G** electrode decreases, and the slope of the straight line in the low frequency region increases obviously. After fitting, different impedance values are obtained, as listed in Table 2, where R_s reflects the transfer resistance of lithium ions in the electrolyte and the interface resistance caused by the SEI film double layer, R_{ct} is the charge-transfer resistance during the electrochemical reaction process, and W_1 represents the Warburg impedance concerning the diffusion of Li^+ into the bulk electrode. After 3 cycles (Fig. 8(a)), the R_s , R_{ct} and W_1 of the **PTCDA-Na-G** electrode are 2.4, 62.8 and 0.0002Ω , respectively, which are lower than those of the **PTCDA-Na** electrode (the R_s , R_{ct} and W_1 are 4.2, 71.9 and 23.8Ω , respectively). The results show that the impedance



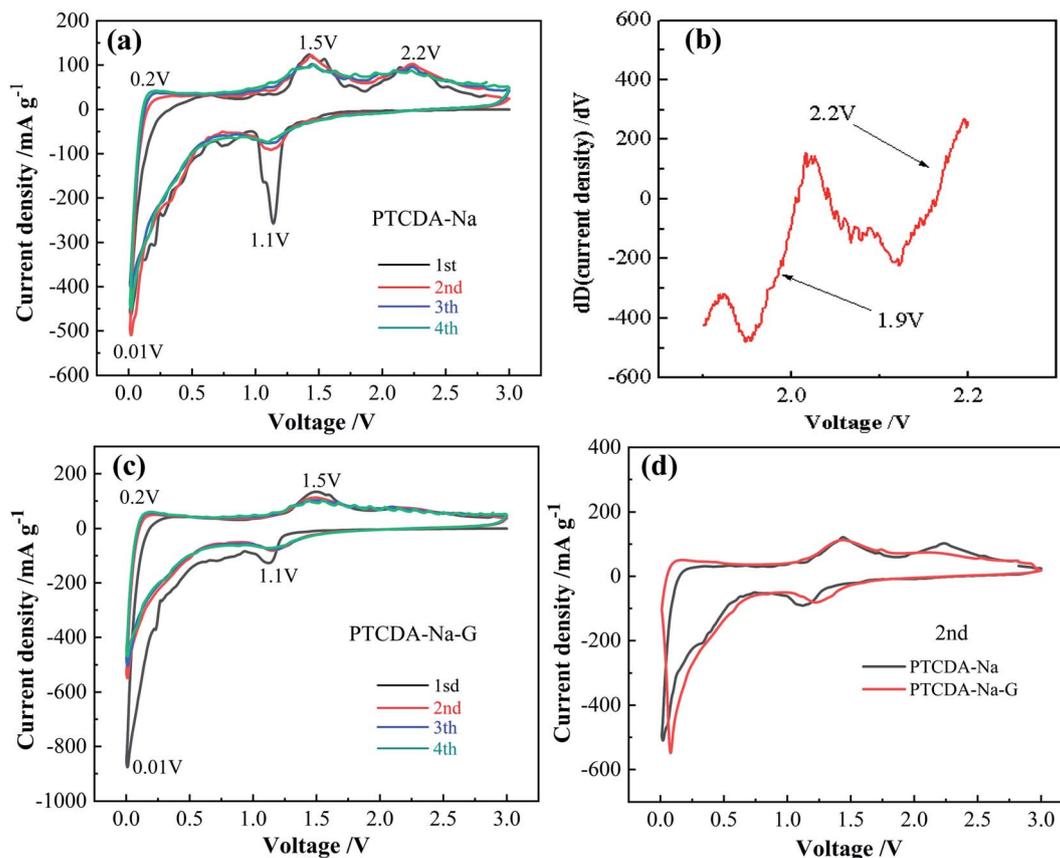


Fig. 7 (a) Cyclic voltammograms curves of the PTCDA-Na electrode; (b) differential representation of the 1st cyclic voltammogram curve of the PTCDA-Na electrode; (c) cyclic voltammograms curves of the PTCDA-Na-G electrode; (d) contrast of the 2nd cyclic voltammograms curves of PTCDA-Na and PTCDA-Na-G electrodes.

of the PTCDA-Na-G electrode decreases obviously after graphene recombination, especially the Warburg impedance (W_1) concerning the diffusion of Li^+ into the bulk electrode, which would be beneficial to the performance of the electrode. After 100 cycles (Fig. 8(b)), the R_{ct} of both the electrodes dropped significantly. Meanwhile, the W_{1-R} of PTCDA-Na-G is still much smaller than that of the PTCDA-Na electrode.

The charge/discharge behaviors of PTCDA-Na and PTCDA-Na-G electrodes were investigated as shown in Fig. 9. It is obvious that the PTCDA-Na-G electrode presents superior cycle stability to the PTCDA-Na electrode. The initial charge capacity of the PTCDA-Na-G electrode is $890.5 \text{ mA h g}^{-1}$; thereafter, the charging capacity decreased in the next 20 cycles. After 20 cycles, the charging capacity began to rise gradually, and the

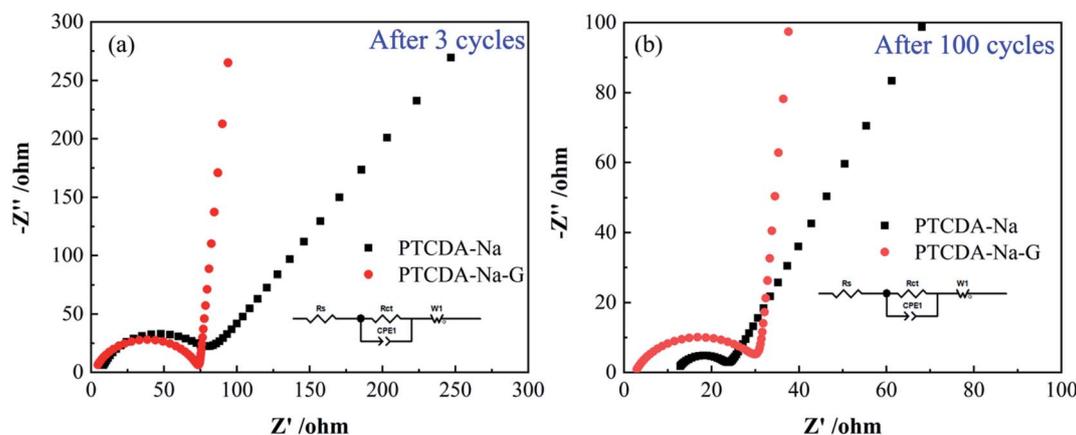


Fig. 8 (a) EIS impedance of PTCDA-Na and PTCDA-Na-G electrodes after 3 cycles; (b) EIS impedance of PTCDA-Na and PTCDA-Na-G electrodes after 100 cycles.



Table 2 EIS impedance fitting parameters of PTCDA-Na and PTCDA-Na-G electrodes

Electrode	R_s/Ω (error)	R_{ct}/Ω (error)	W_{1-R}/Ω (error)
PTCDA-Na (3 rd cycle)	4.21 (67.7%)	62.82 (21.2%)	23.8 (202.3%)
PTCDA-Na-G (3 rd cycle)	2.42 (208.0%)	71.98 (0.22%)	0.0002 ($7.71 \times 10^{-6}\%$)
PTCDA-Na (100 th cycle)	12.14 (1.52%)	11.51 (2.32%)	3.84 (7.28%)
PTCDA-Na-G (100 th cycle)	2.53 (2.02%)	27.16 (11.36%)	3.43 (254.59%)

charging capacity remained at $840.0 \text{ mA h g}^{-1}$ after 200 cycles with a capacity retention of 94.3%. The PTCDA-Na-G electrode has good stability and extremely high reversibility of electrochemical reactions. During the 200 cycles of the charge and discharge process, the coulombic efficiency of the electrode still remains above 98%. Meanwhile for the PTCDA-Na electrode, the initial charge capacity is $680.3 \text{ mA h g}^{-1}$, which is much lower than that of the PTCDA-Na-G electrode (Fig. 9(a)). After 200 cycles, the capacity gradually decreases to $401.8 \text{ mA h g}^{-1}$ with a capacity retention of 59.1%, which are also much lower than those of the PTCDA-Na-G electrode. After compositing with graphene, the rate capability is also obviously improved, as shown in Fig. 9(b). At different current densities, the PTCDA-Na-G electrode displays stable rate cycle performance, while for the PTCDA-Na electrode, the cycle capacities are very unstable, and

display a declining trend at different current densities. The results indicate that the rate capacities of the PTCDA-Na-G electrode are much higher and more stable than those of the PTCDA-Na electrode, which clearly reflect that graphene plays very important roles in the performance improvement of PTCDA-Na molecules.

In addition, an investigation was made on the electrochemical performance of the graphene/Li battery in order to clarify the reasons for the improvement of PTCDA-Na active materials after compositing with graphene. As shown in Fig. 9(c), the initial charge capacity of the graphene electrode is 235 mA h g^{-1} , which is much lower than that of the PTCDA-Na electrode (600 mA h g^{-1}) and PTCDA-Na-G electrode (800 mA h g^{-1}). Moreover, as the cycling progresses, the capacity of graphene decreases (Fig. 9(d)). The results show that

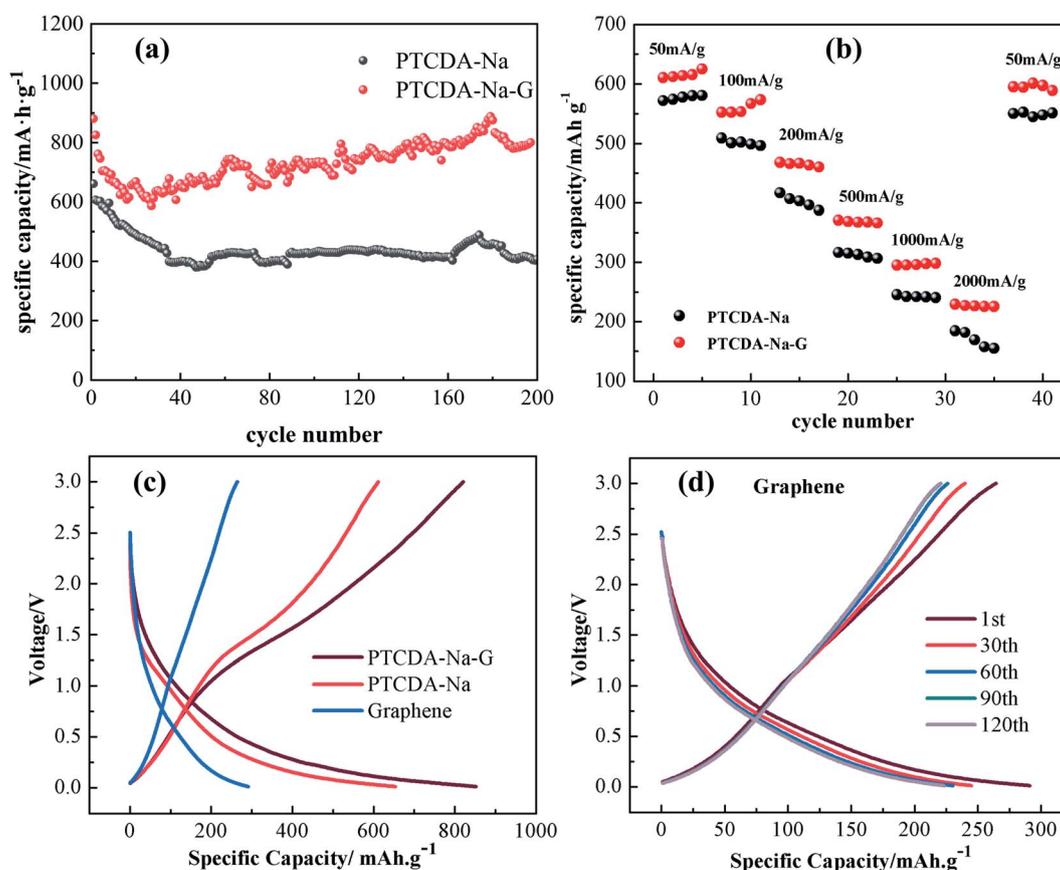


Fig. 9 (a) Cycle performance of PTCDA-Na and PTCDA-Na-G electrodes; (b) rate capacity of PTCDA-Na and PTCDA-Na-G electrodes; (c) initial charge/discharge curves of PTCDA-Na and PTCDA-Na-G and graphene electrodes; (d) charge/discharge curves of the graphene electrode in different cycles.



Table 3 Cycling performance data of TNTCDA-NiPc and previously reported similar electrode materials

Electrodes	Initial capacity (mA h g ⁻¹)	200 th capacity (mA h g ⁻¹)	Capacity retention	References
PTCDA-Na	680.3	401.8	59.1%	In this work
PTCDA-Na-G	890.5	840.0	94.3%	In this work
Sodium tartrate (ST)	182.3	295.3	162%	46
Sodium pyromellitate (SP)	182.4	246.1	135%	46
Sodium oxalate (SO)	187.9	354.2	188%	46
Sodium citrate (SC)	186.8	188.8	101%	46

graphene is not the main capacity contributor in **PTCDA-Na-G** composites, while the **PTCDA-Na** active material is the main capacity contributor. On the one hand, graphene can effectively improve the dispersion of **PTCDA-Na** active compounds and expose more active sites in **PTCDA-Na-G** composites. On the other hand, graphene with good conductivity can further enhance the conductivity of the composites, making the capacity of active compounds easier to show and effectively improving the rate cycling performance at a high current. On the one hand, graphene can effectively improve the dispersion of **PTCDA-Na** active compounds and expose more active sites in **PTCDA-Na-G** composites. On the other hand, graphene with good conductivity can further enhance the conductivity of the composites, making the capacity of active compounds easier to show and effectively improving the rate cycling performance at a high current. And that's why the **PTCDA-Na-G** electrode has better charge-discharge energy storage performance than the **PTCDA-Na** electrode. According to reported literature,^{44,45} annealing at a certain temperature can improve the performance of **PTCDA**, increase electrical conductivity, and reduce thermal decomposition.

In order to illustrate the advantage of **PTCDA-Na** electrodes, a systematic comparison between the performance of **PTCDA-Na** electrodes in this work and that of previously reported similar sodium carboxylate electrodes is provided as shown in Table 3. It can be seen that for large particles of sodium carboxylic acid active compounds,⁴⁶ their initial capacities are relatively low, but as the cycling goes on, their capacities gradually increase. This is due to the gradual pulverization of the active substances during the cycle process, which exposes more active sites. Nevertheless, the **PTCDA-Na** active compounds investigated in this work are composed of porous honeycomb-like sheet-like particles with a larger specific surface area and more exposed active sites than previously published bulk sodium carboxylic acid salt active materials, resulting in a larger initial capacity of the **PTCDA-Na** electrode. After graphene recombination, the dispersion and conductivity of **PTCDA-Na-G** active substances are further improved, and more active sites

are exposed, and larger initial capacity and better rate performance are displayed. As a result, **PTCDA-Na** with a honeycomb structure and its graphene composites prepared in this work exhibit superior electrochemical energy storage properties.

4 Conclusions

In this work, graphene composite 3,4,9,10-perylenetetracarboxylic sodium salts (**PTCDA-Na-G**) were prepared; after the **PTCDA-Na** active substances with a porous honeycomb structure are encapsulated by graphene, their dispersion and specific surface area are further effectively improved, so that more active sites are exposed. In addition, graphene with a thin layer structure further improves the conductivity of composite electrode materials. As a result, the electrochemical energy storage performance (such as initial capacity, cycle stability and rate capability) of the **PTCDA-Na-G** composite electrode is greatly improved. Furthermore, compared with previously reported sodium carboxylic acid salts (ST, SP, SO and SC) with a large size bulk structure, the **PTCDA-Na** with a porous honeycomb structure prepared in this work and especially its graphene complexes (**PTCDA-Na-G**) show superior electrochemical energy storage properties due to their large specific surface area, high dispersion, more exposed active sites and large electrical conductivity. Furthermore, compared with previously reported sodium carboxylic acid salt compounds with a large size bulk structure, the **PTCDA-Na** with a porous honeycomb structure prepared in this work and its graphene complexes show superior electrochemical energy storage properties due to their large specific surface area, high dispersion, more exposed active sites and large electrical conductivity. The strategy of sodium salinization of an organic compound and its graphene recombination technology will provide new ideas for the design of novel organic electrode materials with high performance for lithium-ion batteries.

Conflicts of interest

There are no conflicts to declare.

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