

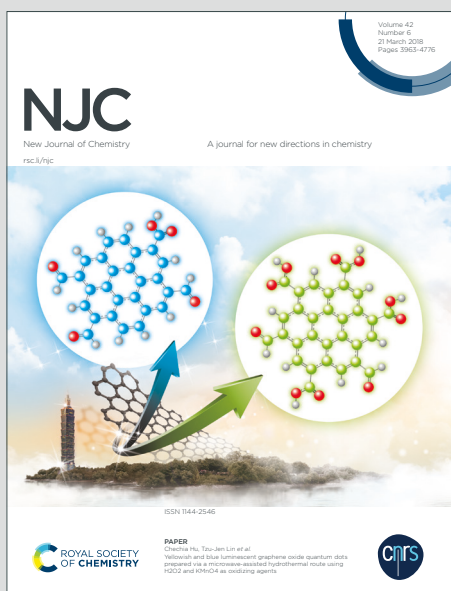
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Ultrastable Conductive Microporous Covalent Triazine Frameworks Based on Pyrene Moieties Provide High-Performance CO₂ Uptake and Supercapacitance

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Covalent organic triazine frameworks (CTFs) are a subclass of covalent organic frameworks and conjugated microporous polymers that exhibit heteroatom effects and possess high surface areas and excellent chemical and thermal stabilities; they have applications in energy storage and gas adsorption. In this study, we prepared two microporous polymers the pyrene-functionalized CTFs Pyrene-CTF-10 and Pyrene-CTF-20—through ionothermal treatment of 1,3,6,8-cyanopyrene (TCNPy) in the presence of molten zinc chloride (ZnCl₂) at 500 °C (at ZnCl₂-to-TCNPy molar ratios of 10:1 and 20:1, respectively). Pyrene-CTF-10 and Pyrene-CTF-20 had specific BET surface areas of 819 and 1019 m² g⁻¹, respectively, with pore size distributions of 1.10 and 1.35 nm, respectively, and high char yields (up to 70%). Furthermore, these pyrene-CTF polymeric frameworks exhibited excellent specific capacitances at a current density of 0.5 A g⁻¹: 380 F g⁻¹ for Pyrene-CTF-10 and 500 F g⁻¹ for Pyrene-CTF-20. In addition, both Pyrene-CTFs had excellent cycling stability, retaining 97% of their capacitances after cycling 2000 times at a current density of 10 A g⁻¹. Moreover, Pyrene-CTF-10 displayed good CO₂ uptake capacity (2.82 and 5.10 mmol g⁻¹ at 298 and 273 K, respectively).

Introduction

Supercapacitors, also known as electrochemical capacitors or electric double-layer capacitors (EDLCs), are being used more widely as replacements for fuel cells and batteries in energy storage applications, military devices, electric vehicles, and electrochemical devices because of their long lifetimes, rapid rates of charging and discharging, and high power densities.^{1,2} The electrochemical mechanisms behind the operation of supercapacitors can be divided into two categories: nonfaradaic processes (e.g., in EDLCs) depending on the accumulation of electrostatic charge at the electrode–electrolyte interface and faradaic processes (e.g., in pseudocapacitors) occurring at the surfaces of conjugated polymers, heteroatom-doped carbon materials, and metal oxide/hydroxide systems that feature fast redox reactions.^{2,3} Thus, the construction of materials for use as electrodes in energy storage applications requires that they have large specific surface areas, high nitrogen contents, large surface areas available to the electrolyte, and hierarchical porous structures.^{3–5} Anthropogenic carbon dioxide (CO₂) emissions will likely have many serious negative effects on human life and the environment because of increasing global temperatures and changes to weather events.^{6–9} Therefore, lowering emissions and the levels of CO₂ through sequestration and capture has become a challenge for both academia and industry. The wet cleaning method has been the most commonly used chemical process for capturing and separating CO₂ with aqueous alkanolamine solutions (amine scrubbing), but this method has the disadvantages of high energy consumption, low efficiency, difficulties of renewal, amine-mediated corrosion of equipment,

solvent loss, and high cost.^{10–15} As a result, we need new and effective methods for decreasing the emission and increasing the capture of CO₂ during industrial processes. Covalent organic polymers (COPs) are attractive porous materials for many applications, including sensors; the removal of pollutants; CO₂ capture, storage, and separation; drug delivery; catalysis; SO₂ adsorption; gene therapy; Li-S batteries; energy conversion; and light-harvesting. They have many interesting properties: outstanding thermal stability, high BET surface areas, accessible porosity, large pore volumes, chemical resistance, biocompatibility, and excellent electrical conductivity when compared with metal-organic frameworks.^{16–26} The many types of COP materials include covalent organic frameworks (COFs), covalent triazine frameworks (CTFs), porous aromatic frameworks, benzimidazole-linked polymers, carbazole-based microporous polymers, and porous imine-linked networks.^{27–34} Amine scrubbing adsorbs CO₂ through chemisorption, as opposed to the physisorption mechanism of COPs.^{10,12,34} Post-synthesis or bottom-up approaches can be used to incorporate CO₂-philic functional groups [e.g., carboxyl (COOH), hydroxyl (OH), imine (–N=CH), benzimidazole, carbazole, and triazine units] into covalent organic porous materials to improve their CO₂ capture.^{34–38} CTFs are both COFs and conjugated microporous polymers that display large heteroatom effects, possess permanent nanopores with high Brunauer–Emmett–Teller (BET) surface areas, have excellent chemical and thermal stabilities, and feature tunable electronic structures.^{39–43} Most CTF framework materials have been prepared through ionothermal trimerization of nitrile groups, superacid catalysis, and phosphorous pentoxide (P₂O₅) catalysis.^{28,43–45} Many recent reviews have highlighted the great potential of CTF materials in gas separation and storage, heterogeneous catalysis, energy storage, and photocatalysis.^{46–}

⁴⁸ Porous polymeric materials containing pyrene units have been tested recently in many different fields—including chemical sensing, photoelectric devices, gas storage and separation, and photocatalysis—because of their high propensity for forming excimers, distinct solvatochromic phenomena, ready functionalization, delayed fluorescence, and tunable photophysical properties.^{49–51}

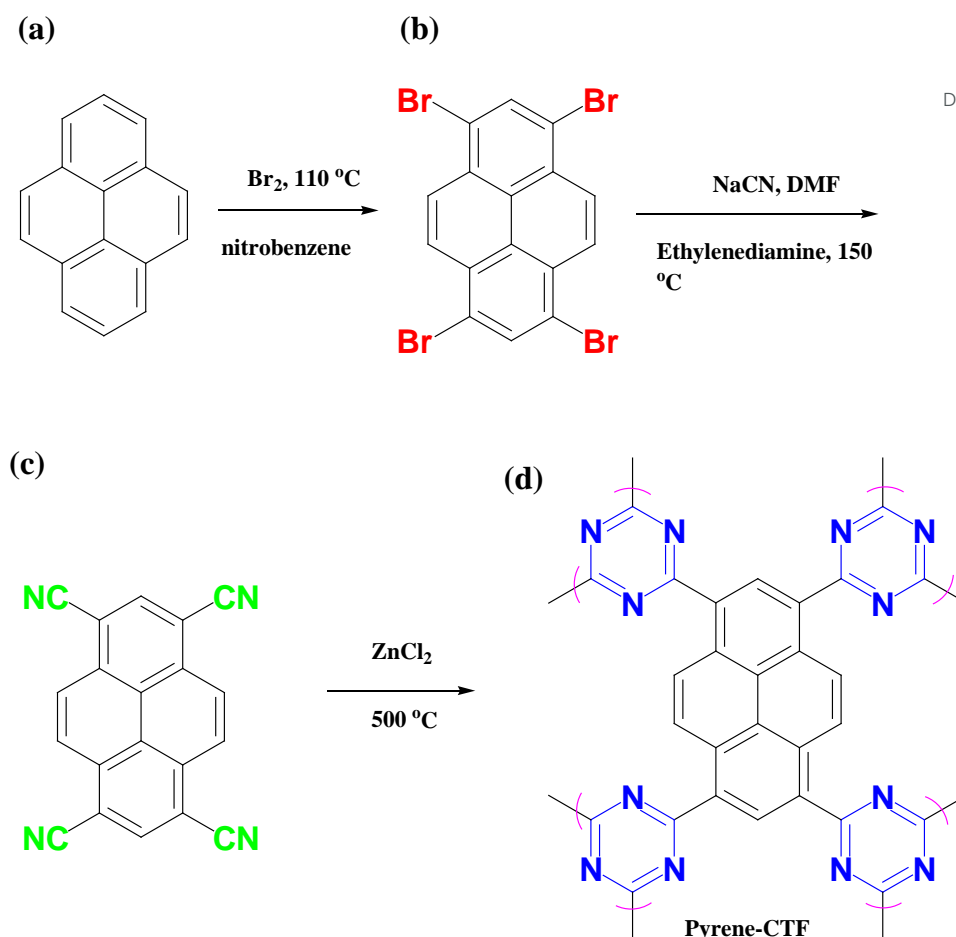
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Scheme 1. Synthesis of (d) Pyrene-CTF from (a) Pyrene, (b) TBrPy, and (c) TCNPy.

In a previous study, we prepared a series of conductive bicarbazole-based CTFs (Car-CTFs) through the trimerization of [9,9'-bicarbazole]-3,3',6,6'-tetracarbonitrile (Car-4CN) with molten zinc chloride (ZnCl_2); these materials exhibited outstanding thermal stability, excellent energy storage, and CO_2 uptake.²⁸ Janiak et al. prepared the CTFs PCTF-1 and PCTF-2 through reactions of tetra(4-cyanophenyl)ethylene with ZnCl_2 at 400 °C; these two materials captured CO_2 at levels of 3.30 and 1.87 mmol g^{-1} , respectively, at 273 K.⁵² Voort et al. prepared CTF materials from 4,4',4'',4'''-(1,4-phenylenebis(pyridine-4,2,6-triyl))tetrabenzonitrile in anhydrous ZnCl_2 under ionothermal conditions at 400 and 500 °C, with the former material (CTF-20-400) providing a CO_2 uptake of 3.48 mmol g^{-1} at 273 K.⁸ Han et al. prepared the CTF TPC-1 through defluorination carbonization and trimerization of the nitrile groups of tetrafluoroterephthalonitrile in molten ZnCl_2 ; this polymeric framework had a high specific surface area (1940 $\text{m}^2 \text{g}^{-1}$) and displayed excellent CO_2 adsorption (4.9 mmol g^{-1} at 273 K). In this study, we prepared pyrene-based CTFs (Pyrene-CTFs) through trimerization of the nitrile groups of 1,3,6,8-cyanopyrene (TCNPy) in the presence of molten ZnCl_2 at 500 °C (Scheme 1). We used wide-angle X-ray diffraction (WAXD), Fourier transform infrared (FTIR) spectroscopy, X-ray photoelectron spectroscopy (XPS), Raman spectroscopy, transmission electron microscopy (TEM), thermogravimetric analysis (TGA), and the Brunauer–Emmett–Teller (BET) method to examine the properties of these Pyrene-CTFs

(crystallinity BET surface areas, porosities, chemical surface compositions, thermal stability, and chemical structures). We performed electrochemical measurements and CO_2 uptake analyses to examine their potential applications as high-performance electrode materials and for gas adsorption.

Experimental Section

Materials

Pyrene, nitrobenzene, HCl solution (37%), ethanol (EtOH), and methanol (MeOH) were purchased from Acros. Ethylenediamine (99%), tetrahydrofuran (THF), *N,N*-dimethylformamide (DMF), dichloromethane (DCM), acetone, and copper(I) cyanide (CuCN) was purchased from Sigma–Aldrich.

1,3,6,8-Tetrabromopyrene (TBrPy)^{54,55}

A solution of bromine (1.15 mL, 22 mmol) in nitrobenzene (5 mL) was added dropwise to a solution of pyrene (1 g) in nitrobenzene (15 mL) and then the mixture was heated at 120 °C for 4 h. Cooling to room temperature provided a pale-green solid, which was filtered off, washed three times with EtOH, and dried under vacuum at 100 °C to provide a pale-green solid (2.21 g, 91%). M.p.: >300 °C.

1,3,6,8-Cyanopyrene (TCNPy)

A mixture of TBrPy (3.00 g, 4.65 mmol) and CuCN (9.69, 108 mmol) in dry DMF (100 mL) was heated under reflux under a N_2

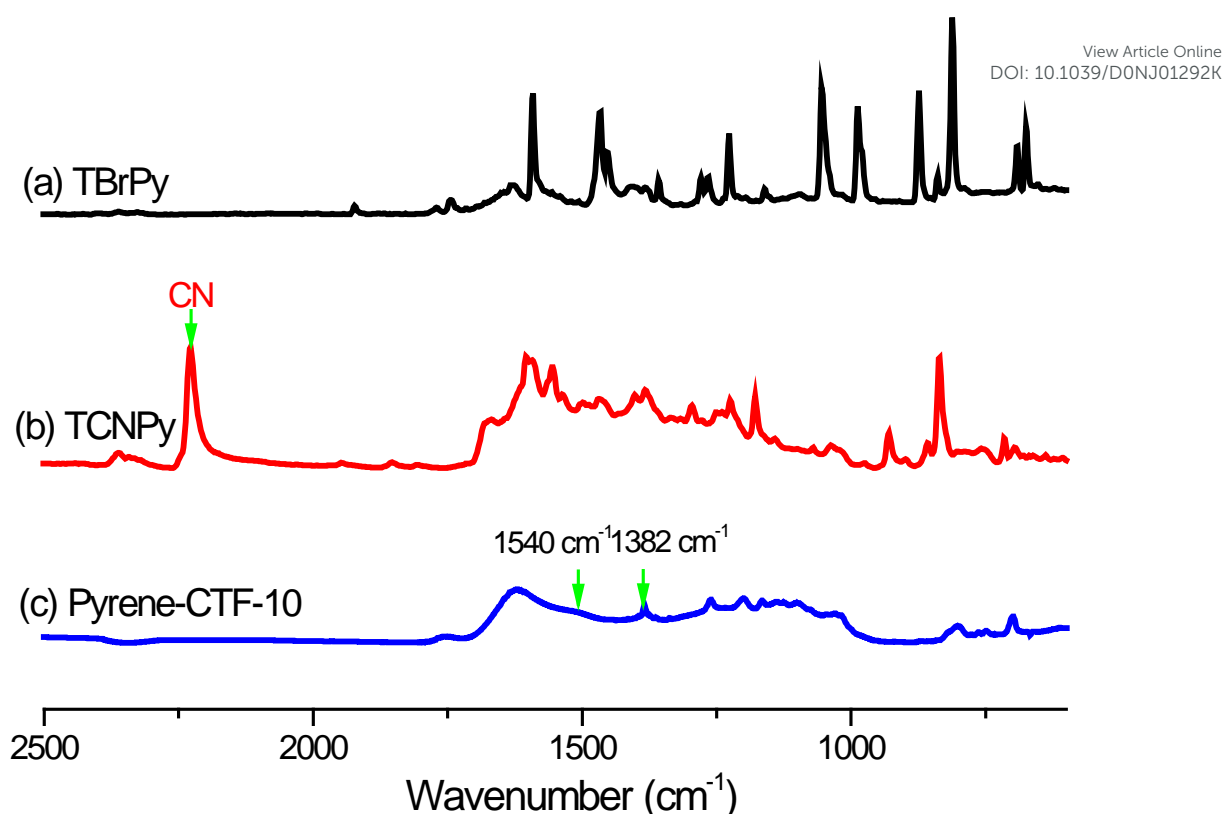


Figure 1. FTIR spectra of (a) TBrPy, (b) TCNPy, and (c) Pyrene-CTF-10.

atmosphere at 140 °C for 3 days. The mixture was opened to the air and cooled to room temperature, at which point it was then filtrated to remove unreacted CuCN. The solution was poured into a mixture of cold water (75 mL) and ethylenediamine (4 mL); this mixture was heated at 50 °C for 2 h. The yellow powder was filtered off and washed with water and MeOH to afford a yellow powder (1.24 g). M.p.: >400 °C; FTIR (KBr, cm^{-1}): 3070 (C–H stretching), 2220 (CN stretching).

Pyrene-CTFs

In an ampoule, TCNPy (200 mg, 0.46 mmol) was mixed with anhydrous ZnCl_2 [63 mg (4.63 mmol) or 126 mg (9.26 mmol)] (Scheme 1). The temperature was raised to 500 °C at a heating rate of 5 °C/min and then the samples were maintained at that temperature for 48 h under the N_2 atmosphere. After cooling, each obtained black solid was washed with 1 M HCl and water (each for 24 h). The resultant black solids (Py-CTF-10, Py-CTF-20) were washed several times with THF, DCM, and MeOH. Finally, the black powders were dried for 24 h at 100 °C.

Results and Discussion

Synthesis of TBrPy, TCNPy, and Pyrene-CTFs

The monomer 1,3,6,8-cyanopyrene (TCNPy) was synthesized in two steps (Scheme 1). First, pyrene was reacted with bromine (Br_2) in nitrobenzene at 120 °C for 5 h to afford TBrPy as a pale-green solid, which was reacted with CuCN in DMF at 140 °C for 48 h to give TCNPy as a yellow powder. We then used an ionothermal method for trimerization of the nitrile groups of TCNPy in the presence of molten anhydrous ZnCl_2 (at weight ratios of 1:10 and 1:20) at 500 °C

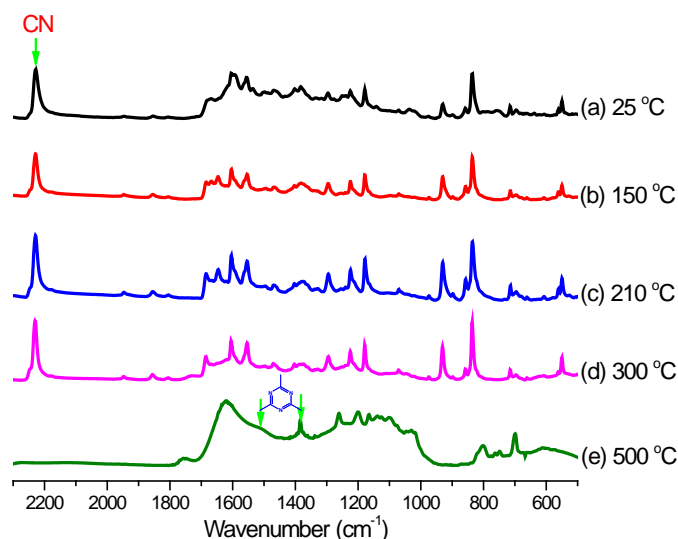


Figure 2. FTIR spectral profile of Pyrene-CTF-10, recorded at temperatures from room temperature to 500 °C.

to give Pyrene-CTF-20 and Pyrene-CTF-20, respectively (Scheme 1, Table 1). Figure 1 displays the FTIR spectra of TBrPy, TCNPy, and the Pyrene-CTFs. The spectrum of TBrPy [Figure 1(a)] featured characteristic absorption bands centered at 3050 and 804 cm^{-1} , assigned to the stretching of aromatic C–H^{56–59} and C–Br bonds, respectively. The FTIR spectrum of TCNPy [Figure 1(b)] contained an absorption band at 2220 cm^{-1} representing the stretching of CN triple bonds, confirming its synthesis. As displayed in Figures 1(c), the spectra of the Pyrene-CTFs featured absorption bands

Table 1. Thermal stabilities, porosity parameters, Raman spectral data, and electrochemical data of the Pyrene-CTFs.

Sample	TCNPy/ ZnCl ₂	<i>T</i> _{d10%} (°C)	Char yield (%)	<i>S</i> _{BET} (m ² g ⁻¹)	Pore size (nm)	Pore volume (cm ³ g ⁻¹)	<i>I</i> _D / <i>I</i> _G (Raman)	CV (F g ⁻¹)
Pyrene CTF-10	1:10	500	72	819	1.10	0.88	1.07	380
Pyrene-CTF-20	1:20	474	72	1019	1.35	1.12	1.47	500

at 1565 and 1365 cm⁻¹ representing the triazine rings, but no signals for CN groups, confirming the successful trimerization/cyclization reactions of the cyano groups.^{8,28,60} In addition, the intensities of the characteristics absorption bands of the triazine ring in Pyrene-CTF-20 gradually decreased at 500 °C, due to the occurrence partial carbonization [Figure S1].^{61,62} We performed FTIR spectral analyses to monitor the trimerization of the nitrile groups in the preparation of Pyrene-CTF-10 from room temperature to 500 °C (Figure 2). For temperatures between 25 and 300 °C, the FTIR spectrum of Pyrene-CTF-10 featured the characteristic absorption band at 2220 cm⁻¹ for stretching of the CN triple bonds. When the temperature reached 500 °C, however, this characteristic band disappeared completely and two new absorption signals appeared at 1565 and 1360 cm⁻¹, confirming the complete trimerization of the nitrile groups and the formation of triazine rings.²⁸

We used solid state ¹³C CP/MAS NMR spectroscopy (Figure S2) to confirm the chemical structures of TBrPy and TCNPy and the formation of triazine rings in the Pyrene-CTFs materials. The solid state ¹³C NMR spectrum of TBrPy [Figure S2(a)] displays signals at 127, 116 and 110 ppm at for the aromatic carbon atoms. Figure S2(b) reveals that the spectrum of TCNPy features signals at 132 and 108 ppm for the aromatic carbon nuclei and signal at 116 ppm for CN group. The solid state ¹³C NMR spectra [Figures S2(c) and (d)] of Pyrene-CTF-10 and Pyrene-CTF-20 exhibited signals at 168, 136, 123 and 116 ppm which we assign to the carbon nuclei of the triazine [-C(Ar)=N-] and the pyrene units.^{8,28,63} We determined the thermal stabilities of TCNPy, Pyrene-CTF-10, and Pyrene-CTF-20 through TGA under a N₂ atmosphere (Figure 3). Although Pyrene-CTF-10 and Pyrene-CTF-20 were synthesized in molten ZnCl₂ at 500 °C, the decomposition temperature (*T*_{d10%}) of Pyrene-CTF-10 (500 °C) was higher than that of Pyrene-CTF-20 (474 °C); they had the same char yield of 71 wt.%. These values for the TCNPy monomer were 346 °C and 4 wt.%, respectively. Thus, these Pyrene-CTFs displayed outstanding thermal stability.^{8,28}

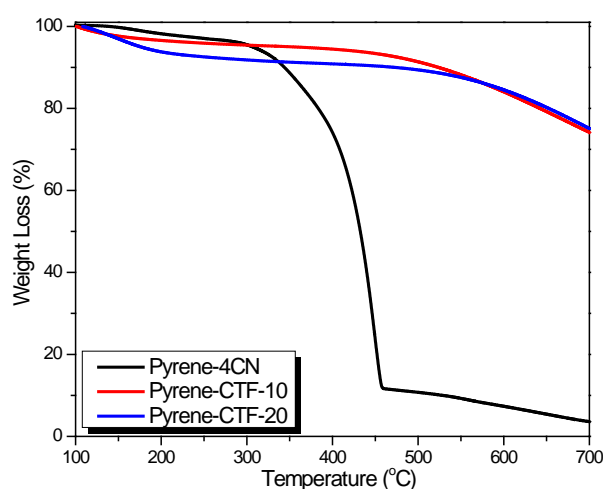
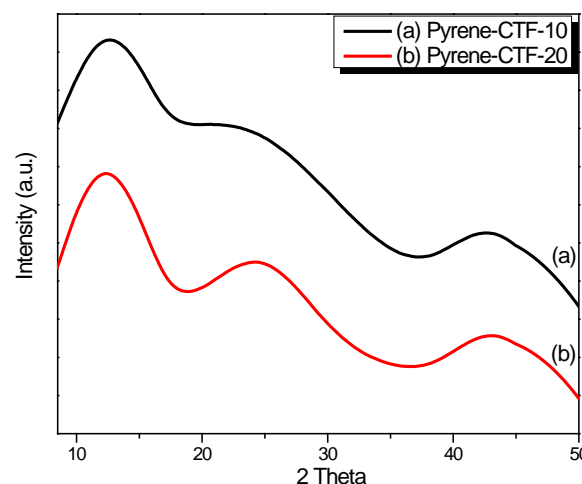
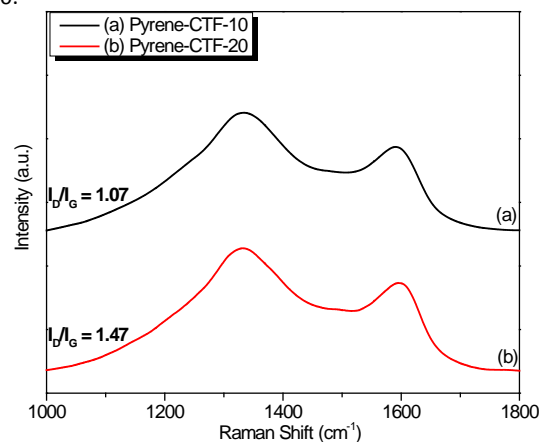
**Figure 3.** TGA analyses of TCNPy, Pyrene-CTF-10, and Pyrene-CTF-20, recorded under N₂.**Figure 4.** XRD patterns of Pyrene-CTF-10 and Pyrene-CTF-20.

Figure 4 provides the XRD patterns of Pyrene-CTF-10 and Pyrene-CTF-20, revealing that they had partially crystalline structures with two broad peaks at 13 and 25° that we assign to the vertical spacing between stacked sheets (001) and the in-plane reflection (100), respectively.^{15,64} Figure 5 presents the Raman spectra of Pyrene-CTF-10 and Pyrene-CTF-20 in the range from 1000 to 1900 cm⁻¹. The spectra of both Pyrene-CTFs materials featured two strong signals characteristic of D and G bands, suggesting the existence of graphitic carbonized structures. In general, the D and G bands represent the graphitic carbon atoms: second-order Raman scattering with sp³ hybridization and first-order Raman scattering with sp² hybridization, respectively.^{15,28,64,65} From the Raman spectra of Pyrene-CTF-10 and Pyrene-CTF-20, the D bands were centered at 1342 and 1335 cm⁻¹, respectively, while the G bands appeared at 1584 and 1586 cm⁻¹, respectively. The *I*_D/*I*_G ratios for Pyrene-CTF-10 and Pyrene-CTF-20 were 1.07 and 1.47, respectively. Thus, Pyrene-CTF-10 possessed a lower degree of graphitization and featured less defects in its structure when compared with Pyrene-CTF-20.

**Figure 5.** Raman spectra of Pyrene-CTF-10, and Pyrene-CTF-20.

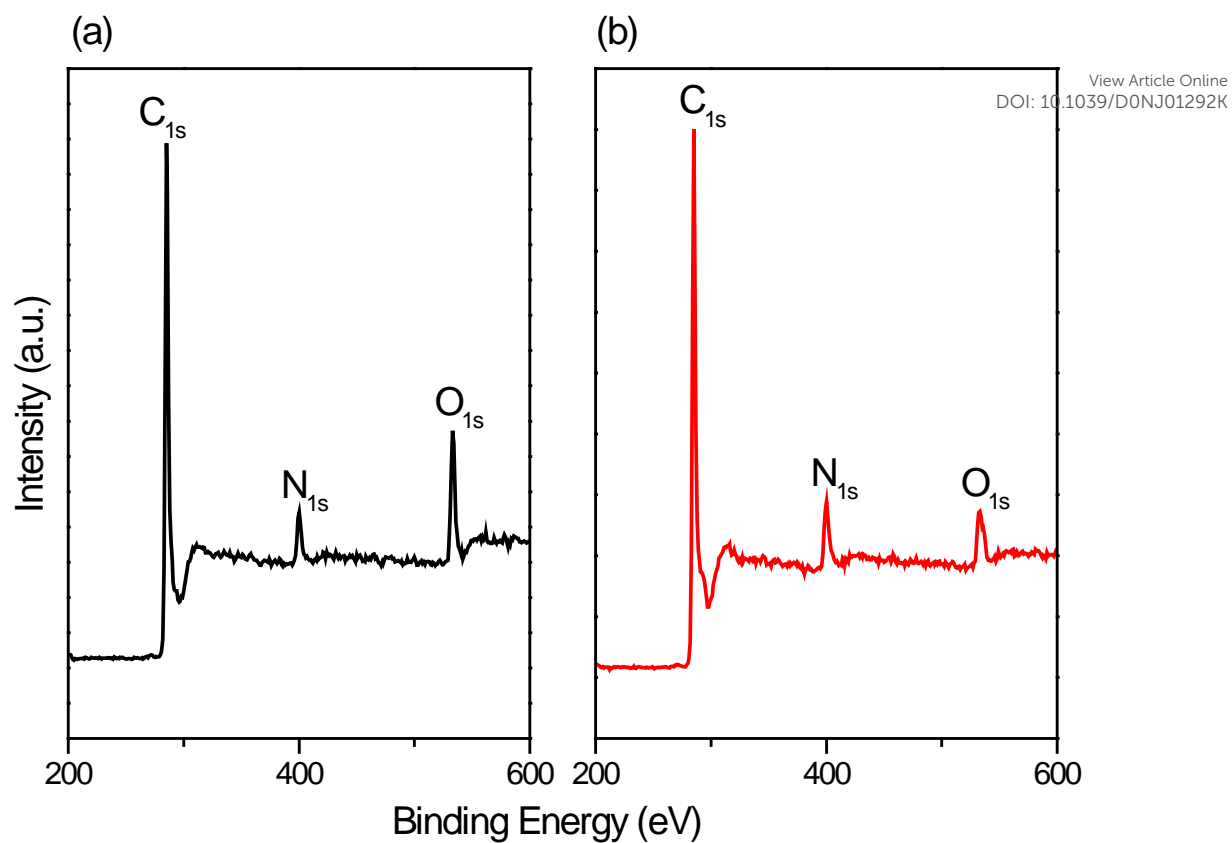


Figure 6. XPS survey spectra of (a) Pyrene-CTF-10 and (b) Pyrene-CTF-20.

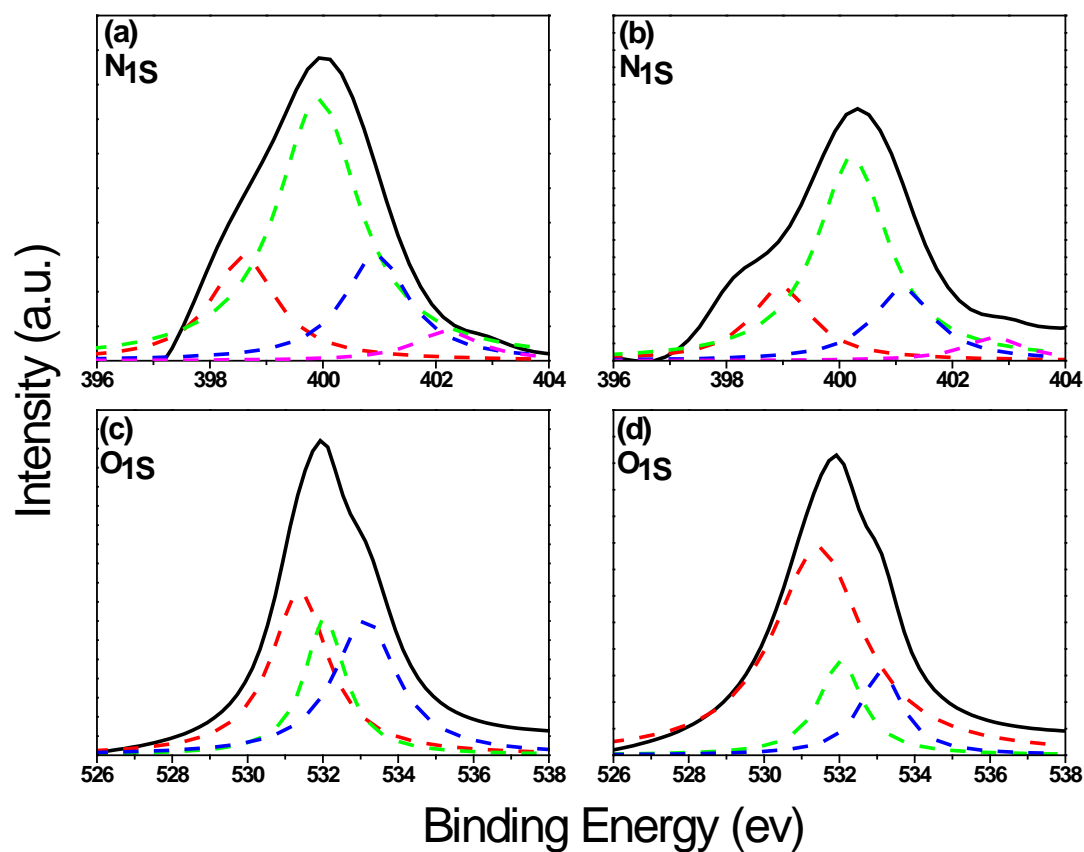


Figure 7. XPS fitting curves of N 1s and O 1s orbitals: (a, c) Pyrene-CTF-10; (b, d) Pyrene-CTF-20.

Table 2. Curve-fitting data and area fractions of the signals in the N1s and O1s plots in **Figure 7**.

Sample	N species			O species		
	N-6	N-5	N-Q	C-O	C-OH	H ₂ O
Pyrene-CTF-10	68.24	21.44	10.32	45.29	15.61	39.10
Pyrene-CTF-20	59.92	23.97	16.12	69.16	15.58	30.84

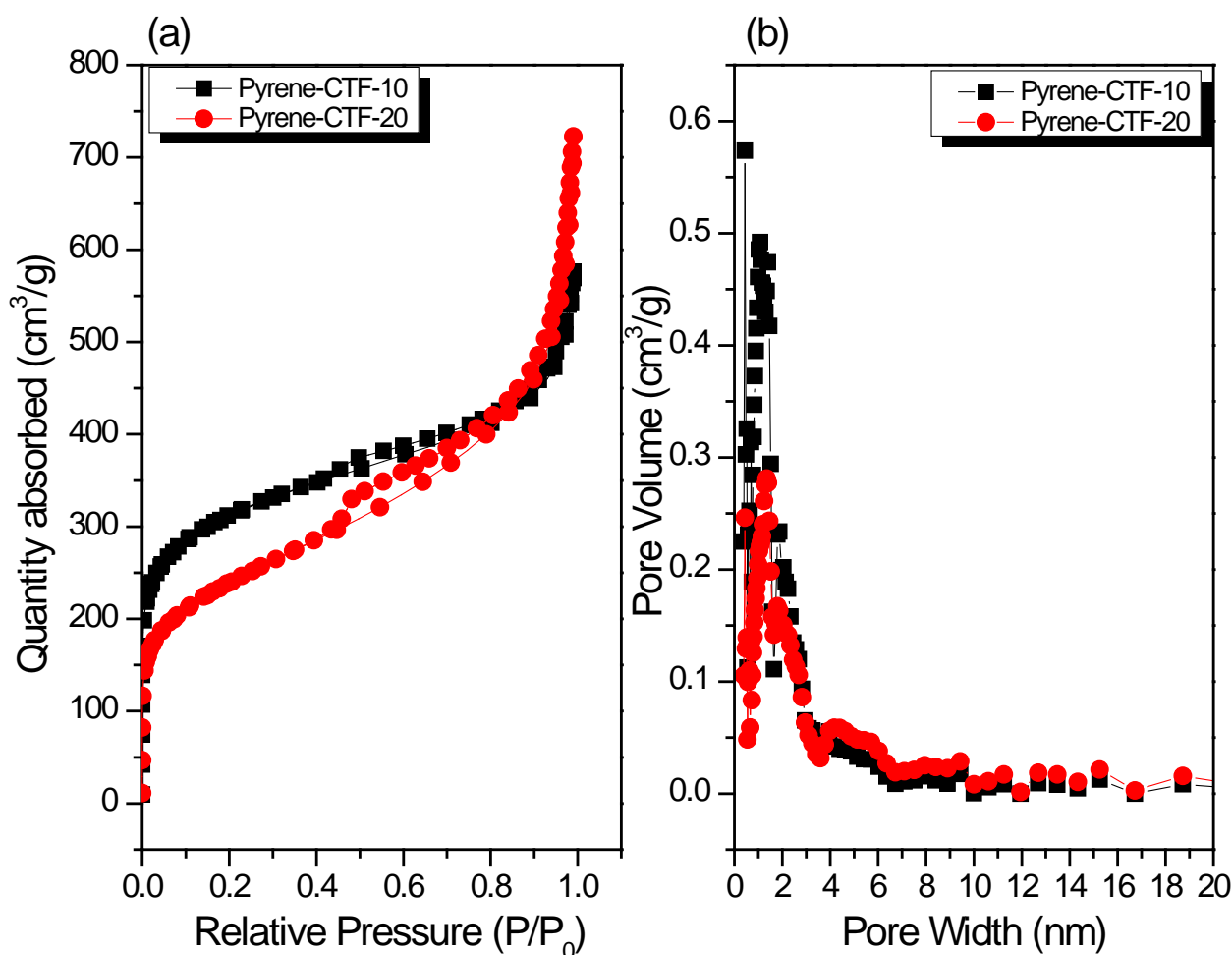
**Figure 8.** (a) N₂ adsorption/desorption measurements and (b) pore size distributions of Pyrene-CTF-10 and Pyrene-CTF-20, recorded at 77 K.

Figure 6 displays the XPS spectra of the Pyrene-CTFs. Three peaks appeared in each spectrum: at 284 eV (for carbon atoms in the polymeric CTF framework), 400 eV (C–N bonds for the N 1s orbitals in the triazine units), and 530 eV (O 1s orbitals for absorbed water and oxygen), respectively.^{28,53,66,67}

We fitted the XPS curves for the N 1s and O 1s orbitals to investigate the chemical compositions on the surfaces of these Pyrene-CTFs (Figure 7, Table 2). The fitting results for the N 1s orbitals [Figures 7(a) and 7(b)] revealed the presence of three N species, including quaternary-N species (401 eV), pyrrolic species (400 eV), and hexagonal pyridinic N atoms from the triazine ring (398.7 eV). According to quantitative analysis, the pyrrolic N and pyridinic species were the most abundant in the Pyrene-CTF frameworks. Furthermore, the surfaces of the Pyrene-CTF materials contained three O species [Figures 7(c) and 7(d)], with signals at 531, 532, and 533 eV corresponding to polarized C–O bonds, water, and absorbed oxygen.^{28,53,66,67}

We measured N₂ adsorption/desorption isotherms (Figure 8, Table 1) at 77 K to examine the porosities of our Pyrene-CTFs. Figure 8(a) reveals that Pyrene-CTF-10 had microporous characteristics, with a specific surface area of 819 m² g^{−1} and a pore volume of 0.88 cm³ g^{−1}. The BET curve of Pyrene-CTF-20 featured a type IV isotherm, with a high specific surface area of 1019 m² g^{−1} and a pore volume of 1.12 cm³ g^{−1}. Thus, the specific BET surface area of Pyrene-CTF-20 was higher than that of Pyrene-CTF-10, suggesting the presence of closely packed nanoparticles and a larger number of defects in the Pyrene-CTF-20 structure.^{8,28} Furthermore, we used NL-DFT theory to calculate the pore diameters in Pyrene-CTF-10 and Pyrene-CTF-20 [Figure 8(b), Table 1] to be 1.10 and 1.35 nm, respectively. TEM images (Figure 9) revealed that Pyrene-CTF-10 and Pyrene-CTF-20 both possessed irregular microporous structures.

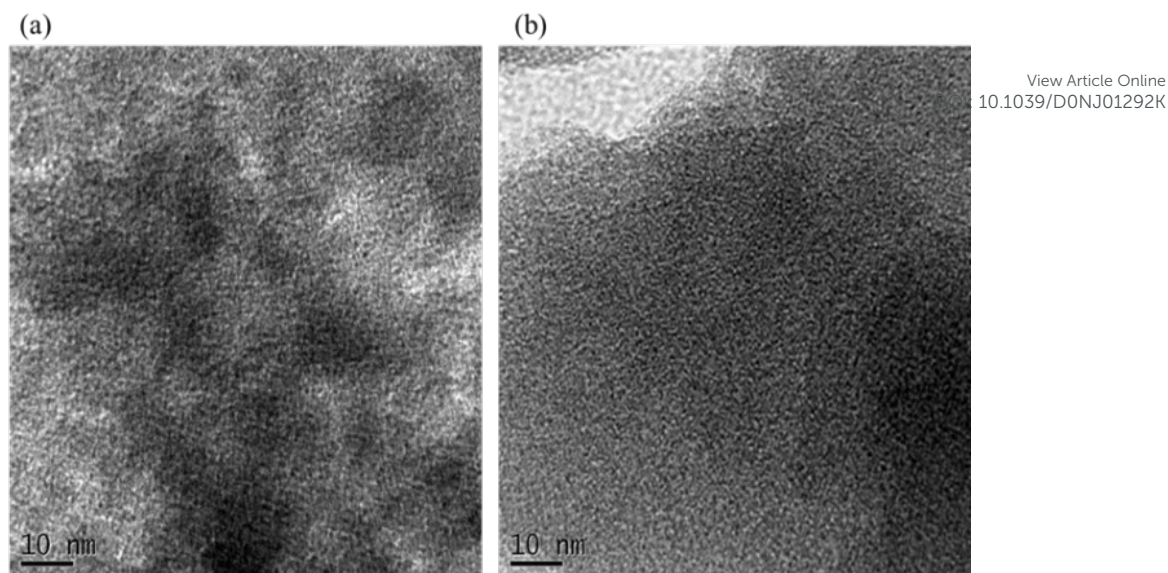


Figure 9. TEM images of (a) Pyrene-CTF-10 and (b) Pyrene-CTF-20.

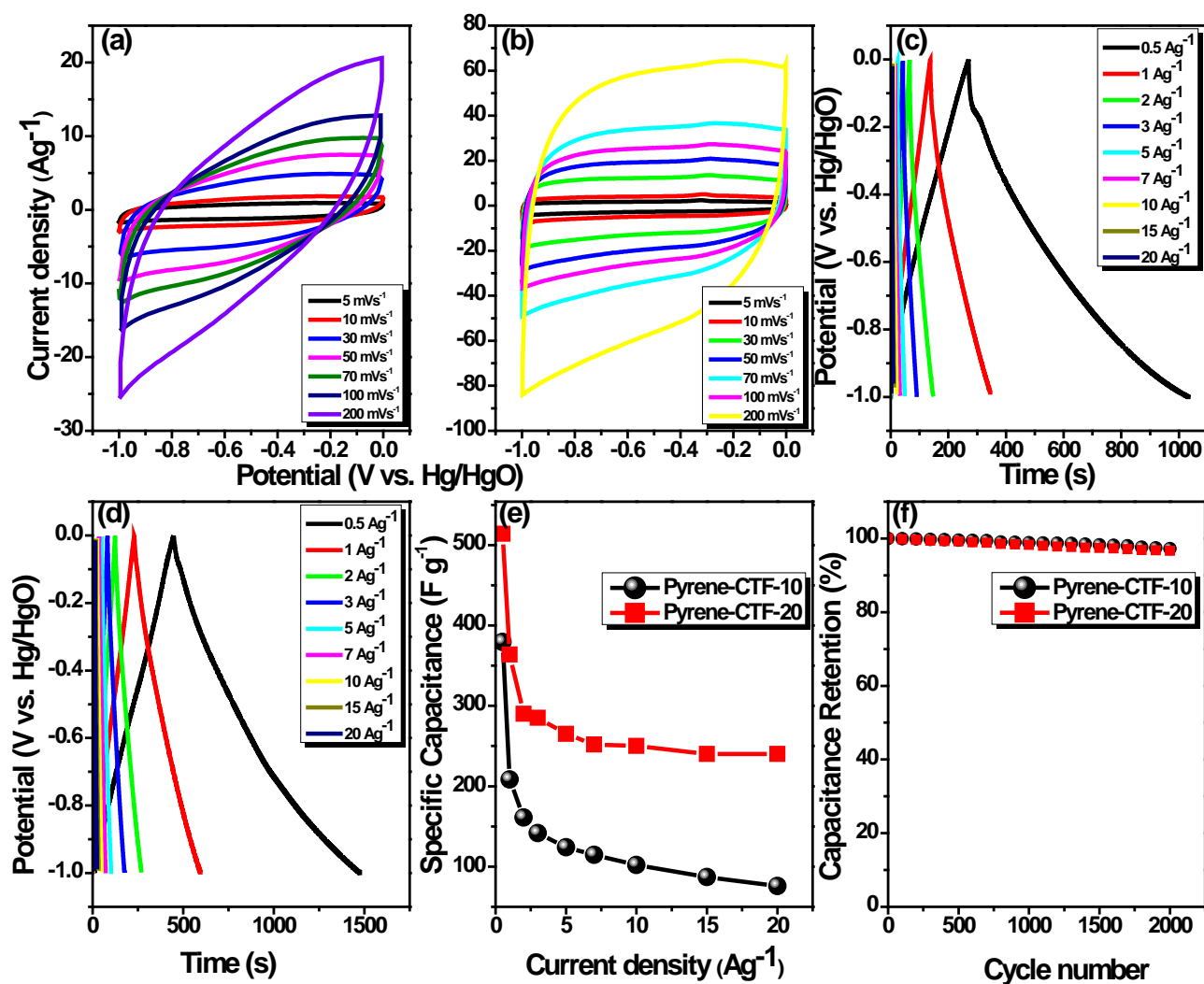


Figure 10. (a, b) CV curves of Pyrene-CTF-10 and Pyrene-CTF-20. (c, d) GCD profiles of Pyrene-CTF-10 and Pyrene-CTF-20, recorded at various currents. (e) Specific capacitances of Pyrene-CTF-10 and Pyrene-CTF-20 recorded at current densities from 0.5 to 20 A g⁻¹. (f) Cycling stabilities of Pyrene-CTF-10 and Pyrene-CTF-20 electrodes, recorded at a current density of 10 A g⁻¹ over 2000 cycles.

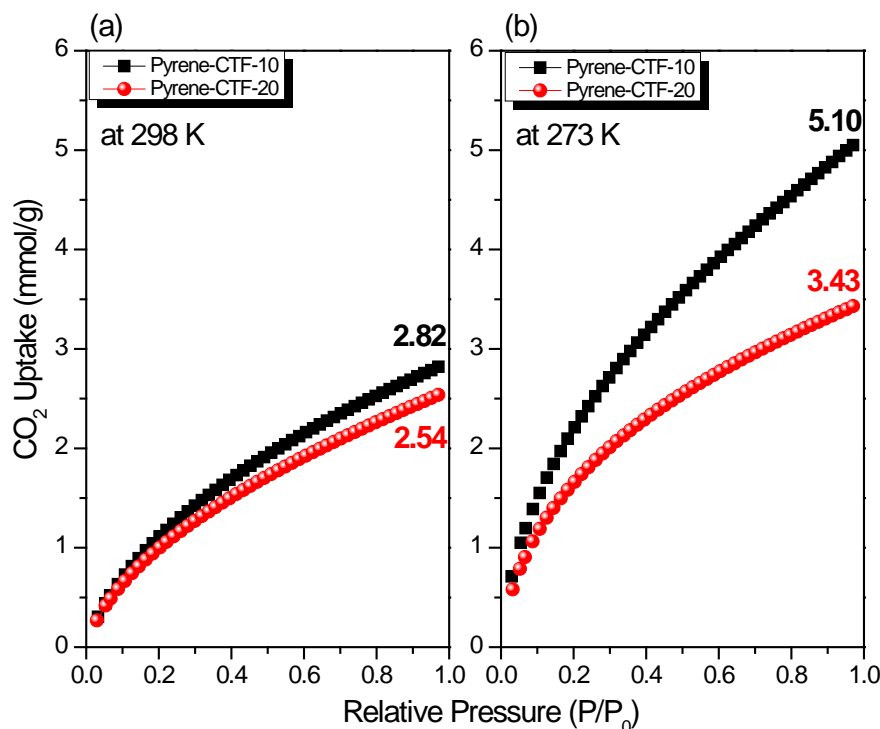


Figure 11. CO₂ uptake of (a) Pyrene-CTF-10 and (b) Pyrene-CTF-20, measured at 298 and 273 K.

Energy Storage and CO₂ Uptake of Pyrene-CTFs

From the BET and XPS data, we knew that our Pyrene-CTF materials had high specific surface areas, N-heteroatom frameworks (primarily pyridinic and pyrrolic N species), and mesoporous and microporous properties. Therefore, we expected these materials to have great potential applications in energy storage and gas uptake.²¹ We measured the electrochemical properties of the Pyrene-CTFs through cyclic voltammetry (CV) at various scan rates (5–200 mV s⁻¹) in 1 M aqueous KOH [Figures 10(a) and 10(b)]. The CV profiles for the Pyrene-CTFs had rectangular (“rice-like”) shapes.^{2,4,28} The appearance of rectangular CV curves for these CTF materials based on pyrene units suggested a combination of electric double-layer capacitance and pseudocapacitance, arising from the presence of various types of nitrogen and oxygen species (heteroatom effects, based on XPS data) in both of these polymeric CTF materials.^{2,4,28} In addition, the galvanostatic charge/discharge (GCD) profiles of Pyrene-CTF-10 and Pyrene-CTF-20 recorded at various current densities [Figures 10(c) and 10(d)] had triangular shapes, again suggesting the presence of both EDLC and pseudocapacitance arising from the graphitic carbon structures presenting various types of functionalized units (e.g., pyrrolic species; hexagonal pyridinic N atoms in triazine rings; C=O and phenolic OH groups).^{2,21} As displayed in Figure 10(e), at a current density of 0.5 A g⁻¹, Pyrene-CTF-10 and Pyrene-CTF-20 displayed excellent specific capacitances of 380 and 500 F g⁻¹, respectively. The higher specific capacitance of Pyrene-CTF-20 compared with that of Pyrene-CTF-10 presumably arose from its graphitic structure, microporosity, and high N (7.19 wt%) and O atom (9.62 wt%) contents. In addition, the specific capacitances of the Pyrene-CTFs decreased upon increasing the scan rate, due to the adsorption of electrolyte ions on the electrode surfaces.^{8,68,69} In addition, we investigated the cyclic stabilities of Pyrene-CTF-10 and Pyrene-CTF-20 through GCD measurements over 2000 cycles at a current density of 10 A g⁻¹

[Figure 10(f)]. Pyrene-CTF-10 and Pyrene-CTF-20 both retained 97% of their initial capacitance after 2000 cycles, indicating their excellent cycling stability in 1 M KOH as the electrolyte. Deng et al. prepared conductive CTFs through ionothermal reactions of tetracyanoquinodimethane (TCNQ) in the presence of molten ZnCl₂ at various temperatures. They found that the specific capacitances of TCNQ-CTF-600, TCNQ-CTF-700, TCNQ-CTF-800, and TCNQ-CTF-900 at a current density of 0.2 A g⁻¹ were 284, 350, 380, and 356 F g⁻¹, respectively. The highest specific capacitance (380 F g⁻¹) was that of TCNQ-CTF-800, presumably because of its high N-atom content (8.13%) and large surface area (3663 m² g⁻¹).² In addition, Zhi et al. derived amorphous CTFs from terephthalonitrile that displayed a specific capacitance of 298 F g⁻¹ at 0.2 A g⁻¹.⁷⁰ In addition, we have previously reported a series of Car-CTFs formed through trimerization of Car-4CN with molten ZnCl₂; among them, Car-CTF-5-500 provided a specific capacitance of 545 F g⁻¹ at 5 mV s⁻¹.²⁸ In comparison with these other CTF materials, Pyrene-CTF-20 displays excellent supercapacitor performance (Table S1), presumably because of its graphitic structure, microporosity, and N and O atom contents. Figure S3 shows that the calculated coulombic efficiency (CE) with the current density. It shows that the CE is very high at current density of 0.5 and 1 A g⁻¹, this is due to the very small applied charging and discharging currents. While at current density >1 A g⁻¹, the CE is almost ~100%. We recorded CO₂ adsorption isotherms for both Pyrene-CTFs at 25 and 0 °C [Figures 11(a) and 11(b)]. The CO₂ uptake of Pyrene-CTF-10 was 2.82 and 5.10 mmol g⁻¹ at 25 and 0 °C, respectively; for Pyrene-CTF-20, these values were lower at 2.54 and 3.43 mmol g⁻¹, respectively. Thus, Pyrene-CTF-10 captured more CO₂ at 25 °C (2.82 mmol g⁻¹) than other previously reported CTF materials at the same temperature, including Car-CTF-10-500 (2.25 mmol g⁻¹),²⁸ CTF-1 (2.77 mmol g⁻¹),⁷¹ PCTF-1-3 (2.02–3.00 mmol g⁻¹),⁷² CTF-20-400 (3.48 mmol g⁻¹),³ CTF-0 (1.54–2.34 mmol g⁻¹),⁷³ and CTF-DCN-500 (2.70 mmol g⁻¹),⁶² due to Pyrene-CTF-10 having a specific surface area of 819 m² g⁻¹ and a high N-atom content (9.85 wt%) arose from the presences of triazine units, which leads to increase the binding affinity interaction between CO₂ molecules and Pyrene-CTF-10. Table S2

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provides a performance comparison Pyrene-CTFs and other reported CTFS for CO₂ uptake. Thus, our CV and CO₂ uptake measurements suggest that Pyrene-CTFs have great potential for use in energy storage and gas storage.

Conclusions

We have synthesized two pyrene-containing conductive CTFS, Pyrene-CTF-10 and Pyrene-CTF-20, through reactions of TCNPY with molten ZnCl₂ (as catalyst) at 500 °C. These Pyrene-CTFs exhibited outstanding thermal stabilities (high char yields), partial crystallinity, and high specific surface areas, based on BET, XRD, and TGA analyses. In addition, Pyrene-CTF-10 displayed gas uptake capacities for CO₂ of 2.82 mmol g⁻¹ at 25 °C and 5.10 mmol g⁻¹ at 0 °C. More interestingly, the electrochemical performances of our two Pyrene-CTFs were highly stable and efficient, with highest capacitances at 0.5 mV s⁻¹ of 380 F g⁻¹ for Pyrene-CTF-10 and 500 F g⁻¹ for Pyrene-CTF-20.

Characterization

FTIR spectra were recorded using a Bruker Tensor 27 FTIR spectrophotometer and the conventional KBr disk method; 32 scans were collected at a spectral resolution of 4 cm⁻¹. The films used in this study were sufficiently thin to obey the Beer-Lambert law. Wide-Angle X-ray diffraction (WAXD) pattern was obscured from the wiggler beamline BL17A1 of the National Synchrotron Radiation Research Center (NSRRC), Taiwan. A triangular bent Si (111) single crystal was used to obtain a monochromated beam having a wavelength (λ) of 1.33 Å. Cross-polarization with MAS (CPMAS) was used to acquire ¹³C NMR spectral data at 75.5 MHz. The CP contact time was 2 ms; ¹H decoupling was applied during data acquisition. The decoupling frequency corresponded to 32 kHz. The MAS sample spinning rate was 10 kHz. Transmission electron microscope (TEM) images were obtained with a JEOL JEM-2010 instrument operated at 200 kV. BET surface area and porosimetry measurements of the prepared samples (ca. 40-100 mg) were performed using a Micromeritics ASAP 2020 Surface Area and Porosity analyzer. Nitrogen isotherms were generated through incremental exposure to ultrahigh-purity N₂ (up to ca. 1 atm) in a liquid nitrogen (77 K) bath. Surface parameters were determined using BET adsorption models in the instrument's software. TGA was performed using a TA Q-50 analyzer under a flow of N₂ atmosphere. The samples were sealed in a Pt cell and heated from 40 to 800 °C at a heating rate of 20 °C min⁻¹ under a flow of N₂ atmosphere at a flow rate of 60 mL min⁻¹. The Raman spectra were investigated using Horiba Jobin-Yvon HR800 Raman Spectrometer with 633 nm laser, 10 sec accumulated scans repeated for 20 times and 50x magnification lens. The electrochemical performances were performed using Zahner Zennium E electrochemical workstation with three electrodes configurations using Hg/HgO as a reference electrode and 1.0 M KOH solution as an aqueous medium. The working electrode was the glassy carbon electrode (GCE) coated by the slurry of the tested material (The slurries were prepared by dispersing the Pyrene-CTFs (45 wt. %), carbon black (45 wt. %), and Nafion (10 wt. %) in EtOH (1 mL)). The cyclic voltammetry (CV) was tested over the range between 0.0 and -1.0 V and the scan rates were investigated from 5 mV/s up to 200 mV/s. Similarly, the charge and discharge (GCD) cycles were tested at the same potential

range with different current densities from 0.5 to 20 Ag⁻¹. The capacitances (Cs) were calculated using the reported equations for CV and GCD respectively.^{74,75}

Conflicts of interest

There are no conflicts to declare.

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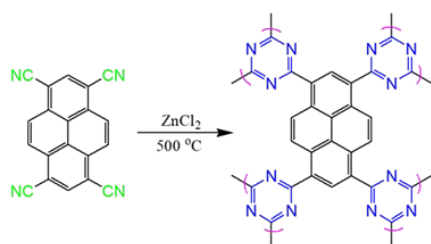
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TOC Graphic



Specific surface area: 1000 m²/g
CO₂ uptake at 0 °C: 5.10 mmol/g
Specific capacitance: up to 380 F g⁻¹

Two pyrene-functionalized CTFs through ionothermal treatment TCNPY in the presence of molten zinc chloride at 500 °C, which displayed high-performance CO₂ uptake and supercapacitance