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Insight into Sulfur Confined in Ultramicroporous Carbon

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ABSTRACT: Here, we provide a deeper insight into the state of sulfur confined in ultramicroporous carbon (UMC) and clarify its electrochemical reaction mechanism with lithium by corroborating the results obtained using various experimental techniques, such as X-ray photoelectron spectroscopy, electron energy loss spectroscopy, in situ Raman spectroscopy, and in situ electrochemical impedance spectroscopy. In combination, these results indicate that sulfur in UMC exists as linear polymeric sulfur rather than smaller allotropes. The electrochemical reactivity of lithium with sulfur confined in UMC (pore size ≤0.7 nm) is different from that of sulfur confined in microporous carbon (≤2 nm, or ultramicroporous carbon containing significant amount of micropores) and mesoporous carbon (>2 nm). The observed quasi-solid-state reaction of lithium with sulfur in UMC with a single voltage plateau during the discharge/charge process is due to the effective separation of solvent molecules from the active material. The size of carbon pores plays a vital role in determining the reaction path of lithium with sulfur confined in UMC.

INTRODUCTION

Lithium–sulfur (Li–S) batteries have been the most researched electrochemical energy-storage system recently. Sulfur is an attractive positive electrode material for lithium-ion batteries, mainly due to its high specific capacity (1675 mAh g⁻¹), high specific energy (2600 Wh kg⁻¹), and low cost. On the downside, sulfur and its discharge product (Li₂S) are electrical insulators, and the lithiation of sulfur results in a solid/liquid transformation (Li₂S₃/Li₂S) which are soluble in electrolyte. The dissolved polysulfides diffuse to the anode and get reduced at the anode. The solubility of polysulfide gradually reduces the amount of sulfur in the cathode, thereby limiting the cycle life of Li–S batteries. Several strategies have been proposed to overcome these issues related to Li–S batteries, such as: (i) by combining sulfur with various conducting matrices like carbon nanotubes (CNTs), graphene, and metal oxides; (ii) by modifying the electrolyte composition; or by opting to ionic liquid electrolytes, polymer electrolytes, or solid-state electrolytes; and (iii) by modifying the cell configuration.

Ultramicroporous carbon (UMC) exhibits unique advantages for the development of sustainable Li–S batteries and other applications. Since first reports, it has been speculated that UMC confines smaller allotropes of sulfur, thereby eliminating the formation of higher-order lithium polysulfides. Due to space constraints in UMC, the direct contact between sulfur and organic liquid electrolyte solvents is prevented, thereby eliminating the dissolution of active species. This allows the use of carbonate-based solvents in Li–S batteries, which are otherwise considered unsuitable. Despite these advantages, little is known about the state of sulfur confined in UMC and its reactivity toward lithium. Few reports investigated Li reactivity toward sulfur confined in UMC. In our earlier report, we have attempted to investigate the state of sulfur in UMC by transmission electron microscopy (TEM), electron energy loss spectroscopy (EELS), and X-ray photoelectron spectroscopy (XPS) techniques. From XPS analysis, direct transformation of sulfur to lower-order polysulfide Li₂S₃/Li₂S was concluded. However, XPS technique is limited to the surface or subsurface region. Hence,

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the state of sulfur confined in UMC and its reactivity toward Li are still unclear. Combining the results obtained using various experimental techniques, such as XPS, EELS, in situ Raman spectroscopy, and in situ electrochemical impedence spectroscopy (EIS), we here aim at a detailed understanding of the state of sulfur in UMC and its reaction mechanism with Li in carbonate-based electrolytes.

## RESULTS AND DISCUSSION

### Structure and Morphology of UMC and UMC–S Composite

Structure and Morphology of UMC and UMC–S Composite. The UMC–S composite was prepared by the melt-infusion method under vacuum. The synthesis of the UMC host was detailed in our earlier report. UMC and sulfur were mixed at a 1:1 weight ratio, heated at 155 ± 1 °C for 12 h, and cooled to room temperature (RT) under vacuum in a glass oven. Figure 1 shows the N2 adsorption–desorption isotherm and pore-size distribution (inset) of UMC.

![Figure 1. Nitrogen adsorption–desorption isotherm and pore-size distribution (inset) of UMC.](image_url)

isotherm and pore-size distribution (inset) of the UMC host. The N2 adsorption–desorption isotherms of UMC exhibit type I isotherm. The Brunauer–Emmett–Teller (BET) surface area and pore volume are 1600 m² g⁻¹ and 0.66 cm³ g⁻¹, respectively. The density functional theory (DFT) pore-size distribution curve shows exclusively micropores, most of them with a diameter of 0.53 nm. Figure 2a shows XPS spectra recorded in the S 2p region on a UMC.

Nitrogen adsorption

Figure 1 shows the proposed synthesis scheme for the infusion of sulfur into the UMC host material. During heating, sulfur undergoes various changes. Initially, the room-temperature-stable orthorhombic S8 converts to monoclinic sulfur (β-S) at 95.5 °C. On further heating to 119 °C, monoclinic sulfur melts, and at above 150 °C, cyclo-S8 undergoes thermal scission and simultaneously radical polymerization, producing linear, polymeric sulfur chains. Since the glass oven temperature was maintained at 155 °C under reduced pressure, we hypothesize that linear sulfur chains are infused into ultramicropores, driven by capillary forces. The infused linear polymeric sulfur chains tend to align the carbon lattice (graphitization) along the length of the carbon pore. In fact, this conclusion is supported by EELS. EELS spectra of the UMC–S composite recorded at the C-K edge exhibited an increase in π* peak intensity, indicating ordering (graphitization) after sulfur incorporation. Due to size constraints, the infused linear polymeric sulfur maintains its linear form inside UMC even at RT. The operando X-ray absorption spectroscopy (XAS) measurements of UMC–S electrodes showed the formation of higher-order polysulfide (Li2S)n at the very start of discharge. This also indicates that sulfur exists as a linear polymeric chain in UMC and not as smaller allotropes. Sulfur is known to exist as single-walled carbon nanotubes with an inner diameter of 1.1 nm and double-walled carbon nanotubes with an inner diameter of 0.6 nm. It is interesting to note that, although sulfur was infused at 600 °C under vacuum (<1 Pa) into the CNTs, it preferred to exist in linear form at room temperature. (sulfur exists as short chains at 600 °C) From the data of the different methods, we conclude that sulfur, in fact, exists as a linear polymeric chain inside UMC, which has pores predominantly with a diameter of 0.5 nm.

### Electrochemical Performance of UMC–S in Carbonate and Ether Electrolytes

Electrochemical Performance of UMC–S in Carbonate and Ether Electrolytes. Figure 4a shows the discharge/charge voltage profiles of the UMC–S composite cathode with carbonate electrolyte (1.0 M LiPF6 in ethylene carbonate (EC)/dimethyl carbonate (DMC) (1:1 v/v)) cycled at C/20. The first discharge curve consists of two plateaus. The short plateau at 2.38 V (vs Li⁺/Li) is due to the reaction of lithium with the surface functional groups of the carbon host in the composite electrode (it disappeared after the first cycle). Figure 4b shows the first discharge voltage profile for the as-synthesized carbon host (UMC) with no sulfur infiltrated. A similar plateau at 2.38 V (vs Li⁺/Li) is observed in the UMC host due to the reaction of Li with surface functional groups of carbon, which confirms that the plateau observed at 2.38 V corresponds to the reaction of lithium with the carbon host, as it is very similar in the first 100 mAh g⁻¹ discharge capacity.
The second discharge plateau at 1.8 V corresponds to the reaction of sulfur in UMC host with Li. Further cycling of the UMC–S composite cathode resulted in a single discharge and charge plateau at 1.8 and 2.2 V (vs Li+/Li), respectively. A more detailed discussion of the electrochemical characterization results of the UMC–S composite with a carbonate electrolyte was given in our previous report.\textsuperscript{16}

The UMC–S cathode with carbonate electrolyte exhibited a large irreversible capacity loss (ICL) in the first cycle (Figure 4a), which could be due to the reaction of polysulfides expelled to the surface. During the discharge process, sulfur is converted to polysulfides, which results in volume changes. Due to this, a fraction of polysulfide might have been expelled to the surface, which reacts irreversibly with carbonate molecules and reduces the amount of sulfur available for the second cycle. Therefore, a possible way to reduce the ICL in the first cycle would be to decrease the content of sulfur in the UMC to counter the volume change. The reduced amount of sulfur in the composite would reduce the areal capacity, which could be recovered, however, by increasing the thickness of the electrode.

Figure 5 shows the discharge voltage profiles of the UMC–S composite cathode in an ether-based electrolyte (free of LiNO\textsubscript{3} additive). It should be emphasized that the reaction mechanism of the UMC–S electrode with lithium in ether-based electrolytes is slightly different. The linear ether molecules have better access to the sulfur in UMC compared to the cyclic carbonate molecules. Hence, the sulfur accessible to solvent molecules will react similar to the sulfur confined in the mesoporous carbon.\textsuperscript{1} Sulfur that is not reachable to solvent molecules will follow the quasi-solid-state reaction. In the first few cycles, the UMC–S composite features three voltage plateaus during discharge with ether-based electrolyte, in accordance with an earlier report.\textsuperscript{39} The first two plateaus are due to the reduction of sulfur to higher-order polysulfides (Li\textsubscript{2}S\textsubscript{x}, x = 8–4) at 2.4–2.3 V and further to lower-order polysulfides (Li\textsubscript{2}S\textsubscript{y}/Li\textsubscript{2}S) at 2.0–2.1 V. The third plateau at lower potential is assigned to the reduction of sulfur confined in ultramicropores, suggesting a solid-to-solid phase transition. From the sixth cycle, the discharge pattern constitutes only a quasi-solid-state behavior, as observed in the case of carbonate-based electrolyte. During the initial five cycles, the polysulfide might have dissolved in ether molecules and reacted at the
anode similar to that of sulfur confined in mesoporous carbon. A single voltage plateau is attained in discharge and charge profiles (Figures 4a and 5a), although the cells were operated between 3.5 and 1 V (after five cycles in ether-based electrolytes), which is above the solid electrolyte interphase (SEI) formation potential and irrespective of the electrolytes used. This underlines the importance of the carbon structure. Thus, without the necessity for a stable SEI, the size constraints in the UMC host reduced the contact between sulfur and solvent, thereby leading to quasi-solid-state behavior.
X-ray Photoelectron Spectroscopy (XPS) Analysis. Figure 6 compares the XPS spectra in the C 1s and O 1s region recorded on a UMC–S as-prepared electrode and after the first discharge. The C 1s XPS spectra of the as-prepared electrode exhibited three peaks at binding energies (BEs) of 287.0, 289.0, and 291.0 eV corresponding to C−O, C=O, and C−F bonds. During discharge, the peak at 289.0 eV, corresponding to C=O species, lost intensity considerably and almost disappeared due to the reaction with lithium. Consequently, the peak at 287.0 eV, corresponding to C−O, gained intensity after discharge. The peak at 291.0 eV, which corresponds to C−F bonds, is due to the binder poly(vinylidene difluoride) (PVDF). Considering the various plausible SEI components, an increase of the intensity of the peaks related to both C=O and C−O would be expected upon the growth of an SEI layer. Instead, a decrease in C=O intensity is observed upon discharge from 3.5 to 1 V, which can be attributed to the reduction of the corresponding functional groups at the surface of the carbon host by lithium. Therefore, SEI formation was not initiated at this point. This is in line with previous reports that, in the absence of additives in the carbonate-based or ether-based electrolyte, SEI formation is expected to occur only below 0.9 V (vs Li+/Li).

In Situ Raman Spectrum Analysis. To probe the role of carbon (UMC) in transporting lithium to sulfur (as sulfur infused in UMC is isolated from the electrolyte molecules), in situ Raman spectroscopy measurements were performed on a UMC–S cathode at C/20. Here, we used 1.0 M LiPF₆ in EC/DMC (1:1 v/v) as electrolyte and an ECC-Opto-Std (EL-CELL GmbH) electrochemical cell. The voltage range was 1–3.5 V (vs Li⁺/Li). Figure 7 shows the first discharge profile, and the inset shows the evolution of the D and G bands corresponding to the host UMC during discharge. The lithiation of the carbon–sulfur composite was expected to affect the Raman D-band (disordered or defective structures) and G-band (graphitic structures) of the carbon host. During discharge, there was no weakening or broadening of the D-band. However, the position of the G-band was shifted to slightly lower wavenumbers during the initial discharge and remained unchanged until full discharge. Initially, carbon experienced a charge transfer, resulting in the reduction of the surface functional groups, which led to a red shift of the G-band. From the in situ Raman spectroscopy measurements, we conclude that there is no insertion of Li into the carbon (UMC) host in the potential range of 1–3.5 V vs Li⁺/Li and that the role of UMC is similar to that of a carbon-coated cathode material for Li-ion batteries.

In Situ Electrochemical Impedance Spectroscopy (EIS). To get further insight into the reactivity of UMC–S with lithium, in situ electrochemical impedance spectroscopy (EIS) measurements were performed on a UMC–S/Li cell during the first cycle. Impedance spectra were recorded at
various points of discharge and charge, as depicted in Figure 8a (EIS1−EIS7). Initially, impedance spectra were recorded at the open-circuit voltage potential (EIS1). Subsequently, the electrode was galvanostatically discharged for 5 h at a C/20 rate, followed by 1 h of equilibration. Then, the next impedance spectrum was recorded (EIS2). All other impedance spectra were recorded similarly in the frequency range of 200 kHz to 1 mHz, with a voltage amplitude of 5 mV. During post-processing (not shown here), it turned out that the as-recorded impedance spectra suffered from significant noise at frequencies below 0.1 Hz. Thus, the recorded spectra were cut at this frequency. From the Nyquist plot of the preprocessed data (Figure 8b), it is evident that the cell impedance was highest at EIS1. We attribute this high initial impedance to interfacial resistance. After discharging the sample for 5 h, the interfacial resistance was considerably reduced. The significant reduction of the interfacial resistance points toward less resistive products at the surface. Further discharging of the cell gradually reduced the impedance to a minimum at point EIS5, followed by an increase during subsequent charging (EIS7). The gradual decrease and increase of the impedance during the discharge and charge processes, respectively, indicate that the discharge products are less resistive than the charge products.

We have also analyzed the distribution of relaxation times for the measured impedance spectra. This method is well known for the identification of polarization losses in fuel cells and batteries and allows the deconvolution of the losses hidden in the impedance spectrum. Each point in the obtained distribution function represents the fraction of the polarization at a certain characteristic frequency and thus equals the resistance of a single resistor−capacitor circuit with the time constant \( \tau \) in the measurement model shown in Figure 8c. The distribution functions of the preprocessed impedance spectra are shown in Figure 8d. Apparently, there are two processes, \( P_1 \) and \( P_2 \) (represented by two peaks in the distribution), which dominate the impedance response in the measured frequency range. Interestingly, these two processes follow different trends in the course of the first cycle: for \( P_1 \), constant expansion and shift to lower frequencies of the associated peak is observed, whereas both the shift and the dimension of the peak belonging to \( P_2 \) seem to be reversible with discharge and charge of the UMC−S electrode. For insertion materials, the frequency ranges observed for \( P_1 \) and \( P_2 \) (10 Hz to 1 kHz and 1−10 Hz, respectively) have already been identified in the literature and are commonly ascribed to losses from contact resistances (interparticle or at the particle−current collector interface) and faradic resistances associated with the charge-transfer reaction at the electrode−electrolyte interface, respectively. In this context, the observed reversible dependency of \( P_2 \) on the open-circuit potential fits perfectly to the assignment mentioned above because the charge-transfer resistance is well known to depend on the lithiation degree of the insertion compound. The steady increase of the peak associated with process \( P_1 \) points toward a steady increase of the contact resistance, which might be caused by volume changes of the sulfur compound during the discharge and charge processes.

**Overall Electrochemical Reaction Mechanism of Lithium with UMC−S Composite.** In general, the reaction...
of lithium with a carbon–sulfur composite cathode in a Li–S cell using different electrolytes can follow two types of reaction mechanisms, namely, a solid−liquid−solid reaction and/or a quasi-solid-state reaction. Electrochemically, these are reflected by two different types of voltage profiles. This difference in reaction mechanism is mainly due to the difference in the accessibility of solvent to the active component sulfur. If the solvent molecules are accessible to sulfur, then the reaction follows a solid−liquid−solid reaction mechanism; if the solvent access to sulfur is restricted, then the reaction follows a quasi-solid-state reaction mechanism. The former mechanism has been identified in mesoporous carbon (<2 nm) or microporous carbon (>0.7 nm) sulfur composites. However, the quasi-solid-state reaction mechanism is followed even in microporous carbon, provided it is pretreated by cycling at lower potentials to form a stable SEI.32

![Figure 9](https://example.com/figure9.png)

**Figure 9.** Schematic illustration of carbon–sulfur composites with carbon pores of varying size: (a) ultramicropore, (b) micropore, and (c) mesopore, and the related reactivity toward lithium in Li–S batteries.

**Figure 9** illustrates the difference in solvent accessibility by sulfur in carbon pores of varying size (UMC, microporous and mesoporous carbon) in carbon–sulfur composites.

(i) In the case of ultramicropores (<0.7 nm in Figure 9a), solvated ions tend to be desolvated at the surface when the size of the pore is smaller than the size of the solvated ions.7 Thus, ultramicropores can mitigate or prevent the dissolution of active material, as the solvent concentration is very low or nearly zero inside the pore. This results in a quasi-solid-state reaction mechanism, irrespective of the electrolyte used (carbonate- or ether-based).39,52,53

(ii) In the case of microporous (0.7 to ≤2 nm) carbon–sulfur composites, the solvated ions can still have access to the active material (Figure 9b).

(iii) In the case of mesoporous carbon–sulfur composites, the sulfur is even more accessible by the solvent molecules (Figure 9c), leading to the dissolution of active material and intermediate polysulfides. This results in two discharge plateaus in the ether-based electrolyte, while the cells were not cyclable in carbonate-based electrolytes.21

**Origin of the Voltage Plateau.** Several studies on sulfur confined in UMC displayed a single plateau during the discharge/charge process.14,16,39,52–54 This is different from the two-plateau discharge and charge behavior of typical cyclo-S8.55,56 It was speculated that the smaller allotropes (S8 to S4) enable the direct formation of lower-order polysulfides, resulting in a single plateau in the discharge curve. Nevertheless, no evidence was reported to support the existence of smaller allotropes of sulfur in UMC. However, operando XAS on UMC–S electrodes suggests the gradual conversion of higher-order polysulfide (Li2S4) to lower-order polysulfides (Li2S2/Li2S) during the electrochemical discharge.36 In fact, for the first time, the higher-order and lower-order polysulfides were clearly differentiated. The observation of higher-order polysulfides during the initial stage of the discharge suggests that sulfur exists as linear polymeric sulfur rather than as smaller allotropes in the UMC host. Lower-order polysulfides are unlikely to combine and form higher-order polysulfides during discharge. This leads to the question of why a single plateau rather than two plateaus are found in the electrochemical discharge curve. The solid-state conversion potential of S8 to Li2S is 2.24 V vs Li+/Li, considering the free energy change for the formation of Li2S from S8.52 The conversion potential was calculated assuming cyclic S8, and the potential may change depending on the sulfur polymorph.

\[
\frac{1}{8} S_8 + 2Li^+ + 2e^- \leftrightarrow Li_2S, E^o = 2.24 V
\]

This suggests that the solid-state conversion of S8 to Li2S should result in a single plateau below 2.24 V. Smaller allotropes are not required for a single plateau. Indeed, the single plateau was also observed for solid-state Li–S batteries. Hence, the electrochemical reaction of lithium with sulfur infused in UMC could be considered as a solid-state conversion, where the electrolyte molecules have no direct access to the sulfur—a quasi-solid-state reaction. In the case of sulfur confined in mesoporous carbon, where sulfur is accessible to the electrolyte molecules,36 the conversion of S8 to Li2S shows different thermodynamics (due to the dissolved sulfur or polysulfide), similar to iodine redox process, where the redox potential depends on the reaction medium.57,58

**CONCLUSIONS**

Among various sulfur hosts, UMC exhibits unique advantages toward the development of sustainable Li–S batteries, particularly in conventional carbonate-based electrolytes. The effective separation of solvent molecules and sulfur or polysulfide confined in ultramicropore-based electrolytes reduces the side reactions between polysulfide and electrolyte. The elimination of polysulfide in the electrolyte also reduces the side reactions at the anode side. So far, it was postulated that sulfur confined in ultramicropores is present in the form of smaller sulfur allotropes, due to size constraints. However, our experimental results suggest that sulfur in UMC exists as linear polymeric sulfur rather than as smaller allotropes. We also showed that the electrochemical reactivity of lithium with sulfur confined in UMC is different from that of sulfur confined in microporous carbon, ultramicroporous carbon containing a significant amount of micropores, or mesoporous carbon. The size of the carbon pores plays a critical role in determining the reaction path of lithium with sulfur confined in it.

**EXPERIMENTAL SECTION**

Synthesis and Characterization of UMC and UMC–S Composite. Ultramicroporous carbon (UMC) was prepared
by KOH activation of the carbon derived from coconut shells. The coconut shell powder was carbonized at 600 °C for 2 h under Ar gas flow. The carbon obtained was mixed with KOH at a ratio of 1:4 (wt %) and activated under Ar flow. Initially, the mixture was heated to 400 °C and maintained at that temperature for 1 h, and then the temperature was raised to 800 °C and kept for 2 h. The activated carbon obtained was neutralized using dil. HCl and washed with water and dried. N₂ adsorption–desorption isotherms were obtained using a Micromeritics ASAP 2020 analyzer. The specific surface area was calculated using the Brunauer–Emmett–Teller (BET) method, and the pore-size distributions were calculated by the density functional theory (DFT) method. X-ray photoelectron spectroscopy (XPS) measurements were carried out with a PHI 5800 MultiTechnique ESCA System using Al Kα (1486.6 eV) monochromatized radiation. The measurements were done at a detection angle of 45°, using pass energies of 93.9 and 29.35 eV for survey and detail spectra, respectively. For binding energy calibration, the main C (1s) peak was set to 284.8 eV. The sample surfaces were sputtered for 3 and 10 min for survey and detail spectra, respectively. For Raman measurements, on a He–Ne laser with a wavelength of 632.8 nm as the excitation source. TEM and EELS spectra were obtained using a Cs-corrected FEI Titan 80-300 microscope equipped with a GIF Quantum 965 energy filter and operated at 80 kV to avoid knock-on damage and to increase the energy resolution.

Electrochemical Measurements. The UMC–S composite (45.8 wt % of S) and PVDF binder were mixed in a weight ratio of 90:10 using N-methyl-2-pyrrolidinone as a solvent. The slurry obtained was coated on a stainless steel foil and dried at 90 °C overnight. The fabricated electrode contained around 2 mg cm⁻² of sulfur loading. Li foil (Aldrich, 99.9%) was used as the anode. A borosilicate glass fiber sheet was used as the separator. Either 1 M LiPF₆ in 1:1 ethylene carbonate (EC)/dimethyl carbonate (DMC) mixture (LP30, BASF) or 1.0 M LiTFSI in dioctyl (DOL)/dimethoxyethane (DME) (1:1 v/v) mixture was used as the electrolyte. The cell assembly and electrode fabrication were done in an Ar-filled glovebox. Electrochemical studies were performed at RT using Swagelok type cells. The cells were cycled between 1.0 and 3.5 V vs Li⁺/Li at different current densities using an Arbin battery cycling unit BT2000. For EIS measurements, an electrochemical workstation (Bio-Logic) was used. The amplitude was set at 5 mV in the frequency range of 200 kHz to 1 mHz.

In Situ Raman Studies. In situ Raman measurements on UMC–S electrodes were performed using an ECC-Opto-Std (EL-CELL GmbH) electrochemical cell. A thick slurry containing UMC–S and PVDF in the ratio of 90:10 was deposited onto an stainless steel current collector (16 mm) with a 1 mm hole in the middle. The electrodes were dried in an oven at 90 °C for 12 h. In situ Raman cell was assembled using the UMC–S–coated current collectors as a positive electrode, lithium foil as an anode, and a borosilicate glass fiber soaked with 1 M LiPF₆ in 1:1 ethylene carbonate (EC)/dimethyl carbonate (DMC) (LP30, BASF) as separator and electrolyte. The assembled cell was connected to an IM6 electrochemical system (ZAHNER-Elektrik GmbH) for galvanostatic cycling. In situ Raman cell was discharged and charged between 1.0 and 3.5 V vs Li⁺/Li at a rate of C/20. Raman spectra were collected at RT using a confocal Raman microscope (InVia, Renishaw) in the spectral range of 1000–2500 cm⁻¹ using a He–Ne laser with a wavelength of 632.8 nm. The laser power was set to ≈1.0 mW, and the spectrum acquisition time was 20 s.

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Notes
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