Critical Role of Functional Groups Containing N, S, and O on Graphene Surface for Stable and Fast Charging Li-S Batteries

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1. Introduction

New electrochemical energy storage systems are urgently needed to meet the ever-growing needs of the portable electronics market. Among emerging energy storage technologies, lithium-sulfur (Li-S) batteries are considered the best candidate to replace commercial lithium-ion (Li-ion) batteries (LiBs). Despite its higher theoretical energy density (2600 Wh kg\(^{-1}\)), abundance, low cost, and non-toxicity, sulfur has been restricted by concerns related to the lithium/sulfur reaction.\(^{[3,4]}\) Sulfur and its end-discharge products, that is, Li\(_2\)S\(_2\)/Li\(_2\)S, are intrinsic insulators and poorly soluble in the battery electrolyte.\(^{[5]}\) By contrast, the intermediate discharge products such as Li-polysulfides (LiPs) are soluble in the electrolyte, enabling more efficient use of...
the active material, but with possible migration to the surface of the Li anode and partially back to the cathode. This is called the "shuttle effect," which causes the loss of active material and subsequent reduction of battery cycle life.[6,7]

In the last decades, many studies have attempted to integrate sulfur into carbon-based nanostructures to address the problems of low electronic conductivity and shuttle effect.[8–12] The specific capacity of Li-S was increased in the first few cycles because of the improved conductivity obtained by integrating sulfur into the carbon nanostructure. However, the nonpolar surface of carbons was not successful in retaining the dissolution of polar LiPs upon long cycling time.[9] Although S8 is nonpolar and can be adsorbed relatively well by carbon materials, Li2S and LiPs are polar and adsorb poorly in hydrophobic hosts.[13] Furthermore, the inner confinement of sulfur in the carbon matrix limited the diffusion of Li+ to a small portion of active material.[14]

Free-standing porous carbon hosts have been proposed to exploit a larger portion of the active material at the sulfur cathode, increase the conductivity of the sulfur cathode, and avoid the use of non-conductive binders (e.g., polyvinylidene fluoride (PVDF)), which modifies the porous structure of carbon, which is essential for enduring volume changes (~80%) upon cycling.[15–17] The hydrophobic surface of carbon should be modified to improve its interaction with soluble LiPs. Metals or metal oxides are polar groups that can have a strong chemical interaction with LiPs.[18,19] Although S8 is nonpolar and can be adsorbed relatively well by carbon materials, Li2S and LiPs are polar and adsorb poorly in hydrophobic hosts.[13] Furthermore, the inner confinement of sulfur in the carbon matrix limited the diffusion of Li+ to a small portion of active material.[14]

Direct grafting of small and polar functional groups on the surface of the host would be more favorable for building the Li-S cell at the practical energy density level. This method permits the diversification of the organic functionalities attached with no large modification of the host conductivity, structure, and porosity.[20,21] In the next step, the size of the functional groups can be reduced and simplified to a single atom similar to doping of carbon materials. Previous studies showed the benefits of heteroatom doping on carbon, in particular using N atom, leading to strong interactions with LiPs.[14] Nevertheless, there are only a limited number of elements (e.g., N, S, and O) that can be included as dopants into carbon materials, and it is still a challenge to reach a high doping content (typically doping is <10%), which guarantees the presence of the dopant on the carbon surface.[22–25] By taking advantage of the tunable chemistry and variety of potential functional groups, we can instead tune the surface properties of carbon materials by covalently grafting the most suitable one to their surface. Thus, understanding the interactions of LiPs with a specific functional group at the molecular level is important for selecting suitable molecules that can improve both the energy density and cycling stability of Li-S batteries.

In this study, the performance of Li-S batteries was enhanced by directly functionalizing the surface of the graphene-based substrate with versatile molecules. A benzene ring with variable functional groups was covalently grafted on the surface of expanded reduced graphene oxide (eRGO) using diazonium chemistry. Nitrobenzene and benzene sulfonate were selected as model molecules to understand their interaction with LiPs by combining electrochemical tests and density functional theory (DFT) calculations at the molecular level. The results demonstrate how the presence of vertical nitro- or sulfonate benzene molecules (~1 nm high) on the surface of graphene facilitates the binding with LiPs and thus stabilizes the performance of Li-S cells reaching practical energy and power densities.

2. Results and Discussion

2.1. Modification of Surface Properties of Expanded Reduced Graphene Oxide Host

Graphene is a highly conductive nanomaterial, which has been extensively reported as a conductive additive and support for the preparation of electronic compounds.[26] In this study, eRGO was used as a conductive substrate to investigate the role of specific functional groups in the electrochemical performance of Li-S batteries. The nitrobenzene and benzene sulfonate were grafted on the eRGO surface by diazonium chemistry. We used 4-nitrobenzene diazonium tetrafluoroborate and 4-sulfonic acid phenyl diazonium tetrafluoroborate to obtain nitrobenzene functionalized eRGO (NG) and benzene sulfonate functionalized eRGO (SG), respectively (Figure 1a). The porous structure of eRGO with the 3D network was preserved after functionalization (Figure S1, Supporting Information). Among potential functionalities, we investigated the effect of the nitro group and sulfonate group on the performance of Li-S cells because 1) nitrogen doping has been demonstrated to effectively enhance the interactions between carbon and sulfur upon battery cycling; and 2) sulfonate group has an affinity for LiPs species formed during the electrochemical process. Further details related to the grafting and synthesis procedure are presented in the experimental section.

To confirm the success of the covalent functionalization of graphene, surface chemical compositions of synthesized NG and SG together with the precursor eRGO were characterized by X-ray photoelectron spectroscopy (XPS) (Figure 1b). No traces of N 1s were observed in the eRGO sample, while two peaks at 406 and 400.5 eV appeared in NG (middle graph of Figure 1b). The main N 1s peak at 406 eV was assigned to nitrogen functional groups in the electrochemical performance of Li-S battery; and 2) sulfonate group has an affinity for LiPs species formed during the electrochemical process. Further details related to the grafting and synthesis procedure are presented in the experimental section.
high reactivity of aryl radicals generated from the diazonium salt.\[28\] The XPS analysis at the S 2p core level for the SG sample also showed the presence of an intense peak at 168 eV (bottom of Figure 1b) (4.6% in atomic content), demonstrating the covalent grafting of sulfonate benzene group on the pristine sample (1 benzene sulfonate group per 9C atoms of eRGO).\[29\]

The loading of nitrobenzene and benzene sulfonic acid functional groups on NG and SG was calculated using thermogravimetric analysis (TGA) (Figure 1c). No obvious weight loss was observed at 150–300 °C for NG, SG, and eRGO. This is a typical temperature range where graphene oxide (GO) loses oxygen-containing functional groups. The lack of any peak in TGA indicates that eRGO had lower defectivity of GO.\[30\]

Functionalized NG and SG samples showed a larger weight loss (≈10%) than eRGO during TGA between 350 and 600 °C due to the decomposition of nitrobenzene and benzene sulfonic acid functional groups from NG and SG, respectively. A similar functionalization degree obtained with both molecules indicates that the nitro or sulfonate group has a minor effect on the grafting process, which is instead governed by the diazonium tetrafluoroborate moiety present in both molecules. Weight loss in all three samples above 600 °C was caused by the gradual decomposition of the eRGO defective graphene structure. The uniform distribution of both nitrobenzene and benzene sulfonate functional groups on the surface of graphene was also confirmed by energy dispersive spectroscopy (EDS) mapping (Figures S2–S4, Supporting Information).

The surface area and porous structure are important factors that can affect not only the sulfur loading, but also the final energy density; thus, we investigated whether grafting of both nitrobenzene and benzene sulfonate groups on the eRGO substrate would affect the surface area and porosity of the substrate.\[31\] We characterized eRGO, NG, and SG using Brunauer–Emmett–Teller (BET) analysis (Figure S5, Supporting Information), which yielded specific surface areas of 430, 170, and 180 m² g⁻¹, respectively. The smaller surface area was attributed to the presence of functional groups and to the collapse of macropores during solution processing. A large surface area can be beneficial in Li-S cells to increase the interaction of the electrolyte with active materials, while a small surface area enables reduction of the volume of the electrolyte required for wetting the electrode with benefit for the galvanostatic cycling performance of Li-S cells at a practical level.

2.2. Li/S Cells: Cycling and Rate Performance

Our approach aims in designing sulfur electrodes at a practical level. Thus, instead of infiltrating elemental sulfur (S₈) which is
complicated and require the use of thermal treatment, we infiltrated a fixed amount of Li_2S_n polysulfides in the eRGO, NG, and SG samples by simple solvent casting technique, to achieve a high sulfur loading (i.e., 4.5 mg cm\(^{-2}\)) in the cathode and low sulfur to electrolyte (S/E) ratio (i.e., 1:5) in the Li-S cell. We then performed galvanostatic tests of Li-S cells at the C/10 current rate (Figure 2). During the first cycle when the Li-S cells were assembled by using eRGO, we discharged/charged NG and SG at a relatively low current, that is, C/20, to convert Li_2S_\(n\) into Li_2S and further facilitate the recovery back to S_8 during charging. At the first cycle, the Li/S cell built with pristine eRGO showed a Coulombic efficiency of 86% while the cells made by NG and SG revealed higher Coulombic efficiencies of 92 and 93%, respectively (Figure S6, Supporting Information). This difference was due to the larger conversion of Li_2S_\(n\) present in the functionalized eRGO into Li_2S.

All Li-S cells were then cycled at the C/10 current rate. The one using eRGO as the sulfur host delivered an initial specific capacity of \(\approx 900\) mAh g\(^{-1}\), which decreased to lower values upon cycling, reaching 450 mAh g\(^{-1}\) at cycle 250 (Figure 2a). The cell-based on the NG host delivered a higher initial specific capacity of 1200 mAh g\(^{-1}\) decreasing to \(\approx 750\) mAh g\(^{-1}\) at cycle 250 (Figure 2b). The Li/S cell using the SG electrode exhibited stability similar to that using NG but with a slightly different behavior during the first few cycles where the capacity quickly dropped and stabilized to values ranging between 800 and 700 mAh g\(^{-1}\) from cycle 100 to 250 (Figure 2c). It has to be highlighted that no one of the reported Li/S cells could reach a delivered capacity as close as to the theoretical value of Li/S, that is, 1675 mAh g\(^{-1}\). That is mostly related to the fact that the Li/S cells are built using low sulfur to electrolyte ratio (1:5), which causes lower amount of active material reacting with Li, with consequent value of delivered specific capacity lower than the theoretical.\[15,17,32\]

The effect of NG and SG groups was even more apparent when rate capability tests were used. We started from a current density of C/10 and subsequently increased the rate in four steps up to 3C. The capacity of the Li/S cell using the eRGO electrode decayed quickly especially at high current rates, reaching \(\approx 150\) mAh g\(^{-1}\) at 3C (Figure 2d,g). The voltage/capacity profile (Figure 2d) shows a large increase in voltage polarization.

A final test at the C/10 rate showed a capacity of 750 mAh g\(^{-1}\), less than 70% of the initial one, indicating irreversible damage to the device. In contrast, the two Li-S cells using the functionalized graphene showed capacity values in the range of battery applications at high C rates. Indeed, the NG and SG electrodes delivered a capacity of 500–450 mAh g\(^{-1}\) even at 3C, (Figure 2e,f).

Note that the charging time at 3C is 20 min, which is important for the application in the automotive industry. Moreover, when the current density was switched back to the initial value of C/10, both electrodes recovered almost 95% of the capacity delivered at cycle 5, further demonstrating high cycling stability...
at high C rates (Figure 2h,i). The improved performance of NG and SG as compared to eRGO was attributed to the presence of nitro and sulfonate functional groups on the surface of graphene, fostering stronger interaction with LiPs and faster Li diffusion to the electrode, improving the utilization of the active material at higher C rates, even for prolonged cycles at 2C, see Figure S7, Supporting Information.

2.3. Binding Interactions Between Functionalized Graphene and Li-Polysulfides Polar Species

The experiments showed that the presence of N- and S-based functional groups improved the stability and cyclability of the material, which could be attributed to the strong polar–polar interaction between LiPs and vertical nitrobenzene and benzene sulfonate functional groups. To confirm this assumption and explain the differences observed in the experiments, the interactions of sulfur and LiPs with eRGO, NG, and SG hosts were evaluated via DFT calculations. The unmodified carbon surface was modeled with a C_{54}H_{20} system consisting of 18 aromatic rings. The details of the bonding motifs, methods, and models are presented in the experimental section. We studied the BEs of eRGO, NG, and SG with LiPs with different Li:S ratios. The different chemical species present in the device during charging/discharging, from S_8 to different Li_2S_n with 1 ≤ n ≤ 8 were investigated systematically (Figure 3).

DFT calculations demonstrated that the NG and SG groups are...
oriented vertically on the graphene surface and can interact with polysulfides. Although the eRGO substrate is expected to have the highest BE with S\textsubscript{8} because both eRGO and S\textsubscript{8} are nonpolar species, eRGO and SG showed comparable binding with S\textsubscript{8} (1.23 and 1.27 eV, respectively), while BE of S\textsubscript{8} with NG was slightly lower (1.14 eV) (Figure 3). DFT showed that this is related to the weak interaction between sulfur and OH groups, which results in lower energy achieved for the NG system.\[34\]

The BE of interaction of Li\textsubscript{2}S\textsubscript{8} with NG (= 1.72 eV) and SG (= 1.46 eV) were significantly larger than the BE with eRGO substrate (= 1.25 eV), (Figure 3a–c). The difference was more striking for interaction with Li\textsubscript{2}S\textsubscript{n} having higher Li content (n = 6, 4, 2, and 1) and thus higher polarity. Indeed, eRGO showed weaker interactions with polar Li\textsubscript{2}S\textsubscript{n} species (40–50% lower BEs weaker than Li\textsubscript{2}S\textsubscript{8}) (Figure 3a). Instead, NG and SG highlighted stronger interactions, suggesting that the functional groups may improve the stability of such species on the substrate surface. The added functional groups actively participate in the immobilization of Li\textsubscript{2}S\textsubscript{n} species with the coordination of lithium cation by strongly electronegative atom(s); main interactions were with two oxygen atoms from the SO\textsubscript{3} group in the SG sample, and with the nitrogen atom of NO\textsubscript{2} in the NG sample, respectively. Therefore, a larger fraction of sulfur can be electrochemically active during the redox reaction, avoiding undesired LiPs “shuttling” to the anode surface, which agrees with the better response to the galvanostatic cycling shown in Figure 2. The Li\textsubscript{2}S\textsubscript{n} adsorption of the carbon materials has been also tested visually by soaking 1 g of eRGO, NG, and SG in 10 mL of 1m Li\textsubscript{2}S\textsubscript{8} dimethoxyethane (DME) solution, Figure S8, Supporting Information. The Figure highlights how the NG and SG carbons are able to fully adsorb the polysulfide contained in the solution, while the bottle containing the eRGO still evidences the presence of polysulfide species in the solution. The binding of polysulfides to functionalized eRGO was further analyzed using the Fukui functions, (Figure S9, Supporting Information). The Figure highlights interactions of Li\textsubscript{2}S\textsubscript{x} polysulfide mostly with functional groups (i.e., -NO\textsubscript{2} and SO\textsubscript{3}), rather than graphene layer, in the case of the functionalized carbons. Increases in electron density at Li-bonding oxygen atoms is observed, together with the decrease of electron density at the aromatic ring, indicating a shift of electron density within the functional group attached to the carbon.

2.4. Specific Energy and Power Densities at Practical Level

Figure 4 shows the specific energy and power density delivered at different current rates. Differently from previous studies, we calculated all the values considering the total weight of the built Li-S cells produced, including electrodes, electrolytes, separators, and current collectors.\[35\] Energy density and power density are often calculated by considering only part of the cell components, which generates higher apparent density. We think that it is more appropriate to use the total weight of the
device based on Table S1, Supporting Information. Figure 4a shows the Ragone plots of the eRGO electrode; specific power and energy densities of eRGO are worse than those of commercial Li-ion systems both at lower and higher current rates.\(^{[16]}\)

By contrast, Li-S cells made by NG and SG carbon hosts show a Ragone plot with enhanced specific power and energy densities (Figure 4b,c). The performance of both cells is comparable to that of Li-metal cells at low C rates\(^{[37]}\) and better than that of Li-metal cells at high C rates.\(^{[38]}\)

To understand whether this improvement was caused by an improvement of reaction kinetics, we carried out cyclic voltammetry (CV) tests at different scan rates (Figure S10, Supporting Information). The current values for different redox peaks (A, B, and C in CV tests) were plotted as a function of the square root of the scan rate (Figure 4d–f). Then, a linear fitting was applied and the Randles–Sevick equation was used to calculate the Li\(^+\) diffusion coefficient (\(D_{\text{Li}^+}\)) at different stages of discharge/charge.\(^{[39]}\) According to the Randles–Sevick equation, the \(D_{\text{Li}^+}\) values at the cathodic (A and B) and anodic (C) peaks of the Li-S cell using the pristine eRGO electrode were of the order of \(10^{-9}\) cm\(^2\) s\(^{-1}\). When employing the NG and SG electrodes, the calculated \(D_{\text{Li}^+}\) values increased by one order of magnitude relative to pristine eRGO (Table S2, Supporting Information), showing sharper redox peaks and narrower peak separation (Figure S10, Supporting Information). This indicates that the better battery performance observed in NG and SG (as compared to eRGO) can be attributed to improved redox kinetics and higher Li\(^+\) diffusion coefficients.

2.5. Electrochemical Mechanism of LiPs on Functionalized Graphene Host

To gain more insight into electrochemical stability, XPS was employed to understand the evolution of different chemical species on the graphene surface. Figure 5a shows the voltage profile during the first discharge cycle, with red, and black dots marking the beginning and end of the cycle, respectively, indicating the state where the XPS analysis was performed for all samples.

In the S2ps spectrum, the eRGO electrode showed two main peaks at 159.9 and 162.2 eV, corresponding to the long-, medium-, and short-chain polysulfide species deposited on its surface at the pristine state (Figure 5b left). Similar peaks appeared in the NG electrode (Figure 5c left), while SG showed additional peaks at 159.5 eV owing to the formation of additional Li\(_2\)S\(_n\) species and a peak at 163.2 eV related to the presence of thiosulfate groups on the graphene surface.\(^{[40]}\)

Thus, Li\(_2\)S\(_n\) deposited during the synthesis phase was distributed at different chain lengths as a result of a disproportionation reaction that occurred during the formation of Li\(_2\)S\(_8\) in the DME solvent. After the Li-S cell was fully discharged, the pristine eRGO electrode revealed a shift of both aforementioned peaks to different BEs (one peak from 159.9 to 160.1 eV, and the second from 162.2 to 161.7 eV), confirming the conversion of Li\(_2\)S\(_n\) to Li\(_2\)S\(_2\) and Li\(_2\)S (Figure 5b right). Besides this, the NG and SG samples also showed additional peak at 167.0 eV, related to the formation of Li\(_2\)S-(S)O\(_3\) (Figure 5c right and Figure 5d right).

The S–O bonds on the surface of both NG and SG electrodes can prevent, in part, the migration of soluble Li\(_2\)S\(_n\) to the Li-metal surface, especially during prolonged cycling, thus keeping the active material close to the cathode side.\(^{[40]}\)

The NG electrode, which has the larger formation of Li\(_2\)S-(S)O\(_3\), showed better stability upon cycling (Figure 2b), while the SG electrode, with a lower concentration of Li\(_2\)S-(S)O\(_3\), showed slightly worse cycling stability (Figure 2c). The pristine eRGO electrode, where the Li\(_2\)S-(S)O\(_3\) on the surface was not formed, was unstable, decreasing the stability upon battery cycling. The formation of S–O bonds upon discharge was further confirmed for NG and SG electrodes from the XPS at the O 1s level (Figure S11, Supporting Information), where an additional peak appeared at 527.7 eV, while the same peak was negligible in the eRGO electrode. Overall, the XPS data given are in good agreement and provide direct evidence of what is already suggested by galvanostatic, DFT, and CV results. Furthermore, the use of functionalities at the cathodic side brings to better stability of the Li anode. SEM images of the Li recovered after 250 cycles of discharge/charge at 0.1C, Figure S12, Supporting Information, clearly evidence how the surface of Li-metal is less corroded when NG and SG electrodes are employed at the cathode side, while heavy corrosion and large holes are present at the surface of the Li recovered from a Li/S cell using pristine eRGO.

3. Conclusions

A new pathway to improve the specific energy and power densities of Li/S cells was proposed as an alternative to doping by introducing versatile functional groups (e.g., nitrobenzene and benzene sulfonate) on the surface of highly conductive graphene-based substrates. We demonstrated that the presence of N, S, and O on the surface of an eRGO-based substrate increased the interaction between a wide range of Li\(_2\)S\(_n\) species and the surface of the electrode. The presence of nitrobenzene and benzene sulfonate functional groups suggests that 1) the increased BEs favor the interaction of sulfur, polysulfides, and sulfide species with the surface of the electrode, and 2) the faster Li diffusion on the electrode surface promotes the reaction between Li and S and brings them closer to the surface of the host, even at high current rates. These claims were confirmed by experimental observation of better cycling stability, faster charging/discharging at a current rate as high as 3C, and by higher Li\(^+\) diffusion on functionalized electrodes. By using a practical sulfur loading of 4.5 mg cm\(^{-2}\) with an electrolyte volume of only five times this weight (S/E 1:5), the Li-S cells based on NG and SG delivered capacity, energy, and power densities comparable to commercial LiBs, outperforming them at high current rates. The functionalization strategy of graphene-based sulfur host proposed in this study offers a new concept for designing practical sulfur electrodes with enhanced cycling performance.

4. Experimental Section

Thermally Expanded Reduced Graphene Oxide: Commercial GO from Graphenea was used to prepare thermally eRGO.\(^{[41]}\) The GO powder in a glass container was heated to 320 °C for 3 min. The GO powder was expanded and exfoliated by the removal of oxygen-containing functional
groups on the surface of GO. Apart from the eRGO that was synthesized in the laboratory, the commercially high surface area reduced GO, which was prepared similarly from the graphene supermarket, was also used as a starting material to synthesize NG and SG.

**Synthesis of 4-Sulfonic Acid Phenyl Diazonium Tetrafluoroborate and 4-Nitrobenzene Diazonium Tetrafluoroborate**: The two diazonium salts were synthesized according to the previously reported procedures. First, 6 mL of tetrafluoroboric acid (48% in water) was added to 50 mL of 4-sulfamic acid suspension (6 g); after stirring for 20 min, the mixture was cooled to 0 °C in an ice bath. Saturated sodium nitrite (2 g) in water was then slowly added to the mixture under stirring. After 2 h reaction, the white precipitate (4-sulfonic acid phenyl diazonium tetrafluoroborate) was collected and washed with cold water. The 4-nitrobenzene diazonium tetrafluoroborate was synthesized using the same procedure, but 98% 4-nitroaniline was used as a precursor instead of 4-sulfamic acid.

**Synthesis of NG and SG**: The as-prepared eRGO (300 mg) was first sonicated in 150 mL, 2 mg mL\(^{-1}\) sodium dodecyl sulfate aqueous solution to allow the surfactant to penetrate the pores of eRGO. Then, 100 mg of 4-sulfonic acid phenyl diazonium tetrafluoroborate was added

![Figure 5](image-url). Chemical evolution of LiPs on different graphene-based hosts during battery discharge. a) Galvanostatic voltage profile measured on NG sample during the 1st discharge cycle at a current of C/40 (1C = 1675 mA g\(^{-1}\)). XPS spectrum at the S2p core level for the b) eRGO, c) NG, and d) SG at the pristine state (red spot); after full discharge (black spot).
stepwise to eRGO. After stirring at room temperature in the dark for 3 h, the black powder (SG) was filtrated and thoroughly washed with water and ethanol. The NC was obtained following the same procedure but by using 4-nitrobenzene diazonium tetrafluoroborate instead of 4-sulfonic acid phenyl diazonium tetrafluoroborate.[33]

Characterization: XPS spectra were collected using PHI VersaProbe III with a monochromatic Al Ka source (1486.6 eV); the spot size was 100 μm. The discharged samples were transferred directly from the glove box to the XPS chamber without exposing them to the air. The Multipak Spectrum was used for the analysis of the spectra. The BEs were calibrated using the C 1s peak at 285 eV. Surface area, pore volume, and pore size distribution were measured by nitrogen-sorption using Micromeritics TriStar 3000. Unfunctionalized sample was degassed for a minimum of 12 h at 120 °C under a N2 flow. SG and NG samples were degassed for a minimum of 16 h at 120 °C under a N2 flow. The surface area was calculated using the BET algorithm. Scanning electron microscopy was performed using a JEOL JSM-7800F Prime microscope equipped with EDS. TGA was performed using a Mettler Toledo TGA/DSC3+ System at a heating rate of 5 °C min⁻¹ under Ar.

Density Functional Theory Calculations: The pristine carbon surface was simulated using a C₆H₆ model, consisting of 18 aromatic rings. The interactions of the resulting surfaces with different sulfur and (poly-) sulfide species (S₂, Li₂S, Li₂S₂, Li₂S₃, Li₂S₄, and Li₂S₅) were studied by testing at least five different starting geometries for each, prior to geometry optimization to determine the most favorable geometries. To reduce the computation time, all structures were pre-optimized using a smaller basis set (6-31G(d)) prior to the final geometry optimization using 6–31+G(d,p). The M06-2X functional, as implemented in Gaussian16 B.01, was used and shown to be appropriate for studies of graphene and carbon nanotubes.[34] The BEs were obtained by subtracting the total energy of the carbon layer-Li₂S system from the sum of the isolated carbon layer and Li₅S₆ species optimized separately. To reduce error related to the difference in the basis set size with or without the carbon layer, the basis set superposition error correction was considered by calculating the optimized geometries of separated species in the basis set of the entire system, and the error was determined to be around 0.1 eV.

Electrolyte and Electrodes Preparation: The electrolyte was prepared by dissolving 0.4 mol LiNO₃ (Aldrich) and 1 mol of LiTFSI (Solvionic, France) in 1 L of a 1:1 volume ratio mixture of Dioxolane (DOL, Aldrich)/DME (Aldrich). Then, 1 mL Li₂S₅ solution was prepared by dissolving 2 mol of Li (Chemetall, thickness of 250 μm) and 8 mol of sulfur (Aldrich, 99.9%) in 1 L of DME (Aldrich, 99.9%) at 80 °C. The prepared solution was stirred for about 4 h until a red-colored solution was obtained. The film electrodes were prepared on carbon-coated Al foil (Aldrich) using three powders: eRGO, NG, and SG. Seventy percent of active material (eRGO, NG, SG) was mixed with 10% PVDF (Arkema) and 20% multiwalled carbon nanotubes (MWCNTs, Aldrich) (final weight ratio of 70:10:20) and drop-casted on an Al foil (4.2 mg cm⁻²) using a doctor blade (film casting knife, MTI, EQ-se-KTQ-150D). The resulting 40 μm thick films were cut into disks and dried at 80 °C under vacuum to remove the residual solvent prior to the assembly of Li-S cells. The electrode had a final weight of 6.42 mg cm⁻², of which the PVDF-carbon loading was 2.22 mg cm⁻² (MWCNTs and corresponding carbon hosts: eRGO, NG, or SG). The 1 mL Li₂S₅ DME solution was subsequently drop-casted on the surface of eRGO, NG, and SG electrodes at 80 °C to obtain a final Li₂S₅ loading of 4.71 mg cm⁻² (sulfur = 4.47 mg cm⁻²). The total electrode weight was 11.15 mg cm⁻².

Electrochemical Characteristics: The as-prepared electrodes were used in coin cell configurations together with the Celgard separator (12 mm), which was soaked with 20 μL cm⁻² of electrolytes (sulfur:to-electrolyte ratio (S/E) 1:5). The cells were cycled at 1.8–2.8 V using a current density of 167 mA g⁻¹ = 0.1 C (1 C = 1675 mA g⁻¹) for prolonged cycling tests, and 0.1, 0.5, 1, 2, and 3C for rate capability tests. All the cycling tests were performed using a Scribner 585 battery test system. CV measurements were performed using a potentiostat/galvanostat VSP (BioLogic, France) instrument in the voltage range of 1.6–2.8 V at a scan rate of 0.3, 0.4, and 0.5 mV s⁻¹. For detecting the diffusion coefficient of the different carbon materials, the current values at different redox peaks from CV analysis were plotted as a function of the scan rate0.5, following a linear fitting as shown in Figure 4d–f. The Li-ion diffusion coefficient was calculated using the Randles–Sevick equation: D Li = 268 600 × e−1.5 × area × D Li × conc Li × rate0.5, where D Li is the current of the peak detected from CV analysis, e is the number of the electron involved in the electrochemical process, area is the cathode area, conc Li is the Li-ion concentration in the electrolyte, and rate is the CV scan rate.

Supporting Information
Supporting Information is available from the Wiley Online Library or from the author.

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Conflict of Interest
The authors declare no conflict of interest.

Data Availability Statement
Research data are not shared.

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