ABSTRACT: Graphene has received much attention as a supercapacitor electrode material due to its chemical inertness in preventing reaction with electrolytes and the large surface area due to its two-dimensional nature. However, when graphene sheets are processed into electrodes, they tend to stack together and form a turbostratic graphite material with a much reduced surface area relative to the total surface area of individual graphene sheets. Separately, electrochemical exfoliation of graphite is one method of producing single-layer graphene, which is often used to produce graphene for supercapacitor electrodes, although such exfoliated graphene still leads to reduced surface areas due to stacking during electrode fabrication. To utilize the large surface area of graphene, graphene must be exfoliated in situ within a supercapacitor device after the device fabrication. However, graphitic electrodes are typically destroyed upon exfoliation, which is largely due to the loss of electrical connectivity among small exfoliated graphene flakes. Here, we report successful in situ exfoliation of graphene nanostripes, a type of quasi-one-dimensional graphene nanomaterial with large length-to-width aspect ratios, as the anode material in supercapacitors. We find that the in situ exfoliation leads to over 400% enhancement in capacitance as the result of retaining the electrical connectivity among exfoliated quasi-one-dimensional graphene nanostripes in addition to increasing the total surface area, paving ways to fully realizing the benefit of graphene electrodes in supercapacitor applications.

INTRODUCTION

Extensively studied active materials for supercapacitors include pure graphene,1 heteroatom-doped graphene,2,3 and graphene composites (e.g., with metal oxide nanoparticles).4 Graphene is attractive as a supercapacitor electrode material due to its large surface area as a two-dimensional material and its high carrier mobility and quantum capacitance.5,6 An additional advantage is that graphene, as well as other carbon-based nanomaterials, can be deposited on surfaces in precise configurations to make interesting devices such as microsupercapacitors.7 However, processing of graphene sheets into supercapacitor electrodes typically leads to stacking of graphene sheets,8 reducing the total surface area and decreasing the carrier mobility9 such that the advantages of graphene are often not realized in practical supercapacitor devices. Therefore, it is desirable to develop methods to obtain single-layer graphene in practical supercapacitor devices. Indeed, many reports have focused on developing methods to increase the surface area of nanocarbons.10

Methods of producing single-layer graphene in high yield include oxidative unzipping of carbon nanotubes,11 oxidative chemical exfoliation,12 liquid-phase exfoliation,13 and electrochemical exfoliation.14 Electrochemical exfoliation is the process of separating graphite or multilayer graphene into single-layer graphene sheets by applying a voltage to intercalate large ions, separating graphite sheets to single-layer graphene. This has been done in organic,14 aqueous,15 and ionic liquid16 solutions and has been used to produce graphene for field-effect transistors,17 energy storage,18 transparent conductive electrodes,19 and gas sensors.20 However, if single-layer graphene is desired for these applications, researchers should be cognizant that packaging the exfoliated graphene into a device typically results in stacking of graphene sheets.21 Of these, we find electrochemical exfoliation particularly interesting because it can potentially be applied in situ in a packaged supercapacitor electrode.

Although in situ exfoliation of graphitic electrodes typically catastrophically degrades the electrode performance due to loss of electrical contact as individual graphene sheets separate,21−23 we demonstrate in this contribution that by employing quasi-one-dimensional graphene nanostripes
GNSPs as the active electrode material, the electrode supercapacitor performance can be enhanced upon in situ exfoliation due to the electrical conductivity maintained by the percolating nature of one-dimensional materials. These concepts, i.e., electrical percolation by quasi-one-dimensional materials and in situ electrochemical modification of electrodes, have actually been explored in other materials but have not been simultaneously studied in carbon nanomaterials to the best of our knowledge. GNSPs are a graphene nanomaterial previously developed in our group that demonstrate chemical purity, good crystallinity, high carrier mobility, and quasi-one-dimensionality (e.g., GNSP dimensions are $\sim 400 \text{ nm} \times 60 \text{ \mu m}$). In our study of in situ exfoliated GNSPs, we employ X-ray diffraction (XRD) to study structural changes in the material and electrochemical impedance spectroscopy (EIS) to study the behavior of the electrode at various degrees of exfoliation and conjecture that the improved capacitance is due to an increase in surface area while maintaining the electrical connectivity in the electrode.

**RESULTS**

The quasi-one-dimensional nature of GNSPs is demonstrated in Figure 1. Figure 1a is a normal-incidence scanning electron microscopy (SEM) image of GNSPs on the growth substrate without further modification. Here, the one-dimensional nature is not apparent, but the fabricated material can be seen as a vertically oriented graphene material, where graphene sheets grow vertically with respect to the growth substrate and form a dense interconnected network. Detailed reviews of vertically oriented graphene can be found in refs 28, 29. When dispersed onto a substrate (Figure 1b), our GNSPs, which are a particular class of vertically oriented graphene nanomaterial, become apparently quasi-one-dimensional with large length-to-width aspect ratios. Several GNSPs are visible in Figure 1b, but a wider area image (Figure 1c) shows that the GNSP dispersion contains a large amount of GNSPs (highlighted in Figure 1c) with varying dimensions. An analysis of 177 individual GNSPs reveals that the average width and length of GNSPs are $\sim 450 \text{ nm}$ and $\sim 12 \text{ \mu m}$, respectively. A histogram of analyzed GNSPs aspect ratios is provided in Figure 1d, where the average aspect ratio is $\sim 34:1$, and the most frequent aspect ratio is $\sim 10:1$. (Note: these GNSPs do not exhibit quantum confinement effects as their typical width is $\sim 450 \text{ nm}$ while quantum confinement effects begin at widths below $\sim 40 \text{ nm}$.) The Brunauer–Emmett–Teller (BET) surface area was $64 \text{ m}^2/\text{g}$.

The successful fabrication of a graphene material was confirmed by Raman spectroscopy and X-ray photoelectron spectroscopy (XPS). The Raman spectrum is presented in Figure 2a. The D (1361 cm$^{-1}$), G (1589 cm$^{-1}$), and 2D (2704 cm$^{-1}$) peaks are characteristic of graphene, and the D’ (1609 cm$^{-1}$) and D + D’ (2945 cm$^{-1}$) peaks together with the G and D’ peaks being well resolved classify the sample as a crystalline graphene nanomaterial according to the three-stage defect model. We also acknowledge that the D peak is very intense in this nanomaterial, which may be partially due to the abundant edges.

The XPS survey spectrum (Figure 2b) demonstrates the chemical purity of the graphene material, as the only visible peak (at $\sim 285 \text{ eV}$) corresponds to C 1s. To further investigate
the nature of the carbon in the material, we studied the C 1s peak by a high-resolution narrow scan (Figure 2c). We fit the C 1s spectrum with two components corresponding to sp² hybridized C-C bonds (284.1 eV) and a π-π* satellite (290.5 eV) with a Tougaard background. The sp² peak was fit to a finite Lorentzian line shape with an asymmetry parameter of 0.18 and a full-width at half-maximum of 0.87 eV, consistent with previous reports. 33,34 Here, the finite Lorentzian line shape refers to a convolution of the Lorentzian and Gaussian functions, with the Lorentzian function asymmetrically raised to different exponents for binding energies above and below the peak center. This approach therefore creates asymmetric tails, which is useful for modeling conducting materials. We further note that the finite Lorentzian line shape is an established method for modeling the C 1s sp² component, whose use and practical applications have been discussed in various sources, 35-37 and that the specific function used for this spectral analysis was defined as LF(0.53, 1.2, 240, 250,3) in the CasaXPS software. The π-π* satellite was fit with a symmetric Gaussian–Lorentzian function with a width of 3.6 eV. Given that the sp² peak and the satellite were fairly well resolved, we did not impose constraints on the peak widths and peak positions. The fitted peak positions (i.e., the energy separation of 6.4 eV between the sp² and the satellite peaks 38) as well as the width of the sp² peak thus obtained were found to be consistent with the literature, which confirmed the validity of our fit. The residuals spectrum in Figure 2c further demonstrated that there was no need to consider additional peaks in the fitting. The fit yielded an Abbe criterion of 0.67, which was not ideal compared with the Abbe criterion of 1 for perfect noise around a fit. However, a linear fit to the spectral background (Figure 2c, inset) gave an Abbe criterion of 0.69, suggesting that the Abbe criterion of the peak fit was reasonable based on the intrinsic noise in this spectrum. 39

The strong sp² peak and the presence of the π-π* peak verify the graphene crystallinity, i.e., domains with delocalized p-orbitals. 34,36 The combined results of Raman spectra and XPS data confirmed that the GNSPs are highly crystalline and chemically pure graphene material. Further characterization of the GNSPs, including transmission electron microscopy, energy-dispersive X-ray spectroscopy, selected area electron diffraction, and ultraviolet photoelectron spectroscopy can be found in ref 26.

The GNSPs were packaged into a supercapacitor in a coin cell configuration, and the capacitance was measured by cyclic voltammetry at various scan rates between 2 and 4 V, as shown in Figure 3a1. The cyclic voltammograms were nearly rectangular, indicating a fairly ideal electrochemical double-layer capacitor process with little contribution from redox processes. 41 The nonidealities, such as the positive slope while charging and negative slope while discharging, may arise from slight changes in the intercalation state of Li⁺ or PF₆⁻ (e.g., Li⁺ intercalation could occur at voltages as high as 2.5 V in disordered graphene materials; for further discussion on the matter, see Supporting Information Note 1), which had also been reported previously in graphene supercapacitors. 1 In addition, these electrodes demonstrated the classic capacitor behavior of increasing current response with increasing scan rate. 1

Graphene electrodes were electrochemically exfoliated in situ by ramping the cell voltage to 1 V, holding the cell voltage at 1 V for 3 h, then ramping the cell voltage to a high voltage, holding the cell voltage at the high voltage for 3 h, then ramping down to 2 V. The high voltages ranged from 5 to 10 V. The rationale for this voltage program was that a high voltage caused intercalation of PF₆⁻, 44 which, as a large molecule, induced separation of graphene sheets (i.e., exfoliation), and the low voltage (1 V) caused intercalation of a Li⁺–propylene carbonate complex, which also exfoliated the graphene. 15 We also performed these experiments without the 1 V exfoliation step, and the capacitance of these electrodes did not enhance as well as those that included the 1 V exfoliation step (see Figure S2).

Märkle et al. 21 performed a similar exfoliation procedure using the same electrolyte (1 M LiPF₆ in propylene carbonate/ dimethyl carbonate), the same counter/reference electrode (lithium metal), and similar cycling between relatively low (3 V) and high voltages (5.5 V) except employing a regular (not quasi-one-dimensional) graphitic material in the working electrode, but observed “[e]lectrical contact loss between the graphite particles themselves and/or particles and the current collector”. The only significant difference between the system
described in this contribution and the system studied by Märkle et al. is that we are employing quasi-one-dimensional GNSPs while they did not, and, as demonstrated herein, instead of observing catastrophic device failure, we observe significantly enhanced capacitance due to exfoliation.

Before proceeding further, we remark that using high exfoliation voltages may initially seem concerning since the stability window of the electrolyte in this configuration is reported as only 6 V.\textsuperscript{45,46} Indeed, we conjecture that this high voltage may cause some decomposition that affects ion transport. Further, such high voltages may be dangerous. However, in our experiments, we did not observe damages or danger on a macroscopic scale, and the microscopic damages, although present, were slight. We believe that these findings were due to the relatively small voltage gradient at any point within the cell despite the fact that the total voltage across the entire coin cell may be large. This behavior is typical of supercapacitors, as ions migrate to biased electrodes and screen the total voltage,\textsuperscript{47} and ion intercalation into the bulk of the GNSPs electrode may also provide additional screening.

After exfoliation, the capacitance was measured again by cyclic voltammetry under the same conditions. The cyclic voltammograms for graphene exfoliated at 10 V are shown in Figure 3a2. The voltammograms of the electrode exfoliated at 10 V (Figure 3a2) revealed two differences from the preexfoliated graphene (Figure 3a1): (i) the current at a given voltage was much higher, demonstrating a higher capacitance, and (ii) the nonideal behavior, i.e., the slope, was more uniform through the voltammogram. The latter phenomenon may be due to intercalation behavior through the entire voltage range, which may be made possible by an increase in interlayer spacing and a decrease in intercalation potential. The cyclic voltammograms for graphene electrodes exfoliated at 5, 6, 7, 8, and 9 V as well as the reduced-scale cyclic voltammograms (to visualize lower scan rates) of all electrodes are provided in Figures S2–S9.

We calculated the specific capacitance, $C$, of the electrode at each degree of exfoliation (the preexfoliation capacitance and the capacitance after each exfoliation step) according to eq \ref{eq:cap}.

$$C = \frac{\int I \, dV}{2\mu m \Delta V}$$

(1)

where $\int I \, dV$ is the integrated area of the current response ($I$) of the cyclic voltammetry curve with the voltage differential ($dV$), $\nu$ is the scan rate, $m$ is the mass of the graphene, and $\Delta V$ is the voltage window of the cyclic voltammetry scan. The capacitances are shown in Figure 3b1. These data follow the classic supercapacitor behavior of decreasing capacitance with increasing scan rate.\textsuperscript{41} For the lowest scan rate, the capacitance increased steadily with increasing exfoliating voltage. For the highest scan rate; however, exfoliation at 8 V resulted in the best performance, while exfoliation at 9 and 10 V resulted in lower capacitance for high scan rates. This decrease in performance for exfoliation at 9 and 10 V may be due to solvent decomposition on the electrode surface during exfoliation,\textsuperscript{45,46} producing byproducts that impaired ion transport to and from the electrode such that the capacitance was impaired for high scan rates but not low scan rates.

To quantitatively compare the performance of each exfoliated electrode with respect to the preexfoliated electrode, we calculated the capacitive enhancement at each scan rate ($E_{\nu}$) according to eq \ref{eq:enh}.

$$E_{\nu} = \frac{C_{\text{exfoliated}, \nu}}{C_{\text{initial}, \nu}} \times 100\%$$

(2)

where $C_{\text{exfoliated}, \nu}$ is the specific capacitance at a scan rate, $\nu$, of the exfoliated graphene and $C_{\text{initial}, \nu}$ is the specific capacitance at a scan rate, $\nu$, of the preexfoliated graphene. Figure 3b2 plots the capacitive enhancement at each scan rate and demonstrates an enhancement of 418\% for graphene exfoliated at 10 V. In addition, for graphene electrodes exfoliated at 5 and 6 V, the enhancement increases with increasing scan rate, whereas for graphene electrodes exfoliated at 9 and 10 V, the enhancement decreases with increasing scan rate. For graphene electrodes exfoliated at 7 and 8 V, the enhancement first decreases and then increases with increasing scan rate.

To further understand this increase in capacitance, we investigated whether its origin was associated with an increase in nonfaradic charge storage (i.e., double-layer capacitance) or in faradic charge storage (i.e., redox reactions). An increase in the double-layer capacitance would result from an increase in the effective surface area, while an increase in faradic charge storage would result from chemical activation of the graphene material such that redox-active sites were created. Nonfaradic and faradic processes can be distinguished by comparing the cyclic voltammograms with fast and slow scan rates. Generally, nonfaradic (capacitive) current dominates at fast scan rates, and faradic current dominates at slow scan rates. Therefore, if a faradic process is taking place at an electrode, then a redox peak at some voltage will be visible at low scan rates but will be hidden by the nonfaradic current at high scan rates.\textsuperscript{47} To determine whether the increased capacitance observed here involved faradic processes, we normalized the cyclic voltammogram current response for each exfoliation step at the slowest scan rate (Figure 4a) and the highest scan rate (Figure 4b). In both plots, we observe that the current curves were similar and quasi-rectangular for all electrodes exfoliated at 6–10 V for the slowest scan rate and at 5–10 V at the highest scan rate.

![Figure 4. Normalized current response for each electrode at 3.16 mV/s (a) and 1000 mV/s (b).](https://dx.doi.org/10.1021/acsomega.0c06048)
suggested that the electrochemical processes did not change significantly with exfoliation. For further discussion of the evidence for the increase of surface area during exfoliation, see Supporting Information Note 2.

Next, we investigated the effect of the number of exfoliation cycles on capacitance enhancement. For these experiments, the capacitance was measured then the GNSP electrode was held at 1 V for 1 h, then at 6 V for 1 h, then returned to 2 V for a cyclic voltammogram capacitance measurement. This process was repeated for 20 cycles. Similar measurements were also repeated using 10 V instead of 6 V. These voltages (6 and 10 V) were chosen because 6 V was the electrochemical stability window of the electrolyte, \(^{45,46}\) and previous measurements demonstrated the greatest enhancement at 10 V. The cycle-dependent exfoliation using a high voltage of 6 and 10 V is shown in Figure 5. In both cases, exfoliation always enhanced the capacitance, but the number of cycles for maximum enhancement differed: For exfoliation at 6 V, the capacitance enhancement plateaued after two cycles and resulted in \(\sim 300\%\) capacitance enhancement, whereas for exfoliation at 10 V, enhancement maximized at \(\sim 250\%\) after nine cycles, then decayed with increasing cycles.

The data shown in Figure 5 contrast with the results presented in Figure 3 in two aspects: In Figure 3, exfoliation at 10 V resulted in higher capacitance than exfoliation at 6 V; and exfoliation at 10 V resulted in \(\sim 420\%\) capacitance enhancement in Figure 3 rather than \(\sim 250\%\) enhancement in Figure 5. We resolve this discrepancy by considering that the electrolyte was not stable under a voltage gradient of 10 V \(^{45,46}\) so that exfoliating at 10 V may have caused solvent decomposition that was destructive to the capacitance. In addition, exfoliating at lower voltages before exfoliating at 10 V may enable better ion transport into the bulk of the electrode (e.g., via pores) that helped screen the total voltage such that when a voltage was applied to the electrode, there was a voltage gradient across the electrode, and the voltage gradient at the solution–electrode interface was not too intense. In other words, if the electrode was partially exfoliated before applying a high voltage, then counter ions in the bulk of the electrode would help screen the voltage for the electrode–solution interface when a high voltage was applied, leading to reduced electrolyte decomposition while enabling further exfoliation.

To better understand the cause of increased capacitance in exfoliated electrodes, we measured the X-ray diffraction (XRD) patterns of our GNSP electrodes exfoliated at different voltages (Figure 6). Specifically, we investigated the XRD spectra of an as-grown GNSP electrode, a GNSP electrode exfoliated at 6 V for 20 cycles, and a GNSP electrode exfoliated at 10 V for 20 cycles. Interestingly, for both the as-grown electrode and the electrode exfoliated at 6 V, the position of the [002] peak was similar, but for the electrode exfoliated at 10 V, the [002] peak disappeared. According to Bragg’s law \(d_{002} = n\lambda/2 \sin \theta\), where \(n\) is a positive integer, \(\lambda\) is the incident X-ray wavelength, and \(\theta\) is the diffraction angle, the interlayer spacings in the as-grown graphene and graphene exfoliated at 6 V were 3.40 and 3.37 Å, respectively. Given that the [002] peak indicated the interlayer spacing, the presence of the [002] peak implied that applying a voltage of 6 V did not fully exfoliate the graphene (i.e., separate it into individual monolayers) even though the peak intensity was reduced and linewidth broadened (suggesting partial exfoliation). In contrast, applying a voltage of 10 V fully exfoliated the GNSP electrode so that there was no longer ordered c-axis stacking, hence the complete disappearance of the [002] peak.

Finally, we used electrochemical impedance spectroscopy (EIS) to further understand the changes taking place during the exfoliation that led to increased capacitance. The Nyquist plots for each exfoliation step are shown in Figure 7a. The impedance data were fit to a modified Randles circuit (Figure 7b) with the fit parameters and mean-square errors (MSE) provided in Table 1. In the equivalent circuit, the capacitor and the Warburg diffusion elements are replaced with constant phase elements (CPE\(_1\) and CPE\(_W\), respectively) to account for the nonidealities.

Before interpreting the equivalent circuits and the model parameters, we introduce the mathematical relations of the circuit elements. The impedances of a capacitor (\(Z_C\)), Warburg element (\(Z_W\)), and constant phase element (\(Z_{CPE}\)) are defined, respectively, in eqs 3–5

\[
Z_C = \frac{1}{i\omega C} \quad (3)
\]

\[
Z_W = \frac{1}{Q\sqrt{i\omega}} \quad (4)
\]

\[
Z_{CPE} = \frac{1}{Q(i\omega)^n} \quad (5)
\]

where \(i\) is the imaginary number, \(\omega\) is the radial frequency, \(C\) is the capacitance, \(R\) is the resistance, and \(n\) and \(Q\) are constants.\(^{41–43}\) In the cases of \(n = 1\) and 0.5, the constant phase element (CPE) becomes mathematically identical to the capacitor and Warburg element, respectively. Therefore, a CPE is often used to model nonideal capacitors and Warburg elements.\(^{41,48}\) For a CPE that is modeling a capacitor, the extent to which \(n\) deviates below 1 can be interpreted as the “leakiness” of the capacitor, and for a CPE modeling a Warburg element, values of \(n\) deviating above 0.5 indicate diffusion within a porous three-dimensional structure (as opposed to diffusion near a perfect two-dimensional plane).\(^{41,48,49}\)
As the material becomes sufficiently exfoliated such that pores were large enough to behave quasi-two-dimensionally, exfoliating at or above 7 V nearly restored the di-fusion behavior and an increase in porosity of the material, respectively.

Exfoliating at or above 7 V nearly restored the diffusion behavior to a Warburg element, which we interpret as the capacitance increased and the intercalation resistance decreased, suggesting better electrolyte contact with the material. In addition, the “leakage” from CPE may represent some intercalation behavior.

We provide the following interpretation of the modeling parameters shown in Figure 7b. In the equivalent circuit, CPE represents the double-layer capacitance, R1 represents intercalation charge transfer resistance, and CPEW represents diffusion behavior. In addition, the “leakage” from CPE may represent some intercalation behavior.

Exfoliating to 5 and 6 V increased Q1 (which is often interpreted as the capacitance), decreased the intercalation resistance (R1), and increased nW (implying three-dimensional diffusion behavior). We physically interpret these parameter changes as an increase in the surface area and decrease in the intercalation energy barrier due to better electrolyte contact and an increase in porosity of the material, respectively. Exfoliating at or above 7 V nearly restored the diffusion behavior to a Warburg element, which we interpret as the material becoming sufficiently exfoliated such that pores were large enough to behave quasi-two-dimensionally.

For exfoliating voltages above 7 V, R1 increased substantially with increasing exfoliating voltage. This may be due to electrolyte decomposition, which impeded ion transport and likely responsible for the degraded performance at high scan rates for graphene electrodes exfoliated above 7 V (see Figure 3). In addition, n1 decreased to 0.67, indicating a very leaky capacitor. It is interesting, however, that these electrodes demonstrated an increased capacitance (at low scan rates, see Figure 3) despite electrolyte decomposition. On the other hand, for exfoliating voltages above 7 V, the MSE of the fit increased substantially, indicating that the device no longer complied with a Randles circuit model, which may be attributed to nonidealities caused by electrolyte decomposition.

### DISCUSSION

Our capacitance measurements demonstrate that GNSP electrodes can be exfoliated in situ to substantially enhance their capacitance (Figure 3), which is made possible by the quasi-one-dimensional nature of GNSPs (Figure 1). Comparison of the current responses of each electrode at both fast and slow scan rates indicates that the charge storage processes in the GNSP electrodes are nonfaradic (Figure 4), which suggests that the increase in capacitance is due to structural rather than chemical changes (e.g., chemical functionalization). Comparing the cycle-dependent exfoliated GNSP electrode (Figure 5) with the gradually exfoliated GNSP electrode (Figure 3) demonstrates that when exfoliating at 10 V, initial exfoliation at a lower voltage results in better capacitance. Our XRD data (Figure 6) reveal that an applied voltage of 6 V does not fully exfoliate the GNSP electrode, whereas applying a voltage of 10 V completely exfoliates the electrode. Finally, our EIS data (Figure 7) suggest that the electrochemical processes have fundamentally altered with exfoliation: the diffusion behavior first shifts from a two-dimensional to a three-dimensional diffusion behavior, suggesting that the material becomes porous upon exfoliation. In addition, the double-layer capacitance increases and the intercalation resistance decreases, suggesting better electrolyte contact with the material. As the material becomes sufficiently exfoliated, the diffusion returns to a two-dimensional behavior. The EIS data also suggest that some intercalation charge transfer has taken place. However, noting that the electrochemical processes are nonfaradic based on the cyclic voltammetry data, the intercalation charge transfer is likely minimal, although it may account for the observation of nonideal capacitive charge storage.

We acknowledge that our reported capacitances (55 F/g, see Figure 3b1) are not particularly good, even for the best-
performing devices (Liu et al., for example, report a capacitance of 154 F/g for a graphene electrode). However, maintaining the electrical connectivity with GNSPs. We are also interested in using exfoliated GNSPs to form percolating networks among high capacitance metal oxide nanoparticles. Overall, we consider in situ exfoliation of electrode materials together with the incorporation of quasi-one-dimensional GNSPs a promising research direction for achieving substantially improved supercapacitors.

## EXPERIMENTAL SECTION

**GNSP Fabrication.** GNSPs were fabricated by plasma-enhanced chemical vapor deposition (PECVD) described previously. Specifically, a microwave-induced hydrogen/methane plasma with trace 1,2-dichlorobenzene (Alfa Aesar, 99%) yielded GNSPs on a ∼(0.75 cm × 1.25 cm) copper foil (McMaster-Carr, 99.9%) in a 1/2” outer diameter quartz tube. The PECVD growth system was custom built and consisted of eight parallel deposition chambers each fitted with an Evenson Cavity (Optos Instruments Inc., Frederick, MD) excited by 70 W of 2.45 GHz microwave power source (ENS 4 × 200 W CPS, Sairem, Décines-Charpieu, France). The plasma volume was ∼1 cm³. H₂ (MATHESON, 99.999%) and CH₄ (MATHESON, 99.999%) gases were introduced to the chambers by mass flow controllers (MC series, Alicat Scientific, Tuscan, AZ), and 1,2-dichlorobenzene was placed in a vacuum-sealed vial and introduced to the chamber through a leak valve. The pressure in the chamber before splitting into eight chambers was held at 4.8 Torr, the total flow rates of H₂ and CH₄ were 48 and 5 sccm, respectively, and the ratio of CH₄ to 1,2-dichlorobenzene (3-chloropyridine) was ∼2:1 as measured by a residual gas analyzer (RGA; XT300M, Extorr Inc., New Kensington, PA) placed upstream of the deposition chamber and connected via a capillary. The plasma was maintained for ∼3 h to synthesize sufficient graphene material for use as a supercapacitor electrode.

**GNSP Characterization.** The resulting GNSP material was characterized by scanning electron microscopy (SEM), Brunauer–Emmett–Teller (BET) surface area method, Raman spectroscopy, X-ray photoelectron spectroscopy (XPS), and X-ray diffraction (XRD). SEM was performed using a Hitachi S-4100 (Hitachi, Tokyo Japan) with an accelerating voltage of 5 kV. Individual GNSPs were obtained for imaging by sonicating the copper growth substrate with GNSPs grown on it in N-methyl-2-pyrrolidinone (NMP, Sigma-Aldrich, 99.5%) for 30 min, centrifuging the dispersion to remove undispersed chunks of graphitic material at 850 rpm for 90 min, extracting one drop of dispersion, and evaporating it over a silicon wafer in low vacuum at 70 °C. The BET surface area was measured in a BELSORP-max volumetric instrument (BEL-Japan, Inc.) via equilibrium N₂ adsorption isotherm at 77 K. Raman spectroscopy was performed in a Renishaw M-1000 Micro-Raman (Renishaw, Gloucestershire, U.K.) spectrometer operating with a 514.5 nm argon-ion laser with a spectral resolution of 1 cm⁻¹ and a spot size of ∼20 µm. A dual-wedge polarization scrambler was inserted to depolarize the laser. XPS data were collected using a Kratos AXIS Ultra spectrometer (Kratos Analytical, Manchester, U.K.). The instrument was equipped with a hybrid magnetic and electrostatic electron lens system, a delay-line detector, and a monochromatic Al Kα X-ray source (1486.7 eV). Data were collected at a pressure of ∼5 × 10⁻⁹ Torr with photoelectrons collected at 0° with respect to the sample. The analyzer pass energy was 80 eV for the survey spectrum and 10 eV for all other spectra. The instrument energy scale and work...
function were calibrated using clean Au, Ag, and Cu standards. The instrument was operated by Vision Manager software v. 2.2.10 revision 5. The data were analyzed using CasaXPS software (CASA Software Ltd). XRD data were collected in a PANalytical X’Pert Pro X-ray powder diffractometer using the Cu Kα1 line (λ = 1.5406 Å) with a tube voltage and current of 40 kV and 20 mA, respectively.

Coin Cell Preparation. Supercapacitor working electrodes consisted of GNSP material and polyvinylidene fluoride (PVDF, MTI Corporation, ≥ 99.5%) binder in an 88:12 ratio and were mixed in N-methyl-2-pyrrolidinone (NMP, Sigma-Aldrich, 99.5%) in a centrifugal mixer (AR-100 Thinky U.S.A., Inc., Laguna Hills, CA) at 5000 rpm for 10 min. A thin layer of the resulting slurry was spread across a stainless-steel spacer (MTI Corporation) with a spatula and dried at 120 °C in vacuum for 16 h. Two-electrode 2032-coin cells were assembled in an argon-filled glovebox (O2: < 0.1 ppm, H2O: < 0.1 ppm). The counter/reference electrode was lithium foil (Sigma-Aldrich, 99.9%, 0.75 mm, mechanically cleaned immediately before cell assembly), and the separator was a propylene separator (Celgard 2400). The electrolyte was 1 M LiPF6 in propylene carbonate/dimethyl carbonate (1:1 mixture by volume, both Sigma-Aldrich, ≥99%), stored over molecular sieves, 3 Å, Beantown Chemical), and about eight drops of electrolyte were used in each coin cell. The electrolyte was mixed in a dried HDPE bottle.

Electrochemical Characterization. All electrochemical measurements were performed on a Reference 600 (Gamry Instruments, Warminster, PA). Capacitance was measured by cyclic voltammetry between 2 and 4 V at scan rates of 3.16, 10, 31.6, 100, 316, and 1000 mV/s. Electrochemical impedance spectroscopy measurements were taken potentiostatically at 2 V with an rms voltage of ±5 mV. All voltages are with respect to Li/Li+. Lithium intercalation in graphene; capacitance and enhancement of GNSP electrodes exfoliated without the 1 V exfoliation step; cyclic voltammogram capacitance measurements of GNSPs exfoliated with high voltages ranging from 5 to 10 V (Note 1 and Figure S1); cyclic voltammograms for graphene electrodes exfoliated at 5, 6, 7, 8, and 9 V, as well as the reduced-scale cyclic voltammograms of all electrodes (Figures S2–S9); and extrapolation of double-layer capacitance and solution resistance from cyclic voltammograms (Note 2 and Figure S10) (PDF)

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Author Contributions
J.D.B. designed the experiment; fabricated the GNSP materials; carried out Raman spectroscopy, XPS, XRD, and electrochemical measurements; and participated in preparing the manuscript. D.R.D. carried out GNSP dispersion and SEM imaging. N.-C.Y. coordinated and oversaw the research project and participated in preparing the manuscript.

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Notes
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