Graphene/PVDF Composites for Ni-rich Oxide Cathodes toward High-Energy Density Li-ion Batteries

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Abstract: Li-ion batteries (LIBs) employ porous, composite-type electrodes, where few weight percentages of carbonaceous conducting agents and polymeric binders are required to bestow electrodes with electrical conductivity and mechanical robustness. However, the use of such inactive materials has limited enhancements of battery performance in terms of energy density and safety. In this study, we introduced graphene/polyvinylidene fluoride (Gr/PVdF) composites in Ni-rich oxide cathodes for LIBs, replacing conventional conducting agents, carbon black (CB) nanoparticles. By using Gr/PVdF suspensions, we fabricated highly dense LiNi0.85Co0.15Al0.05O2 (NCA) cathodes having a uniform distribution of conductive Gr sheets without CB nanoparticles, which was confirmed by scanning spreading resistance microscopy mode using atomic force microscopy. At a high content of 99 wt.% NCA, good cycling stability was shown with significantly improved areal capacity (Q\textsubscript{areal}) and volumetric capacity (Q\textsubscript{vol}), relative to the CB/PVdF-containing NCA electrode with a commercial-level of electrode parameters. The NCA electrodes using 1 wt.% Gr/PVdF (0.9:0.1) delivered a high Q\textsubscript{areal} of ~3.7 mAh cm\textsuperscript{-2} (~19% increment) and a high Q\textsubscript{vol} of ~774 mAh cm\textsuperscript{-3} (~18% increment) at a current rate of 0.2 C, as compared to the conventional NCA electrode. Our results suggest a viable strategy for superseding conventional conducting agents (CB) and improving the electrochemical performance of Ni-rich cathodes for advanced LIBs.

Keywords: lithium-ion batteries; Ni-rich cathodes; graphene; PVDF; electrode density; scanning probe microscopy

1. Introduction

In an attempt to meet the global energy demand while maintaining sustainable development of society, rechargeable batteries have been considered to be effective solutions for renewable energy [1,2]. Lithium-ion batteries (LIBs) have been used as major power sources for a variety of applications, including portable electronics and electric vehicles, since having been devised in the 1990s [3–7]. Lower energy cost LIBs are desirable (i.e., <$125 kWh\textsuperscript{-1} for pack cost for electric vehicles) for providing a longer lifespan at an affordable cost [1,8]. Accordingly, most research has focused on the development of electrode materials and the electrode engineering via tuning the electrode parameters toward cycling-stable LIBs with high-energy density [4–9].

LIB electrodes are composed of a porous, composite-type film coated on a current collector foil, where the film contains active materials, carbonaceous conducting agents, and polymeric binders. Due to their low electrical conductivities, typical cathodes for LIBs require few percentages of nanoscale conductive additives, such as carbon blacks (CBs). However, employing CB nanoparticles in electrodes causes intrinsic challenges in
Materials manufacturing processes, such as slurry formulation and coating/drying steps, due to their agglomeration behavior [10,11]. In addition, the low tap density of CB restricts an improvement of volumetric capacity (energy density) of electrodes. In practice, the slurry preparation step is decisive in a subsequent slurry coating process and the resulting electrochemical performance of electrodes [10]. Many studies on slurry processing have been conducted to improve rheological behavior and resulting electrode performance by modifying the methods and sequences of constituent mixing [10,12–14]. Nevertheless, the uniform dispersion of CB nanoparticles within a slurry and hence, their effective distribution in electrodes for ensuring the conductive pathways still remain challenging, even with smaller CB contents.

In this study, we used Gr nanosheets as a conducting agent in a Ni-rich oxide cathode, LiNi_{0.85}Co_{0.15}Al_{0.05}O_2 (NCA), replacing conventional CB nanoparticles for LIBs. By simply mixing an electrochemically exfoliated Gr dispersion and a polyvinylidene fluoride (PVdF) binder solution, we fabricated uniform Gr/PVdF suspensions with controlled Gr contents; this enabled uniform distribution of the Gr nanosheets within NCA electrodes due to the intimate interactions of the functional groups of Gr and the electronegative fluorine in PVdF. Surface probe microscopy on a cross section of the electrodes was conducted to differentiate the microstructure and intraparticle resistances. Using 1 wt.% Gr/PVdF suspension (Gr:PVdF = 0.9:0.1 in wt.%) without a conventional CB, we achieved highly dense NCA cathodes (99 wt.% NCA; electrode density ($\rho$)~3.8 g cm$^{-3}$) that showed good cycling stability and higher areal capacity ($Q_{\text{areal}}$) and volumetric capacity ($Q_{\text{vol}}$) than those of a counterpart with a commercial-level of electrode parameters (96 wt.% NCA; $\rho$~3.3 g cm$^{-3}$) at a current rate of 0.2 C.

2. Materials and Methods

2.1. Preparation of Gr/PVdF Suspensions

Gr nanosheets were prepared using an electrochemical exfoliation method. As the electrolyte, an aqueous solution containing 0.5 M ammonium sulfate ((NH$_4$)$_2$S$_2$O$_8$) was used, where two graphite foil electrodes were immersed in parallel at a fixed distance of 10 mm. When applying a DC voltage (+10 V) between the two graphite electrodes for about 10 min, a positively charged electrode was gradually exfoliated into Gr nanosheets, which was then collected through vacuum filtration, followed by washing with deionized water and ethanol. Gr/PVdF suspensions with various Gr contents (10–90 wt.%) were prepared in the following two steps. First, the Gr dispersion and PVdF (Solef® 6020, Solvay, Brussels, Belgium) solution were prepared at concentrations of 2 wt.% and 10 wt.% in N-methyl-2-pyrrolidone (NMP), respectively. The two separate solutions were then prepared with a vortex mixer to form Gr/PVdF suspensions at different Gr contents (10–90 wt.%).

2.2. Electrochemical Testing

The cathode slurries were prepared by mixing 99 wt.% NCA powder (Samsung SDI, Yongin, Korea) and 1 wt.% Gr/PVdF suspensions, where the Gr content was varied from 0.1 to 0.9 wt.%. The as-prepared slurries were cast on a 20 μm-thick Al foil using a doctor blade method and dried in a convection oven at 80 °C for 2 h, followed by vacuum drying at 120 °C overnight. For comparison, NCA electrodes with commercial-level electrode parameters of NCA:CB (Super P®, Imerys Graphite & Carbon, Bironico, Switzerland):PVdF = 96:2:2 were prepared. The NCA electrode was set as 3.8 (±0.1) g cm$^{-3}$ and 3.3 (±0.1) g cm$^{-3}$ for Gr-containing electrodes and CB-containing electrodes, respectively. The mass loading was set as ~16 mg cm$^{-2}$ and it was increased to ~19 mg cm$^{-2}$, to fabricate thicker electrodes. The electrodes were punched into discs with a diameter of 12 mm. Coin-type 2032 cells were assembled for electrochemical testing in an Ar filled glove box, with a lithium metal counter-electrode (16Φ) and polypropylene separator (Celgard 2400, Celgard, Charlotte, NC, USA). LiPF$_6$ (1 M) in a solvent mixture of ethylene carbonate, ethyl methyl carbonate, and dimethyl carbonate at a volume ratio of 2:4:4 with 1.5% vinylene carbonate additive was used as the electrolyte.
Electrochemical properties were measured using a WBCS3000L battery cycler (WonAT-tech, Seoul, Korea). The formation process was conducted at 0.2 C (1C = 200 mAh g$^{-1}$) in the voltage range of 2.75–4.3 V vs. Li$^+$/Li for two cycles with a CC-CV charge mode using a 0.05 C cut-off. The discharge rate capability was measured at various current rates of 0.2–10 C after charging at 0.2 C with a CC-CV protocol using a 0.05 C cut-off. The cycle performance was evaluated at 30 and 60 °C in the voltage range of 2.75–4.3 V or 2.75–4.5 V vs. Li$^+$/Li at a current rate of 0.5 C with a CC-CV charging protocol using a 0.05 C cut-off after two cycles of the formation process. Electrochemical impedance spectroscopy (EIS) was carried out using a biologic VSP-100 instrument in the frequency range of 1 MHz to 10 mHz with a voltage perturbation of 10 mV. The internal pressure change was monitored using a home-made gas cell system containing a cathode charged to 4.5 V vs. Li$^+$/Li and 10 mL of a fresh electrolyte at 60 °C for 24 h in an Ar atmosphere.

2.3. Characterizations

The microstructure and Gr distribution were observed using field-emission scanning electron microscopy (SEM, JSM-7600F, Jeol Ltd., Tokyo, Japan). The samples were cross-sectioned using a cross-section polisher (SM-09010, Jeol Ltd., Tokyo, Japan). The in-plane resistance of NCA electrodes was measured using an electrode resistance measurement system (RM2610, Hioki, Nagano, Japan). The electrical resistance distribution inside the cross-sectioned electrodes was measured in scanning spreading resistance microscopy (SSRM) mode using atomic force microscopy (AFM, NX-10, Park systems, Suwon, Korea) equipped with a diamond-coated probe (CDT-NCHR, NANOSENSORS™, Neuchatel, Switzerland) with a nominal force constant of 80 N m$^{-1}$ over a scan area of 10 × 10 µm$^2$. By applying appropriate bias voltages (~1.5 V), the current passing through samples (between the tip and sample base, Al current collector) was recorded using a logarithmic current amplifier in SSRM mode.

3. Results and Discussion

We prepared Gr dispersions by an electrochemical exfoliation method with commercially available graphite foils [15–17]. By applying a DC voltage (+10 V), a graphite foil was exfoliated into Gr sheets in an aqueous ammonium persulfate solution (0.5 M), due to the evolution of gases, such as SO$_2$ and O$_2$. The as-exfoliated Gr sheets were then collected by vacuum filtration and washing processes, followed by dispersion in a NMP solvent, as schematically depicted in Figure 1a. The exfoliated Gr sheets have lateral sizes of 0.1–2 µm and an average thickness of 4.6 nm (refer to Figure S1). The Gr surface/edges appeared to be partially oxygen-functionalized (C–OH/C–O/C=O) due to the oxidation of graphite by OH$^-$ ions during the electrochemical process on the basis of XPS C 1s spectra (Figure S2) [16,17]. We performed confocal Raman mapping on the exfoliated Gr sheets, which showed the high-intensity G peak and 2D peak at ~1580 and ~2700 cm$^{-1}$, respectively, with an intensity ratio of D to G ($I_D/I_G$) of ~0.3 (Figure S3), which is much lower than that of chemically or thermally reduced graphene oxides ($I_D/I_G = 1.2–1.5$) [15,18,19].
Gr/PVdF suspensions were fabricated by simply mixing the Gr dispersion (2 wt.% Gr) and PVdF solution (10 wt.% in NMP) at different Gr contents (10–90 wt.%), speculating that two types of interactions between the Gr sheets and PVdF occur:—(i) hydrogen bonding between the oxygen functional groups (–O–C=O/–OH) on Gr and –CF2/–CH2 dipoles in PVdF and (ii) π–π interactions between the delocalized π electrons in Gr and –CH2 dipoles in PVdF—(see the right side of Figure 1a) [20–25]. A Gr/PVdF suspension (50:50 in wt.%) was cast on an Al foil and the film microstructure was compared to that of the film coated with a CB/PVdF suspension (50:50 in wt.%). As shown in Figure 1b,c, the Gr/PVdF film appeared to be highly homogeneous without noticeable protrusions and the plan-view SEM image showed uniform microstructure (Figure 1b). On the other hand, local protrusions and microcracks were found on the surface of the CB/PVdF film, as indicated by red arrows in Figure 1c. This difference in the microstructure between the two different type films could stem from the specific interactions between the Gr sheets and PVdF and the intrinsic agglomeration of CB in NMP (refer to Figure S4). The agglomeration of CB can cause intrinsic issues in slurry formation and coating/drying processes for electrode fabrication.

Typically, inactive components, such as CB and PVdF binder, have been used to build Ni-rich oxide cathodes; however, they should be minimized to enhance the energy density (volumetric capacity) of electrodes [26,27]. We fabricated electrodes having a high NCA content (99 wt.%) using Gr/PVdF and CB/PVdF suspensions (Figure 1d–g). The NCA electrodes composed of Gr/PVdF were robust without noticeable delamination from the Al foil current collector, even for the electrode containing a low PVdF content (0.1 wt.%).
By contrast, the NCA electrodes using the CB/PVdF were easily detached after electrode punching: the electrode containing 0.1 wt.% PVdF binder was completely delaminated from the current collector (Figure 1g).

We assessed rate capability of NCA electrodes that were fabricated using Gr/PVdF suspensions with different Gr contents (x in NCA:Gr:PVdF = 99:x:1−x) and compared a NCA electrode with a commercial-level electrode setting with NCA:CB:PVdF = 96:2:2 and ρ=3.3 g cm$^{-3}$, which is hereafter denoted as the Ref. electrode. The discharge rate capability of the NCA electrodes measured at 0.2−10 C are shown in Figure 2a and the corresponding voltage profiles measured in the voltage range of 2.75–4.3 V vs. Li$^+$/Li are shown in Figure S5. The NCA electrode containing 0.9 wt.% Gr exhibited good rate performance, comparable to that of the Ref. electrode: the two electrodes had ~200 and ~183 mAh g$^{-1}$ at 0.2 C and 2 C, respectively. The NCA electrodes with less Gr contents (0.5 and 0.7 wt.%) showed slightly inferior performance, which could be attributed to the low electrical conductivities. These results were corroborated by in-plane conductivity and EIS measurements. As shown in Figure 2b, only the NCA electrode containing 0.9 wt.% Gr exhibited a comparable electrical conductivity with that of the Ref. electrode, whereas the others had low conductivities. In addition, we compared Nyquist plots of the NCA electrodes in configurations of a symmetric cell (NCA|NCA) and an asymmetric cell (Li|NCA), as depicted in Figure S6. The data for symmetric cells before the formation process (Figure S6a) revealed that the contact resistance between NCA particles and current collector and the interparticle resistance within electrodes was decreased with increasing Gr content [28,29]: the Nyquist plot of 0.9 wt.% Gr-NCA electrode was similar to that of the Ref. electrode. Similarly, for asymmetric cells after formation (Figure S6b), 0.9 wt.% Gr-NCA electrode and Ref. electrode had almost similar trends in the Nyquist plots, two depressed arcs at high frequencies and a straight line inclined at a constant angle to the real axis at low frequencies [30].

Next, we evaluated the cycle performance of NCA electrodes with different electrode compositions, measured in the voltage ranges of 2.75–4.3 V or 2.75–4.5 V vs. Li$^+$/Li at a current rate of 0.5 C and temperatures of 30 °C and 60 °C. As shown in Figure 3a,b, the NCA electrode with 0.9 wt.% Gr exhibited a similar capacity retention to the Ref. electrode at 30 °C and 60 °C, respectively. With the decreasing Gr content, the capacity decay was severe. Under the operation at high-voltage cut-off condition (2.75–4.5 V vs. Li$^+$/Li) and 30 °C, both the 0.9 wt.% Gr-NCA electrode and the Ref. electrode had similar behaviors during 100 cycles (Figure 3c). These results confirm that even with a high NCA content (99 wt.%), but low contents of inactive components (total 1 wt.%), the NCA electrodes show excellent cycling stability, comparable to the Ref. electrode containing high contents.
of CB/PVdF (total 4 wt.%). Note that the Gr-NCA electrodes had a considerably higher $\rho$ ($\sim 3.8$ g cm$^{-3}$) than that of the Ref. electrode ($\sim 3.3$ g cm$^{-3}$), featuring dense electrode architectures. By considering the electrode volume, we replotted the $Q_{\text{vol}}$ retention data in Figure 3d–f. The NCA electrode with 0.9 wt.% Gr exhibited a considerably large $Q_{\text{vol}}$, compared to that of the Ref. electrode. For example, at 30 °C (Figure 3d), the 0.9 wt.% Gr-NCA electrode had a $Q_{\text{vol}}$ of $\sim 790$ mAh cm$^{-3}$ at the 1st cycle and delivered a $Q_{\text{vol}}$ of $\sim 580$ mAh cm$^{-3}$ at the 100th cycle ($\sim 73\%$ retention), whereas the Ref. electrode had a $Q_{\text{vol}}$ of $\sim 610$ mAh cm$^{-3}$ and $\sim 460$ mAh cm$^{-3}$ at 1st and 100th cycles, respectively, with capacity retention of $\sim 75\%$.

Figure 3. (a–c) Specific capacity retention and (d–f) volumetric capacity retention of NCA electrodes with different electrode compositions at current rate of 0.5 C, measured in the voltage range of (a,d) 2.75–4.3 V at 30 °C, (b,e) 2.75–4.3 V at 60 °C, and (c,f) 2.75–4.3 V at 30 °C.

To differentiate the microstructure and components distribution for NCA electrodes with different compositions (1 wt.% Gr/PVdF vs. 4 wt.% CB/PVdF), we performed CP-SEM observation for 0.9 wt.% Gr-NCA and Ref. electrodes. As shown in Figure 4a1,b1, a distinct difference in the electrode thicknesses between the 0.9 wt.% Gr-NCA electrode and Ref. electrode was observed due to their different $\rho$. The Gr sheets were uniformly distributed within the Gr-NCA electrode, overall (Figure 4b2–b4). On the other hand, the Ref. electrode showed local agglomeration of CB nanoparticles between the NCA particles.
(Figure 4a2–a4). It is noteworthy that the 0.9 wt.% Gr-NCA electrode had no notable micro-cracks, even after a high-pressure calendaring process for high electrode density (3.8 g cm⁻³). We further conducted SSRM measurements combined with AFM on the cross section for the 0.9 wt.% Gr-NCA and Ref. electrodes, to visualize spatial heterogeneity of the resistance within NCA particles (see Figure 4c,d). Comparison between the resistance mapping images revealed that the Gr-NCA electrode had relatively uniform distributions of the intraparticle resistance, relative to those of the Ref. electrode. This result indicates that the use of Gr/PVdF suspension affords the uniform distribution of the conducting agent, Gr and hence, effective conducting pathways within NCA electrodes, even with a significantly small Gr amount (0.9 wt.%).

![Figure 4. (a1–a4, b1–b4) Cross-sectional SEM images and (c,d) AFM and SSRM images of NCA electrodes with (a,c) CB/PVdF and (b,d) Gr/PVdF. The red arrows in b3 indicate Gr sheets between the NCA particles.](image-url)

Ni-rich oxide cathodes have been reported to cause safety issues due to structural instability arising from the phase transition during repeated cycles, accompanied by the evolution of O₂ and CO₂ gases [31,32]. This behavior is accelerated under battery operation with overcharging (>4.3 V vs. Li⁺/Li) and/or at elevated temperatures in the charged state. In addition, inactive materials such as CB and PVdF can release gaseous CO₂ through oxidation reactions with reactive oxygen species generated from charged NCA particles [33]. Such gaseous byproducts could induce severe thermal runaway by highly exothermic reaction with flammable electrolytes. We monitored the internal pressure change with respect to testing time using a home-made gas cell system, which contained a NCA electrode fully charged to 4.5 V vs. Li⁺/Li and 10 mL of a fresh electrolyte, at 60 °C in Ar atmosphere (refer to the inset of Figure 5). An empty cell and a cell containing only electrolyte were also examined as references. As shown in Figure 5, as the testing proceeded, the internal pressure monotonically increased for all the cells. The NCA electrode containing Gr/PVdF exhibited a relatively small increase of the pressure, as compared to that of the Ref. electrode, which appeared to be a result of the use of less contents of inactive materials, confirming the efficacy of Gr/PVdF suspension.
Lastly, we evaluated discharge \( Q_{\text{areal}} \) and \( Q_{\text{vol}} \) of NCA electrodes containing 1 wt.% Gr/PVdF. We fabricated three types of Gr-NCA electrodes (Gr-A, Gr-B, and Gr-C) with different electrode parameters, such as mass loading \( (m_{\text{areal}}; 16 \text{ mg cm}^{-2} \text{ vs. } 19 \text{ mg cm}^{-2}) \) and electrode density, \( \rho \) \( (3.3 \text{ vs. } 3.8 \text{ g cm}^{-3}) \), to compare with the Ref. electrode containing 4 wt.% CB/PVdF \( (m_{\text{areal}}; 16 \text{ mg cm}^{-2} \text{ vs. } 3.3 \text{ g cm}^{-3}) \). Figure 6a−d show the cross-sectional SEM images of the four electrodes with different electrode parameters. For the Ref. electrode (Figure 6a), the presence of intraparticle microcracks was prominent. On the other hand, overall, the microstructures of Gr-NCA electrodes with a high \( \rho \) \( (3.8 \text{ g cm}^{-3}) \) while reducing inactive components (1 wt.% Gr/PVdF), which enabled for an enhanced \( \rho \) and hence, a large \( Q_{\text{vol}} \) (thinner electrode), as well as a large \( Q_{\text{areal}} \) (high mass-loading). 

**Figure 5.** Internal pressure changes of in-situ gas cells, which contained NCA electrodes with CB/PVdF and Gr/PVdF (number of measurements: 5 times). Graphs for an empty cell and a cell containing only electrolyte as references are also shown.
Electrochemically exfoliated Gr nanosheets were employed as conducting agent in NCA cathodes for LIBs. Gr/PVdF suspensions were fabricated by mixing Gr dispersion (in NMP) and PVdF solution (in NMP) at Gr contents of 10–90 wt.%. The Gr/PVdF suspensions appeared to be considerably homogeneous, compared to the CB/PVdF suspensions. Consequently, the cast Gr/PVdF composite films had uniform microstructures, which was thought to be due to specific interactions between the Gr sheets and PVdF, such as hydrogen bonding and π–π interactions. By using Gr/PVdF suspensions, we achieved highly packed NCA electrodes featuring a high NCA content of 99 wt.% and a high density of 3.8 g cm$^{-3}$ without notable microcracks. The CP-SEM and AFM-SSRM analyses confirmed that the use of the Gr/PVdF suspension enabled the uniform distribution of Gr nanosheets, ensuring effective conducting pathways within NCA electrodes, even with an extremely small Gr amount (0.9 wt.%). In addition, the home-made gas cell measurements proved the efficacy of using a small content of Gr/PVdF (1 wt.%), as compared to the case of conventional CB/PVdF (4 wt.%), in terms of battery safety. The NCA electrodes using a small content of Gr/PVdF (0.9:0.1, total 1 wt.%) showed a larger $Q_{\text{vol}}$ of ~790 mAh cm$^{-3}$ than that of the Ref. electrode ($Q_{\text{vol}}$=610 mAh cm$^{-3}$) with a commercial-level of electrode setting (96 wt.% NCA; $\rho$=3.3 g cm$^{-3}$) with good capacity retentions during 100 cycles, measured in the voltage ranges of 2.75–4.3 V and 2.75–4.5 V vs. Li$^+/\text{Li}$ and at temperatures of 30 °C at a current rate of 0.5 C. Furthermore, by enhancing the $m_{\text{areal}}$ (19 mg cm$^{-2}$) of the 0.9 wt.% Gr-containing NCA electrode, we demonstrated a high $Q_{\text{areal}}$ of ~3.7 mAh cm$^{-2}$ and a high $Q_{\text{vol}}$ of ~774 mAh cm$^{-3}$ at 0.2 C. These findings suggest that the use of Gr/PVdF suspensions is a promising solution for advanced Ni-rich oxide cathodes.

4. Conclusions

Figure 6. (a–d) Cross-sectional SEM images of CB- or Gr-containing NCA electrodes with different electrode parameters (Ref., Gr-A, Gr-B, and Gr-C) and (e,f) $Q_{\text{areal}}$ and $Q_{\text{vol}}$ with respect to areal current (at 0.2–10 C).
Supplementary Materials: The following are available online at https://www.mdpi.com/article/10.3390/ma14092271/s1, Figure S1: AFM data of Gr sheets, Figure S2: XPS C 1s of Gr sheets, Figure S3: Raman data of Gr sheets, Figure S4: Digital photographs of dispersions, Figure S5: Voltage profiles of NCA electrodes, Figure S6: Nyquist plots of NCA electrodes.

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References
1. Cano, Z.P.; Banham, D.; Ye, S.; Hintennach, A.; Lu, J.; Fowler, M.; Chen, Z. Batteries and fuel cells for emerging electric vehicle markets. Nat. Energy 2018, 3, 279–289. [CrossRef]
2. Li, M.; Lu, J.; Chen, Z.; Amine, K. 30 Years of Lithium-Ion Batteries. Adv. Mater. 2010, 22, 587–603. [CrossRef]
3. Lim, B.-B.; Myung, S.-T.; Yoon, C.S.; Sun, Y.-K. Comparative Study of Ni-Rich Layered Cathodes for Rechargeable Lithium Batteries: Li[Ni0.85Co0.11Al0.04]O2 and Li[Ni0.94Co0.06Mn0.09Al0.01]O2 with Two-Step Full Concentration Gradients. ACS Energy Lett. 2016, 1, 283–289. [CrossRef]
4. Ritchie, A.G. Recent developments and likely advances in lithium rechargeable batteries. J. Power Sour. 2004, 136, 285–289. [CrossRef]
5. Goodenough, J.B.; Kim, Y. Challenges for Rechargeable Li Batteries. J. Am. Chem. Soc. 2013, 135, 81–90. [CrossRef] [PubMed]
6. Liu, C.; Li, F.; Ma, L.P.; Cheng, H.M. Advanced materials for energy storage. Adv. Mater. 2010, 22, E28–E62. [CrossRef]
7. Schmuch, R.; Wagner, R.; Hörpel, G.; Placke, T.; Winter, M. Performance and cost of materials for lithium-based rechargeable automotive batteries. Nat. Energy 2018, 3, 267–278. [CrossRef]
8. Wang, Y.; Fu, X.; Zheng, M.; Zhong, W.H.; Cao, G. Strategies for Building Robust Traffic Networks in Advanced Energy Storage Devices: A Focus on Composite Electrodes. Adv. Mater. 2019, 31, e1804204. [CrossRef] [PubMed]
9. Bitsch, B.; Dittmann, J.; Schmitt, M.; Scharfer, P.; Schabel, W.; Willenbacher, N. A novel slurry concept for the fabrication of lithium-ion battery electrodes with beneficial properties. J. Power Sour. 2014, 265, 81–90. [CrossRef]
10. Scherlieder, W.; Ivanov, S.; Tran, H.Y.; Theil, S.; Froboese, L.; Westphal, B.; Wohlfahrt-Mehrens, M.; Kwade, A. Influence of formulation method and related processes on structural, electrical and electrochemical properties of LMS/NCA-blend electrodes. Prog. Solid State Chem. 2014, 42, 157–174. [CrossRef]
11. Kraysberg, A.; Ein-Eli, Y. Conveying Advanced Li-ion Battery Materials into Practice The Impact of Electrode Slurry Preparation Skills. Adv. Energy Mater. 2016, 6, 1600655. [CrossRef]
12. Lee, G.-W.; Ryu, J.H.; Han, W.; Ahn, K.H.; Oh, S.M. Effect of slurry preparation process on electrochemical performances of LiCoO2 composite electrode. J. Power Sour. 2010, 195, 6049–6054. [CrossRef]
13. Cho, K.Y.; Kwon, Y.I.; Youn, J.R.; Song, Y.S. Evaluation of slurry characteristics for rechargeable lithium-ion batteries. Mater. Res. Bull. 2013, 48, 2922–2926. [CrossRef]
14. Park, C.W.; Lee, J.-H.; Seo, J.K.; Jo, W.Y.; Whang, D.; Hwang, S.M.; Kim, Y.-J. Graphene collage on Ni-rich layered oxide cathodes for advanced lithium-ion batteries. Nat. Commun. 2021, 12, 2145. [CrossRef] [PubMed]
15. Mattevi, C.; Eda, G.; Aaglini, S.; Miller, S.; Mkhoyan, K.A.; Celik, O.; Mastrogiovanni, D.; Granovitzi, G.; Garfunke, E.; Chhowalla, M. Evolution of Electrical, Chemical, and Structural Properties of Transparent and Conducting ChemicallyDerived Graphene Thin Films. Adv. Funct. Mater. 2009, 19, 2577–2583. [CrossRef]
19. López, V.; Sundaram, R.S.; Gómez-Navarro, C.; Olea, D.; Burghard, M.; Gómez-Herrero, J.; Zamora, F.; Kern, K. Chemical Vapor Deposition Repair of Graphene Oxide: A Route to Highly-Conductive Graphene Monolayers. *Adv. Mater.* 2009, 21, 4683–4686. [CrossRef]

20. Karan, S.K.; Mandal, D.; Khatua, B.B. Self-powered flexible Fe-doped RGO/PVDF nanocomposite: An excellent material for a piezoelectric energy harvester. *Nanoscale* 2015, 7, 10655–10666. [CrossRef] [PubMed]

21. Ataur Rahman, M.; Chung, G.-S. Synthesis of PVDF-graphene nanocomposites and their properties. *J. Alloy. Compd.* 2013, 581, 724–730. [CrossRef]

22. Zhang, X.-J.; Wang, G.-S.; Cao, W.-Q.; Wei, Y.-Z.; Cao, M.-S.; Guo, L. Fabrication of multi-functional PVDF/RGO composites via a simple thermal reduction process and their enhanced electromagnetic wave absorption and dielectric properties. *RSC Adv.* 2014, 4, 19594–19601. [CrossRef]

23. El Achaby, M.; Arrakhiz, F.Z.; Vaudreuil, S.; Essassi, E.M.; Qaiss, A. Piezoelectric β-polymorph formation and properties enhancement in graphene oxide—PVDF nanocomposite films. *Appl. Surf. Sci.* 2012, 258, 7668–7677. [CrossRef]

24. Huang, L.; Lu, C.; Wang, F.; Wang, L. Preparation of PVDF/graphene ferroelectric composite films by in situ reduction with hydrobromic acids and their properties. *Rsc Adv.* 2014, 4, 45220–45229. [CrossRef]

25. Xu, X.-L.; Yang, C.-J.; Yang, J.-H.; Huang, T.; Zhang, N.; Wang, Y.; Zhou, Z.-W. Excellent dielectric properties of poly(vinylidene fluoride) composites based on partially reduced graphene oxide. *Compos. Part B Eng.* 2017, 109, 91–100. [CrossRef]

26. Han, J.; Li, H.; Kong, D.; Zhang, C.; Tao, Y.; Li, H.; Yang, Q.-H.; Chen, L. Realizing High Volumetric Lithium Storage by Compact and Mechanically Stable Anode Designs. *ACS Energy Lett.* 2020, 5, 1986–1995. [CrossRef]

27. Zhang, C.; Lv, W.; Tao, Y.; Yang, Q.-H. Towards superior volumetric performance: Design and preparation of novel carbon materials for energy storage. *Energy Environ. Sci.* 2015, 8, 1390–1403. [CrossRef]

28. Gaberscek, M.; Moskon, J.; Erjavec, B.; Dominko, R.; Jamnik, J. The Importance of Interphase Contacts in Li Ion Electrodes: The Meaning of the High-Frequency Impedance Arc. *Electrochem. Solid State Lett.* 2008, 11, A170. [CrossRef]

29. Thorat, I.V.; Joshi, T.; Zaghib, K.; Harb, J.N.; Wheeler, D.R. Understanding Rate-Limiting Mechanisms in LiFePO4 Cathodes for Li-Ion Batteries. *J. Electrochem. Soc.* 2011, 158, A1185–A1193. [CrossRef]

30. Hwang, S.M.; Lim, Y.-G.; Kim, J.-G.; Heo, Y.-U.; Lim, J.H.; Yamauchi, Y.; Park, M.-S.; Kim, Y.-J.; Dou, S.X.; Kim, J.H. A case study on fibrous porous SnO2 anode for robust, high-capacity lithium-ion batteries. *Nano Energy* 2014, 10, 53–62. [CrossRef]

31. Li, W.; Erickson, E.M.; Manthiram, A. High-nickel layered oxide cathodes for lithium-based automotive batteries. *Nat. Energy* 2020, 5, 26–34. [CrossRef]

32. Myung, S.-T.; Maglia, F.; Park, K.-J.; Yoon, C.S.; Lamp, P.; Kim, S.-J.; Sun, Y.-K. Nickel-Rich Layered Cathode Materials for Automotive Lithium-Ion Batteries: Achievements and Perspectives. *ACS Energy Lett.* 2016, 2, 196–223. [CrossRef]

33. Bak, S.-M.; Nam, K.-W.; Chang, W.; Yu, X.; Hu, E.; Hwang, S.; Stach, E.A.; Kim, K.-B.; Chung, K.Y.; Yang, X.-Q. Correlating Structural Changes and Gas Evolution during the Thermal Decomposition of Charged Li(x)Ni0.8Co0.15Al0.05O2 Cathode Materials. *Chem. Mater.* 2013, 25, 337–351. [CrossRef]

34. Kim, J.; Lee, H.; Cha, H.; Yoon, M.; Park, M.; Cho, J. Prospect and Reality of Ni-Rich Cathode for Commercialization. *Adv. Energy Mater.* 2018, 8, 1702028. [CrossRef]

35. Park, K.Y.; Lim, J.M.; Luu, N.S.; Downing, J.R.; Wallace, S.G.; Chaney, L.E.; Yoo, H.; Hyun, W.J.; Kim, H.U.; Hersam, M.C. Concurrently Approaching Volumetric and Specific Capacity Limits of Lithium Battery Cathodes via Conformal Pickering Emulsion Graphene Coatings. *Adv. Energy Mater.* 2020, 10, 2001216. [CrossRef]