Electronic Supplementary Information

Crumpled N-doped Carbon Nanotubes Encapsulated with Peapod-like Ge Nanoparticles for High-rate and Long-life Li-ion Battery Anodes

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Experimental details

Synthesis of Inorganic-organic hybrid GeO\textsubscript{x}/EDA nanowires

GeO\textsubscript{x}/EDA nanowires were prepared by a facile and low-cost solvothermal process\textsuperscript{1,2}. In a typical procedure, 1 g GeO\textsubscript{2} and 0.25 g Fe\textsubscript{2}O\textsubscript{3} were mixed together and then transferred to a 50 ml Teflon-lined stainless-steel autoclave filled with 8 ml H\textsubscript{2}O and 5 ml EDA. After hydrothermal treatment at 200 °C for 60 h, the products were collected, then thoroughly washed with ethanol for three times, and finally vacuum dried at 60 °C for 3 h.

Synthesis of 1D peapod-like Ge/CN\textsubscript{x} nanomaterials

The as-synthesized GeO\textsubscript{x}/EDA nanowires were annealed in the air at 500 °C for 2 h to remove the EDA. The as-synthesized GeO\textsubscript{2} nanowires were further coated with polypyrrole (PPy) and then thermally treated under Ar/H\textsubscript{2} at 850 °C to produce peapod-like Ge/CN\textsubscript{x} nanomaterials. In a typical synthesis, 300 mg GeO\textsubscript{2} nanowires was firstly dispersed in anhydrous methylacetate (20 mL) containing 100 mg anhydrous FeCl\textsubscript{3} as an oxidant and 20 mg poly(vinyl acetate as a stabilizing agent. Then monomer pyrrole (20 μL; 0.97 mg/μL) was added dropwise. After stirring for overnight reaction at room temperature, the PPy coated GeO\textsubscript{2} nanowires were collected and washed three times with methyl acetate. After vacuum dried at 60 °C overnight, the GeO\textsubscript{2}@PPy nanowires were thermally treated in a tube furnace under Ar/H\textsubscript{2} atmosphere (10% H\textsubscript{2}) at 850 °C for 2 h with the heating ramp of 5 °C min\textsuperscript{-1}.

Structural and Electrochemical Characterizations

The morphology, structures and composition of samples were characterized by X-ray diffraction (GAXRD, Philips X’Pert Pro), field-emission scanning electron microscopy (FE-SEM, FEI nanoSEM 450), transmission electron microscopy (TEM, FEI Titan 60-300Cs), high-resolution
TEM (HR-TEM, JEM-2010F) and energy-dispersive X-ray spectroscopy (EDS, Oxford INCA 200, Oxford Instruments, and Oxfordshire, U.K.). XRD were measured using a Philips X’ Pert Pro diffractometer with copper Kα (λ = 1.5416 Å) radiation. X-ray photoelectron spectroscopy (XPS) measurements were conducted on a Kratos Axis Ultra DLD spectrometer using a monochromated Al Kα radiation. Raman scattering measurement was measured in LabRam HR with Ar+ laser excitation in back scattering geometry (514.5 nm). The TG results were obtained via TGA (NETZSCH; TG 209 F3) measurements.

The Li storage properties of Ge/CNx peapods are investigated in the half cell in an electrolyte consisting of 1:1 vol/vol mixture of ethylene carbonate (EC) and diethyl carbonate (DEC) with 5 wt.% Fluorinated ethylene carbonate (FEC). To prepare working electrodes, peapod-like Ge/CNx, poly(vinyl difluoride) (PVDF, Sigma Aldrich) and carbon black (conducting agent) with the ratio of 80:10:10 were mixed in N-Methyl pyrrolidone (NMP, Sigma Aldrich) forming a slurry and then pasted on a copper foil. The mass loading of Ge active materials is typically about 1.5 mg/cm². A Celgard 2400 film was used as the separator and pure Li foil as the counter electrode. The 2025 coin-like cells were assembled in an argon-filled glove box. The charge and discharge measurements of the batteries were conducted with an MTI automatic battery cycler with the voltage cutoffs between 0.01 and 1.5 V vs Li/Li⁺. The electrochemical impedance spectroscopy (EIS) tested was carried out on Autolab Electrochemical workstation from 10 mHz to 10⁵ Hz.
Figure S1. FESEM image (a) and the corresponding XRD pattern (b) of a single inorganic-organic hybrid GeO$_x$/EDA nanowires with rectangular cross section. The XRD pattern in Fig. S1b is the same as the one acquired from previously-reported GeO$_x$/EDA, $^{1,2}$ suggesting the as-synthesized nanowires are hybrid GeO$_x$/EDA nanowires.

Figure S2. TG curve of inorganic-organic hybrid GeO$_x$/EDA nanowires under Air.
Figure S3. FESEM image and corresponding XRD pattern of GeO₂ nanowires.

Figure S4. FE-SEM (a), TEM (b) and FTIR spectrum (c) of PPy coated GeO₂ nanowires. The thickness of PPy shell is about 15-20 nm. In Figure c, the IR absorption peaks centered at 549.6 and 874.9 cm⁻¹ band indicate the existence of GeO₂ and the peaks at 3430.2, 1548.7, 1450.4 and 1253.7 cm⁻¹ confirm the presence of PPy shell.
Figure S5. Different magnified SEM (a, b) and TEM(c, d) images for Ge/CNx nanopeapods, confirming that the isolated NPs are incorporated into the nanotubular shells resembling peapods.

Figure S6. Raman Spectrum of Ge/CNx in the range of 800-1900 cm\(^{-1}\).

Different from the Raman spectrum of carbon nanotubes with two distinct D and G bands, \(^3\)\(^4\) Raman scattering spectrum in Fig. 2c demonstrates a dramatic broadening and overlap of D (1350) and G (1850 cm\(^{-1}\)) bands for CNx shell. The peak
at G-band is attributed to the vibration of sp\(^2\)-bonded carbon atoms in a 2D hexagonal lattice, namely, the stretching modes of C-C bonds of typical graphite, while the peak at D-band is associated with vibrations of carbon atoms with dangling bonds in plane terminations of the disordered graphite and is related to the defects and disorders of structures in carbon materials. The presence of N in the CN\(_x\) shell result in an asymmetric tailing of D band extending out to about 1000 cm\(^{-1}\) as well as two broadening and overlap of D and G bands. In our case, the I\(_D\)/I\(_G\) in the Raman scattering spectrum is estimated to be ~0.95, meaning that the CN\(_x\) nanotubes is partially graphitized, which is in accordance with previous reported results.\(^3\)\(^-\)\(^5\)

**Figure S7.** The X-ray photoelectron spectroscopy (XPS) results of Ge/CN\(_x\) nanopeapods. (a) The high-resolution N 1s spectrum can be deconvoluted into pyridinic N (398.2 eV), pyrrolic N (400.1 eV), and graphitic N (401.1 eV); (b) Fine XPS of C 1s; (c) The fine XPS of Ge 2p shows an intense peak at 1217.4 and weak peak at 1219.8 eV, corresponding to the Ge and GeO\(_2\), respectively.
Figure S8. TG profile for peapod-like Ge/CNx. Assuming the Ge was fully transformed into GeO₂, the weight ratio of CNₓ in Ge/CNx peapods is calculated to be 18.2%.

Figure S9. Nitrogen adsorption-desorption isotherms and pore distribution of peapod-like Ge/CNx. The measured BET surface area is about 19.5 m² g⁻¹.
**Figure S10.** Delithiation capacity of the Ge/CN\textsubscript{x} nanopeapods as a function of cycle numbers based on the mass loading of Ge active materials, revealing that the capacity is as high as 1200 mA h g\textsuperscript{-1} after initial 5 cycles at a 0.5 C rate.

**Fig. S11.** Cycling performance of Ge/CN\textsubscript{x} nanopeapods at high rates of 8 and 10 C. The reversible capacity at a 10 C rate is measured to be about 810 mA h g\textsuperscript{-1}. 


Figure S12. The differential plots (dQ/dV vs. V) for the 1\textsuperscript{st}, 5\textsuperscript{th} and 50\textsuperscript{th} cycles. The three peaks at 0.17, 0.36, and 0.49 V are related to the different Li\textsubscript{x}Ge alloys formed during lithiation. The peaks between 0.3 and 0.7 V in the oxidation scan can be ascribed to delithiation of Li\textsubscript{x}Ge.

Figure S13. Low-magnification TEM image of the Ge/CN\textsubscript{x} nanopeapods after 1200 cycles
Figure S14. The EIS results of the Ge/CN\textsubscript{x} after the 1\textsuperscript{st} and 1200\textsuperscript{th} cycle and the corresponding equivalent circuit. The semicircle at high frequencies can be assigned to the SEI resistance and contact resistance (R\textsubscript{f}) and the charge transfer impedance on electrode–electrolyte interface (R\textsubscript{ct}). After 1200\textsuperscript{th} cycle, the R\textsubscript{f} slightly enlarges from 101 to 119 Ω while the R\textsubscript{ct} decreases from 98 to 30 Ω, suggesting stable SEI and structural stability of Ge/CN\textsubscript{x} for the continuous cycles.
Table S1. Comparison of electrochemical performance found in the present work with those of reported Ge-based materials.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Cycle Stability</th>
<th>Rate performance</th>
<th>$R_{ct}$ (Ω)</th>
<th>Ref.</th>
</tr>
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<tr>
<td>C-N/Ge</td>
<td>906. mAh g$^{-1}$@50 cycles@600 mAg$^{-1}$</td>
<td>500 mAh g$^{-1}$@4A g$^{-1}$</td>
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<tr>
<td>Ge/RGO/C</td>
<td>993 mAh/g@600cycles@1000 mAg$^{-1}$</td>
<td>750 mAh g$^{-1}$@9.6A g$^{-1}$</td>
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<tr>
<td>Ge@C-N</td>
<td>813 mAh g$^{-1}$@90cycles@500 mA g$^{-1}$</td>
<td>406 mAh g$^{-1}$@3A g$^{-1}$</td>
<td>23.3</td>
<td>8</td>
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<tr>
<td>Ge@C</td>
<td>328.6mAh g$^{-1}$@90cycles@500 mA g$^{-1}$</td>
<td>200 mAh g$^{-1}$@3A g$^{-1}$</td>
<td>45.7</td>
<td>8</td>
</tr>
<tr>
<td>P-Ge@C</td>
<td>1099mAh g$^{-1}$@100cycles@160mA g$^{-1}$</td>
<td>708 mA h g$^{-1}$@1.6Ag$^{-1}$</td>
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<tr>
<td>Ge-C</td>
<td>900mAh g$^{-1}$@50cycles@150mA g$^{-1}$</td>
<td>613 mAh g$^{-1}$@0.9Ag$^{-1}$</td>
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<tr>
<td>Pc-Ge NWs</td>
<td>789mAh g$^{-1}$@50cycles@160mA g$^{-1}$</td>
<td>450 mAh g$^{-1}$@1.6Ag$^{-1}$</td>
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<tr>
<td>Ge/N-RGO</td>
<td>700mAh g$^{-1}$@200cycles@500mA g$^{-1}$</td>
<td>210 mAh g$^{-1}$@10Ag$^{-1}$</td>
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<tr>
<td>Hollow Ge@C</td>
<td>1137 mA h g$^{-1}$@200cycles@320mAg$^{-1}$</td>
<td>360 mA h g$^{-1}$@40Ag$^{-1}$</td>
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<tr>
<td>Ge–MWCNT</td>
<td>800 mA h g$^{-1}$@200cycles@320mAg$^{-1}$</td>
<td>490 mA h g$^{-1}$@8.115Ag$^{-1}$</td>
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<tr>
<td>Ge-CN$_x$</td>
<td>1080mAh g$^{-1}$@1200cycles@800 mAg$^{-1}$</td>
<td>874 mA h g$^{-1}$@12.8Ag$^{-1}$</td>
<td>30</td>
<td>This work</td>
</tr>
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</table>

References


