High-Performance Quasi-Solid-State MXene-Based Li–I Batteries

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ABSTRACT: Lithium–iodine (Li–I) batteries have attracted tremendous attention due to their high energy and power densities as well as the low cost of iodine. However, the severe shuttle effect of iodine species and the uncontrollable lithium dendrite growth have strongly hindered their practical applications. Here we successfully develop a quasi-solid-state Li–I battery enabled by a MXene-based iodine cathode and a composite polymer electrolyte (CPE) containing NaNO₃ particles dispersing in a pentaerythritol-tetraacrylate-based (PETEA-based) gel polymer electrolyte. As verified by experimental characterizations and first-principle calculations, the abundant functional groups on the surface of MXene sheets provide strong chemical binding to iodine species, and therefore immobilize their shuttling. The PETEA-based polymer matrix simultaneously suppresses the diffusion of iodine species and stabilizes the Li anode/CPE interface against dendrite growth. The NaNO₃ particles act as an effective catalyst to facilitate the transformation kinetics of LiI₃ on the cathode. Owing to such synergistic optimization, the as-developed Li–I batteries deliver high energy/power density with long cycling stability and good flexibility. This work opens up a new avenue to improve the performance of Li–I batteries.

INTRODUCTION

In modern society, the development of new-generation rechargeable batteries with high energy density, high power density, and low cost is highly desired due to soaring demand for utilizing renewable energy and reducing air pollution.† As a promising alternative to lithium-ion (Li-ion) batteries, rechargeable lithium–iodine (Li–I) batteries based on a conversion mechanism (I₂ + 2Li → 2LiI) have drawn intensive attention owing to their high theoretical capacity (1040 mA h cm⁻² and 211 mA h g⁻¹), superior rate performance, degassing-free battery chemistry, and the low cost of iodine.‡,§ However, currently, the development of Li–I batteries has been stunted by many intrinsic obstacles. Iodine-based electrodes usually suffer from poor electrical conductivity of the elemental iodine (I₂).¶ More severely, iodine species (i.e., I₂, I₃⁻, and I⁻) are highly dissoluble in organic electrolyte solvents. This not only results in serious self-discharge due to their shuttle effect, but also leads to increased internal resistance together with decreased Li⁺ transport kinetics.¶‖ Furthermore, uncontrollable dendrite growth on the Li anode during cycling deteriorates the Li/electrolyte interface and causes safety hazards.¶⊥

To improve the performance of Li–I batteries, many efforts have been devoted to confine iodine in carbon (C) matrices (such as graphene,¶‴ carbon microtubes,¶⊥ porous carbon,¶⊥⊥ and heteroatom-doped carbon¶⊥⊥⊥) with the aim of enhancing electronic conductivity and suppressing the dissolution/diffusion of iodine species. However, the physical adsorption of iodine species in these carbon matrices is insufficient to prohibit the shuttle effect. Therefore, it is highly desired to develop a novel iodine-based cathode simultaneously possessing high iodine loading and strong chemical binding of iodine species. As for the electrolyte, replacing organic liquid electrolyte (LE) with solid electrolyte can synergistically restrain the diffusion of dissolved iodine species and eliminate safety issues (fire and explosion, etc.), which could result from the leakage of flammable electrolyte solvents.¶‴ However, the low ionic conductivity of solid electrolyte and the unstable electrode/solid electrolyte interface greatly limit their applications in Li–I batteries.

Supporting Information

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Recently, we have proposed a versatile technique for the preparation of MXene-based electrode materials, and an intensive investigation on the liquid/polymer electrolytes for conversion mechanism-based alkali metal batteries. Based on these, for the first time, we successfully develop a flexible quasi-solid-state Li-I battery integrated with a Ti₃C₂Tₓ (T represents surface functional group, and x is number of such groups) MXene-wrapped carbon cloth–iodine (CC–I/MXene) cathode and a composite polymer electrolyte (CPE) composed of catalytic NaN₃ particles dispersing in a pentaerythritol-tetraacrylate-based (PETEA-based) gel polymer electrolyte. As verified by theoretical calculations and experimental investigations, the Ti₃C₂Tₓ MXene sheets with abundant surface functional groups can effectively confine the iodine species inside the cathode, and therefore restrain the shuttle effect. In the in situ prepared CPE, the PETEA-based polymer matrix can efficiently suppress the diffusion of iodine species, and benefit for the formation of a stable Li anode/CPE interface without dendrite growth. Meanwhile, the NaN₃ particles not only greatly enhance the mechanical strength of CPE, but also effectively increase the kinetic transformation of LiI in the cathode. The as-developed quasi-solid-state Li–I batteries exhibit a high energy density with stable cycling performance and excellent flexibility.

**RESULTS AND DISCUSSIONS**

Figure 1 illustrates the preparation of the Li|CPE|CC–I/MXene batteries. Iodine can be facilely loaded onto the porous carbon cloth (CC) by a simple “solution—adsorption” method to obtain the flexible carbon cloth–iodine (CC–I) electrodes. After the iodine-absorption, the specific surface area of carbon cloth significantly decreased, indicating a successful loading of iodine into the porous structure of carbon cloth (shown in Figure S1). Delaminated Ti₃C₂Tₓ MXene was prepared by selectively etching Al atoms in Ti₃AlC₂ (MAX) precursor materials followed by sonication (Figure S2). By adopting a “drop-filtration” process, the CC–I electrodes can be tightly wrapped by the MXene sheets (Figure 1, upper panels), which can greatly improve the stability of iodine due to the strong interaction (Figure S3).

To further enhance the electrochemical performance of CC–I/MXene cathodes in batteries, optimization of electrolyte has been conducted. Based on previous research, it is believed that the addition of LiNO₃ in ether-based electrolytes benefits for the formation of a passive film on the Li anode, and therefore suppresses its corrosion from dissolved iodine species. However, as shown in Figure S4, a Li anode precycled in a LiNO₃-containing liquid electrolyte (LE) cannot suppress the serious self-discharge, while Li–I cells using a suspension of insoluble NaN₃ in LE can deliver a negligible self-discharge, which is comparable to that of the LiNO₃-containing LE. Therefore, we present a new understanding on the role of nitrates in Li–I batteries. We suggest that NaN₃ can catalyze the conversion from I₃⁻ to elemental I₂ near the end of the charging process. Thus, we prepared a composite polymer electrolyte consisting of 2 wt % NaN₃ homogeneously dispersing in a PETEA-based gel polymer electrolyte. First, NaN₃ particles were dispersed in 1 M bis-(trifluoromethane) sulfonamide lithium (LiTFSI) in 1,2-dioxolane (DOL):dimethoxymethane (DME) (1:1, v/v) electrolyte (in which a small amount of PVDF-HFP (poly-(vinylidene fluoride-hexafluoropropylene)) was dissolved to

![Figure 1. Schematic illustration of the preparation of MXene-wrapped carbon cloth–iodine cathodes and composite polymer electrolyte.](image-url)
improve the dispersion of NaNO₃ particles] to form a suspension, and then PETEA monomer together with 2-hydroxy-2-methyl-1-phenyl-1-propanone (HMPP) photoinitiator were dissolved into the suspension to obtain a precursor solution. After that, the precursor solution was subjected to ultraviolet-light-irradiation (UV-irradiation) to initiate a radical polymerization of C≡C bonds on the monomers, and thereafter the cross-linked CPE was in situ constructed on the surface of the CC−I/MXene cathode (Figure 1, lower panels). In such CPE, the PETEA-based polymer matrix with strong chemical interaction with iodine species can efficiently suppress their shuttle effect and simultaneously stabilize the Li anode/CPE interface. Furthermore, NaNO₃ particles can not only greatly enhance the mechanical strength of the CPE (Figure S5), but also act as an effective catalyst on the iodine cathode/CPE interface. Such collaborative optimization on electrode and electrolyte is expected to achieve a superior cycling stability for the Li−I batteries.

As shown in Figure S6a, the carbon cloth shows a morphology of tidily knitted carbon fibers. No structural change is observed after absorbing iodine (Figure S6b). For the CC−I/MXene electrode, it is clearly seen that the iodine-loaded carbon fibers are well-wrapped by the MXene sheets (Figure 2a and Figure S7a,b). This is further confirmed by the uniform distribution of elemental titanium (Ti), carbon (C), and iodine (I) in the energy dispersive spectrometer (EDS) images of a CC−I/MXene electrode (Figure 2b). Furthermore, no characteristic peak of iodine can be observed in the X-ray diffraction (XRD) pattern (Figure S7c) and Raman spectrum (Figure S7d) of the MXene-wrapped electrode, indicating the formation of amorphous iodine during the solution−adsorption process, which is consistent with previous reports.¹³ X-ray photoelectron spectroscopy (XPS) was applied to investigate the chemical composition of the CC−I/MXene electrode. The Ti 2p spectrum (Figure 2c) shows two Ti 2p₃/2 doublets at 455.1 and 458.7 eV, and two Ti 2p₁/2 doublets at 461.1 and 464.4 eV, respectively. As for the Ti 2p₃/2, three deconvoluted peaks at 455.1, 456.5, and 458.7 eV can be assigned to Ti−C bonds, Ti−X bonds, and Ti−O surface bonds, respectively. Similarly, three deconvoluted peaks at 460.9, 461.9, and 464.4 eV can be observed in the Ti 2p₁/2.¹⁶,²¹ Such a Ti 2p spectrum result confirms the successful wrapping of MXene sheets on the electrode without any compositional change. Figure 2d exhibits the XPS spectrum of I 3d in the CC−I/MXene electrode, in which peaks located at around 619.9 and 631.2 eV are associated with I 3d₅/2 and I 3d₃/2, respectively.²² Noticeably, a peak located at around 623 eV can be ascribed to the chemical adsorption between iodine and the

Figure 2. Characterizations of CC−I/MXene electrodes. (a) FE-SEM images and (b) EDS mappings of elemental Ti, C, and I on a CC−I/MXene electrode. High-resolution XPS spectra of (c) Ti 2p and (d) I 3d in a CC−I/MXene electrode. (e) UV−vis spectra and corresponding digital images of LiI₃ solution before and after loading MXene sheets. (f) Initial discharge/charge curves of LiI/CC−I and LiI/CC−I/MXene cells at 0.5 C.
−O/−OH terminations of MXene. Such chemical bonding is further validated by the O 1s XPS spectrum (Figure S8). The interaction between iodine and the −O/−OH terminations is expected to confine the iodine in the cathode and suppress its dissolution. To estimate the adsorption ability of MXene for iodides, ultraviolet−visible (UV−vis) spectra of a 2 mL 0.005 M LiI3 (prepared by dissolving LiI and I2 with a stoichiometric ratio of 1:1) DOL solution were measured. As shown in Figure 2e, after adding 1 mg of MXene, the characteristic peaks of I3− at around 286 and 365 nm decrease gradually with the aging time. After 2 days adsorption, the solution turned clear (the inset in Figure 2e), which illustrates a strong adsorption of iodides by the MXene. The initial discharge/charge curves of Li−I cells using CC−I and CC−I/MXene electrodes in nitrate-free LE (1 M LiTFSI in DOL/DME) are shown in Figure 2f. The CC−I cathodes show extremely low Coulombic efficiency and cannot be charged to above 3 V, which can be attributed to the severe shuttling of dissolvable iodine species. In sharp contrast, after wrapping with MXene sheets, the CC−I/MXene cathodes exhibit a significantly enhanced Coulombic efficiency (∼73%) and reversible capacity (∼219 mA h g−1), indicating an effective immobilization of iodine species by the MXene sheets.

The Fourier transform infrared spectroscopy (FTIR) spectra of the PETEA monomer and the PETEA-based polymer matrix separated from CPE are shown in Figure S9, which verifies that the monomers have been successfully polymerized in the NaNO3 suspension with a high degree of conversion. The ionic conductivities of the LE, LE suspended with NaNO3, and CPE as a function of temperature from 0 to 90 °C are shown in Figure 3a. The plots of log σ versus T−1 for all electrolyte samples deliver a nonlinear relationship, which can be well-described by the Vogel−Tamman−Fulcher (VTF) empirical equation:24

$$\sigma = \sigma_0 T^{-1/2} \exp \left( -\frac{E_a}{RT T_0} \right)$$

where $E_a$ is the activation energy, $\sigma_0$ is the pre-exponential factor, $R$ is the ideal gas constant, and $T_0$ is a parameter correlated to the glass transition temperature. It is worth noting that the $E_a$ value (which indicates the barrier for ion
The Li-ion transference number ($t_{Li^+}$) is a crucial parameter for electrolytes, since a low $t_{Li^+}$ results in an increase in electrode polarization. It is seen that the $t_{Li^+}$ of CPE reaches 0.52 (Figure 3b), which is much higher than that of the LE (0.23, Figure S10). This can be ascribed to the fact that the polymerized PETEA framework significantly immobilizes the anions. Such enhanced $t_{Li^+}$ is expected to not only diminish the polarization of batteries, but also restrain the dendrite growth on Li anodes. This is confirmed by galvanostatic cycling measurements on a symmetric Li|Li cell at a current density of 0.5 mA cm$^{-2}$. For the Li|LELi cell, the voltage hysteresis between the voltages of Li stripping and plating dramatically increases after ~240 h, demonstrating a deteriorated Li|LE interface, resulting from an accumulated thick solid electrolyte interface (SEI) and Li dendrite growth (Figure 3c). The LiCPEiLi cell, in sharp contrast, delivers a much lower voltage fluctuation and maintains stability up to 300 h. This validates a uniform Li deposition with a stable Li/CPE interface. Therefore, the safety risks caused by dendrite growth have been successfully mitigated.

The durability and safety of the CPE are elucidated by measuring its weight loss in an open environment as a function of aging time at room temperature. As shown in Figure 3d, the CPE exhibits a slight weight loss with aging time (17 wt % after 12 h) due to the volatilization of electrolyte solvent, which is much lower than that of the LE (61 wt % after 12 h). Therefore, the CPE shows a much enhanced safety characteristic compared to LE.

To investigate the electrochemical characteristics of the as-assembled Li−I batteries, the performances of CC−I/MXene electrodes using LE, NaNO$_3$-containing LE, and CPE were tested. As shown from the self-discharge phenomena (Figure S11), the potential of the Li|LE|CC−I/MXene cell decreases much slower after adding the iodine (17 wt % after 12 h) due to the volatilization of electrolyte solvent, which is much lower than that of the LE (61 wt % after 12 h). Therefore, the CPE shows a much enhanced safety characteristic compared to LE. To directly characterize the dissolution and diffusion of iodine species in electrolytes, we performed visual observation on the same amount of iodine powder soaked in LE and CPE. The LE immediately turns dark brown after adding the iodine (Figure 3e, left), implying that the iodine powder massively dissolves in the LE. However, the CPE remains milky white with a light-yellow skin layer during the aging process (Figure 3e, right), indicating that the dissolution of iodine has been significantly alleviated in the CPE.
rapidly due to the dissolution of active material and shuttle effect, while the cell with CPE achieves a negligible self-discharge after aging for 48 h, indicating the strong immobilization of iodine species by the CPE. The LiI/LE CC−1/MXene cell can deliver an initial discharge capacity of 219 mA h g$^{-1}$ (based on the mass of iodine) at 0.5 C with a Coulombic efficiency of 73% (Figure 4a). It should be noticed that such a capacity value is higher than the theoretical capacity of Li−I batteries (211 mA h g$^{-1}$), since the MXene sheets as well as porous carbon cloth substrates can also impart capacity due to the capacitive behavior (Figure S12). After adding NaNO$_3$ into the LE, an enhanced capacity (301 mA h g$^{-1}$) and Coulombic efficiency (∼92%) are achieved at 0.5 C, suggesting that NaNO$_3$ can greatly enhance the transformation kinetics due to catalytic effect (Figure S13a). The capacity of CC−1/MXene electrodes in Li−I batteries can be further improved to ∼330 mA h g$^{-1}$ at 0.5 C by using CPE, since the cross-linked monomers in electrolyte can effectively immobilize the iodine species and prevent loss of active materials. The LiIPEICCC−1/MXene cells also exhibit superior long-term cycling stability with 85% capacity retention after 1000 cycles, which is much higher than those with LE (57%, 50 cycles) and NaNO$_3$-containing LE (67%, 1000 cycles) (Figure 4b and Figure S13b).

Cyclic voltammetry (CV) was also performed for understanding the chemistry of LiIPEICCC−1/MXene, LiI/LE+NaNO$_3$1 CC−1/MXene, and LiIPEICCC−1/MXene cells. As shown in Figure 4c and Figure S13c, a Li−I cell with the NaNO$_3$-containing LE exhibits enhanced current density and reduced potential gap between the main anodic and cathodic peaks than that of the cell with the LE, which indicates the catalytic effect of NaNO$_3$ particles. Moreover, the Li−I cell using CPE shows the highest peak current density, signifying a lower polarization contributed by the synergistic optimization of the catalytic NaNO$_3$, and PETEA-based polymer matrix. For the LiIPEICCC−1/MXene cells, the capacity contribution from different behaviors can be studied by analyzing the CV curves at different scan rates. By applying the capacity separation method,$^6$ the noncapacitive capacity contribution can be obtained as displayed in the shadowed area of the CV curve (Figure S14). It is seen that, during the cathodic scan, a small hump emerges at ∼3.3 V vs Li/Li$^+$, indicating the conversion of I$_2$ to I$_3^−$. The peak at ∼2.8 V can be assigned to the reaction from I$_3^−$ to I$^−$. The anodic scan is a reversible process, in which the peak at ∼3.0 V indicates that I$^−$ was oxidized to I$_3^−$, and I$_3^−$ was further oxidized to I$_2$ at ∼3.3 V.$^4$ Noticeably, the current sharply increased from ∼3.5 V vs Li/Li$^+$, which could be attributed to the oxidation of the NO$_3^−$ anions, as described in the following equation:

$$\text{NO}_3^− \rightarrow e \rightarrow \text{NO}_3^− $$ (2)

Such NO$_3^−$ has been reported to widely exist in nature with a short life-span$^{28,29}$ and usually forms as a reaction intermediate in electrochemical synthesis.$^{30}$ It is speculated that, in Li−I batteries, the NO$_3^−$ radicals tend to be consumed instantly by the I$^−$ anions as described in the following equation:

$$2\text{NO}_3^− + 2\text{I}^− \rightarrow 2\text{NO}_3^− + 3\text{I}_2$$ (3)

Thus, NaNO$_3$ as the catalyst can effectively enhance the conversion kinetics and the transformation of I$^−$ to I$_2$. This is well consistent with the corresponding ex situ Raman spectra in Figure S15a, in which the Raman peak of I$^−$ (at ∼120 cm$^{-1}$)$^{31,32}$ disappears at the end of the charge process (3.6 V), demonstrating a complete oxidation of I$^−$ to I$_2$. For comparison, due to poor kinetics, the peak of I$^−$ in the LiI/LECC−1/MXene cell still exists at the end of the charge process, which means that I$^−$ cannot fully revert to I$_2$ (Figure S15b), resulting in the limited Coulombic efficiency as shown in Figure 4a.

Electrochemical impedance spectroscopy (EIS) was conducted to further investigate the interfacial behavior and reversibility of Li−I cells. Figure S16a,b presents the EIS results for Li−I cells after different cycles at 0.5 C. The EIS spectra can be well-simulated via an equivalent circuit as shown in Figure S16c, and the simulation results are summarized in Table S2. For the LiIPEICCC−1/MXene cell, the solid electrolyte interface resistance ($R_i$) sharply rises from 98.1 to 187.2 Ω after 50 cycles, which correlates with the unstable SEI film and dendrite growth on the anode as well as the severe shuttle effect in the cathode.$^{35}$ In sharp contrast, after replacing the LE with CPE, the values of $R_i$ are much lower (21.7 Ω after 1 cycle), and maintain steady with limited variation during cycling (40.0 Ω after 50 cycles). This implies an inhibition of the shuttle of iodine species and a stable electrode/CPE interface, which contribute to the significantly improved cycling performance in Figure 4b. The Li anodes disassembled from Li−I cells after 50 cycles were observed by FE-SEM. Massive dendrite structures and microsized precipitates can be found on the surface of the Li anode disassembled from the LiIPEICCC−1/MXene cell, and the iodine content on such an anode is as high as 13.71 wt % (Figure 4d, corresponding to the digital image in Figure S17). The Li anode from the LiI/LE+NaNO$_3$/CC−1/MXene cell shows an iodine content of 6.92 wt % (Figure S13e). With the CPE, the surface of the Li anode tends to be smooth, and the growth of Li dendrite has been remarkably suppressed. The iodine content of this Li anode is as low as 1.25 wt % (Figure 4e), confirming the successful suppression of the shuttle effect.

The rate performances of Li−I batteries are shown in Figure 4f and Figure S13d. Meanwhile, the corresponding voltage profiles are presented in Figure S18. The LiIPEICCC−1/MXene cell delivered specific capacities of 358, 332, 303, 268, and 227 mA h g$^{-1}$ at 0.2, 0.5, 1, 2, 5 C, respectively. Even at a high current density of 10 C, the cell can still maintain a capacity of 182 mA h g$^{-1}$, which is higher than LiIPEICCC−1/MXene and LiI/LE+NaNO$_3$/CC−1/MXene cells. These values are also higher than those of LiIPEICCC−1 cells (Figure S19), and comparable to those of Li−I cells using the CC−1/MXene electrode and LiNO$_3$-containing polymer electrolyte (Figure S20). Furthermore, the charge/discharge potential gap of the cell with CPE is smaller than that of the cell with LE, which agrees very well with CV curves in Figure 4c. LiIPEICCC−1/MXene cells with different areal mass loadings of iodine in the cathodes were further cycled at 0.5 C. With high iodine loadings of 7.5 and 10 mg cm$^{-2}$, the cell still can achieve capacities of 243 and 160 mA h g$^{-1}$ after 350 cycles, respectively (Figure S21). Ragone plots (Figure 4g) have been used to compare the power densities and energy densities of the as-developed LiIPEICCC−1/MXene battery with previously reported Li−I batteries and representative Li-ion batteries.$^{33}$ Obviously, the novel MXene-based Li−I battery delivers high energy densities (1050 W h kg$^{-1}$ at 0.2 C and 485 W h kg$^{-1}$ at 10 C) and power densities (125 W kg$^{-1}$ at 0.2 C and 5700 W kg$^{-1}$ at 10 C; Table S3), which are much higher.
than most of the previously reported Li–I batteries and lithium-ion batteries.

To explore the potential application of the CC–I/MXene electrode and CPE in flexible devices, a soft packed Li–I battery was fabricated (Figure 4h). The LiI/CPE/CC–I/MXene battery can readily power a red light-emitting diode (LED) lamp under a repeated flat–bend–flat test. This demonstrates that the CC–I/MXene electrode can maintain structural integrity, and the electrode/CPE interfaces can keep tight adhesion under a dramatic shape deformation.

Theoretical calculations were conducted to further investigate the interaction between iodides (LiI was selected as a representative of iodides) and MXenes as well as electrolyte. Figure 5a,b reveals that the binding energies of DOL-LiI3 and DME-LiI3 are calculated to be −0.99 and −1.16 eV, respectively, which are stronger than that of graphene–LiI3 (−0.79 eV). Thus, the LiI3 molecules prefer to dissolve into these electrolyte solvents and shuttle to the anode. In contrast, the binding energy of PETEA–LiI3 is around −1.86 eV. Consequently, LiI3 molecules are preferentially immobilized by the functional groups in the polymer matrix of CPE, leading to its low solubility and slow diffusion in the CPE. This is highly consistent with the experimental results shown in Figure 3e.

Furthermore, as shown in Figure 5d–f, binding energies of LiI with Ti3C2T_x MXenes [−1.76 eV for Ti3C2F2 (Figure 5d), −1.95 eV for Ti3C2O2 (Figure 5e), and −2.1 eV for Ti3C2(OH)2 (Figure 5f)] are also much stronger than those of LiI3 with electrolyte solvents, indicating that LiI3 molecules prefer to adhere to the cathode/CPE surface rather than dissolve into the electrolyte. Notably, the difference in charge density of LiI3 adsorbed on Ti3C2T_x MXenes (Figure 5g,h and Figure S22) shows apparent charge transfer from MXenes to LiI3, verifying the strong chemical interactions between LiI3 and MXene, which leads to the enhanced binding energy as mentioned above. Therefore, the above-mentioned theoretical calculation results convincingly prove that MXene and CPE can cooperatively suppress the shuttle of iodine species and therefore improve the performance of Li–I batteries.

### CONCLUSION

In summary, we have discovered that the synergistic optimization of a CC–I/MXene cathode in conjunction with a composite polymer electrolyte can significantly suppress the self-discharge and enhance the reversible capacity of Li–I batteries. The MXene sheets wrapped on the CC–I electrode possess a high binding energy with iodine species, which successfully restrains them in the cathode. The PETEA-based polymer matrix in the in situ prepared CPE not only immobilizes the iodine species against dissolution and diffusion, but also forms a stable Li anode/CPE interface to suppress the growth of Li dendrites. NaNO3 particles act as a catalyst to efficiently improve the Coulombic efficiency of batteries by enhancing the transformation kinetics of LiI3 in the cathode, and also improve the mechanical strength of CPE. The CC–I/MXene and CPE cooperatively enable a quasi-solid-state Li–I battery with high energy/power density, excellent cycling performance, and good flexibility. This work opens a new path to inspire the development of high-performance Li–I batteries and other novel Li-based rechargeable batteries (e.g., Li–oxygen, Li–sulfur, and Li–selenium batteries).
Graphene interlayer. Energy Storage Materials

Detailed experimental procedures and additional materials characterization (PDF)

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Notes

The authors declare no competing financial interest. Safety statement: no unexpected or unusually high safety hazards were encountered.

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