

Cite this: *Chem. Sci.*, 2025, 16, 13723

All publication charges for this article have been paid for by the Royal Society of Chemistry

Stabilizing the electrode–electrolyte interface for high-voltage Li||LiCoO₂ cells using dual electrolyte additives†

Jiwei Ding,^a Chao Yang,^a Wenxi Hu,^a Xiaowei Liu,^a Anran Zhang,^b Deda Peng,^a Jin Han^{*b} and Ya You^{id*abc}

Rechargeable Li||LiCoO₂ batteries are attractive due to their high energy density. However, the growth of lithium dendrites and the degradation of cathode at high potential hinder their practical application. Herein, we propose an advanced fluorinated carbonate-based electrolyte consisting of *trans*-4,5-difluoro-1,3-dioxolan-2-one (DFEC) and tri-(trimethylsilyl) phosphite (TMSPi) as dual additives to construct a stable interface on both the anode and cathode. The results show that DFEC can promote the formation of a stable solid-electrolyte-interface (SEI) layer on the lithium anode to inhibit the growth of dendrites. Additionally, TMSPi is conducive to the production of an inorganic-rich cathode–electrolyte interface (CEI) layer on the LiCoO₂ cathode to inhibit the dissolution of cobalt. Finally, the Li||LiCoO₂ cells with this electrolyte could obtain an initial capacity of 211.6 mAh g^{−1} (846.4 Wh kg^{−1}, active substances based on cathodes) at 1C (1C = 274 mA g^{−1}) with a high-capacity retention of 81.6% after 200 cycles at a high upper cut-off voltage of 4.6 V. This work provides valuable insights into the development of electrolytes for high-voltage Li||LiCoO₂ cells.

Received 29th April 2025
Accepted 18th June 2025

DOI: 10.1039/d5sc03120f

rsc.li/chemical-science

Introduction

Rechargeable Li||LiCoO₂ cells have broad application prospects because of their high energy density. Unfortunately, LiCoO₂ cathodes can only deliver a relatively low reversible capacity (~140 mAh g^{−1}) at the upper cut-off voltage of 4.20 V, despite their high theoretical capacity (274 mAh g^{−1}).^{1–3} Raising the upper cut-off voltage of LiCoO₂ has been proven to be an effective way to increase the practical capacity. However, once the upper cut-off voltage approaches 4.55 V, the electrolyte can suffer from severe decomposition in the absence of a stable interface on the cathode, leading to continuous capacity degradation of LiCoO₂.^{4–6} Additionally, lithium metal anodes are attractive due to their outstanding theoretical capacity (3860 mAh g^{−1}) and low redox potential (−3.04 V vs. SHE).^{7–11} However, lithium-dendrite growth and the accumulation of dead lithium lead to a short cycle life and low Coulomb

efficiency, hindering the practical application of lithium metal anodes. To address the above-mentioned issues with both the cathode and anode of Li||LiCoO₂ cells, electrode/electrolyte interface engineering seems to be an effective strategy.^{12–17}

Electrode/electrolyte interface engineering can be mainly classified into artificial interface and electrolyte engineering.^{18–21} However, artificial interfaces often require a relatively complex preparation process, which is not conducive to large-scale production. In contrast, electrolyte engineering through the use of additives is a simple and feasible approach. For instance, 2,3-dimethylmaleic anhydride was used as a film-forming additive for constructing a dense and uniform CEI film on an LiCoO₂ cathode, thereby enhancing the interface stability of the LiCoO₂ cathode.²² 2-Fluoro-5-iodopyridine has been proposed as a film-forming additive for lithium metal anodes, and addressed the issue of dendrite growth on lithium metal anodes.²³ However, these electrolyte formulations can only solve the problem with the LiCoO₂ cathode or the lithium metal anode individually.^{24–26} Therefore, there is an urgent need to explore novel electrolyte formulations capable of simultaneously stabilizing both the electrodes.

In this work, we adopted a dual-additive strategy and selected DFEC with a low lowest unoccupied molecular orbital (LUMO) energy level and TMSPi with a high highest occupied molecular orbital (HOMO) energy level as the film-forming additives for the lithium metal anode and the LiCoO₂ cathode, respectively. The results show that DFEC can contribute to the generation of a stable SEI layer on the lithium

^aState Key Laboratory of Advanced Technology for Materials Synthesis and Processing, Wuhan University of Technology, Wuhan 430070, P. R. China. E-mail: youya@whut.edu.cn

^bInternational School of Materials Science and Engineering, School of Materials Science and Microelectronics, Wuhan University of Technology, Wuhan 430070, P. R. China. E-mail: jinhan@whut.edu.cn

^cHubei Longzhong Laboratory, Wuhan University of Technology, Xiangyang Demonstration Zone, Xiangyang 441000, China

† Electronic supplementary information (ESI) available: Experimental section, test parameters and additional figures. See DOI: <https://doi.org/10.1039/d5sc03120f>

metal anode to inhibit the lithium-dendrite growth. TMSPI can induce the formation of an inorganic-rich CEI layer on the LiCoO_2 cathode to inhibit the dissolution of cobalt and enhance the cycling stability. Finally, the proposed electrolyte enabled $\text{Li}||\text{LiCoO}_2$ cells with an initial capacity of 211.6 mAh g^{-1} at 1C and a high-capacity retention of 81.6% after 200 cycles at a high upper cut-off voltage of 4.6 V.

Results and discussion

Characterization of the electrolyte

The solvation structure of the electrolyte TFDT (1 M LiPF_6 dissolved in a blend of bis(2,2,2-trifluoroethyl) carbonate (TFEC), fluoroethylene carbonate (FEC), DFEC and TMSPI (4 : 1 : 0.05 : 0.05, vol%)) was revealed using molecular dynamics (MD) simulation. The selected snapshots and locally enlarged structures (Fig. 1a) show that the Li^+ cations are closely coordinated with TFEC, FEC, and PF_6^- anions. Additionally, the snapshot results show that the additives DFEC and TMSPI are not involved in the solvation structure. This conclusion was further supported by the subsequent nuclear magnetic resonance (NMR) characterization. In the ^1H NMR spectra (Fig. 1b), no peak shift was observed between the spectra of TFDT with and without the additives DFEC and TMSPI, indicating that these two additives are not involved in the solvation structure. The local coordination environment around the Li^+ cations was obtained by analyzing the radial distribution function (RDF) (Fig. 1d). The average coordination number of the Li^+ cations

with PF_6^- , TFEC and FEC was 3.3, 0.97, and 0.95, respectively. However, the average coordination number with the additives DFEC and TMSPI is negligible, indicating that they are not involved in the solvation structure.^{27–29}

To investigate the redox reactivity of the included solvents and additives in the electrolytes, the corresponding HOMO and LUMO were calculated and analyzed using frontier molecular orbital theory (Fig. 1c). Due to the introduction of fluorine, the fluorinated organic solvents TFEC and FEC show higher LUMO levels and lower HOMO levels than the commercial carbonate solvent, indicating their superior oxidation stability. Notably, the additive DFEC possesses a lower LUMO level than the main solvents TFEC and FEC, indicating that DFEC can be preferentially reduced on the lithium metal anode before TFEC. Additionally, the HOMO level of TMSPI (-5.84139 eV) is higher than that of TFEC and FEC (-7.66346 eV), indicating that TMSPI is preferentially oxidized before TFEC and FEC on the LiCoO_2 cathode. Thus, the additives DFEC and TMSPI could potentially be decomposed to produce an interface on the lithium metal anode and LiCoO_2 cathode simultaneously. To evaluate the electrochemical stability of the electrolytes, LSV was conducted at a scanning rate of 0.1 mV s^{-1} on the anodic side in a coin cell containing stainless steel as the working electrode and Li metal as the counter and reference electrode. 1 M LiPF_6 dissolved in a blend of ethylene carbonate (EC), diethyl carbonate (DEC) and vinylene carbonate (VC) (3 : 7 : 0.1, vol%)—denoted as EDV—was chosen as the reference electrolyte. The results in Fig. 1e show that the current rises

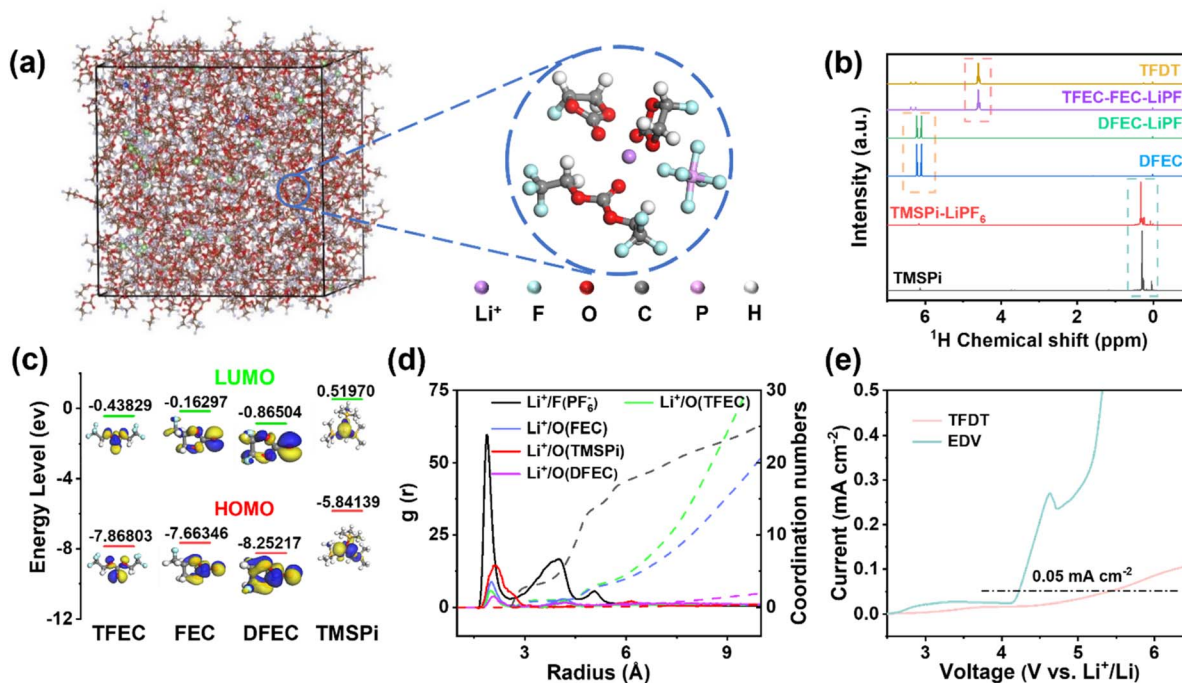


Fig. 1 (a) Snapshot and locally enlarged solvation structure acquired from MD simulations of the electrolyte TFDT. (b) ^1H NMR of various solvents and electrolytes. (c) LUMO/HOMO energy level diagram for the molecules TFEC, FEC, DFEC, and TMSPI. (d) RDF results for Li^+ acquired from MD simulations for TFDT. (e) Electrochemical stability window of the electrolytes EDV and TFDT on the anodic side determined by linear sweep voltammetry (LSV) in a coin cell comprising stainless steel as the working electrode and Li metal as the counter electrode at a scanning rate of 0.1 mV s^{-1} .

sharply in EDV when the potential is higher than 3 V (vs. Li/Li^+). In comparison, for TFDT, the current remains relatively stable until the potential is higher than 5.0 V, demonstrating its prominent anodic electrochemical stability.^{30,31}

Electrochemical performance and interface characterization of the LiCoO_2 cathode

To further evaluate the effect of the electrolytes on the electrochemical performance, the cycle performance of $\text{Li}||\text{LiCoO}_2$ cells with the two electrolytes was measured with an upper cut-off voltage 4.6 V at 1C after it had been activated at 0.1C for the first three cycles (Fig. 2a and b). An initial discharge capacity of 195.9 mAh g^{-1} was delivered by $\text{Li}||\text{LiCoO}_2$ cells with the electrolyte EDV, while the current rapidly decreased to $108.57 \text{ mAh g}^{-1}$ after the 100th cycle. In comparison, $\text{Li}||\text{LiCoO}_2$ cells with TFDT exhibited an initial capacity of 211.6 mAh g^{-1} and maintained a specific capacity of $190.37 \text{ mAh g}^{-1}$ after the 100th cycle. The corresponding coulombic efficiency for the $\text{Li}||\text{LiCoO}_2$ cells with the electrolytes TFDT and EDV is 95.94% and 87.8%, respectively. Notably, TFDT endowed the cell with a high-capacity retention of 81.6% after 200 cycles, which was much higher than that achieved using EDV (7%) (Fig. 2c). Throughout the entire 200 cycles, the coulombic efficiency of the cell with the TFDT was consistently higher than that of the EDV cell. The reason behind the quite different cycling

performances of the electrolytes EDV and TFDT was elucidated using XRD (Fig. S1†). Although the LiCoO_2 electrode cycled in the EDV has similar characteristic peaks to the pristine LiCoO_2 electrode, after cycling in TFDT, the peak intensities of the 003 and 012 crystal planes of the LiCoO_2 were reduced to a much lesser extent compared to those cycled in EDV (Table S1†), indicating that the CEI layer generated in TFDT maintains the structural stability of LiCoO_2 . In addition, the first cycle of the cell with TFDT shows obvious fluctuations in the curve when the voltage is around 4.0 V, while these disappear in the second cycle. This phenomenon can likely be attributed to the decomposition of the electrolyte additive in the first cycle (Fig. S2†) and is not present in EDV (Fig. S3†). However, the charging and discharging curves suffer dramatic variations. A high average discharge potential of 4.0 V can be delivered by $\text{Li}||\text{LiCoO}_2$ cells with TFDT throughout 200 cycles, while the value drops from 4.0 V to 3.5 V with EDV (Fig. S4†).

The rate performance of $\text{Li}||\text{LiCoO}_2$ cells using EDV and TFDT was evaluated under cycling from 0.1C to 5C in the range of 3.0–4.6 V (Fig. 2d). The $\text{Li}||\text{LiCoO}_2$ cell with TFDT can attain capacities of 214.2, 209.4, 206.1, 201.9, 195.7, and 187.3 mAh g^{-1} at 0.1, 0.2, 0.33, 0.5, 1, 2, and 5C, respectively. When the current density is returned to 0.1C, the cell can recover to 202.9 mAh g^{-1} (94.7% retention rate) and maintain stable cycling performance in subsequent cycles. In

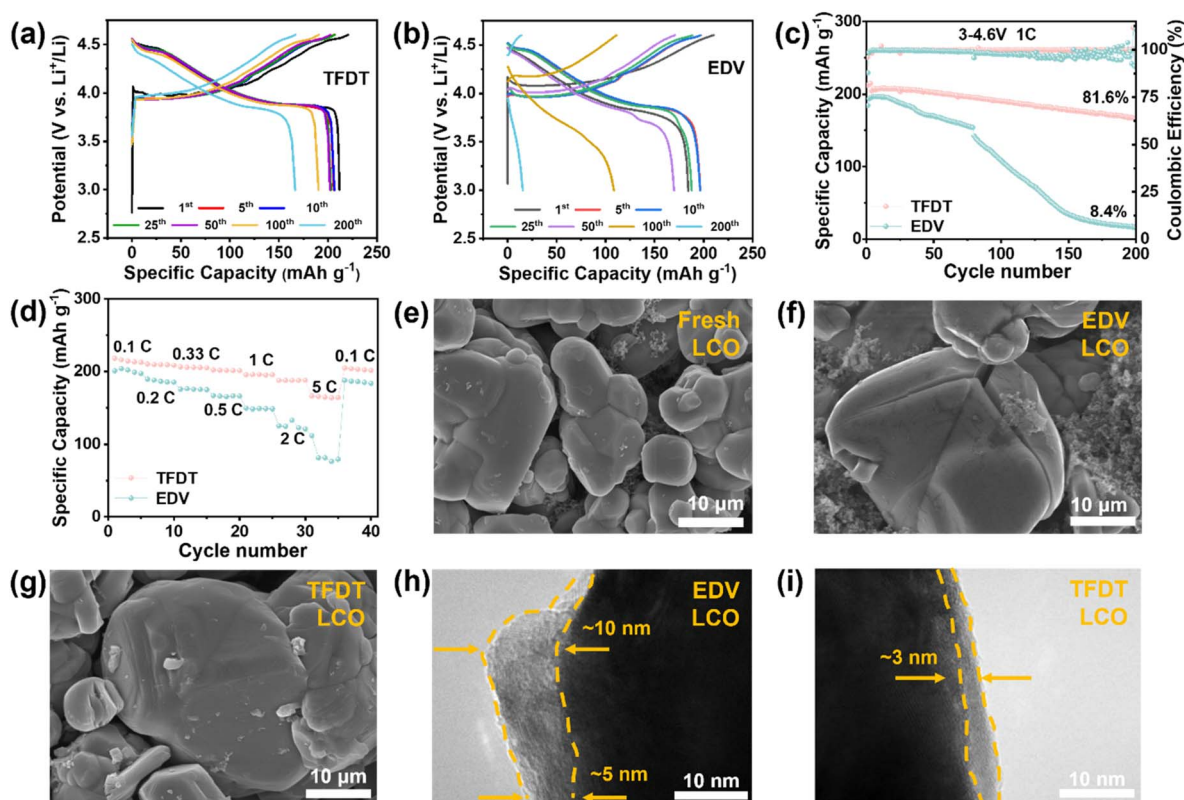


Fig. 2 Galvanostatic charge and discharge curves of $\text{Li}||\text{LiCoO}_2$ cells with the electrolytes TFDT (a) and EDV (b) at different cycles. (c) Cycle performance of $\text{Li}||\text{LiCoO}_2$ cells with both electrolytes at 1C rate. (d) Rate performance of $\text{Li}||\text{LiCoO}_2$ cells with EDV and TFDT. SEM images of fresh LiCoO_2 (e) and the LiCoO_2 cathodes cycled in EDV (f) and TFDT (g) for 50 cycles at 1C with an upper charge cut-off voltage of 4.6 V. TEM images of the LiCoO_2 cathodes cycled in EDV (h) and TFDT (i) under the same measurement conditions.



comparison, the capacity of the $\text{Li}||\text{LiCoO}_2$ cell with EDV was much lower than that of the TFDT at the corresponding current density, indicating the enhanced lithium-ion transport dynamics endowed by TFDT. Furthermore, a constant current intermittent titration test (GITT) was conducted to measure the Li^+ diffusion coefficient when using the electrolytes TFDT and EDV (Fig. S5†). The results indicate that the Li^+ diffusion coefficient in the $\text{Li}||\text{LiCoO}_2$ cell with TFDT is higher than that in the $\text{Li}||\text{LiCoO}_2$ cell with EDV.

The surface morphology of the LiCoO_2 cathode was probed using SEM after 50 cycles in the different electrolytes at 3–4.6 V, as shown in Fig. 2e–g. The results show that large cracks appear on the surface of the LiCoO_2 particles after 50 cycles in EDV (Fig. 2f), which are not found for the fresh LiCoO_2 electrode (Fig. 2e) or the LiCoO_2 electrode cycled in TFDT (Fig. 2g). To further inspect the LiCoO_2 electrode interface, transmission electron microscopy (TEM) was performed on the LiCoO_2 cathodes cycled in EDV and TFDT. It was found that an inhomogeneous CEI layer with a varying thickness (5–10 nm) was formed on the LiCoO_2 cathode in EDV (Fig. 2h), whereas a dense and thin (3–4 nm) CEI layer was generated on the LiCoO_2 cathode in TFDT (Fig. 2i). The above results could indicate that

the severe degradation of LiCoO_2 particles in EDV is related to the absence of a stable CEI layer, while the structural stability of LiCoO_2 particles in TFDT benefits from the protection of a robust CEI layer.

Characterization of the Li metal anode. To inspect the lithium-dendrite growth on lithium anodes in $\text{Li}||\text{LiCoO}_2$ coin cells, the morphology of Li metal anodes cycled in different electrolytes was examined using scanning electron microscopy (SEM) after 50 cycles at 3–4.6 V. Irregular multi-dendrite porous structures were clearly observed on the Li anode with the baseline electrolyte EDV (Fig. 3a). This porous structure implies the production of dead lithium and lithium dendrites in the electrolyte. In contrast, the morphology of the plated Li is relatively flat without obvious dendrites in TFDT (Fig. 3d), indicating that the uniform lithium deposition induced by the SEI layer was derived from the TFDT. The morphology of the lithium metal anode surface after cycling in the different electrolytes was further evaluated using atomic force microscopy (AFM) characterization. As shown in Fig. 3b and c, the surface of lithium metal cycled in EDV is relatively rough. The longitudinal profile of the lithium metal surface is located between –173 nm and 133 nm, and the root mean square roughness (R_q)

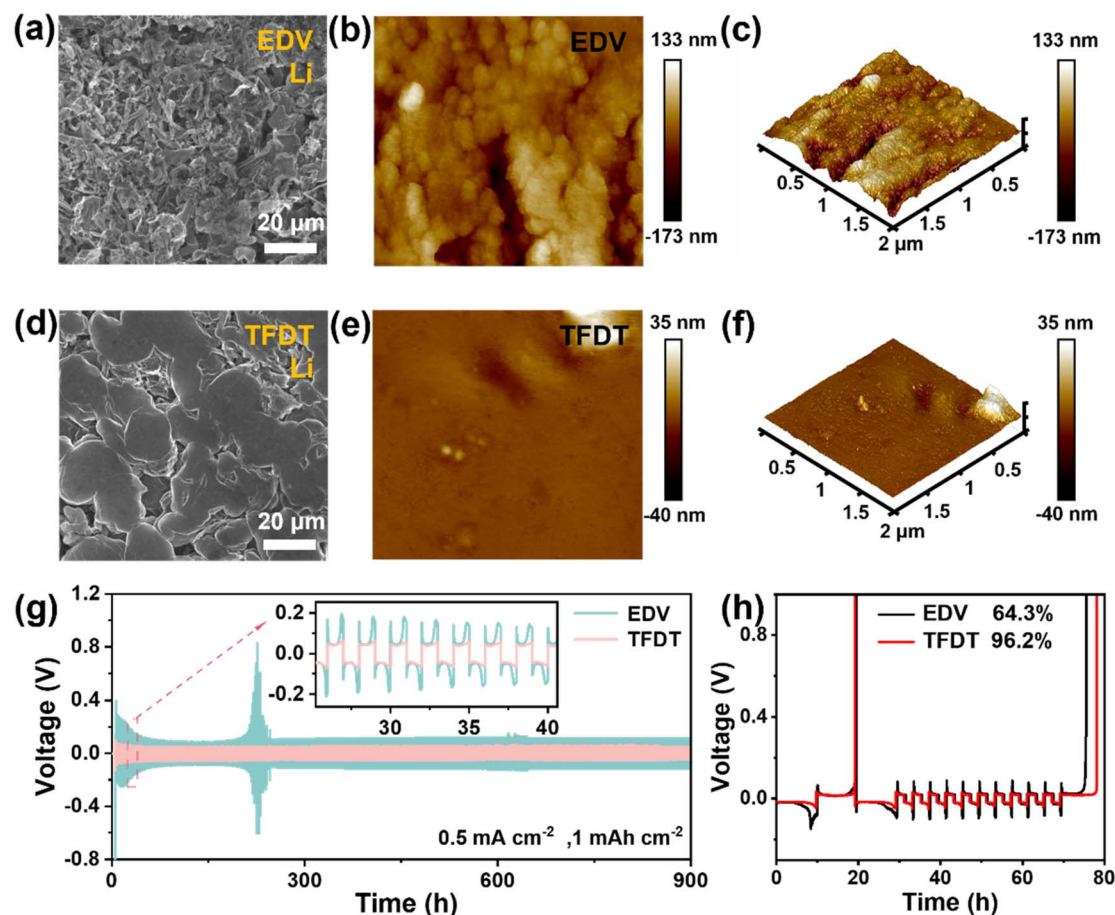


Fig. 3 SEM images of lithium metal anodes cycled in the electrolytes EDV (a) and TFDT (d) after 50 cycles at 1C with an upper charge cut-off voltage of 4.6 V in $\text{Li}||\text{LiCoO}_2$ coin cells. AFM images of lithium metal anodes cycled in EDV (b and c) and TFDT (e and f) under the same measurement conditions. (g) Plating/stripping voltage profiles of the $\text{Li}||\text{Li}$ symmetrical cells in EDV and TFDT. (h) Voltage curves of plating/stripping for $\text{Li}||\text{Cu}$ cells with EDV and TFDT.

and arithmetic average roughness (R_a) are 38.7 nm and 28.7 nm (Table S2†), respectively. In contrast, the surface of the lithium metal cycled in TFDT is relatively smooth, as shown in Fig. 3e and f. The corresponding longitudinal profile of the surface is located between -40 nm and 35 nm, and the roughness results of R_q and R_a are 3.95 nm and 2.01 nm, respectively. In short, TFDT effectively suppressed the dendrite growth and guided a flat lithium deposition by generating a robust SEI.

Li/Li and Li/Cu cells were assembled and measured to evaluate the effect of the electrolyte TFDT on lithium plating and stripping. In the symmetric Li/Li cells setup at a current density of 1 mAh cm^{-2} , TFDT endowed the cell with a relatively low polarization voltage (~ 51 mV) and a long cycle life of 900 h (Fig. 3g), while the commercial electrolyte EDV resulted in a high polarization voltage (~ 118 mV) with an unstable phenomenon. The Auerbach method was used to detect the stability of electrolytes, and the average coulomb efficiency (CE) during plating/stripping was recorded for Li/Cu asymmetric cells, as shown in Fig. 3h.³² TFDT endowed the Li/Cu cells with a higher average CE (96.2%) than EDV (64.3%), indicating that TFDT is more compatible with lithium metal. The exchange current density (J_0) of Li||LiCoO₂ cells with the electrolytes TFDT and EDV was assessed using Tafel curves during the plating/

stripping process at 20°C . The results show that TFDT generates a higher J_0 than EDV (0.116 vs. 0.007 mA cm^{-2}) (Fig. S6†), indicating faster kinetics and better reversibility for lithium plating/stripping in TFDT.³³

Interfacial chemistry. To further reveal the influence of the TFDT electrolyte on the interface, X-ray photoelectron spectroscopy (XPS) analysis was first performed on the lithium metal anode (Fig. 4a and b). The C 1s spectra in Fig. 4a reveal a C–C peak at 284.8 eV , which was assigned to the conductive carbon. The C–O peak at 285.6 eV and O–C=O peak at 289.3 eV originate from Li₂CO₃ as a result of the decomposition of the organic carbonate solvents. In the F 1s spectra (Fig. 4b), the lithium metal anode in TFDT reveals a stronger signal of inorganic LiF (284.8 eV) and Li_xPO_yF_z (284.8 eV) than that of EDV. The intensity of all elements was normalized to evaluate their relative contents in the SEI layer (Fig. 4c). The relative proportions of F and P in the SEI formed in TFDT are significantly higher than those in the SEI formed in EDV; in turn, the proportions of C and O in the SEI formed in TFDT were lower. This indicates that the SEI layer formed on the lithium metal anode in TFDT contains inorganic-rich substances (LiF and Li_xPO_yF_z), thereby enhancing the interface stability of the lithium metal anode.

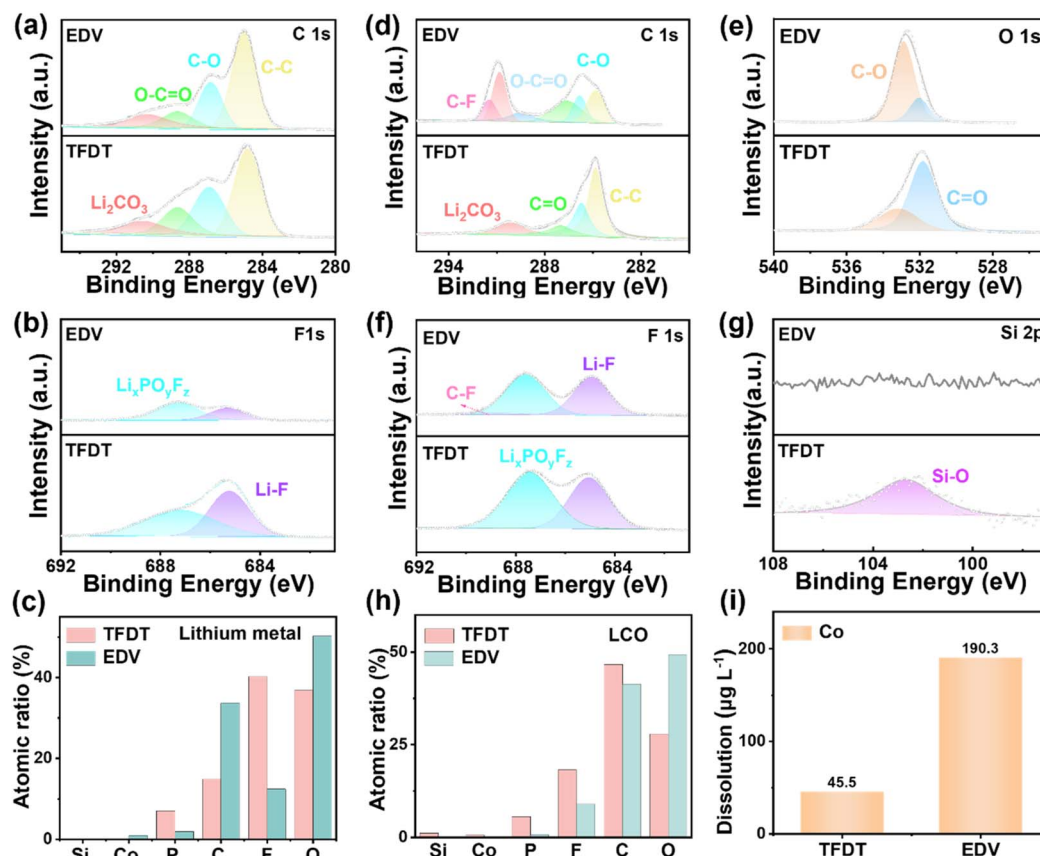


Fig. 4 High-resolution C 1s (a) and F 1s (b) XPS spectra with deconvolution for lithium metal anodes after 50 cycles in the electrolytes EDV and TFDT. (c) Relative contents of the elements in the SEI layer formed on the surfaces of the lithium metal anodes in the different electrolytes. High-resolution C 1s (d), O 1s (e), F 1s (f), and Si 2p (g) XPS spectra with deconvolution for LiCoO₂ cathodes after 50 cycles in the electrolytes EDV and TFDT. (h) Relative contents of the elements in the CEI layer on the surface of the LiCoO₂ cathode. (i) Amount of Co that escaped from the LiCoO₂ cathodes after 50 cycles in the different electrolytes.

To probe the interface chemistry on the LiCoO_2 cathodes cycled in the electrolytes TFDT and EDV, the XPS spectra for the LiCoO_2 cathode in a charged state (4.6 V) are presented in Fig. 4d–g. In the C 1s spectra (Fig. 4d), the spectra peak at 285.6 eV, 287.1 eV, and 289.3 eV are ascribed to C–O, C=O, and O–C=O functional groups, respectively. The peaks at 290.1 eV and 291.4 eV were potentially due to Li_2CO_3 and polyvinylidene fluoride (PVDF).⁴ It is speculated that the CEI on the LiCoO_2 surface in EDV is much richer in organic species than the TFDT one. Regarding the O 1s spectra in Fig. 4e, the peaks at 531.8 eV and 533.0 eV are assigned to C=O and C–O functional groups, respectively.³⁴ When comparing the inorganic fluorine-containing species in the CEIs (Fig. 4f), the CEI formed in TFDT shows a higher signal for $\text{Li}_x\text{PO}_y\text{F}_z$, which is mainly due to the decomposition of the additive TMSPi during the cycling process. Furthermore, the Si–O peak at 103 eV for the CEI on the LiCoO_2 cycled in TFDT is ascribed to the decomposition of the additive TMSPi (Fig. 4g). The Si–O bond has a relatively high bond energy, which can form a stable chemical adsorption layer on the electrode surface and enhance the mechanical strength of the interface, maintaining the stability of the interface.³⁵ The XPS spectra were normalized to calculate the relative proportions of different substances in the CEI (Fig. 4h). The relative proportions of C, F, P and Si in the CEI on the LiCoO_2 cathode in TFDT were significantly higher than those on LiCoO_2 cathode with EDV, while the proportion of O was lower for the LiCoO_2 with TFDT. These collective findings indicate that the CEI formed in TFDT mainly contains inorganic substances, which is beneficial for stabilizing the material structure of the LiCoO_2

cathode.³⁶ In addition, the effective influence of the CEI interface on the LiCoO_2 was investigated using inductively coupled plasma optical emission spectroscopy (ICP), as shown in Fig. 4i. It was found that the amount of cobalt leached in TFDT was $45.5 \mu\text{g L}^{-1}$, which was just 23.9% the amount leached in EDV ($190.3 \mu\text{g L}^{-1}$), indicating the enhanced structural stability of the LiCoO_2 cathode in TFDT.

Thermal safety test for EDV and TFDT electrolytes. Organic electrolytes under deep delithiation will undergo intense exothermic reactions; therefore, thermal safety is crucial for lithium-ion cells.^{37,38} The flammability of the electrolytes was assessed using a conventional ignition test, as shown in Fig. 5a. The TFDT electrolyte could not be ignited after being exposed to the flame of a methane lighter for 6 s, nor was it ignited in two subsequent attempts, while the EDV electrolyte caught fire rapidly and burned violently until the electrolyte had burned off. Differential scanning calorimetry (DSC) was used to measure the thermal abuse tolerance of the electrolytes when paired with a charged-state LiCoO_2 electrode at 4.6 V (Fig. 5b). It can be seen that the LiCoO_2 /EDV mixture showed an exothermic reaction with a total exothermic heat of 444 J g^{-1} . However, the total heat release of the LiCoO_2 /TFDT mixture was 241 J g^{-1} , which was 54.3% that of the LiCoO_2 /EDV mixture. These results indicate that TFDT can reduce the heat production of exothermic reaction for the electrode and electrolyte mixture.

To further investigate the heat-related behavior associated with these electrolytes, a cone calorimeter was used to collect some parameters of the combustion process, such as the self-extinction time (SET), total heat release (THR), and maximum

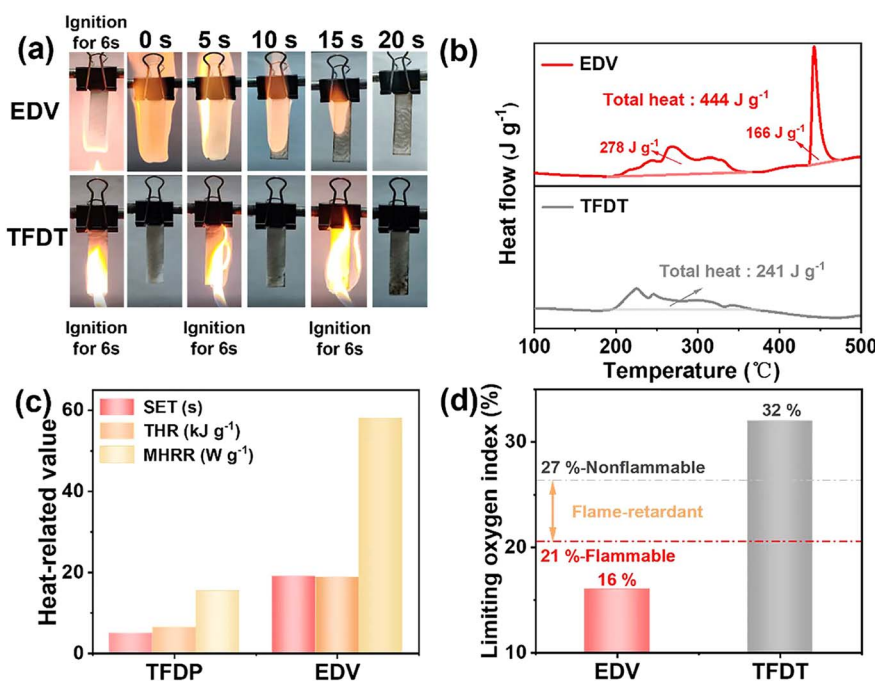


Fig. 5 (a) Ignition tests for EDV and TFDT electrolytes. (b) DSC profiles of charged-state LiCoO_2 (4.6 V) mixed with EDV or TFDT in a weight ratio of 6.5 : 3.5 at a scan rate of 5°C min^{-1} between 100 and 500°C . (c) Self-extinguishing time (SET), time to ignite (TTI), and total heat release (THR) of EDV and TFDT as determined using a cone calorimeter. All samples were exposed to heat radiation until they had thoroughly ignited. (d) Limiting oxygen indices of EDV and TFDT.

heat release rate (MHRR). All samples were exposed to a constant thermal radiation value of 25 kW m^{-2} until fully ignited. The TFDT electrolyte has a lower MHRR (15.4 W g^{-1} vs. 58.1 W g^{-1}), lower SET (5 s vs. 19 s) and lower THR (6.54 kJ g^{-1} vs. 18.78 kJ g^{-1}) than EDV electrolyte, as shown in Fig. 5c. In addition, the gas produced during the combustion of TFDT has a higher CO concentration and lower CO_2 content than that of EDV, which indicates the incomplete combustion and notable flame retardance of TFDT (Table S3†). To quantify the flammability of the electrolytes, the limiting oxygen index (LOI) test was used to further measure their flammability. As shown in Fig. 5d, TFDT has a much higher LOI (32%) than EDV (16%). Based on the classification zones in Fig. 5d, TFDT has flame-retardance characteristics, while EDV is flammable. In summary, TFDT has better flame retardance and self-extinguishing characteristics compared with EDV. Additionally, the lower heat release during combustion may be related to the higher concentration of fluorine-containing solvents in TFDT.

Conclusions

In summary, a novel electrolyte containing dual additives was designed, and its corresponding solvation structure was studied. The electrolyte has properties that optimize the formation of electrolyte interfaces. The TMSPI additive enhances the interface stability of the LiCoO_2 cathode by forming stable CEI layers, thereby further preventing the structural collapse of LiCoO_2 and the leaching of cobalt under high operation potential. Additionally, the additive DFEC generates an inorganic-rich SEI layer on the surface of the lithium metal anode and inhibits the growth of lithium dendrites. Therefore, the tailored electrolyte endowed Li/LiCoO_2 cells with an initial capacity of 211.6 mAh^{-1} (846.4 Wh kg^{-1}) and a high-capacity retention of 81.6% after 200 cycles at a high cut-off voltage of 4.6 V. This work broadens the perspective on electrolyte development for high-voltage Li/LiCoO_2 cells through the use of electrolyte additives.

Data availability

The data supporting this article have been included as part of the ESI.†

Author contributions

Y. Y. and J. H. conceived and supervised the project; J. D. designed the experiments; J. D., W. H., and D. P. conducted the experiments; X. L. and A. Z. conducted the DFT calculations and MD simulations. J. D. and C. Y. conducted the characterizations and analysis. J. D. drafted the manuscript with the help of Y. Y. and J. H.; all authors agreed on the manuscript and approved submission.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by National Key Research and Development Program of China (2022YFB3803400).

Notes and references

- 1 E. Cha, J. H. Yun, R. Ponraj and D. K. Kim, *Mater. Chem. Front.*, 2021, **5**, 6294–6314.
- 2 S. Awasthi, *J. Energy Storage*, 2024, **100**, 113439.
- 3 Z. Chang, C. Ma, R. Wang, B. Wang, M. Yang, B. Li, T. Zhang, Z. Li, P. Zhao, X. Qi and J. Wang, *ACS Appl. Mater. Interfaces*, 2024, **16**, 18980–18990.
- 4 K. Guo, C. Zhu, H. Wang, S. Qi, J. Huang, D. Wu and J. Ma, *Adv. Energy Mater.*, 2023, **13**, 2204272.
- 5 M. Ma, Z. Zhu, D. Yang, L. Qie, Z. Huang and Y. Huang, *ACS Appl. Mater. Interfaces*, 2024, **16**, 12554–12562.
- 6 C. Luo, Q. Liu, X. Wang, Y. Tian, Z. Liu, F. Kang and B. Li, *Nano Energy*, 2023, **109**, 108323.
- 7 X.-B. Cheng, R. Zhang, C.-Z. Zhao and Q. Zhang, *Chem. Rev.*, 2017, **117**, 10403–10473.
- 8 J.-F. Ding, Y.-T. Zhang, R. Xu, R. Zhang, Y. Xiao, S. Zhang, C.-X. Bi, C. Tang, R. Xiang, H. S. Park, Q. Zhang and J.-Q. Huang, *Green Energy Environ.*, 2023, **8**, 1509–1530.
- 9 H. Kang, H. Kang, M. Lyu and E. Cho, *EcoMat*, 2024, **6**, e12498.
- 10 X. Duan, J. Sun, L. Shi, S. Dong and G. Cui, *Interdiscip. Mater.*, 2024, **4**, 217–234.
- 11 J. Xu, X. Cai, S. Cai, Y. Shao, C. Hu, S. Lu and S. Ding, *Energy Environ. Mater.*, 2023, **6**, e12450.
- 12 O. B. Chae and B. L. Lucht, *Adv. Energy Mater.*, 2023, **13**, 2203791.
- 13 Y. Yan, S. Weng, A. Fu, H. Zhang, J. Chen, Q. Zheng, B. Zhang, S. Zhou, H. Yan, C.-W. Wang, Y. Tang, H. Luo, B.-W. Mao, J. Zheng, X. Wang, Y. Qiao, Y. Yang and S.-G. Sun, *ACS Energy Lett.*, 2022, **7**, 2677–2684.
- 14 H. Yang, T. Zhao, K. Huang, Y. Shen, Y. Huang and W. Luo, *Adv. Funct. Mater.*, 2023, **33**, 2302203.
- 15 J. Yang, X. Liu, Y. Wang, X. Zhou, L. Weng, Y. Liu, Y. Ren, C. Zhao, M. Dahbi, J. Alami, D. A. Ei-Hady, G. L. Xu, K. Amine and M. Shao, *Adv. Energy Mater.*, 2021, **11**, 2101956.
- 16 Y. You, A. Dolocan, W. Li and A. Manthiram, *Nano Lett.*, 2018, **19**, 182–188.
- 17 Q. Ding, Z. Jiang, K. Chen, H. Li, J. Shi, X. Ai and D. Xia, *Carbon Energy*, 2024, **6**, e602.
- 18 S. Chen, J. Zheng, D. Mei, K. S. Han, M. H. Engelhard, W. Zhao, W. Xu, J. Liu and J. G. Zhang, *Adv. Mater.*, 2018, **30**, 1706102.
- 19 S. Dai, W. Fang, T. Wang, Y. Gao, T. Zhang, Z. Qin, G. Chen and X. Zhou, *Chem. Eng. J.*, 2024, **500**, 157269.
- 20 M. Ma, R. Huang, M. Ling, Y. S. Hu and H. Pan, *Interdiscip. Mater.*, 2023, **2**, 833–854.
- 21 Z. Wang, L. P. Hou, Z. Li, J. L. Liang, M. Y. Zhou, C. Z. Zhao, X. Zeng, B. Q. Li, A. Chen, X. Q. Zhang, P. Dong, Y. Zhang, J. Q. Huang and Q. Zhang, *Carbon Energy*, 2022, **5**, e283.



- 22 A. Fu, C. Xu, J. Lin, Y. Su, H. Zhang, D.-Y. Wu, X. Zhang, M. Xia, Z. Zhang, J. Zheng and Y. Yang, *J. Mater. Chem. A*, 2023, **11**, 3703–3716.
- 23 E. Zhang, H. Tian, M. Li, S. Le, B. Li, L. Wu, Q. Zhang and L. Fan, *Chem. Commun.*, 2023, **59**, 10996–10999.
- 24 A. Zhang, Z. Bi, G. Wang, S. Liao, P. Das, H. Lin, M. Li, Y. Yu, X. Feng, X. Bao and Z.-S. Wu, *Energy Environ. Sci.*, 2024, **17**, 3021–3031.
- 25 C. Zhang, S. Gu, D. Zhang, J. Ma, H. Zheng, M. Zheng, R. Lv, K. Yu, J. Wu, X. Wang, Q.-H. Yang, F. Kang and W. Lv, *Energy Storage Mater.*, 2022, **52**, 355–364.
- 26 S. D. Zhang, M. Y. Qi, S. J. Guo, Y. G. Sun, X. X. Tan, P. Z. Ma, J. Y. Li, R. Z. Yuan, A. M. Cao and L. J. Wan, *Small Methods*, 2022, **6**, 2200148.
- 27 H. Zhu, J. Ma, H. Ding, H. Wu, C. Zhang, L. Ni, X. Fang, L. Lao and X. Wang, *Appl. Surf. Sci.*, 2024, **677**, 161100.
- 28 Y. Wang, Z. Li, Y. Hou, Z. Hao, Q. Zhang, Y. Ni, Y. Lu, Z. Yan, K. Zhang, Q. Zhao, F. Li and J. Chen, *Chem. Soc. Rev.*, 2023, **52**, 2713–2763.
- 29 J. Wu, Z. Gao, Y. Wang, X. Yang, Q. Liu, D. Zhou, X. Wang, F. Kang and B. Li, *Nano-Micro Lett.*, 2022, **14**, 147.
- 30 Y. Wu, Z. Zeng, H. Zhang, M. Liu, S. Lei, W. Zhong, S. Cheng and J. Xie, *Energy Storage Mater.*, 2024, **70**, 103499.
- 31 W. Xi, X. Xia, J. Zhu, D. Yang and S. Xie, *Nano Res.*, 2024, **17**, 7230–7241.
- 32 F. Li, J. Liu, H. Wang, Y. Ren, X. Tang and G. Li, *ACS Energy Lett.*, 2025, **10**, 779–787.
- 33 C.-C. Su, M. He, M. Cai, J. Shi, R. Amine, N. D. Rago, J. Guo, T. Rojas, A. T. Ngo and K. Amine, *Nano Energy*, 2022, **92**, 106720.
- 34 Z. Zhang, C. Qin, K. Wang, X. Han, J. Li, M. Sui and P. Yan, *J. Energy Chem.*, 2023, **81**, 192–199.
- 35 R. Tian, O. Seitz, M. Li, W. Hu, Y. J. Chabal and J. Gao, *Langmuir*, 2010, **26**, 4563–4566.
- 36 Z. Zhang, C. Qin, X. Cheng, J. Li, Y. Zhang, W. Zhao, L. Wang, Y. Du, M. Sui and P. Yan, *Electron*, 2023, **1**, e9.
- 37 A. Yang, C. Yang, K. Xie, S. Xin, Z. Xiong, K. Li, Y.-G. Guo and Y. You, *ACS Energy Lett.*, 2023, **8**, 836–843.
- 38 Z.-H. Wu, Y. Wu, Y. Tang, J.-C. Jiang and A.-C. Huang, *Process Saf. Environ. Prot.*, 2023, **169**, 285–292.

