

Supplementary Information

Revolutionizing Lithium-Ion Batteries: Exploiting Liquid Crystal Electrolytes

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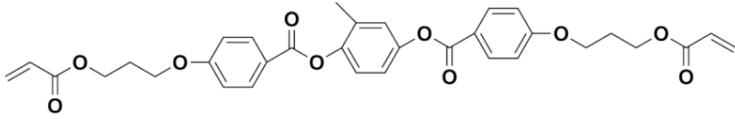
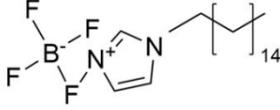
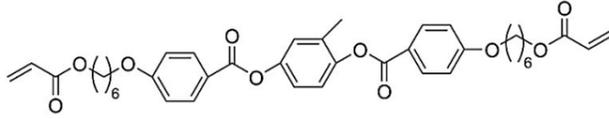
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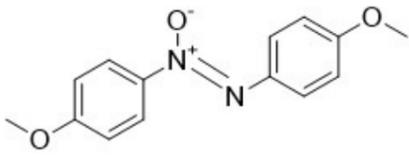
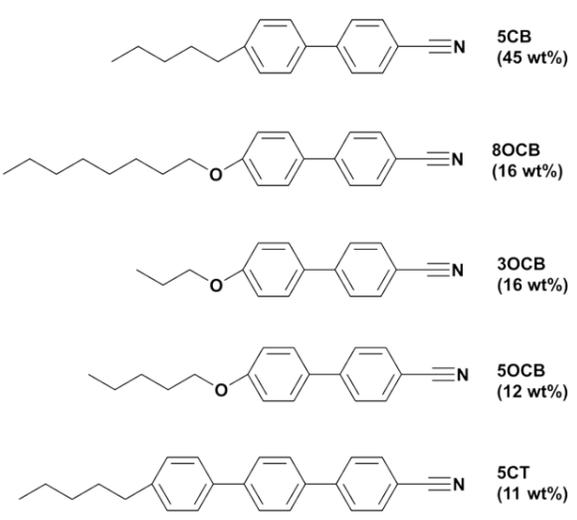
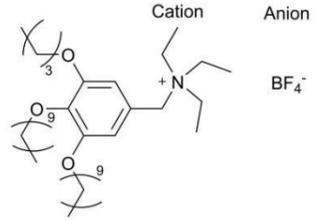
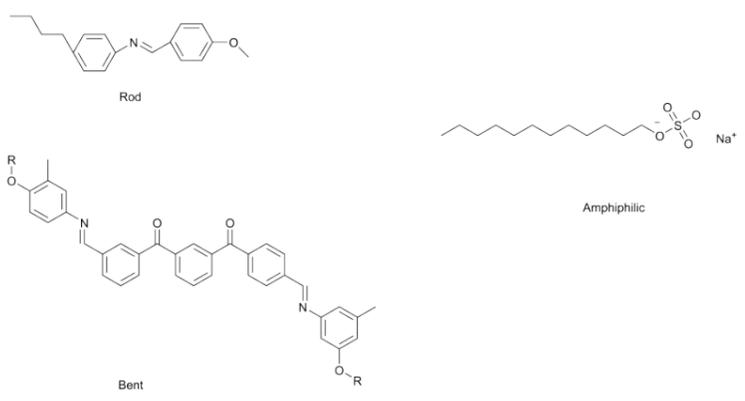
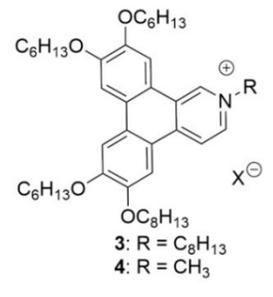
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Table S1. Nanostructure dimensionality of LCEs

Phase Type	Ion Channel Dimension	Ionic Conductivity	Key Advantages
Cubic (3D)	3D interconnected	Highest ($>10^{-3}$ S cm^{-1})	Isotropic ion transport; stable at low temperatures
Smectic (2D)	2D layered	Moderate ($\sim 10^{-4}$ S cm^{-1})	Anisotropic conduction; tunable layer spacings
Columnar (1D)	1D channel	Lowest ($\sim 10^{-5}$ - 10^{-8} S cm^{-1})	Ease of alignment via external fields, temperature or surface

Table S2 The commercialization process of LCEs

Number	Chemical Formula	Ion Transport Mechanism	Ionic/Non-ionic	Ionic Conductivity (σ)	Remarks under Specific Conditions	Commercialization process	ref.
1.		2D	non-ionic	$3 \pm 0.5 \times 10^{-3}$ S cm ⁻¹ at room temperature		RM257 is an industrial-grade UV liquid crystal monomer for LCEs, used in smart devices and lithium batteries. It dissolves easily, has low viscosity, and is mass-produced.	1
2.		2D	ionic	1.4×10^{-3} S cm ⁻¹ at 30 °C	It combines the amphiphilic feature of imidazolium 1-hexadecyl-3-methylimidazolium tetrafluoroborate with propylene carbonate and LiBF ₄ , distinguishing it from other conventional electrolytes	It is classified as ionic LCEs, with potential applications in electrochemical devices (e.g., fuel cells, supercapacitors, and fluorination media).	2
3.	<i>poly-(methoxy-poly(ethylene glycol) methacrylate)</i>	2D	non-ionic polymer	1.0×10^{-7} S cm ⁻¹ at 25 °C	It has combined with LC or hydrophobic blocks to further improve its properties, setting it apart from other conventional polymer electrolytes.	It is a classical compound in the liquid crystal display (LCD) field, first commercialized in the 1970s by companies such as Merck & Co., Inc. C6M is a custom-synthesized monomer that has not yet been commercialized on a large scale. Its high-purity synthesis is costly (estimated to be over \$500 per gram), and it requires strict inert environment handling, which limits its industrial applications.	3
4.		3D	non-ionic	1.79×10^{-3} cm ⁻¹ at 20 °C	It combines with ionic liquids (IL) to form polymer networks, which exhibits enhanced electrochemical properties compared to other electrolytes.		4

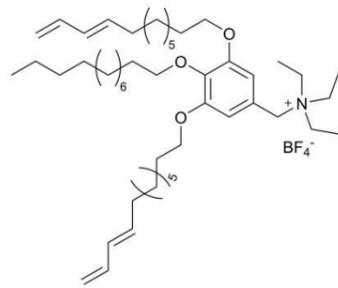
5.		2D	non-ionic	$1.3 \times 10^{-4} \text{ S cm}^{-1}$ at room temperature	It is a laboratory-grade innovative electrolyte, designed to address the interface compatibility and dendrite issues of SSBs. It has not yet been commercialized, but is described as a promising strategy.	5
6.		2D	non-ionic	$10^{-6} \text{ S cm}^{-1}$ at 72 °C	As a new type of solid electrolyte, it is targeted for applications in rechargeable metal ion batteries ($\text{Li}^+/\text{Na}^+/\text{Mg}^{2+}$), aiming to solve problems such as leakage of liquid electrolytes and dendrite growth.	6
7.		3D	ionic	-	It is a prototype compound developed in the laboratory and can be used as a candidate material for SSEs and fuel cells.	7
8.		2D	ionic	-	The electro-optic effect of graphene oxide LCs is applied to optical switch devices, but has not yet been commercially utilized on a large scale.	8
9.	 <p>3: R = C₈H₁₃ 4: R = CH₃</p>	2D	ionic	-	There are no direct commercialization cases, but basic research has been active (since year 2010 to present), and the target applications include: organic semiconductors (photovoltaics, OLEDs, transistors), ionic conductive electrolytes (LIBs, fuel	9

cells), and fluorescent sensors (anion-selective luminescence)

It is currently used in the laboratory research stage and has not yet been commercially scaled up. It applies potential as a new type of functional material in areas such as separation membranes, ionic conductors, and adsorbents. It can be used for virus filtration and is a new material at the laboratory stage, with potential for industrial applications.

It is targeted at proton exchange membranes (PEMs) used in fuel cells and electrochemical devices, but it is currently still in the proof-of-concept stage.

10.



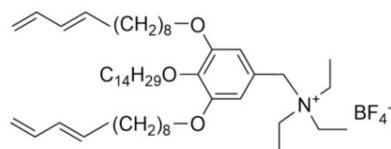
3D

ionic

$3.1 \times 10^{-4} \text{ S cm}^{-1}$ at 90°C ,
 $7.75 \times 10^{-5} \text{ S cm}^{-1}$ at 90°C ,
 $8.8 \times 10^{-6} \text{ S cm}^{-1}$ at 90°C

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11.



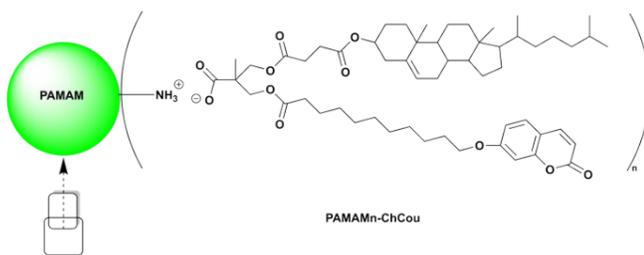
2D

ionic

-

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12.



1D or 2D

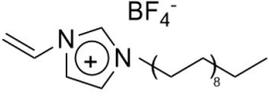
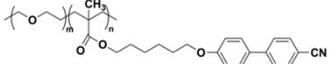
ionic

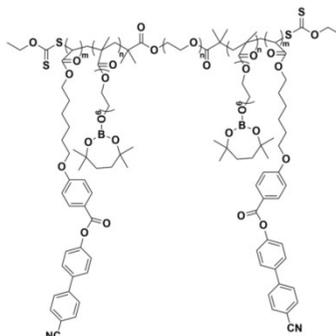
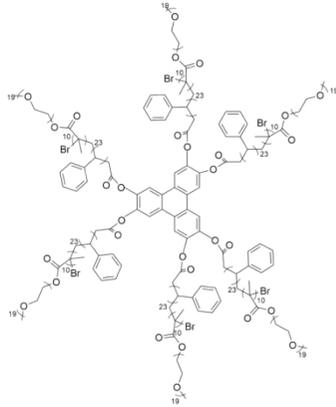
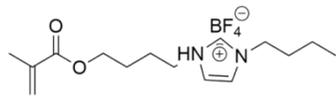
$10^{-4} \sim 10^{-3} \text{ S cm}^{-1}$ from 25°C to 100°C

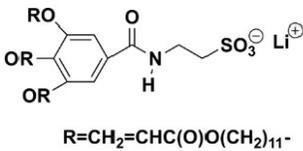
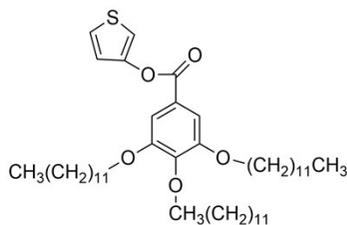
The material uses ionic LC dendrimers to create tunable 1D/2D proton-conductive channels stabilized by coumarin photocrosslinking.

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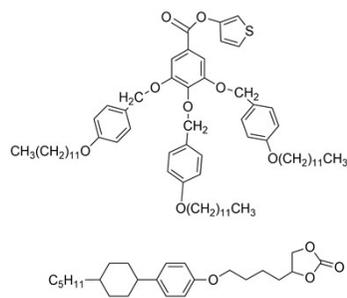
Table S3 Performance summarization of LCEs

Number	Chemical Formula	Electrochemical stability	Thermal stability	Mechanical properties	Ease of fabrication	Interface compatibility	Safety	Cost	Scalability	ref.
1.		The electrochemical window is 5.2 V. It is superior to liquid electrolyte.	Decomposition temperature of inoic LC monomer: 268 °C. Meeting the safety requirements of lithium batteries (>200 °C), superior to flammable liquid electrolytes.	Self-supporting flexible membrane (free-standing and flexible), capable of bending, with a cross-linked structure (PEGDA photopolymerization) providing mechanical strength	One-pot photopolymerization (a mixture of ILC/PEGDA/PEGDE/LiBF ₄ + UV curing)	Suppressing lithium dendrites, low interfacial resistance	The all-solid-state structure eliminates the risk of liquid leakage. No volatile solvents, high thermal stability.	High cost	Solution processing is conducive to scaling up.	13
2.		-	High decomposition temperature (290 °C), stable interlamellar ordered structure.	The star-shaped structure enhances flexibility, while the interlamellar units provide strength.	The film was prepared by atom transfer radical polymerization (ATRP) synthesis and induced to form a lamellar ordered structure	The hydrophilic poly-(methoxy poly (ethylene glycol) methacrylate) (PPEGMA) block promotes contact with the electrode, and the PEO side chain has a strong ability to dissolve lithium salts.	Fully solid-state intrinsically safe structure	High cost	The yield of star-shaped polymers is relatively low (requiring multiple purification steps), and there are challenges in large-scale productions.	3

3.		<p>The electrochemical window is wide, measured at 4.85 V.</p>	<p>Decomposition temperature of 390 °C, with no leakage risk</p>	<p>Optimize the interface through the use of fluorinated ethylene carbonate (FEC) additives</p>	<p>Solution casting method is compatible with existing processes</p>	<p>Adding FEC to the lithium metal interface can inhibit the side reaction between the cyano-group and lithium.</p>	<p>The solid electrolyte is non-flammable and shows no thermal runaway during puncture/cutting tests.</p>	<p>High cost</p>	-	14
4.		<p>The electrochemical window is 5.1 V, meeting the requirement of high voltage.</p>	<p>Decomposition temperature of 390 °C, no leakage risk, superior to liquid electrolyte</p>	<p>It has a flexible self-supporting membrane that can restore its shape after being bent.</p>	<p>Solution casting (in THF) and annealing (at 120 °C) induce an ordered structure.</p>	<p>Low polarization, no dendrites</p>	<p>Fully solid-state, without risk of combustion or explosion</p>	<p>High cost</p>	-	15
5.		<p>Electro-chemical window > 4.2 V, meeting the requirement of</p>	<p>The decomposition temperature is greater than</p>	<p>The composite electrolyte forms a self-</p>	<p>UV curing is fast and efficient.</p>	<p>The cyclic stability is good (> 85%), and the interface is</p>	<p>Non-flammable and no leakage</p>	<p>High cost</p>	-	16

6.	 <p>R=CH₂=CHC(O)O(CH₂)₁₁-</p>	high voltage.	290 °C. The ionic liquid characteristics eliminate the risk of volatile leakage of the traditional liquid electrolyte.	supporting film, with the cross-linking agent PEGDA providing the network framework, and the ordered structure of the LC phase enhancing the mechanical strength. Self-supporting films possess the characteristics of flexibility, transparency and high mechanical strength.	relatively stable.	Solid-state encapsulation of liquid electrolyte, eliminating the risks of leakage and combustion.	High cost	-	17
7.		The electrochemical window is 4.0 V, meeting the requirement of high voltage.	Good thermal stability	-	Multi-step synthesis, solvent-assisted coating	Solvent-free design reduces the risk of explosion	High cost	-	18

8.



The electrochemical window is as wide as 3.9 V, and the cycling stability is excellent, thus meeting the requirements for high voltage applications.

This phase persists in a LC state within the temperature range from room temperature up to either 80 °C or 53 °C.

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The solution processing approach is viable; however, precise control of components poses significant challenges.

The electrode interface exhibits good compatibility with both the Li metal anode and the LiFePO₄ cathode, characterized by a low interface impedance.

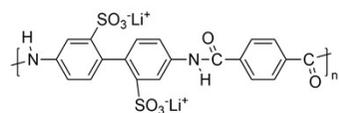
Minimize the risks of leakage and volatilization

High cost

-

19

9.



The electrochemical window measures 5.6 V.

The thermal stability is satisfactory. The LiFSI nanocrystals commence to degrade at 140 °C.

The mechanical properties far exceed those of traditional SPEs (typically < 10 MPa), approaching the rigidity of ceramic electrolytes, and can effectively prevent lithium dendrite penetration.

Two-step process: RMIC preparation and LiMIC preparation

The interface of lithium metal is capable of forming a stable solid electrolyte interphase (SEI) layer, and the corresponding interfacial impedance remains relatively low.

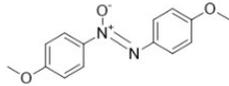
Without volatile solvents, it completely eliminates the risk of leakage and combustion.

High cost

-

20

10.



The electrochemical window is 4.9 V, meeting the requirement of high voltage.

-

The 3D fiber network structure enhances mechanical strength and effectively blocks dendrite penetration.

In-situ polymerization: A solvent-free approach is necessitated. It involves the direct polymerization of the precursors (PEGDMA /ETFP/ LC) within the battery.

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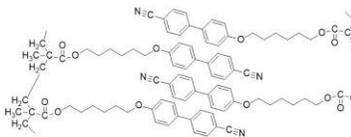
Solid-state structure, tested through bending and pricking methods

High cost

-

5

11.



The electrochemical window is 4.5 V, meeting the high-voltage requirements.

It has good thermal stability. It does not shrink after being heated at 150 °C for 60 minutes. It begins to decompose at 350 °C.

PEGDA cross-linking provides the supporting framework, while the liquid crystal copolymer induces microphase separation.

Solution casting, UV curing, room temperature drying, vacuum annealing

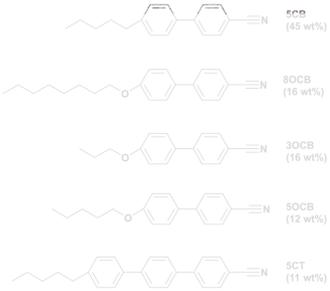
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It is flame-retardant and has no leakage risk.

High cost

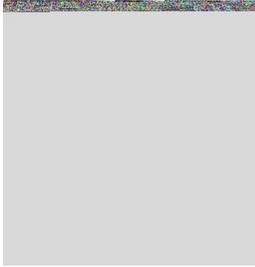
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21

12.		-	The thermal stability is poor and the melting point of the material is relatively low.	With self-supporting flexible membrane and capable of being bent and shaped.	Solution casting method	-	Solid-state system, no leakage, low flammability, inhibition of dendrite growth	Moderate to high costs	-	6
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13.	$n-C_{16}H_{33}-N \begin{array}{c} \diagup \\ \diagdown \end{array} \begin{array}{c} + \\ - \end{array} \begin{array}{c} \diagdown \\ \diagup \end{array} N-CH_3$ PF_6^-	Limited PF_6^- hydrolysis, with a relatively narrow voltage window.	The phase transition temperature is moderate (<125 °C)	Dependent on porous substrates, it is flexible but has limited strength.	Alumina film surface treatment (PVA/C16TAB) → Capillary filling (isotropic phase at 140 °C)	-	The PF_6^- anion will hydrolyze upon contact with water, generating HF (a highly toxic and corrosive substance), and a strictly dry environment is required.	High cost	-	22
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14.



Excellent thermal stability

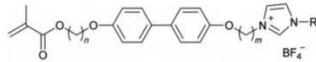
Solution-based film formation (THF solution spin coating) achieves uniform films

Low volatility but with the risk of fluorine-containing anions

High cost

9

15.



The operating temperature for water treatment is within the range of room temperature to 100 °C, but high-temperature applications are limited.

The ultrathin film (100 - 150 nm) is supported on a polysulfone substrate.

Photopolymerization (UV-LED) + Sacrificial Layer Transfer Method (PVA-coated PET film → Polysulfone substrate)

No leakage risk

High cost

The spin-coating and UV polymerization process have the potential for large-scale applications, but the large-scale production of ultra-thin and defect-free films remains to be verified.

23

16.		<p>The polymer film remains stable up to 120 °C.</p>	<p>It has a self-supporting film that is insoluble in water and organic solvents, indicating that the cross-linked network is stable.</p>	-	-	<p>Fully solid-state, with no risk of solvent leakage</p>	<p>High cost</p>	-	10
17.		<p>Forming a highly cross-linked LCNs through photopolymerization/thermal curing, enhancing stability</p>	<p>Introducing covalent "nanosticks" or increasing the crosslinking density can maintain the integrity of the pore channels.</p>	<p>Self-assembly reduces costs and is processable by solution method.</p>	-	<p>No heavy metals or volatile solvents. Theoretical safety is high.</p>	<p>High cost</p>	-	24
18.		-	-	<p>Solution casting and vacuum drying</p>	-	<p>The water-free system avoids the risk of high-temperature steam pressure. Benzenesulfonic acid is less corrosive</p>	<p>High cost</p>	-	25

19.



The phase transition temperature is relatively low.

After crosslinking, it forms polymer networks, enhancing mechanical stability

The PAMAM dendrimer macromolecules (G0-G4) and the coumarin-cholesterol dendrasylic acid (Ac-ChCou) self-assemble through ions. Solvent evaporates to form a film, followed by UV irradiation

than strong inorganic acids.

UV curing without the use of initiators results in fewer side reactions and is relatively safer.

High cost

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