

单离子聚合物快离子导体

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A Single-Ion Polymer Superionic Conductor

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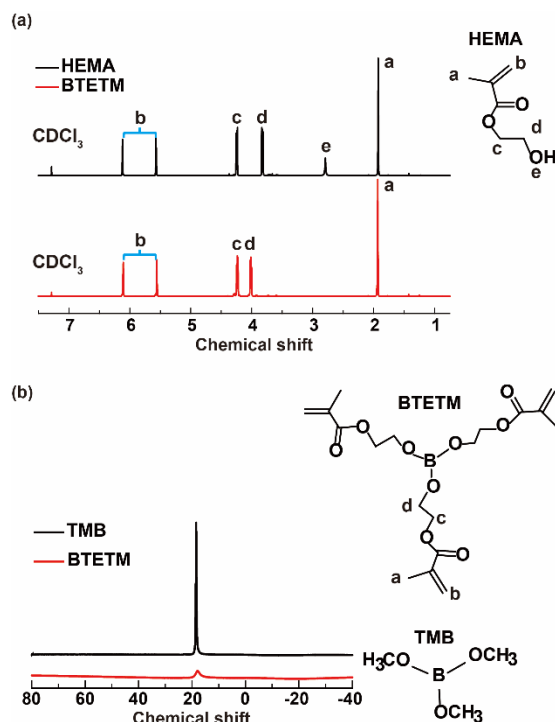


Fig. S1 (a) ^1H -NMR spectra of the HEMA and BTETM samples. (b) ^{11}B -NMR of the TMB and BTETM samples.

The structure of the synthesized BTETM was characterized by nuclear magnetic resonance (NMR). It is obvious that when the hydroxyl groups of the HEMA are substituted, the hydroxyl resonance peak of e disappears, while the resonance peak of d shifts from the original 3.82 to 4.01 due to the lack of electrons in the boron atom. In addition, both TMB and BTETM exhibit a resonance peak near 18.0, which belongs to the tri-coordinated boron atoms, but the resonance peak of the BTETM is much broader than that of the TMB due to the different coordination molecules of the tri-coordinated boron atoms. These results confirm the successful synthesis of the BTETM.

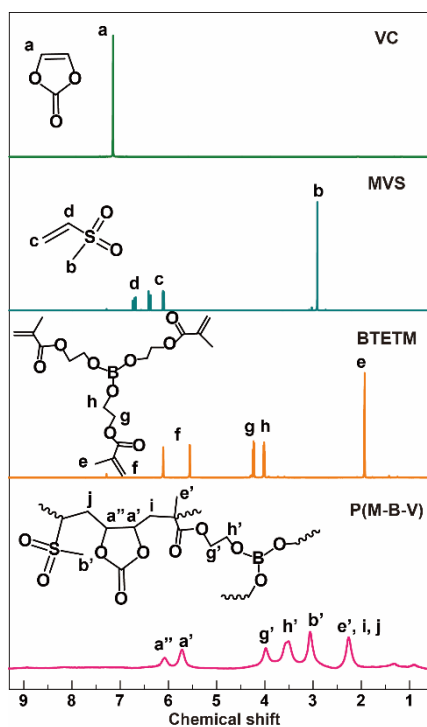


Fig. S2 ^1H -NMR spectra of the VC, MVS, BTETM, and P(M-B-V).

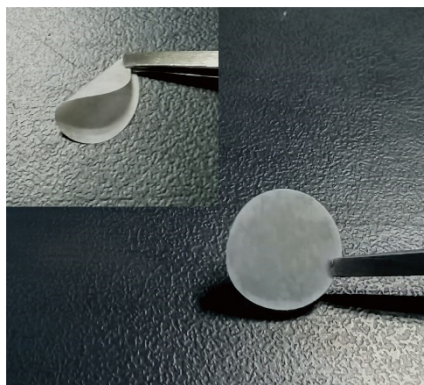


Fig. S3 Optical image of the P(M-B-V) membrane.

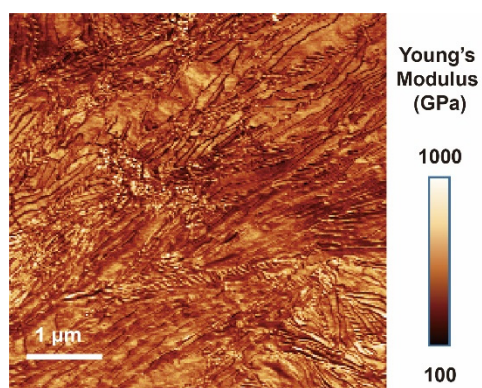


Fig. S4 AFM Young's modulus mapping of the PEO SPE.

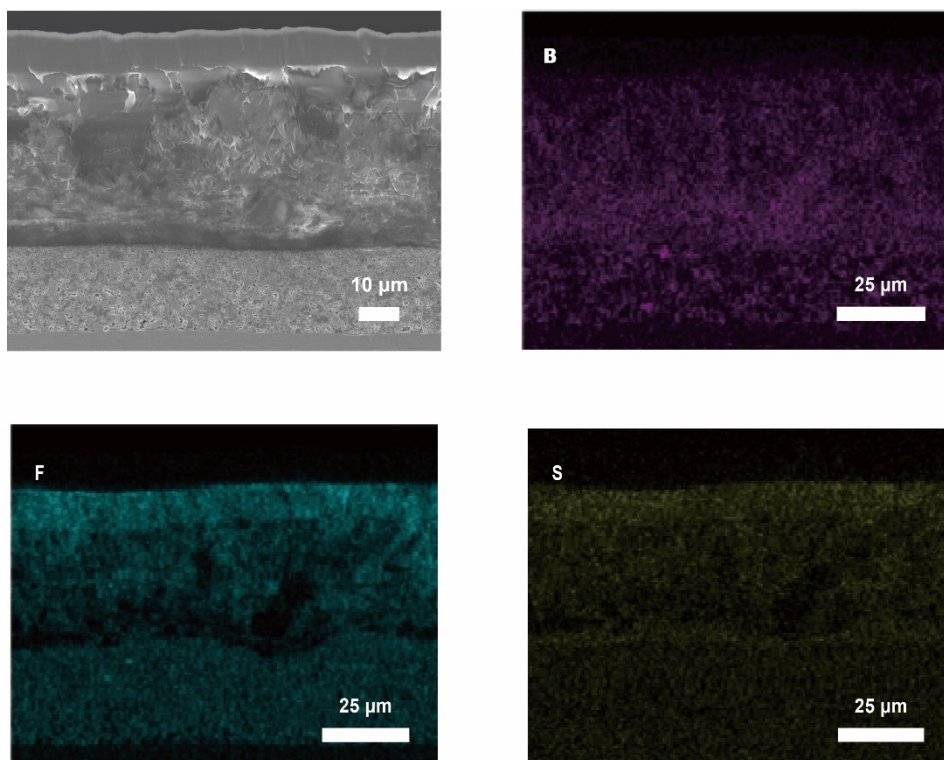


Fig. S5 The cross-sectional SEM image of the P(M-B-V) and corresponding EDX elemental mapping (B, F, and S).

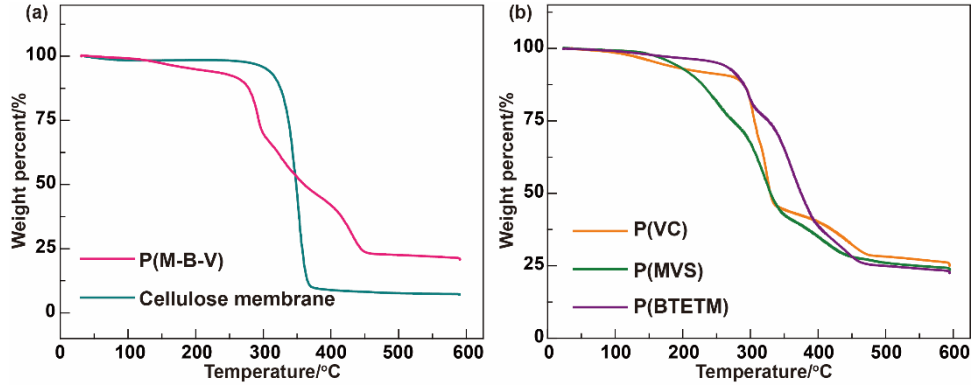


Fig. S6 (a-b) Thermogravimetric curves of the cellulose membrane, P(M-B-V), P(VC), P(MVS) and P(BTETM).

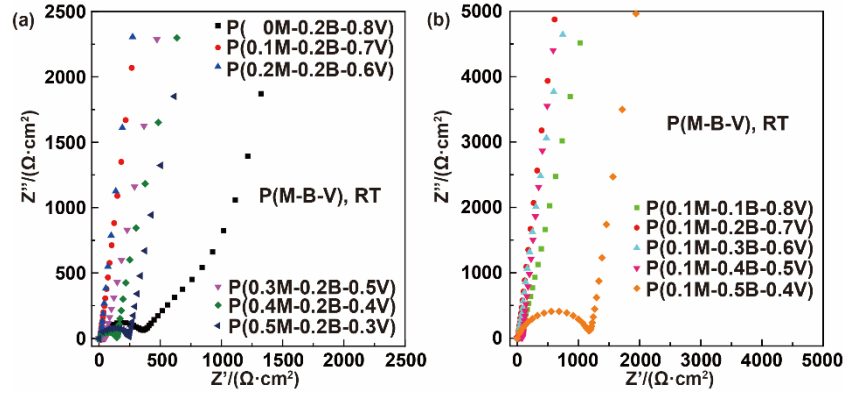


Fig. S7 (a-b) Nyquist plots of the P(M-B-V) SPEs with different weight fractions of MVS, BTETM and VC measured at RT.

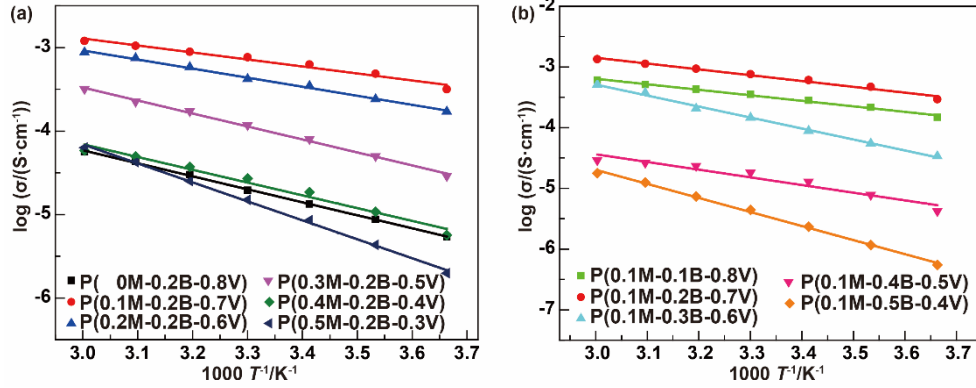


Fig. S8 (a-b) Arrhenius plots at temperatures from 0 to 60 °C for the P(M-B-V) SPEs with different weight fractions of MVS, BTETM, and VC.

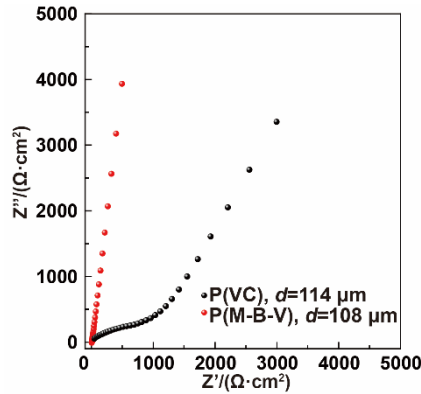


Fig. S9 Nyquist plots of the P(VC) and P(M-B-V) measured at RT.

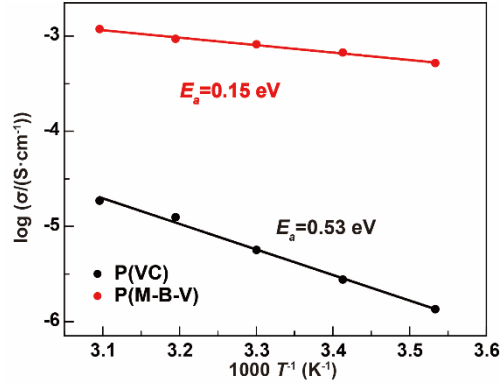


Fig. S10 Arrhenius plots at temperatures from 10 °C to 50 °C for the P(VC) and P(M-B-V).

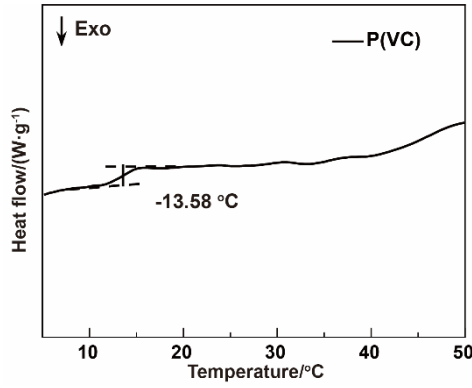


Fig. S11 DSC curve of the P(VC).

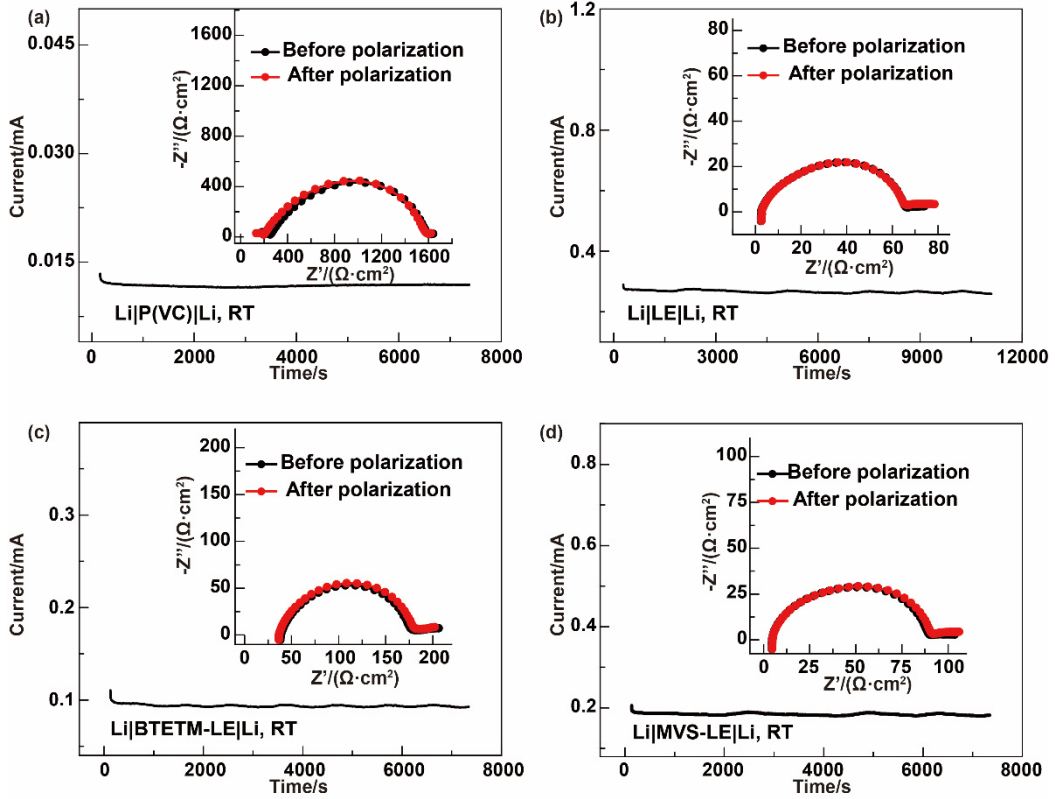


Fig. S12 (a–d) Current-time plots with Nyquist plots before and after polarization (inset) of symmetric Li cells with P(VC), liquid, BTETM-liquid and MVS-liquid electrolytes.

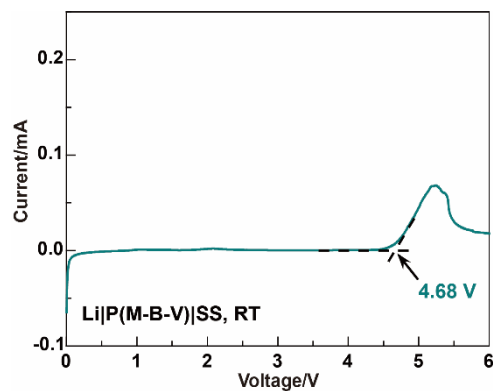


Fig. S13 Linear sweep voltammetry (LSV) curve of the P(M-B-V).

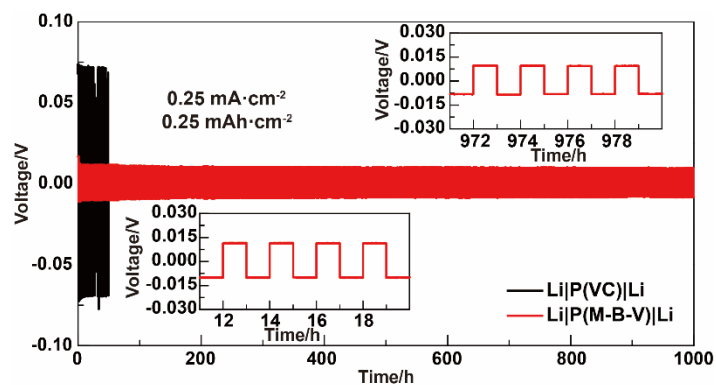


Fig. S14 Li plating/stripping cycling of symmetric Li cells containing P(VC) (black) and P(M-B-V) SISC (red) at a current density of $0.25 \text{ mA}\cdot\text{cm}^{-2}$ for 1 h at RT. The insets show magnified curves at the indicated time periods.

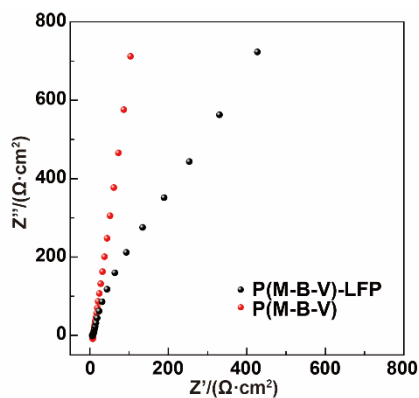


Fig. S15 Nyquist plots of the P(M-B-V)-LFP (the weight fraction of P(M-B-V) in the cathode is approximately 0.22%–0.25%) and the P(M-B-V) SISC.

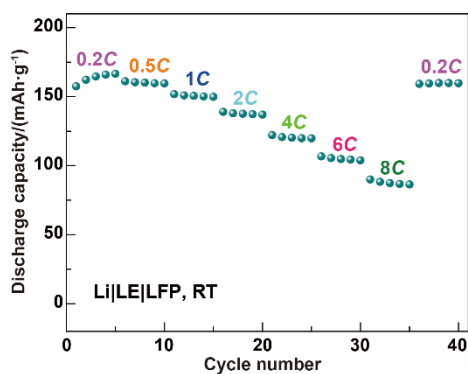


Fig. S16 Rate capability of Li|LFP batteries with liquid electrolyte at RT.

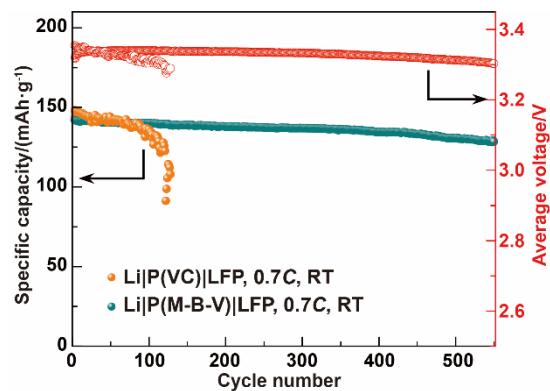


Fig. S17 Cycling performance of Li||LFP batteries with the P(M-B-V) SISC (dark cyan) and the P(VC) (orange) at 0.7C, RT.

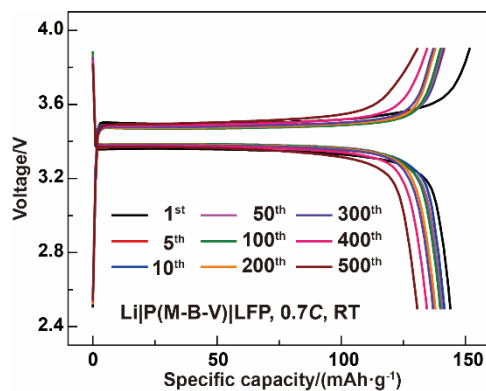


Fig. S18 Voltage profile evolution of the Li|P(M-B-V)|LFP battery cycled 0.7C, RT.

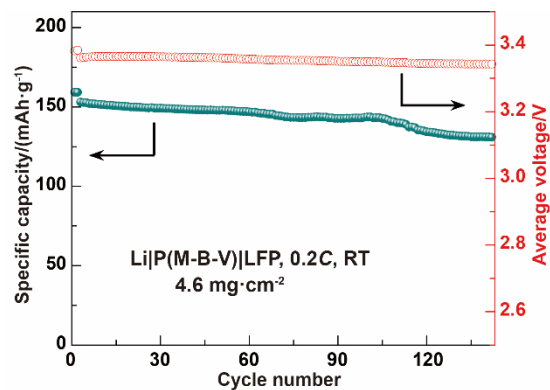


Fig. S19 Cycling performance of Li|P(M-B-V)|LFP battery with cathode areal loading as high as 4.6 mg·cm⁻² at 0.2C, RT.

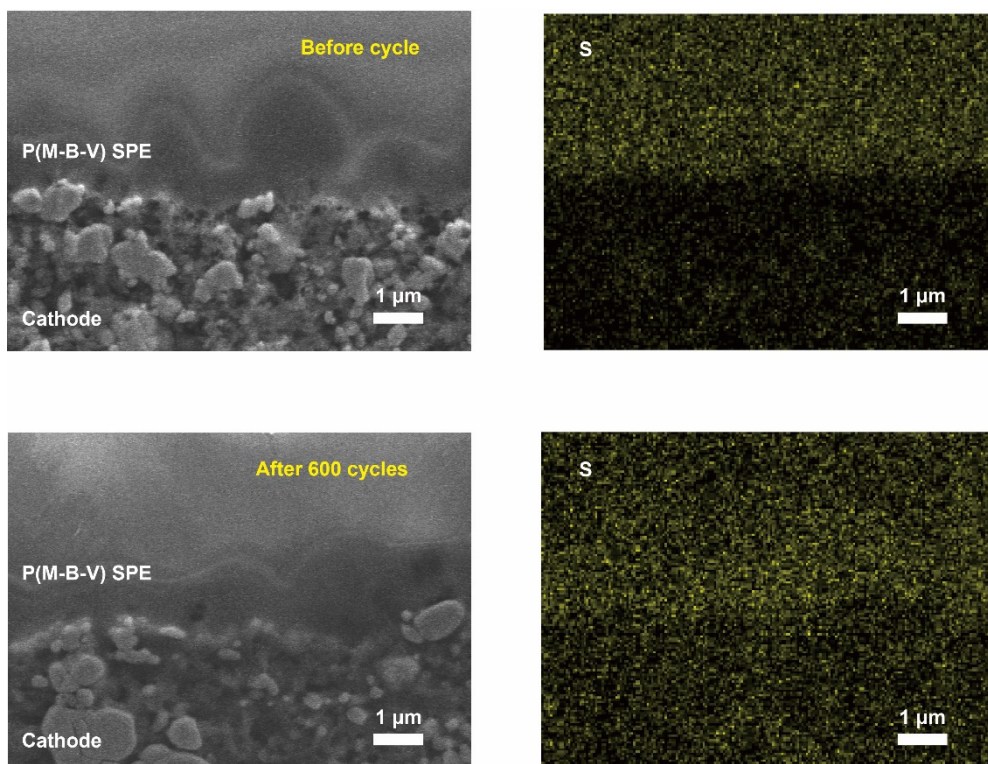


Fig. S20 SEM image and EDX elemental mapping (S) of the interface between the LFP cathode and the P(M-B-V) SISC before and after 600 cycles.

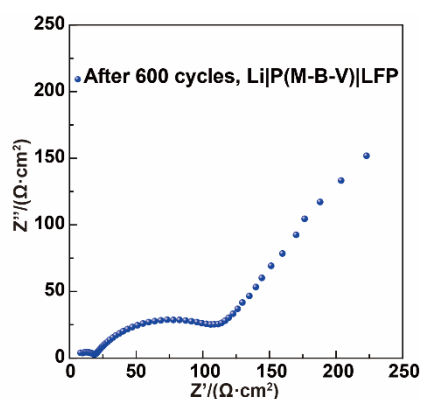


Fig. S21 Nyquist plot of the Li|P(M-B-V)|LFP battery after 600 cycles.

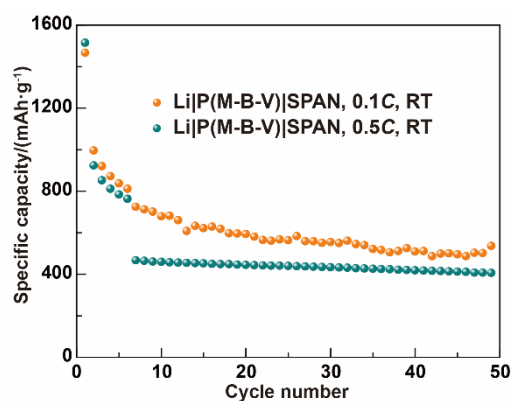


Fig. S22 Cycling performance of the Li|P(M-B-V)|SPAN batteries at 0.1C and 0.5C (activating at 0.05C for 6 cycles before the cycling).

Table S1 Lithium-ion transference number and the ambient ionic conductivity of the P(M-B-V) SPEs with different weight fractions of MVS, BTETM and VC.

	BTETM	MVS	VC	$\sigma/(\text{mS}\cdot\text{cm}^{-1})$	t_{Li^+}
1	0.2	0.0	0.8	0.021	0.63
2	0.2	0.1	0.7	1.290	0.94
3	0.2	0.2	0.6	0.353	0.67
4	0.2	0.3	0.5	0.096	0.88
5	0.2	0.4	0.4	0.055	0.78
6	0.2	0.5	0.3	0.038	0.82
7	0.1	0.1	0.8	0.945	0.90
8	0.3	0.1	0.6	0.489	0.80
9	0.4	0.1	0.5	0.119	0.68
10	0.5	0.1	0.4	0.005	0.58

Table S2 Lithium-ion transference number of the P(M-B-V), P(VC), liquid, BTETM-liquid and MVS-liquid electrolytes.

	R_0/Ω	R_s/Ω	I_0/mA	I_s/mA	t_{Li^+}
P(M-B-V)	333.1	354.6	0.0297	0.0279	0.94
P(VC)	674.0	692.0	0.0134	0.0119	0.48
LE	32.22	32.22	0.2890	0.2580	0.36
B-LE	69.79	72.72	0.1100	0.0926	0.60
M-LE	43.96	44.95	0.2050	0.1820	0.48

Table S3 Performance comparison of the P(M-B-V) SISC and the P(M-B-V) based solid state battery with recent works.

Material	Electrolyte type	$\sigma \times 10^4$ at room temperature ($\text{S}\cdot\text{cm}^{-1}$)	t_{Li^+}	Tensile stress/MPa	Cycle performance(battery-C rate-temperature-cycles-capacity retention)	Ref
B-PEG@DMC	Solid polymer electrolyte (SPE)	0.23	—	—	LFP/Li-0.2C-25 °C-100 th -73%	1
SLIMs	Gel polymer electrolyte (GPE)	0.27	0.62	0.40	LFP/Li-0.1C-25 °C -150 th -80%	2
NPEs	SPE	0.32	0.47	1.60	LFP/Li-0.5C-25 °C -100 th -92%	3
CPSHPE	SPE	0.89	0.26	0.43	LFP/Li-0.1C-30 °C-70 th -87.7%	4
PBI-g-LiPSTFSI	GPE	1.50	0.91	14.50	LFP/Li-1C-25 °C-500 th -87%	5
ANP-5	GPE	1.50	0.95	—	LFP/Li-0.5C-30 °C-100 th -84%	6
PVDF/PP	SPE	1.53	0.24	7.05	LFP/Li-0.3C-25 °C-180 th -97.8%	7
LITFSI/PUA	SPE	0.03	—	11.1	LFP/Li-0.03C-25 °C-20 th	8
PAS/PS	SPE	0.54/60 °C	0.96	—	LFP/Li-0.2C-25 °C-60 th -88.1%	9
IN-SCPEs	SPE	1.90	0.90	0.50	LFP/Li-0.5C-28 °C-200 th -83.2%	10
PIL-PEO	SPE	6.00	0.22	—	LFP/Li-0.5C-55 °C-50 th -77%	11
SHCPE	Composite polymer electrolyte (CPE)	0.80	0.43/60 °C	0.12	LFP/Li-0.2C-60 °C-60 th -95.8%	12
P(M-B-V) SISC	SPE	12.90	0.94	13.10	LFP/Li-0.5C-30 °C-630 th -90%	This work

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