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Supporting Information

Gel Electrolytes with Wide Potential Window for High-rate Al-ion Batteries

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Fig. S1. Raman spectra of the positive electrodes: pristine, charged-2.4 V and charged-2.6 V.



Fig. S2. C 1s peak of the graphitic positive electrodes: pristine, charged-2.4 V and charged-2.6 V.



Fig. S3. EDS analysis of the graphitic positive electrodes: (a) Pristine, (b) Charged-2.4 V, (c) Charged-2.6 V and (d) Discharged-0.3 V. The insets are the corresponding EDS mapping images.

Element	Weight (%)	Atom (%)
C K	100.0	100.0

Table. S1. EDS element analysis results of the original positive electrode.

Element	Weight (%)	Atom (%)
C K	76.75	90.04
Al K	5.76	3.01
CI K	17.49	6.95

Table. S2. EDS element analysis results of the charged-2.4 V positive electrode.

Element	Weight (%)	Atom (%)
C K	69.58	86.21
Al K	7.72	4.26
Cl K	22.70	9.53

Table. S3. EDS element analysis results of the charged-2.6 V positive electrode.

Element	Weight (%)	Atom (%)
СК	86.08	94.5
Al K	2.72	1.33
Cl K	11.20	4.17

Table. S4. EDS element analysis results of the discharged-0.3 V positive electrode.



Fig. S4. CV curves of the solid-state AIB at a scan rate of 1 mV s⁻¹ in the voltage range of 0–2.70 V.



Fig. S5. The charge/discharge curves of the Et₃NHCl-based batteries with different bending states: (a) Liquid system, (b) Quasi-solid system. (c) The AIB with Et₃NHCl-based gel electrolytes operating stably under different bending conditions.



Fig. S6. (a) Galvanostatic charge and discharge profiles of the solid-state batteries at different current densities. (b) The initial three charge and discharge curves. (c) The charge and discharge curves at different cycles. (d) The charge and discharge curves at the charge current density of 500 mA g⁻¹ and the discharge current density of 3000 mA g⁻¹.



Fig. S7. (a) Cycling stability of the solid-state AIBs at lower temperature (current density of 200 mA g^{-1}). (b) The charge and discharge curves at different temperatures (current density 200 mA g^{-1}).



Fig. S8. The comparison of the LSV curves of the two kinds of gel-polymer electrolyte systems.

with corresponding temperatures							
T / °C	-20	0	25	40	60	80	100
σ / S cm ⁻¹	2.95×10 ⁻⁴	1.04×10 ⁻³	4.52×10 ⁻³	7.48×10 ⁻³	1.07×10 ⁻²	1.47×10 ⁻²	1.83×10 ⁻²

Table. S5. The values of the ionic conductivities (σ) of Et₃NHCl-based gel-polymer electrolyte with corresponding temperatures



Fig. S9. (a) Raman spectra (b) ₂₇Al NMR spectra (c) FTIR spectra of the samples during preparation.



Fig. S10. (a) TG curves of monomer and polymer skeleton of the electrolyte. (b) The mass change during the heating process of the Et₃NHCl-based gel electrolyte.



Fig. S11. The load-displacement curve of the Et₃NHCl-based gel electrolyte.



Fig. S12. The electrochemical impedance spectroscopy measurements. (a,b) The Nyquist plots of the solid-state battery after 0, 1, 50 and 100 cycles at a frequency range of 100 kHz to 0.01 Hz with an amplitude of 5 mV. (c) Equivalent circuit model of the studied system.