

Supporting Information

Quasi-solid electrolyte developed on hierarchical rambutan-like γ -AlOOH microspheres with high ionic conductivity for lithium ion batteries

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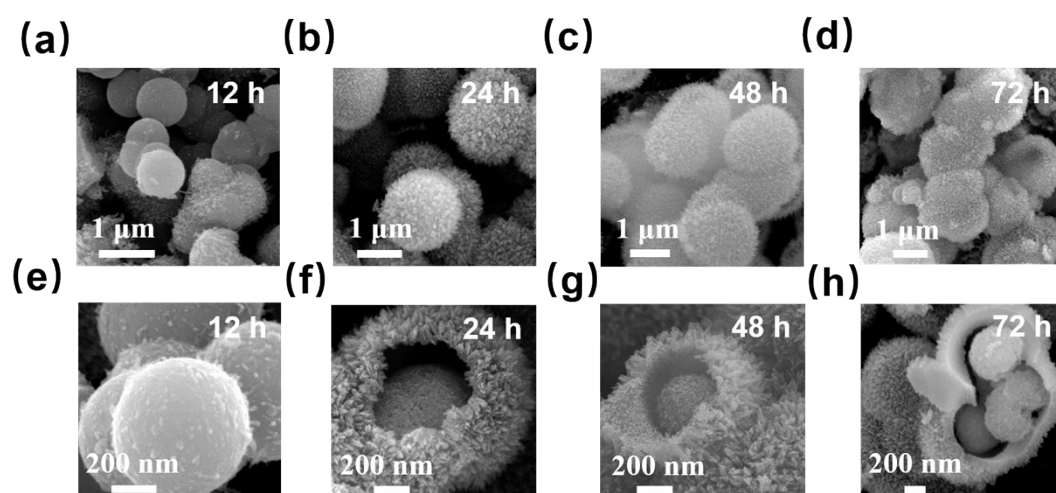


Fig. S1 SEM images for the products at different reaction times.

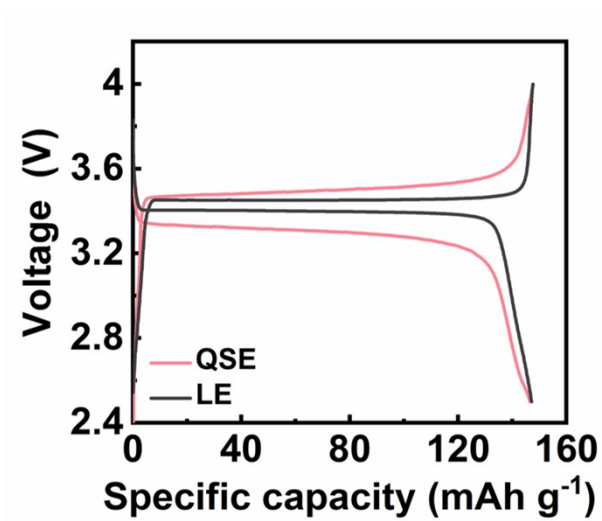


Fig. S2 The initial charge/discharge profiles of LFP/ γ -AlOOH-QSE/Li and LFP/QSE/Li cells.

The initial discharge specific capacities of the LFP/ γ -AlOOH-QSE/Li and LFP/LE/Li cells are similar at 0.1 C (146.6 vs. 147.1 mAh g⁻¹), though the latter one exhibits slightly a higher capacity retention (99.3% vs. 99.7%).

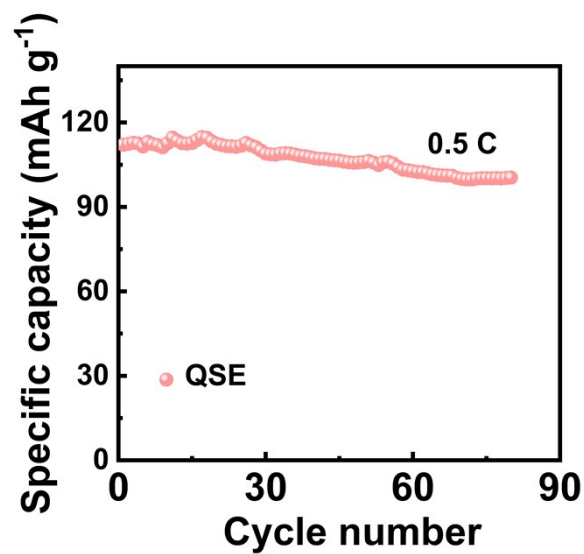


Fig. S3 Cycling performance of the assembled LFP/ γ -AlOOH-QSE/Li cell at 0.5 C.

Although the specific capacity of LFP/AlOOH-QSE/Li degrades faster at high current rate compared with that of LE (Fig. 2b), the capacity retention could still be remained 95.6% after 80 cycles under high current density of 0.5 C.

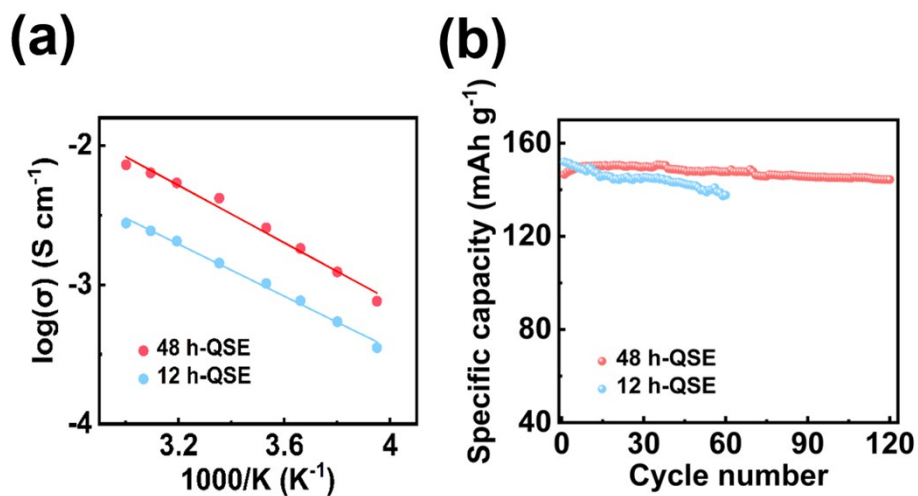


Fig. S4 (a) Ionic conductivities of 48 h-QSE, 12 h-QSE symmetrical cells within the temperature range of -20 – 80 °C; (b) Cycling performances of the assembled cells with 48 h-QSE and 12 h-QSE as electrolytes.

According to Fig. S4b, long-term cycling performance of the assembled cells with 48 h-QSE electrolytes is better than that with 12 h-QSE, further verifying the important influence of morphologies of γ -AlOOH microspheres on the properties of QSE materials.

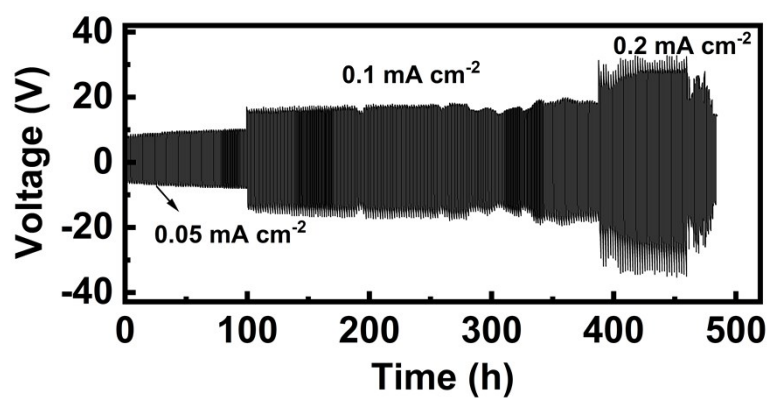


Fig. S5 Time-dependent voltage profiles for lithium plating/stripping experiment for Li/LE/Li symmetric coin cell under the current densities of 0.05, 0.1, and 0.2 mA cm⁻².

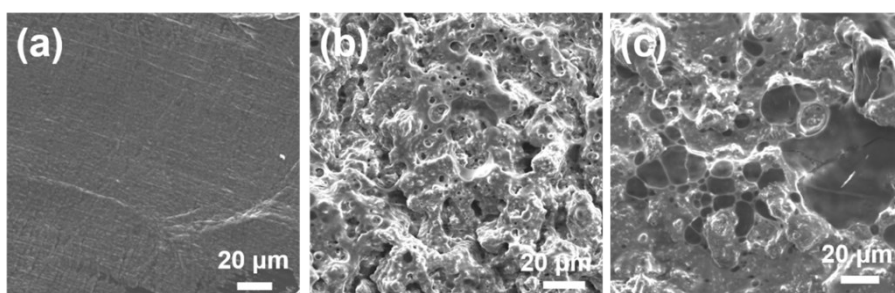


Fig. S6 Surface morphologies of Li anodes. (a) The fresh pristine Li anode; (b) LFP/ γ -AlOOH-QSE/Li cell; (c) Conventional LFP cell with LE.

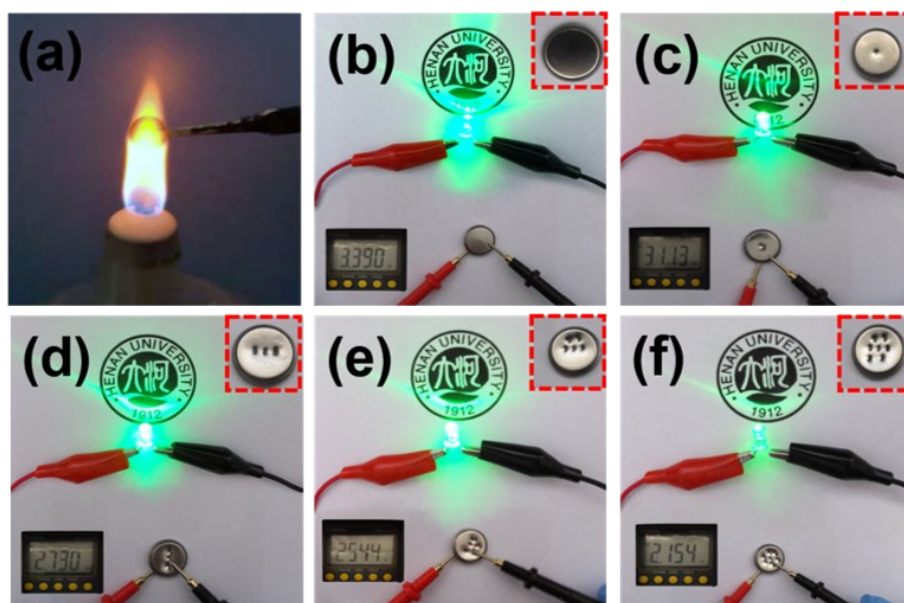


Fig. S7 (a) Combustion test of the γ -AlOOH-QSE sheet; (b-f) Illustration of lighting LED lamp under continuous puncture treatment.

Table S1 Impedances values for assembled LFP/ γ -AlOOH-QSE/Li cell at different cycles under 0.1 C.

	20th	60th	100th
$R_{ct}(\Omega)$	2489	2858	3030
$W_1(\Omega)$	81	78	83

Table S2 The ionic conductivities of previously proposed QSEs.

QSE	Ionic conductivity ($S\ cm^{-1}$)	Ref.
hollow silica (HS)	2.50×10^{-3} (25 °C)	1
ZIF-8	1.05×10^{-4} (25 °C)	2
Porous molecular cage	1.00×10^{-3} (25 °C)	3
$Mg_2B_2O_5/PEO$	1.53×10^{-4} (40 °C)	4
$Li_{1.3}Al_{0.3}Ti_{1.7}(PO_4)_3$	8.05×10^{-5} (25 °C)	5
MCM-41	6.37×10^{-4} (30 °C)	6
SiO_2	1.74×10^{-4} (25 °C)	7
$\gamma-Al_2O_3$	1.10×10^{-3} (25 °C)	8
MOF	3.10×10^{-4} (25 °C)	9
PVDF-HFP/ Al_2O_3	1.00×10^{-3} (25 °C)	10
PVDF- PEO/ $Li_{1.4}Al_{0.4}Ti_{1.6}(PO_4)_3$	5.24×10^{-4} (25 °C)	11
PAN/ $Li_{1.4}Al_{0.4}Ti_{1.6}(PO_4)_3$	6.50×10^{-4} (60 °C)	12
SSZ-13/PEO	1.91×10^{-4} (60 °C)	13
COF/ $LiClO_4$	0.26×10^{-3} (25 °C)	14
γ -AlOOH-QSE	4.00×10^{-3} (25 °C)	This work
PVDF-HFP / $Li_7La_3Zr_{12}O_{12}$	1.10×10^{-4} (25 °C)	15

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