

Supporting Information

Highly Efficient Multicomponent Gel Biopolymer Binder Enables Ultrafast Cycling and Applicability in Diverse Battery Formats

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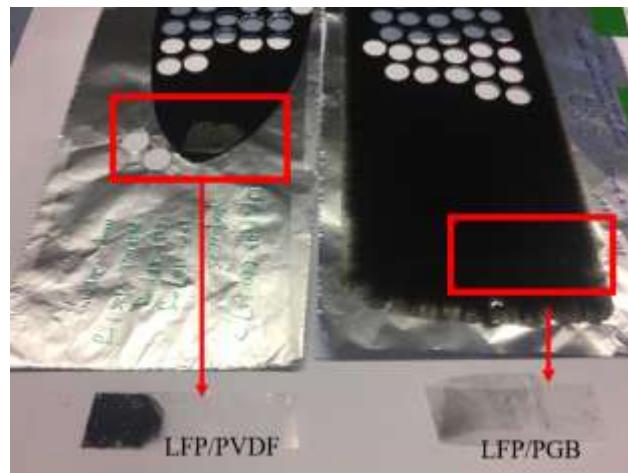


Figure S1. Simplified peeling test using tapes on LFP/PVDF and LFP/PGB electrodes.

PGB: polymer gel binder.

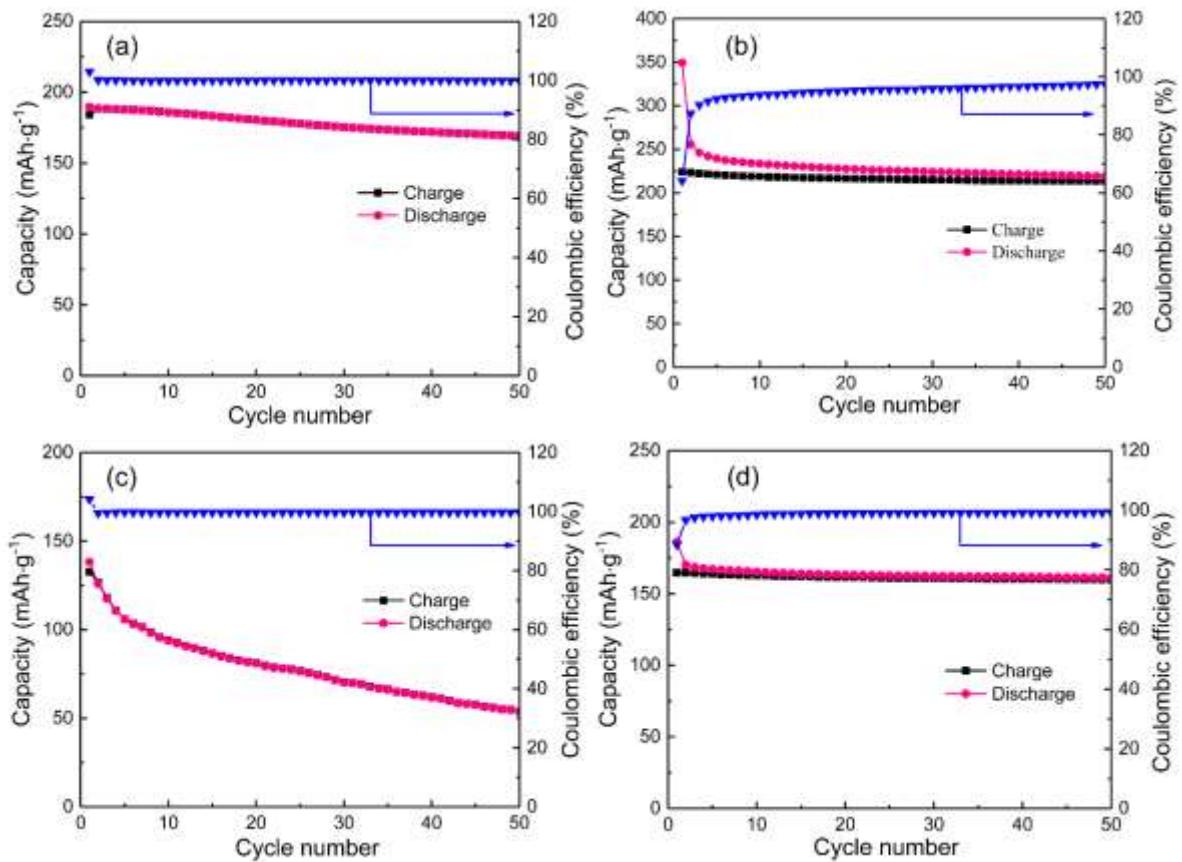


Figure S2. Cycling performance and Coulombic efficiency of (a) LFP/PGB (b) LTO/PGB (c) LFP/PVDF and (d) LTO/PVDF half cells cycled at low rate of 0.5C.

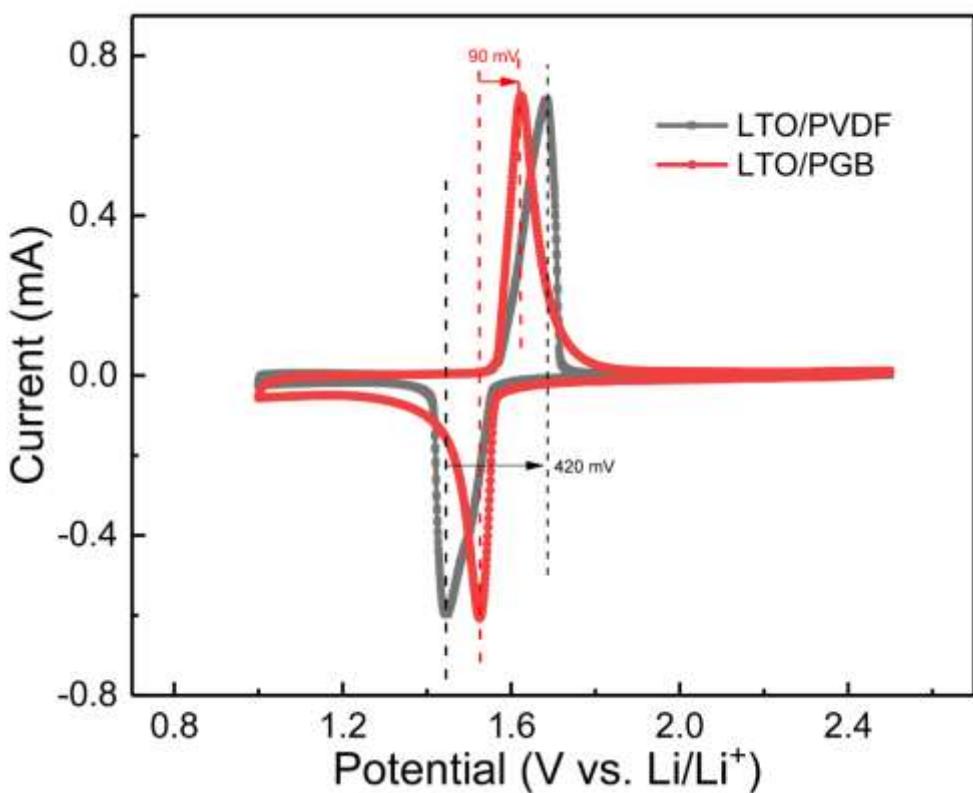


Figure S3. CV profiles of LTO/PGB and LTO/PVDF electrodes. The tests were performed in a voltage range of 1-2.5 V at a scanning rate of $0.1 \text{ mV}\cdot\text{s}^{-1}$.

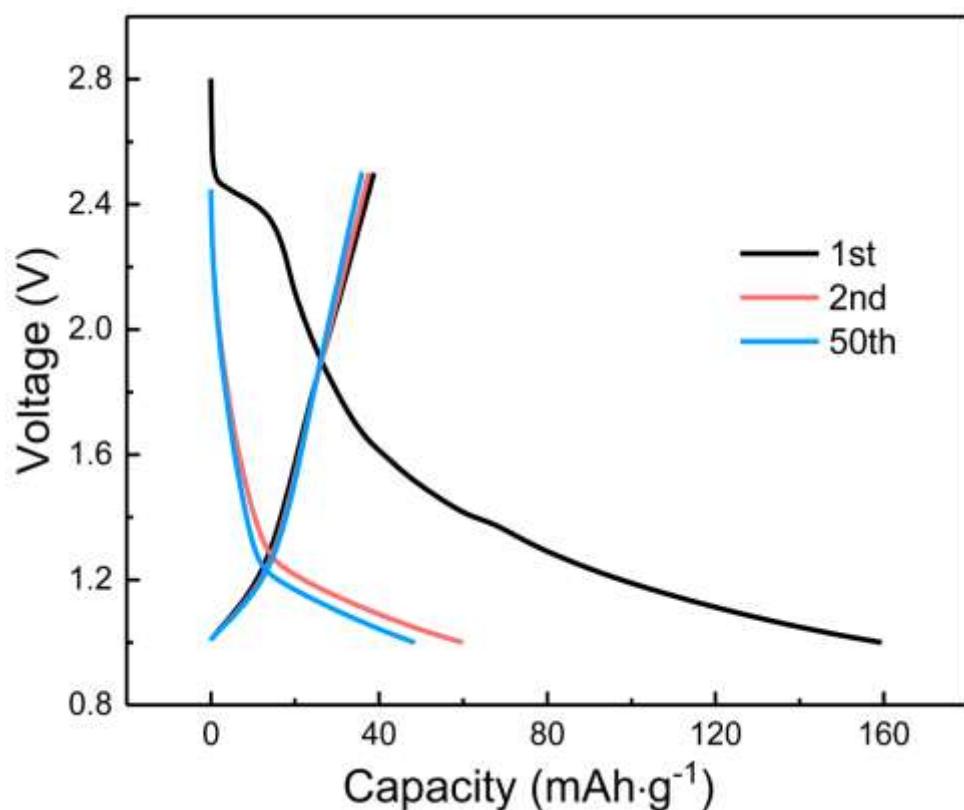


Figure S4. Charge/discharge voltage profiles of the electrode containing PGB and carbon black (prepared with a weight ratio of 1:1) cycled with a current density of 50 mA·g⁻¹.

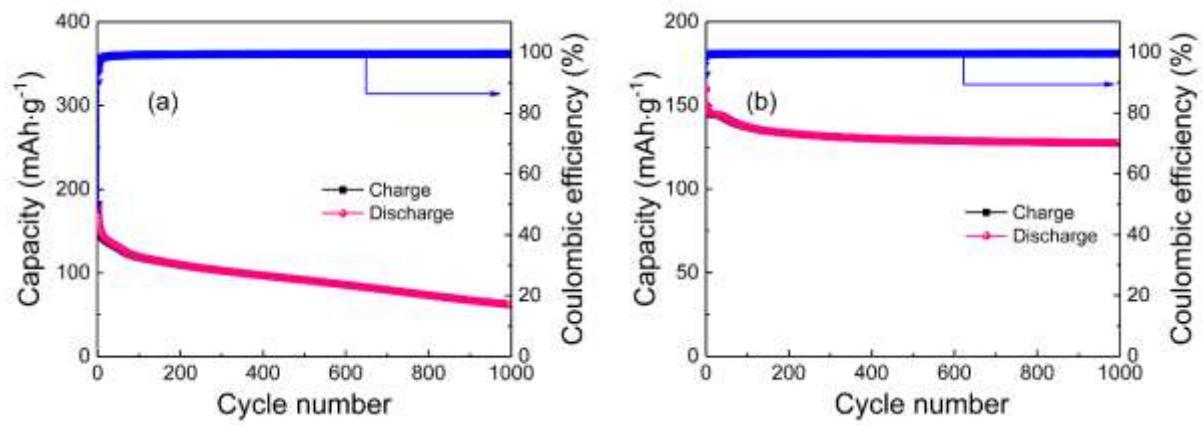


Figure S5. Cycling performance and Coulombic efficiency of (a) LTO/PGB and (b) LTO/PVDF half cells cycled at high rate of 10C.

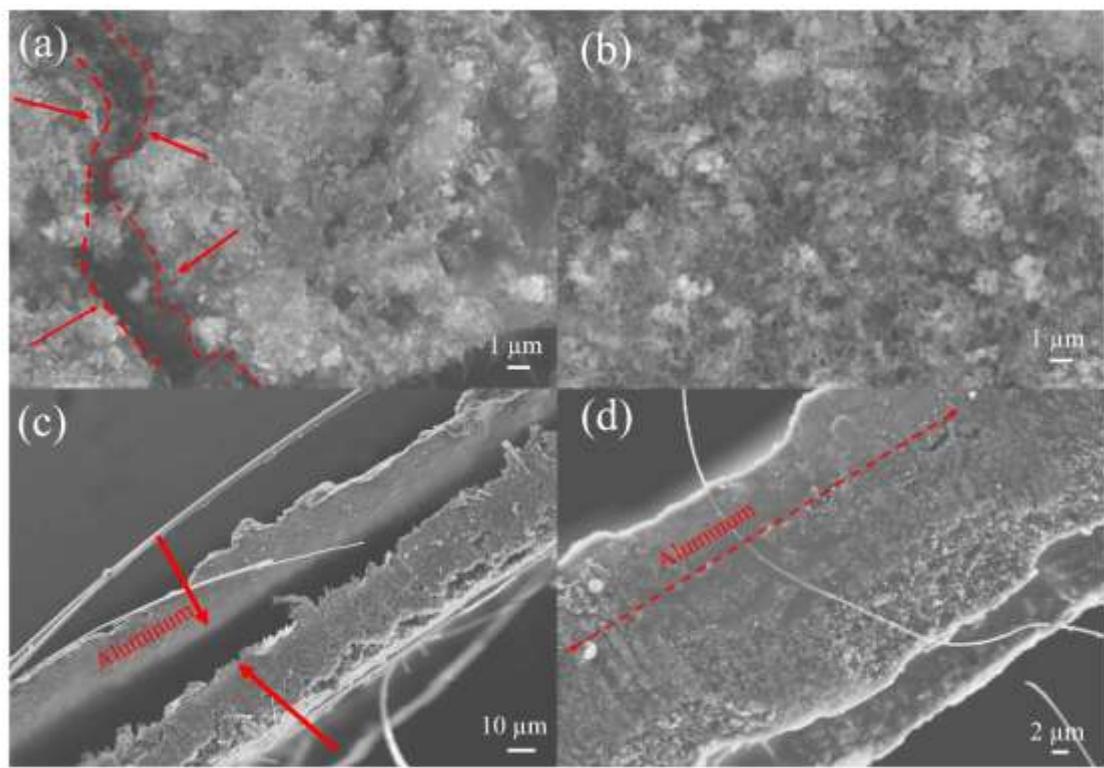


Figure S6. SEM images of (a) LFP/PVDF, (b) LFP/PGB surface after 200 cycles at 10C. Cross-sectional SEM images of (c) LFP/PVDF, (d) LFP/PGB after 200 cycles at 10C.

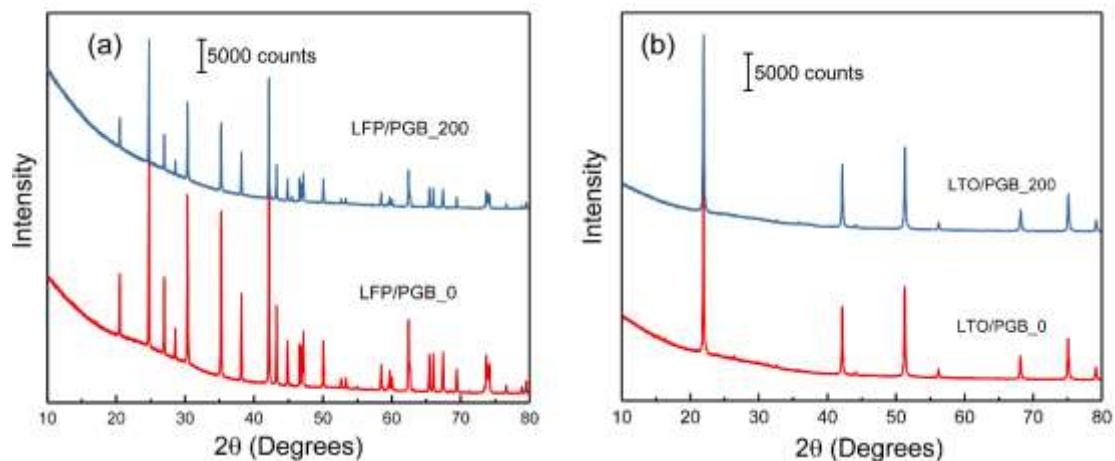


Figure S7. XRD patterns of electrodes before and after 200 cycles at 10C (a) LFP/PGB half cell (b) LTO/PGB half cell (performed using Co K_{α1}, $\lambda=1.7889$ Å).

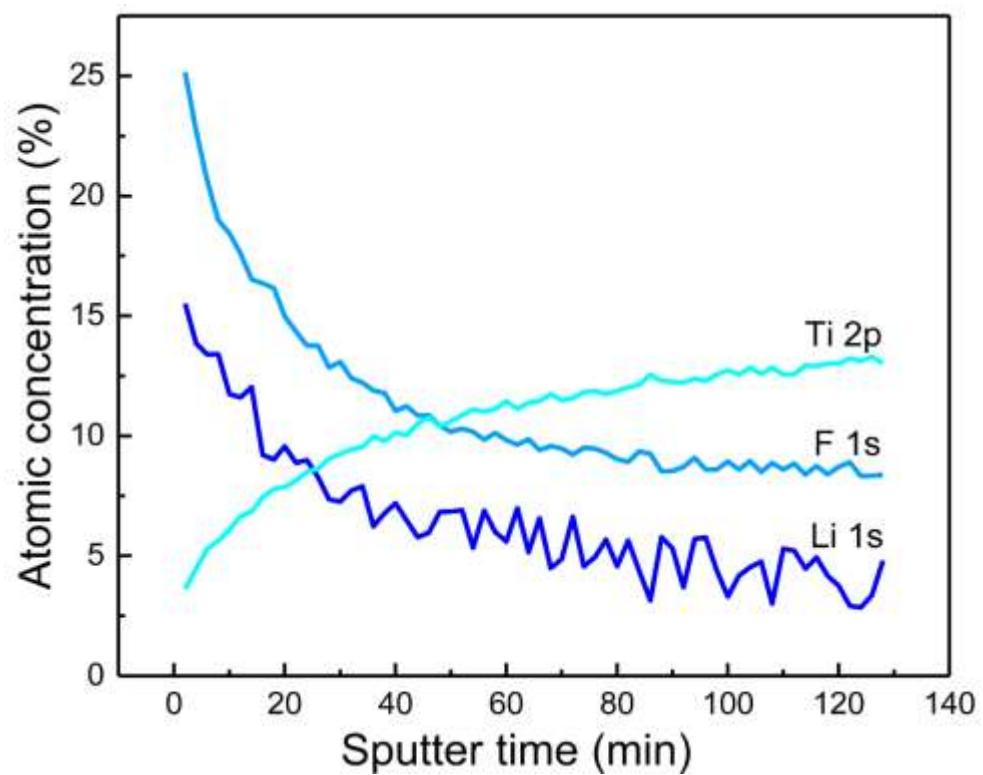


Figure S8. Depth profiling of a LTO/PGB electrode after 50 cycles at 10C vs. Li/Li⁺.

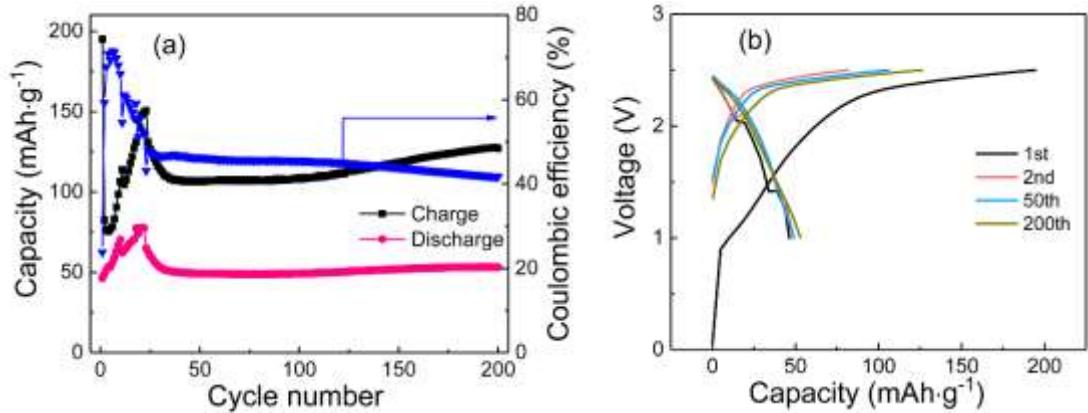


Figure 9. (a) Cycling performance and (b) charge/discharge voltage profiles of LFP||LTO-PVDF full cell at 10C.

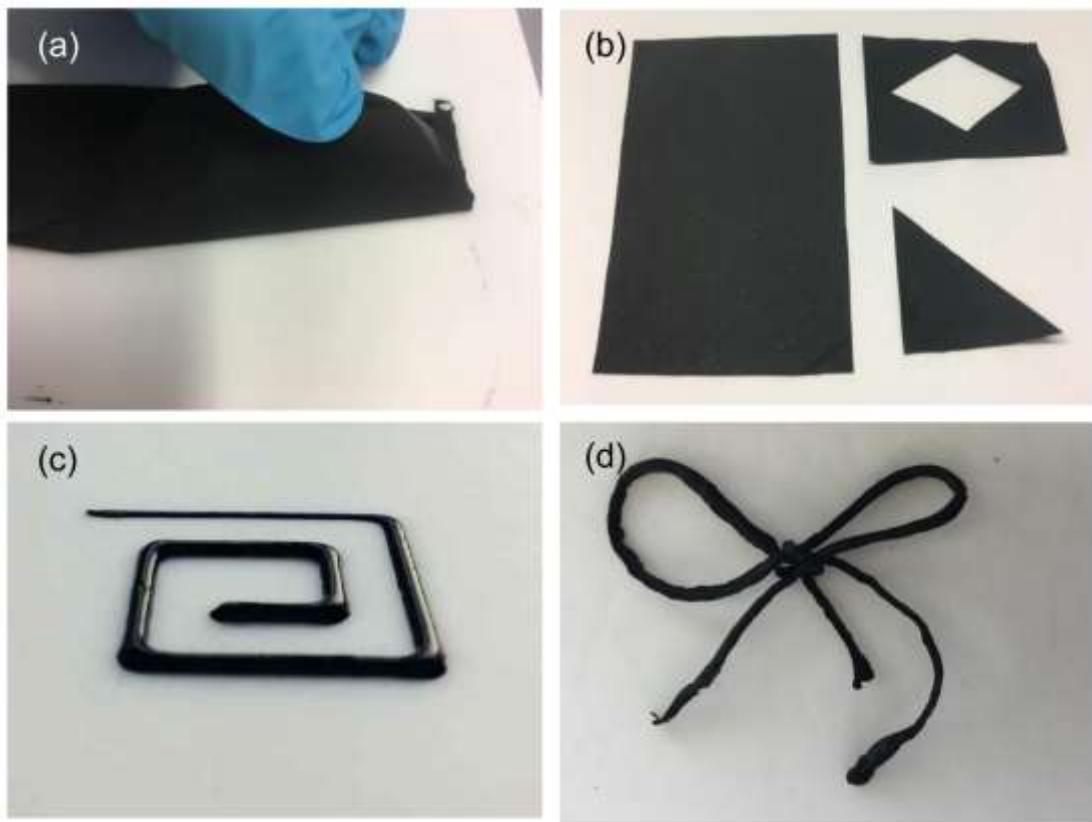


Figure S10. (a) The freestanding electrodes can be easily peeled off from the Teflon substrate and (b) can be cut into designed structures; (c) The electrode slurry can be directly printed into intricate structures as well as (d) extruded into flexible fibers which can be made into various shapes easily.

Table S1. Comparison of electrochemical performance between LFP/PGB and LTO/PGB with previously reported literature utilizing different modifications of LFP and LTO.

Modifications	Capacity (mAh·g ⁻¹)	Rate	Particle size
LFP/PAB-C ¹	137.5 (300 th)	10C	
C-LFP ²	139.8 (50 th)	1C	~90 nm
Nano-LFP ³	123 (1 st)	10C	<65 nm
C-LFP ⁴	130 (1 st)	5C	146 nm
LFP Nanoplates ⁵	148 (1 st)	10C	<200 nm
LFP@G ⁶	139.8 (600 th)	10C	<200 nm
<i>LFP/PGB (this work)</i>	<i>110 (200th)</i>	<i>10C</i>	<i>~400 nm</i>
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N-LTO@TiC/C ⁷	143 (1 st)	10C	
LTO-RTO/C ⁸	121.6 (200 th)	2C	
LTO-TiO ₂ ⁹	94.2 (1 st)	15C	
Hollow LTO microspheres ¹⁰	139 (1 st)	10C	
Microporous LTO microbars ¹¹	141 (1 st)	10C	
<i>LTO/PGB (this work)</i>	<i>112 (200th)</i>	<i>10C</i>	

Table S2. Elemental concentration measured by XPS on three LTO/PGB electrodes

prepared under different conditions.

	Li 1s	C 1s	N 1s	O 1s	F 1s	P 2p	S 2p	Ti 2p	Cu 2p
Pristine	0.66	61.96	5.59	25.52	1.90	0.18	0.11	2.06	2.02
Soaked	1.34	54.68	6.71	21.73	9.84	0.77	0.23	2.47	2.26
After 50 cycles	11.63	37.94	1.05	14.49	30.22	3.89	0.48	0.13	0.18

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