

Supporting Information

Spatially Confined Synthesis of Flexible and Hierarchically Porous Three Dimensional Graphene/FeP Hollow Nanospheres Composite Anode for Highly Efficient and Ultrastable Potassium Ion Storage

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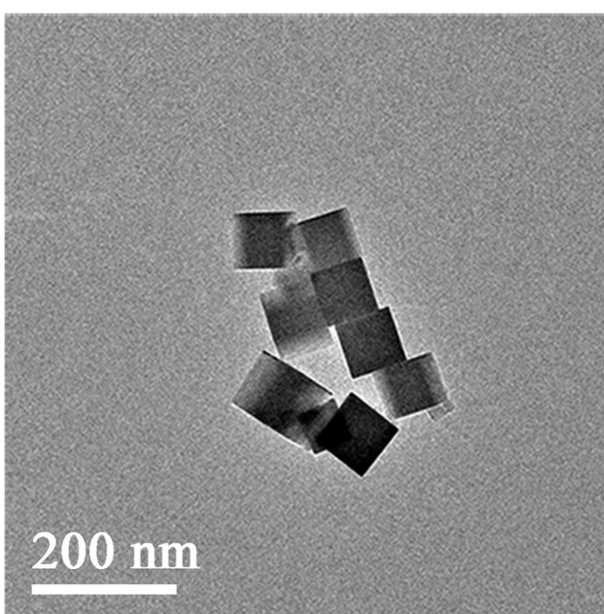


Fig. S1. Pure PB synthesized in the absence of GO under the same procedure as the GO/PB composites.

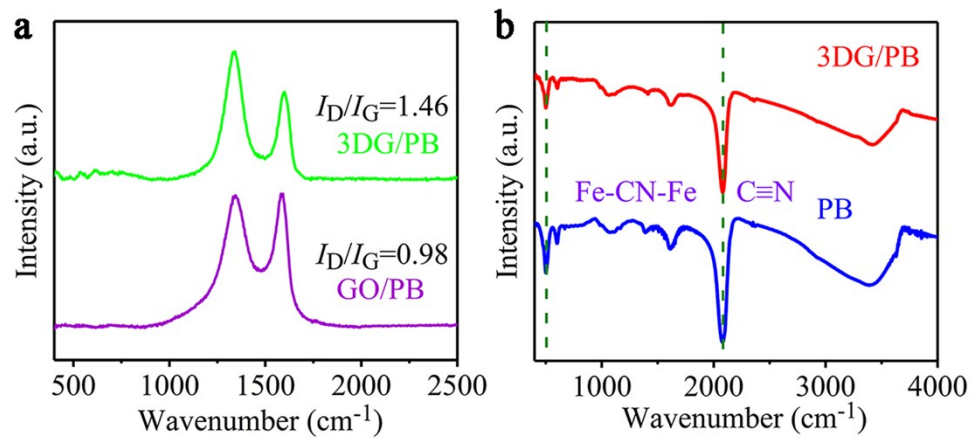


Fig. S2. (a) Raman spectra of GO/PB and 3DG/PB. (b) FTIR spectra of PB and 3DG/PB.

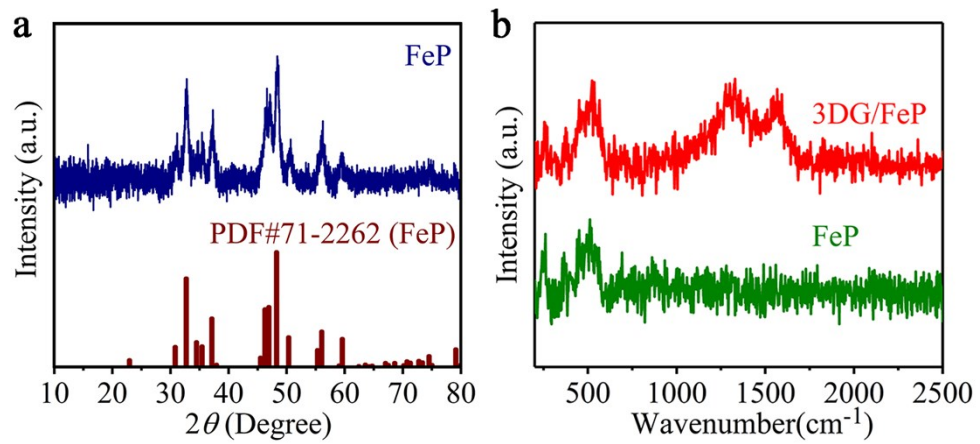


Fig. S3. The XRD pattern of (a) FeP, (b)3DG/FeP and FeP.

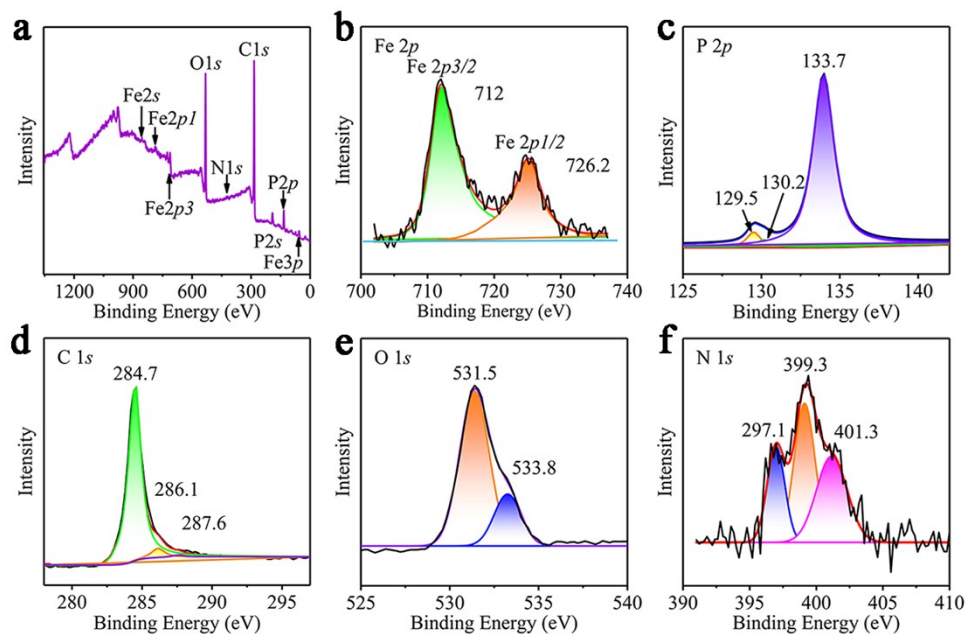


Fig. S4. (a) XPS survey spectrum, (b) Fe 2p, (c) P 2p, (d) C 1s, (e) O 1s and (f) N 1s spectra of the 3DG/FeP.

In the XPS survey spectrum of the 3DG/FeP, the elements of Fe, N, C, P could be detected, which indicated the successful phosphorization of 3DG/PB (Fig. S2a). Peaks at 712 and 726.2 eV corresponds to the Fe 2p, and peaks located at 129.5 and 130.2 eV are assigned to P 2p_{1/2} and P 2p_{3/2} of P-Fe respectively (Fig. S2b and c). Furthermore, the P 2p peaks at 133.4 eV can be related to the P=O. The peaks at 284.7, 286.1, and 287.6 eV of C 1s show the existence of C-C (sp² C), C-N, and C=O bonds, respectively (Fig. S2d and e). The high intensity C-C peak indicates that the GO has been well reduced to rGO. In addition, N signals (397.1, 399.3, and 401.3 eV) are also observed for 3DG/FeP (Figure S2f), which comes from the -C≡N- in PB.

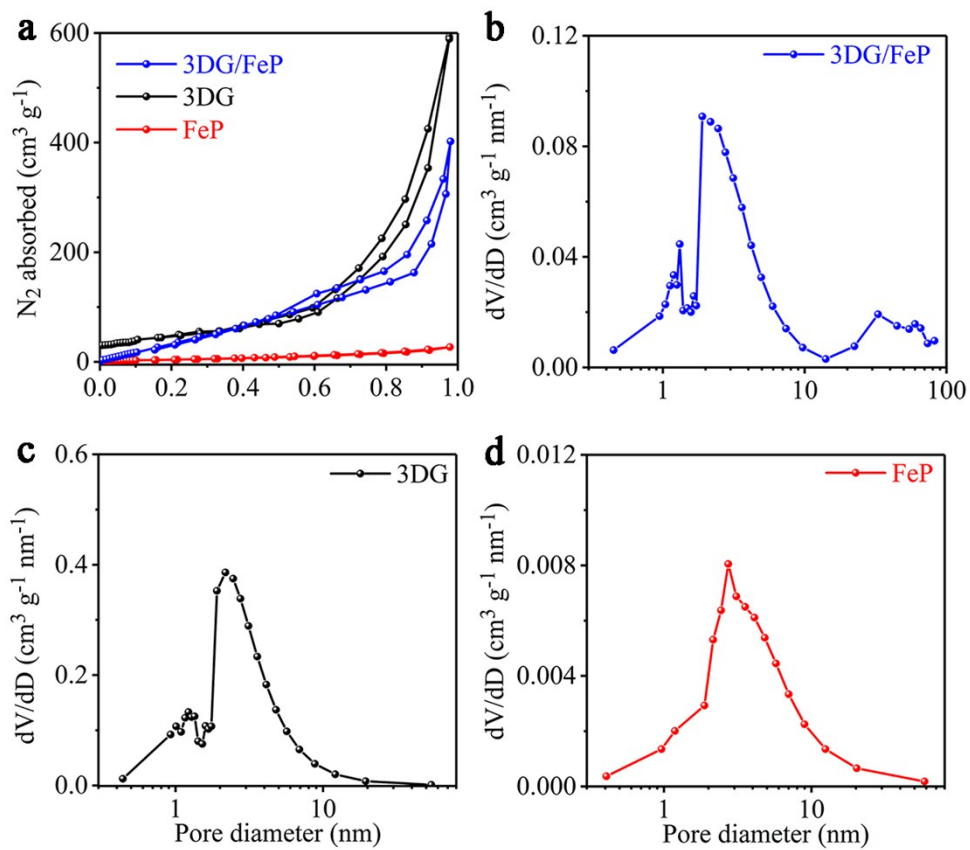


Fig. S5. (a) The N_2 adsorption-desorption isotherms, the pore-size distribution of (b) 3DG/FeP, (c) 3DG, and (d) FeP.

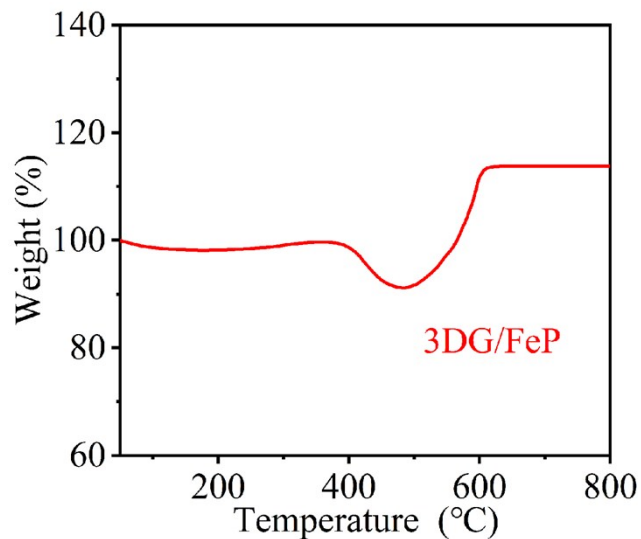
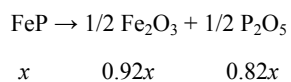


Fig. S6. The TGA curve of 3DG/FeP in air at a heating rate of 10 °C min⁻¹.

FeP content was measured in detail according to the calculation method of the reported carbon-based FeP.^{S1} The weight content of FeP in samples were estimated as follows:



Assuming the FeP content is x , the weight fractions for FeP was calculated as the following formula: $0.92x + 0.82x - (1-x) = 114\%$. Therefore, the FeP content in the 3DG/FeP was about 78 %.

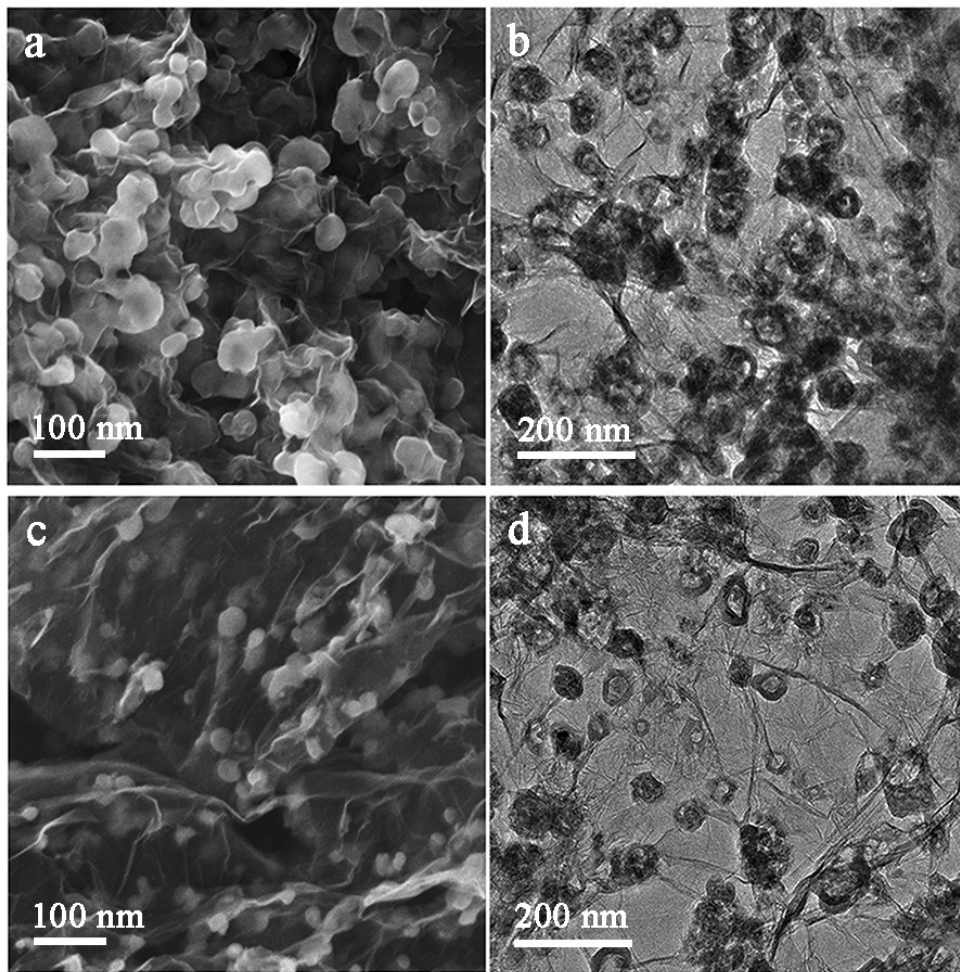


Fig. S7. SEM images of 3DG/FeP-H (a), 3DG/FeP-L (c). TEM images of 3DG/FeP-H (b), 3DG/FeP-L (d) (about 84% of FeP in 3DG/FeP-H and 68% of FeP in 3DG/FeP-L)

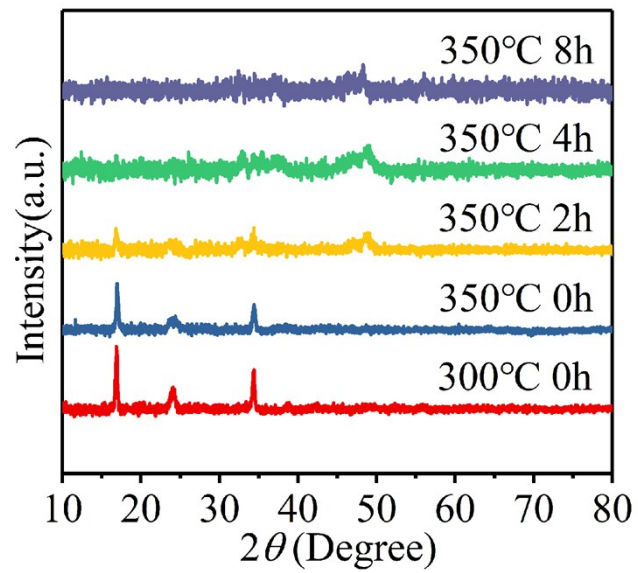


Fig. S8. XRD patterns of the obtained products collected at different reaction temperature for different reaction time, (a) 300 °C 0 h, (b) 350 °C 0 h, (c) 350 °C 2 h, (d) 350 °C 4 h, (e) 350 °C 8 h.

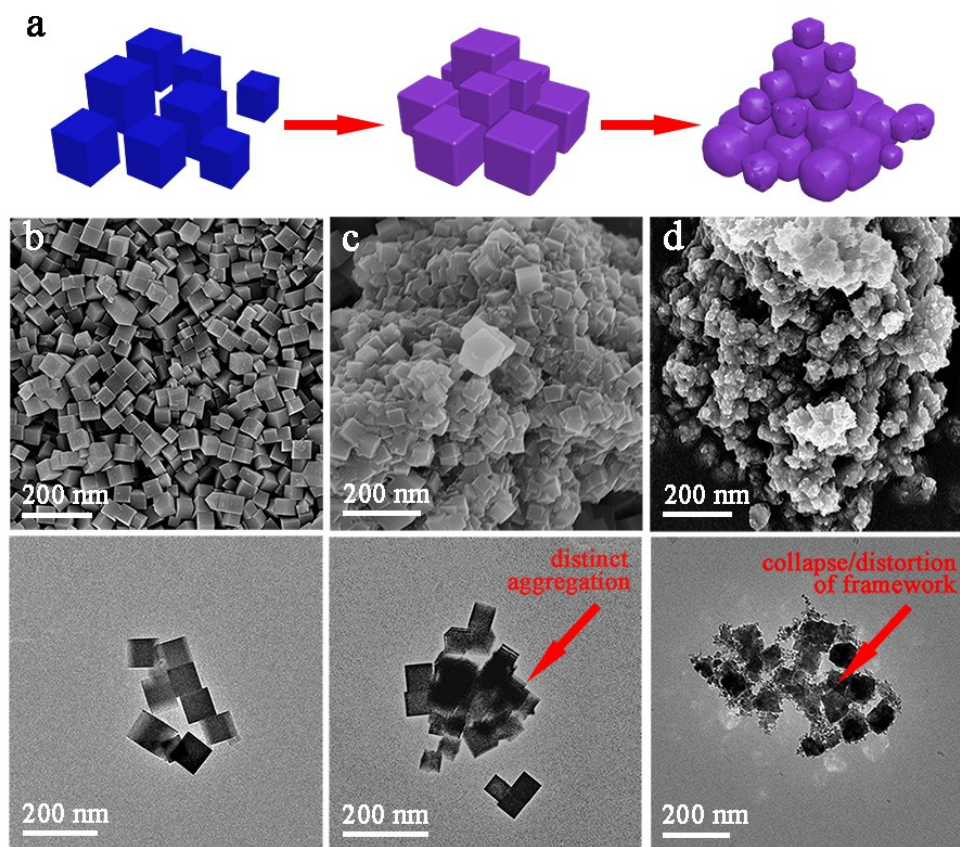


Fig. S9. (a) Thermal conversion diagram of PB. The SEM and TEM images of pure PB (b), FeP after 2 h (c) and 8h (d) thermal doping treatment.

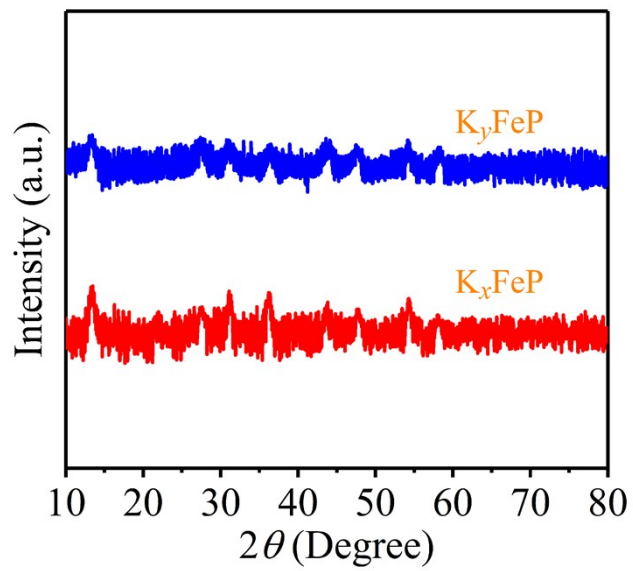


Fig. S10. XRD patterns of the $K_x\text{FeP}$ (0.9 V) and $K_y\text{FeP}$ (0.6 V) during the first discharge process.

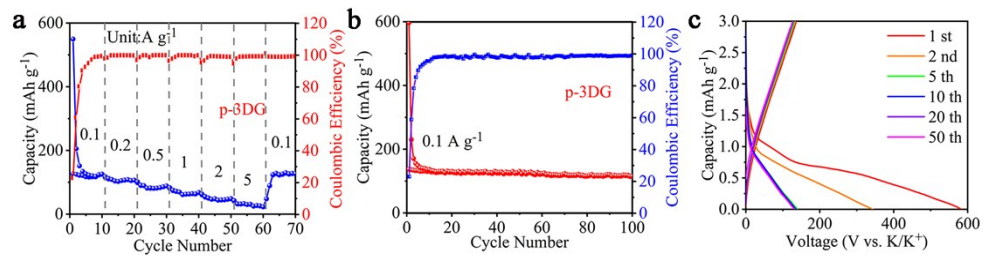


Fig. S11. (a) The rate performance (b) cycling performance of p-3DG anode at 0.1 A g⁻¹ for 100 cycles. (c) The charge/discharge curves of p-3DG at 0.1 A g⁻¹.

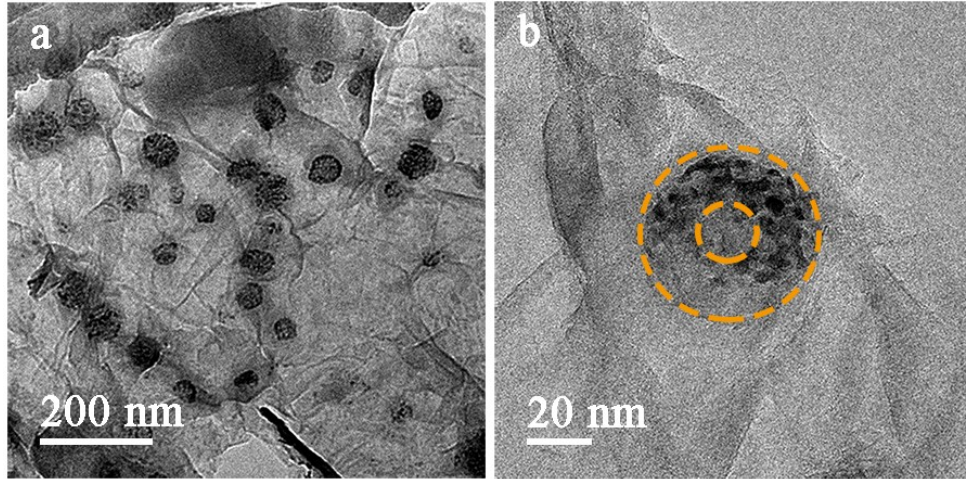


Fig. S12. The TEM images of 3DG/FeP anode after 200 cycles.

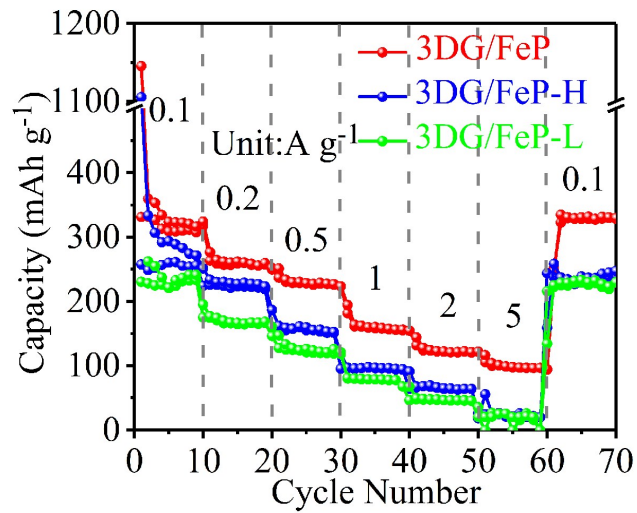


Fig. S13. The rate performance of 3DG/FeP with different mass content of FeP.

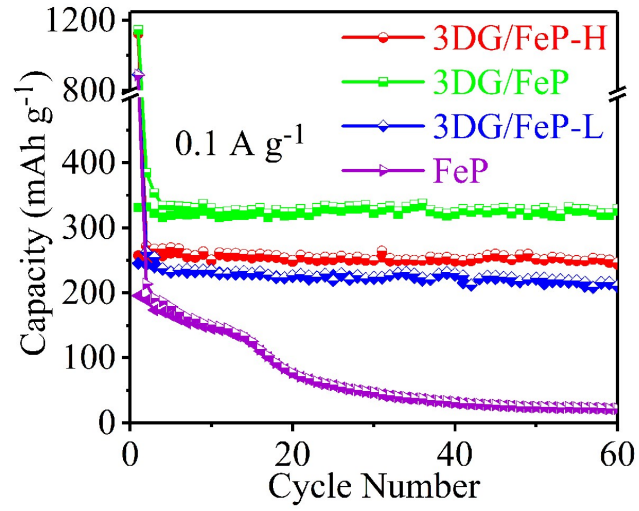


Fig. S14. The cycling performance of 3DG/FeP with different mass content of FeP and pure FeP at 0.1 A g⁻¹.

Table S1. Comparison of recently reported typical FeP-based anode materials for potassium ion batteries.

Materials	Ratio of Active Material	Current Density /mA g ⁻¹	Capacity /mAh g ⁻¹	Capacity (By Electrode) /mAh g ⁻¹	Cycle Life /Capacity Retention @Cycle Number@mA g ⁻¹	Ref
CoP/C	80%	50	126.72	101.4	~75%@200@200 mA g ⁻¹	S2
		1000	54.4	43.52		
FeP@C	80%	100	220	176	92%@300@100 mA g ⁻¹	S3
		2000	40	32		
ZnS@C@rGO	90%	50	320	297	~25%@300@500 mA g ⁻¹	S4
		500	240	187.2		
MoS ₂ /rGO	80%	100	380	304	100@100 mA g ⁻¹	S5
MoSe ₂ /N-C	80%	100	258	206.4	88%@200@100 mA g ⁻¹	S6
Sn/C	80%	25	150	120	75%@30@25 mA g ⁻¹	S7
FeCl ₃ -EG	80%	50	269.5	215.6	88.82%@500@100 mA g ⁻¹ 70%@1300@5000 mA g ⁻¹	S8
VS ₂ /NSA	60%	1000	175.4	105.24	105%@100@1000 mA g ⁻¹	S9
		5000	133.1	79.86		
K ₂ Ti ₈ O ₁₇	70%	100	400	280	85%@50@20 mA g ⁻¹	S10
		2000	100	70		
Hybrid Co ₃ O ₄ -Fe ₂ O ₃ /C	66.6%	20	120	79.92	93%@180@50 mA g ⁻¹	S11
		500	50	33.3		
Ordered Mesoporous Carbon	80%	50	257.4	205.92	99%@20@50 mA g ⁻¹	S12
		1000	146.5	117.2		
Soft Carbon	70%	500	200	140	71.4%@1000@350 mA g ⁻¹	S13
Oak-based Hard Carbon	80%	100	135	108	98%@150@100 mA g ⁻¹	S14
Semi-hollow Micro Soft Carbon	70%	100	312	218.4	82%@500@200 mA g ⁻¹	S15
		500	180	126		

Hard-soft Composite Carbon	80%	100	261	208.8	93%@450@200 mA g ⁻¹	S16
		10000	120	96		
Graphitic Carbon Networks	80%	100	220	176	75%@500@50 mA g ⁻¹	S17
		2000	100	80		
Graphite	90%	20	246	221.4	98%@200@20 mA g ⁻¹	S18
		500	~0	~0		
Reduced Graphene Oxide	80%	5	280/205	224	100%@175@10 mA g ⁻¹	S19
		100	50	40		
FFGF (F-Doped Graphene Foam)	70%	50	326.1	228.27	90.91%@50@60 mA g ⁻¹	S20
		500	165.9	116.13	88.24%@500@500 mA g ⁻¹	
N-doped Graphene	100%	50	350	350	77.78%@100@100 mA g ⁻¹	S21
This Work	100%	100	323	323	97.3%@300@100 mA g ⁻¹	
		2000	127	127	97.6%@2000@2000 mA g ⁻¹	

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