## Supporting Information

## An amorphous Zn-P/graphite composite with chemical bonding for ultrareversible lithium storage

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Figure S1. a), b) FESEM images of the ZnP2 and Zn3P2 powders, respectively.



**Figure S2.** The Raman fingerprint peaks of the alfa- $ZnP_2$  differing from the milled P counterparts.



Figure S3. The initial three discharge/charge profiles of ZnP<sub>2</sub>.



Figure S4. The XRD pattern of  $Zn_3P_2/C$  (10 h) milled which degraded into a mixture containing crystalline Zn.



Figure S5. The crystalline Zn within the 3Zn-2P/C composite.



Figure S6. The XPS spectra of survey spectrum for  $ZnP_2$  and  $ZnP_2/C$ .



**Figure S7.** The XPS spectra of P 2p for  $ZnP_2$  and  $ZnP_2/C$ .



Figure S8. The XPS spectra of Zn 2p for  $ZnP_2$  and  $ZnP_2/C$ .



Figure S9. The Raman Spectra: a) ZnP2 and its carbon composite; b) Zn3P2 and its carbon composite



Figure S10. The electrochemical impedance spectra of the pristine  $ZnP_2$  electrode (pink), the pristine  $ZnP_2/C$  electrode (red), the  $ZnP_2/C$  electrode (blue) after 200 cycles.



Figure S11. The electrode morphology after 50 cycles: a) the ZnP<sub>2</sub>/C electrode; b) the ZnP<sub>2</sub> electrode.



**Figure S12.** The cycle stability of the Li-ion full cell  $\text{LiCoO}_2//\text{ZnP}_2/\text{C}$  at a current density of 200 mA h g<sup>-1</sup>.