

## Crystallization and electrochemical properties of $K_xV_2O_5$ nano-ribbons obtained via solvothermal process as a promising cathode for PIBs

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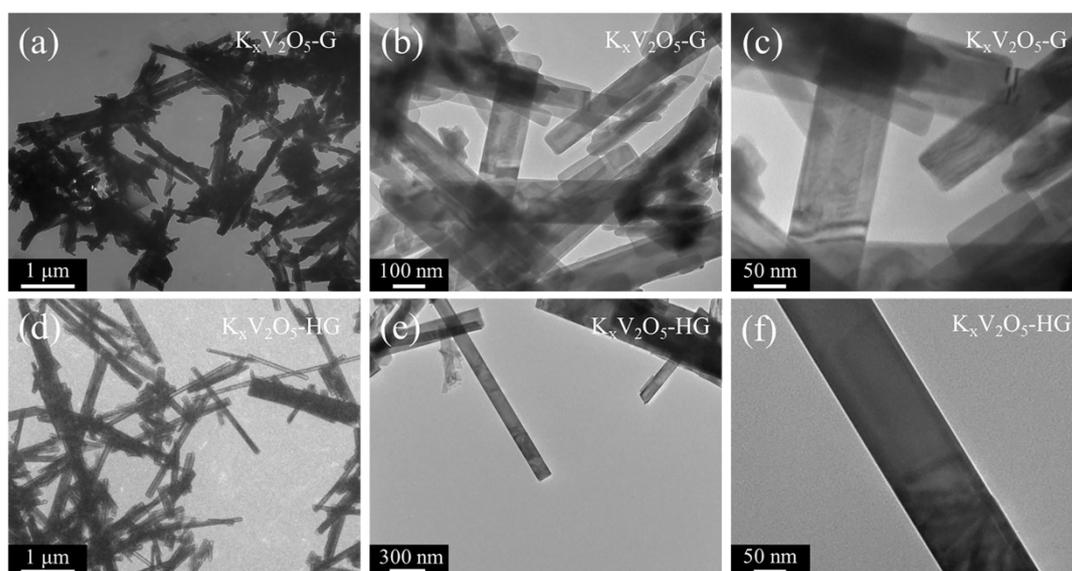


Fig. S1 TEM images of  $K_xV_2O_5$ -G (a, b, c) and  $K_xV_2O_5$ -HG (d, e, f) at different magnifications.

As depicted in Fig. S1,  $K_xV_2O_5$ -HG (d, e, f) (that is,  $K_xV_2O_5$ -HG1 mentioned in experimental section) prepared by sol-gel process owns more uniform nano-ribboned morphology. It is conducive to the potassium storage performance.  $K_xV_2O_5$ -HG has a relatively longer length of  $\sim 2$ -5  $\mu\text{m}$  and smoother surface compared to  $K_xV_2O_5$ -G (a, b, c). However, the length of  $K_xV_2O_5$ -G is shorter and its agglomeration is also more

obvious, which is synthesized by introducing glucose only in hydrothermal condition.

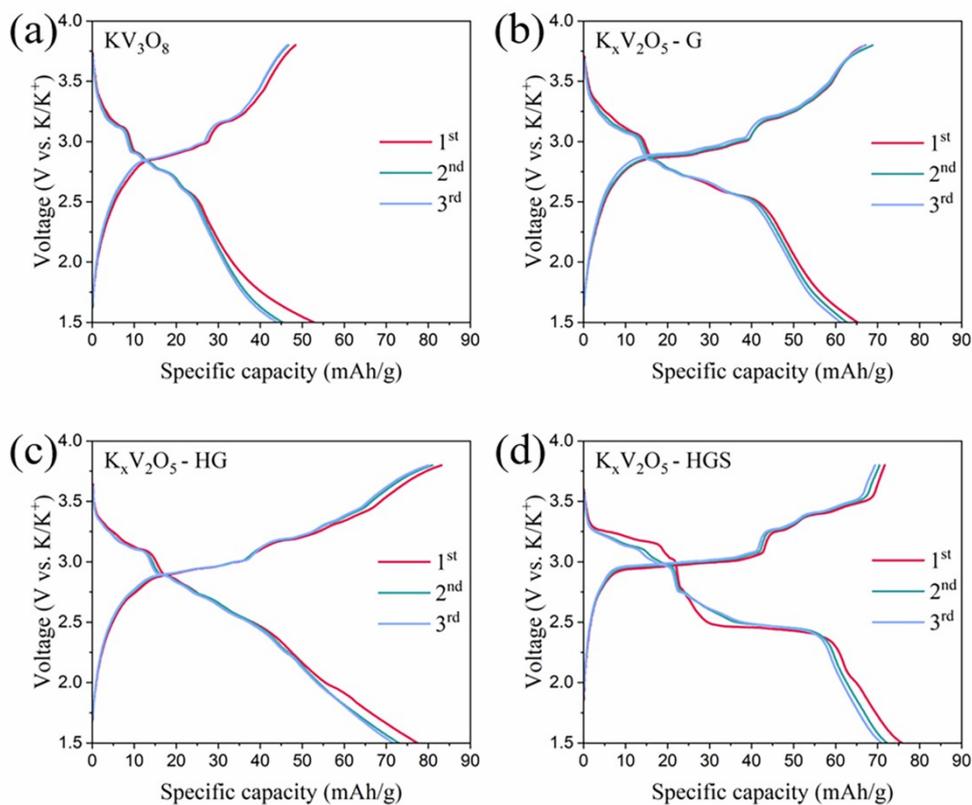


Fig. S2 Initial three galvanostatic charge/discharge (GCD) profiles for KV<sub>3</sub>O<sub>8</sub> (a), K<sub>x</sub>V<sub>2</sub>O<sub>5</sub>-G (b), K<sub>x</sub>V<sub>2</sub>O<sub>5</sub>-HG (c) and K<sub>x</sub>V<sub>2</sub>O<sub>5</sub>-HGS (d) at 50 mA/g between 1.5-3.8 V.

It can be seen from Fig. S2 that the voltage plateaus and curve profiles of KV<sub>3</sub>O<sub>8</sub> (a), K<sub>x</sub>V<sub>2</sub>O<sub>5</sub>-G (b), K<sub>x</sub>V<sub>2</sub>O<sub>5</sub>-HG (c) and K<sub>x</sub>V<sub>2</sub>O<sub>5</sub>-HGS (d) only display little changes for their initial three cycles. So, the significant changes in plateaus and curve profiles are primarily caused by subsequent long-term cycling.

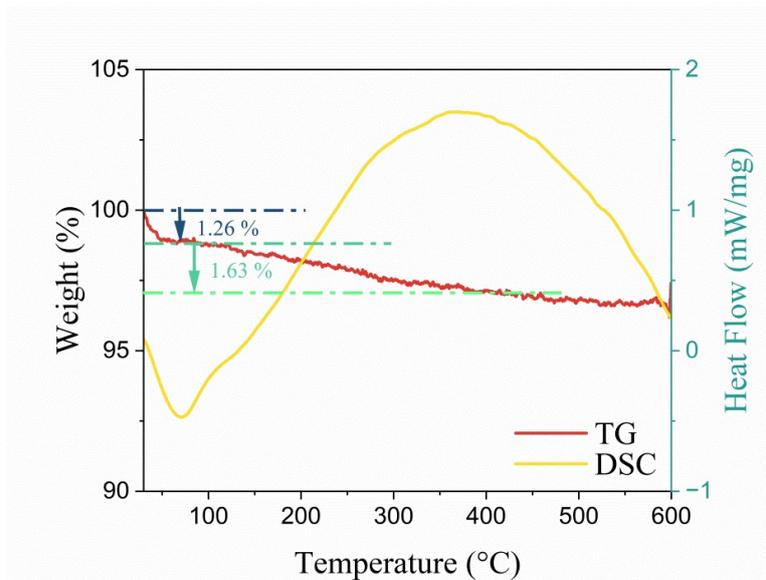


Fig. S3 TG-DSC of  $K_xV_2O_5$ -HG in Ar atmosphere.

TG-DSC was conducted at the temperature range of  $\sim 30$  to  $600$   $^{\circ}C$  with a heating rate of  $10$   $^{\circ}C/min$  under inert Ar atmosphere in order to determine the rough content of crystalline water in  $K_xV_2O_5$ -HG. It can be observed that a weight loss of  $1.26\%$  occurs between  $\sim 30$  and  $100$   $^{\circ}C$  due to the evaporation of adsorbed water, accompanying by a downward endothermic peak in the region of  $50$ - $100$   $^{\circ}C$ . Besides, another weight loss of  $1.63\%$  happens between  $100$  and  $400$   $^{\circ}C$  owing to the release of deep crystalline water along with further crystallization of  $K_xV_2O_5$ -HG. Meanwhile, an upward exothermic peak appears between  $200$  and  $600$   $^{\circ}C$ , which is caused by the crystallization of  $K_xV_2O_5$ -HG. So, TG-DSC test manifests that  $K_xV_2O_5$ -HG in our study contains a certain amount of bound water ( $\sim 2.89$  wt.%). Based on the analyses of XRD and XPS in this work, we believe that the value of  $x$  in  $K_xV_2O_5$ -HG is  $\sim 0.5$ . Hence, it can be estimated that  $K_xV_2O_5$ -HG contains  $\sim 0.33$  of bound  $H_2O$ , which can be ultimately written as  $K_{0.5}V_2O_5 \cdot 0.33H_2O$ -HG. As the writing of this chemical formula involves several approximations, it is more universal by utilizing  $K_xV_2O_5$ -HG in the main manuscript.