Electronic Supplementary Information (ESI) for

Enhanced electrochemical performance of lithium ion batteries using

Sb₂S₃ nanorods and graphene composites as anode materials

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Synthesis of graphene oxide

First, the graphite powder was oxidized using 0.9 g NaNO₃ and 37 mL of concentrated H₂SO₄ that were added to 1 g graphite powder cooled in an ice bath. This mixture was continuously stirred while 5 g of KMnO₄ was added slowly over 1 h. It was left to stir for 2 h in the ice bath, and then removed and left for 4 days under continuous stirring. A black viscous liquid was obtained and 100 mL of deionized water was added over 30 min while stirring continuously. The mixture was stirred for a further 2 h, 10 mL H₂O₂ (30wt% aqueous solution) was slowly added and then the mixture was left to stir for another 2 h. The resulting oxidized material was washed three times by 10wt% diluted hydrochloric acid and then washed by deionized water till the pondus hydrogenii (pH) value was close to 7. A light yellow GO powder was obtained after freeze drying for 12 h.



Fig. 1S XRD pattern of Sb₂S₃ crystals.



Fig. 2S TGA curve of Sb₂S₃/G composites.



Fig. 3S SEM images of Sb₂S₃/G composites with half amount of GO with different magnification.



Fig. 4S SEM image of Sb₂S₃ crystals.



Fig. 5S CV of Sb₂S₃ crystals at a scan rate of 0.1 mV s⁻¹ in the voltage range of 0.01-3 V (vs. Li^+/Li).



Fig. 6S Cyclic stability of Sb₂S₃/HG composites at various current rates.



Fig. 7S (a) Rate capacity of Sb₂S₃ crystals at various current rates; and (b) cyclic stability of Sb₂S crystals.



Fig. 8S Cyclic stability of Sb_2S_3/G composites tested at various current rates.



Fig. 9S SEM image of Sb_2S_3/G composites with different magnification after 100 cycles at a current rate of 0.2C