

Supplementary material:

Bismuth–Antimony Alloy Embedded in Carbon Matrix for Ultra-Stable Sodium Storage

Wensheng Ma, Bin Yu, Fuquan Tan, Hui Gao * and Zhonghua Zhang *

Key Laboratory for Liquid-Solid Structural Evolution and Processing of Materials (Ministry of Education), School of Materials Science and Engineering, Shandong University, Jingshi Road 17923, Jinan 250061, China

* Correspondence: hgao199306@gmail.com (H.G.); zh_zhang@sdu.edu.cn (Z.Z.)

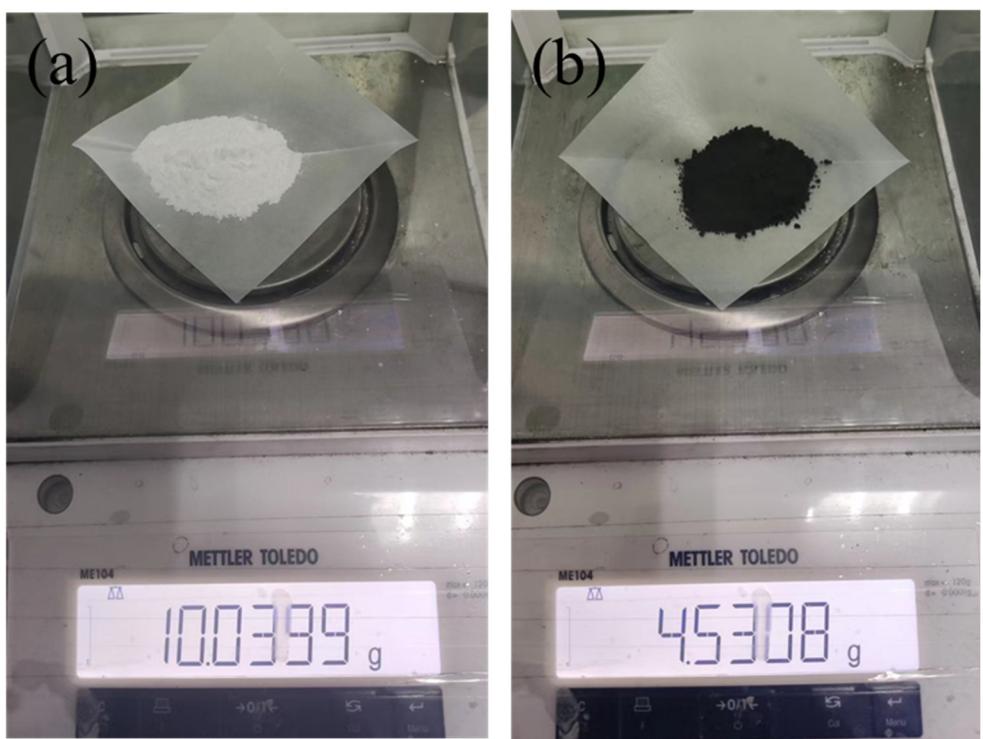


Figure S1. The photographs of (a) a mixture of bismuth citrate and antimony potassium tartrate in a molar ratio of 1:1 and (b) the obtained $\text{Bi}_1\text{Sb}_1@\text{C}$ sample after cleaning and drying.

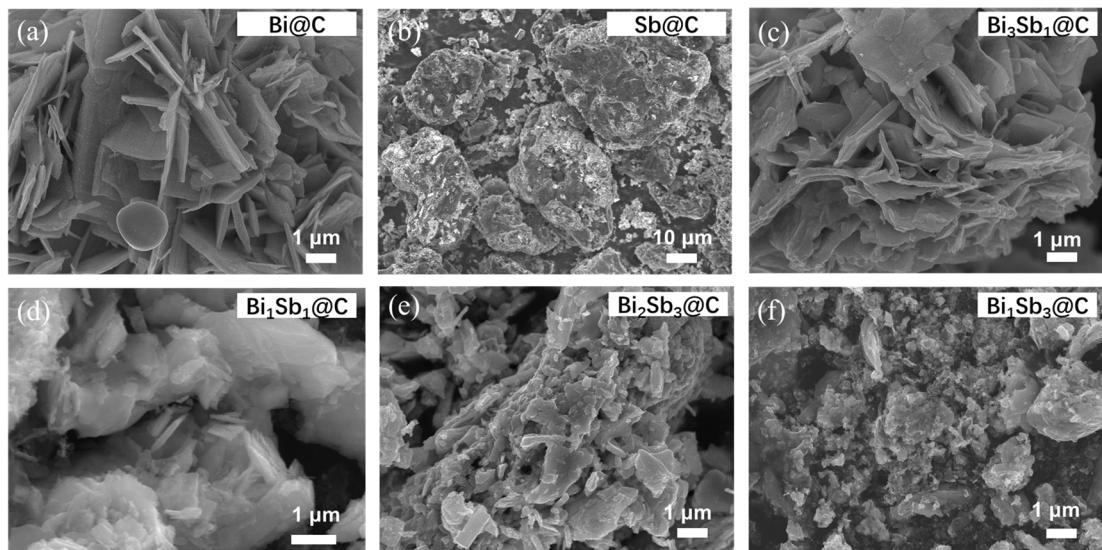


Figure S2. The SEM images at low magnifications of the (a) Bi@C, (b) Sb@C, (c) Bi₃Sb₁@C, (d) Bi₁Sb₁@C, (e) Bi₂Sb₃@C and (f) Bi₁Sb₃@C samples.

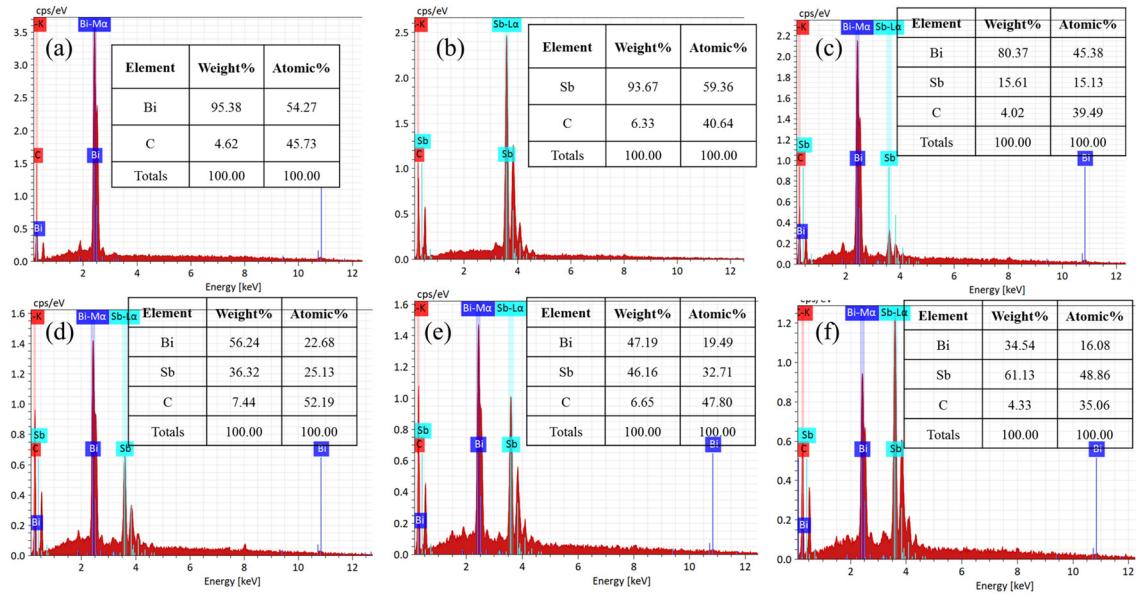


Figure S3. The corresponding EDX results of the (a)Bi@C, (b) Sb@C, (c) Bi₃Sb₁@C, (d) Bi₁Sb₁@C, (e) Bi₂Sb₃@C, and (f) Bi₁Sb₃@C samples.

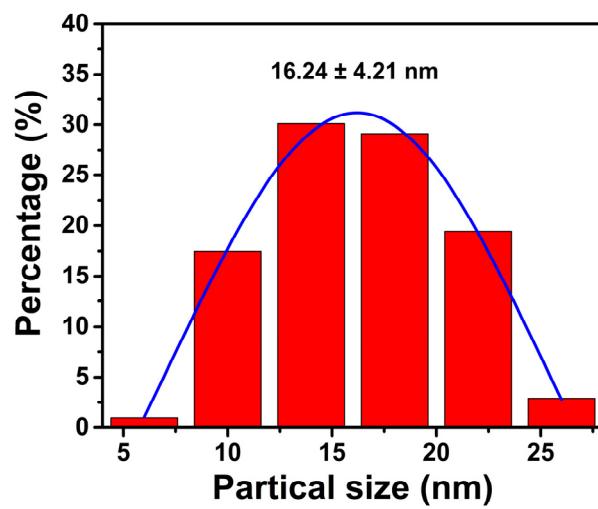


Figure S4. The size distribution pattern of the Bi_1Sb_1 alloy for the $\text{Bi}_1\text{Sb}_1@\text{C}$ sample.

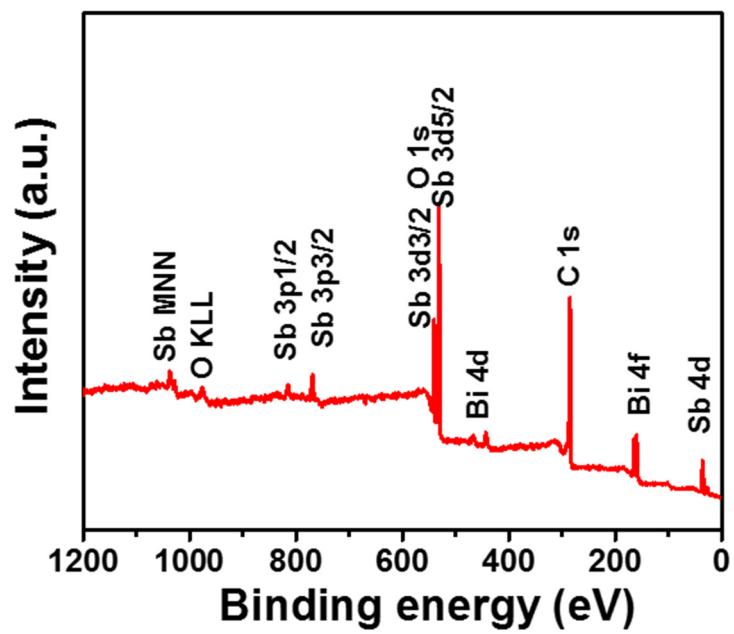


Figure S5. The low-resolution XPS spectrum of the $\text{Bi}_1\text{Sb}_1@\text{C}$ sample.

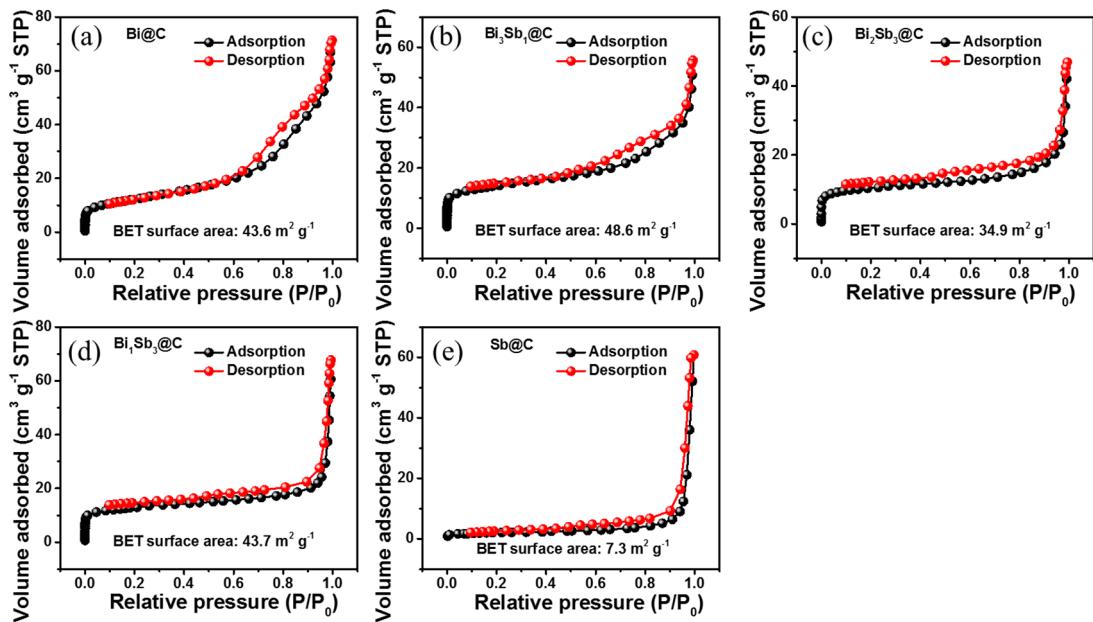


Figure S6. N_2 adsorption-desorption isotherms of the (a) $\text{Bi}@\text{C}$, (b) $\text{Bi}_3\text{Sb}_1@\text{C}$, (c) $\text{Bi}_2\text{Sb}_3@\text{C}$, (d) $\text{Bi}_1\text{Sb}_3@\text{C}$, and (e) $\text{Sb}@\text{C}$ samples.

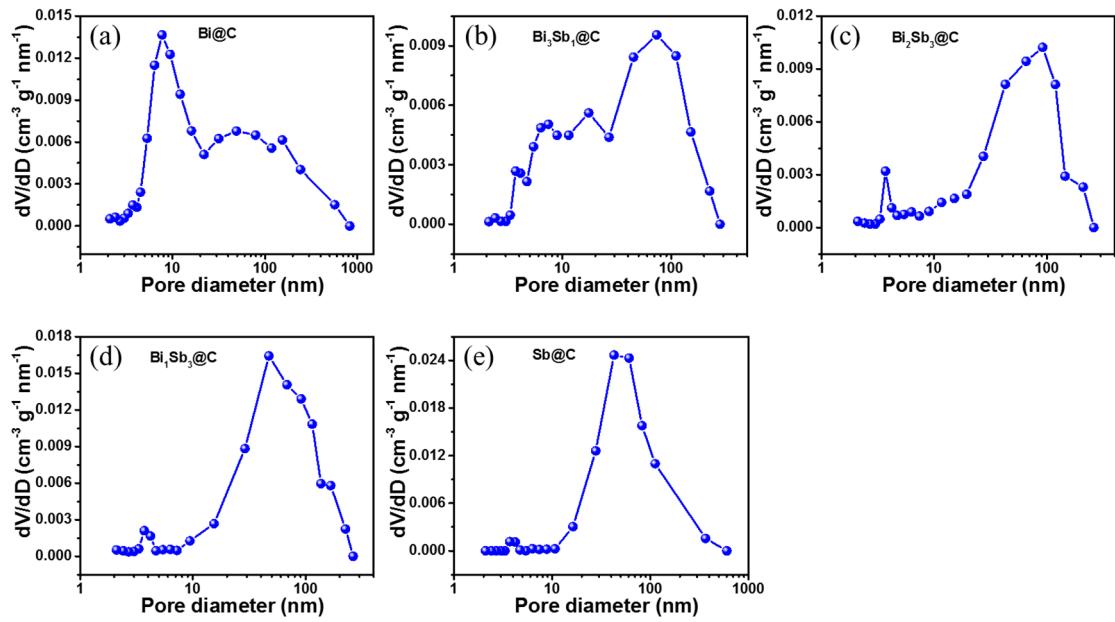


Figure S7. The pore size distribution curves of the (a) $\text{Bi}@\text{C}$, (b) $\text{Bi}_3\text{Sb}_1@\text{C}$, (c) $\text{Bi}_2\text{Sb}_3@\text{C}$, (d) $\text{Bi}_1\text{Sb}_3@\text{C}$, and (e) $\text{Sb}@\text{C}$ samples.

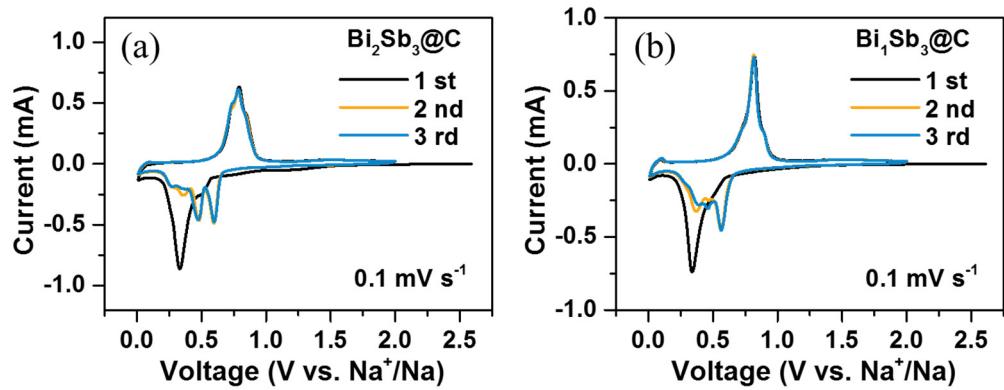


Figure S8. Initial three cycles of CV curves at a scan rate of 0.1 mV s^{-1} of the (a) $\text{Bi}_2\text{Sb}_3@\text{C}$ and (b) $\text{Bi}_1\text{Sb}_3@\text{C}$ samples.

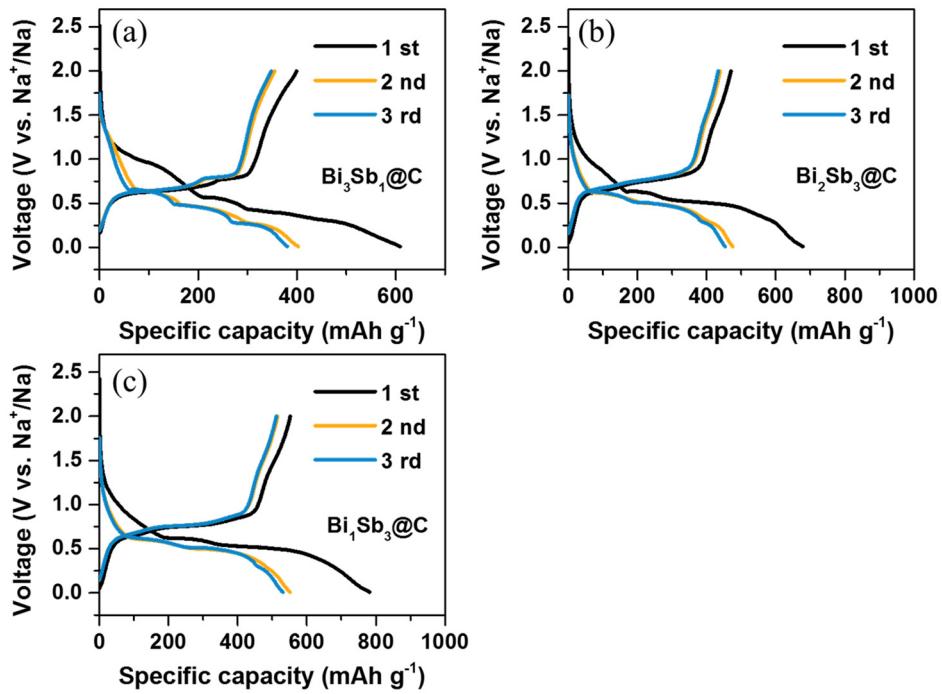


Figure S9. The discharge-charge curves of the (a) $\text{Bi}_3\text{Sb}_1@\text{C}$, (b) $\text{Bi}_2\text{Sb}_3@\text{C}$, and (c) $\text{Bi}_1\text{Sb}_3@\text{C}$ samples during the initial three cycles.

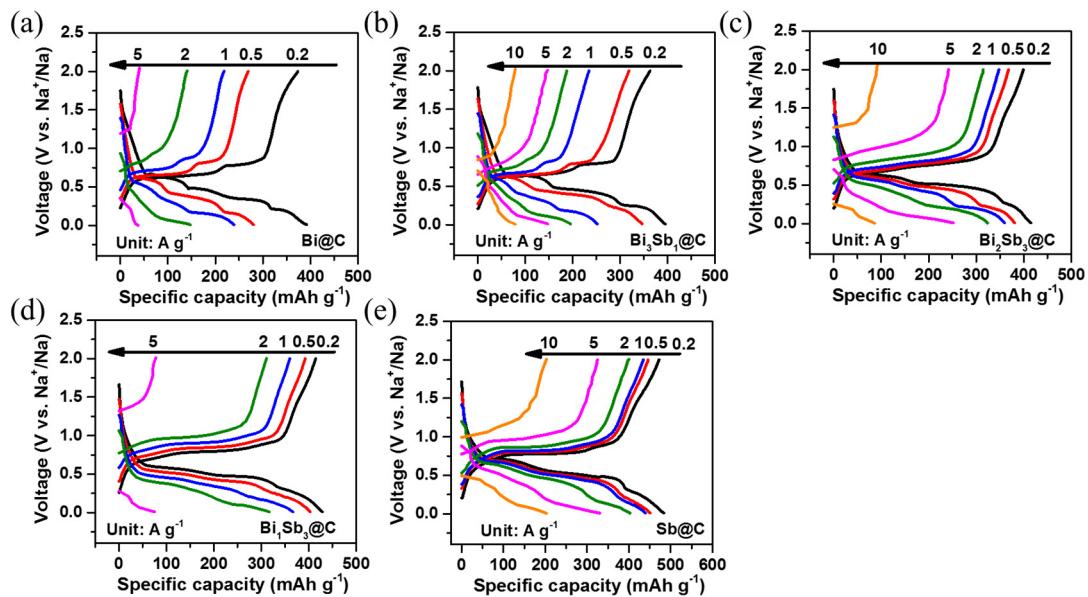


Figure S10. The discharge-charge profiles at different current densities of the (a) Bi@C, (b) Bi₃Sb₁@C, (c) Bi₂Sb₃@C, (d) Bi₁Sb₃@C, and (e) Sb@C samples.

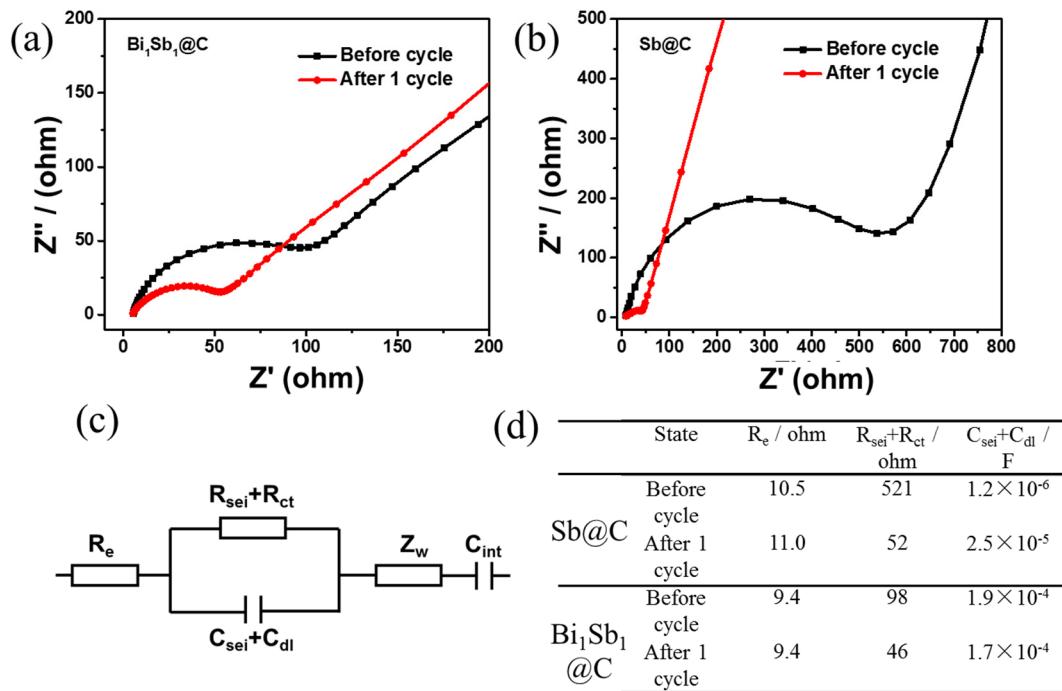


Figure S11. Nyquist plots of (a) $\text{Bi}_1\text{Sb}_1@\text{C}$ and (b) $\text{Sb}@\text{C}$ before/after the first cycle. (c, d) Fitted circuit diagram and corresponding fitting results.

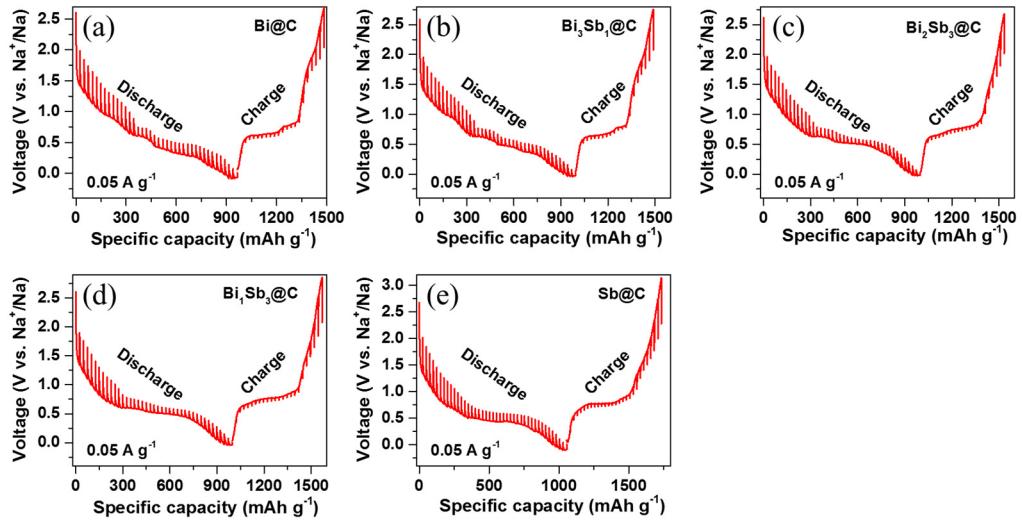


Figure S12. The GITT curves of the (a) $\text{Bi}@\text{C}$, (b) $\text{Bi}_3\text{Sb}_1@\text{C}$, (c) $\text{Bi}_2\text{Sb}_3@\text{C}$, (d) $\text{Bi}_1\text{Sb}_3@\text{C}$, and (e) $\text{Sb}@\text{C}$ samples during the first discharge and charge processes.

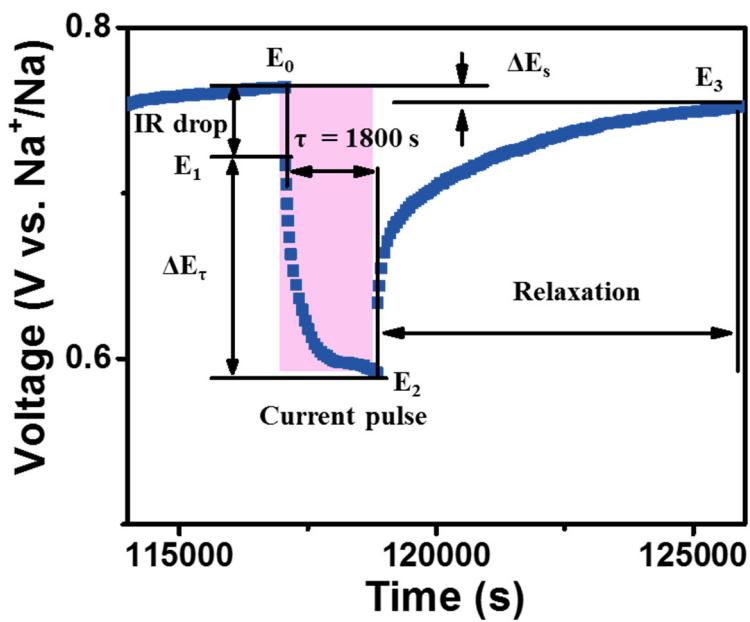


Figure S13. Schematic of one discharge pulse at $\sim 0.75 \text{ V}$ (vs. Na^+/Na) for the $\text{Bi}_1\text{Sb}_1@\text{C}$ electrode during discharge process. ΔE_s and ΔE_τ represent the steady-state voltage change and transient voltage change, respectively. During a pulse discharge, the voltage rapidly drops from E_0 to E_1 due to ohmic and charge transfer resistance ($\text{IR} = E_0 - E_1$). Then the voltage gradually decreases to E_2 due to the sodiation process of electrode. During the relaxation process, the voltage returns to the steady state (E_3). The values of ΔE_s and ΔE_τ are calculated by $\Delta E_s = E_0 - E_3$ and $\Delta E_\tau = E_1 - E_2$, respectively.

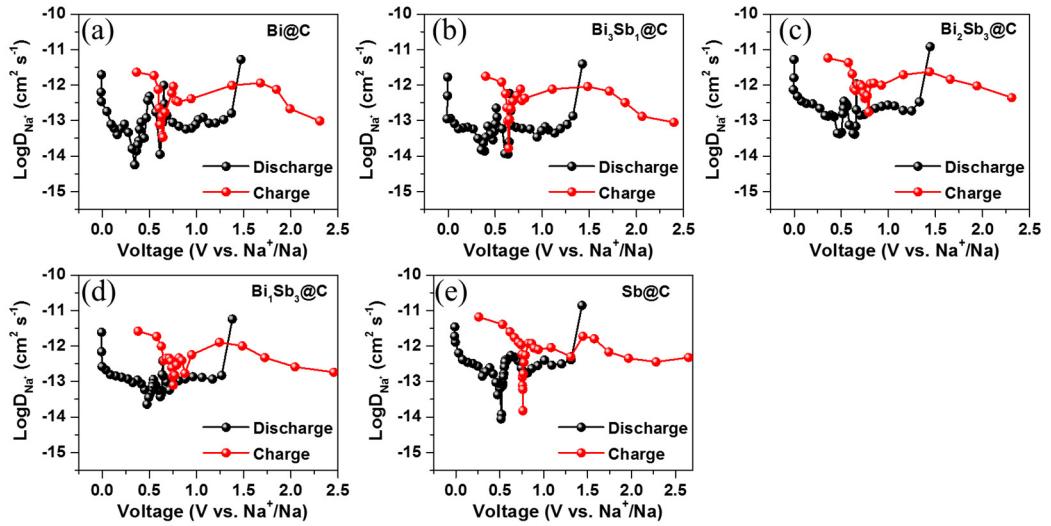


Figure S14. The diffusion rate for Na^+ of the (a) $\text{Bi}@\text{C}$, (b) $\text{Bi}_3\text{Sb}_1@\text{C}$, (c) $\text{Bi}_2\text{Sb}_3@\text{C}$, (d) $\text{Bi}_1\text{Sb}_3@\text{C}$, and (e) $\text{Sb}@\text{C}$ samples during the first discharge and charge processes.