Supporting Information

Comparison of Perovskite and Perovskite Derivatives for use in Anion-based Pseudocapacitor Applications

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Figure S1. Nitrogen sorption isotherms for BET surface area analysis of unreduced CaMnO_{3- δ} (CMO) perovskite and Ca₂MnO_{4- δ} (CMO RP) Ruddlesden-Popper oxides.



Figure S2. Additional SEM images of CaMnO₃₋₆ (CMO) perovskite oxide.



Figure S3. Additional SEM images of $Ca_2MnO_{4-\delta}$ (CMO RP) Ruddlesden-Popper oxide.



Figure S4. Mn 3p XPS analysis of CaMnO_{3- δ} (CMO) perovskite and Ca₂MnO_{4- δ} (CMO RP) Ruddlesden-Popper oxides before and after reduction in a reducing atmosphere of 7% H₂ in Ar at 325° C.



Figure S5. Electrochemical characterization of unreduced CaMnO_{3- δ} and Ca₂MnO_{4- δ} materials as pseudocapacitors in a 3-electrode cell. (a) CVs of CMO performed at multiple scan rates over a 1 V window. (b) CVs of CMO RP performed at multiple scan rates over a 1 V window.



Figure S6. PXRD pattern of SrFeO_{2.5} (SFO). Reduction of SrFeO₃ perovskite was performed in an atmosphere of 7% H_2 /Ar at 325°C to achieve the Brownmillerite phase.



Figure S7. Gravimetric capacitances determined from CVs at various scan rates using SrFeO_{2.5} as the anode material and either r-CMO or r-CMO RP as the cathode material in two-electrode cell asymmetric pseudocapacitor experiments.

Table S1. Electrochemically measured oxygen diffusion rates in $CaMnO_{3-\delta}$ (CMO) perovskite and $Ca_2MnO_{4-\delta}$ (CMO RP) Ruddlesden-Popper oxides.

Sample	Composition	Intercalation Oxygen Diffusion Rate (cm ² s ⁻¹)	Deintercalation Oxygen Diffusion Rate (cm ² s ⁻¹)
CMO	CaMnO _{3.11}	1.88E-11 <u>+</u> 5.7E-12	2.44E-11 <u>+</u> 3.6E-12
r-CMO	CaMnO _{2.53}	2.64E-11 <u>+</u> 2.8E-12	3.71E-11 <u>+</u> 1.2E-12
CMO RP	Ca ₂ MnO _{3.92}	3.54E-12 <u>+</u> 1.4E-12	1.68E-12 <u>+</u> 3.6E-13
r-CMO RP	Ca ₂ MnO _{3.61}	1.60E-11 <u>+</u> 8.6E-14	1.44E-11 <u>+</u> 4.9E-14