

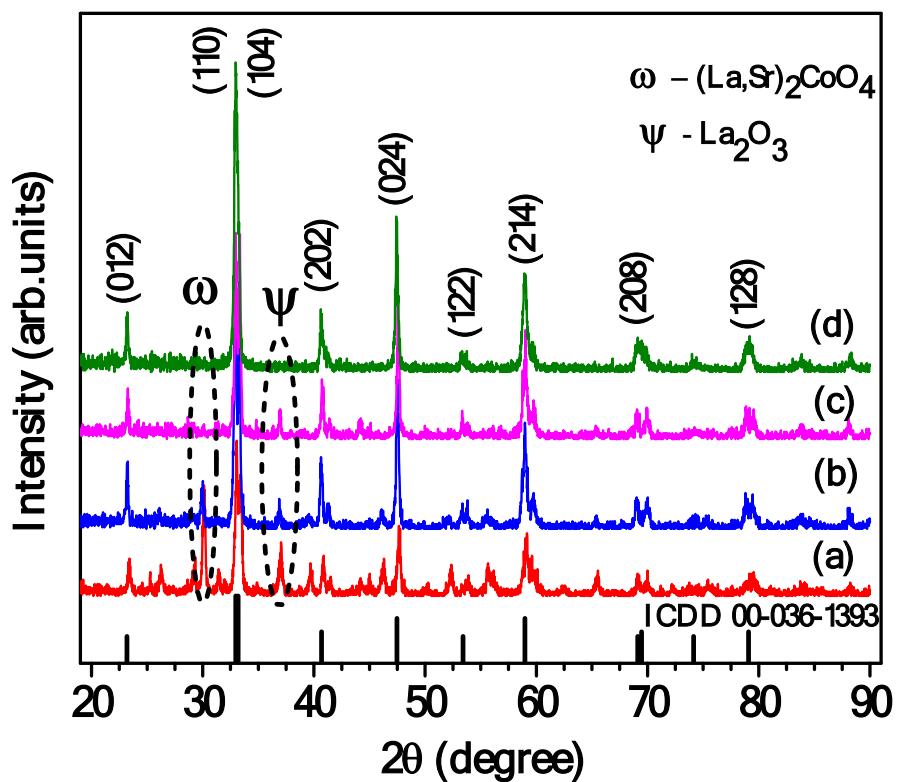
### Supporting information

**Table ST1:** Comparison the  $\text{La}_{1-x}\text{Sr}_x\text{CoO}_3$  perovskite oxide catalyst based aprotic  $\text{Li-O}_2$  battery.

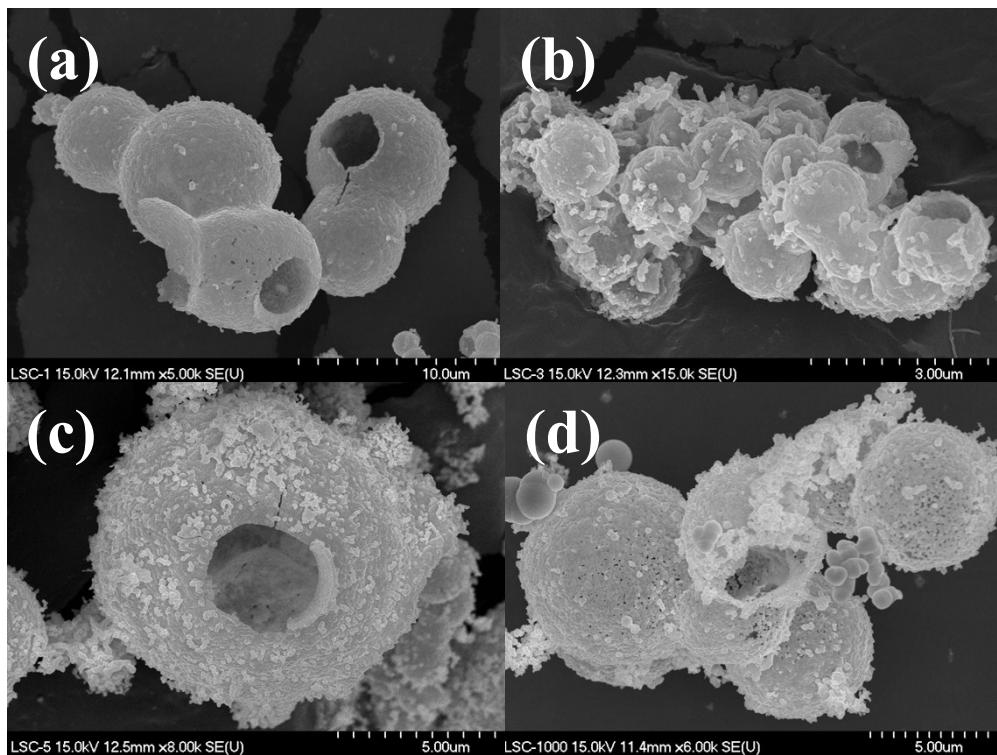
Catalyst	Current rate	Discharge capacity (mAh/g)	Overpotential (V)	Reference
HS $\text{La}_{0.6}\text{Sr}_{0.4}\text{CoO}_{3-\delta}$	<b>100 mA g<sup>-1</sup></b>	<b>4895</b>	<b>1.016</b>	<b>Present work</b>
$\text{La}_{0.6}\text{Sr}_{0.4}\text{CoO}_{3-\delta}$	100 mA g <sup>-1</sup>	4701	1.1	R1
$\text{La}_{0.6}\text{Sr}_{0.4}\text{CoO}_{3-\delta}$	100 mA g <sup>-1</sup>	3256	0.85	R2
$\text{La}_{0.6}\text{Sr}_{0.4}\text{CoO}_3$	30 mA g <sup>-1</sup>	3672	~1.4	R3
$\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.9}\text{Mn}_{0.1}\text{O}_3$	200 mA g <sup>-1</sup>	3107	~1.4	R4
$\text{La}_{0.5}\text{Sr}_{0.5}\text{CoO}_{3-x}$ Nanotubes	25 mA g <sup>-1</sup>	5799	1.14	R5
$\text{La}_{0.8}\text{Sr}_{0.2}\text{Mn}_{0.6}\text{Ni}_{0.4}\text{O}_3$	50 mA g <sup>-1</sup>	5364	1.33	R6
$\text{La}_{0.4}\text{Sr}_{0.6}\text{MnO}_3$	50 mA g <sup>-1</sup>	5624	1.45	R7

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- R2. J.J. Lee, M.Y. Oh, K.S. Nahm, Effect of Ball Milling on Electrocatalytic Activity of Perovskite  $\text{La}_{0.6}\text{Sr}_{0.4}\text{CoO}_{3-\delta}$  Applied for Lithium Air Battery, Journal of The Electrochemical Society, 163 (2016) A244-A250.
- R3. N. Sun, H. Liu, Z. Yu, Z. Zheng, C. Shao, The  $\text{La}_{0.6}\text{Sr}_{0.4}\text{CoO}_3$  perovskite catalyst for  $\text{Li-O}_2$  battery, Solid State Ionics, 268 (2014) 125-130.

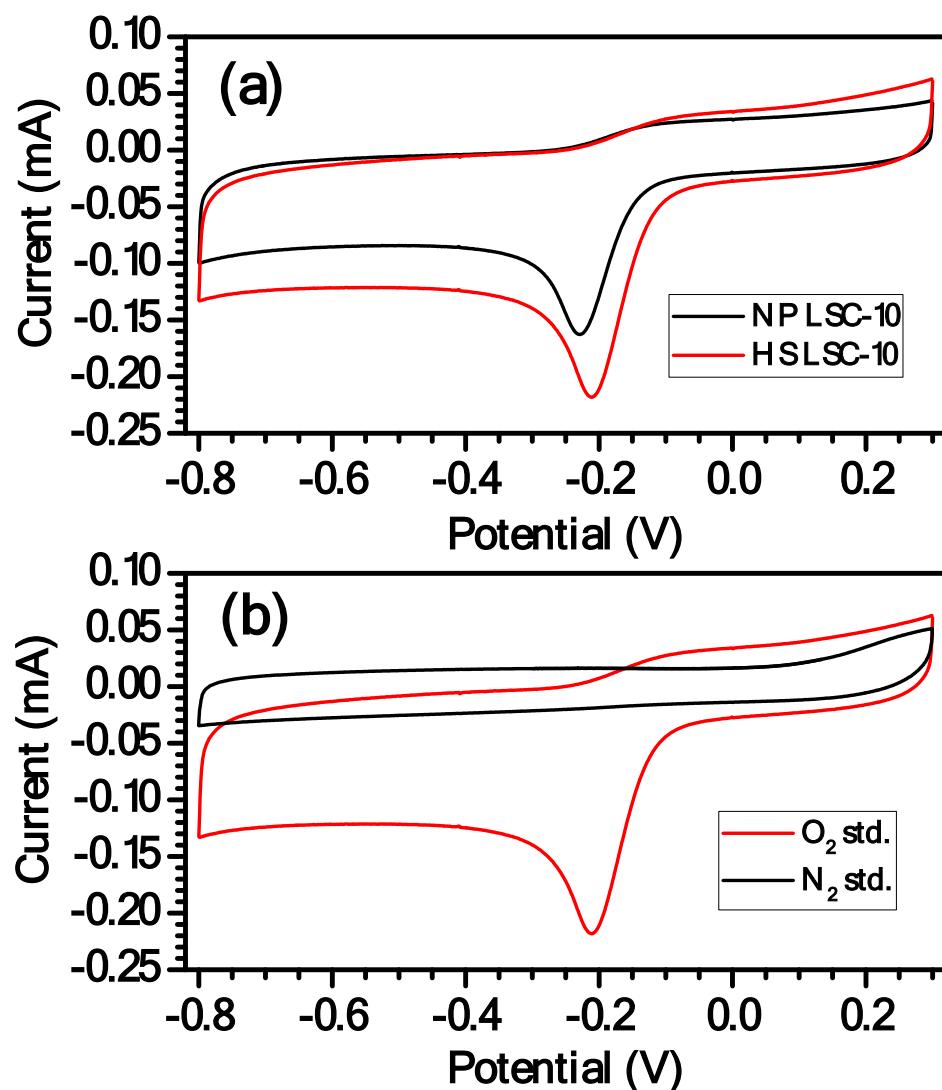
- R4. N. Sun, H. Liu, Z. Yu, Z. Zheng, C. Shao, Mn-doped  $\text{La}_{0.6}\text{Sr}_{0.4}\text{CoO}_3$  perovskite catalysts with enhanced performances for non-aqueous electrolyte  $\text{Li-O}_2$  batteries, *RSC Advances*, 6 (2016) 13522-13530.
- R5. G. Liu, H. Chen, L. Xia, S. Wang, L.-X. Ding, D. Li, K. Xiao, S. Dai, H. Wang, Hierarchical Mesoporous/Macroporous Perovskite  $\text{La}_{0.5}\text{Sr}_{0.5}\text{CoO}_{3-x}$  Nanotubes: A Bifunctional Catalyst with Enhanced Activity and Cycle Stability for Rechargeable Lithium Oxygen Batteries, *ACS Applied Materials & Interfaces*, 7 (2015) 22478-22486.
- R6. Z. Wang, Y. You, J. Yuan, Y.-X. Yin, Y.-T. Li, S. Xin, D. Zhang, Nickel-Doped  $\text{La}_{0.8}\text{Sr}_{0.2}\text{Mn}_{1-x}\text{Ni}_x\text{O}_3$  Nanoparticles Containing Abundant Oxygen Vacancies as an Optimized Bifunctional Catalyst for Oxygen Cathode in Rechargeable Lithium–Air Batteries, *ACS Applied Materials & Interfaces*, 8 (2016) 6520-6528.
- R7. Y. Zhao, Y. Hang, Y. Zhang, Z. Wang, Y. Yao, X. He, C. Zhang, D. Zhang, Strontium-doped perovskite oxide  $\text{La}_{1-x}\text{Sr}_x\text{MnO}_3$  ( $x=0, 0.2, 0.6$ ) as a highly efficient electrocatalyst for nonaqueous  $\text{Li-O}_2$  batteries, *Electrochimica Acta*, 232 (2017) 296-302.



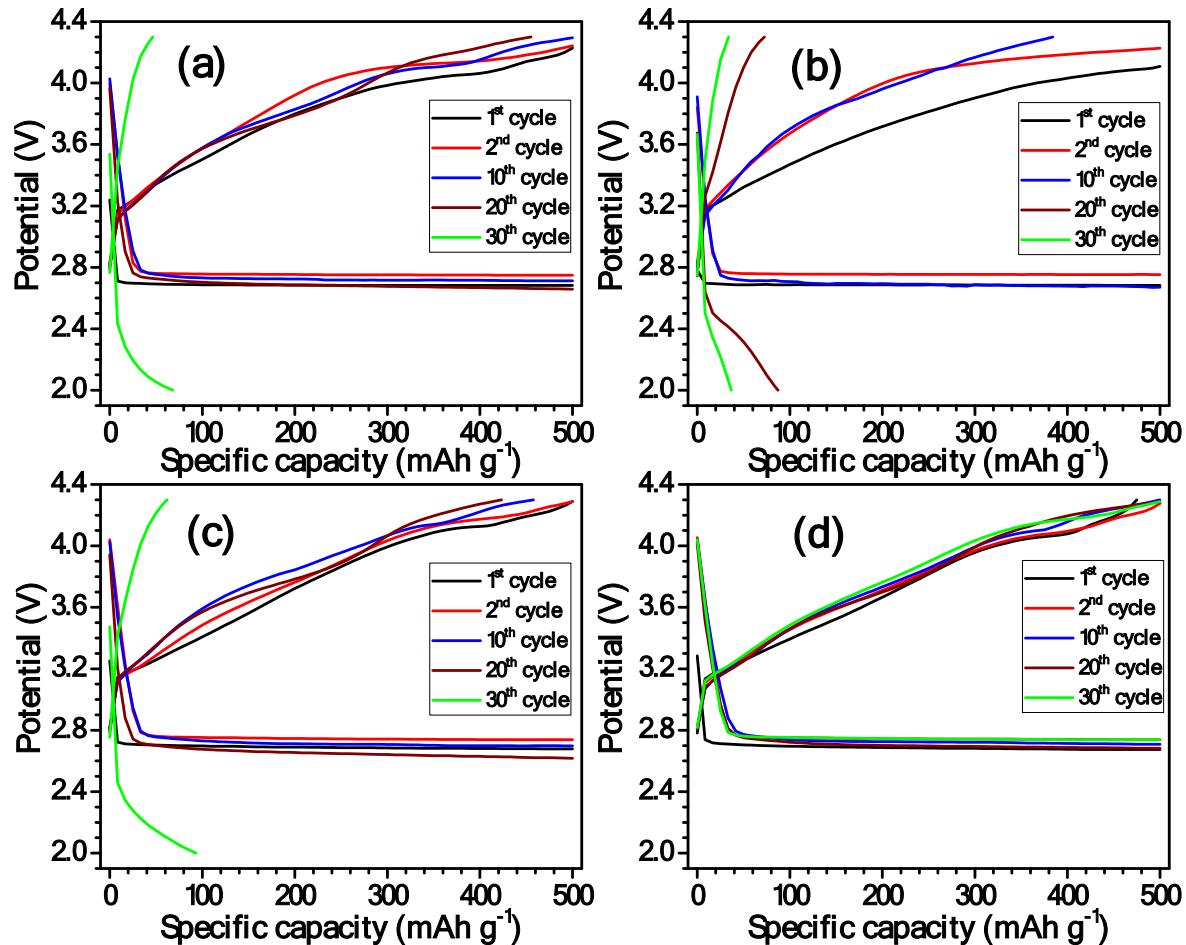
**Figure S1.** The XRD pattern of the synthesised  $\text{La}_{0.6}\text{Sr}_{0.4}\text{CoO}_3$  (HS LSC) at different temperatures (a) 700 °C (HS LSC-7), (b) 800 °C (HS LSC-8), (c) 900 °C ( HS LSC-9), and (d) 1000 °C ( HS LSC-10).



**Figure S2.** Scanning electron microscope images of pristine HS LSC-7 (a), HS LSC-8 (b), HS LSC-9 (c) and HS LSC-10 (d).



**Figure S3.** (a) Comparison cyclic voltammetry of NP-LSC-10 and HS-LSC-10 in O<sub>2</sub> saturated 0.1 M KOH electrolyte and (b) comparison CV of HS-LSC-10 in O<sub>2</sub> and N<sub>2</sub> saturated 0.1 M KOH solution at 5 mV s<sup>-1</sup> scan rate.



**Figure S4.** The limited capacity range charge/discharge curves of (a) HS LSC-7, (b) HS LSC-8, (c) HS LSC-9 and (d) HS LSC-10 catalyst at  $100 \text{ m g}^{-1}$  between 2 to 4.3 V.

### Calibration of Hg/HgO reference electrode:

The calibration of Hg/HgO reference electrode was performed in a standard three-electrode system with polished Pt wire as the working and counter electrodes, and the Hg/HgO electrode used as reference electrode. The standard electrode potential of Hg/HgO/1M KOH was 0.118 V vs. SHE based on the manufacture's specification (ALS co., Ltd). 0.1 M KOH electrolyte is pre-purged and saturated with high purity H<sub>2</sub> gas for 20 minutes. Linear scanning voltammetry is then run at a scan rate of 0.5 mV s<sup>-1</sup>, and the potential at which the current crossed zero is taken to be the thermodynamic potential for the hydrogen electrode reactions. Here, the zero-current point is at -0.886 V, so ( $E(\text{RHE}) = E(\text{Hg/HgO}) + 0.886 \text{ V}$ ).

