Supplementary data

Enhanced Catalytic Centres by RuO₂ addition to CuFe₂O₄ Cathode Catalyst towards Rechargeable Lithium-Air Battery: Influence of CO₂ on the Li-O₂ battery performances

Pamangadan C Sharafudeen and Perumal Elumalai*

Electrochemical Energy Storage Lab, Department of Green Energy Technology, Madanjeet School of Green Energy Technologies, Pondicherry University, Puducherry – 605014, India

Materials used

Copper nitrate trihydrate (Cu(NO₃)₂·3H₂O, Sigma-Aldrich), ferric nitrate nonahydrate (Fe(NO₃)₃·9H₂O, Loba chemicals), Ruthenium oxide (RuO₂, Sigma-Aldrich) and glycine (C₂H₅NO₂, Avra chemicals) were used for the synthesis of the RuO₂@CuFe₂O₄. The non-aqueous electrolyte was comprised of lithium triflate (LiCF₃SO₃, Sigma-Aldrich) in tetraethylene glycol dimethyl ether (TEGDME, Sigma-Aldrich). The separator used in the batteries were made of Whatman glass microfiber filter paper (Alfa Aesar).



Figure S1. Raman spectrum recorded for the $CuFe_2O_4$ powder sample.



Figure S2. XPS (a) survey spectrum, and (b-e) Deconvoluted XPS profiles of Cu2p, Fe2p, O1s and C 1s scan profiles recorded for the CuFe₂O₄ (CFO) sample.



Figure S3. SEM images at different magnifications recorded for the pristine CuFe₂O₄



sample.

Figure S4. GCD curves recorded at a current density of 500 mA g^{-1} for the Li-Air battery containing each of the pristine CuFe₂O₄ and the RCFO-5 composite catalysts.

Figure S5. OCV profile recorded in the ambient atmosphere for the Li-Air battery having the RCFO-5 as the cathode catalyst.





Figure S6. The equivalent circuit used to fit the obtained Nyquist plots in the Li-CO₂ battery.

State	Voltage (V)	R _{ct} (Ω)
First discharge	3.1	271
	2.5	310
	2.0	366
Fifth discharge	3.1	435
	2.5	445
	2.0	600

Table S1. The R_{ct} resistance calculated at different depth of discharge in first and 5th discharge cycles.