Electronic Supplementary Information

Rheological phase reaction synthesis and electrochemical performance of rufigallol anode for lithium ion batteries

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Materials	Electrode composition (active material /conductive additive/binder)	Stable charge capacity (mAh g ⁻¹) /Current density (mA g ⁻¹)	Initial capacity (mAh g ⁻¹) /Current density (mA g ⁻¹)	*Reference /year
Lithium Perylenetetracarboxylates $(Li_4C_{24}H_8O_8)$	70:20:10	195 (25) 170 (50) 110 (400)	470 and 200 (25)	[1]/2013
Silver terephthalate $(Ag_2C_8H_4O_4)$	60:30:10	171 (71) 161 (140) 122 (564)	650 and 210 (14)	[2]/2016
Zinc terephthalates $(ZnC_8H_4O_4)$	60:30:10	330 (118)	626 and 237 (118)	[3]/2017
Aluminum fumarates (AlCl ₃ -FumA)	50:40:10	392 (37.5)	899 and 401 (37.5)	[4]/2017
Metal formates (Zn ₃ (HCOO) ₆)	70:15:15	550 (60)	1344 and 693 (60)	[5]/2010
Nickel naphthalenetetracarboxylates $(Ni_2C_{14}H_4O_8)$	60:30:10	500 (100)	1823 and 982 (100)	[6]/2012
Lithium perylenetetracarboxylates $(Li_4C_{24}H_8O_8)$	55:35:10	120 (24) 90 (240)	210 and 130 (24)	[7]/2015
Rufigallol-Li/Ni (Li ₄ NiC ₁₄ H ₂ O ₈)	60:35:5	530 (100) 420 (200) 300 (500)	1214 and 560 (100)	This work

Table S1. The most-advanced works summarized for carbonyl metal-organic materials as anodes for lithium ion batteries.



Figure S1. SEM images of fresh electrode (a) and the retrieved electrode (b) after 200 cycles at 500 mA g^{-1} of rufigallol; SEM images of fresh electrode (c) and the retrieved electrode (d) after 200 cycles at 500 mA g^{-1} of R-LN.



Figure S2. XRD patterns of fresh electrode and the retrieved electrode after 200 cycles at 500 mA g^{-1} of rufigallol (a) and R-LN (b). There are three typical peaks at 43.8°, 50.9° and 74.8°, corresponding to the stainless steel mesh current collector (shown in c).

*References:

- 1 R.R. Zhao, Y.L. Cao, X.P. Ai and H.X. Yang, J. Electroanal. Chem., 2013, 688, 93.
- 2 J. Xue, C. Fan, Q.J. Deng, M.J. Zhao, L.P. Wang, A.J. Zhou and J.Z. Li, *Electrochim. Acta*, 2016, 219, 418.
- 3 L.P. Wang, J. Zou, S.L. Chen, J.Y. Yang, F.Z. Qing, P.Gao and J.Z. Li, *Electrochim. Acta*, 2017, 235, 304.
- 4 Y. Wang, Q.T. Qu, G. Liu V.S. Battaglia and H.H. Zheng, Nano Energy, 2017, 39, 200.
- 5 K. Saravanan, M. Nagarathinam, P. Balaya and J.J. Vittal, J. Mater. Chem., 2010, 20, 8329.
- 6 X.Y. Han, F. Yi, T.L. Sun and J.T. Sun, Electrochem. Commun., 2012, 25, 136.
- 7 M. Veerababu, U.V. Varadaraju and R. Kothandaraman, *Int. J. Hydrogen Energy*, 2015,40, 14925.

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