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Li–air battery and ORR activity of nanocarbons produced with good synthesis rate by solution plasma process†

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Nanocarbons were successfully synthesized from benzene (BZ), nitro-benzene (BZ-NO₂) and aniline (BZ-NH₂) by solution plasma process (SPP). The SPP was generated by a bipolar pulsed power supply between two tungsten electrodes at room temperature. The synthesized nanocarbons were investigated. The highest synthesis rate, 40 mg min⁻¹, was for the BZ nanocarbon. The transmission electron microscopy (TEM) morphology showed that the nanocarbon sizes were 15–25 nm. The Brunauer–Emmett–Teller (BET) analysis shows a highest surface area of 220 m² g⁻¹, pore size of 0.45 cm³ g⁻¹, and average pore diameter of 20.0 nm for the BZ nanocarbon. Cyclic voltammetry (CV) in an acidic medium exhibited the oxygen reduction reaction (ORR) of the nanocarbons. The nanocarbon from BZ-NH₂ obtained a high special capacity of 15 500 mA h per g of carbon at the discharge rate of 0.1 mA cm⁻² with 1.0 mg carbon loading for the lithium (Li)–air battery. The ORR is an important reaction in Li–air batteries and fuel cells for the application of next-generation batteries and energy conversion devices.

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1. Introduction

Recently, Li–air batteries have been used in energy conversion and storage devices for various applications.^{1–4} These face critical challenges that result from poor cycle life, low practical energy density, low round-trip efficiency, and high manufacturing costs. To improve the energy storage properties, catalytic activity and reduce the cost, ORR is demanded.^{5–7}

Presently, platinum (Pt) is commonly used as a catalyst for ORR but it has high cost in the production and long term instability.^{8–10} To introduce alternatives to Pt catalysts, nanocarbons and heteroatom-doped nanocarbon have attracted attention. The advantages of nanocarbons and heteroatom-doped nanocarbon are the low production cost, excellent chemical properties and higher electrical conductivity.^{11,12}

Nanocarbons such as carbon black, carbon nanotubes, porous and nonporous carbon materials and graphene have shown large specific surface areas, high conductivity, electrocatalytic activity, mechanical capacity, ultra-light characteristics, and chemical and thermal resistance.^{13–18} Heteroatom (nitrogen, boron, and phosphorus) doped nanocarbon has shown the higher ORR activity.^{19–23}

Previously, the nanocarbon materials have been successfully synthesized by catalytic or noncatalytic chemical vapor deposition, pyrolysis, electric arc discharge, and direct-current (DC) discharge plasma from various organic solvents.^{24–29} However, these synthesis techniques are complicated due to their high process temperatures, low production yield, and the processing time.^{30–33}

Recently an energy saving and non-equilibrium liquid phase plasma named solution plasma process (SPP) has been used to synthesize nanocarbon materials at room temperature.^{34–38} In our laboratory, we conducted various experiments to produce carbon materials from organic solvents by SPP.^{39–42}

In this research, nanocarbons were synthesised by SPP with different frequencies and pulse widths at an electrode distance of 0.75 mm for 30 min. The synthesis rate is the main target in the synthesis of the nanocarbons along with their electrocatalytic activity (such as in Li–air batteries and fuel cells) without thermal treatment compared to the commercial Pt/C catalyst. The catalytic activity for ORR was obtained in acidic medium. The crystallinity, structure, surface area, morphology, elemental composition,

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electrical conductivity, ORR activity and Li-air battery performance of the nanocarbons were investigated. In the case of the Li-air battery, the N-containing nanocarbon (nanocarbon of BZ-NH₂) obtained a high special capacity of 15 500 mA h per g of carbon compared with the nanocarbon of palm oils.

2. Experimental procedure

2.1. Materials

Benzene (>99%; Kanto Chemical Co., INC., Japan), nitrobenzene (>99%; Kanto Chemical Co., INC., Japan), aniline (>99%; Kanto Chemical Co., INC., Japan), polyvinylidene fluoride (PVDF, Sigma-Aldrich, Japan), *N*-methyl-2-pyrrolidone (NMP, Kanto Chemical Co., INC., Japan), acetone (Kanto Chemical Co., INC., Japan) and ethanol (Kanto Chemical Co., INC., Japan) were purchased.

2.2. Experimental setup and production procedure

The experimental setup for SPP is shown in Fig. 1. The plasma was generated by a bipolar pulsed power supply (MPS-R06K01C-WP1-6CH, Kurita Co. Ltd, Japan) between two tungsten (W) electrodes (wire type, purity 99.9%, The Nilaco Corporation, Japan) with a 1 mm diameter, covered with an insulating ceramic tube. The electrode gap was 0.75 mm in the beginning. The concentrations of BZ, BZ-NO₂, and BZ-NH₂ are 50, 18, and 50 mg L⁻¹, respectively. The applied high voltage had different pulse widths (0.5, 1.0 and 2 μs) and repetition frequencies (50, 100, and 150 kHz). The process time was 30 min. The current and voltage waveforms were observed using an oscilloscope (DLM 2022, Yokogawa Meters & Instruments Corporation, Japan) in the range of 0–5 A and 0.6–1.0 kV, respectively.

During SPP, the electrodes were adjusted several times, especially at high frequencies of SPP operation (*i.e.*, 100 and 150 kHz) to keep a constant gap distance between the electrodes. Due to

erosion, the electrode gap distance increases. This adjustment also has the role in maintaining a stable plasma during the process. If the electrode gap distance increases from 0.75 mm to 1 mm or more, the discharge stops. After the nanocarbon materials synthesis, the remaining liquid and solid phases were separated by filtration then dried at 90 °C for five hours in an oven (DX301, Yamato Co. Ltd, Japan). The weight of nanocarbon was measured using a balance (EK2000i, AS-ONE Co. Ltd, Japan). The carbon yield (Y in %) and synthesis rate (R in mg min⁻¹) were calculated from the amount of carbon powders as:

$$Y = (M/M_0) \times 100$$

where M is the mass of the carbon powder, and M_0 is the initial mass of BZ, BZ-NO₂ and BZ-NH₂. However, the synthesis rate was calculated as $R = M/T$. Where T is total process time (*i.e.*, 30 min).

2.3. Characterization

The structural properties were investigated by X-ray diffraction (XRD, Rigaku Smart lab, Rigaku Corporation, Japan) with a Cu K α X-ray radiation source at $\lambda = 0.154$ nm in the 2θ range from 3.0 to 90.0 degrees and a scan speed of 3.0 degrees; and Raman spectroscopy (NRS-100, JASCO Corporation, Japan) used a laser with a wavelength of 532.5 nm and 1 mW power to determine the types of nanocarbons. The morphology, shape, and size of the nanocarbons were observed using transmission electron microscopy (TEM, JEM-2500SE, JEOL Ltd, Japan) with an electron gun at 200 kV. The surface area, pore volume, and average pore diameter were observed using the Brunauer–Emmett–Teller (BET, BELSORP-mini-II, BEL JAPAN, INC., Japan) method from the nitrogen adsorption–desorption isotherms at 77 K. The elemental compositions were observed by X-ray photoelectron spectroscopy (XPS, PHI 5000 Versa Probe II; Ulvac-Phi physical Electronics INC., Japan) using a Mg K α X-ray source with a photon energy of 1253.6 eV. The electrical resistivity and electrical conductivity were determined using a four-point probe method (Laresta-GP, MCP-T610, Mitsubishi Chemical Analytech Co., Ltd, Japan). A concentration of 60 wt% of nanocarbons was mixed with 40 wt% polymer binder (PVDF) and then sonicated in NMP for 30 min. The emulsion was dropped on a glass substrate, then dried in an electric oven at 120 °C for 20 min to form a uniform film.

2.4. Electrochemical measurements

Cyclic voltammetry (CV) measurements were conducted using a computer-controlled ALS-CH model 704 ES electrochemical analyzer (CH Instrument Inc.). The ink made by ultrasonication of the catalyst (5.0 mg), ultrapure water (480 mL), ethanol (480 mL), and Nafion[®] aqueous solution (40 mL) was obtained. The Ag/AgCl solution was applied as the counter and reference electrodes with 0.5 mol L⁻¹ H₂SO₄ electrolyte under N₂ and O₂ gas saturation. The scan rate is 20 mV s⁻¹ at -0.2 V to 1.2 V (*vs.* Ag/AgCl).

2.5. Fabrication of Li-air battery

A Li-air battery was fabricated as shown in Fig. 2. The air electrode was fabricated by a mixture of nanocarbon and polyvinylidene difluoride (PVDF) as binder in a weight ratio of 70:30 at 8 MPa.

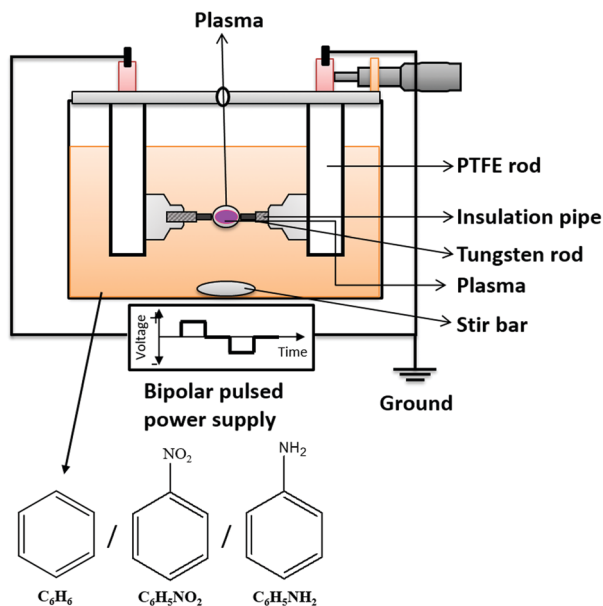


Fig. 1 Schematic diagram of experimental setup with precursors BZ, BZ-NO₂ and BZ-NH₂ for solution plasma process (SPP).



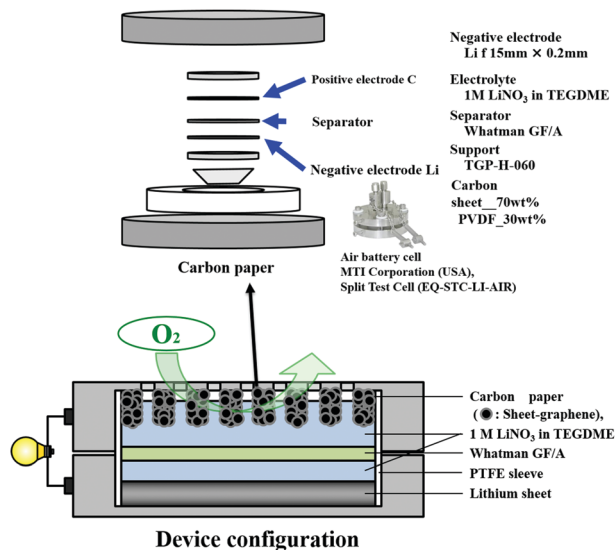


Fig. 2 Image of the Li-air battery fabricated from nanocarbon.

Lithium foil (thickness: 1 mm) was purchased (Honjo Chemical Corp., Japan). Air battery cells (MTI Corporation, USA) and the electrode of 1 M LiNO_3 in tetraethylene glycol dimethyl ether (TEGDME) were used. Battery tests were conducted with a Solar Tron SI 1280 B at current density of 0.1 mA cm^{-2} for discharge and 0.5 mA cm^{-2} for charge. The cut-off voltage in the discharge was set to 2.0 V.

3. Results and discussion

3.1. Synthesis rate

Nanocarbons were synthesized from BZ, BZ- NO_2 and BZ- NH_2 using SPP. After SPP, the synthesized nanocarbons were separated from the liquid, dried and the synthesis rates calculated (Fig. 3). The highest carbon yields (γ) for BZ, BZ- NO_2 and BZ- NH_2 are 1.92, 0.32 and 0.48%, respectively, at 150 kHz (see Table S1, ESI[†]). The nanocarbons were showed the lowest synthesis rates

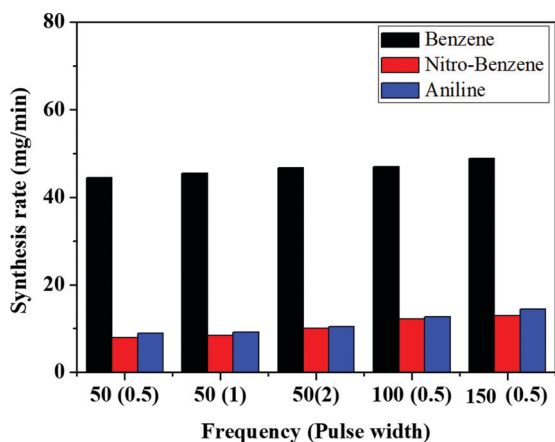


Fig. 3 Graph of the synthesis rate vs. frequency (pulse width) of the synthesized nanocarbons.

(42, 6 and 7 mg min^{-1}) at 50 kHz, 0.5 μs , and highest synthesis rates (50, 13 and 15 mg min^{-1}) at 150 kHz, 0.5 μs , for BZ, BZ- NO_2 and BZ- NH_2 respectively. It confirms that when the frequencies were gradually increased, the synthesis rate also increased with them.⁴³ In the case of BZ, the synthesis rate of the nanocarbon was higher than for BZ- NO_2 and BZ- NH_2 due to their structures. It means that only the π -conjugated bond is successfully enhanced in the nanocarbon synthesis by SPP. The schematic of nanocarbon formation from BZ using SPP is shown in Fig. S1 (ESI[†]). In this study, the highest synthesis rates of the nanocarbons at 150 kHz, 0.5 μs are described for the future investigation. So far in the paper, we have selected for the discussion only the data corresponding to the frequency of 150 kHz with a pulse width of 0.5 μs .

3.2. Morphology and surface area

The TEM images confirmed the morphology. The morphologies of the nanocarbons from BZ, BZ- NO_2 and BZ- NH_2 by SPP are seen in Fig. 4(a)–(c). The average diameters were in the 15–25 nm range with carbon in a ball-like shape.⁴⁴ The nanocarbons were agglomerated with a few on the surface. The high-resolution TEM (HRTEM) images are shown in Fig. 4(a')–(c'). Graphite-like structures with a good nanocarbon crystallinity were observed in the corresponding SAED images in Fig. 4(a'')–(c''). The SAED images show the (002), (100), and (101) planes of the nanocarbon. These data are confirmed in the XRD patterns.

The nanocarbon characteristics such as surface area, pore size, and diameter were determined using the Brunauer–Emmett–Teller (BET) method (Table S2, ESI[†]). For the BZ, BZ- NH_2 and BZ- NO_2 nanocarbons, the BET surface areas were 220, 230 and

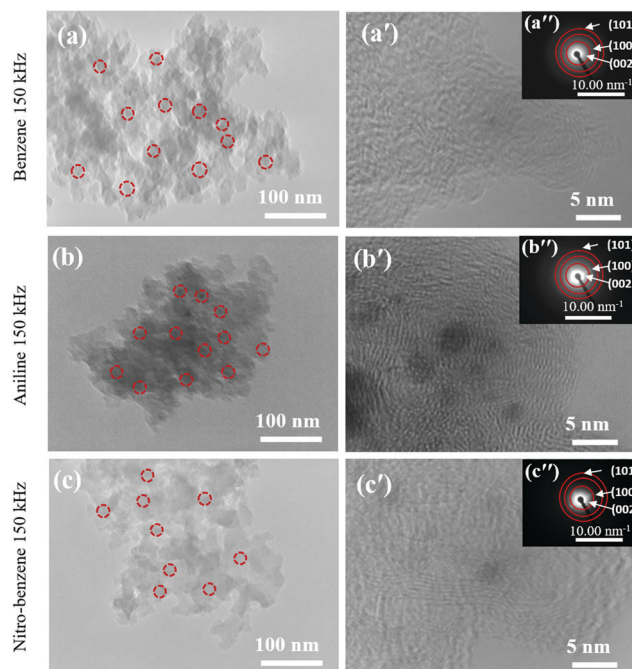


Fig. 4 (a–c) TEM images of the nanocarbons; (a'–c') HRTEM images; and (a''–c'') SAED patterns (inset) for BZ, BZ- NH_2 and BZ- NO_2 at 150 kHz and 0.5 μs .



222 m² g⁻¹, the pore volumes were 0.45, 0.49 and 0.46 cm³ g⁻¹, and the average pore diameters were 20.0, 17.0 and 15.0 nm, respectively. The nanocarbon from BZ-NH₂ had an enhanced surface area of 230 m² g⁻¹.

3.3. Structural properties

The measured XRD patterns and Raman spectra are shown in Fig. 5. The XRD patterns of the BZ, BZ-NH₂ and BZ-NO₂ nanocarbons (Fig. 5(a)) show a large peak at 23° and a small broad peak at 44° which correspond to the (002) and (100)/(101) planes, respectively, of the nanocarbon.^{45,46} The small and broad carbon shapes suggested that the graphite plane (JCPDS 75-1621) had quite good crystallinity. The sharp peaks of tungsten carbide (WC) (JCPDS #25-1047) are observed in all the samples at 36°, 42°, 62° and 74° corresponding to the (111), (200), (220), (311) and (222) planes respectively.⁴⁷ In the case of the BZ-NH₂ and BZ-NO₂ nanocarbons, the WC peaks were increased due to a higher sputtering of the tungsten electrode and the chemical structures compared with the nanocarbon from BZ. The chemical structures of BZ (C-C, C=C and C-H), BZ-NO₂ (C-C, C=C, C-N, N=O and C-H) and BZ-NH₂ (C-C, C=C, C-N, N-H and C-H) are different. The sputtering rate

depends on the chemical structure of the solvents even under the same plasma conditions. During the solution plasma process, the solution conversion pathway has an effect on the results.

Fig. 5(b) shows that the D (1350 cm⁻¹) and G (1600 cm⁻¹) band Raman peaks were observed in all samples. The D-band showed the structural disorders and defects,⁴⁸ and the G band referred to sp² graphite sheet-like structure nanocarbon. The I_D/I_G ratios of (0.80, 1.10 and 0.9) for the BZ, BZ-NH₂ and BZ-NO₂ nanocarbons were respectively calculated from the peaks. The I_D/I_G values indicated the defect sites with degree of graphitization for the nanocarbons.⁴⁹

The I_G/I_D ratio is related to the crystallite size *L_a* using the Tuinstra-Koenig relationship as follows:⁵⁰

$$L_a \text{ (nm)} = (2.4 \times 10^{-10}) \lambda^4 (I_G/I_D),$$

where λ is the Raman excitation wavelength at 532 nm. The *L_a* values are correlated to the inter-defect distance of the surface. The *L_a* value of the BZ-NH₂ nanocarbon showed the BZ-NH₂ nanocarbon to have more defect sites due to its nitrogen molecules.

The XRD and Raman results corresponded with the TEM images and showed that the nanocarbons were graphite-like structures with crystallinity.

3.4. Electrical conductivity and composition

The four-point probe method was applied to the nanocarbons to identify resistance (Ω), surface resistance ($\Omega \text{ cm}^{-2}$), resistivity ($\Omega \text{ cm}^{-1}$) and electrical conductivity (S cm^{-1}) (see Tables S3–S5, ESI† and Fig. 6). So, we observed that when the resistivity ($\Omega \text{ cm}^{-1}$) of the nanocarbon was decreased, the conductivity was increased. In the case of the BZ nanocarbon, it shows the highest electrical conductivity compared with the nanocarbons from BZ-NH₂ and BZ-NO₂. The elemental composition was found using XPS spectroscopy, as shown in Fig. S2 (ESI†). The BZ nanocarbon (C1s 99.6, and W4f 0.4 atom%), BZ-NH₂ nanocarbon (C1s 98.2, N1s 1.6, and W4f 0.2 atom%), and BZ-NO₂ nanocarbon (C1s 97.1, O1s 15.9, N1s 5.0 and W <0.1, atom%) were identified at 150 kHz and 0.5 μs . The XPS spectroscopy

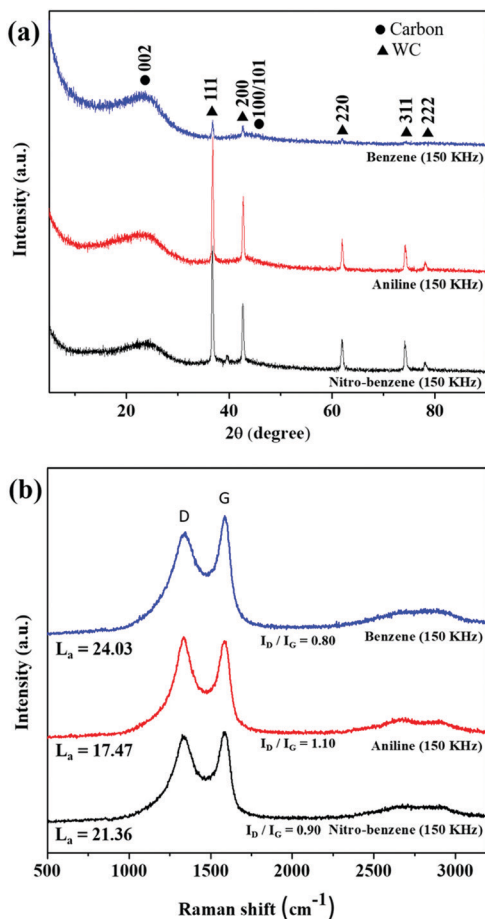


Fig. 5 The structural properties of the nanocarbons. (a) XRD patterns, and (b) Raman spectra for the nanocarbons from BZ, BZ-NH₂ and BZ-NO₂ at 150 kHz and 0.5 μs .

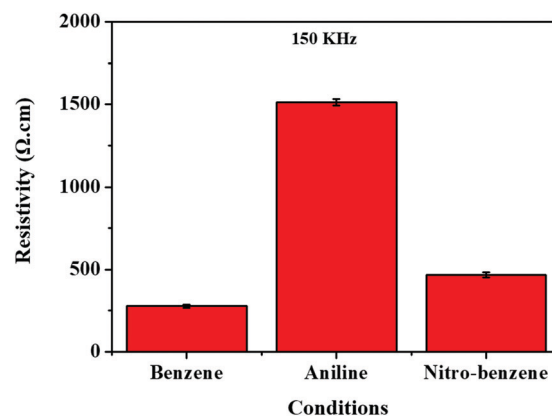


Fig. 6 Electrical resistivity using the four probe method for the nanocarbons under various conditions.



data showed that BZ nanocarbon contained carbon (C) and tungsten (W); BZ-NH₂ nanocarbon contained carbon (C), nitrogen (N) and tungsten (W); and BZ-NO₂ nanocarbon contained carbon (C), oxygen (O), nitrogen (N) and tungsten (W), corresponding with the XRD patterns.

3.5. Catalytic activity

The ORR electrocatalytic activity was also tested in 0.5 mol L⁻¹ H₂SO₄ solution with N₂ and O₂ saturation at 20 mV s⁻¹ scan rate, as shown in Fig. 7(a)–(c). The anodic and cathodic peaks are shown in the potential range of -0.2 to 1.2 V. Under N₂-saturation featureless CV curves can be observed for the BZ and BZ-NO₂ nanocarbons as shown in Fig. 7(a) and (b). On the other hand, the CV curve with O₂-saturated electrolyte shows a cathodic peak around 0.4 V corresponding to ORR. The ORR catalytic activity has use in fuel-cell applications as a cathode. Fig. 7d presents the LSV curves of BZ, BZ-NO₂ and BZ-NH₂ in O₂-saturated 0.5 mol L⁻¹ H₂SO₄ solution at a scan rate of 10 mV s⁻¹ with a rotation speed of 1600 rpm. The LSV curve was obtained with an onset potential of 0.39 V for BZ-NH₂. Moreover, the current density of BZ-NH₂ was larger than for BZ and BZ-NO₂ and was investigated. The BZ-NH₂ nanocarbon showed a synergistic effect of N and WC on the ORR activity. The specific capacity shows such a difference to various nanocarbons due to the properties of the tungsten (W), nitrogen (N) and oxygen (O) containing nanocarbons. In the case of BZ-NH₂, the synergistic effect of N and WC doping on the ORR activity was compared with BZ and BZ-NO₂. BZ-NH₂ showed enhanced carbon properties, such as surface area, resistivity, nitrogen configuration and degree of graphitization, which played a role in the improved ORR activity in the acid media compared with BZ and BZ-NO₂.

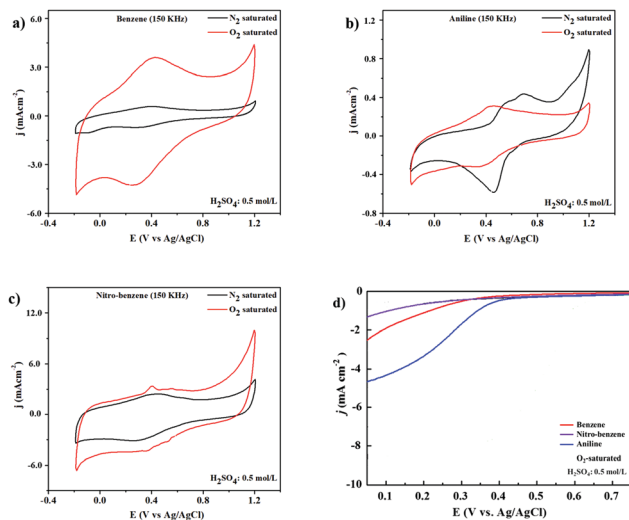


Fig. 7 CV measurements of (a) benzene (BZ), (b) aniline (BZ-NH₂) and (c) nitrobenzene (BZ-NO₂) in 0.5 mol L⁻¹ H₂SO₄ solution at a scan rate of 20 mV s⁻¹ under N₂ (black lines) and O₂ (red lines) saturation. LSV measurements: (d) presents the LSV curves of BZ, BZ-NH₂, and BZ-NO₂ in O₂-saturated 0.5 mol L⁻¹ H₂SO₄ solution at a scan rate of 10 mV s⁻¹ with a rotation speed of 1600 rpm.

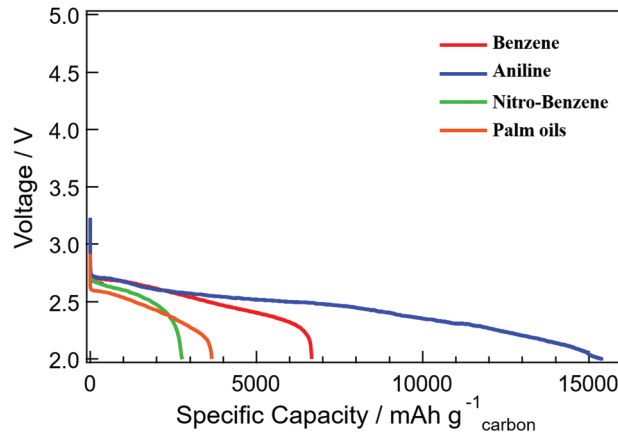


Fig. 8 The discharge curves of the Li-air batteries with 1.0 mg carbon loadings at 0.1 mA cm⁻² rates for nanocarbons from BZ, BZ-NH₂, BZ-NO₂ and palm oils.

3.6. Li-air battery test

The electrochemical performances of the cell voltage and capacity of the Li-air batteries, the discharge of 1.0 mg cm⁻² carbon loading in 1.0 atom of dry oxygen at 0.1 mA cm⁻² rates, are shown in Fig. 8. The specific capacities of 2800, 3600, 6700 and 15000 mA h per g of carbon were identified for the nanocarbons from BZ-NO₂, palm oils, BZ and BZ-NH₂ respectively at the rate of 0.1 mA cm⁻² with the 1.0 mg cm⁻² carbon loading. The highest special capacity of 15500 mA h per g of carbon was obtained for the BZ-NH₂ nanocarbon, compared with the palm oils nanocarbon.

4. Conclusions

We have synthesized the nanocarbons by solution plasma process. The highest synthesis rate was observed for the nanocarbon from BZ. The graphite-like structure and the 15–25 nm sized nanocarbons were investigated using the TEM images supported by XRD and Raman spectroscopy measurements. In the ORR electrocatalytic activity of the nanocarbons, the cathodic peak at 0.4 V was clearly observed for all samples. The performance of the Li-air battery, with the highest special capacity of 15500 mA h per g of carbon for the BZ-NH₂ nanocarbon, was compared with the nanocarbon of palm oils. We believe that SPP could be able to synthesise nanocarbons with electrocatalytic activity for applications such as Li-air batteries and fuel cells.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 S. W. Lee, N. Yabuuchi, B. M. Gallant, S. Chen, B.-S. Kim, P. T. Hammond and Y. Shao-Horn, *Nat. Nanotechnol.*, 2010, **5**, 531.
- 2 A. Chen and P. Holt-Hindle, *Chem. Rev.*, 2010, **110**, 3767.
- 3 X. Ma, H. Meng, M. Cai and P. Shen, *J. Am. Chem. Soc.*, 2012, **134**, 1954.
- 4 Y. Zhao, L. Yang, S. Chen, X. Wang, Y. Ma, Q. Wu, Y. Jiang, W. Qian and Z. Hu, *J. Am. Chem. Soc.*, 2013, **135**, 1201.
- 5 B. Winther-Jensen, O. Winther-Jensen, M. Forsyth and D. R. MacFarlane, *Science*, 2008, **321**, 671.
- 6 Y. Zheng, Y. Jiao, J. Chen, J. Liu, J. Liang, A. Du, W. Zhang, Z. Zhu, S. C. Smith, M. Jaroniec, G. Q. Lu and S. Z. Qiao, *J. Am. Chem. Soc.*, 2011, **133**, 20116.
- 7 Z. Schnepp, Y. Zhang, M. J. Hollamby, B. R. Pauw, M. Tanaka, Y. Matsushita and Y. Sakka, *J. Mater. Chem. A*, 2013, **1**, 13576.
- 8 H. Wang, T. Maiyalagan and X. Wang, *ACS Catal.*, 2012, **2**, 781.
- 9 C. Soldano, A. Mahmood and E. Dujardin, *Carbon*, 2010, **48**, 2127–2150.
- 10 A. Manikandan, L. Lee, Y.-C. Wang, C.-W. Chen, Y.-Z. Chen, H. Medina, J.-Y. Tseng, Z. M. Wang and Y.-L. Chueh, *J. Mater. Chem. A*, 2017, **5**, 13320–13328.
- 11 Z. Yang, W. Sun, L. Wang, Z. Wan, J. Wang, S. Wang and G. Liu, *Corros. Sci.*, 2020, **175**, 108860.
- 12 M. Topsakal, H. Şahin and S. Ciraci, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2012, **85**, 155445.
- 13 J. Liang, Y. Jiao, M. Jaroniec and S. Z. Qiao, *Angew. Chem., Int. Ed.*, 2012, **51**, 11496–11500.
- 14 Q. Lv, W. Si, J. He, L. Sun, C. Zhang, N. Wang, Z. Yang, X. Li, X. Wang and W. Deng, *Nat. Commun.*, 2018, **9**, 1–11.
- 15 S. Chen, A. Zehri, Q. Wang, G. Yuan, X. Liu, N. Wang and J. Liu, *ChemistryOpen*, 2019, **8**, 58–63.
- 16 W. Lei, H. M. Barnes, J. Zhang and Z. Cai, *Wood Fiber Sci.*, 2017, **49**, 22–32.
- 17 S. W. Lee, B. M. Gallant, H. R. Byon, P. T. Hammond and Y. Shao-Horn, *Energy Environ. Sci.*, 2011, **4**, 1972.
- 18 S. W. Lee, B.-S. Kim, S. Chen, Y. Shao-Horn and P. T. Hammond, *J. Am. Chem. Soc.*, 2009, **131**, 671.
- 19 A. Watthanaphanit, G. Panomsuwan and N. Saito, *RSC Adv.*, 2014, **4**, 1622.
- 20 G. Panomsuwan, A. Watthanaphanit, T. Ishizaki and N. Saito, *Phys. Chem. Chem. Phys.*, 2015, **17**, 13794.
- 21 T. Ishizaki, S. Chiba, Y. Kaneko and G. Panomsuwan, *J. Mater. Chem. A*, 2014, **2**, 10589–10598.
- 22 T. Morishita, T. Ueno, G. Panomsuwan, J. Hieda, A. Yoshida, M. A. Bratescu and N. Saito, *Sci. Rep.*, 2016, **6**, 36880.
- 23 X. Hu, X. Shen, O. Takai and N. Saito, *J. Alloys Compd.*, 2013, **552**, 351–355.
- 24 S. Chae, G. Panomsuwan, M. A. Bratescu, K. Teshima and N. Saito, *ACS Appl. Nano Mater.*, 2019, **2**, 1350–1355.
- 25 C. Chokradjaroen, S. Kato, K. Fujiwara, H. Watanabe, T. Ishii and T. Ishizaki, *Sustainable Energy Fuels*, 2020, **4**, 4570–4580.
- 26 O. L. Li, S. Chiba, Y. Wada, G. Panomsuwan and T. Ishizaki, *J. Mater. Chem. A*, 2017, **5**, 2073–2082.
- 27 M. A. Bratescu, O. Takai and N. Saito, *J. Alloys Compd.*, 2013, **562**, 74–83.
- 28 K. Hyun, T. Ueno, O. L. Li and N. Saito, *RSC Adv.*, 2016, **6**, 6990–6996.
- 29 C. H. Choi, S. H. Park and S. I. Woo, *J. Mater. Chem.*, 2012, **22**, 12107–12115.
- 30 G. Panomsuwan, N. Saito and T. Ishizaki, *J. Mater. Chem. A*, 2015, **3**, 9972–9981.
- 31 N. Thongwichit, O. L. H. Li, W. Yaowarat, N. Saito and U. Suriyapraphadilok, *Jpn. J. Appl. Phys.*, 2016, **55**, 01AE10.
- 32 J. Kim, J. Chun, S.-G. Kim, H. Ahn and K. C. Roh, *J. Electrochem. Sci. Technol.*, 2017, **8**, 338–343.
- 33 D.-w. Kim, O. L. Li and N. Saito, *Phys. Chem. Chem. Phys.*, 2014, **16**, 14905–14911.
- 34 O. Takai, *Pure Appl. Chem.*, 2008, **80**, 2003–2011.
- 35 G. Panomsuwan, N. Saito and T. Ishizaki, *Phys. Chem. Chem. Phys.*, 2015, **17**, 6227.
- 36 W. Wang, S. Chakrabarti, Z. Chen, Z. Yan, M. O. Tade, J. Zou and Q. Li, *J. Mater. Chem. A*, 2014, **2**, 2390–2396.
- 37 T. Sudare, T. Ueno, A. Watthanaphanit and N. Saito, *Phys. Chem. Chem. Phys.*, 2015, **17**, 30255–30259.
- 38 K. Hyun, T. Ueno, O. L. Li and N. Saito, *RSC Adv.*, 2016, **6**, 6990–6996.
- 39 T. Morishita, T. Ueno, G. Panomsuwan, J. Hieda, A. Yoshida, M. A. Bratescu and N. Saito, *Sci. Rep.*, 2016, **6**, 36880.
- 40 J. Kang, O. L. Li and N. Saito, *Nanoscale*, 2013, **5**, 6874.
- 41 D. Kim, O. L. Li, P. Pootawang and N. Saito, *RSC Adv.*, 2014, **4**, 16813.
- 42 S. Lee, Y. Heo, M. A. Bratescu, T. Ueno and N. Saito, *Phys. Chem. Chem. Phys.*, 2017, **19**, 15264–15272.
- 43 O. L. Li, H. Hayashi, T. Ishizaki and N. Saito, *RSC Adv.*, 2016, **6**, 51864.
- 44 D. Kim, O. L. Li and N. Saito, *Phys. Chem. Chem. Phys.*, 2015, **17**, 407.
- 45 G. Panomsuwan, S. Chiba, Y. Kaneko, N. Saito and T. Ishizaki, *J. Mater. Chem. A*, 2014, **2**, 18677.
- 46 X. Zhang and Z. Lix, *Nanoscale*, 2012, **4**, 707–714.
- 47 T. Murphy, R. Hayes, S. Imberti, G. G. Warr and R. Atkin, *Phys. Chem. Chem. Phys.*, 2014, **16**, 13182–13190.
- 48 K. F. Adekunle, *Open J. Polym. Chem.*, 2015, **5**, 34–40.
- 49 N. Taufiqurrahmi and S. Bhatia, *Energy Environ. Sci.*, 2011, **4**, 1087–1112.
- 50 L. G. Cançado, K. Takai, T. Enoki, M. Endo, Y. A. Kim, H. Mizusaki, A. Jorio, L. N. Coelho, R. M. Paniago and M. A. Pimenta, *Appl. Phys. Lett.*, 2006, **88**, 163106.

