Nitrogen-enriched carbon electrodes in electrochemical capacitors: investigating accessible porosity using CM-SANS†

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Carbon electrochemical capacitor electrodes containing nitrogen groups were studied with respect to their electrochemical behaviour, chemical composition and physical characteristics. Thermal treatment of nitrogen-enriched carbon materials in different atmospheres was used to control the specific type and concentration of nitrogen groups present, while importantly retaining similar pore size distributions. Pyridinic nitrogen is shown to be most likely responsible for increased values of surface area normalized specific capacitance, although the mechanisms by which this occurs are poorly understood. Contrast matched-small angle neutron scattering (CM-SANS) was employed to probe the electrode porosity accessible to an electrolyte and indicates that there is no appreciable difference between the materials studied. Cyclic Voltammetry showed no evidence of electrode reactions occurring over the operating potential range. Therefore a greater amount of charge is displaced at pyridinic sites during the charge–discharge process. This may occur due to a specific adsorption mechanism, coupled with enhanced electron conductivity through the carbon matrix.

Introduction

Electrochemical capacitors (ECs), also referred to as supercapacitors, are electrical energy storage devices that absorb and release charge at relatively high rates when compared with electrochemical cells. In addition to high power density, they exhibit exceptionally long cycle lives and high efficiency, enabling them to complement batteries and fuel cells in systems where there is a substantial difference between peak and average power demands (e.g. in electric vehicles).

Charge is stored through two main mechanisms; electrical double-layer capacitors (EDLCs) store charge electrostatically at the interface between electrode surfaces and electrolyte ions, whereas pseudocapacitance is more akin to the processes that occur in battery electrodes. The most commonly used electrode materials in EDLCs are activated carbons due to their combination of electrical conductivity, high specific surface area (SSA) and the porous structure of carbon materials can be tailored to better accommodate specific electrolyte ions. In addition, the capacitance associated with some activated carbons has been shown to include a significant contribution through pseudocapacitive reactions involving oxygen containing functional groups, which has inspired research into the effects of incorporating alternative heteroatoms into the carbon matrix.

Several studies have concluded that the inclusion of nitrogen atoms into the carbon electrode structure can enhance the performance of ECs although it has been noted that the mechanism by which this occurs are unclear. Proposed mechanisms include pseudocapacitive reactions involving electrolyte species, a larger surface area being rendered electrochemically active (through enhanced wettability of the electrode surface) and superior electron conduction through the carbon matrix due to the electron donor effect of certain nitrogen groups.

Reports generally normalize specific capacitances by a measure of surface area to account for changes in the double-layer component of the overall capacitance however this approach is problematic when comparing materials with different pore and structural characteristics as normalized values for activated carbons are known to vary significantly. Also, capacitance measured on carbon surfaces can vary by up to an order of magnitude depending on the orientation of graphene planes, and it has been reported that the pore characteristics of the electrode exert substantial influence over specific capacitance.

Nitrogen-enriched carbon materials (NECMs) are normally synthesized by impregnating a carbon (or carbon precursor) with nitrogen-containing organic compounds that subsequently decompose during thermal treatment. A further thermal treatment...
step under a reducing atmosphere was employed in this work to control both the concentration and type of nitrogen groups present, while retaining similar pore and structural characteristics. This approach substantially reduces the uncertainties associated with the process of surface area normalization. Furthermore this method removes the influence of electrochemically active surface oxygen groups, such as quinone, which can complicate the interpretation of electrochemical behaviour. Herein, we present work showing that the increased capacitance associated with NECMs is not due to increased surface accessibility through the use of contrast-matched small angle neutron scattering (CM-SANS). This technique is proposed as a unique tool for the investigation of inaccessible porosity in porous electrodes.

**Experimental**

Nitrogen-enriched carbon materials (NECMs) were produced by carbonisation, activation and further thermal treatment of a resorcinol-formaldehyde xerogel impregnated with m-phenylene diamine. The NECM produced after treatment under hydrogen is denoted as ‘C–H’, the material treated under an argon atmosphere as ‘C–Ar’, and that treated under a mixed atmosphere as ‘C–HAr’.

Nitrogen adsorption–desorption isotherms at 77 K were obtained using a Micromeritics ASAP 2420 instrument; samples were degassed at 200 °C for 10 h before analysis. Elemental analysis was performed using a Perkin Elmer 2400 Series II CHNS/O Elemental Analyser, and proximate analysis was performed using a Mettler Toledo TGA/DSC1 STARe instrument. XPS was performed in an ion pumped VG Microtech CLAM 4 MCD analyser system using 200 W unmonochromated Mg X-ray excitation (1253.6 eV).

Small-angle neutron scattering (SANS) was performed using the LOQ small-angle diffractometer at the ISIS Pulsed Neutron Source (STFC Rutherford Appleton Laboratory, Didcot, U.K.). Deuterium oxide was identified as a suitable contrast matching liquid by calculating the scattering length density of the NECMs.

Electrodes were manufactured by mixing the NECM under study with polymer binder (KynarFlex) and milled carbon black (Vulcan XC72), in an 80/10/10 ratio by mass. Symmetric two-electrode EC cells were constructed using a version of a test cell used in previous investigations, and employing potassium hydroxide solution (28.5 wt%) as electrolyte.

Experimental tests were carried out using a Solartron Analytical 1470E Multichannel Potentiostat/Galvanostat with 1455A Frequency Response Analyser. Cell temperature was maintained at 25 °C (±0.1 °C). Cells were cycled between 0.0 and 1.0 V (vs. OCP) Galvanostatically at various rates between 0.1 and ca. 12 A g⁻¹. Electrochemical impedance spectra were obtained at a potential bias of 0.1 V (vs. OCP) using a 10 mV perturbation over the frequency range 100 kHz to 10 MHz. Cyclic Voltammetry was performed between 0.0 and 1.0 V (vs. OCP) at sweep rates of 10 mV s⁻¹ and 100 mV s⁻¹. Specific capacitance values are based on the mass of dry, ash-free active material and are expressed on a three-electrode basis.

More details of the synthesis method, physical characterization, and electrochemical testing of the NECMs is given as [ESI†].

**Results and discussion**

Nitrogen adsorption–desorption isotherms for the NECMs studied are shown in Fig. 1. The isotherms are a mixture of ‘type I’ and ‘type IV’ form although the uptake at low relative pressures differs for each sample indicating that micropore volume is dependent on the hydrogen concentration during thermal treatment. SSAs and pore characteristics for each sample derived from the isotherms are given in Table 1.

Although there is an increase in SSA and total pore volume with increasing hydrogen concentration during thermal treatment, these materials have similar pore characteristics, which allows for comparisons between their capacitances to be made on a surface area normalised basis. The chemical composition of the samples determined using both combustive elemental analysis and X-ray photoelectron spectroscopy (XPS), a surface specific technique, are given in Table 2.
Nitrogen content is seen to decrease with increasing hydrogen concentration indicating that thermal treatment under hydrogen is a useful technique by which to control the amount of nitrogen groups in NECMs. Table 2 also shows that C–H possesses the lowest nitrogen content while displaying the largest micropore volume, therefore the abstraction of nitrogen groups is accompanied by the evolution of accessible porosity. Nitrogen content was then divided into the groups illustrated in Fig. 2 by deconvolution of detailed XPS scans and is expressed as an absolute surface concentration in Table 3.

Table 3 shows that the gas used in the thermal treatment process not only controls the amount of nitrogen in the NECMs but can also be used to control the concentration of specific nitrogen groups. By comparing the composition of C–H and C–Ar it is seen that the reduction is attributable to the removal of N-5 and N-6 type groups. Table 2 also shows that a considerable amount of oxygen atoms are present in the materials, however detailed XPS scans show that the oxygen content is not in the form of electrochemically active quinone groups.

Small-angle neutron scattering (SANS) has been employed to investigate the pore characteristics of carbon materials used in ECs and is especially suited to the study of materials that possess microporosity. Additionally, the technique of contrast-matched SANS has been used to identify the degree of closed porosity in porous carbon materials and has been applied in this work to investigate the amount of porosity accessible to an electrolyte. It is proposed that any increase in electrochemically active surface area resulting from the presence of nitrogen groups would appear as a difference in the scattering behaviour of a sample that has been impregnated with a liquid of the same scattering length density. For the NECMs studied, the calculated scattering length density of the samples almost identical to that of deuterium oxide, which was employed as a suitable analogue to the aqueous electrolyte used in electrochemical testing.

The difference in scattering cross section between the dry and contrast matched carbon materials (of up to three orders of magnitude) seen in Fig. 3 confirms that an excellent degree of contrast matching was obtained.

For each of the NECMs, the SANS plots in Fig. 3 are of a similar form which confirms that their pore characteristics are comparable. For values of scattering wave vector (q) greater than 0.1 Å⁻¹, corresponding to the micropore region, both C–H and C–HAr exhibit a greater amount of scattering in comparison with C–Ar, which correlates well with the micropore volumes presented in Table 1. Further analysis of the SANS data confirms the trend in SSA seen in Table 1 (see ESI†). Where scattering occurs purely due to micropores there remains a difference between the dry and contrast-matched scattering curves, which confirms that even the smallest pores in each NECM are open and filled by the aqueous electrolyte used in electrochemical testing.

Symmetrical two-electrode cells were constructed using the NECMs and aqueous potassium hydroxide solution electrolyte. Cells were tested using electrochemical impedance spectroscopy, obtained spectra are given as ESI†. Parameters derived from the spectra are presented in Table 4.

C–H could be expected to produce the largest specific capacitance through the EDLC mechanism by possessing the highest SSA, however C–Ar exhibited the highest specific capacitance confirming that nitrogen heteroatoms can have a beneficial effect on the capacitance. However, polarization resistance also increases with nitrogen content and therefore a trade-off exists between increased capacitance and an increased equivalent series resistance (and therefore reduced power density).

When specific capacitance is normalized on the basis of SSA, the influence of nitrogen content becomes clearer as the two samples that contain a low concentration of edge nitrogen groups have a similar value, whereas C–Ar, which has a significantly larger concentration of edge nitrogen groups, has a value of 20.3 μF cm⁻². The finding that an increased surface area normalized specific
capacitance is associated with greater nitrogen contents is in agreement with several reports.

Moreover, by considering the concentration of the nitrogen groups present, the best correlation is found with the pyridinic form of nitrogen (N-6). It is therefore likely that the increase in capacitance is specifically associated with pyridinic nitrogen (N-6). This can be asserted due to the different nitrogen groups classified as N-5 type, which are found at a binding energy of 400.6 eV. The situation whereby pyrollic nitrogen and the tautomeric forms of pyridonic nitrogen found at this binding energy behave in an identical manner electrochemically, is far less likely than the situation where pyridinic nitrogen is responsible for the larger surface area normalized specific capacitance observed.

Specific capacitance was also determined under Galvanostatic discharge over a range of current densities and the results are presented in Fig. 4. Capacitance values determined at relatively low rates are in agreement with those determined using EIS. The surface area normalized values of specific capacitance in Fig. 4 are higher for C–Ar at all of the rates studied than the samples possessing very little pyridinic nitrogen. The fact that the there is no rate dependence of this effect could indicate that reactions involving N-6 groups do contribute to the overall capacitance of the electrode.

In order to determine whether any potential dependent reactions influence the electrochemical behaviour of the samples Cyclic Voltammetry (CV) was performed in symmetrical two-electrode cells. Electrodes exploiting known pseudocapacitive reactions (e.g. metal oxides,1 quinone/hydroquinone10,26) usually produce peaks in CVs over a voltage range of roughly 400 mV.

**Table 4** Characteristics determined by fitting Randles’ circuit to EIS data for NECMs. (NECM treated under hydrogen ‘C–H’, argon ‘C–Ar’, mixed atmosphere ‘C–HAr’)

<table>
<thead>
<tr>
<th></th>
<th>(c^e [\text{F g}^{-1}])</th>
<th>(c_{\text{SSA}} ^b [\text{F cm}^{-2}])</th>
<th>(R_s ^c [\Omega \text{ cm}^{-2}])</th>
<th>(R_p ^c [\Omega \text{ cm}^{-2}])</th>
</tr>
</thead>
<tbody>
<tr>
<td>C–H</td>
<td>85.4</td>
<td>14.7</td>
<td>0.07</td>
<td>0.03</td>
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<tr>
<td>C–HAr</td>
<td>76.1</td>
<td>14.2</td>
<td>0.07</td>
<td>0.04</td>
</tr>
<tr>
<td>C–Ar</td>
<td>94.5</td>
<td>20.3</td>
<td>0.07</td>
<td>0.06</td>
</tr>
</tbody>
</table>

*a* Specific capacitance determined at 10 MHz and expressed on a three-electrode basis using the active mass of NECM. *b* Specific capacitance normalized on the basis of surface area SSA given in Table 2. *c* Series resistance \( (R_s) \) and polarization resistance \( (R_p) \) determined by fitting Randles’ circuit28 to the impedance spectra (see ESI).

**Fig. 3** Small angle neutron scattering (solid lines) and contrast matched small angle neutron scattering (dashed lines) plots for (a) NECM treated under argon (C–Ar), (b) NECM treated under a hydrogen–argon mixed atmosphere (C–HAr) and (c) NECM treated under hydrogen (C–H).

**Fig. 4** Normalized Galvanostatic discharge capacitance against specific discharge current. Circles: NECM treated under hydrogen (C–H), squares: mixed hydrogen–argon atmosphere (C–HAr), triangles: argon (C–Ar).

**Fig. 5** Cyclic Voltammograms at (a) 10 mV s\(^{-1}\) and (b) 100 mV s\(^{-1}\). Current is normalized based on the mass of NECM. Circles: NECM treated under hydrogen (C–H), squares: mixed hydrogen–argon atmosphere (C–HAr), triangles: argon (C–Ar).

**Conclusions**

NECMs treated in the presence of hydrogen were used to investigate the mechanisms by which nitrogen groups influence the electrochemical behaviour of EC electrodes. CM-SANS experiments indicate that there is no discernable difference in electrolyte
accessibility between the NECMs studied and CV showed no evidence of electrode reactions occurring over the operating potential range. CM-SANS could be an extremely useful technique by which to study electrolyte accessibility in systems where there is interaction between electrodes and liquid electrolytes (e.g. batteries, dye-sensitized solar cells etc.). Electrochemical methods confirmed that nitrogen groups can have a beneficial effect on specific capacitance, and it was concluded that pyridinic nitrogen (N-6) in particular is likely to be responsible for this effect. It is proposed that a greater amount of charge is displaced at pyridinic sites during the charge–discharge process, and that a specific adsorption mechanism coupled with enhanced electron conductivity through the carbon matrix is responsible for this effect.

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Notes and references

8 V. V. N. Obreja, Physica E, 2008, 40, 2596–2605.
31 http://www.isis.stfc.ac.uk/.