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Overcoming diffusion limitations in supercapacitors using layered electrodes

R. Drummond, C. Huang, P.S. Grant, S.R. Duncan

Department of Engineering Science, University of Oxford, 17 Parks Road, OX1 3PJ, Oxford, United Kingdom
Department of Material Science, University of Oxford, Parks Road, OX1 3PH, Oxford, United Kingdom

**HIGHLIGHTS**

- An electrochemical supercapacitor model for layered electrodes is pro-posed.
- The model is validated against experimental impedance data.
- Electrode layering is shown to increase ion mobility across the electrode.
- Layered electrodes extend the frequency range for charge storage in supercapacitors.

**ABSTRACT**

The impact of multi-layered electrode microstructures on the dynamic capacitance of electrochemical double layer supercapacitors is investigated. An electrochemical model that describes ion diffusion and double layer dynamics across the layered electrodes is first developed and then matched to experimental data. With TiO$_2$ particulate and carbon nanotube layered electrodes, two knee frequencies were observed in the real and imaginary capacitance plots in both experiment and model simulations. These two knee frequencies resulted in an increase in real capacitance at high frequencies ($\omega \approx 10^9 - 10^{10}$ rad s$^{-1}$) but a reduction at lower frequencies ($\omega \approx 10^2$ rad s$^{-1}$), with the response being largely insensitive to the relative layer thicknesses. The increased capacity at high frequencies was due to increased ion mobility across the electrodes caused by the layering, allowing diffusion limitations of identical homogeneous electrodes to be overcome. These results imply the suitability of layered electrodes for applications with highly dynamic charge profiles and/or relatively thick electrodes.

**1. Introduction**

Supercapacitors with approximately ten times higher power densities (5–50 kW kg$^{-1}$) and significantly longer cycle lives ($>10^4$ cycles) but lower energy densities (3–40 Wh kg$^{-1}$) than Li ion batteries are attractive for alternating current (AC) ripple filtering, hybrid electric vehicles and meeting peak power demands in electrical grid storage [21, 24, 27]. Consisting of a porous separator sandwiched between two identical electrodes and immersed in a liquid electrolyte, supercapacitors predominantly store charge by fast electrostatic adsorption of electrolyte ions onto the surface of the oppositely charged electrodes, as illustrated in Fig. 1. The main benefit of storing charge on the electrode surface in this way is that it leads to high power densities, being principally limited only by the mobility of the ions across the separator and through the porous electrode. However, surface only energy storage leads to relatively low energy densities, which has motivated the development of active materials with exceptionally high surface areas (500–2000 m$^2$ g$^{-1}$) for supercapacitor electrodes, including activated carbon [5], carbon nanotubes (CNTs) [33], graphene [27] and metal oxide nanomaterials [19].

In contrast to conventional supercapacitor electrodes that are random mixtures of active, electron conducting and binder materials, this paper considers the modelling and design of supercapacitors with structured electrodes. The term structured electrodes here refers to spatially heterogeneous electrodes composed of layers with different materials [20], as illustrated in Fig. 2. Recent experimental data has

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* Corresponding author.

E-mail addresses: ross.drummond@eng.ox.ac.uk (R. Drummond), ann.huang@materials.ox.ac.uk (C. Huang), patrick.grant@materials.ox.ac.uk (P.S. Grant), stephen.duncan@eng.ox.ac.uk (S.R. Duncan).

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shown that layering can give attractive combinations of energy and power density for solid-state supercapacitors [20] and Li-ion batteries [8], although unambiguously resolving the underlying reasons for improved behaviour is complex. Here, a model is developed to better understand and probe the underlying ionic mobility in structured electrodes and its links to energy storage performance, revealing the benefits of structured electrodes for applications where ionic diffusion limitations dominate, such as in high frequency charging and the filtering of ripple current fluctuations at approximately 120 Hz.

AC ripple filtering is a particularly relevant application for the consideration of structured electrodes and involves the filtering of high order harmonics to protect against electrical power surges and spikes [26]. Such filtering has been shown to be critical for the efficient exploitation of electrical energy generated by photovoltaics or wind turbines with intermittent renewable sources, with the ripples being caused by uncontrollable sunshine or wind flow [22]. To date, aluminium electrolytic capacitors (AECs) are the most common ripple filters due to their extremely high power densities (100–1000 kW kg\(^{-1}\)), although their low energy densities (0.01–0.1 Wh kg\(^{-1}\)) results in bulky AECs taking up large volume in electronic circuits and packs [32]. On the other hand, although there have been attempts to use supercapacitors for AC ripple filtering, their frequency response is generally considered too slow. For example, commercial supercapacitors using activated carbon electrodes behave comparably to a resistor rather than a capacitor at > 1 Hz frequency, far below the required operating frequency [17,25], and has led to recent work on the synthesis of supercapacitors using contact glow discharge electrolysis [2,23] and unstructured mixes of carbon nanofibres [31] for power line applications.

The restricted capacity performance at high frequencies is principally due to impeded mobility of the diffusing ions that carry charge through the tortuous pore structure of the electrode: overcoming or delaying this problem is addressed in this paper. In particular, it is noted that since the ionic current flow across an electrode is non-uniform (following from the diffusive nature of the electrochemical equations), a non-uniform electrode diffusivity could be exploited to promote overall current flow across an electrode. We develop a general framework to describe structured (or layered in this case) electrodes, which is material independent, and based on a consideration of the fundamental electrochemical dynamical equations. It is expected that any benefits of structured electrodes for supercapacitor dynamics will be most apparent for thick electrodes (> 100 \(\mu\)m) where diffusion limitations are most pronounced but are otherwise advantageous for volumetric energy density.

Most studies on structured electrodes are simulation based due to a previous lack of an appropriate controlled manufacturing process. However, such processes are now emerging at the laboratory and translational length-scale, based on variations of slurry casting or spray printing [20,23]. Simulation-based results have focused on the optimal design of layered electrodes [12,16,18,28–30,34], and a typical result is the potential for a 61% reduction in Ohmic drop [16]. Similarly, a multi-objective framework has been used in Ref. [29] to discuss the merits of structured electrodes, with metrics such as the standard deviation in the overpotential and resistance being compared [29]. Some of the claimed merits of structured electrodes were refuted in Ref. [11], where simulations of constant current and pulse charging were used to show that many of the aforementioned benefits could be obtained by a judicious choice of porosity in a conventional single layer electrode. In contrast in this paper, the benefits of electrode layering lies in their ability to promote ion mobility across an electrode under high-frequency dynamic currents where diffusion limitations dominate deliverable capacity.

This paper compares experimental supercapacitor impedance and other data with simulation results to explain the mechanisms and limitations of charge storage in layered electrodes. The novelty of the work lies in: (1) An improved electrochemical model for electric double layer capacitors allowing for structured electrodes composed of layers of different material properties; (2) The comparison of the model predictions with experimental electrical impedance spectroscopy (EIS) data; and (3) detailed interpretation of real and imaginary capacitance plots for layered electrodes. We conclude: (1) Layering introduces two knee frequencies in the real capacitance that can be manipulated advantageously; (2) Layering can increase the capacity for high frequency operation; (3) Layering decreases capacity at lower frequencies but may also reduce losses; and (4) Electrode response is surprisingly insensitive to some of the details of the layer configuration.

2. Electrochemical modelling of structured electrodes

The model develops the electrochemical equations of [13,36] to account for electrodes comprised of two discrete electrode sub-layers.

2.1. Electrochemical equations

The model’s spatio-temporal variables and parameters are given in Tables 1 and 2. The model equations are given in Table 3 and the boundary conditions are given in Table 4. The electrochemical assumptions of the model are outlined in Ref. [36] and include the
Table 1: Nomenclature of the structured electrode supercapacitor model.

| Spatio-Temporal Variables | Ionic concentration in the electrolyte. |
|---------------------------|----------------------------------------|
| $c_i(x,t)$                | Potential in the electrolyte.          |
| $\phi_i(x,t)$             | Potential in the electrolyte.          |
| $\phi_d(x,t)$             | Double layer potential difference.     |
| $i_i(x,t)$                | Electron current in the electrode.     |
| $i_s(x,t)$                | Ionic current in the electrolyte.      |
| $i(t)$                    | Applied current density.               |
| $v(t)$                    | Measured voltage.                      |

Parameters

- $\varepsilon$: Porosity coefficient.
- $D$: Effective Diffusion coefficient.
- $C^\text{ep}$: Specific capacitance.
- $\sigma$: Electrode conductivity.
- $\kappa$: Electrolyte conductivity.
- $R$: Universal gas constant.
- $F$: Faradays constant.
- $T$: Temperature.
- $F = \frac{RT}{\kappa}$: A coefficient.
- $K$: A coefficient.
- $\left(\frac{t_s - t_e}{t_e - t_s}\right)$: Transference numbers ($t_s + t_e = 1$).
- $\frac{dq}{dt} = \frac{1}{2}$: Change in $j = \{-1\}$ electrode surface concentration with charge $q$.
- $S$: Electrode surface area.
- $i(t) - I(t)/S$ where $i(t)$ is current.
- $j$: Parameter subscript for the $j^{th}$ layer.

Table 2: Nomenclature for the lengths and positions of the electrodes.

| Length Notation | Description |
|-----------------|-------------|
| Electrode length. | Length of the electrode closest to the current collector. |
| $L$             | Supercapacitor length $L = 2L_{\text{elec}} + L_{\text{sep}}$. |
| $L_{\text{elec}}$ | Length of the collector layer closest to the current collector. |
| $L_{\text{sep}}$ | Collector length. |
| $L_{es}$        | Electron separator length. |
| $L = L_{\text{elec}} + L_{\text{sep}}$ | \|

Table 3: Equations for the electrochemical supercapacitor model with structured electrodes. The equations for the $j^{th}$ layer are denoted by the subscript $j$.

Equations for Electrodes and Separators

- Nernst-Planck expression for the ionic flux and dilute solution theory to describe ion movement within the electrode pores. Pseudo-capacitance effects are ignored in the model, however, these effects are negligible for the high frequencies of interest to this paper ($\approx 100 \text{ Hz}$).

- Consider the supercapacitor cross-section in Fig. 2, where $x$ is the spatial co-ordinate traversing the domain from one current collector to the other. The dynamics of each electrode layer are described by the same electrochemical equations, but with each layer having distinct parameter values. The electrochemical equations are adopted from Ref. [36], with the subscript $j$ referring to the $j^{th}$ electrode layer. For instance, $\sigma_j$ is the electrode conductivity in the $j^{th}$ layer. In the electrodes, the electrochemical equations [36] are:

  \[
  \psi \frac{d\phi_j}{dx} + \frac{C_j^\text{ep}}{F} \left( t_a \frac{dq_{j+1}}{dx} + t_e \frac{dq_{j-1}}{dx} \right) \frac{d(\phi_j - \phi_s)}{dx} = D \frac{d^2 \phi_j}{dx^2} \tag{1a}
  \]

  \[
  C_j^\text{ep} \frac{d(\phi_j - \phi_s)}{dx} = \frac{d\phi_j}{dx} \tag{1b}
  \]

  \[
  \frac{d\phi_j}{dx} = -i - \frac{i_s}{\sigma_j} \tag{1c}
  \]

  \[
  \frac{d\phi_j}{dx} + \frac{t_s - t_e}{f} \frac{dln(\psi_j)}{dx} = \frac{i_s}{\kappa_j} \tag{1d}
  \]

  The first equation (1a) describes the diffusion of the electrolyte ions with a source term due to the double layer. The second equation (1b) accounts for the relaxation of the double layer. The third equation (1c) is Ohm’s law for the electrode and the fourth equation (1d) is the Nernst-Planck equation for the electrolyte. These four equations can be simplified by substituting for $i_i(x,t)$:

  \[
  \psi \frac{d\phi_j}{dx} + \frac{C_j^\text{ep}}{F} \left( t_a \frac{dq_{j+1}}{dx} + t_e \frac{dq_{j-1}}{dx} \right) \frac{d(\phi_j - \phi_s)}{dx} = D \frac{d^2 \phi_j}{dx^2} \tag{2a}
  \]

  \[
  C_j^\text{ep} \frac{d(\phi_j - \phi_s)}{dx} = \theta_j \frac{d^2 \phi_j}{dx^2} \tag{2b}
  \]

  \[
  \sigma_j \frac{d^2 \phi_j}{dx^2} + \frac{\kappa_j}{\sigma_j + \kappa_j} \frac{d\phi_j}{dx} + \frac{t_s - t_e}{f} \frac{dln(\psi_j)}{dx} + i = 0. \tag{2c}
  \]

  But, the algebraic constraint of (2c) complicates any analysis of a model synthesized from this set of equations, making the layered electrode design problem intractable. To eliminate this constraint, a further substitution is introduced by adding and subtracting $\kappa_j \partial \phi_j / \partial x$ to (2c) giving:

  \[
  \frac{d\phi_j}{dx} + \frac{\kappa_j}{\sigma_j + \kappa_j} \frac{d\phi_j}{dx} = \frac{\kappa_j}{\sigma_j + \kappa_j} \frac{t_s - t_e}{f} \frac{dln(\psi_j)}{dx} + i = 0, \tag{3}
  \]

  which can be differentiated in $x$:

  \[
  \sigma_j \frac{d^2 \psi_j}{dx^2} = \frac{\sigma_j k_j}{\sigma_j + \kappa_j} \frac{d^2 (\phi_j - \phi_s)}{dx^2} - \frac{\sigma_j}{\sigma_j + \kappa_j} \frac{t_s - t_e}{f} \frac{d^2 ln(\psi_j)}{dx^2}. \tag{4}
  \]
Substituting (4) back into the RHS of (2b) puts the electrochemical supercapacitor model into the state-space form

$$\frac{d}{dt} \psi_c = \frac{c_e}{\varepsilon_c} \left( \frac{d \psi_c}{dt} + \frac{d \psi_e}{dt} \right) \frac{d(\psi_e - \psi_c)}{dt} = D_e \frac{d^2 \psi_c}{dx^2} \quad (5a)$$

$$\frac{d \psi_e}{dt} = \frac{\sigma_i \psi_e}{C_s^{\sigma} (\sigma_s + \kappa_s)} \frac{d^2 \psi_e}{dx^2} - \frac{\sigma_i}{C_s^{\sigma} (\sigma_s + \kappa_s)} \left( t_e - t_s \right) \frac{d^2 (\ln(c_e))}{dx^2} \quad (5b)$$

This set of equations describes the same set of dynamics as [36], but does not contain algebraic equations. It has a state-space of \((c_e, \psi_e)\) and can be readily analysed and simulated [13]. In the separator, there is only one dynamic equation, that for electrolyte diffusion

$$\frac{d c_s}{dx} = D_s \frac{d^2 c_s}{dx^2} \quad (6)$$

with the subscript \(s\) indicating the separator, since the double layer does not exist in the separator as it is an electrical insulator.

### 2.2. Boundary conditions

The model boundary conditions are also inferred from Ref. [36] and are given in Table 4, with the exception of those at the electrode sub-layer to sub-layer interface, which are introduced here. The boundary conditions on the electrolyte ionic concentration \(c_e\) ensure that the supercapacitor is a closed system, with no ions leaving “through” the current collector and a constant ionic flux is imposed at the separator and across the electrode layer interfaces. The boundary conditions on the double layer potential \(\psi_e\) were inferred from those on the potentials in the solid (active and electronic conductive materials) electrode \(\psi_s\) and those in the electrolyte \(\psi_e\). The boundary conditions on the solid electrode potential \(\psi_e\) enforced all the current to be flowing through the solid phase at the current collectors (with \(i_L = 0\)) and in the liquid phase at the separator (\(i_L = 0\)). Across the electrode sub-layers, it was assumed that current was conserved i.e. \(i_{L, \text{left}} = i_{L, \text{right}}\), where Left and Right are respectively subscripts denoting currents flowing from the left and right across the separator layer. Likewise, the boundary conditions on \(\psi_e\) ensured that the current in the electrolyte was zero at the current collectors (\(i_L = 0\)), equalled the full current at the separator (\(i_L = 0\)) and was conserved across the electrode layers \(i_{L, \text{left}} = i_{L, \text{right}}\). These inter-layer boundary conditions contrast with those from Ref. [29] where no current conservation was imposed.

### 2.3. Voltage

The voltage \(v\) of the supercapacitor is defined as the solid phase potential difference between the two current collectors

$$v(t) = \psi_s(0, t) - \psi_e(L, t). \quad (7)$$

But, this expression is not a function of the state-space \((c_e, \psi_e)\). To obtain such an expression, the voltage equation (7) was first written as the sum of the potential differences across each layer

$$v(t) = -\sum_{i=0}^{L} \psi_s(L_{i+1}, t) - \psi_e(L_{i}, t). \quad (8)$$

The positions \(L_i\) correspond to the layer interfaces and are defined in Table 2 with \(L_0, L_L\) at the current collectors, \(L_1, L_4\) at the layer interface positions and \(L_2, L_3\) at the end points of the separator. This expression assumes that the sub-lowering configuration in each electrode is mirrored but could be easily modified if this was not the case. Equation (8) can be represented as a path integral of the potential gradient

$$v(t) = -\sum_{i=0}^{L} \int_{L_i}^{L_{i+1}} \frac{d \psi_s(x, t)}{dx} dx. \quad (9)$$

Substituting the expression for \(d \psi_e(x, t)/dx\) from (3) into this summation, gives

$$v(t) = \sum_{i=0}^{L} \int_{L_i}^{L_{i+1}} \frac{k_i}{(\sigma_i + \kappa_i)} \frac{d \psi_e(x, t)}{dx} + \frac{k_j}{(\sigma_j + \kappa_j)} \left( t_e - t_s \right) \frac{d \ln(c_e(x, t))}{dx} + i \frac{\sigma_i}{\sigma_j + \kappa_j} \int_{L_i}^{L_{i+1}} \frac{d \ln(c_e(x, t))}{dx}$$

$$+ i \frac{\sigma_i}{\sigma_j + \kappa_j} \int_{L_i}^{L_{i+1}} \frac{d \ln(c_e(x, t))}{dx} \quad (10)$$

which can be evaluated exactly

$$v(t) = \sum_{i=0}^{L} \left[ -\frac{k_e}{(\sigma_e + \kappa_e)} \frac{d \psi_e(x, t)}{dx} + \frac{k_j}{(\sigma_j + \kappa_j)} \left( t_e - t_s \right) \ln(c_e(x, t)) \right] + i \frac{\sigma_i}{\sigma_j + \kappa_j} \int_{L_i}^{L_{i+1}} \frac{d \ln(c_e(x, t))}{dx} \quad (11)$$

to give the voltage as a function of the states \((\psi_e, c_e)\).

### 3. Parameter estimation of the electrochemical model

Accurate parameter estimates were required for the model simulations to represent experimental data and to give predictive power. For parameter identification and simulation purposes, the PDE equations in (5) were first discretised in space using the spectral collection method of [13], with the discretised model parameters then identified using the procedure outlined in Ref. [14].  

The parameter estimation problem can be cast as the solution to the least squares optimisation

$$\theta^* = \min_\theta ||\hat{Z}(\theta) - Z||_2^2 \quad (12)$$

where \(\theta = \{ \kappa_{j}, C_{j}, \sigma_{j}, L_{\text{elec}}, L_{\text{dep}}, L_{\text{lay}}, S_{\text{t}}, \varepsilon_{j}, D_{j} \}\) are the models parameters, \(\theta^*\) is the optimal parameter choice that minimises the distance between simulation and data, \(\hat{Z}(\theta)\) is the simulated response and \(Z\) is a set of data points of bi-layered supercapacitors synthesized with the method outlined in Section II-A.

In this paper, electrical impedance spectroscopy (EIS) data is used for the identification, with EIS giving the frequency domain mapping from current \(I(j\omega)\) to voltage \(V(j\omega)\) as

$$V(j\omega) = G(j\omega)I(j\omega). \quad (13)$$

A local impedance function for the electrochemical model was obtained by linearising the spatially discretised model equations around an equilibrium point and applying the Laplace transform. EIS data was used for the identification because in comparison with time domain constant-current/constant voltage data, it gives a more complete characterisation of supercapacitor dynamics. Solving (12) with \(\hat{Z}(\theta)\) as the impedance function gave a reasonable match to experimental data, with the estimated parameters given in Tables 5 and 6.

### 3.1. Experimental set-up

The electrochemical model parameters were identified from experimental data of bi-layered supercapacitors. The bi-layer electrodes were synthesized using a scalable suspension atomization and automatic spray deposition processing technique developed previously [20,37]. Two types of aqueous suspension were used, one containing TiO2 nanoparticles and another containing carbon nanotubes (CNTs), were prepared separately through sonication. The choice of CNTs and TiO2 as the electrode materials was based on their known supercapacitor behaviour and good performance from previous experimental studies [19,21]. Besides having distinctly different material properties that may make layering effects identifiable, the two materials are also interesting from a manufacturing point of view, since the TiO2 is significantly
composed primarily of TiO₂ formed in a cross flow of air in a fume cupboard. To spray an area up to 20 cm to minimize. The results presented here then represent a means to cheaper than CNTs, so in general the CNT fraction in an electrode should be minimised. The suspensions were pumped into separate nozzles, atomized into micro-droplets by compressed air, and consecutively sprayed to form two layers of the electrode on a current collector maintained at 100 °C on a heated vacuum stage-chuck. Any fugitive water in the spray suspensions evaporated continuously on the heated stage as the electrodes formed. The nozzles moved in a pre-programmed zig-zag pattern along X and Y directions at 20 mm s⁻¹ to spray an area up to 20 cm × 20 cm. The spray processing was performed in a cross flow of air in a fume cupboard.

The electrode layer thicknesses were chosen as given in setup 1 in Table 5, with the electrode sub-layer closest to the current collector composed of CNTs while the sub-layer closest to the separator was composed primarily of TiO₂ particulate. The electrode mass loading was 2.1–2.6 mg cm⁻², the electrolyte was 1 M H₂SO₄ and stainless-steel foil (type 304, Zapp Precision Metals) current collectors were used. Sulphuric acid was chosen as the electrolyte primarily because it is one of the most common aqueous protic electrolytes for supercapacitors, with relatively high ionic conductivity of 0.35 S cm⁻¹ at 1 M [7]. EIS was conducted on a Reference 600/EIS300 Gamry potentiostat/galvanostat operated within the 2 × 10⁻³ - 10⁶ Hz frequency range with a ± 5 mV rms AC voltage, operated at open circuit voltage at room temperature.

### 3.2. EIS simulations: comparison to data

A comparison between the impedance functions of the electrochemical model and the experimental data under ideal conditions is shown in Fig. 4. Here, Fig. 4a plots the Argand diagram of the impedance function of the bi-layered electrodes while Fig. 4b and c respectively show the corresponding real and imaginary parts as a function of frequency. There was a tolerably good fit between the data and simulation across the frequency range 10⁻² – 10² rad s⁻¹. To show that the identification avoided over-fitting, the model was validated in Fig. 8 against data with the position of the sub-layers within the electrode swapped around, showing that the model was able to maintain a good fit to experimental data.

### 3.3. Equivalent circuit synthesis

In supercapacitor modelling, it is often desirable to extract an equivalent circuit from the impedance function to characterise the response. Such circuits are now synthesized directly from the impedance function of the electrochemical model G(ω) using the method from Ref. [15]. In particular, circuits with the structure of Fig. 3a are generated for different RC pair numbers n_RC. Note that since the impedance of a diffusing system is transcendental, fractional capacitances are often used in these circuits [4], although fractional components are not realisable from the discretised electrochemical model. Solving the coupled linearised electrochemical PDEs directly for the analytic impedance function would enable these fractional capacitances to be computed, but this is left to future work.

The considered circuits admit the state-space realisation

\[
\begin{bmatrix}
    \dot{x}_c \\
    \dot{x}_1 \\
    \vdots \\
    \dot{x}_n
\end{bmatrix} =
\begin{bmatrix}
    0 \\
    -1/R_1/C_1 \\
    \vdots \\
    -1/R_nC_n
\end{bmatrix}
\begin{bmatrix}
    x_c \\
    x_1 \\
    \vdots \\
    x_n
\end{bmatrix} + \begin{bmatrix}
    1/C_1 \\
    1/C_1 \\
    \vdots \\
    1/C_n
\end{bmatrix} i(t),
\]

(a) Equivalent circuit with two RC pairs n_RC = 2.
Fig. 4. Experimental and simulated impedance function plots for a supercapacitor with structured bi-layer electrodes in the set-up 1 configuration, with a layer of carbon nanotubes (CNTs) close to the current collector and a layer of TiO$_2$ particulate close to the separator.

(a) Argand diagram of $G(j\omega)$.

(b) Real component of $G(j\omega)$ as a function of frequency.

(c) Imaginary component of $G(j\omega)$ as a function of frequency.
to generate an impedance

\[ G_{\text{imp}}(j\omega) = \frac{1}{C_{j\omega}} + R + \sum_{j=1}^{n} \frac{1}{C_j(j\omega + R)} \]  

(15)

To extract the circuit parameters, balanced truncation [15] is applied to \( G(s) \) and the resulting state-space system is then transformed into (14). In this manner, the circuit components are directly linked to the electrochemical parameters, for example the porosity and electrolyte conductivity. The circuit of Fig. 3a with \( n_{\text{RC}} = 2 \) can then be directly synthesized, with \( C = 0.0573 \ \text{F}, \ C_1 = 0.0138 \ \text{F}, \ C_2 = 0.0277 \ \text{F}, \ R_1 = 213.924 \ \Omega, \ R_2 = 4.740 \times 10^3 \ \Omega \) and \( R_0 = 3.801 \ \Omega \).

To evaluate the diminishing importance of the number of RC pairs to the circuit accuracy, the following error metric is introduced:

\[ \| \frac{G_{\text{circ}}}{G_{\text{imp}}} \|_2 = \frac{1}{n_{\text{circ}}} \left( \| \Re(G(j\omega)) - G_{\text{imp}}(j\omega) \|_2 + \| \Im(G(j\omega)) - G_{\text{imp}}(j\omega) \|_2 \right) \]  

(16)

In essence, this metric measures the average error between \( G_{\text{circ}} \) (for a given number of RC pairs \( n_{\text{circ}} \)) and \( G \) evaluated over the frequency points \( \omega \in \mathbb{R}^+ \), with \( n_{\text{circ}} \) being the number of evaluated frequencies. Fig. 3b plots the error \( \| \frac{G_{\text{circ}}}{G_{\text{imp}}} \|_2 \) as a function of the number of circuit RC pairs \( n_{\text{RC}} \), showing that low order circuits gave sufficient accuracy.

4. Real and imaginary capacitance

This section investigates how the introduction of the sub-layers within the electrodes affected the real and imaginary capacitance [24, 35]. These capacitances are often used as performance metrics for supercapacitors and are obtained from the EIS response by decomposing the impedance function into real and imaginary parts.

\[ G(j\omega) = C'(\omega) + jC''(\omega) \]  

(17)

Primes are used to denote real components and double primes denote imaginary components. This function can be represented as a supercapacitance \( C(\omega) \) such that

\[ G(\omega) = \frac{1}{j\omega C(\omega)} \]  

(18)

The reason for expressing \( G(j\omega) \) in this manner is to obtain a frequency domain form of the capacitance equation \( Q = Cv \), where \( Q \) is the stored charge and \( C \) is the device capacitance. Through some rearranging [35], it follows that the supercapacitance has the form

\[ C(\omega) = C'(\omega) - jC''(\omega) \]  

(19)

where \( C'(\omega) \) is the real capacitance

\[ C'(\omega) = \frac{G''(\omega)}{\omega |G(\omega)|^2} \]  

(20)

and \( C''(\omega) \) is the imaginary capacitance

\[ C'(\omega) = \frac{G'(\omega)}{\omega |G(\omega)|^2}. \]  

(21)

According to Ref. [35], the real part \( C'(\omega) \) is effectively the capacitance while the imaginary capacitance \( C''(\omega) \) corresponds to energy.
dissipated by irreversible losses. Optimising over these capacitances is often said to be equivalent to optimising over the Pareto front of the Ragone plot [9].

4.1. Real & imaginary capacitance: comparison to data

Fig. 5 shows simulated real and imaginary capacitance as a function of frequency for various supercapacitors. For these simulations, set-up 1 had a 32 μm thick sub-layer closest to the current collector composed primarily of TiO$_2$ particulate and a 5 μm thick sub-layer closest to the separator primarily comprising of CNT. Set-up 2 had the position of the sub-layers swapped, with the 5 μm thick CNT sub-layer closest to the current collector and the 32 μm thick TiO$_2$ sub-layer closest to the separator. Experimental data for set-up 1 is shown in black and
corresponding simulation results are shown in blue. Equivalent supercapacitors with electrodes comprising only of TiO$_2$ and only of CNT are shown in red.

The most noticeable feature in the capacitances plots of the structured bi-layer electrodes in Fig. 5 was the presence of two peaks in the imaginary capacitance plots and the corresponding two knee frequencies in the real capacitance plots. While there was only a single peak in the TiO$_2$-only and CNT-only electrodes. The additional peak is however consistent with the recent result in Ref. [10] which also mapped electrochemical equations to EIS results, but for electrodes with a linearly varying porosity fraction through the electrode thickness. The authors in Ref. [10] warned that such peaks may be misinterpreted as additional electrochemical effects [6], but are actually diffusion based, and the results presented here reinforce this conclusion. We verify the presence of two peaks in the case of sub-layered electrodes which are often associated to “constant phase elements” [35], with a physical explanation for such elements remaining a topic of dispute [1]: explanations include surface roughness, distributions of reaction rates and non-uniform current distributions within the electrode. The simulations and experiments here suggest that the presence of the peaks was due to the coupled diffusion between/within the sub-layers.

Fig. 5 also shows the impact on the frequency dependent capacitance caused by layering. Focusing first in the high frequency region $10^0 \leq \omega \leq 10^2$ shown in more detail in Fig. 6, layering increased the capacity in this region. This was due to the two knee frequencies of the sub-layered electrodes and corresponding to the frequency region where diffusion limitations in the electrolyte dominate performance. The presence of two knee frequencies was most likely due to two different effective diffusion coefficients from the electrode sub-layers and their coupling via the boundary conditions at the internal sub-layer interface. This gives encouragement for the use of structured electrodes in applications with high frequency charging. However, the double knee profile also resulted in a drop in capacity in the frequency region $10^{-3} \leq \omega \leq 10^0$, albeit with lower losses. Finally, for DC charging at $\omega \approx 0$, where diffusion effects are insignificant, the capacity of the layered electrode sensibly was an average of the two single layer cases. This suggests that sandwiching relatively cheap TiO$_2$ based electrodes with a smaller fraction of more expensive high performance CNTs could increase capacity at DC over a TiO$_2$ electrode and provide additional capacity in the high frequency region.

This high frequency capability of the structured or bi-layer electrodes explained in terms of increased ion mobility in Fig. 7. This figure shows simulated ionic concentration distributions in space and time for both a supercapacitor with bi-layer TiO$_2$/CNT-structured electrodes and a supercapacitor with CNT-only electrodes. Relatively thick electrode sub-layers of 100 $\mu$ m were used for these simulations to draw out differences more clearly, with single material electrodes of 200 $\mu$ m thickness and a 20 $\mu$ m thick separator in all cases. The bi-layered supercapacitors were charged by a sinusoidal current of amplitude $10^{-3}$ A at frequencies of $5 \times 10^{-2}$ rad s$^{-1}$, $5 \times 10^{-1}$ rad s$^{-1}$ and 5 rad s$^{-1}$.

Fig. 7 illustrates the increased ion mobility in the bi-layered electrodes seen in the model simulations. At low charging frequencies, the response of the bi-layered electrodes was similar to that of the CNT-only electrodes, with the electrolyte ions diffusing across the whole electrode. But, as the charging frequency was increased, the effect of the sub-layers became more pronounced. The layering created a region around the separator where ion mobility was amplified and reduced the effective length of the layered electrode to that of the inner layer. This thin pseudo-electrode explained the improved capability at high frequencies. Also, since the voltage expression (11) contained contributions from the electrolyte concentration $c_e$ at each of the layer interfaces, this increased ion mobility affected the voltage response. These simulations explain the double knee frequencies (due to the two diffusion time constants from the two sub-layers) and the increased performance at high charging frequencies (due to the increased ion mobility across the supercapacitor) in the experimental impedance plots. It is conjectured that this effect can be extrapolated to n sub-layers where the effect may be amplified further.

The model framework described here has sufficient flexibility that it can be used to guide designs for improved functionality, through the considerably widened supercapacitor design space facilitated by structured electrodes and the corresponding manufacturing technologies. Critically, changes in model parameters can be directly linked to changes in energy storage performance. Fig. 8a and b explore this idea by showing device capacitances as a function of the ratios of the electrode sub-layer thicknesses. Noticeably, for all sub-layer thickness ratios, the capacitance plots were qualitatively similar, with the most significant difference in capacitance being when the layers were the same thickness but had their position swapped, as was shown earlier in Fig. 5a and b. This relative insensitivity to the sub-layer thickness ratios, seen in both the model simulations and the experimental data, implies that there is no distinct optimal thickness ratio, although this conclusion may not hold for different electrode configurations, especially as electrode thicknesses increase.
5. Conclusions
A model describing ionic diffusion and electric double layer capacitance dynamics was developed, tuned and validated against EIS data, with the tuned model parameters obtained from the solution of a least mean squares problem. Real and imaginary capacitances for the bilayered structured electrodes contained two knee frequencies, which could be manipulated by changes in the electrode design to increase the capacitance at high frequencies. Model simulation results showed that these increased capacitances derived from increased ion mobility facilitated by sub-layering and the nature of the resulting coupled diffusion between sub-layers.

Extensions of this work will focus on Li-ion battery electrodes, where the design flexibility of the layering could be exploited to promote reaction rate (insertion/de-insertion of Li-ions) homogeneity across the thickness of the electrodes. As well as increasing the capacity during fast charging, homogenisation of electrode charge distribution may reduce capacity degradation.

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