Asymmetric Supercapacitors Based on Transition Metal Oxides Induced Pseudocapacitance

Mak Wai Fatt

School of Materials Science & Engineering

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Abstract

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Abstract

Pseudocapacitor exhibiting higher capacitance and energy density than electrochemical double layer capacitor (EDLC) and delivering greater power density than batteries represent unique charge storage devices that bridge the gap of EDLC and rechargeable battery. In particular, RuO$_2$ based material has received much attention in past decades. However, they are expensive due to their scarce resources. This study reports on alternative pseudocapacitive material such as iron oxide as an electrode material for supercapacitor (SCs) applications. Also, an asymmetrical device configuration was conceptualized to extend the working voltage so as to achieve higher energy density.

The role of nanofibers in electrical double layer formation, pseudocapacitive properties, charge retention and cycling performance has been investigated for various materials. Physical characterization techniques such as Field Emission Scanning Electron Microscopy, X-ray Diffraction, Nitrogen Adsorption Isotherm, and Fourier Transform Infra Red Spectroscopy were used to study the physical properties of various materials. Electrochemical characterization such as cyclic voltammetry, galvanostatic charge-discharge, and electrochemical impedance spectroscopy were employed to investigate the electrochemical performance of different materials. These studies provide insight into the effect of morphology, surface area, electrode-electrolyte interaction on charge storage and transport mechanisms in SCs. In addition, the effect of material
porosity on capacitance contribution was investigated. Higher content of micropores is found to increase the capacitance as more ions can be accommodated in these micropores.

Systematic study on the effect of electrolytes on supercapacitor performance was carried out, including correlation with the size of cations and comparison between aqueous and organic electrolytes and the morphological effect of the electrode material. The size of the ions also plays a substantial part in electrical double layer contribution. With smaller size, higher quantity of charges can be stored and subsequently utilized.

Reversible chemical reactions are also found to have overwhelming contribution to the charge storage via pseudocapacitance. Solvated sodium ions have additional pseudocapacitance which greatly contributes to the charge storage capability. Choice of electrode/electrolyte pair is therefore critical to for obtaining higher charge storage/capacitance in supercapacitors.

An asymmetrical device configuration was conceptualized to extend the working voltage so as to achieve higher energy density. To realize asymmetrical supercapacitor, different electrodes of nanofibers have been judiciously coupled. The effect of asymmetrical supercapacitor architecture on the voltage window has been investigated. The performance of the devices was then evaluated using Ragone plot. Our studies has shown that such asymmetrical supercapacitors proposed increases the energy density of supercapacitors and would better bridge the gap of EDLC and rechargeable batteries.
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Chapter 1 Introduction

1.1 Energy Storage Systems

Escalating oil prices has generated a growing global concern over the state of world energy consumption and supply. There is increased interest in further developing technology utilizing electrical devices as sources of energy supply, especially in the area of transportation. Alternative renewable energy sources is being heavily researched, especially the field of photovoltaic and fuel cell technology. Hybrid Electric Vehicles (HEV) and Fuel Cell Vehicles (FCV) are available for consumer from various manufacturers, albeit at a significant higher cost than the conventional Internal Combustion Engine (ICE). These HEVs and FCVs use a combination of technologies to cater to the energy requirement. The hybridization is attained with the integration of fuel cells, rechargeable batteries, and supercapacitors. The system design needs to ensure maximum energy output with minimal fuel consumption.

People still expect to drive around at times when the sun does not shine (rainy days and nights). Thus energy storage systems (ESS) play a crucial role in practical implementation of renewable clean energy in our daily lives. Batteries and electrochemical capacitors are at the forefront among electric ESS. However, the performance of present ESS needs to be improved to meet higher energy and power demand of future applications ranging from portable electronic to HEVs, FCVs and large industrial equipments.
Figure 1.1: Ragone plot of various electrical storage systems depicting the power and energy densities. For comparison sake, the envisioned asymmetric supercapacitor performance studied in this work is also inserted.

Pseudocapacitor, being a sub-class of supercapacitor represents a rising class of energy storage device capable of bridging the gap between rechargeable batteries and capacitors. Besides, pseudocapacitor possesses higher energy density and thus can be used to complement or replace batteries in some energy storage applications, such as for load leveling and uninterruptible power supplies.
1.2 Motivation of Project

Supercapacitors have been around for several decades with the first patent dated back to 1597 in which a high surface area carbon capacitor was described by Becker [2]. However, it was only until the nineties that supercapacitors became famous in the context of HEV. It is currently used to complement the batteries, owing to the latter higher costs and cumbersomeness. In the same way, using the supercapacitors for power delivery application of more than 10 seconds required over-sizing. Thus SC can only complement battery in electrical system like the HEVs. Such combination will allow fast acceleration and braking energy recovery, and extension of battery life.

There are fundamental differences between batteries and supercapacitors although both are based on electrochemistry. Lithium batteries store energy through intercalation of Li ion, whereas SC store energy in the form of electrical field. Others include cycle life, power density, energy density etc as discussed in literature [3, 4].

Supercapacitors plays important role in complementing batteries in Energy and Storage System ESS such as Uninterruptible Power Supply (UPS), load leveling and emergency doors (16 per plane) on Airbus 380. The 2007 report of US department of Energy assigns equal importance to SC and batteries for future ESS [5]. The capability of important ESS can be depicted in Ragone plot shown in Figure 1.1. These devices must be used in their respective time constant domains. Using Li ion battery for high power application will degrade cycle life drastically.
Introduction

The key to high power supercapacitors lies in using a high specific surface area (SSA) blocking and electronically conducting electrode. High SSA carbon satisfies the requirement for this application. Templated, activated carbon and carbide derived carbons [6], fiber, nanotubes [7] have been investigated for EDLC applications. However, their low energy density often limits their wide application.

Another type of supercapacitors called pseudo capacitor having a higher capacitance utilizes fast, reversible redox reactions at the surface of active materials to achieve greater energy storage. Transitional metal oxides such as RuO2, MnO2, NiO, Co3O4 [8] as well as conducting polymer [9] have been extensively researched as a potential electrode candidate in last decade. These pseudocapacitors have exhibited higher capacitance than that of carbon based EDLC.

Currently, two different approaches are being explored to increase the energy density of supercapacitors. One is the combination of pseudo capacitive metal oxide with a capacitive carbon electrode. The other is the combination of Li-ion intercalation electrodes with a capacitive carbon electrode. Although both can attain greater energy storage, they suffer from low cycle life due to the degradation of the faradic and lithium intercalating electrode. Thus the selection of electrode pair (balance positive and negative electrodes’ capacities) is crucial so as to avoid transforming a high performance supercapacitors into a mediocre battery.

Transition metal oxide (TMO) capacitor has shown high capacitance, however, they also suffer from low electronic conductivity. Often, TMO are co-precipitated or
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electrodeposited onto a metal substrate. However, the complexity and time involved undermine the possibility of up-scaling the fabrication process, thus limit their real life applications. Currently, effort is lacking in addressing this issue. In order to fill this technology gap, this research will focus on the incorporating TMO nanostructure with electrically conducting material.

In addition, fundamental gaps are still present in understanding atomic and molecular level process that governs pseudocapacitor operation, limitation and failure. Fundamental research is required to uncover the underlying principles that govern the complex process. With a better understanding of the interrelated process, new devices can then be fabricated to address current ESS technology gap and fulfilling future energy storage requirements.
Introduction

1.3 Research Objectives and Scope

1. Development of electrically conducting pseudo capacitive composites:
   - Integrating binder, conductivity enhancer and active material in a single component.

2. Study of parameters that affect device performance
   - Varying the content of active nanomaterial.
   - Studying the effect of electrolytes.
   - Investigating the viability of asymmetrical configuration.

3. Investigations of nanomaterial contribution towards device performance
   - Research on charge storing mechanism of device through cyclicvoltametry and galvanostatic charge-discharge curve.
   - Investigate internal resistance of device through electrochemical impedance spectroscopy.
   - Correlate charging/discharging mechanism with morphology, electrode materials.
Chapter 2 Literature Review

2.1 Overview of Electrochemical Capacitor

Electrochemical capacitors also known as supercapacitor, are generally classified into electrochemical double layer capacitors (EDLCs) and pseudocapacitors based on their charge storage mechanism. The first invention was reported in 1957 when Becker [2] filed a patent describing a concept of an electrochemical capacitor utilizing high surface area carbon coated on current collector immersed in sulphuric acid. In 1969, researchers at Standard Oil of Ohio (SOHIO) accidentally rediscovered it while working on experimental fuel cell designs, created a prototype of modern version of electrochemical capacitor. Their cell design which is basically made up of two layers of activated carbon separated by a thin porous insulator still remains as the basis of today’s EDLCs. However, SOHIO did not commercialized their invention, only licensed it to Nippon Electrical Company (NEC), which subsequently initiated the commercialization of the first electrochemical capacitor in 1978. Nevertheless, its energy density still remained low for commercial applications until mid 1990s with the up rise of nanomaterial. Various advances in material science together with refinement of existing systems have drastically improved the performance of EDLC with an equally rapid reduction in cost. Subsequently, EDLC was integrated into commercial plane (Airbus 380) to power emergency evacuation slides in 2005. Nonetheless, the energy density still remains low for application in most electronic gadgets.
Literature Review

In this chapter, the working principle and architecture of capacitor are reviewed. Current theoretical models are briefly reviewed. The various classes of electrode materials including carbon, electrically conducting polymers and transitional metal oxides based are discussed. Their application as electrode material for supercapacitor is reported. Their advantages and limitation are also examined.
2.2 Fundamentals and Applications of Electrochemical Capacitor

2.2.1 Types of Electrochemical Capacitor

As aforementioned, supercapacitor is a charge storage device which can be classified into two categories depending on the charge storage mechanism. As shown in Figure 2.1, a supercapacitor consists of two electrodes, a separator that electrically insulates the two electrodes, and an electrolyte. It differs from the conventional capacitor as charges are electrically stored on the electrode/electrolyte interface. This greatly reduced the distance between the charges, and hence tremendously enhances the capacitance of the electrode according to equation 2.1 which will be discussed in following section.
In general, supercapacitor can store their charges via two types of mechanisms:

(a) Electrical double layer capacitor, where energy is stored in the double layer formed under external potential source. The process is purely electrostatic which involves no faradic charge transfers i.e. No electron transfer across the electrode and electrolyte interface.

(b) Pseudocapacitor, where energy is stored through the fast reversible redox reactions occurring at the electrode surface. Such process involves faradic charge transfers in which electrons are exchanged.

Pseudocapacitor often commands higher capacitance but they also have drawback such as low chemical cyclic activity, higher resistance and lower power density. Their capacitance can be viewed as the superposition of non-faradic double layer capacitance ($C_{dl}$) and faradic pseudocapacitance ($C_{pc}$). The first element ($C_{dl}$) is determined by the total accessible surface area while the latter ($C_{pc}$) is regulated by facile reversible redox reactions.
2.2.1.1 Electrical Double Layer Capacitor (EDLC)

When an electronically conducting electrode is placed in an ionic conducting electrolyte, layer of charges are accumulated on both sides of the solid/electrolyte interface. This parallel layer of charges is termed as double layer. There is no charge transfer occurring across the double layer, and such electrodes are generally referred to as ideally polarizable electrode (non-faradic)[10].

The fundamental concept of EDLCs revolves around 2 ideally polarizable electrodes immersed in an electrolyte. It utilizes double layer capacitance that arises at the electrodes and electrolyte interfaces. A schematic of the electric double layer is shown in Fig 2.1. Charging up the electrode results in double layer formation and the energy is stored in the electrical field instituted by these double layers on both electrodes. Since no electrochemical reaction is involved, the reversibility and cycle life of EDLC is much better than in rechargeable batteries.

2.2.1.1.1 Theoretical Models of Double Layer

2.2.1.1.1 Helmholtz Double Layer

Double layer was first coined by Helmholtz in 1887.[11] He proposed a model with two layers of charges; one resides on the metal surface, and the counter charge in solution at the solid/electrolyte interface. These two sheets of charges of opposite polarity are separated by a distance of molecular order. Figure 2.1 illustrates the Helmholtz model. The capacitance of electrical double layer is given by equation 2.1.
Where $\varepsilon_0$ is the permittivity of free space having a constant value ($8.854 \times 10^{-12}$ J$^{-1}$ m$^{-1}$), $\varepsilon_r$ is the dielectric constant of electrolyte, $A$ the surface area of electrode, and $d$ distance between the parallel plates of charges.

This model predicts that the capacitance is independence of applied voltage and concentration of electrolyte. Thus, it fails to explain the variation of capacitance with potential and concentration of electrolyte at low electrolyte concentration ($<0.1$ mol dm$^{-3}$). Nonetheless, the Helmholtz model seems to be appropriate for polarizable electrodes in sufficiently high concentration of electrolyte ($>1$ mol dm$^{-3}$).

### 2.2.1.3.2 Gouy-Chapman Model

Between 1910-1913, Gouy and Chapman proposed a diffuse double layer model which take into account the dependency of capacitance on both potential and electrolyte concentration.[11] They suggested that the random thermal motion will prevent the counter ions from accumulating on the surface of electrode. In this model, the ions are viewed as point charges contained within the diffuse layer. Taking into account both Poisson and Boltzmann equations, the capacitance can be given by equation 2.2 [11]:

$$C = \frac{\varepsilon_0 \varepsilon A}{d}$$

$$C = \left(\frac{2\varepsilon \varepsilon_r \varepsilon^2 \varepsilon_0}{kT}\right)^\frac{1}{2} \cosh\left(\frac{z e\varphi_m}{2kT}\right)$$

12
Where \( z \) is the magnitude of ionic charge, \( c_0 \) is the concentration of ions in bulk solution, \( k \) is the Boltzmann constant, \( T \) is the absolute temperature, and \( \varphi_m \) is the potential at the metal surface.

The Gouy-Chapman model has addressed the drawback of Helmholtz model, recognizing the dependence of capacitance on both the potential and concentration. However, this model is only valid for very dilute solutions (<0.001 mol dm\(^{-3}\)) and at potential close to point of zero charge.[11] Nevertheless, the model has contributed to a better understanding of the double layer, and find it uses in understanding the stability of colloids.

2.2.1.1.3 Stern Model

Stern subsequently combined these two aforementioned models (Helmholtz and Gouy-Chapman) in 1924, and proposed that some ions are adhered to the electrode, while some are scattered in the diffuse layer due to thermal disarray.[11] This model take into account the fact that ions have a finite size, and consequently do not come closer to the electrode than the order of ionic radius. It also assumed two region of charge separation. One part of solution charges is immobilized in the compact layer, and the rest is dispersed in the diffuse layer as shown in Figure 2.2.

In this model, the double layer is effectively made up of two capacitances in series, \( C_H \) due to the Helmholtz layer and \( C_G \) due to the diffuse layer. The total capacitance is therefore given by equation 2.3,
In very dilute solution and close to the point of zero charge, $C_G$ becomes very small according to equation 2.3; with $C_H$ remains constant, overall capacitance can be approximated to $C_G$. Correspondingly, overall capacitance can be approximated to $C_H$ in concentrated solutions. However, Stern model inapplicable in certain solutions where ions (e.g. Cl, I, Br) come close or being adsorbed onto the electrode.

2.2.1.1.4 Grahame Model

In 1947, Grahame proposed that some ionic or neutral species can be adsorbed onto the electrode although the closest approach to the electrode is occupied by solvent molecules.[11] This could come from de-solvated ions or ions which do not possess salvation shell. Such ions which adsorbed to the electrode are termed specifically adsorbed ions.[11] Grahame suggested the existence of three regions, namely, inner Helmholtz plane (IHP), outer Helmholtz plane (OHP) and diffuse layer as shown in Figure 2.3.

2.2.1.1.5 Bockris, Devananthan and Muller Model

In 1963, Bockris, Devananthan and Muller proposed a model that considered the interaction of polar solvent molecules with electrode.[11] They suggested that there would be interaction between electrode and dipole of water molecules in dipole solvent, such as water. Since solvent molecules exist in overwhelming quantity, there would be a layer of solvent molecules adhered to the surface of electrode. The dipole of these solvent
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molecules would have a fixed orientation due to charge interaction with electrode. Occasionally, some solvent molecules would be displaced by specifically adsorbed ions. There will also be subsequent layers with orientation intermediate of the first layer and that in the bulk solution as shown in Figure 2.3.
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Conductor Electrolyte

Helmholtz model

Gouy-Chapman model

Stern model

Figure 2.2: Schematic depicting the Helmholtz model, Guoy-Chapman model and Stern model.
Figure 2.3: Schematic depicting the Grahame model and Bockris, Devanathan and Muller models
2.2.2 Pseudocapacitor

A different kind of capacitance called pseudocapacitance arises on electrodes when application of external potential induces faradic current. This is fundamentally different from the EDLC due to the way the charge is stored. In principal, pseudocapacitance can increase the effective capacitance of electrochemical capacitor by as much as two orders over that of EDLC. [12] Such pseudocapacitance is derived from reactions such as electro-sorption of atoms, electrochemical doping-dedoping, electrochemical redox charge transfers, and intercalation. [13, 14] Electrosorption occurs when ions such as $H^+$ are reversibly adhered to the surface of electrode such as platinum. Doping-dedoping process takes place in electronically conducting polymers like polyaniline, poly(3,4-ethylenedioxythiophene), polypyrrole, polythiophene etc. Redox charge transfers take places when electro-active species donates (or receives) electrons and gets oxidized (or reduced). Materials undergoing such redox reactions generally include transitional metal oxides such as RuO$_2$, MnO$_2$, V$_2$O$_5$, SnO$_2$, Fe$_3$O$_4$, Co$_3$O$_4$ and etc. Formally, lithium intercalation exhibits a pseudocapacitance though lithium ion systems are generally referred as battery devices also known as lithium ion battery. [13]


2.2.2.1 Pseudocapacitor Mechanism

2.2.2.1.1 Electronically Conducting Polymers

Electronically conducting polymer such as polyaniline, polypyrrole and polythiophene derivatives also exhibit pseudocapacitance. They are rendered conductive through the conjugated $\pi$ bond system along polymer backbone. Such polymers are attractive as they have good electrical conductivity from a few S cm$^{-1}$ to 500 S cm$^{-1}$ in the doped state, [15] coupled with their relatively fast charge-discharge kinetics and rapid doping-dedoping process. [16] Energy can be stored and retrieved as the delocalized $\pi$ electrons are accepted and released through electrochemical doping-dedoping process. Conducting polymers can be p-doped with anions when oxidized and n-doped when reduced. The general representations for the charging-discharging process are as follows:

$$C_p \leftrightarrow C_p^{n+} (A^-)_n + n e^- \quad (\text{forward, p-doping})$$

$$C_p + n e^- \leftrightarrow C_p^{n-} (C^+)_n \quad (\text{forward, n-doping})$$

Where $C_p$ represents the electronically conducting polymer, $A$ is the anions, and $C$ is the cations. The forward reaction represents the charging process, and vice versa.

Supercapacitor based solely on electronically conducting polymers can be subdivided into three categories, [17-21] namely, Type I, II and III as follow:

Type I: Having the same p-dopable or n-dopable polymer for both electrodes. (Symmetrical)
Type II: Having two different of either p-dopable or n-dopable polymer with different range of electro-activity. (Asymmetrical)

Type III: Having the same polymer for both electrodes with the p-doped form working as the positive electrode, and the n-doped form working as the negative electrode. (Symmetrical)

Thus far, Type III supercapacitor has been the most attractive. In this configuration, the charge is released at higher potential (up to 3V) as compared with both type I and II. [9] This higher operating voltage translates into both higher power and energy density. Supercapacitor based on electronically conducting polymers generally exhibit high capacitance value ranging from 90 to 530 F g⁻¹. [22] However, their performances are often undermined by difficulty of the n-doping process and their poor stability during cycling due to volumetric changes during doping-dedoping process. In addition, the electrochemical activity of conducting polymer is often restricted by its working potential range and low polymer degradation potential. [23]

2.2.2.1.2 Transitional Metal Oxides

Transition metal oxides are generally considered an attractive alternative as electrode material for as they exhibit high specific capacitance. They process a variety of oxidation states readily available for redox charge transfer, thus add on to the existing double layer storage.
2.2.3 Materials for Electrochemical Capacitor

2.2.3.1 Materials for EDLC

2.2.3.1.1 Activated Carbon

Amorphous carbon is the most widely researched electrode material, with activated carbon being the most common. Their reported specific capacitance is in the range of 30 to 100 F g\(^{-1}\).[24] Other carbonaceous material such as graphite powder, carbon black, carbon aerogel and carbon fibers have capacitance of 35, 10, 23 and 6.9 µF cm\(^{-2}\), respectively.[25] High electrical conductivity and specific surface area are the crucial factors determining the specific capacitance of EDLC.

2.2.3.1.2 Carbon Nanotubes

CNT being another form of carbon is also being invested intensively due to its high surface area to volume ratio, suitable mesoporous pore size, good chemical stability and excellent electronic conductivity.[26-28] The entanglement of nanotubes has enabled better ions diffusion to the active surface sites, and percolation of active electrolytes is also made more efficient.[29] CNT also have great cycling performance as nanotubular architecture generally has a high resilience to volumetric changes during charge and discharge.

Untreated SWNTs generally have specific capacitance ranging from 50-100 F g\(^{-1}\).[30] Substantial works have also been done to increase the volumetric capacitance by using aligned CNTS as electrodes for EDLCs.[31, 32] The aligned nanotubes known as
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nanotubes “forest” has a finer dimension and more uniform distribution exhibited greater energy storage. However, the high cost and complexity of the fabrication process limits their commercial utilization presently.

2.2.3.1.3 Graphene

Graphene, a two dimensional single layer material comprises of an atom thick hexagonal carbon lattice with delocalized electron, is emerging as a promising candidate for supercapacitor. Its high theoretical surface area of 2630 m$^2$ g$^{-1}$ coupled with good electronic conductivity, excellent thermal conductivity, outstanding strength and reasonable chemical stability have rendered itself useful for charge transport and storage application.[33-36]

Although high specific capacitance of 200 F g$^{-1}$ has been reported for graphene in aqueous electrolyte, commonly cited values are much lower. The performance is strongly dependant on the synthesis routes, porosity and the surface functional groups.

2.2.3.2 Electrically conducting polymers

Electrically conducting polymers make good pseudocapacitive material for electrode of supercapacitors. They possess good electronic conductivity, have low cost, and adjustable redox activity through chemical modification.[37, 38] They are capable of pseudocapacitive storage through the redox process. Ions are transferred to the polymer
backbone when oxidation occurs and vice versa. The processes have good reversibility as the redox reactions do not alter the structures.[39]

ECPs can be either positively or negatively charged. The injected charges can be balanced by the insertion of ions in the polymer matrix. Electronic conductivity can thus be induced via such oxidation and reduction reactions which generate delocalized electrons. Such oxidation and reduction processes are termed as doping. Oxidation induced positively charged polymer are termed as p-doped, while reduction induced negatively charged polymer are termed as n-doped.

Table 2-1 depicts a comparison of capacitance and voltage windows of various electrically conducting polymers and composites in different electrolytes for supercapacitors.
Table 2-1: A comparison of capacitance and voltage windows of various electrically conducting polymers and composites in different electrolytes for supercapacitor.

<table>
<thead>
<tr>
<th>Material</th>
<th>Electrolyte</th>
<th>Working Voltage (V)</th>
<th>Specific Capacitance (F/g)</th>
<th>Scan Rate (mV s(^{-1})) or Current Rate (A g(^{-1}))</th>
<th>Reference Electrode</th>
<th>System</th>
<th>Cycles</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>PANI-20wt% MWNT</td>
<td>1M H(_2)SO(_4)</td>
<td>-0.8-0.4</td>
<td>670</td>
<td>2 mV s(^{-1})</td>
<td>Hg/Hg(_2)SO(_4)</td>
<td>3E</td>
<td>-</td>
<td>[40]</td>
</tr>
<tr>
<td>Ppy20wt% MWNT</td>
<td>1M H(_2)SO(_4)</td>
<td>-0.6-0.2</td>
<td>506</td>
<td>5 mV s(^{-1})</td>
<td>Hg/Hg(_2)SO(_4)</td>
<td>3E</td>
<td>-</td>
<td>[40]</td>
</tr>
<tr>
<td>PEDOT/Ppy (5:1)</td>
<td>1M KCl</td>
<td>-0.4-0.6</td>
<td>290</td>
<td>2 mV s(^{-1})</td>
<td>SCE</td>
<td>3E</td>
<td>1000</td>
<td>[41]</td>
</tr>
<tr>
<td>Ppy10%-Carbon</td>
<td>6M KOH</td>
<td>-1-0</td>
<td>235</td>
<td>1 mV s(^{-1})</td>
<td>Hg/HgO</td>
<td>3E</td>
<td>2000</td>
<td>[42]</td>
</tr>
<tr>
<td>Ppy21%-Carbon</td>
<td>6M KOH</td>
<td>-1-0</td>
<td>322</td>
<td>1 mV s(^{-1})</td>
<td>Hg/HgO</td>
<td>3E</td>
<td>2000</td>
<td>[42]</td>
</tr>
<tr>
<td>Ppy63%-Carbon</td>
<td>6M KOH</td>
<td>-1-0</td>
<td>390</td>
<td>1 mV s(^{-1})</td>
<td>Hg/HgO</td>
<td>3E</td>
<td>2000</td>
<td>[42]</td>
</tr>
<tr>
<td>Ppy35%-Carbon</td>
<td>6M KOH</td>
<td>-1-0</td>
<td>433</td>
<td>1 mV s(^{-1})</td>
<td>Hg/HgO</td>
<td>3E</td>
<td>2000</td>
<td>[42]</td>
</tr>
<tr>
<td>Ppy-MWNT</td>
<td>1M H(_2)SO(_4)</td>
<td>-0.4-0.6</td>
<td>427</td>
<td>5 mV s(^{-1})</td>
<td>Ag/AgCl</td>
<td>3E</td>
<td>-</td>
<td>[43]</td>
</tr>
<tr>
<td>Ppy-RuO(_2)</td>
<td>1M H(_2)SO(_4)</td>
<td>-0.2-0.7</td>
<td>302</td>
<td>0.5 mA/cm(^2)</td>
<td>Hg/HgO</td>
<td>3E</td>
<td>-</td>
<td>[44]</td>
</tr>
<tr>
<td>Ppy</td>
<td>0.5M H(_2)SO(_4)</td>
<td>-0.5-0.5</td>
<td>250</td>
<td>50 mV s(^{-1})</td>
<td>Ag/AgCl</td>
<td>3E</td>
<td>5000</td>
<td>[45]</td>
</tr>
<tr>
<td>Ppy-MnO(_2)</td>
<td>0.5M H(_2)SO(_4)</td>
<td>-0.5-0.5</td>
<td>602</td>
<td>50 mV s(^{-1})</td>
<td>Ag/AgCl</td>
<td>3E</td>
<td>5000</td>
<td>[45]</td>
</tr>
<tr>
<td>Material</td>
<td>Electrolyte</td>
<td>Potential Range (V)</td>
<td>Current Density (A g(^{-1}))</td>
<td>Reference</td>
<td></td>
<td></td>
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<tr>
<td>PANI-F</td>
<td>2M H(_2)SO(_4)</td>
<td>0.2-0.8</td>
<td>0.1 A g(^{-1})</td>
<td>3E</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PANI80wt%-graphene</td>
<td>2M H(_2)SO(_4)</td>
<td>0.2-0.8</td>
<td>0.1 A g(^{-1})</td>
<td>3E</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PANI50wt%-graphene</td>
<td>2M H(_2)SO(_4)</td>
<td>0.2-0.8</td>
<td>0.1 A g(^{-1})</td>
<td>3E</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PANI10wt%-graphene</td>
<td>2M H(_2)SO(_4)</td>
<td>0.2-0.8</td>
<td>0.1 A g(^{-1})</td>
<td>3E</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PANI-CNT</td>
<td>1M H(_2)SO(_4)</td>
<td>0.2-0.7</td>
<td>5.9 A g(^{-1})</td>
<td>SCE</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PEDOT:PSS-RuO(_2)</td>
<td>0.5M H(_2)SO(_4)</td>
<td>0-0.9</td>
<td>50 mV s(^{-1})</td>
<td>3E</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PEDOT-MWNT</td>
<td>1M H(_2)SO(_4)</td>
<td>0-0.8</td>
<td>2 mV s(^{-1})</td>
<td>2E</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PEDOT-MWNT (70:30wt %)</td>
<td>1M TEABF(_4) in AN</td>
<td>0-0.8</td>
<td>2 mV s(^{-1})</td>
<td>2E</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PEDOT-MWNT (70:30wt %)</td>
<td>1M TEABF(_4) in AN</td>
<td>0-0.8</td>
<td>2 mV s(^{-1})</td>
<td>2E</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ppy-Graphene</td>
<td>1M NaCl</td>
<td>0-1.0</td>
<td>1 A g(^{-1})</td>
<td>2E</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

2.2.3.3 Transitional Metal Oxides

Ruthenium Oxides

Among the transition metal oxides, RuO$_2$ is the most widely researched due to its high specific capacitance and good electrical conductivity.[51] Its hydrous form, RuO$_2$.H$_2$O has shown a high specific capacitance of up to 768 F g$^{-1}$.[52] Higher capacitance is ascribed to the additional proton (H$^+$) intercalation into the hydrated amorphous lattice whereas crystalline RuO$_2$ involve only surface and near surface redox reaction. The author also reported that water is essential for proton (H$^+$) diffusion into the amorphous lattice. Such structure also process excellent chemical stability and good electrical conductivity of the order of 10 S cm$^{-1}$ necessary for good capacitor performance.[53] Accordingly, the reaction for the redox reaction is given by [54],

$$H_aRuO_2.xH_2O + \delta H^+ + \delta e^- \rightleftharpoons H_a + \delta RuO_2.xH_2O$$

Recent studies have revealed that nanostructures enhance the capacitance of RuO$_2$. RuO$_2$ in various forms, nanorods [55], exfoliated nanosheets [56], nanoparticles [57], and nanotubes array [58] have shown improved specific capacitance. However, their scarcity and high cost severely hindered its application in commercial supercapacitor. Although making ruthenium containing composite might reduce the cost, such approach often undermines the capacitance.[44, 59, 60] Hence, considerable effort was spent researching alternative cost effective transitional metal oxides system capable of redox reaction.
Manganese Oxides

Manganese oxide such as MnO$_2$ has shown attractive capacitance besides having a lower toxicity and cheaper cost. Composites based on MnO$_2$ have been extensively investigated for supercapacitor application. Goodenough et al.\cite{61} initially reported amorphous MnO$_2$ composite electrode exhibiting a pseudocapacitive response in mild aqueous electrolyte with a decent specific capacitance of 200 F g$^{-1}$. There are several oxidation states available for manganese oxides, including Mn(0), Mn(I), Mn(II), Mn(III), Mn(IV), Mn(V), Mn(VI), Mn(VII).\cite{62} It has been postulated that charge storage processes in MnO$_2$ is accompanied by the intercalation and de-intercalation of both proton (H$^+$) and surface adsorption of alkali cations (M$^+$).\cite{63,64} The respective electrode reactions are shown as follow.\cite{65,66}

$$\text{MnO}_2 + \delta \text{H}^+ + \delta e^- \rightleftharpoons \text{H}_\alpha + \delta \text{MnOOH}_6 (0 < \delta < 1)$$

$$\text{(MnO}_2)_\text{surface} + M^+ + e^- \rightleftharpoons \text{H}_\alpha + (\text{MnOO}^{-}\text{M}^+)_{\text{surface}}$$

Since only the outer surface of electrode is used for charge storage, a decrease in particles size will result in better utilization of material. Also, the diffusion of cations will be eased along with better proton (H$^+$) intercalation and de-intercalation. Toupin et al.\cite{63} has demonstrated that nano-metric amorphous MnO$_2$ thin film exhibits an ultrahigh specific capacitance of 1380 F g$^{-1}$. However, due to the low diffusivity of protons and alkali cations in conventionally prepared MnO$_2$ particles in presence of binder, the specific capacitance is often below 200 F g$^{-1}$. One shortcoming of MnO$_2$ symmetrical device is lack of oxidation states in manganese at negative potential.\cite{8,67} Thus,
research effort was spent on utilizing MnO₂ as a positive electrode in asymmetrical device. [8, 68]

**Tin Oxides**

Tin oxide based (SnO₂ and SnOₓ) materials being applied in supercapacitor application have also been reported.[69] Alkali metal cations play a critical role charge-discharge mechanism in SnOₓ.[70] It is proposed that surface adsorption (desorption) of cations (M⁺) along with proton (H⁺) and cations (M⁺) intercalation (de-intercalation) occur simultaneously as follow,

\[
(SnO_x)_{surface} + M^+ + e^- \rightleftharpoons H_a + (SnO_xM^+)_{surface}
\]

\[
SnO_x + H^+ + \delta e^- \rightleftharpoons SnO_xH
\]

\[
SnO_x + M^+ + \delta e^- \rightleftharpoons SnO_xM
\]

**Vanadium Oxides**

Amorphous vanadium oxides (V₂O₅) have also received considerable attention as candidate for supercapacitor material. Initially, Lee et al. has reported its supercapacitive behavior in aqueous KCl electrolyte. [71] They proposed that the K⁺ cations in the KCl electrolyte actually contribute to supercapacitive redox reaction. Similar to MnO₂, charges reside mainly on the surface, with limited ions diffusion distance of ca 30 to 50 nm. [72] Often, conducting matrices such as carbon and CNT are incorporated into V₂O₅ to enhance pseudocapacitive yield. Indeed, nano-metric thick amorphous hydrated
vanadium oxide (V$_{2}$O$_{5}$xH$_{2}$O) thin film electrodeposited on CNT has shown improved specific capacitance.[73, 74] Sofiane et al. have shown that specific capacitance actually falls off with increasing thickness of V$_{2}$O$_{5}$. [74]

**Iron Oxides**

Iron oxide (α-Fe$_{2}$O$_{3}$) is an attractive candidate for supercapacitor due to its low cost, abundance and environmentally benign. The specific capacitances of iron based materials are dependent on the surface morphology and generally reported to be 5-210 F g$^{-1}$.[75-78] Electrochemical capacitive behavior is reported to be enhanced by switching to nano-sized materials. Large capacitances have also been reported in sulfites solution with capacitive current deriving entirely from the EDL alone.[79] However, their poor electronic conductivity has restricted their applications in high power devices. CNTs have been added to improve the conductivity with moderate success. Zhao et al. [80] have reported a hybrid supercapacitor made of MWNTs and α-Fe$_{2}$O$_{3}$/MWNTs thin film having a high energy density of 50 Wh kg$^{-1}$ at a specific power density of 1 kW kg$^{-1}$ over a potential range of 0-2.8V. However, the specific capacitance was reduced by 92% when the scan rate was increase from 2 to 200 mV s$^{-1}$ due to sluggish kinetics and poor electron transport within the α-Fe$_{2}$O$_{3}$ film.

Another form of iron oxide (Fe$_{3}$O$_{4}$) has also been investigated for supercapacitor application. Yet, the low specific capacitance limits its practical application.[81, 82] Chen et al. [77] has reported a moderately low specific capacitance of 118.2 F g$^{-1}$, which
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is among the highest reported value so far. Fe$_3$O$_4$ also suffers low capacitance retention of 88.8% after 500 cycles.
Table 2-2: A comparison of capacitance and voltage windows of various Transitional metal oxides and composites in different electrolytes for supercapacitor.

<table>
<thead>
<tr>
<th>Material</th>
<th>Electrolyte</th>
<th>Working Voltage (V)</th>
<th>Specific Capacitance (F/g)</th>
<th>Scan Rate (mV s(^{-1})) or Current Rate (A g(^{-1}))</th>
<th>Reference Electrode</th>
<th>System</th>
<th>Cycles</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>RuO$_2\cdot$xH$_2$O</td>
<td>0.5M H$_2$SO$_4$</td>
<td>0-1.0</td>
<td>720</td>
<td>2 mV/sec</td>
<td>SCE</td>
<td>3E</td>
<td>-</td>
<td>[52]</td>
</tr>
<tr>
<td>Fe$_2$O$_3$</td>
<td>1M NaOH</td>
<td>-0.6-0.1</td>
<td>178</td>
<td>5 mV/sec</td>
<td>SCE</td>
<td>3E</td>
<td>-</td>
<td>[83]</td>
</tr>
<tr>
<td>MnO$_2$</td>
<td>0.5M H$_2$SO$_4$</td>
<td>-0.5-0.5</td>
<td>225</td>
<td>50 mV/sec</td>
<td>Ag/AgCl</td>
<td>3E</td>
<td>5000</td>
<td>[45]</td>
</tr>
<tr>
<td>SnO$_2$/C</td>
<td>1M H$_2$SO$_4$</td>
<td>0-1</td>
<td>37.8</td>
<td>5 mV/sec</td>
<td>Ag/AgCl</td>
<td>3E</td>
<td>-</td>
<td>[84]</td>
</tr>
<tr>
<td>Co$_3$O$_4$</td>
<td>6M KOH</td>
<td>-0.1-0.5</td>
<td>456</td>
<td>1 A/g</td>
<td>Ag/AgCl</td>
<td>3E</td>
<td>500</td>
<td>[85]</td>
</tr>
<tr>
<td>V$_2$O$_5$/C</td>
<td>2M KCl</td>
<td>-0.2-0.7</td>
<td>295</td>
<td>5 mV/sec</td>
<td>SCE</td>
<td>3E</td>
<td>-</td>
<td>[86]</td>
</tr>
<tr>
<td>MoO$_2$/C</td>
<td>1M Na$_2$SO$_4$</td>
<td>0-1.2</td>
<td>93</td>
<td>0.1 A/g</td>
<td>SCE</td>
<td>3E</td>
<td>1500</td>
<td>[87]</td>
</tr>
<tr>
<td>Fe$_3$O$_4$</td>
<td>1M Na$_2$SO$_4$</td>
<td>-1-0.1</td>
<td>118</td>
<td>6mA</td>
<td>SCE</td>
<td>3E</td>
<td>500</td>
<td>[77]</td>
</tr>
<tr>
<td>NiO/CNT</td>
<td>1M KOH</td>
<td>0-0.65</td>
<td>260</td>
<td>20A/g</td>
<td>Hg/HgO</td>
<td>3E</td>
<td>1000</td>
<td>[88]</td>
</tr>
<tr>
<td>IrO$_2$/MWNT</td>
<td>0.1M KOH</td>
<td>-0.5-0.5</td>
<td>74</td>
<td>10 mV/sec</td>
<td>Ag/AgCl</td>
<td>3E</td>
<td>200</td>
<td>[89]</td>
</tr>
<tr>
<td>MWNT</td>
<td>1M H$_2$SO$_4$</td>
<td>0-1</td>
<td>69</td>
<td>10mA</td>
<td>-</td>
<td>2E</td>
<td>-</td>
<td>[90]</td>
</tr>
<tr>
<td>SnO$_2$/MWNT</td>
<td>1M H$_2$SO$_4$</td>
<td>0-1</td>
<td>91</td>
<td>10mA</td>
<td>-</td>
<td>2E</td>
<td>-</td>
<td>[90]</td>
</tr>
<tr>
<td>RuO$_2$/MWNT</td>
<td>1M H$_2$SO$_4$</td>
<td>0-1</td>
<td>133</td>
<td>10mA</td>
<td>-</td>
<td>2E</td>
<td>-</td>
<td>[90]</td>
</tr>
<tr>
<td>TiO$_2$/MWNT</td>
<td>1M H$_2$SO$_4$</td>
<td>0-1</td>
<td>148</td>
<td>10mA</td>
<td>-</td>
<td>2E</td>
<td>-</td>
<td>[90]</td>
</tr>
<tr>
<td>MnO$_2$</td>
<td>1M KNO$_3$</td>
<td>0-0.6</td>
<td>160</td>
<td>-</td>
<td>-</td>
<td>2E</td>
<td>-</td>
<td>[91]</td>
</tr>
<tr>
<td>MnO$_2$</td>
<td>0.1M K$_2$SO$_4$</td>
<td>0-1.0</td>
<td>36</td>
<td>-</td>
<td>-</td>
<td>2E</td>
<td>-</td>
<td>[8]</td>
</tr>
<tr>
<td>V$_2$O$_5$</td>
<td>2M KCl</td>
<td>0-0.9</td>
<td>190</td>
<td>2mV/sec</td>
<td>-</td>
<td>2E</td>
<td>-</td>
<td>[92]</td>
</tr>
</tbody>
</table>
2.2.4 Asymmetrical Supercapacitor

An aqueous based supercapacitor has particular advantages over its organic counterpart in terms of higher ionic conductivity, lower cost, lower hazards and better power capability. Such devices have lower sensitivity to oxygen and moisture, hence, require less complicated production processes. Convention supercapacitors are made up of two symmetrical electrodes, and have a narrow stable voltage window of 1V resulting in a low energy density.

Recently, asymmetrical supercapacitors which couple electrodes operating reversibly in different potential ranges have been proposed as a promising way to increase voltage in aqueous electrolytes. This asymmetric supercapacitor can reliably operate above 1.23V (the breakdown voltage of water) without gas evolution in aqueous electrolyte. Operation is made possible because the reaction kinetics for gas evolution is slow although thermodynamically feasible. This results in higher energy density according to equation 2.5. Such asymmetrical supercapacitor allows an operating voltage that may approach twice that of EDLC and thus greatly improves the energy density.
Table 2-3: A comparison of capacitance and voltage windows of various asymmetrical supercapacitors in different electrolytes for supercapacitor.

<table>
<thead>
<tr>
<th>Positive Electrode (Cathode)</th>
<th>Negative Electrode (Anode)</th>
<th>Electrolyte</th>
<th>Working Voltage (V)</th>
<th>Specific Capacitance (F/g)</th>
<th>Energy (Wh/kg)</th>
<th>Power (kW/kg)</th>
<th>Cycles</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>MnO₂</td>
<td>AC</td>
<td>1M LiOH</td>
<td>0-1.5</td>
<td>62.4</td>
<td>19.5</td>
<td>-</td>
<td>1500</td>
<td>[93]</td>
</tr>
<tr>
<td>MnO₂</td>
<td>AC</td>
<td>0.5M K₂SO₄</td>
<td>0-1.5</td>
<td>-</td>
<td>7</td>
<td>10</td>
<td>23000</td>
<td>[94]</td>
</tr>
<tr>
<td>MnO₂</td>
<td>AC</td>
<td>0.5M K₂SO₄</td>
<td>0-1.8</td>
<td>31</td>
<td>17</td>
<td>2</td>
<td>23000</td>
<td>[95]</td>
</tr>
<tr>
<td>MnO₂</td>
<td>AC</td>
<td>1M KNO₃</td>
<td>0-2.0</td>
<td>140</td>
<td>21</td>
<td>123</td>
<td>1000</td>
<td>[91]</td>
</tr>
<tr>
<td>MnO₂</td>
<td>AC</td>
<td>0.1M K₂SO₄</td>
<td>0-2.0</td>
<td>21</td>
<td>11.7</td>
<td>16</td>
<td>195000</td>
<td>[68]</td>
</tr>
<tr>
<td>MnO₂</td>
<td>AC</td>
<td>1M KCl</td>
<td>0-2.0</td>
<td>52</td>
<td>28.8</td>
<td>8</td>
<td>100</td>
<td>[96]</td>
</tr>
<tr>
<td>MnO₂</td>
<td>AC</td>
<td>0.1M K₂SO₄</td>
<td>0-2.2</td>
<td>31</td>
<td>17.3</td>
<td>19</td>
<td>10000</td>
<td>[8]</td>
</tr>
<tr>
<td>MnO₂</td>
<td>PANI</td>
<td>1M KNO₃</td>
<td>0-1.2</td>
<td>-</td>
<td>5.86</td>
<td>42.1</td>
<td>500</td>
<td>[97]</td>
</tr>
<tr>
<td>MnO₂</td>
<td>Ppy</td>
<td>1M H₂SO₄</td>
<td>0-1.4</td>
<td>-</td>
<td>7.37</td>
<td>62.8</td>
<td>500</td>
<td>[97]</td>
</tr>
<tr>
<td>MnO₂</td>
<td>PEDOT</td>
<td>1M KNO₃</td>
<td>0-1.8</td>
<td>-</td>
<td>13.5</td>
<td>120.1</td>
<td>500</td>
<td>[97]</td>
</tr>
<tr>
<td>MnO₂/C</td>
<td>V₂O₅ NW/MWNT</td>
<td>1M Na₂SO₄</td>
<td>0-1.6</td>
<td>45</td>
<td>16</td>
<td>3.75</td>
<td>100</td>
<td>[98]</td>
</tr>
<tr>
<td>MnO₂/MWNT</td>
<td>SnO₂/MWNT</td>
<td>2M KCl</td>
<td>0-1.7</td>
<td>38</td>
<td>20.3</td>
<td>143.7</td>
<td>1000</td>
<td>[99]</td>
</tr>
<tr>
<td>MnO₂</td>
<td>Fe₃O₄</td>
<td>0.1M K₂SO₄</td>
<td>0-1.8</td>
<td>21.5</td>
<td>8.1</td>
<td>10.2</td>
<td>5000</td>
<td>[8]</td>
</tr>
<tr>
<td>Fe₃O₄</td>
<td>AC</td>
<td>6M KOH</td>
<td>0-1.2</td>
<td>37.9</td>
<td>7.6</td>
<td>0.07</td>
<td>500</td>
<td>[82]</td>
</tr>
<tr>
<td>LiMn₂O₄</td>
<td>AC</td>
<td>1M Li₂SO₄</td>
<td>0-1.8</td>
<td>56</td>
<td>10</td>
<td>2</td>
<td>20000</td>
<td>[100]</td>
</tr>
<tr>
<td>Hydrous V₂O₅</td>
<td>AC</td>
<td>0.5M K₂SO₄</td>
<td>0-1.8</td>
<td>64.4</td>
<td>29</td>
<td>2</td>
<td>10</td>
<td>[101]</td>
</tr>
<tr>
<td>NiO</td>
<td>C</td>
<td>6M KOH</td>
<td>0-1.3</td>
<td>38</td>
<td>10</td>
<td>10</td>
<td>1000</td>
<td>[102]</td>
</tr>
<tr>
<td>Ni(OH)₂/MWNT</td>
<td>AC</td>
<td>6M KOH</td>
<td>0-1.5</td>
<td>96</td>
<td>32</td>
<td>1.5</td>
<td>2000</td>
<td>[103]</td>
</tr>
<tr>
<td>MWNT</td>
<td>Fe$_2$O$_3$-MWNT</td>
<td>1M LiPF$_6$ in EC:DMC</td>
<td>0-2.8</td>
<td>80</td>
<td>50</td>
<td>1</td>
<td>500</td>
<td>[80]</td>
</tr>
<tr>
<td>--------</td>
<td>------------------</td>
<td>----------------------</td>
<td>-------</td>
<td>----</td>
<td>----</td>
<td>----</td>
<td>-----</td>
<td>-------</td>
</tr>
<tr>
<td>MWNT</td>
<td>TiO$_2$ NW</td>
<td>1M LiPF$_6$ in EC:DEC:DMC</td>
<td>0-2.8</td>
<td>11.5</td>
<td>12.5</td>
<td>1.2</td>
<td>600</td>
<td>[104]</td>
</tr>
<tr>
<td>AC</td>
<td>TiO$_2$</td>
<td>1M LiClO$_4$ in EC:DMC</td>
<td>1.2-3.5</td>
<td>44</td>
<td>80</td>
<td>0.35</td>
<td>600</td>
<td>[105]</td>
</tr>
<tr>
<td>RuO$_2$/TiO$_2$</td>
<td>AC</td>
<td>1M KOH</td>
<td>0-1.4</td>
<td>46</td>
<td>12.5</td>
<td>0.15</td>
<td>1000</td>
<td>[106]</td>
</tr>
<tr>
<td>PEDOT-C</td>
<td>AC</td>
<td>1M TEABF$_4$ in AN</td>
<td>0-1.5</td>
<td>160</td>
<td>-</td>
<td>-</td>
<td>20000</td>
<td>[49]</td>
</tr>
</tbody>
</table>

2.2.5 Performance of Supercapacitor

The performance of supercapacitor depends on various attributes. Figure 2.4 shows that a supercapacitor can be treated as two capacitors in series.

\[ \frac{1}{C_T} = \frac{1}{C_p} + \frac{1}{C_n} \]  

(2.4)

Figure 2.1: Schematic of an electrical double layer capacitor and its circuit equivalent.
Literature Review

If the two electrodes are the same, that is capacitance of positive and negative electrode is identical, the overall capacitance \( C_T \) would be half of the capacitance of one electrode. Such is the case in symmetrical supercapacitor. In the case of asymmetrical supercapacitor, where the electrodes are made of different materials, \( C_T \) is mainly dominated by the one with smaller capacitance.

The energy density \( E \) and theoretical maximum power density \( P_M \) can be expressed as:

\[
E = \frac{1}{2} C_T V^2 \tag{2.5}
\]

\[
P_M = \frac{V^2}{4R_{ESR}} \tag{2.6}
\]

Where \( C_T \) is the overall capacitance or device capacitance, \( V \) is the operating voltage of the supercapacitor, \( R_{ESR} \) is the equivalent series resistance of the supercapacitor.

The operating voltage is determined by the material used for electrode and stability window of the electrolyte involved.
Chapter 3 Experimental Setup

3.1 Fabrication of Electrode Materials

3.1.1 Filtration of CNT Buckypaper

SWNT was prepared by dispersing 30 mg of commercial SWCNT (P3-CNT, Carbon Solution Inc.) 200 ml deionized water, and sonicated for 30 minutes using a bath sonicator. Figure 3.1 illustrates the procedures taken to obtain a bucky paper. The SnO₂ mixed P3-CNT composite was prepared by adding 6 weight percentage of SnO₂ hollow sphere particles to the CNT suspension and stirred for 12 hours to obtain homogenous solution. The suspension was then filtered through a filter membrane (Whatman, 20 nm pore size, 47 mm diameter) using a filter flask and vacuum pump. This mesh was subsequently air dry at room temperature for 24 hours to peel off as a freestanding thin film with a thickness of approximately 20 μm. The film is then sized appropriately to be used for characterization. An interconnected network of SWNT (pristine or mixed with SnO₂ nanomaterials) is depicted at bottom of Figure 3.1.
Experimental Setup

Dispersed SWNT solution
+ TMO

Figure 3.1: (Top) Obtaining CNT bucky paper via vacuum filtration. (Bottom) Photograph of a flexible CNT bucky paper.
3.1.2 SnO₂ and P3-CNT mixture

SnO₂ hollow nanospheres was synthesized in accordance to a method described by X. W. Lou et al. [107]. Briefly, certain amounts of urea and potassium stannate trihydrate (K₂SnO₃·3H₂O, Aldrich, 99.9%) were dissolved in a mixed solvent of ethanol/water (37.5% ethanol by volume) to obtain a concentration of 0.1M. The solution was then transferred to a teflon lined stainless steel hydrothermal autoclave and heated in an air flow oven at 150°C for 24 hours. The white precipitate was then obtained by centrifugation at 3000 rpm, washed with ethanol and deionized water. The product was then dried at room temperature in vacuum oven.
3.1.3 Electrospinning of Nanofibers

Electrospinning is a simple facile and economical way to fabricate interwoven nanofibers. Figure 3.2 illustrates the general electrospinning setup. The setup is highly versatile equipped with various parameters such as flow rate, voltage, size of needle and distance between plate and needle to better control the electrospinning process and aspect ratio of the electrospun fibers.

![Schematic illustration of the Electrospinning Process](image)

**Figure 3.2**: Schematic illustration of the Electrospinning Process
Experimental Setup

3.1.3.1 $\alpha$-Fe$_2$O$_3$ Nanofibers

For synthesis of Fe(acac)$_2$/PVP composite nanofibers, 1 g of Polyvinylpyrrolidone (PVP) (Mw 3,600,000 Fluka) was dissolved in mixture of absolute ethanol (Sigma-Aldrich) and glacial acetic acid (Tedia Company Inc.). Then 0.6 g of iron (II) acetylacetonate (Fe(acac)$_2$, 98%, Sigma-Aldrich) was added, and stirred for 24 h to obtain a homogenous solution. All chemical reagents were used as received. In the electrospinning procedure, the Fe(acac)$_2$/PVP solution was transferred into a 10 ml glass syringe mounted onto an electrospinning apparatus. The feeding rate of precursor solution was controlled at 1 ml h$^{-1}$ by a syringe pump (KDS Scientific Model 210). A positive potential of 10 kV was applied to the stainless steel needle tip of ~200 μm diameter (SS 304) through a high voltage source (Gamma High Voltage Research). Distance between the needle tip and collector (aluminium foil) was maintained around 10 cm. Under the strong electric field, fibers were ejected from the Taylor cone formed at the needle tip and deposited on the collector. Electrospun fibers were subsequently heated at a ramping rate of 4 °C min$^{-1}$ to 500 °C, annealed for 5 hours, and cooled naturally to room temperature. The obtained fibers were reddish mats in appearance indicating the Fe$^{3+}$ oxidation state of iron.
Experimental Setup

3.1.3.2 V$_2$O$_5$ Nanofibers

In making VO(acac)$_3$/PVP composite nanofibers, 1 g of Polyvinylpyrrolidone (PVP) (Mw 3,600,000 Fluka) was dissolved in mixture of absolute ethanol (Sigma-Aldrich) and glacial acetic acid (Tedia Company Inc.). Then 0.6 g of vanadyl acetylacetonate (VO (acac)$_3$, 98%, Sigma-Aldrich) was added, and stirred for 24 h to obtain a homogenous solution. All chemical reagents were used as received. In the electrospinning procedure, the VO(acac)$_3$/PVP solution was transferred into a 10 ml glass syringe mounted onto an electrospinning apparatus. The feeding rate of precursor solution was controlled at 1 ml h$^{-1}$ by a syringe pump (KDS Scientific Model 210). A positive potential of 10 kV was applied to the stainless steel needle tip of ~200 μm diameter (SS 304) through a high voltage source (Gamma High Voltage Research). Distance between the needle tip and collector (aluminium foil) was maintained around 10 cm. Under the strong electric field, fibers were ejected from the Taylor cone formed at the needle tip and deposited on the collector. Electrospun fibers were subsequently heated at a ramping rate of 2 °C min$^{-1}$ to 500 °C, annealed for 30 minutes, and cooled at 2 °C min$^{-1}$ to room temperature. The obtained fibers were yellowish in colour indicating the V$^{5+}$ oxidation state of vanadium.
3.1.4 Rapid Polymerization of Polyaniline Nanofibers

PANF was synthesized by rapid mixing polymerization [108]. The oxidant dopant solution was prepared by dissolving 25 mmol of ammonium persulfate (98% Sigma-Aldrich) in 50 ml deionized water, and stirred for 30 min. Aniline dopant solution was made by adding 20 mmol of aniline (Sigma-Aldrich) into 50 ml deionized water, and stirred for 30 min. The oxidant solution is then quickly added to the aniline solution and stirred vigorously for a minute, and left to polymerized for an hour. After the polymerization, the dark green residue was washed with 300 ml of 0.2 M HCl solution followed by 300 ml of acetone to remove residual ammonium persulfate and aniline monomers respectively. The dark green residue was then dried at 50 °C in vacuum for 12 hours.
Experimental Setup

3.2 Fabrication of Composite Electrode

The test electrodes were prepared by dispersing VNFs or PANFs, super P (99%, Alfa Aesar) and poly-vinylidene fluoride (PVdF) (Sigma-Aldrich) in N-methylpyrrolidone (NMP) (Sigma-Aldrich) with a weight ratio of 75:15:10, respectively to form slurry. Resultant slurry was then coated onto a 1 cm\(^2\) area of graphite paper and subsequently dried at 60°C for an hour. The mass loading of electrode material is ca. 2 mg.

![Figure 3.3: Photograph of electrodes coated with composites](image-url)
Experimental Setup

3.3 Physical Characterizations

3.3.1 Field-Emission Scanning Electron Microscopy (FESEM)

The surface morphologies of the nanostructured electrode materials were investigated using a field-emission scanning electron microscope (JSM 6340 F) operating with accelerating voltages of 5 kV to 10 kV. Samples were prepared by pasting the subject onto the stainless steel holder using carbon tapes. All samples were coated with ca. 15-25 nm of using a platinum sputter (JEOL JFC-1600) prior to microscopy to avoid charging effect.

3.3.2 Powder X-ray Diffraction (XRD)

Powder X-ray diffraction (XRD) was performed using a Shimadzu diffractometer (Cu-kα) with step scanning (0.01°, 0.5 s dwell time) over a 2θ range of 10° to 140°. The patterns were analyzed by applying the Rietveld method using fundamental parameters approach contained within the software TOPAS V3 (Bruker-AXS).

3.3.3 High Resolution Field Emission Transmission Electron Microscopy

The high resolution TEM (HRTEM) images were performed at 200kV by using a JEOL JEM-2100F transmission electron microscope equipped with high contrast objective aperture (20μm in diameter) equivalent to a point to point resolution of 1.7Å.
Experimental Setup

3.3.4 Nitrogen adsorption-desorption isotherm

The nitrogen adsorption-desorption isotherms were obtained at 77 K using a NOVA 3200e system. The BET surface area of the samples was determined by investigating the nitrogen desorption isotherms using a NOVA 3200e system at 77 K. The samples were degassed at 373 K under nitrogen flow for 24 hours prior to measurement. Pores size distributions were investigated by applying Barrett-Joyner-Halenda (BJH) analysis on the nitrogen desorption isotherm obtained.

3.3.5 Thermo gravimetric Analysis (TGA)

The experiment was carried out with a heating rate of 2 °C min⁻¹ or 10 °C min⁻¹ from room temperature to 800°C using the TGA Q500 Series from TA. Gas flow consisting of a mixture of synthetic air and nitrogen gas in the percentage ratio of 40:60 was maintained.

3.3.6 Fourier Transform Infrared Spectroscopy (FT-IR)

In a typical experiment, potassium bromide and a small amount of the nanofibers are mixed homogenously using a mortar and pestle. The mixture was then transferred to a mould and subjected to a pressure of 10 kPa for 5 minutes. A highly transparent disc containing less than 2% weight ratio of the testing subject (nanofibers) was obtained. Infrared spectroscopy was then performed on the disc, scanned sixteen times over the range of 4000-400 cm⁻¹ using Perkin-Elmer Spectrum GX FTIR.
Experimental Setup

3.3.7 X-ray Photoelectron Spectroscopy (XPS)

X-ray photoelectron spectroscopy (XPS) spectra were recorded using a Kratos AXIS Ultra spectrometer with the monochromatic Al Kα X-ray radiation. Deconvolution and peak fitting were performed with computer assisted surface analysis XPS software (CASA) using a Voigt function with 30% Lorentzian character and a Shirley background.

3.4. Electrochemical Characterization

3.4.1 Cyclic Voltammetry (CV)

The electrochemical behaviors of the electrode material were investigated using cyclic voltammetry measurements carried out in various configurations, including three electrode configuration, two electrode symmetrical and asymmetrical device configurations. A standard three electrode cell configuration is consisted of a working electrode (electrode material under evaluation), platinum foil (5 cm x 5 cm) as counter electrode, and saturated calomel electrode (SCE) or silver chloride electrode (Ag/AgCl in 3M KCl) as reference electrode. All the experiments were conducted using a computer controlled potentiostat (Solartron 1470E).
Experimental Setup

3.4.2 Galvanostatic Charge-Discharge (GCD)

Galvanostatic charge-discharge performances were conducted in a two electrode configuration. The measurements were taken using a computer controlled potentiostat (Solartron, 1470E). Different current densities were employed to investigate the rate capability of the various materials. Effect of organic electrolyte such as LiClO$_4$ was also studied.

3.4.3 Electrochemical Impedance Spectroscopy (EIS)

Electrochemical impedance for all the devices was measured using a frequency response analyzer from Solartron (SI 1255B impedance/gain-phase analyzer with Zview software) in potentiostatic mode with AC amplitude of 10 mV, from 10 kHz to 10 mHz.
Chapter 4 Results and Discussions

This chapter details the results of various experiments undertaken in this thesis. Firstly, P3-CNTs being commercially available are first explored as a working prototype, underpinning subsequent research work. It was subsequently mixed with materials such as SnO₂ in attempt to increase the capacitance via pseudocapacitance.

Electrospinning was employed to synthesize nanofibers to investigate the effect of morphology in capacitive performance. TMO nanomaterial such as Fe₂O₃ and V₂O₅ were electrospun and their performances investigated. The effect of conducting additive such as carbon black is also studied. Also, the performance of Fe₂O₃ in various electrolytes is looked into. Their superior electrochemical behaviors brought about by the nanostructure were also investigated. In addition, V₂O₅ nanofibers and other class of materials like electronically conducting polymer (ECP) such as polyaniline were also studied.

To increase the energy density, asymmetrical configuration was employed to increase the working potential. A common electrolyte was judiciously selected in which both the electrode material can function as supercapacitor electrode. Their electrochemical behaviors and performances were being investigated and compared. A correlation of the charge storage mechanism of the supercapacitor with the conducting additive, morphology, surface area and interfacial properties had been investigated through various electrochemical characterization techniques.
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4.1 Investigation of Electrode Materials for Symmetrical Supercapacitor Application

4.1.1 CNT based Supercapacitors

Firstly, the use of CNTs as both active electrode material and the current collector was explored. CNTs have high surface area, conductivity and aspect ratio which make them a good candidate for electrode material in supercapacitor application. As P3-CNTs are commercially available, attempts to fabricate supercapacitors were made using P3-CNTs.

![Figure 4.1](image)

**Figure 4.1** : (a) FESEM image of P3-CNT (Carbon solutions) bucky paper. (b) HR-TEM image of a bundle of P3 CNT.
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![Graph showing TGA of P3-CNT (Carbon solutions).](image)

Figure 4.2 illustrates the thermal gravimetric analyses of P3 CNTs. There is a gradual weight loss observed for P3 CNT at lower temperature (ca. 100 °C) due to moisture left behind by the filtration process. There is some weight loss of ca. 10 wt% over the range of 200 to 400 °C, probably due to the decomposition of the functional groups such as carboxylic, and hydroxyl. The significant drop in wt % occurring from 400 to 700 °C can be ascribed to the decomposition of carbon. No observable weight change is noted beyond 750 °C, indicating the completion of carbon combustion. The incombustible residual can be ascribed to the metal catalysts impurities used during the
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growth process of the CNTs. The final residual weight of 9.8 wt% was obtained for P3-CNT suggesting a purity of more than 90%.

Bucky papers of P3-CNT were tested in various electrolytes of 1M H₂SO₄, 1M Na₂SO₄ and 3M KCl in three electrode configuration with SCE as reference.

Figure 4.3 illustrates the electrochemical CV and GCD curves of P3-CNT bucky paper tested in 1M H₂SO₄ electrolyte. The peak observed in the CV curve at ca. 0.25 V can be attributed to faradic contribution from the oxygenated functional groups. In general, the deviation from an ideal box-shape CV curve of a supercapacitor without internal resistance (dotted line) can be explained by the presence of functional groups, impurities, and a decreased conductivity due to the absence of a dedicated current collector.

Figure 4.3 : (a) 2-electrode Cyclic Voltammogram (b) 2-electrode Galvanostatic charge-discharge of P3-CNT bucky paper in 1M H₂SO₄ electrolyte.
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The specific capacitance of the device from the GCD test is calculated to be 87 F g\(^{-1}\). Besides CV measurements, GCD test is another device characterization that can be employed to evaluate the specific capacitance and internal resistance of a SC device. Figure 4.3 (b) shows GCD curve of the same device measured at a constant current of 1 mA. Each branch of the entire curve starts with a voltage drop (IR drop). As P3-CNT exhibits electrical double layer capacitive property, ways to improve the capacitance will be investigated.
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4.1.2 SnO$_2$-CNT mixture based Supercapacitors

Figure 4.4: (a) FESEM image of SnO$_2$ hollow spheres (b) HR-TEM image of SnO$_2$ hollow spheres (c) FESEM image of P3-CNTs mixed SnO$_2$ hollow spheres.

Figure 4.4: (a) FESEM image of SnO$_2$ hollow spheres (b) HR-TEM image of SnO$_2$ hollow spheres (c) FESEM image of P3-CNTs mixed SnO$_2$ hollow spheres.
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Pseudocapacitive materials such as transition metal oxides were introduced to increase the capacitance thereby enhancing the energy density. The feasibility of direct mixing CNT with SnO_2 was investigated. Figure 4.4 depicts (a) FESEM image, (b) HR-TEM image of SnO_2 spheres and (c) FESEM image of SnO_2 mixed P3-CNT. From Figure 4.4 (b) image, it can be observed that that SnO_2 spheres are hollow which significantly reduce the mass of SnO_2 required and may enable a higher surface area for charge transfer process. The SnO_2 spheres are well embedded into the P3-CNT network as seen in Figure 4.4 (c). This will allow better transportation of electrons to and from the sites of redox processes.

![Figure 4.5](image)

**Figure 4.5:** (a) 3-electrode Cyclic Voltammogram (b) 2-electrode Galvanostatic charge-discharge of P3-CNT bucky paper and SnO_2 mixed P3-CNT bucky paper in 1M H_2SO_4 electrolyte.
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Figure 4.5 depicts the electrochemical performance of both the P3-CNT and physical mixture of SnO$_2$ P3-CNT electrode in 1 M H$_2$SO$_4$ electrolyte. Apparently, addition of SnO$_2$ does not significantly contribute to the electrochemical performance. The minute increment in pseudocapacitive contribution observed in SnO$_2$-CNT devices had led to the out-phasing of direct mixing of pseudocapacitive material nanomaterials with CNTs in subsequent investigation.

To continue the effort of improving the energy density of capacitance of materials, other TMO as well as electrically conducting polymers (ECP) were investigated. Candidate materials need to either possess multiple oxidation states or be capable of being dope/de-doped.
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4.1.3 \( \text{Fe}_2\text{O}_3 \) Nanofibers Supercapacitors

Bulk chemical materials have already been intensively researched in many applications. However, the chemistry and performance of many nanomaterials have not yet been widely explored and understood. Nanostructures with high surface area will certainly lead to better capacitance. Currently, there are many routes to synthesize nanomaterials, including hydrothermal [109] and electrospinning [110]. Electrospinning is an economical, easy and reliable method to fabricate high surface area nanofibers with diameters ranging from tens of nanometers to several micrometers with good mechanical strength. It also offers advantages like control over morphology, composition and porosity with simple equipment. Additionally, these nanofibers can often become porous after calcination at suitable temperatures and heating rate [111]. Hence, electrospinning was employed to fabricate nanostructured material in this work.

4.1.3.1 Physical Characterization of \( \text{Fe}_2\text{O}_3 \) Nanofibers

\( \text{Fe}_2\text{O}_3 \) are reported to exhibit pseudocapacitive behavior. Their electrochemical performance is strongly influenced by the surface morphology of the materials, which depends on the methods of preparation. They are also chemically stable, cheap and environmentally benign. As previously mentioned, electrospinning technique was employed to fabricate \( \text{Fe}_2\text{O}_3 \) fibers. \( \text{Fe}(	ext{acac})_2 \) was selected as metal acetylacetonate can be decomposed at moderate temperature (\(< 800 \degree \text{C} \)) to become metal oxide. The electrospun \( \text{Fe}(	ext{acac})_2 \) fibers were subsequently peeled off and analyzed using TGA to find the optimum calcination temperature to obtain \( \text{Fe}_2\text{O}_3 \) nanofibers.
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Figure 4.6 shows the thermal gravimetric analysis (TGA) of as spun Fe(acac)$_2$ nanofibers and PVP respectively. The optimized calcination temperature range of electrospun fibers was studied. The TGA helps to illustrate the thermal behavior of the as spun fibers. Both samples exhibit thermal weight loss below 100°C which can be attributed to the evaporation of moistures present in the PVP. The weight percentage of the FNF continues to decrease gradually in the temperature range of ~100-350 °C, and dropped rapidly at ca. 400 °C. The weight loss from 180°C to 350 °C might be due to the decomposition of Fe(acac)$_2$ and its transformation into α-Fe$_2$O$_3$. The subsequent drop in weight at 360 to 400°C is due to the decomposition of PVP into carbon dioxide and water vapor, as evident from the decomposition curve of pristine PVP. The weight stabilization after 450 °C signifies the end of PVP decomposition and complete formation
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of Fe$_2$O$_3$ phase. These results suggested that the calcination temperature to obtain Fe$_2$O$_3$ after calcinations falls slightly beyond 450 $^\circ$C. Therefore, electrospun fibers for subsequence studies were calcined in the furnace at 500 $^\circ$C, for a constant duration of 5 hours.

![Figure 4.7: FTIR of PVP, as spun FNF-PVP, and calcined FNF respectively.](image)

FT-IR was employed to investigate the bonds present in calcined Fe$_2$O$_3$. Figure 4.7 compares the FT-IR spectra of PVP, as spun composite fibers and Fe$_2$O$_3$ nanofibers calcined at 500 $^\circ$C. The broad peak in the general range of 3600–3100 cm$^{-1}$ may be assigned to anti-symmetrical and symmetrical O–H bonding stretching vibrational modes for O-H.[112] Bonding in the region of 1730–1600 cm$^{-1}$ also relates to O–H bonding.
bending vibrational modes. [112] The bands at 460 and 532 cm\(^{-1}\) observed in Fe\(_2\)O\(_3\) calcined at 500 °C can be attributed to stretching vibrational modes of Fe-O. [112]

The absence of observable change of FT-IR spectra of PVP and as spun composite nanofibers suggests that Fe\(^{2+}\) may have weak or no interactions with PVP. This agrees with the observation in Figure 4.6 showing that the as spun composite nanofibers was having the same decomposition temperature as PVP powder. After calcination at 500 °C for 5 hours, those bands corresponding to organic components was still observed but at a lower intensity. This suggests that some carbon may be left behind from the decomposition of PVP, which may enhance the electrical conductivity of the Fe\(_2\)O\(_3\) nanofibers.
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Figure 4.8: FESEM images of as-spun Fe(acac)$_2$/PVP composite fibers at (a) lower magnification, (b), (c) and (d) nano-porous α-Fe$_2$O$_3$ fibers after calcination at 500 °C.

Morphology of the as spun Fe(acac)$_2$/PVP composite fibers and Fe$_2$O$_3$ nanofibers calcined at 500°C was investigated by FESEM. Figure 4.8 shows FESEM images of Fe(acac)$_2$/PVP composite fibers and Fe$_2$O$_3$ nanofibers. It was observed that the as spun nanofibers were randomly oriented with diameter of 200 – 400 nm and length up to hundreds of micrometers. After undergoing calcination, the nanofibers shrunk to about 4 to 6 times in diameter while maintaining the same fiberous morphology. The annealed Fe$_2$O$_3$ nanofibers have diminished overall diameters (40 to 100 nm). The TEM image
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(inset of Figure 4.8 (d)) further confirms the shrinkage of the fibers, which also exhibits partial crystallinity. This change in fiber aspect ratio was derived from the elimination of PVP during sintering. This removal of the PVP renders more porous surfaces which are beneficial to the diffusion of ions which is crucial to both the power and energy density of supercapacitors.

Figure 4.9: XRD pattern of α-Fe₂O₃ nano-porous fibers.

Identification of the sintered Fe(acac)₂ was investigated with XRD. Figure 4.9 shows the X-ray diffraction patterns of electrospun α-Fe₂O₃ nanofibers after annealing at 500°C for 5 hrs. The XRD reveals reflections belonging to space group R-3CH (JCPDS 62).
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card no. 33-0664). Rietveld refinement was carried out using TOPAS. Lattice parameters of Fe₂O₃ were calculated as $a = 5.036(3)$ Å, $c = 13.76(3)$ Å, fitted with R-Bragg value of 0.664. This XRD indicates that a single phase of hematite α-Fe₂O₃ has been obtained [113, 114].

![Figure 4.10](image.png)

**Figure 4.10**: (Top) Nitrogen adsorption–desorption isotherms and (Bottom) BJH desorption isotherm of the nano-porous α-Fe₂O₃ fibers.

The surface properties of the α-Fe₂O₃ nanofibers were further investigated using the nitrogen isotherm. Figure 4.10 (top) presents nitrogen adsorption/desorption isotherms of α-Fe₂O₃ nanofibers at 77 K. Multipoint Brunauer-Emmett-Teller (BET) analysis performed on α-Fe₂O₃ nanofibers gives a specific surface area of 33.95 m² g⁻¹. The adsorption isotherm exhibited type IV isotherm with type H3 hysteresis between adsorption and desorption isotherms and a steep slope at higher relative pressure,
suggesting mesoporosity. Mesoporous structure is reported to be beneficial to capacitive performance by allowing fast ion diffusion for both double layer formation and surface redox reaction, thus enhancing rate performance. Barrett, Joyner and Halenda (BJH) method was employed to investigate the pore size distribution in $\alpha$-$\text{Fe}_2\text{O}_3$ and depicted in Figure 4.10 (bottom). It is observed that $\alpha$-$\text{Fe}_2\text{O}_3$ nanofibers have pore distribution between 10 to 60 nm with mode at 20 nm. These pores enable smooth traffic of ions from and to the inner active material which may not be accessible otherwise, hence enhancing the power capability of the supercapacitor.

### 4.1.3.2 Effect of Conducting Addictive

![Specific Capacitance Contribution](image)

**Figure 4.11:** Plot of specific capacity contribution from the composite with different carbon black added taken at a current density of 0.1 A g$^{-1}$ in 1 M $\text{Na}_2\text{SO}_4$ electrolyte.
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The effect of carbon black on the pseudocapacitive behavior of $\alpha$-Fe$_2$O$_3$ is also studied. Figure 4.11 depicts the relationship of specific capacitance with current density taken with different weight percentage of $\alpha$-Fe$_2$O$_3$ and carbon black. It is observed that the pseudocapacitance contributed by $\alpha$-Fe$_2$O$_3$ nanofibers is significantly enhanced with addition of carbon black. Although $\alpha$-Fe$_2$O$_3$ is electrochemically active, it suffers a low electrical conductivity of $10^{-14}$ S cm$^{-1}$ [76]. Such low conductivity inevitably hinders the utilization of the pseudocapacitive storage of $\alpha$-Fe$_2$O$_3$. Herein, the carbon black functions as an electrical conductor facilitating electrons transfer to and from the sites of redox reactions, thereby improving the utilization of pseudocapacitive sites of $\alpha$-Fe$_2$O$_3$. Also, having more carbon black content ensures better distribution of $\alpha$-Fe$_2$O$_3$ nanofibers. This allows greater amount $\alpha$-Fe$_2$O$_3$ being able to be used for charge transfer and hence realizing greater charge storage.

It is observed that the specific contribution from carbon black is trivial as compared to $\alpha$-Fe$_2$O$_3$. This is because it is only capable of storing a thin layer of charges in the form of double layer typically of less than 10 nm, hence their limited charge storage capacity.
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Figure 4.12: Plot of specific capacitance of α-Fe₂O₃ nanofibers with different weight percentage of carbon black added against current density in 1 M Na₂SO₄ electrolyte.

The effect of current density on the specific capacitance of α-Fe₂O₃ was also studied. Figure 4.12 shows the relationship of specific capacitance of α-Fe₂O₃ nanofibers of different weight percentage with current density. Generally, specific capacitance is observed to reduce with increasing current density regardless of the amount of carbon black added. As with higher current density, fewer ions are able to diffuse to the redox active sites resulting in less charge transfers and storage. Greater proportion of the total charges is then stored on the surface (non faradic double layer storage). On the other hand, at low current density, more charge transfer is able to proceed given the longer
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charge/discharge duration. This translates into greater pseudocapacitance which is progressively absent in higher current rate.

4.1.3.3 Investigation of Electrolytes Effect

![Graphs showing charge-discharge cycles for different electrolytes](image)

**Figure 4.13**: Galvanostatic charge-discharge cycles of the nano-porous $\alpha$-Fe$_2$O$_3$ fibers electrode at different current densities in (a) 1 M Na$_2$SO$_4$, (b) 1 M (NH$_4$)$_2$SO$_4$ and (c) 1 M Li$_2$SO$_4$ electrolytes. (d) Galvanostatic charge-discharge cycles of the nano-porous $\alpha$-Fe$_2$O$_3$ fibers electrode at 0.2 A g$^{-1}$. 

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To evaluate the electrochemical characteristics of $\alpha$-Fe$_2$O$_3$ nanofibers as electrode materials for supercapacitors, cyclic voltammetry and galvanostatic charge-discharge measurements were employed. The effect of different electrolytes on the electrochemical performance of $\alpha$-Fe$_2$O$_3$ nanofibers was investigated using aqueous 1 M Na$_2$SO$_4$, 1M (NH$_4$)$_2$SO$_4$ and 1 M Li$_2$SO$_4$ as electrolyte.

The galvanostatic charge-discharge performance of $\alpha$-Fe$_2$O$_3$ nanofibers in different electrolytes at various current densities (in two electrode configuration) are shown in Figure 4.13(a), (b) and (c). To better understand the effect of electrolyte on the electrochemical performance of, comparison of different electrolyte at same current density is depicted in Figure 4.13(d). It is clearly seen that the galvanostatic charge/discharge time is longest in Na$_2$SO$_4$ electrolyte, followed by (NH$_4$)$_2$SO$_4$ and Li$_2$SO$_4$. This implies that capacity is the highest in Na$_2$SO$_4$ electrolyte, followed by (NH$_4$)$_2$SO$_4$ and Li$_2$SO$_4$. Important factors influencing the capacity and hence capacitance include the charge and size of the individual electrolytes’ ions. The solvated cationic sizes of NH$_4^+$, Na$^+$, and Li$^+$ are 0.25 nm, 1.02 nm, 0.76 nm respectively. [115, 116] Accordingly, the conductivity of electrolytes can be arranged in decreasing order which is (NH$_4$)$_2$SO$_4$ > Li$_2$SO$_4$ > Na$_2$SO$_4$. System consisting of smaller ions is capable of up taking more surface area, hence, higher capacity can be derived. Also, smaller ions are able to move through the electrolyte more rapidly than larger ions, hence their conductivity is higher [10]. Therefore, the expected capacity of the various electrolytes in descending order of is (NH$_4$)$_2$SO$_4$ > Li$_2$SO$_4$ > Na$_2$SO$_4$. This contradicts with the result
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discussed above. To better understand this phenomenal, a three electrode cyclic voltammetry was conducted.

![Cyclic voltammogram](image)

**Figure 4.14**: Cyclic voltammogram of the nano-porous α-Fe$_2$O$_3$ fibers electrode in 1 M Na$_2$SO$_4$, 1M (NH$_4$)$_2$SO$_4$ and 1 M Li$_2$SO$_4$ electrolyte respectively.

The cyclic voltammogram recorded at the scan rate of 1 mV s$^{-1}$ in three electrode configurations are shown in Figure 4.14. It is clearly seen that an oxidation peak is obtained at +0.45 V vs SCE, coupled with corresponding reduction peak at +0.31V vs SCE in Na$_2$SO$_4$ electrolyte. This pair of redox peaks can be assigned to reversible pseudocapacitive behavior originates from the fast faradic reaction at the electrode
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surface. In the process, Na\textsuperscript{+} cations could have been adsorbed/desorbed to and from the surface of Fe\textsubscript{2}O\textsubscript{3}. No redox peak is seen in the other two electrolytes which suggest that such redox reaction is absent. Therefore, α-Fe\textsubscript{2}O\textsubscript{3} exhibits additional charge transfer reaction that is absent in the other electrolytes studied. Hence, the higher capacity observed in Na\textsubscript{2}SO\textsubscript{4} can be ascribed to this addition pair of redox peaks.

It is also noted that cyclic voltammogram are approximately rectangular, notwithstanding the redox peaks in Na\textsubscript{2}SO\textsubscript{4}, indicating good charge propagation in the electrodes. It is worth mentioning that an oblique symmetric cyclic voltammetry loop indicates a high contact resistance within the electrode, so only high conducting electrodes approximate an ideal rectangular geometry.

The specific capacitance ‘C\textsubscript{s}’ can be calculated from charge/discharge curves according to the following equation 4.1,

\[
C_s = \frac{2I}{m \frac{dv}{dr}}
\]  

(4.1)

Where, \(C_s\) is the specific capacitance, \(I\) is the applied current, \(m\) is the total mass of the active material and \(dv/dt\) is the slope of the discharge curve.
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Figure 4.15: (a) Specific capacitance of nano-porous α-Fe$_2$O$_3$ fibers at different current density in 1 M Na$_2$SO$_4$, 1M (NH$_4$)$_2$SO$_4$ and 1 M Li$_2$SO$_4$ electrolytes respectively. (b) Cycling behavior of nano-porous α-Fe$_2$O$_3$ fibers at different current density in 1 M Na$_2$SO$_4$, 1M (NH$_4$)$_2$SO$_4$ and 1 M Li$_2$SO$_4$ electrolytes respectively.

The values of specific capacitances calculated from galvanostatic discharge curves at different current densities are shown in Figure 4.15. The calculated specific
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capacitance values at constant current density of 0.02 A g\(^{-1}\) for different electrolytes was 316 F g\(^{-1}\), 115 F g\(^{-1}\) and 63 F g\(^{-1}\) in 1 M Na\(_2\)SO\(_4\), 1 M (NH\(_4\))\(_2\)SO\(_4\) and 1 M Li\(_2\)SO\(_4\) electrolytes, respectively. The observed difference of specific capacitance in all three electrolytes is ascribed to the different properties of electrolytes and varying redox mechanisms previously discussed. The highest values of specific capacitance in case of Na\(_2\)SO\(_4\) electrolyte is due to additional charge transfer reaction which is absent in the other electrolytes tested. It is also observed that differences in specific capacitance between the various electrolytes in more pronounced at lower current density. At high current density, the ions have lesser time to travel to electroactive sites in the nanoporous \(\alpha\)-Fe\(_2\)O\(_3\) nanofibers, thus fewer charges can be surface adsorbed. Additionally, the ions are less orderly arranged which further diminish charge storage capacity. Higher density also impedes the transportation of Na\(^+\) cations, resulting in lesser Na\(^+\) adsorption onto the surface of \(\alpha\)-Fe\(_2\)O\(_3\) nanofibers. This further support the conclusion made on origin of redox peak observed.
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4.1.3.4 Effect of Morphology

The surface morphology plays an important role in the characteristics of the pseudocapacitor devices, particularly the ability to store charges. The exposed surface area of the electrode determines the capacity, energy density and power density of the pseudocapacitors. Clearly, materials with higher surface area exhibit greater capacitances and better energy densities consistently.

Figure 4.16: FESEM image of Commercial α-Fe₂O₃

Figure 4.16 shows a FESEM image of commercial α-Fe₂O₃ powder. As compared to morphology depicted in Figure 4.8, commercial α-Fe₂O₃ powder contains agglomeration of particles of diverse shapes and sizes. This may result in lower utilization of active sites available for charge transfers.
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Figure 4.17: (a) Nitrogen adsorption-desorption isotherms of commercial $\alpha$-Fe$_2$O$_3$ at 77 K. (b) Derived BJH plot depicting pores sizes distribution of commercial $\alpha$-Fe$_2$O$_3$.

Figure 4.17 (a) depicts the nitrogen adsorption-desorption isotherms of commercial $\alpha$-Fe$_2$O$_3$ at 77 K and the derived BJH plot. Multipoint Brunauer-Emmett-Teller (BET) analysis performed on $\alpha$-Fe$_2$O$_3$ nanofibers yields a specific surface area of 4.98 m$^2$g$^{-1}$. This value is sustainably lower than the BET value obtained for FNF which was 33.95. This may have an impact of the resultant specific capacitance and hence the energy density of the $\alpha$-Fe$_2$O$_3$. Derived BJH from the desorption isotherm is shown in Figure 4.17 (b). It is observed that the BJH analysis did not yield fruitful result on the pore size distribution of the commercial $\alpha$-Fe$_2$O$_3$ powder. The pores size analyzed lies outside the limit of nitrogen adsorption-desorption isotherm. However, it can be concluded that the pore sizes are either less than 3 nm or more than 100 nm or both. Therefore, it is clear that the $\alpha$-Fe$_2$O$_3$ powder is not mesoporous.
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Figure 4.18: (a) Galvanostatic charge-discharge cycles of the α-Fe₂O₃ nanofibers and commercial α-Fe₂O₃ electrode. (b) Plot of specific capacitance with current density.

Figure 4.18(a) depicts the GCD of α-Fe₂O₃ nanofibers and commercial α-Fe₂O₃ powder. Clearly, the electrospun α-Fe₂O₃ nanofibers have exhibited longer charge-discharge time profile at the same current density. The relationship between specific capacitance and current density was also investigated. The specific capacitance ‘Cₛ’ can be calculated from charge/discharge curves at various current densities according to equation 4.1. Figure 4.18 (b) shows the relationship of specific capacitance of both α-Fe₂O₃ nanofibers and commercial powder. It is observed that the α-Fe₂O₃ nanofibers have higher specific capacitance than the α-Fe₂O₃ powder which can be ascribed to the smaller particle’s size and higher surface area as evident from FESEM images and BET data previously discussed. Also, α-Fe₂O₃ nanofibers being more porous allows better traffic of ions to and from the electrode surface, thereby enabling better utilization of active sites for charge transfer.
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4.1.3.5 Charge Storage Mechanism XPS Studies

To better understand the charge storage mechanism, X-ray photoelectron spectroscopy (XPS) was employed. Figure 4.19 shows the XPS spectra of FNF electrodes after charging to 0.6 V and cycling for 1000 cycles. The Fe 2p$_{3/2}$ peak is greater than that of Fe 2p$_{1/2}$, a result of spin-orbit (j-j) coupling. The main peak of 2p$_{3/2}$ is located at 710.7 eV, in agreement with literature [117-119]. An accompanying satellite peak is observed at approximately 8 eV higher than the main Fe 2p$_{3/2}$ peak is clearly distinguishable and does not overlap with either the Fe 2p$_{1/2}$ or Fe 2p$_{3/2}$ peaks [117-119]. This indicates the presence of Fe (III) oxidation state. Therefore, the change of oxidation states of FNF during galvanostatic charging and discharging process, an indication of the redox process was not evidenced by XPS analysis. However, additional peaks of Na 1s
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were detected in the survey scan of FNF after cycling, as shown in Figure. 4.19 (b), suggesting the intercalation of Na$^+$ ions into the oxide lattice of Fe$_2$O$_3$. This agrees well with the three electrode CV test previously discussed in section 4.1.3.2. It is therefore clear that intercalation of Na$^+$ ions process rather than redox process contributes significantly to the charge storage mechanism of FNF.
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4.1.3.6 Performance of Symmetrical Fe₂O₃ Nanofibers Supercapacitor

To characterize the performance of α-Fe₂O₃ nanofibers as electrode material for energy storage device, energy density and average power density evaluated from galvanostatic charge-discharge curves in a voltage window of 0 to 1 V according to Equation 4.2 and 4.3.

\[ E_D = \frac{1}{2} C_D V_D^2 \]  \hspace{1cm} (4.2)

\[ P_D = \frac{E_D}{t} \]  \hspace{1cm} (4.3)

where \( E_D \) is the energy density of the device, \( C_D \) is the specific capacitance of the device, \( V \) is the maximum charging voltage, and \( t \) is the time taken to fully discharge.

Figure 4.20: Ragone plot for the nano-porous α-Fe₂O₃ fibers electrode in 1 M Na₂SO₄, 1M (NH₄)₂SO₄ and 1 M Li₂SO₄ electrolyte respectively.
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The energy and power density is summarized as a Ragone plot in Figure 4.20. The \( \alpha-\text{Fe}_2\text{O}_3 \) nanofibers have exhibited both higher power and energy density in \( \text{Na}_2\text{SO}_4 \) electrolyte. This is ascribed to relatively small cationic size of \( \text{Na}^+ \) and the additional pseudocapacitive behavior of \( \alpha-\text{Fe}_2\text{O}_3 \) in \( \text{Na}_2\text{SO}_4 \) solution previously discussed. The \( \alpha-\text{Fe}_2\text{O}_3 \) nanofibers also have better performance in \( (\text{NH}_4)_2\text{SO}_4 \) as compared to \( \text{Li}_2\text{SO}_4 \) solution due to the smaller cationic size of \( \text{NH}_4^+ \). Highest power density of 0.02 kW/kg and maximum energy density of 9 Wh/kg were achieved in \( \text{Na}_2\text{SO}_4 \) electrolyte.

![Graph showing cyclic performance and coulombic efficiency](image)

**Figure 4.21**: Cyclic performance and coulombic efficiency of the nano-porous \( \alpha-\text{Fe}_2\text{O}_3 \) fibers electrode in 1M \( \text{Na}_2\text{SO}_4 \) at constant current density of 0.02 A g\(^{-1}\).
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Long term cycle stability of electrode material is an important factor to consider in for selection of electrode material for in supercapacitor application. The electrochemical stability is examined by galvanostatic charge-discharge cycling at a constant current density of 0.02A g\(^{-1}\). Figure 4.21 shows the cycling performance of Fe\(_2\)O\(_3\) nanofibers electrodes in two electrode device configuration. The specific capacitance greatly increases in the first 200 cycles, indicating that the Fe\(_2\)O\(_3\) nanofibers were undergoing activation to reach its maximum capacitance. Thereafter, the specific capacitance stabilized around 300 F g\(^{-1}\). The capacitance retention remains as high as 97% at 1000\(^{th}\) cycle, indicating that α-Fe\(_2\)O\(_3\) nanofibers have good electrochemical stability. The columbic efficiency is also close to 99% during cycling, suggesting that no gas evolution occurred in the voltage region.

4.1.3.7 Electrochemical Impedance Spectroscopy (EIS) Studies

EIS is a useful technique to resolve the charge storage components resulting from different mechanism such as EDL or pseudocapacitance as well as different resistance components such as electrolyte resistance, electrode resistance and interfacial electrode-electrolyte resistance. Hence, EIS is used to investigate the electrochemical behavior. The Nyquist plots for FNF supercapacitor device before and after cycling were recorded and are shown in Figure 4.22.
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An intersection observed at the real axis in the high frequency region, followed by a non-vertical slope in low frequency region. This can be explained by two reasons: (1) an interface exists between the electrode and the electrolyte, and (2) a double-layer capacitance connects in parallel with a resistance and leads to the formation of a RC loop [120]. The transformation of the plot to a vertical line takes place at low frequency can be mainly ascribed to the capacitive response of porous carbons. The presence of the non-vertical slope at the low frequency region of the Nyquist plot may be associated with low electronic conductivity of the electrode and different pore size distribution of the carbons [120].
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To better investigate on the electrochemical behavior, an equivalent circuit simulating FNF electrodes is proposed and shown in Figure 4.23. The circuit would involve the following elements, namely, electrolyte resistance, \( R_b \); double layer capacitance, \( C_{dl} \); pseudocapacitance, \( C_p \); interfacial charge transfer resistance, \( R_{ct} \); and Warburg impedance, \( W_o \) due to distributed resistance within the pores of FNF. The experimental data was fitted well with the proposed model in Figure 4.23. The equation equivalent of the model is given by equation 4.4.

\[
Z = R_s + \left( \frac{1}{(j\omega)\alpha C_{dl}} + \frac{1}{R_{ct} + (j\omega C_{pc})^{-1}} \right)^{-1} + R_s \frac{\text{ctnh} (jW_{r} \omega)^{\alpha r}}{(jW_{r} \omega)^{\alpha r}} \tag{4.4}
\]

where \( R_s \) is the electrode resistance, \( C_{dl} \), \( \omega \) and \( \alpha \) are the capacitance of the distributive system, the angular frequency, an exponent that determines the deviation of the capacitance from ideality due to a distribution of relaxations, \( R_W, W_T \), are the diffusion resistances, related to both the diffusion length and the diffusion coefficient, and \( W_P \), the exponent that measures its deviation from ideality, \( j \) is \( \sqrt{-1} \).
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By fitting equation 4.4 to the respective impedance data depicted in Figure 4.22, the values shown in Table 4-1 have been evaluated.

<table>
<thead>
<tr>
<th>Cycle</th>
<th>$R_s(\Omega)$</th>
<th>$C_{dl}(\text{mF cm}^{-2})$</th>
<th>$R_{ct}(\Omega)$</th>
<th>$C_p(\text{mF cm}^{-2})$</th>
<th>$R_w(\Omega)$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial</td>
<td>3.69</td>
<td>21.86</td>
<td>1.162</td>
<td>12.93</td>
<td>225.2</td>
</tr>
<tr>
<td>200th</td>
<td>3.49</td>
<td>10.14</td>
<td>21.97</td>
<td>219.75</td>
<td>1360</td>
</tr>
</tbody>
</table>

Table 4-1: Components of the equivalent circuit fitted for the impedance spectra of $\alpha$-$\text{Fe}_2\text{O}_3$ nanofibers symmetrical device setup in Figure 4.22.

Equation 4.4 in the case here can be simply categorized into three frequency regions. The real axis intersection at the sufficiently high frequency provides the electrolyte resistance, while the mid high frequency informs about the charge transfer resistance. And the low frequency region encompasses most or all the electrochemical activities.

The electrolyte resistance of the electrodes is negligible, indicating that the electrochemical properties of bulk electrolytes remain stable. Both greater charge transfer resistance $R_{ct}$ and higher pseudocapacitance $C_p$ are obtained after undergoing galvanostatically charged and discharged for 200 cycles. This indicates a greater amount of faradic current, a result of more charge transfer reactions taking place. This agrees with the explanation discussing that FNF electrodes have undergone an activation process which enables better accessibility of electrolyte ions to its pores.
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4.1.4 V$_2$O$_5$ Nanofibers Supercapacitors

Vanadium oxides have been investigated for their potential use in energy storage applications, particularly for lithium ion batteries, due to the availability of multiple oxidation states [121]. Herein, V$_2$O$_5$ nanofibers are synthesized via electrospinning route and characterized.

4.1.4.1 Physical Characterization of V$_2$O$_5$ Nanofibers

Figure 4.24 illustrates typical X-ray diffraction (XRD) patterns of electrospun VNF annealed at 470 °C for 30 min in the range of 2θ from 10 to 60°. Reproduced by permission of ECS - The Electrochemical Society.

Figure 4.24: XRD patterns of electrospun VNF annealed at 470 °C for 30 min in the range of 2θ from 10 to 60°. Reproduced by permission of ECS - The Electrochemical Society.

Figure 4.24 illustrates typical X-ray diffraction (XRD) patterns of electrospun VNF after annealing at 470 °C for 30 min. The XRD pattern of VNF revealed highly intense reflections that corresponded to orthorhombic Shcherbinaite structure with $Pmn2_1$ space group (JCPDS card no. 41-1426). Lattice parameters of VNF calculated from
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Rietveld crystal structure refinement was found to be $a = 11.513 \pm 2$ Å, $b = 3.5656 \pm 2$ Å
and $c = 4.3727 \pm 3$ Å.

![Field emission scanning electron microscopic (FE-SEM) images of electrospun VNF, (b) Electrospun VNF annealed at 470 °C for 30 min.](image)

Field emission scanning electron microscopic (FE-SEM) images of as-spun VNF shown in Figure 4.25(a) (with the inset showing the magnified view of the respective images) revealed smooth surface of the fibers with diameters ranging from 300 to 500 nm and fiber length over 200 μm. As spun VNF were annealed at various temperatures between 300 and 500 °C to remove the polymer and to crystallize $V_2O_5$ nanocrystals in the VNF. Heating at higher temperature ($> 500$ °C) leads to destruction of the fibrous structure, while at lower temperature (300-400°C) resulted in incomplete removal of PVP polymer. The fibers heat treated at optimum temperature around 470 °C retained the fibrous morphology with well-defined crystalline structure. Figure 4.23(b) illustrates the VNF after annealing at 470 °C for 30 minutes. The annealed VNF exhibited a more spiral
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like structure, with diameter of 300-500 nm, lower than that of as spun VNF. This change in fibrous morphology is mainly attributed to the decomposition of PVP polymer during sintering. Vaporization of the polymer during this process left behind pores and the surface seems to have a spiral like morphology, which allows facile movement of electrolyte during the electrochemical reaction. It is worth noting that, sintering of as spun VNF was conducted at relatively low temperature (~470 °C) to achieve a high performance V_{2}O_{5} electrode.

Figure 4.26: (a) Nitrogen adsorption-desorption isotherms of VNF at 77 K. (b) BJH plot depicting pores sizes distribution of VNF.

The nitrogen adsorption/desorption isotherms of both VNF at 77 K presented in Figure 4.26 (a) exhibited type II isotherm (according to IUPACS classification [122]) with type H3 hysteresis showing absence of limiting adsorption at high relative pressure.
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for both adsorption and desorption isotherm. Hysteresis loop extending down to low pressure region was observed indicating the presence of micropores which is advantageous for double layer charge storage. Desorption isotherm of PANF shows hysteresis loop at the relative pressure of 0.8–0.9, suggesting meso-porosity (high surface area) which is beneficial for the movement of ions throughout the nanofibers. The gradual increase in adsorption curves of both VNF in the middle relative pressure region can be attributed to multi-layer adsorption of nitrogen in the mesopores and macropores. The leap (plunge) close to the saturation pressure in adsorption (desorption) isotherm suggests pores filling of large mesopores and macropores which is favourable as they allow ions to amass in the proximity of electrode permitting high power capability.

Brunauer-Emmett-Teller (BET) analysis [123] of the isotherms yields specific surface area of 9.5 m² g⁻¹ for VNF. The obtained values are consistent with the previous reports on such material (VNF: 10 m² g⁻¹) [92, 124]. To investigate the pore size distribution, Barrett-Joyner-Halenda (BJH) analysis was applied on the desorption isotherms [125]. Figure 4.24 (b) depicts the pore volume distribution curves obtained by BJH analysis. Much of the surface area can be attributed to pores from the micropores range (<2 nm, IUPAC classification) [126]. These micropores are readily accessible to the hydrated K⁺ (0.3 nm) and Cl⁻ (0.3 nm) ions and would certainly enhance both the electrical double layer and pseudocapacitive contribution of the nanofibers [115].
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4.1.4.2 Chemical Characterization of V$_2$O$_5$ Nanofibers

![Cyclic Voltammogram](image)

**Figure 4.27:** Three electrode cyclic voltammograms of VNF at 10 mV s$^{-1}$ vs. SCE.

Three electrodes setup was utilized to study the electrochemical stability of VNF electrodes, in which platinum sheet and standard calomel electrode (SCE) were used as counter and reference electrodes, respectively. Figure 4.27 shows the three electrode CV curve of VNF electrode. The CV curve of VNF electrode shows good stability under positive and negative polarization with reversibility from 0.2 $-$ 1 V vs. SCE. These results illustrate that VNF is a potential candidate for negative electrode.
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4.1.4.3 Charge Storage Mechanism XPS Studies

![Graphs showing XPS spectra](image)

**Figure 4.28**: (a) high resolution XPS spectra of V 2p peaks (b) XPS survey spectra of V$_2$O$_5$ nanofibers composite.

Figure 4.28(a) shows the high resolution XPS spectra of VNF electrodes after charging to 0.9 V and cycling for 1000 cycles. The spin–orbit splitting obtained for V 2p$_{3/2}$ and V 2p$_{1/2}$ is 7.75 eV, confirming the existence of V (5+) oxidation state and hence, the presence of V$_2$O$_5$ [127] This suggests that the change in oxidation states of VNF during galvanostatic charging and discharging process, an indication of the redox process was not evidenced by XPS analysis. However, additional peaks of K 2s, K 2p$_{1/2}$ and K 2p$_{3/2}$ were detected in the survey scan of VNF after cycling, as shown in Fig. 4.28 (b), which suggest the intercalation of K$^+$ ions into the oxide lattice of V$_2$O$_5$. It is therefore clear that intercalation of K$^+$ ions process rather than redox process contributes significantly to the charge storage mechanism of VNF.
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4.1.4.4 Performance of Symmetrical V$_2$O$_5$ Nanofibers Supercapacitor

Figure 4.29: (a) CV curves for annealed electrodes measured at various scan rates in 3 M KCl in two-electrode symmetric configurations. (b) Galvanostatic charge-discharge curves for annealed VNF electrodes measured at various current densities under the same conditions. (c) Cycling behavior of VNF-VNF cell. Reproduced by permission of ECS-The Electrochemical Society.

Cyclic voltammetry (CV) and galvanostatic charge-discharge studies have been employed to evaluate the supercapacitive behaviour of electrospun VNF in symmetric device configuration. Figure 4.29 (a) illustrates the CV traces of VNF-VNF symmetric
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device as a function of scan rate with respect to its current density in 3 M KCl aqueous solution. It can be observed that charging and discharging processes in VNF electrode are relatively fast. The current which depicts the direction of the charge flow, showed some delay upon the reversal of the potential sweep, suggesting diffusional resistance for the VNF-VNF symmetrical device possibly due to the diffusion process of ions through the pores.

Galvanostatic charge-discharge measurements were carried out for VNF symmetric configurations in 3 M KCl electrolyte and given in Figure 4.29 (b). The symmetry of the charge-discharge curves indicates good capacitive behaviour in symmetric configurations. In such configuration, specific capacitance as high as 213 F g\(^{-1}\) was obtained for VNF electrodes, respectively. The obtained values for VNF electrodes in aqueous medium are also consistent with the previous reports [92]. Such high capacitance can be attributed to the high surface area provided by the porous and fibrous structure of VNFs, which eased the diffusion of ions into the pores thus contributing to the enhanced pseudocapacitance.
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4.1.5 Polyaniline Nanofibers Supercapacitors

Electrically conducting polymers make good pseudocapacitive material for electrode of supercapacitors. They possess good electronic conductivity, have low cost, and adjustable redox activity through chemical modification.[37, 38] They are capable of pseudocapacitive storage through the redox process. Ions are transferred to the polymer backbone when oxidation occurs and vice versa. The processes have good reversibility as the redox reactions do not alter the structures.[39]

Pseudocapacitors employing electronically conducting polymer such as polyaniline [40, 46, 128, 129], polypyrrole [38, 40, 43, 45] and poly(3,4-ethylenedioxythiophene) (PEDOT) [49, 130] derivatives have been reported. Different methods used to fabricate polyaniline nanofibers include electrospinning [86], hard templates, soft template, interfacial polymerization [18], and rapid polymerization [131]. In particular, interfacial polymerization and rapid polymerization have achieved more attention due to its easiness, large-scale production-ability and environmentally benign nature. Furthermore, it has a capability to produce high-quality polyaniline nanofibers with control of their morphology, size and diameter [18]. Herein, rapid chemical polymerization route is used to synthesize polyaniline nanofibers. Also, the performance of symmetrical PANF was characterized which will serve as a comparison for asymmetrical device employing PANF electrode in later section.
4.1.5.1 Physical Characterization of Polyaniline Nanofibers

Figure 4.30: XRD patterns of PANF and electrospun VNF annealed at 470 °C for 30 min in the range of 2θ from 10 to 60°. Reproduced by permission of ECS- The Electrochemical Society.

Figure 4.30 illustrates typical X-ray diffraction (XRD) patterns of electrospun VNF after The XRD pattern of PANF revealed several broad reflection peaks (2θ) at 15°, 20.6°, 25.2° and, 26.8° which confirms the formation of polyaniline by rapid polymerization procedure. These reflections indicate the presence of crystalline order in the bulk PANF samples, which is consistent with literature [132].
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Figure 4.31: (a) FESEM image of PANF by rapid chemical polymerization. (b) TEM image of a single PANF. Reproduced by permission of ECS - The Electrochemical Society.

Figure 4.31 displays the morphological features of the PANF, showing interwoven structure with shorter fibers compared to electrospun VNF. The PANF fiber length and diameter PANF estimated from FE-SEM images with length of ca. 100-600 nm and diameter ranging from 30-200 nm. Transmission electron microscopy (TEM) was conducted to study the morphology of PANF (inset Figure 4.31) clearly depicts nanostructured smooth surface of PANI.
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Figure 4.32: (a) Nitrogen adsorption-desorption isotherms of PANF at 77 K. (b) BJH plot depicting pores sizes distribution of PANF. Reproduced by permission of ECS - The Electrochemical Society.

The nitrogen adsorption/desorption isotherms of both PANF at 77 K presented in Figure 4.32 (a) exhibited type II isotherm (according to IUPACS classification [122]) with type H3 hysteresis showing absence of limiting adsorption at high relative pressure for both adsorption and desorption isotherm. Hysteresis loop extending down to low pressure region was observed indicating the presence of micropores which is advantageous for double layer charge storage. Desorption isotherm of PANF shows hysteresis loop at the relative pressure of 0.8–0.9, suggesting micro-porosity (high surface area) which is beneficial for the movement of ions throughout the nanofibers. The gradual increase in adsorption curves of PANF in the middle relative pressure region can be attributed to multi-layer adsorption of nitrogen in the macropores. The leap (plunge) close to the saturation pressure in adsorption (desorption) isotherm suggests pores filling of large macropores which is favourable as they allow ions to amass in the proximity of
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electrode permitting high power capability.

Brunauer-Emmett-Teller (BET) analysis [123] of the isotherms yields specific surface area of 36.8 m$^2$ g$^{-1}$ for PANF. The obtained values are consistent with the previous reports on such material (PANF: 34.4 m$^2$ g$^{-1}$) [92, 124]. To investigate the pore size distribution, Barrett-Joyner-Halenda (BJH) analysis was applied on the desorption isotherms [125]. Figure 4.32 (b) depicts the pore volume distribution curves obtained by BJH analysis. Much of the surface area can be attributed to pores from the micropores range (<2 nm, IUPAC classification) [126]. These micropores are readily accessible to the hydrated K$^+$ (0.3 nm) and Cl$^-$ (0.3 nm) ions and would certainly enhance both the electrical double layer and pseudocapacitive contribution of the nanofibers [115].

4.1.5.2 Chemical Characterization of Polyaniline Nanofibers

![Three electrode cyclic voltammograms of PANF at 10 mV s$^{-1}$ vs. SCE.](image)

Figure 4.33: Three electrode cyclic voltammograms of PANF at 10 mV s$^{-1}$ vs. SCE.
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A three electrode electrochemical cell setup was also utilized to study the electrochemical stability of PANF electrodes, in which platinum sheet and standard calomel electrode (SCE) were used as counter and reference electrodes, respectively. To use it as a possible candidate in an asymmetrical supercapacitor, we tested PANF in 3 M KCl. It is worthy to mention that PANF works better in H₂SO₄ [133], however such acid media is not suitable for VNF nor FNF. Figure 4.30 shows the three electrode CV curve of PANF electrode. The CV curve of PANF electrode shows good stability under positive and negative polarization with reversibility from 0.2 –1 V vs. SCE. These results illustrate that PANF is a good candidate for positive electrode.

4.1.5.3 Performance of Symmetrical Polyaniline Nanofibers Supercapacitor

![Graphs showing CV curves and galvanostatic charge-discharge curves for PANF electrodes.]

Figure 4.34: (a) CV curves for PANF electrodes measured at various scan rates in 3 M KCl in two-electrode symmetric configurations. (b) Galvanostatic charge-discharge curves for PANF electrodes measured at various current densities. Reproduced by permission of ECS - The Electrochemical Society.
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Galvanostatic charge-discharge measurements were carried out for PANF-PANF symmetric configurations in 3 M KCl electrolyte and given in Figure 4.34 (a) and (b). The symmetry of the charge-discharge curves indicates good capacitive behaviour in symmetric configurations. In such configurations the specific capacitance as high as 176 F g\(^{-1}\) was obtained for PANF electrodes. The obtained values for PANF electrodes in aqueous medium are also consistent with the previous reports [129, 134]. The high capacitance can be attributed to the high surface area provided by the porous and fibrous structure of PANFs, which eased the diffusion of ions into the pores thus contributing to the enhanced pseudocapacitance. A relatively lower specific capacitance obtained for PANF electrodes as compared to those in acidic media like H\(_2\)SO\(_4\) could be attributed to the absence of reversible doping process of PANFs which usually occurred in acidic media [133].
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4.2 Asymmetrical Supercapacitors

Pseudocapacitor electrodes based on either electronically conducting polymers (ECP) (polyaniline (PANI) [128, 129], poly(3,4-ethylendioxythiophene) (PEDOT) [130], and polypyrrole (PPy) [38]) or transition metal oxides (TMO) [135] RuO$_2$ [58, 136], NiO$_x$ [137], Co$_3$O$_4$ [138] and V$_2$O$_5$ [92, 139] have been extensively investigated as electrode materials for supercapacitor applications [140]. Pseudocapacitor based electrodes often provide higher specific capacitance as the charge storage redox mechanism involves the entire electrode active mass as compared to EDLC carbon electrodes wherein primarily only the electrode surface contributes to the specific capacitance via double layer charge formation.

Energy density of supercapacitor is proportional to square of voltage ($E = \frac{1}{2}CV^2$), where $C$ is the specific capacitance (Fg$^{-1}$) and $V$ is the voltage window (Volts). In aqueous electrolytes, both EDLC and pseudocapacitors have lower working voltages (<1.2 V) which severely discounts their energy density. Although pseudocapacitors exhibit higher specific capacitance and hence greater energy density, such increment is often inadequate and cyclability is also questionable.

One approach to further increase the energy density is to widen the working voltage (~2-3 V) of both EDLC and pseudocapacitor through adoption of organic-based (such as acetonitrile) electrolytes instead of aqueous solutions [3]. However, such aprotic organic solvents not only entail detrimental costs from fabrication processes requiring water free environment, but also possesses safety and toxicity issues. Another approach
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to increase the working voltage is to construct an asymmetric supercapacitor (ASC) in an aqueous medium by employing two different kinds of electrodes. Employing electrodes of different nature in ASC for positive and negative polarization allows both electrodes to operate in an optimal voltage range [94, 135, 141-144]. It has been reported that ASC consisted of TMO-AC[68, 82, 135, 145] or ECP-AC[9, 146] pair is capable of relatively higher operating voltage.

Among the transition metal oxides, amorphous hydrous ruthenium oxide (RuO$_2\times$H$_2$O) exhibits pseudocapacitive behaviour with large specific capacitance, but it is too expensive for the commercialization. V$_2$O$_5$ is one of the attractive candidates because of its layered structure, multiple oxidation states of vanadium which allows surface/bulk redox reactions leading to high capacitance in symmetric supercapacitor configuration [92]. Among the conducting polymers, polyaniline (PANI) is considered as a promising electrode materials for supercapacitors owing to its environmental stability, controllable electrical conductivity multiple intrinsic redox states and excellent processability [147].

Nanostructured active materials are capable of providing higher power density than bulk counterparts owing to shorter diffusion pathways, increased accessible surface area leading to facile flow of ions at the electrode/electrolyte interface. Electrospinning is a simple, low cost scalable facile route for making high aspect ratio 1D nanofibers with controllable diameter using electrostatic force [148]. Also, electrospinning often results in highly porous interconnected three dimensional fibrous electrode that provides both a continuous conduction pathway and increased electrode/electrolyte contact surface [149]. Such attributes are crucial to determine the performance of supercapacitors.
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As aforementioned, we have successfully synthesized and fabricated symmetrical supercapacitors from CNT, FNF, PANF and VNF. Their three electrode performance was also characterized and discussed.

Herein, we have constructed an ASC based on a combination of nanostructured TMO and ECP to study the capacitive properties and widen the operating voltage in aqueous medium. In the present work, an attempt has been made to construct an ASC comprising of electrospun vanadium pentoxide nanofibers (VNF) as negative electrode and polyaniline nanofibers (PANF) as positive electrodes in aqueous medium [108]. A schematic illustration of the ASC utilizing electrospun VNF and PANF is shown in Figure 4.36. The structure, surface morphology and diameter of the electrospun fibers was controlled by varying the parameters like applied potential, precursor concentration, viscosity and flow rate of the solution. The detailed electrochemical studies of both VNF and PANF electrodes are conducted and analyzed in both symmetric and asymmetric configurations and described in detail.
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**Figure 4.35**: Schematic illustration of an asymmetrical supercapacitor comprising of VNF and PANF as positive and negative electrodes, respectively. Reproduced by permission of ECS - The Electrochemical Society.

In order to fabricate VNF and PANF electrodes in asymmetric configuration (ASC), a common aqueous electrolyte needs to be utilized. Three electrodes setup was utilized to study the electrochemical stability of individual electrodes, in which platinum sheet and standard calomel electrode (SCE) were used as counter and reference electrodes, respectively. A table summarizing the respective electrode materials, electrolytes and their corresponding electrochemical window is depicted in Table 4-2.
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Table 4-2: Table of electrode material and their respective potential window in various electrolytes.

<table>
<thead>
<tr>
<th>Electrode Material</th>
<th>Electrolyte</th>
<th>Potential Window (vs SCE)</th>
<th>Potential Electrode Candidate</th>
</tr>
</thead>
<tbody>
<tr>
<td>P3-CNT</td>
<td>H₂SO₄</td>
<td>0 to 0.8 V</td>
<td>Positive or Negative</td>
</tr>
<tr>
<td></td>
<td>KCl</td>
<td>0 to 0.8 V</td>
<td>Positive or Negative</td>
</tr>
<tr>
<td>FNF</td>
<td>Na₂SO₄</td>
<td>0 to 0.65 V</td>
<td>Positive</td>
</tr>
<tr>
<td></td>
<td>Li₂SO₄</td>
<td>0 to 0.65 V</td>
<td>Positive</td>
</tr>
<tr>
<td></td>
<td>(NH₄)₂SO₄</td>
<td>0 to 0.65 V</td>
<td>Positive</td>
</tr>
<tr>
<td>VNF</td>
<td>KCl</td>
<td>-1 to 0.2 V</td>
<td>Negative</td>
</tr>
<tr>
<td>PANF</td>
<td>KCl</td>
<td>-0.2 to 1 V</td>
<td>Positive</td>
</tr>
</tbody>
</table>

From Table 4-2, it would be a good choice to assemble an asymmetrical supercapacitor using VNF and PANF as the negative and positive electrode, respectively. Although PANF results in higher capacitance in aqueous electrolyte like H₂SO₄ [140], VNF readily dissolved in such acidic media. Hence, KCl was chosen as the optimum electrolyte after several failed electrochemical trials of VNF-PANF in various aqueous electrolytes including H₂SO₄ and NaOH.
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4.2.1 Electrochemical Characterization of Asymmetrical Supercapacitor

Figure 4.36: Three electrode cyclic voltammograms VNF and PANF at 10 mV s\(^{-1}\) vs. SCE. Reproduced by permission of ECS - The Electrochemical Society.

Figure 4.36 shows the three-electrode CV of both VNF and PANF extracted from Figures 4.27 and 4.33. These CV curves reveal that PANF electrode exhibited high electrochemical stability under positive polarization, giving a high electrochemical reversibility up to +1 V vs. SCE, whereas VNF electrode shows stability under negative polarization with reversibility of -1 V vs. SCE. These results illustrate that VNF is more stable being a negative electrode, while PANF is more stable as a positive electrode. Hence, a working voltage window of 2 V (\(E_{PANF} - E_{VNF} = 1.0\ V - [-1.0\ V] = 2\ V\)) is viable if the two materials are judiciously paired in an asymmetric system with 3 M KCl as the
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common electrolyte. Generally, the aqueous system is stable up to ~1.23 V and said potential electrolysis takes place. However, in the asymmetric combination provides wider operating potential up to 2 V. This extension of the working potential in the aqueous medium is mainly due to the combination of the high hydrogen overpotential offered by VNF under negative polarization together with high oxygen overpotential when PANF is used as positive electrode [145].

Another factor to be considered is the optimization of desired capacity of such asymmetric configuration which is crucial to achieve high energy density. It is important to note that in a symmetric configuration, the applied voltage is split equally between the two electrodes. However, in asymmetric system where the two electrodes are different the voltage will be split depending on the capacitance of individual electrode material. The capacitance of each electrode is related to the respective mass and intrinsic pseudocapacitance. Mass balances between the two electrodes are needed to optimize the cell voltage of the asymmetric capacitor. The capacity balance will enlist the conservation of charges (i.e. \( q^+ = q^- \)) and can be obtained via the following equation,

\[
\frac{m_+}{m_-} = \frac{C_+ \times \Delta E_-}{C_- \times \Delta E_+}\quad (4.7)
\]

where \( m_+ \), \( m_- \), \( C_+ \), \( C_- \), \( \Delta E_+ \), and \( \Delta E_- \) are the respective mass, specific capacitance and potential windows of three electrodes configuration of the individual positive and
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negative electrode. Optimization of the mass ratio was done based on Equation 4.4 to be 1.2:1 for PANF : VNF.

**Figure 4.37**: (a) Cyclic voltammograms (CV) of two-electrode cell of VNF and PANF at 10 mV s\(^{-1}\) in the presence of 3 M KCl electrolyte solution with different operating potentials (b) CV traces of an asymmetrical PANF-VNF device measured between 0-2 V at various scan rates ranging from 2-100 mV s\(^{-1}\). (c) Galvanostatic charge-discharge curves of an asymmetrical PANF-VNF device measured between 0-2 V at various current densities. Reproduced by permission of ECS - The Electrochemical Society.
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Figure 4.37(a) shows the CV curves of VNF-PANF asymmetric device in two electrode configuration measured at different potential windows as shown in the figure at 10 mV s\(^{-1}\). In an aqueous medium, a CV curve of less oblique rectangular geometry is noted for PANF-VNF asymmetric device with working voltage below 2 V. With higher working voltage, the overall energy storage capability of the asymmetrical device is greatly enhanced. However, an undesirable drastic current surge is observed if working voltage exceeds 2.5 V which may be attributed to hydrogen evolution at the negatively polarized VNF electrode. Thus, we chose a potential window of 2 V in our subsequent experiments to investigate the overall electrochemical performance of the asymmetrical device.

The CV of PANF-VNF asymmetric device, measured in two-electrode configuration between 0-2 V at various scan rates ranging from 2 to 100 mV s\(^{-1}\) is depicted in Figure 4.37(b). This figure indicates a fast charge/discharge capability at the measured scan rate. The CV curves resemble an ideal rectangle at low scan rate, and exhibit a slightly distorted trapezium as scan rate increases. Higher deviation is observed at higher scan rate which may be due to both kinetically limited charge storage process and the resistivity of the electrode, which will be verified by EIS measurement in later section. It is worth mentioning that the specific capacitance was calculated based on the total mass of active material instead of that of a single electrode. A device capacitance of 41 F g\(^{-1}\) is obtained at 5 mV s\(^{-1}\) which is slightly lower capacitance than VNF-VNF and PANF-PANF symmetric configurations. However, such asymmetrical configuration enables a working voltage up to 2 V, thereby enhancing energy density according to
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Equation 4.2.

Galvanostatic charge-discharge studies were conducted for VNF-PANF asymmetric device with different current densities. Figure 4.37(c) depicts the typical charge-discharge curves of a PANF-VNF asymmetric supercapacitor. The symmetrical charging and discharging characteristics indicates good capacitive behaviour of ASC.

4.2.2 Comparison of Symmetrical and Asymmetrical Configuration on Supercapacitor Performance

![Graph](image)

**Figure 4.38**: Plot of specific capacitance vs. current density. Reproduced by permission of ECS - The Electrochemical Society.

Figure 4.38 summaries the specific capacitance calculated from galvanostatic charge-discharge curves for both symmetric (VNF-VNF and PANF-PANF) and asymmetric configuration (VNF-PANF). For symmetrical configuration, galvanostatic values are extracted from Figures 4.29 and 4.34, while for asymmetrical configuration,
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values are extracted from Figure 4.37. In general, the specific capacitance gradually decreases with higher current density due to a larger IR drop at high current density. Decrease in specific capacitance values were observed with increasing current density. As charge/discharge rate increases, fewer ions are able to diffuse into the sub-surface resulting in less numbers of redox reactions, which leads to reduced charge storage. Nevertheless, the promising results from both CV and galvanostatic charge-discharge measurements of VNF-PANF asymmetric system indicates a higher working voltage, good pseudocapacitive properties.

![Figure 4.39: A Ragone plot of symmetrical and asymmetrical devices.](image)

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Energy and power densities are used to characterize the performance of energy storage devices. To demonstrate the electrochemical performance of our device, energy discharged and average power densities were evaluated from galvanostatic charge-discharge curves according to Equations (4.2) and (4.3).

Figure 4.39 shows the Ragone plot of both symmetrical and asymmetrical devices. VNF-PANF supercapacitor exhibited much higher energy and power density than its symmetrical configuration counterparts. This is ascribed to the synergistic effect endowed by the combination of PANF and VNF namely; (1) the nanostructure of both PANF and VNF that facilitates diffusion of electrolyte ions and enables higher capacitance; (2) the asymmetrical configuration that widens the operating potential window thus enabling redox process from different oxidations states of vanadium.
4.2.3 Effect of Morphology on Asymmetrical Supercapacitor

![Graph showing power density vs. time](image)

**Figure 4.40**: (a) FESEM image of Polyaniline [136], (b) FESEM image Commercial V$_2$O$_5$ powder (Sigma-Aldrich), (c) Galvanostatic charge-discharge curves for PANF-VNF and PANI-V$_2$O$_5$ electrodes measured at same current density of 0.1 A g$^{-1}$. (d) Performance comparison of nanostructured fibers ASC with commercial micron sized particles counterpart. Reproduced by permission of ECS - The Electrochemical Society.

Figure 4.40 depicts the FESEM image polyaniline, commercial V$_2$O$_5$ (Sigma-Aldrich) and a Ragone plot of ASC constituted of nanofibers with commercial powder of V$_2$O$_5$ and PANI [150]. Nanostructured ASC exhibited higher power and energy densities when compared to their conventional commercial counterparts. This is owing to the better utilization of the active material in the nanostructured ASC [133]. Moreover, nanostructure morphology can facilitate the diffusion of electrolyte ions, while the intrinsic high electroactive surface area enables larger electrode/electrolyte interface thus
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leading to higher specific capacitance. In addition, the optimization process for the fabrication of electrodes by mass loading based on the specific capacitance obtained in the three electrode configuration. However, the obtained energy density depends on the nature of material and its morphology. We believe that owing to the open porous one dimensional nature of fibers, the entire mass is involved in the electrochemical process resulting in high energy density, whereas in commercial powders only surface of the materials are involved owing to the micron sized particles thus providing lower values irrespective of the applied current density.

Additionally, to demonstrate the practical application of the fabricated asymmetric device, a red light emitting diode (1.6 V LED) is connected. During such testing, the asymmetrical device is galvanostatically charged at low current rate and discharged via LED, the lighting of LED is shown in Figure 4.41. Tandem cells of asymmetrical devices can be employed to increase the working voltage to the required level.
Results and Discussions

Figure 4.41: Photo image of red LED connected in series with asymmetrical device.
4.2.4 Cycling Performance of Asymmetrical Supercapacitor

![Graph: Cycle performance of asymmetrical device with a voltage of 2 V at scan rate of 5 mV s⁻¹. Reproduced by permission of ECS - The Electrochemical Society.]

Figure 4.42: Cycle performance of asymmetrical device with a voltage of 2 V at scan rate of 5 mV s⁻¹. Reproduced by permission of ECS - The Electrochemical Society.

Long cycling life is also another important requirement for supercapacitors especially for pseudocapacitors wherein faradic redox process are involved. Figure 4.42 shows the capacity retention of the asymmetrical capacitor cycled at 5 mV s⁻¹ up to 2 V. Capacity of device is normalized with respect to the first discharge capacitance. The asymmetric device displayed an initial specific capacitance of 42.7 F g⁻¹ with good cycling stability up to 2000 cycles, with capacity remaining at ~73% at 2000th cycle.
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Supercapacitors are fundamentally expected to have long cycle life as energy is stored physically via electrical double layer formation. Whereas in the case of pseudocapacitor, cycle life is often limited by the parasitic chemical reactions and mechanical breakdown of structures due to swelling of electrode materials, both of which will be more pronounced at higher voltage. Here, the initial capacity degradation can be ascribed to the stabilization of the crystal structure. Minor fading was observed once the structure was stabilized (after 150 cycles) during electrochemical cycling. High energy density of 21.7 Wh kg\(^{-1}\) is still maintained despite the fade. This finding shed new insights on coupling conducting polymer and nanostructured transition metal oxides to attain high energy density storage devices through careful selection of electrode materials and morphology tuning.

**Figure 4.43:** FESEM images of (a) V\(_2\)O\(_5\) nanofibers, (b) polyaniline nanofibers after cycling for 1000 cycles. Reproduced by permission of ECS - The Electrochemical Society.
Results and Discussions

Figure 4.43 shows the FESEM images of (a) V$_2$O$_5$ nanofibers, (b) polyaniline nanofibers after cycling for 1000 cycles. In comparison to the FESEM images of VNF and PANF depicted in Figure 4.25 and Figure 4.31, respectively, it is clear that the fibrous structures had collapsed after cycling. These observations explain the fading cycling performance previously discussed. A crumbled structure impedes the traffic of ions to and from the electrode materials. Also, less electronic charges can be harvested as more particles are being electrically isolated. All these result in lower active sites utilization and less charge capacity, hence lower capacitance and energy density are observed.

4.2.5 Electrochemical Impedance Studies of Asymmetrical Supercapacitor

To investigate electrochemical performance, such as their frequency dependence and equivalent series resistance (ESR) of the symmetrical and asymmetrical supercapacitors, EIS measurements were carried out in the frequency range 10 kHz to 10 mHz at an alternating current (AC) amplitude of 10 mV. Figure 4.44 illustrates the Nyquist impedance spectra of both VNF and PANF electrodes in symmetrical and asymmetrical configurations in aqueous electrolyte, respectively.
At sufficiently high frequencies, intersection with the real axis \((Z')\) provides the value of the ESR \((R_e)\) and the diameter of semicircle reveals the charge transfer resistance \((R_{ct})\) of electrode, which arises from diffusion of ions at the PANF-VNF electrode/electrolyte interface. The Nyquist impedance spectra in Figure 4.44 consists of a minor high frequency arc and a low frequency spike showing an inclination at an angle 45° indicating that the both VNF and PANF electrodes has good pseudocapacitive (combined contribution of both electrical double layer capacitance and faradic pseudocapacitance) characteristics [92].
Results and Discussions

The transformation of the plot to a vertical line takes place at low frequency can be mainly ascribed to the capacitive response of porous carbons. The presence of the non-vertical slope at the low frequency region of the Nyquist plot may be associated with low electronic conductivity of the electrode and different pore size distribution of the carbons [120].

To better understand the devices, measured impedance spectra were analyzed using the complex nonlinear least squares fitting method fitted with less than 10% error on the basis of the equivalent circuit depicted in Figure 4.23. The respective circuit equation is given in equation 4.4 aforementioned.

The equivalent circuit parameters fitted for the impedance spectra of symmetric device configuration with VNF-VNF and PANF-PANF electrodes and asymmetric device configuration with VNF-PANF electrodes in 3 M KCl are tabulated in Table 4-3. VNF-VNF configuration exhibited a slightly lower ESR (1.82 $\Omega$) compared to PANF-PANF (2.31 $\Omega$) and PANF-VNF (2.65 $\Omega$), probably due to the porous fiberous structure easing the flow of ions.

In the middle to low frequency region, both symmetrical and asymmetrical configurations showed high Warburg impedance values resulting from semi-infinite diffusion resistance of ions in the PANF and VNF electrode. Apparently, PANF-VNF exhibited a larger double layer capacitance ($C_{dl}$), probably resulting from the synergic effect.
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Table 4-3: Components of the equivalent circuit fitted for the Nyquist plots of all symmetrical and asymmetrical device configurations depicted in Figure 4.44.

<table>
<thead>
<tr>
<th>Configuration</th>
<th>$R_b$ ($\Omega$)</th>
<th>$C_d$ (mF cm$^{-2}$)</th>
<th>$R_{ct}$ ($\Omega$)</th>
<th>$C_p$ (mF cm$^{-2}$)</th>
<th>$R_w$ ($\Omega$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>VNF-VNF</td>
<td>1.82</td>
<td>14.65</td>
<td>0.51</td>
<td>7.78</td>
<td>15.21</td>
</tr>
<tr>
<td>PANF-PANF</td>
<td>2.31</td>
<td>16.34</td>
<td>19.17</td>
<td>0.06</td>
<td>16.42</td>
</tr>
<tr>
<td>VNF-PANF</td>
<td>2.65</td>
<td>21.76</td>
<td>21.37</td>
<td>20.56</td>
<td>23.53</td>
</tr>
</tbody>
</table>

The electrolyte resistance of the electrodes is small, indicating that the electrochemical properties of bulk electrolytes remain relatively stable. Both greater charge transfer resistance $R_{ct}$ and higher pseudocapacitance $C_p$ are obtained for the asymmetrical configuration. This indicates a greater amount of faradic current, a result of more charge transfer reactions taking place, analogous to the results revealed from CV and GCD measurements.

4.1.6 Contributions of Surface and Inner Charge Storage Capacity

To better understand the charge storage mechanism for the different nanofibers electrode in aqueous system, the measured current is segregated into two components. (1) capacitive contribution from fast electrical double layer and (2) diffusional controlled pseudocapacitance from charge transfer redox reaction. Based on the work of Trasatti et al. [151, 152], it is able to use the dependence of current on voltammetric scan rate to quantitatively determine the capacitive contribution. Accordingly, the voltammetric charge ($q$) of the electrode materials can be categorized into two parts, namely, surface capacitive charge ($q_s$) and diffusional controlled charge ($q_d$). Such can be described by the following equation,
Where $q_d$ is the diffusional controlled charge storage component, $q_T$ is the total charge storage capacity, and $q_s$ is the surface charge component.

Surface capacitive charge having a faster kinetics can be correlated with double layer capacitance, while diffusional controlled charge depends mainly on the slower pseudocapacitance. Assuming semi-infinite linear diffusion process and within a reasonable range of scan rates, $q_s$ can be derived from the plot of charge density against the inverse square root of potential scan rate ($\nu$) and extrapolating $\nu$ to $\infty$ as given by the following equation,

$$q = q_s + kv^{-0.5}$$

Where $q$ is the charge stored, $q_s$ is the surface charge component, $k$ is an arbitrary constant, $\nu$ is the scan rate.

On the other hand, if ions have ample time for diffusion to the inner and less accessible sites, via a very slow scan rate, theoretical maximum total charge storage ($q_T$) can be derived by using appropriate function of $\nu$. Thus $q_T$ can be extracted from plotting inverse charge storage with square root of scan rate according to

$$\frac{1}{q} = \frac{1}{q_T} + kv^{0.5}$$

Where $q$ is the charge stored, $q_T$ is the total charge storage capacity, $k$ is an arbitrary constant, $\nu$ is the scan rate.
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Figure 4.45 shows the extrapolated curves for FNF, VNF and PANF, respectively. The deviation from linearity of such plot at high scan rate can be ascribed to the polarization effects that are omitted in the above equation [151, 152]. The extrapolated values are extracted and used in subsequent analysis.
Figure 4.45: Variation of charge density (q) with respect to scan rates (v) : (a) Extrapolation of v to $\infty$ (the y-intercept of the q vs. $v^{0.5}$ plot) giving the outer charge for FNF electrodes. (b) Extrapolation of v to 0 (the y-intercept of the l/q vs. $v^{0.5}$ plot) giving the maximum volumetric charge for FNF electrodes. Extrapolation as mentioned in (a) for (c) PANF (e) VNF. Extrapolation as mentioned in (b) for (d) PANF (f) VNF.
Results and Discussions

Figure 4.46: Categorization of contribution from surface capacitive and diffusional controlled capacity of various nanofibers.

Figure 4.46 illustrates the derived surface capacitive and diffusional controlled components. It clearly shows that diffusional controlled charge storage play a significant role in the total capacity of the nanocomposite electrode. It is noted that PANF has the largest surface capacitive charge storage contribution while VNF reaped in the lowest value. This is consistent with the observation in FESEM and BET characterization which shows the highest surface area for PANF which facilities the transport of electrolyte ions and enhances the electrochemical utilization of PANF surface sites. FNF has approximately 8% charge contribution coming from the surface charge component, showing better surface charge storage capability than VNF also in agreement with the observation in BET characterization. Additionally, FNF has shown a superior overall charge storage capability than PANF which can be ascribed to the presence of multiple oxidation states presented in iron.
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On the other hand, VNF has shown a greater overall charge capacity among all the nanocomposites. It is evaluated that surface charge component make up of about 2% of the overall charge capacity, suggesting that only a small portion of the VNF surface sites is accessible to ions. In contrast, diffusional charge storage component make up most of the overall charge capacity, indicating that inner surface or bulk of the VNF provides more active sites for charge storage. Hence, the primary charge storage mechanism in VNF is mainly governed by the diffusional controlled electrochemical reaction rather than the surface driven EDL mechanism.
The relationship of charge storage with scan rate is also studied. Figure 4.46 depicts the plot of charge storage of various nanofibers with scan rate. It is clear that surface capacitive process becomes more accountable with higher scan rate. This is because when the nanofibers are subjected to higher scan rate, the electrolytes’ ions have less time to diffuse to the active sites which results in fewer uptakes of active sites for pseudocapacitive process. This agrees well with the cyclic voltammogram results.
Results and Discussions discussed in previous sections. The increment of diffusional charge storage capability is more prominent in both FNF and VNF than PANF. This can be ascribed to the intercalation of Na\(^+\) and K\(^+\) cations into the lattice of Fe_2O_3 and V_2O_5 nanofibers, respectively.
Chapter 5 Conclusions

This thesis presents a study on the charge storage mechanism and pseudocapacitive behavior of transitional metal oxide nano-composites and application of such nanomaterials in supercapacitor device configuration. Also, an asymmetrical architecture of transition metal oxide and conducting polymer has been successfully employed to increase the working voltage which renders greater energy density storage capability.

Electrochemical behaviors of EDL nanomaterials such as CNT were investigated. Although they show great power density, challenge of enhancement of energy density still remains attributable to the limited accessible surface area. Tin oxides nanospheres were added to the CNT to enhance the capacitance, however, no substantial improvement was observed probably due to the high contact resistance between the SnO₂ and CNT. Another pseudocapacitive material, Fe₂O₃ has been demonstrated to be a promising electrode material for application in supercapacitor. Electrospinning technique was employed to weave interconnected network of nanofibers. The sintering temperature and ramping rate have been optimized to 500 °C and 5 °C/min to obtain α-Fe₂O₃ nanofibers as well as maintaining its fibrous morphology. Such nanostructured fibrous morphology can heighten the specific capacitance by increasing the accessible surface area and facilitates ions transportation. However, the specific capacitance remains mediocre which is attributable to the high resistivity of α-Fe₂O₃. To improve the specific capacitance, carbon black was added as conducting agent to facilitate electronic pathways for current
Conclusions

The specific capacitance was enhanced substantially with the introduction of carbon black. Furthermore, the pseudocapacitive contribution from FNF was significantly increased as more charges can be harvested from the redox-active sites. The observations also revealed that specific capacitance is comprised of (i) non-faradic double layer capacitance and (ii) faradic pseudocapacitance. The former is primarily governed by the accessible conducting surface area whereas the later is greatly influenced by the pseudocapacitive properties of redox materials.

Extending the methodology, other transitional metal oxides such as $V_2O_5$ nanofibers was also synthesized. It exhibited good pseudocapacitive behavior in aqueous KCl solution and shows promise as a good electrode material for supercapacitor application.

To achieve higher energy density, the working voltage needs to be further extended. As such, the asymmetrical device configuration was conceptualized. To implement such architecture, a common electrolyte which works well with both the positive and anode is required. Polyaniline, an electrode material which also exhibits pseudocapacitive behavior in aqueous KCl solution is roped in. Nanostructured polyaniline was also obtained and found to be a good positive electrode material. On the other hand, $V_2O_5$ nanofibers demonstrate better performance as as a negative electrode material. Hence, nanofibers of polyaniline and $V_2O_5$ were coupled and assembled into an asymmetrical supercapacitor device. The product was found capable of having a wider...
Conclusions

working voltage of up to 2 V in aqueous solution. As such, a higher energy density is achieved.

Moreover, XPS analysis is performed to provide further insight into the charge storage mechanism of VNF and FNF suggesting the intercalation of K$^+$ and Na$^+$ ions into the oxide lattice of Fe$_2$O$_3$ and V$_2$O$_5$ lattices, respectively. It is therefore clear that intercalation of ions process rather than redox process contributes significantly to the charge storage mechanism of TMO nanofibers. Also, the surface double layer and diffusional controlled bulk storage ratio is also investigated.

In summary, the contributions of this thesis include investigating the application of $\alpha$-Fe$_2$O$_3$ in a two electrode symmetrical device configuration, the effects of conducting agent on the pseudocapacitive performance, and the employment of asymmetrical device configuration to improve the energy density of the supercapacitor device. Performance enhancement of the devices was demonstrated with nanostructured materials. A thorough understanding of the contribution of nanomaterials towards charge storage and mechanism was demonstrated.
Chapter 6 Future Works

Stemming from the works made in this thesis, further studies are needed in several areas to corroborate as well as to extend current work. Some recommendations are as follow,

6.1 Enhancing Electrical Conductivity

While the ASC benefit from good energy, they carry a low power density rating. It may be good to replace electrodes with a composite which has highly accessible surface area of good conductivity and great pseudocapacitive behavior. To achieve such combination, a conducting pseudocapacitive material is needed. Although electrically conducting polymer is having both the qualities, their conductivity is still insufficient. More needs to be done to further enhance their electrical conductivity. Also, polymers often suffers from short cycle life, hence, efforts need to spend on minimizing degradation of polymer on cycling. Similarly, transitional metal oxides also suffer low electrical conductivity although they have exhibited high capacitance and are capable of greater charge storage. Therefore, more work is required to improve the electrical conductivity. Carbon coating is a good avenue to better electrical conductivity. Such coating can be achieved by hydrothermal method using a carbon source precursor like glucose. Co-electrospinning is also a feasible route to acquire carbon coating. By co-spinning metal oxide precursor with carbon source like polyacrylonitrile and later sinter
to obtain carbon-metal oxide interlinked network, a better connected network can be realized.

**6.2 Investigation of Charge Transfer Mechanism via in-situ Route**

Investigation of redox process in transitional metal oxides is critical to understanding of charge storage process. Most of the research done until now exercises ex-situ characterization such as XPS and XRD. Such routes are unable to better comprehend the transition process of charge storage and utilization. The electrode material may have undergone changes after being exposed to the environment. In-situ XPS, XRD and FTIR characterization could provide a good real time portrayal of the ongoing process and changes. This enables a better understanding of the changes in valence states and/or ions intercalation/de-intercalation occurring in the electrode material.
List of Symbols and Abbreviations

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Description</th>
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<tbody>
<tr>
<td>EDL</td>
<td>Electrical Double Layer</td>
</tr>
<tr>
<td>SC</td>
<td>Supercapacitor</td>
</tr>
<tr>
<td>EC</td>
<td>Electrochemical capacitor</td>
</tr>
<tr>
<td>EDLC</td>
<td>Electrical Double Layer Capacitor</td>
</tr>
<tr>
<td>AC</td>
<td>Activated carbon</td>
</tr>
<tr>
<td>CNT</td>
<td>Carbon nanotubes</td>
</tr>
<tr>
<td>SWNT</td>
<td>Single-walled nanotubes</td>
</tr>
<tr>
<td>MWNT</td>
<td>Multi-walled nanotubes</td>
</tr>
<tr>
<td>NW</td>
<td>Nanowires</td>
</tr>
<tr>
<td>CB</td>
<td>Carbon black</td>
</tr>
<tr>
<td>PANI</td>
<td>Polyaniline</td>
</tr>
<tr>
<td>PEDOT</td>
<td>Poly (4-styrenesulfonate)</td>
</tr>
<tr>
<td>PSS</td>
<td>Poly (styrenesulfonate)</td>
</tr>
<tr>
<td>Ppy</td>
<td>Polypyrrole</td>
</tr>
<tr>
<td>PMeT</td>
<td>Poly (3-methylthiophene)</td>
</tr>
</tbody>
</table>
List of Symbols and Abbreviations

<table>
<thead>
<tr>
<th>Symbol</th>
<th>Description</th>
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<tbody>
<tr>
<td>PVP</td>
<td>Polyvinylpyrrolidone</td>
</tr>
<tr>
<td>PVdF</td>
<td>poly-vinylidenefluoride</td>
</tr>
<tr>
<td>NMP</td>
<td>N-methylpyrrolidone</td>
</tr>
<tr>
<td>α-Fe₂O₃</td>
<td>Alpha iron (III) oxide or hematite</td>
</tr>
<tr>
<td>FNF</td>
<td>Iron (III) oxide nanofibers</td>
</tr>
<tr>
<td>VNF</td>
<td>Vanadium (IV) oxide nanofibers</td>
</tr>
<tr>
<td>AN</td>
<td>Acetonitrile</td>
</tr>
<tr>
<td>EC</td>
<td>Ethylene carbonate</td>
</tr>
<tr>
<td>DEC</td>
<td>Diethylene carbonate</td>
</tr>
<tr>
<td>DMC</td>
<td>Dimethyl carbonate</td>
</tr>
<tr>
<td>CV</td>
<td>Cyclic Voltammetry</td>
</tr>
<tr>
<td>GCD</td>
<td>Galvanostatic charge-discharge</td>
</tr>
<tr>
<td>EIS</td>
<td>Electrochemical Impedance Spectroscopy</td>
</tr>
<tr>
<td>FESEM</td>
<td>Field Emission Scanning Electron Microscopy</td>
</tr>
<tr>
<td>TEM</td>
<td>Transmission Electron Microscopy</td>
</tr>
<tr>
<td>XRD</td>
<td>X-ray Diffraction</td>
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# List of Symbols and Abbreviations

<table>
<thead>
<tr>
<th>Symbol</th>
<th>Description</th>
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<tbody>
<tr>
<td>BET</td>
<td>Brunauer, Emmett and Teller</td>
</tr>
<tr>
<td>BJH</td>
<td>Barrett, Joyner and Halenda</td>
</tr>
<tr>
<td>FTIR</td>
<td>Fourier Transform Infrared spectroscopy</td>
</tr>
<tr>
<td>TGA</td>
<td>Thermo gravimetric Analysis</td>
</tr>
<tr>
<td>$C_T$</td>
<td>Total capacitance</td>
</tr>
<tr>
<td>$C_p$ or $C_+$</td>
<td>Capacitance of positive electrode</td>
</tr>
<tr>
<td>$C_n$ or C.</td>
<td>Negative electrode</td>
</tr>
<tr>
<td>V</td>
<td>Voltage</td>
</tr>
<tr>
<td>$R_{ESR}$</td>
<td>Equivalent Series Resistance</td>
</tr>
<tr>
<td>E</td>
<td>Energy</td>
</tr>
<tr>
<td>P</td>
<td>Power</td>
</tr>
<tr>
<td>$C_D$</td>
<td>Specific capacitance of the device</td>
</tr>
<tr>
<td>$C_{dl}$</td>
<td>Double layer capacitance</td>
</tr>
<tr>
<td>$C_{ps}$</td>
<td>Pseudocapacitance</td>
</tr>
<tr>
<td>Z</td>
<td>Magnitude of ionic charge</td>
</tr>
<tr>
<td>C</td>
<td>Concentration</td>
</tr>
</tbody>
</table>
List of Symbols and Abbreviations

$C_0$  Concentration of ions in bulk solution

$q$  Electronic charge

$m_+$  Mass of positive electrode

$m_-$  Mass of negative electrode

$e$  charge of an electron ($1.60217646 \times 10^{-19} \text{C}$)

$\Delta E_+$  Voltage of positive electrode

$\Delta E_-$  Voltage of negative electrode

$t$  Time

$k$  Boltzmann constant ($1.3806503 \times 10^{-23} \text{ m}^2 \text{kg s}^{-2} \text{K}^{-1}$)

$T$  Absolute temperature in Kelvin

$\varphi_m$  Potential at the metal surface

$\varepsilon_0$  Permittivity of free space ($8.854 \times 10^{-12} \text{J} \text{m}^{-1}$)

$\varepsilon_r$  Dielectric constant of electrolyte

$A$  Surface area

$d$  Distance between parallel plates of charges

$C_H$  Capacitance contribution from the Helmholtz layer
List of Symbols and Abbreviations

<table>
<thead>
<tr>
<th>Symbol</th>
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<tbody>
<tr>
<td>$C_G$</td>
<td>Capacitance contribution from the diffuse layer</td>
</tr>
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List of Publications

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