

# SCIENTIFIC REPORTS

OPEN

## Hierarchical One-Dimensional Ammonium Nickel Phosphate Microrods for High-Performance Pseudocapacitors

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Received: 01 July 2015

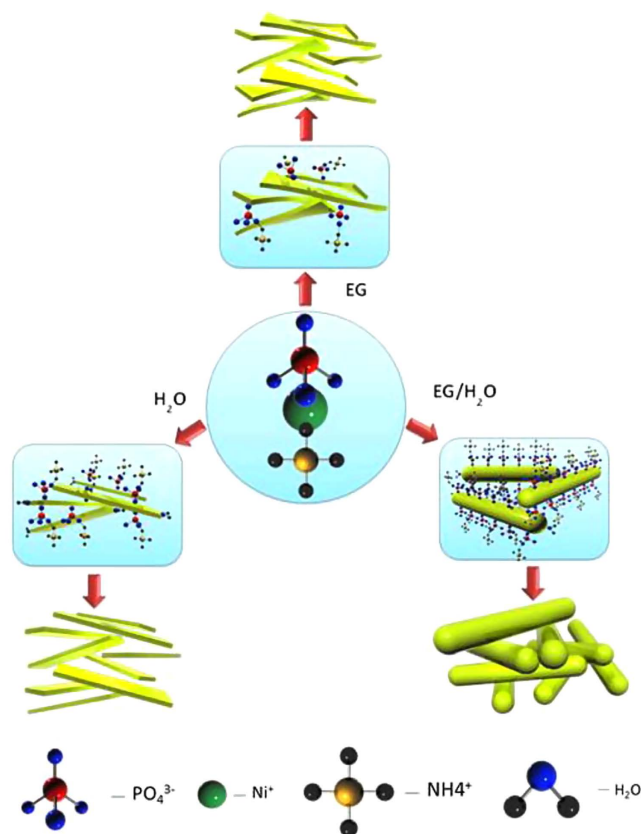
Accepted: 03 November 2015

Published: 03 December 2015

High-performance electrochemical capacitors will drive the next-generation portable, flexible and wearable electronics. Unlike the conventional all-carbon supercapacitors (electric double layer capacitors, EDLC) with high power but poor energy density, pseudocapacitors capitalize the high energy density inherent to reversible redox reactions and provide a facile means to enhancing the energy ratings of supercapacitors. The high length-to-diameter ratio and anisotropic character of 1-D architecture makes them suitable for use in energy storage. For the first time, we report 1-D microrod structures (~ 36 nm width) of ammonium nickel phosphate hydrate (ANP<sub>mr</sub>) as a pseudocapacitor with high energy rating and power handling. To confirm the data, the ANP<sub>mr</sub>-based pseudocapacitor was subjected to various configurations (i.e., half-cell, symmetric, asymmetric, and flexible all-solid-state) and in each case it gave excellent values compared to any accessible literature to date. We clearly demonstrate that a flexible all-solid-state ANP<sub>mr</sub>-based pseudocapacitor achieved high areal capacitance of 66 mF cm<sup>-2</sup> with extra-ordinary energy (21.2 mWh cm<sup>-2</sup>) and power (12.7 mW cm<sup>-2</sup>) densities. This work opens doors for a facile, robust and scalable preparation strategy for low-cost, earth-abundant electrode materials for high-performance pseudocapacitors.

A considerable amount of research attention has continued to be devoted to renewable and clean energy technologies globally. The research on improving the electrical energy storage is crucial for increasing the supply of more energy from renewable sources to curb the present energy challenges<sup>1–5</sup>. There are two main classes of electrochemical capacitors (ECs); the all-carbon based electrical double layer capacitors (EDLCs) that store energy by charge-separation at the solid-electrolyte interface, and the pseudocapacitors that store energy by redox or Faradaic processes. EDLCs (also known as the supercapacitors or ultracapacitors) are important energy storage devices with adequate energy and high power densities compared to conventional electrochemical energy storage systems such as batteries and fuel cells<sup>6,7</sup>. An extraordinary storing of electrical energy with exceptional power has been projected to increase the awareness and development of important technologies such as hybrid electric vehicles, portable electronics and power-saving units. It is well-established that the performance of supercapacitor-driven technologies is dependent on the physicochemical properties of their electrode materials<sup>8</sup>. Flexible and wearable electronics have begun to attract intense research interests due to their reliability, ease of handling, and the great promises to be used as flexible energy storage devices<sup>9</sup>. Pseudocapacitors have emerged as the most successful and substantial electrical energy storage devices for the flexible and wearable electronics. However, one of the major short-comings of the pseudocapacitor over their supercapacitor counterpart is their poor rate-capability compared to supercapacitor<sup>10</sup>. Therefore, the need for the development of high-performance pseudocapacitor materials is urgent.

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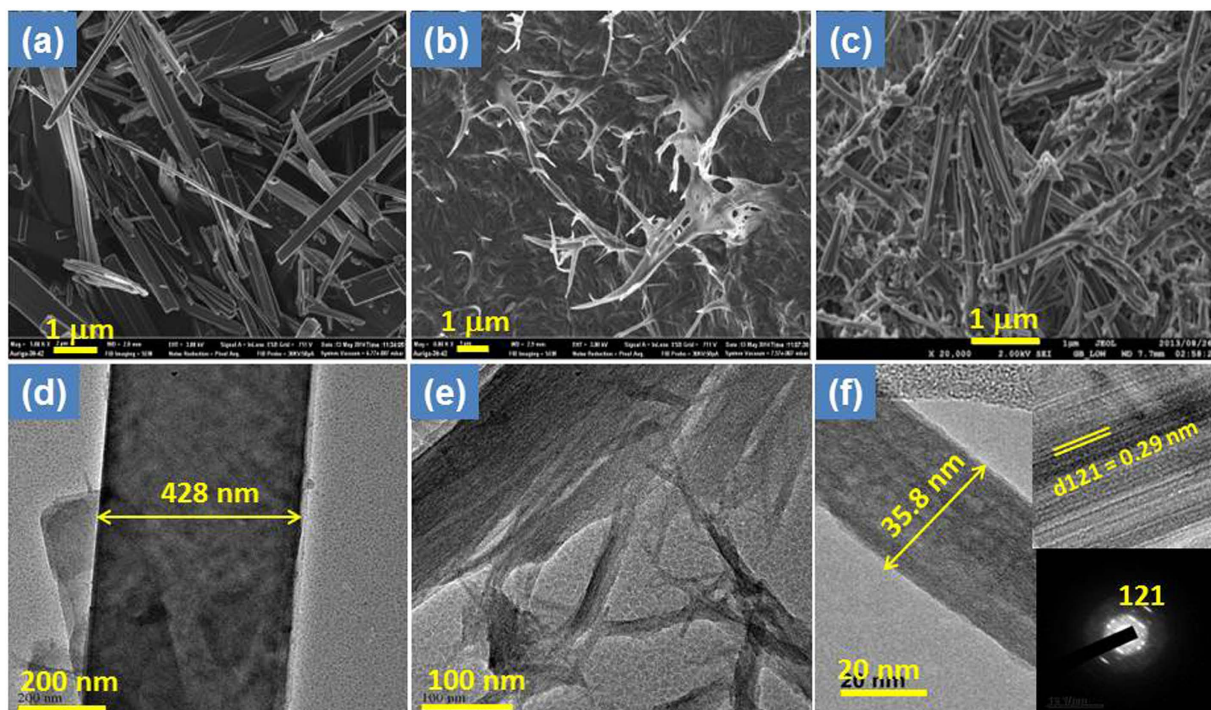
**Figure 1. (Ozoemena):** Schematic representation of various formations of ANP materials.

Hierarchical 1-D and 2-D materials maximize the supercapacitive properties due to their unique ability to permit ion propagations<sup>11,12</sup>. Phosphate-rich materials (PRMs), such as mesoporous  $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$  nanoalmonds<sup>13</sup>, 1-D layered  $\text{NH}_4\text{CoPO}_4 \cdot \text{H}_2\text{O}$  microrods<sup>14</sup> and 2-D  $\text{VOPO}_4$  nanosheets<sup>15</sup> have been reported as high-performance pseudocapacitors. For example, the energy and power densities of these PRMs range between  $30\text{--}140 \text{ Wh kg}^{-1}$  and  $1\text{--}27 \text{ kW kg}^{-1}$  for 3-electrode configurations. One of the major challenges with the PRMs is their low electrical conductivity, which explains why, for example, the  $\text{VOPO}_4$  nanosheets had to be integrated with high-electrical-conducting graphene sheets<sup>15</sup>. There is a need therefore to make PRMs that allow for compatible conduction pathways in their structures for improved redox-activity and pseudocapacitive behaviour. 1-D materials have been known to greatly influence space-confined transport phenomena thereby improving the charge accumulation and redox reactions<sup>11,16</sup>. Indeed, the major attractions of 1-D architecture include their high length-to-diameter ratio and anisotropic character. The small diameter of the 1-D architecture is important as it allows for enhanced accommodation of possible large volume changes, thus preventing possible cracking or fracturing of the structures usually observed in bulk or micron-sized materials. It is possible for ions and electrons to be simultaneously integrated into 1-D architectures, thereby making them ionically and electronically conductive. Importantly, 1-D architecture are characterised by large surface-to-volume ratio that permit efficient contact between the active mass of the electrode and the electrolyte, promoting high-rate capability. It is without doubt therefore that an important strategy for achieving high-performance PRMs is the preparation of their hierarchical 1-D architectures such as the wire-like or rod-like morphology. To our knowledge, rod-like PRMs do not exist.

Motivated by the above advantages of 1-D materials in pseudocapacitor applications, and the current challenges identified in the literature, this work describes the first synthesis of 1-D  $\text{NH}_4\text{NiPO}_4$  microrods (with nanometric width) with a view to increasing the energy and power densities. Herein we introduce a facile strategy to synthesize various morphologies of  $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$  by hydro/solvothermal route in ethylene glycol (EG), water and mixed solvents of EG/ $\text{H}_2\text{O}$  using nickel-based acetate and ammonium phosphate without the use of a template and additives. We show that when subjected to various experimental conditions from half-cell and symmetric to asymmetric and flexible all-solid-state configurations, the  $\text{NH}_4\text{NiPO}_4$  microrods still maintained excellent performance.

## Results and Discussion

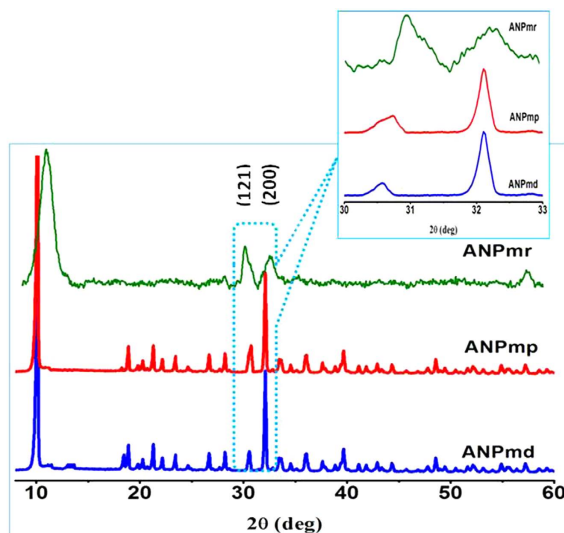
A schematic representation of the synthetic strategy adopted for the formation of three different nanostructures of  $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$  is summarised in Fig. 1. The microdendrites of  $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$  ( $\text{ANP}_{\text{md}}$ ),



**Figure 2. (Ozoemena):** SEM micrograph of as-synthesized various morphologies of (a) ANP<sub>mp</sub> (microplatelets), (b) ANP<sub>md</sub> (microdendrites) and (c) ANP<sub>mr</sub> (microrods), respectively. TEM and HR-TEM images (d) ANP<sub>mp</sub> with diameter of 428 nm, (e) ANP<sub>md</sub> microdendrites with >100 nm diameter and (f) single nanorod of ANP<sub>mr</sub> with 35.8 nm diameter, the inset of 2f showing lattice fringes corresponding to the (121) plane of microrods and their SAED pattern.

microplatelets (ANP<sub>mp</sub>) and microrods (ANP<sub>mr</sub>) were synthesized in EG, water, and EG/water mixture via facile solvo/hydrothermal processes, respectively. The morphology and size of as-synthesized NH<sub>4</sub>NiPO<sub>4</sub>·H<sub>2</sub>O were confirmed with the SEM and TEM analyses. Figure 2 (a–f) shows the SEM and TEM images of ANP<sub>md</sub>, ANP<sub>mp</sub> and ANP<sub>mr</sub> showing the unique formation of different morphologies of microplatelets, microdendrites and microrods. It is interesting to note from Fig. 2 that the obtained microplatelets (Fig. 2a) have average dimensions of 400–600 nm in length and nearly 428 nm in diameter (Fig. 2d). Figure 2b,e illustrate the SEM and TEM images of ANP<sub>md</sub> with average dimension of 100–300 nm in length. As shown in Fig. 2c, the obtained microrods have average dimensions of 200–300 nm in length and 35–40 nm in diameter, with few of them as nanorods. As illustrated by the HRTEM image, (Fig. 2f and insets), the observed microrods (with nanometric width) are polycrystalline in nature with clear lattice fringes. The d-spacings of the lattice fringes is found to be 0.29 nm, corresponding to the (121) plane of ANP<sub>mr</sub> along with SAED pattern corresponding to (121) plane. Note that these microrods could only be formed at 48 h as shown from the SEM and TEM images (Supplementary Information, Figure-S1).

From the results, it is evident that spatial localization of water molecules in solvent mixture is critical in mediating the shape growth; the highly viscous solvent EG ( $\eta = 21$  mPa s, 20°C) restrains the mobility of reactants compared to water ( $\eta = 1.0087 \times 10^{-3}$  mPa s, 20°C). The solubility and mobility of reactants considerably favours the homogeneous nucleation process when an appropriate amount of EG/H<sub>2</sub>O is used<sup>17</sup>. In general, the formation mechanism of the microstructures seems to occur via the hydrolysis of acetate to acetic acid and hydroxide ion in aqueous solution, followed by the reaction between the PO<sub>4</sub><sup>3-</sup>, NH<sub>4</sub><sup>+</sup> and Ni<sup>2+</sup> ions. The diffusion of the active sites of PO<sub>4</sub><sup>3-</sup>, NH<sub>4</sub><sup>+</sup> and Ni<sup>2+</sup> ions increases with a rise in temperature thus enhancing the nucleation process. The viscosity of EG decreased with increase in temperature thereby facilitating fast nucleation by reducing the interlayer spacing thus enhance the anisotropic growth of nanorods<sup>18</sup>. The NH<sub>4</sub>NiPO<sub>4</sub>·H<sub>2</sub>O layers are formed by sharing the highly distorted NiO<sub>6</sub> octahedra corners with cross-linked distorted PO<sub>4</sub><sup>3-</sup> tetrahedra and NH<sub>4</sub><sup>+</sup> ions inserted between the inorganic layers via hydrogen bonding<sup>19</sup>. There are other 1-D materials, such as LnPO<sub>4</sub> and CePO<sub>4</sub>, whose formation is driven by diffusion-controlled growth mechanism, i.e., attachment of infinite linear chains along the axis of its crystalline phase<sup>20</sup>. The structural arrangement of NH<sub>4</sub>NiPO<sub>4</sub>·H<sub>2</sub>O also contains open channels of octahedra along the parallel and perpendicular axis in (010) and (001) planes which may tend to form linear chain extending of octahedral along the axis. Additionally, a drastic change in morphology has also been reported when using different reactants. For instance, Zhao *et al.*<sup>13</sup> recently reported that nickel nitrate gives almond-like NH<sub>4</sub>NiPO<sub>4</sub>·H<sub>2</sub>O whereas

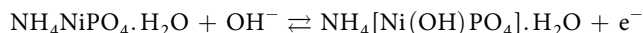


**Figure 3. (Ozoemena):** XRD pattern of ANP samples of ANP<sub>mp</sub>, ANP<sub>md</sub> and ANP<sub>mr</sub>. The inset is an expanded view of (121) and (200) peaks.

nickel acetate gives dendrite-like  $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$  morphology. The nitrate anion, which is a weak base and a good-leaving group, can easily substitute with one another and proceeds to 3-D growth process, whereas strong base of acetate anions surrounding the cation impede the 3-D growth. However, acetate ions and equal mixture of water and EG play a crucial role in the formation of unique morphologies of 1-D  $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$  materials, a facile and scalable approach which is more suitable to be extended to the preparation of various 1-D and 2-D ammonium metal phosphate ( $\text{NH}_4\text{MPO}_4 \cdot \text{H}_2\text{O}$ , where  $\text{M} = \text{Ni}$ ,  $\text{Mn}$ ,  $\text{Fe}$ ,  $\text{Co}$ , etc) materials.

Figure 3 shows the XRD pattern of  $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$  samples of ANP<sub>mp</sub>, ANP<sub>md</sub> and ANP<sub>mr</sub>. The XRD patterns indicate the characteristic peaks of a pure phase of  $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$  and all the observed peaks can be readily indexed to a pure orthorhombic phase (space group:  $\text{Pmn}2_1$ ) with the cell parameters of  $a = 5.425 \text{ \AA}$ ,  $b = 8.77 \text{ \AA}$  and  $c = 4.31 \text{ \AA}$  in accordance with the JCPDS card no 86-0585. Interestingly, an increase in intensity of the (121) peak is observed compared to the (200) peak infers the preferential orientation or anisotropic growth along the  $c$ -axis (insert Fig. 3). The broad peak of the ANP<sub>mr</sub> is a clear indication that its particle size is smaller than those of the ANP<sub>mp</sub> and ANP<sub>md</sub>.

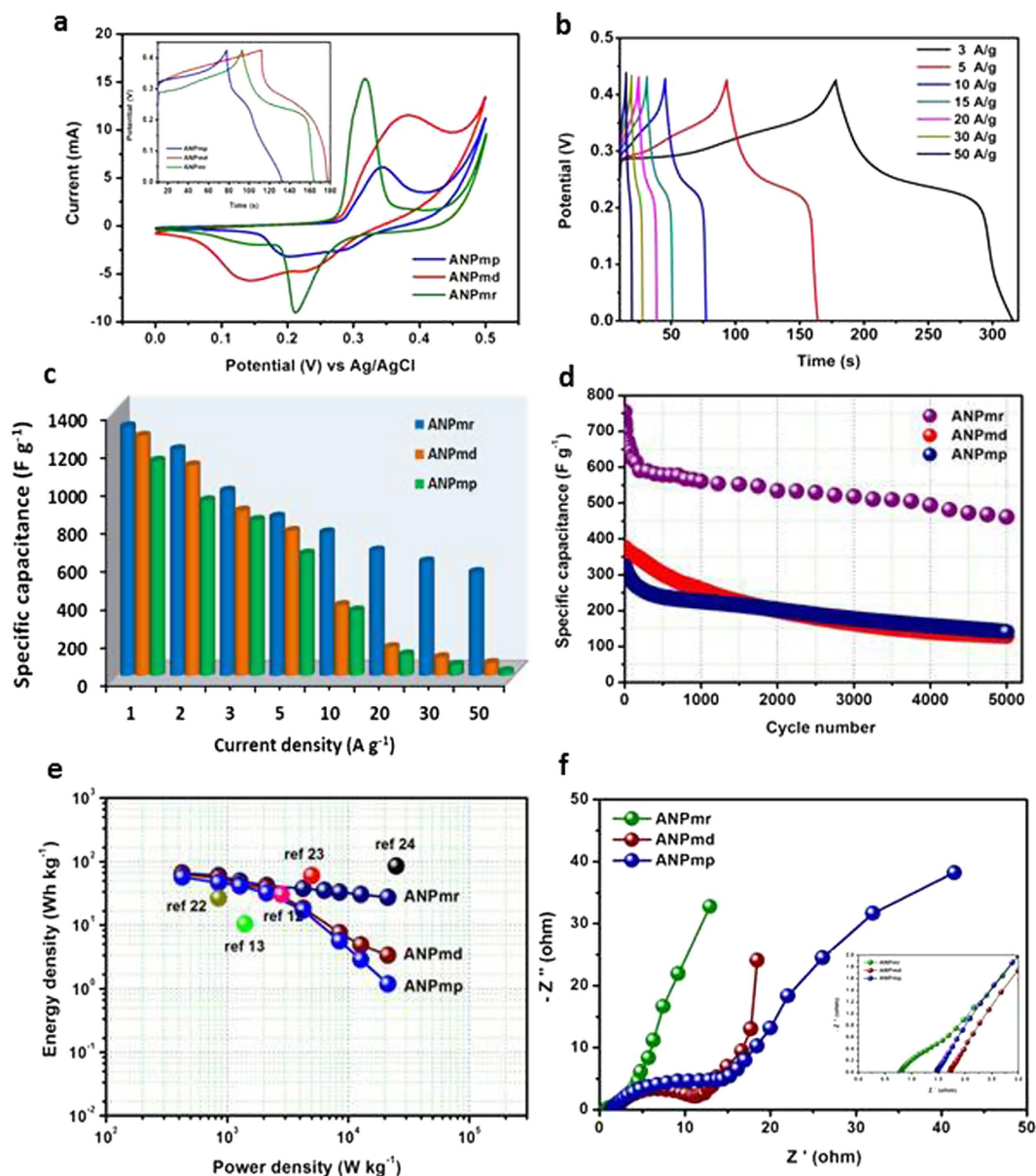
**Electrochemical characterization of three-electrode system.** Figure 4 compares the electrochemical performance of the three-electrode configurations of the three microstructures. The cyclic voltammetric evolutions (Fig. 3a) depict a redox couple arising from the redox-active nickel ( $\text{Ni}^{2+}/\text{Ni}^{3+}$ ), confirming the pseudocapacitive behaviour of the  $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$ . The emergence of the redox couple is represented as follows:<sup>14,21</sup>



The microrods showed well-defined electrochemistry with large current density and narrower peak-to-peak potential ( $\Delta E_p \approx 150 \text{ mV}$  at  $20 \text{ mVs}^{-1}$ ) compared to the  $>200 \text{ mV}$  shown by the microdendrites and microplatelets, meaning that ANP<sub>mr</sub> exhibits better electrochemical reversibility and faster electron transfer kinetics. Also, unlike the others, the microrods showed no additional oxidation peaks, indicating that the only oxidation process is that of  $\text{Ni}^{2+}/\text{Ni}^{3+}$  without any other phase changes.

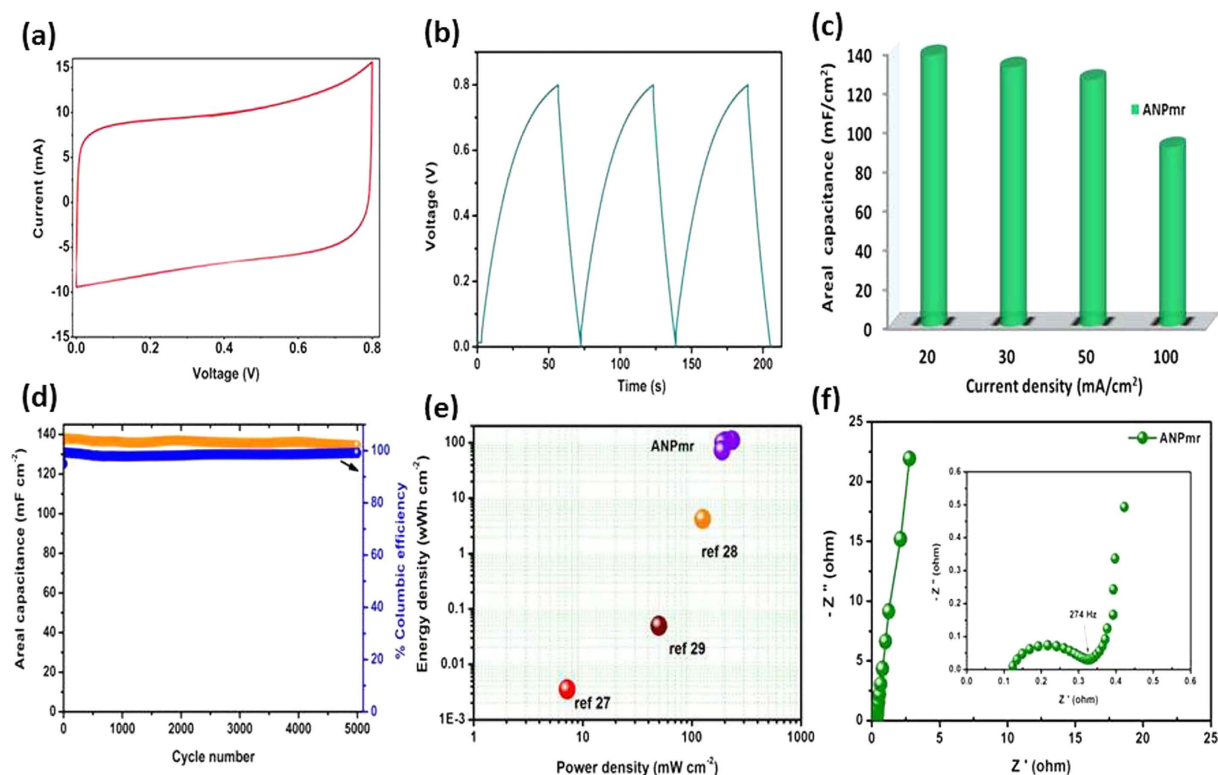
Figure 4b shows typical charge/discharge curves of the ANP<sub>mr</sub> at different current densities ( $1$  to  $50 \text{ A g}^{-1}$ ). At all the current densities investigated (Fig. 4c), the ANP<sub>mr</sub> electrode showed the best performance compared to the ANP<sub>md</sub> and ANP<sub>mp</sub>, achieving remarkable a maximum reversible specific capacity of  $\sim 1400 \text{ F g}^{-1}$ . Interestingly, the ANP<sub>mr</sub> showed an extra-ordinary high rate capability proven by the high capacitance of  $545 \text{ F g}^{-1}$  at a very high current density of  $10 \text{ A g}^{-1}$  which is extremely high in comparison with the capacitance value reported in the literature to date. The high value reflects the effective ion migration even at a higher speed which is influenced by high surface area of microrods with minimum diffusion length of ion accessibility. Upon continuous cycling, the electrode experienced capacitance loss at the initial cycles, stabilized at about 200 cycles and then retained nearly  $\sim 80\%$  of its original specific capacitance after 5,000 cycles (Fig. 4d). Figure 4e highlights the Ragone plot of the calculated power and energy densities. The results are higher than most recently reported supercapacitor nanomaterials based on 3-electrode configurations (Supplementary Information, Table-S1).





**Figure 4. (Ozoemena):** Comparative supercapacitive performance of the three different ANP electrodes using *three-electrode* (half-cell) configurations with nickel foam as the working electrode and 3M KOH as the aqueous electrolyte: **(a)** Typical CV curves at 10 mV s<sup>-1</sup>, insert shows the galvanostatic charge-discharge (GCD) curves at 5 A g<sup>-1</sup>; **(b)** GCD curves between 3 and 50 A g<sup>-1</sup>; **(c)** Specific capacitance vs. current density, i.e., rate capability test; **(d)** durability test for 5,000 continuous GCD cycles at 10 A g<sup>-1</sup>; **(e)** Ragone plots compared with other similar electrode materials reported in literature using 3-electrode configurations<sup>12,13,22,23,24</sup>; **(f)** Nyquist plot of 3 M KOH solution (inset; magnified view).

To provide further insights into the pseudocapacitive behaviour of the ANP materials, EIS experiments were conducted at open-circuit voltage at room temperature. Figure 4f compares the EIS (Nyquist plots) of the three ANP-based electrodes. The EIS data were satisfactorily fitted with the electrical equivalent circuit (EEC) comprising two Voigt RC elements, involving a series resistance ( $R_s$ ), charge-transfer resistance ( $R_{ct}$ ) and constant-phase elements (CPE or Q). As summarised in the Table-S2 (Supplementary Information) the  $R_s$  of the ANP<sub>mr</sub> (ca. 0.8 Ω) is smaller than those of the ANP<sub>mp</sub> (1.48 Ω) and ANP<sub>md</sub> (1.72 Ω). Also, the total  $R_{ct}$  values of the ANP<sub>mr</sub> (ca. 25.6 Ω) is smaller than those of the ANP<sub>md</sub> (ca. 31.4 Ω) and ANP<sub>mp</sub> (ca. 41.4 Ω). These results clearly indicate that the rod-like morphology provides the least internal resistance of the electrode and permits faster charge transportation compared to other morphologies investigated in this study. In addition, from the Fig. 4f, the ANP<sub>mr</sub> showed near-vertical line as expected of a high-performing pseudocapacitance compared to others. The “knee” or “onset” frequency ( $f_o$ ), which is a measure of the power capability of a supercapacitor, decreases as ANP<sub>mr</sub>



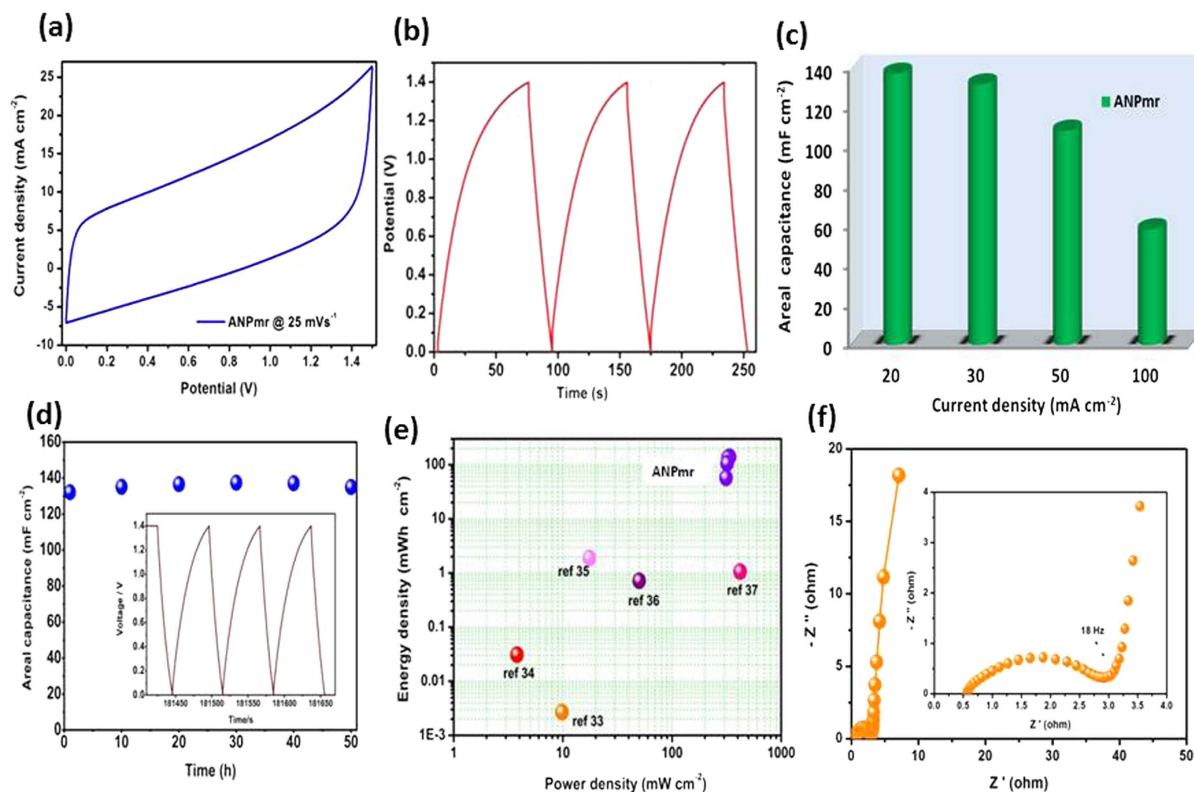
**Figure 5. (Ozoemena):** Electrochemical performances of symmetric pseudocapacitors of ANP<sub>mr</sub> coated on carbon cloth in 3M KOH: (a) Typical cyclic voltammogram obtained at a scan rate of 10 mVs<sup>-1</sup>; (b) galvanostatic CD profiles of ANP<sub>weg</sub> at the current density of 10 mA cm<sup>-2</sup>; (c) Areal capacitance calculated from CD curves as a function of current density; (d) Cycling stability and coulombic efficiency from 5,000 continuous charge-discharge cycles at 10 mA cm<sup>-2</sup>; (e) Ragone plot of A ANP<sub>mr</sub> symmetric supercapacitor compared with other symmetric supercapacitor values reported in literature<sup>27,28,29</sup>; and (f) Nyquist plot of ANP<sub>mr</sub> symmetric supercapacitor (inset; magnified view).

(5 kHz) > ANP<sub>md</sub> (1.7 kHz) > ANP<sub>mp</sub> (1.18 kHz), confirming the higher energy-storage capability of the ANP<sub>mr</sub> over other electrodes. The experimentally observed impedance curve was best fitted with the equivalent circuit and the calculated value of ESR of ANP<sub>mr</sub>, ANP<sub>md</sub> and ANP<sub>mp</sub> were found to be 0.79, 1.72 and 1.48 Ω. It is noted that the impedance of CPE is defined as equation (1):

$$Z_{CPE} = \frac{1}{[Q(j\omega)^n]} \quad (1)$$

where  $Q$  represents the frequency-independent constant relating to the surface electroactive properties,  $\omega$  is the radial frequency, the exponent  $n$  arises from the slope of  $\log Z$  vs.  $\log f$  (and has values  $-1 \leq n \leq 1$ ). If  $n = 0$ , the CPE behaves like a pure resistor;  $n = 1$ , CPE behaves as a pure capacitor;  $n = -1$ , CPE behaves as an inductor; while  $n = 0.5$  corresponds to Warburg impedance ( $Z_w$ ) which is related to the diffusion of the ions. The  $n$  values observed for the ANP<sub>mr</sub> ( $n_1 = 0.58$  and  $n_2 = 0.53$ ) while other electrodes were *ca.* 0.8, which suggests that the rod-like structure allows for improved ion diffusion than others. Further, the impedance curves were measured after 5,000 consecutive charge-discharge cycles (Supplementary Information, Figure-S2).

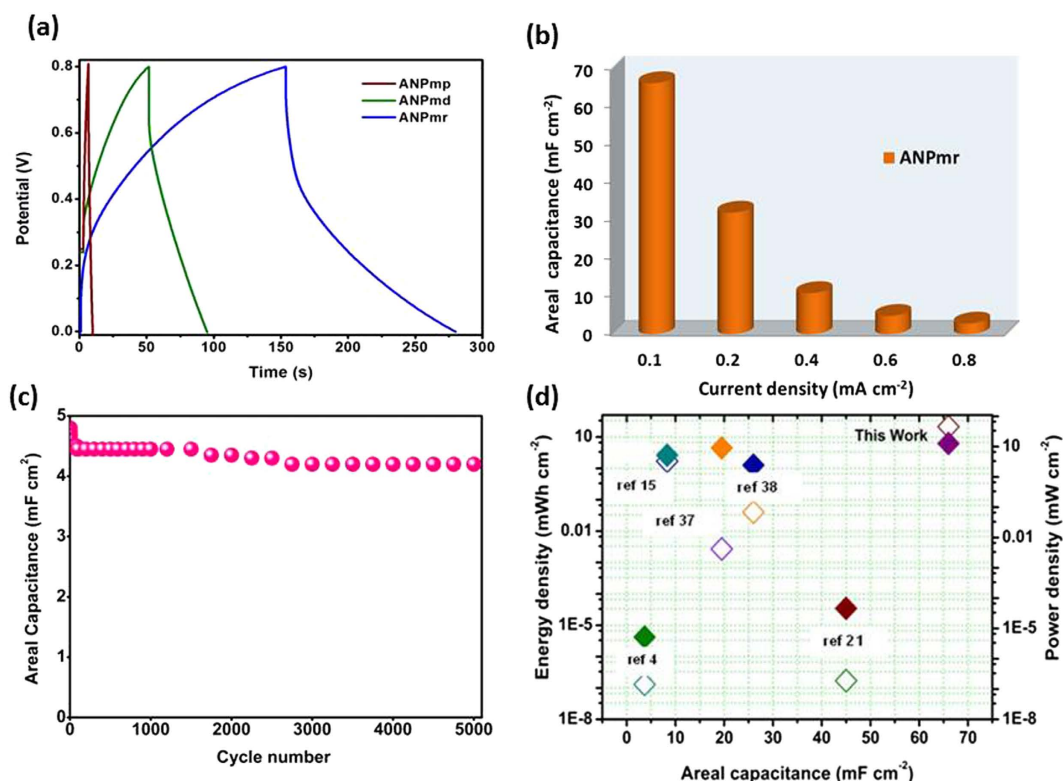
**Symmetric pseudocapacitors in alkaline electrolyte (3 M KOH).** Considering the high-performance of the ANP<sub>mr</sub> electrode at half-cell configuration, subsequent studies on full-cell pseudocapacitor devices were devoted to the ANP<sub>mr</sub>. Figure 5 summarises the performance of the ANP<sub>mr</sub> as a symmetric pseudocapacitor in an aqueous alkaline electrolyte (3 M KOH) using carbon cloth as substrate/current collector, showing typical CV curve at a scan rate of 10 mVs<sup>-1</sup> (Fig. 5a), charge-discharge curves at current density 10 mA cm<sup>-2</sup> (Fig. 5b), areal capacitance (Fig. 5c), cycle stability (Fig. 5d), Ragone plots (Fig. 5e) and Nyquist plot (Fig. 5f) of ANP<sub>mr</sub> symmetric pseudocapacitors. The symmetric pseudocapacitor showed satisfactory rate capability, evident at different scan rate and current densities (Supplementary Information, Figure-S3). Interestingly, the electrode gave high areal capacitance of 138 mF cm<sup>-2</sup> at 20 mA cm<sup>-2</sup>, 92% of which (126 mF cm<sup>-2</sup>) was retained even at a current density of 50 mA cm<sup>-2</sup>. The high rate capability



**Figure 6. (Ozoemena):** Electrochemical performances of asymmetric pseudocapacitors of ANP<sub>mr</sub>//AC coated on carbon cloth 1M Na<sub>2</sub>SO<sub>4</sub>: (a) Typical specific capacitance calculated from CV curve against voltage at a scan rate of 25 mVs<sup>-1</sup>; (b) galvanostatic CD profiles at 10 mA cm<sup>-2</sup>; (c) Areal capacitance from CD curves as a function of current density; (d) Typical voltage-holding (floating) curves acquired for 50 h at the voltage of 1.4 V and at 10 mA cm<sup>-2</sup>, inset is an example of CD curves after 50 h voltage-holding; (e) Ragone plot for this work and similar symmetric supercapacitors reported in literature<sup>33–37</sup>; (f) Nyquist plot of ANP<sub>mr</sub>//AC (inset; magnified view).

can be attributed to the microrods maintaining their excellent structural stability and charge propagation even at higher current densities. It was found that the ANP<sub>mr</sub> has the highest areal capacitance compared with recently reported symmetric supercapacitor materials (*cf.* Supplementary Information, Table-S3). More importantly, the areal capacitance retain more than 97% of its initial values after 5000 continuous charge-discharge cycles with 100% columbic efficiency (Fig. 5d). As shown in Fig. 5e, the ANP<sub>mr</sub> delivered the highest energy and power densities of 69 mWh cm<sup>-2</sup> and 145 mW cm<sup>-2</sup> at a current density of 20 mA cm<sup>-2</sup> and was found that the ANP<sub>mr</sub> has energy values compared to that of literature values. As shown in the EIS spectra (Fig. 5f), the observed high frequency intercept show that ANP<sub>mr</sub> (0.13 Ω) have much smaller ESR with inclined vertical line after the semicircle with the response time of 8 ms was lower than the values reported in liquid electrolyte used supercapacitors, onion-like carbon (26 ms)<sup>25</sup> and bistrilled yarn (17 ms)<sup>26</sup>. As summarised in the Table-S2 (Supplementary Information), the ANP<sub>mr</sub>-based symmetric cell gave very small  $R_s$  (*ca.* 0.13 Ω) and total  $R_{ct}$  ( $R_{ct1} + R_{ct2} \approx 33 \Omega$ ), with each  $n$  value ( $n_1$  or  $n_2$ ) greater than 0.8 but less than 1, clearly confirming the pseudocapacitive properties of the ANP<sub>mr</sub> when deployed in full-cell system.

**Asymmetric pseudocapacitors in neutral aqueous electrolyte (1 M Na<sub>2</sub>SO<sub>4</sub>).** We also prepared asymmetric pseudocapacitors in order to further increase the energy density of the device. Typically, ANP<sub>mr</sub> coated carbon cloth electrodes were used as positive and activated carbon (Norit® supra) coated carbon cloth electrode as negative in 1M Na<sub>2</sub>SO<sub>4</sub> neutral aqueous electrolyte. The cyclic voltammograms of ANP<sub>mr</sub> (Fig. 6a) obtained at a scan rate of 25 mVs<sup>-1</sup> shows rectangular shapes. Figure 6b show the charge-discharge at a current density of 10 mA cm<sup>-2</sup>, the cell gave high areal capacitance of 221 mF cm<sup>-2</sup> at 20 mA cm<sup>-2</sup>, 90% of (201 mF cm<sup>-2</sup>) which was retained even at a current density of 50 mA cm<sup>-2</sup>. It was found that this asymmetric capacitor has best areal capacitances compared to many nanostructured electrodes reported earlier in asymmetric capacitors, such as H-TiO<sub>2</sub> @ MnO<sub>2</sub> (0.9 F cm<sup>-3</sup>)<sup>30</sup>, TiO<sub>2</sub>/NiO nanotube array (2.9 mF cm<sup>-2</sup> at 0.4 mA cm<sup>-2</sup>)<sup>31</sup> and Fe<sub>3</sub>O<sub>4</sub>-SnO<sub>2</sub> core-shell nanorod film (7 mF cm<sup>-2</sup>)<sup>32</sup>.

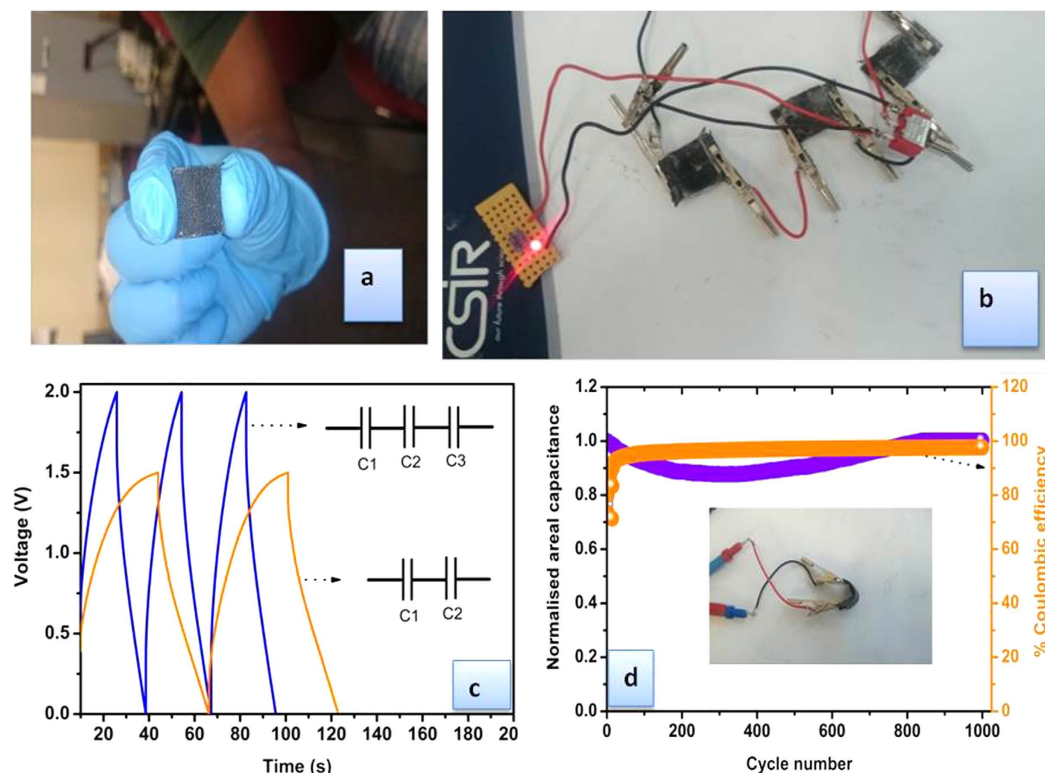


**Figure 7. (Ozoemena):** Electrochemical performances of all solid-state flexible symmetric pseudocapacitors fabricated on a carbon cloth with PVA/KOH polymer electrolyte: (a) Typical galvanostatic CD profiles at  $0.2 \text{ mA cm}^{-2}$ ; (b) Areal capacitance from CD curves as a function of current density; (c) durability test at  $0.6 \text{ mA cm}^{-2}$ ; and (d) Ragone plot of ANP<sub>mr</sub>-based all-solid-state flexible symmetric pseudocapacitors compared with similar systems reported in the literature<sup>4,15,21,37,38</sup>.

As shown in Fig. 6d, 50 h voltage-floating tests show excellent capacity retention for the ANP<sub>mr</sub> cell with areal capacitance of  $135 \text{ mF cm}^{-2}$ , and long cycle stability with an almost 100% columbic efficiency. Ragone plot of ANP<sub>mr</sub> asymmetric capacitors (Fig. 6e) exhibits extraordinary energy ( $134.6 \text{ mWh cm}^{-2}$ ) and power ( $325.6 \text{ mW cm}^{-2}$ ) densities at a current density of  $20 \text{ mA cm}^{-2}$ . These values are much higher when compared with other asymmetric pseudocapacitors (Supplementary Table-S3). Figure 6f shows the Nyquist plot of ANP<sub>mr</sub>/AC asymmetric pseudocapacitor in  $1 \text{ M Na}_2\text{SO}_4$  neutral aqueous electrolyte showed very small ESR ( $0.55 \Omega$ ) with inclined vertical line with the response time of 18 s. From Table-S2 (Supplementary Information), the ANP<sub>mr</sub>-based asymmetric cell showed similar EIS behaviour as its symmetric cell counterpart; very small  $R_s$  (*ca.*  $0.55 \Omega$ ) and total  $R_{ct}$  (*ca.*  $41 \Omega$ ), with each  $n$  value ( $n_1$  or  $n_2$ ) greater than 0.8 but less than 1, further corroborating that the pseudocapacitive behaviour of the ANP<sub>mr</sub> when used in full-cell system. In addition, it may be observed that the CV curve (Fig. 5a) is higher in the positive direction than in the negative side. This behaviour is related to the series resistance of the asymmetric configuration; the  $R_s$  value of the asymmetric configuration is much higher (*ca.*  $0.55 \Omega$ ) than the symmetric counterpart (*ca.*  $0.13 \Omega$ ), which explains why we did not observe this for the symmetric configuration.

**All-solid-state flexible symmetric pseudocapacitors.** Finally, we explored the performance of the as-prepared ANP materials (ANP<sub>mp</sub>, ANP<sub>md</sub> and ANP<sub>mr</sub>) as all-solid-state flexible symmetric pseudocapacitors using PVA-KOH polymer electrolyte and carbon cloth as substrate/current collector (Fig. 7). Also, nickel foam was used as the substrate/current collector for the ANP<sub>mr</sub>-based all-solid-state pseudocapacitors (see Supplementary Information, Figure-S4) but we found carbon cloth to be easier to handle, more flexible, with more improved electrochemical properties than nickel foam. Thus, all further studies were devoted to the use of carbon cloth. From every analysis, the ANP<sub>mr</sub> showed better electrochemical performance than the other two ANP materials (Supplementary Information, Figure-S5). Figure 7a exemplifies typical galvanostatic charge–discharge experiments, while Fig. 6b compares the specific capacitance values obtained at various current densities ( $0.1$ – $0.8 \text{ mA cm}^{-2}$ ). The ANP<sub>mr</sub> gave an excellent specific areal capacitance of  $66 \text{ mF cm}^{-2}$  at  $0.1 \text{ mA cm}^{-2}$  (Fig. 7b), and even at a higher current density of  $0.8 \text{ mA cm}^{-2}$  the capacitance remained as high as  $3 \text{ mF cm}^{-2}$ . This value is much higher compared to the values reported for other all-solid-state symmetric and asymmetric supercapacitors (Supplementary Information, Table-S4). For example, areal capacitance achieved with





**Figure 8. (Ozoemena):** (a) Photograph of the as-prepared ANP<sub>mr</sub>-based flexible all-solid-state symmetric pseudocapacitor (ASSSP); (b) Typical 3 ASSSPs connected in series and lighting up a 1.67 V LED; (c) Charge-discharge profiles of 2 and 3 ASSSPs connected in series giving 1.5 and 2 V, respectively; and (d) cycle stability measured nearly at 120° bending angle and their coulombic efficiency, insert shows picture of an ASSSP bent at 120°.

ANP<sub>mr</sub> was better than the electrochemical double layer microcapacitors which delivered 0.4–2 mF cm<sup>-2</sup> at scan rates of 1–100 mVs<sup>-1</sup><sup>27</sup>, and graphene or carbon nanotube based flexible supercapacitors that showed 3–50 mF cm<sup>-2</sup><sup>28</sup>. From the prolonged cycle stability performed at the scan rate of 0.6 mA cm<sup>-2</sup>, the ANP<sub>mr</sub> retained ~97% of its initial capacitance even after 5000 consecutive cycles (Fig. 7c). The Ragone plot (Fig. 7d) showed significantly higher energy (21.2 mWh cm<sup>-2</sup>) and power (12.7 mW cm<sup>-2</sup>) densities compared to the values reported in the literature to date for all-solid-state SCs (Fig. 6d, and Supplementary Information, Table-S4). From the Nyquist plots of ANP<sub>mr</sub> (Supplementary Information, Figure-S5(e)) the equivalent series resistance (ESR) values obtained before and after 5000 cycles were 4.5 and 23 Ω, respectively. The response time (60 ms) was lower than the reported values for solid electrolytes (80 ms), and activated carbon (700 ms)<sup>22</sup>. Indeed, detailed examination of the EIS data of the various ANP materials in Table-S2 (i.e., ANP<sub>mp</sub>, ANP<sub>md</sub> and the three ANP<sub>mr</sub> obtained at 24, 36 and 48 h) showed an interesting trend on the conductivity of the various morphologies. The three rod-like morphologies gave the least series resistance (i.e.,  $R_s \approx 5 \Omega$ ) compared to the platelet-like ( $R_s \approx 11 \Omega$ ) and dendrite-like ( $R_s \approx 25 \Omega$ ) morphologies. The total  $R_{ct}$  values decrease as follows: ANP<sub>mr@36h</sub> (ca. 140 Ω) > ANP<sub>md</sub> (ca. 115 Ω) > ANP<sub>mp</sub> (ca. 93 Ω) > ANP<sub>mr@24h</sub> (ca. 32 Ω) > ANP<sub>mr@48h</sub> (ca. 24 Ω). This is an interesting result as it clearly corroborates other electrochemical data that shows that our rod-like morphology, obtained at optimized 48 h and 200 °C, gave the best conditions for ion mobility in pseudocapacitor devices.

To further understand the reason for the high-performance of the ANP<sub>mr</sub>, we examined its specific surface area and porosity by performing the Brunauer–Emmett–Teller (BET) measurements (Supplementary Information, Fig. S6 showing the N<sub>2</sub> adsorption–desorption isotherm of the nanorods). The observed BET surface area was 214 m<sup>2</sup> g<sup>-1</sup> with an average pore size distribution of 2–20 nm with a pore volume of 0.7 m<sup>3</sup> g<sup>-1</sup> (BJH desorption), inferring the co-existence of mesoporous and microporous microrods.

As a proof of concept, Fig. 8 describes the bendability of the ANP<sub>mr</sub>-based all-solid-state symmetric pseudocapacitor (ASSSP) and its ability to light up a 1.67 V LED when connected in series. Interestingly, when the ASSSP was bent to nearly 120° and subjected to 1000 charge-discharge cycles, it was able to maintain its performance with ca. 100% coulombic efficiency.

## Conclusions

Novel  $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$  with unique morphologies (microdendrites, microplatelets and microrods) of different supercapacitive properties has been reported. The microrod morphology gave an extraordinarily high specific capacitance, power and energy densities in half-cell and full-cell configurations (i.e., symmetric and asymmetric cells, including all-solid-state flexible pseudocapacitors) in different electrolytes. The well-aligned microrods (*ca.* 35.8 nm diameter) with meso- and microporous surface enhance ion propagation and interfacial interactions compared to the long range plates (428 nm in diameter) with larger thickness or the microdendrites with smaller branched structure (*ca.* 100 nm) with uneven surface. The all-solid-state symmetric pseudocapacitor fabricated from the ANP microrods proved it can generate power even when bent to  $120^\circ$  and can drive an LED when connected in series. The study proves that rod-like morphology (with  $\sim 36$  nm width) provides a significant and promising direction for novel 1-D and 2-D materials to obtain high-performance pseudocapacitors, especially for flexible and wearable electronics. The study has opened new doors of research opportunities for this type of materials. Such opportunities include examination of other redox-active metals other than nickel, and interrogation of the impact of tuning the reported synthesis protocols (e.g., in terms of changes in temperature, solvents, reaction times, work-up conditions, etc) or new synthesis procedure on the final structures and physicochemical properties. These constitute the directions of the on-going investigations in our laboratory.

## Methods

### Synthesis and characterisation of $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$ nanorods (with micrometric length).

Analytical grade chemicals, Nickel (II) acetate tetrahydrate ( $\text{Ni}(\text{CH}_3\text{COO})_2 \cdot 4\text{H}_2\text{O}$ ), ammonium dihydrophosphate ( $\text{NH}_4\text{H}_2\text{PO}_4$ ) and ethylene glycol (EG) were procured from Sigm-Aldrich, and used as received without further purification. In a typical synthesis, 0.5 g of  $\text{Ni}(\text{CH}_3\text{COO})_2 \cdot 4\text{H}_2\text{O}$  and  $\text{NH}_4\text{H}_2\text{PO}_4$  were dissolved thoroughly in 40 ml deionized water. Subsequently, an equal amount of EG was added into the above solution (water and EG volume ratio is 1:1). After vigorous stirring for 1 h, the mixture was then transferred into an autoclave and heated at  $200^\circ\text{C}$  for 48 h. The  $\text{ANP}_{\text{mr}}$  products obtained at  $200^\circ\text{C}$  for 24 and 36 h duration were also tested for comparison. The resulted greenish yellow precipitates were thoroughly washed with deionised water and ethanol to remove any unreacted materials. Finally, the powder was dried slowly at  $50^\circ\text{C}$  in oven or left in air to dry at room temperature overnight. The microrod-like product (evident from SEM and TEM images) is abbreviated herein as  $\text{ANP}_{\text{mr}}$ . The control experiments were also performed using ethylene glycol alone or water alone at the same time period of 48 h and constant temperature of  $200^\circ\text{C}$ . From SEM and TEM, the product from water alone gave platelet-like morphology (abbreviated herein as  $\text{ANP}_{\text{mp}}$ ), while that from ethylene glycol alone gave dendrite-like morphology (abbreviated herein as  $\text{ANP}_{\text{md}}$ ).

The formation of  $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$  was investigated by PANalytical X'Pert PRO diffractometer equipped with Ni-filtered Cu K-alpha radiation ( $\lambda = 1.541841 \text{ \AA}$ ). The morphology of the as-synthesized powders was analysed using JEOL- JSM 7500F scanning electron microscope operated at 2.0 kV. TEM and HRTEM images were obtained from JEOL-Jem 2100 microscope operated at an acceleration voltage of 200 kV. BET measurements were performed to measure the specific surface area and pore size based on the  $\text{N}_2$  adsorption–desorption method by using Micromeritics TriStar II instrument.

**Materials, preparation and pseudocapacitor fabrication.** Nickel foam (Celmet: thickness = 1.6 mm, surface area  $7500 \text{ m}^2$ , cell size = 0.5 mm, 48–52 cells per inch) and carbon cloth (B-1/C, E-TEK) were used as substrates and current collectors in the fabrication of the half-cell (three-electrode system) and full-cell (two-electrode systems), respectively. Prior to use, the nickel foam was properly cleaned as we described before<sup>39</sup> by first sonicating in 1 M HCl solution for 30 min, washed several times with copious amount of de-ionized water, and then dried under vacuum. The electrolyte materials, polyvinyl alcohol (PVA), potassium hydroxide (KOH) and sodium sulphate ( $\text{Na}_2\text{SO}_4$ ) were procured from Sigma-Aldrich and used as received. For the three-electrode system, the electrode materials were prepared by coating a slurry mixture of  $\text{NH}_4\text{NiPO}_4 \cdot \text{H}_2\text{O}$  (ANP), carbon black and polyvinylidene fluoride (PVDF) (80:15:5 weight ratio) on a piece of clean nickel foam and dried overnight in a vacuum oven at  $80^\circ\text{C}$ . The mass of the active material on nickel foam was 0.32 mg for  $\text{ANP}_{\text{mr}}$ , 0.35 mg for  $\text{ANP}_{\text{md}}$  and 0.34 mg for  $\text{ANP}_{\text{mp}}$ . For the symmetric and asymmetric systems, the electrode materials were prepared by coating a slurry mixture of  $\text{ANP}_{\text{mr}}$ , activated carbon (AC, Norit<sup>®</sup> supra 30) and PVDF (50:40:10 weight ratio) on carbon cloth (disc =  $1.6 \text{ cm}^2$ ). The symmetric ( $\text{ANP}_{\text{mr}}//\text{ANP}_{\text{mr}}$ ) and asymmetric ( $\text{ANP}_{\text{mr}}//\text{AC}$ ) pseudocapacitors were performed in 3 M KOH and 1 M  $\text{Na}_2\text{SO}_4$ , respectively. The all-solid-state flexible symmetric pseudocapacitor (square =  $1 \text{ cm}^2$ ) was fabricated in a similar manner as the symmetric cells but using a polymer gel electrolyte (PVA-KOH). In a typical polymer electrolyte preparation, 8 g of PVA and 4 g of KOH were dissolved in deionised water (40 ml) and the mixture was stirred at  $90^\circ\text{C}$  for 1 h to form a gel. Various forms of the ANP systems ( $\text{ANP}_{\text{mp}}$ ,  $\text{ANP}_{\text{md}}$ ,  $\text{ANP}_{\text{mr}}$  as well as the  $\text{ANP}_{\text{mr}}$  obtained at different 24 and 36 h at the same temperature) were investigated as possible candidates as all-solid-state flexible pseudocapacitors.

**Electrochemical measurements and calculations.** All electrochemical tests involving cyclic voltammetry (CV), electrochemical impedance spectroscopy (EIS) and galvanostatic charge-discharge analysis (including voltage-holding experiments) were performed at room temperature using computer-controlled

Multi-channel Potentiostat/Galvanostat Bio-Logic VMP3 work station driven by EC-Lab® v10.40 software with Z-fit tool for EIS data analysis. In a typical three-electrode system, ANP-coated nickel foam was used as the working electrode, platinum mesh as the counter electrode, and Ag/AgCl (3 M KCl) as the reference electrode, in 3 M KOH aqueous solution. EIS measurements were carried out in the frequency ranging from 10 kHz to 10 mHz at the open circuit voltage with AC voltage amplitude of 1.5 mV.

For the half-cells (3-electrode configurations), the specific capacitance ( $C_{sp}$ ), maximum specific power density ( $P_{max}$ ) and specific energy density ( $E_{sp}$ ) was evaluated using the conventional equations (2–4)<sup>40</sup>:

$$C_{sp}(F\ g^{-1}) = \frac{\int i\ dt}{\Delta V\ m} = \frac{i\ \Delta t}{\Delta V\ m} \quad (2)$$

$$P_{max}(W\ kg^{-1}) = \frac{i\Delta V}{m} \quad (3)$$

$$E_{sp}(Wh\ kg^{-1}) = \frac{it\ \Delta V}{m} \quad (4)$$

where:  $i$  (A) is the current,  $\Delta V$  (V)/ $\Delta t$  (s) the slope of the discharge curve, and  $m$  (g) the mass of active materials (ANP and carbon),  $V$  (V) is the voltage obtained during charge. For the full cells (2-electrode configurations), the corresponding parameters were obtained using conventional equations (5–9):

$$C(F) = \frac{i\Delta t}{\Delta V} \quad (5)$$

$$C_{sp}(F\ cm^{-2}) = \frac{4C}{S} \quad (6)$$

$$P_{max}(W\ cm^{-2}) = \frac{V^2}{4SR_{ir}} \quad (7)$$

where,

$$R_{ir}(\Omega) = \frac{\Delta V_{ir}}{2I} \quad (8)$$

$$E(Wh\ cm^{-2}) = \frac{CV^2}{2S} \quad (9)$$

where  $i$  (A) is the applied current,  $\Delta V$  (V)/ $\Delta t$  (s) the slope of the discharge curve and  $S$  (cm<sup>2</sup>) the total geometric surface area of the two electrodes,  $C$  (F) the calculated capacitance,  $V$  (V) is the maximum voltage obtained during charge, and  $R_{ir}$  is the internal resistance which is determined from the voltage drop at the beginning of each discharge, while the  $\Delta V_{ir}$  represents the voltage drop.

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## Acknowledgements

This work was supported by the CSIR (South Africa), the South Africa's Department of Science and Technology (DST) and National Research Foundation (NRF) under the “Nanotechnology Flagship Programme” (*supercapacitors and fuel cell project*, Grant no. 69849). CSIR is a partner in the CREATE-Network funded by the Marie Skłodowska-Curie Research and Innovation Staff Exchange (RISE). The authors are very grateful to Dr John R. Miller (JME, Inc. and Case Western Reserve University, Cleveland, Ohio, USA) for assisting us in validating the initial data of this ANP<sub>mr</sub>-based pseudocapacitor in his laboratory and offering some useful comments. Dr Kumar Raju thanks the CSIR and DST/NRF for post-doctoral research fellowship. We appreciate Drs C. Jafta, P. Ejikeme and K. Makgopa for their useful comments during some of the experiments.

## Author Contributions

Dr. K.R. is a postdoc fellow under the supervision of Prof K. O. The project was conceived by both authors. Kumar carried out the experiments, prepared all the figures. K. O. supervised the work from the experiments to data interpretation, and wrote the main manuscript text assisted by the initial draft submitted by Kumar. All authors reviewed the manuscript.

## Additional Information

**Supplementary information** accompanies this paper at <http://www.nature.com/srep>

**Competing financial interests:** The authors declare no competing financial interests.



**How to cite this article:** Raju, K. and Ozoemena, K. I. Hierarchical One-Dimensional Ammonium Nickel Phosphate Microrods for High-Performance Pseudocapacitors. *Sci. Rep.* **5**, 17629; doi: 10.1038/srep17629 (2015).



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