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# Full organic aqueous battery based on TEMPO small molecule with millimeter-thick electrodes

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**ABSTRACT:** Thick electrodes with sodium and even anions intercalation organic compounds integrated in a neutral-pH aqueous battery offer unique advantages in terms of round trip efficiency, environmental impact, and scalability for off or in-grid renewable energy storage. Herein we report on the first anion-rocking chair / dual-ion organic battery. The latter reaches 35 Wh/kg<sub>materials</sub> at C/8 rate. It shows remarkable cyclability and coulombic efficiency in a cheap and neutral NaClO<sub>4</sub> electrolyte pouch cell with highly loaded millimeter-thick electrodes (5 mAh/cm<sup>2</sup>). This achievement is based on a thorough study of a commercial (2,2,6,6-tetramethylpiperidin-1-yl)oxyl (TEMPO) benzene derivative namely 4-hydroxy TEMPO benzoate and its naphthalene analog (4-carboxy TEMPO naphthalate) as positive electrode materials, and a bipyridinium-naphthalene oligomer as the negative electrode. Combined UV-Vis spectroelectrochemistry and operando XRD account for the much improved cyclability of the hydrophobic 4-carboxy TEMPO naphthalate at the expense of a lower specific capacity. This trend is reversed in the case of the 4-hydroxy TEMPO benzoate derivative. Results show kinetic limitations of the 4-hydroxy TEMPO benzoate are associated with the surrounding composite electrode, while inner-grain ionic and/or electronic transports play a decisive role for the 4-carboxy TEMPO naphthalate.

## Introduction

Aqueous batteries constitute a novel and promising technology towards renewable energy storage as they are inherently safe, minimize cost and environmental impact by comparison to other battery technologies<sup>1-4</sup>, thus offering a low cost and greener alternative to Li-ion systems. This approach is still affiliated however, with energy densities below 50Wh/kg<sub>cell</sub>. Furthermore, the relatively narrow electrochemical window available in aqueous media together with costs and abundance issues makes even more challenging the development of appropriate materials. The latter should possess optimal potential as well as high chemical and electrochemical stability. As a result, the choice of greener and more abundant organic materials appear as a logical step. So far only two families of organic compounds have been demonstrating encouraging performance as positive electrode active materials in aqueous media: i) the p-type conjugated tertiary amine PTPAm showing interesting experimental capacity and good cyclability in 21 m LiTFSI water-in-salt electrolyte<sup>5</sup> and ii) the nitroxyl radicals, more precisely the 2,2,6,6-tetramethylpiperidinyloxy-4-yl (TEMPO) derivatives. The latter is the most studied p-type redox systems in aqueous electrolyte<sup>6-11</sup>. TEMPO undergoes a one electron oxidation process to give the oxoammonium cation, the charge being counterbalanced by the uptake of one anion from the electrolyte (Figure 1).

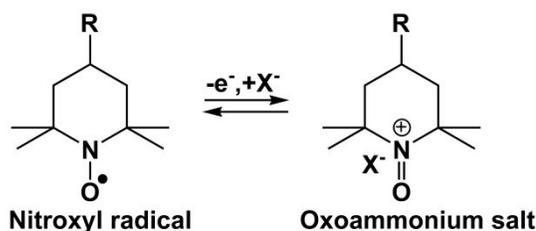


Figure 1. Schematic representation of the electrochemical reaction of TEMPO derivatives.

Up to now, TEMPO derivatives in aqueous batteries have been mainly developed in the form of radical polymers. In this regard, Oyaizu and Nishide's group have demonstrated that the achievement of high molecular weight and proper volume swell ratio is fundamental for the good performance of the polymer<sup>12</sup>. They extensively investigated TEMPO's derivatives<sup>6-10,12,13</sup>, highlighting the difficulties in obtaining proper water compatible TEMPO polymer. As a result, the poly(2,2,6,6-tetramethylpiperidinyloxy-4-yl acrylamide) (referred to as PTAm) revealed the most interesting performance, enabling the development of two anionic rocking chair batteries. Noteworthy, until recently, electrodes were based on thin films of these polymers (from 50 nm to 3.2 μm)<sup>12</sup>. However, because low cost ( $\leq 100$  USD per kWh<sub>installed</sub>) is a sine qua non condition to insure the economic viability of electrochemical systems for grid storage, aqueous batteries need to be built with ultra-thick

(>mm) electrodes. In this context, the same group has recently proposed thick composite electrodes of 3 mAh/cm<sup>2</sup> by hybridizing PTAm with a 3D self-assembled mesh of single-walled carbon nanotubes (SWCNT)<sup>7</sup>. Even though the cost of SWCNT can potentially be an issue, impressive performance has been obtained with only 1 to 5 wt.% SWCNT. Although all these approaches are effective in that they successfully convey organic materials with interesting performance as positive electrode material for neutral aqueous battery, a full cell with thick organic based electrodes of several mAh/cm<sup>2</sup> still remain to be demonstrated to the best of our knowledge. A different approach is proposed in the present work based on “small TEMPO molecules”, rather than polymeric structures. We note that an example has been recently reported in the field of supercapacitor by Itoi et al. who adsorbed the HTB (1-Piperidinyloxy,2,2,6,6-tetramethyl-4-(2-oxo-2-phenylethyl)) in the mesoporosity of active carbon (AC)<sup>14</sup>. Thanks to the huge contact area between the finely dispersed HTB molecules and the conductive AC surface, fast redox reactions of HTB could be achieved in aqueous 1 M H<sub>2</sub>SO<sub>4</sub>. Moreover, the strong adsorption capability of AC counteracts the dissolution of HTB in the electrolyte. On the contrary, it is worth pointing out that this method is inherently restricted to the supercapacitor field because of the high loading of conductive agent. The best performance (81% of capacity retention after 2000 cycles at 1 A/g) were typically obtained with 70 wt.% AC. Importantly, this kind of strategy, that is well known especially in the Li-S battery field since the work of Nazar<sup>15</sup>, can only mitigate dissolution but not suppress it. Here we report on two different strategies that have been followed with the aim to produce a low cost and safe aqueous battery including p-type “small TEMPO molecules”. Our approach will be supported by in-depth study of electrochemical behaviors as well as demonstration of potentialities and limitations of some of these materials in full cells of up to 5 mAh/cm<sup>2</sup> using a new bipyridinium-naphthalene diimide oligomer as the negative electrode.

## Experimental section

General procedure for the synthesis of TEMPO-bearing derivatives (note that all characterizations relative to synthesis of materials including, <sup>1</sup>H NMR, FTIR, TGA/DSC and XRD are reported at the end of the supplementary information file (Figure S20 to S42) to facilitate the reading of the article):

All commercially obtained solvents and reagents were used without further purification unless noted below. 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide (EDC), 4-dimethylaminopyridine (DMAP), 4-hydroxyTEMPO benzoate, 4-carboxyTEMPO, 4-hydroxyTEMPO, phenol, 2-hydroxynaphthalene, benzoic acid and 2-naphthalic acid were purchased from Sigma Aldrich. The PTAm polymer has been prepared and supplied by Nishide and Oyaizu's group.

**Synthesis procedure of 4CTs (4-carboxyTEMPO derivatives):**  
To a stirred solution of 4-carboxyTEMPO (1 eq.) DMAP (0.3 eq.) and the corresponding alcohol (0.7 eq) in dichloromethane cooled to 0°C in an ice bath, was added EDC (1 eq), and the mixture was warmed to ambient temperature over several hours and stirred for 48-72 h, following the reaction advancement through thin layer chromatography. The

solution was diluted with dichloromethane and was washed with water (2 ×), a saturated aqueous solution of sodium bicarbonate (1 ×), and brine (1 ×). The organic layer was dried over sodium sulfate, concentrated in vacuum and purified by silica gel column chromatography to afford the compounds as an orange powder.

**4CT-Benzene:** (yield 75%) <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 7.36 – 7.47 (m, 3H), 7.21 – 7.29 (m, 2H), 2.95 (tt, J = 12.9 Hz, 1H) 1.96-1.97 (m, 2H), 1.58 (t, J = 12.8 Hz, 2H), 1.12 (s, 6H), 1.09 (s, 6H). HRMS (ESI)<sup>+</sup> m/z calculated for C<sub>16</sub>H<sub>22</sub>NO<sub>3</sub>Na 299.1497 [M+Na]<sup>+</sup> found 299.1503. FTIR (cm<sup>-1</sup>): 2830-3000 (rocking CH<sub>2</sub>), 1744 (C=O ester), 1366 (N-O stretch).

**4CT-Napthalene:** (yield 80%) <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 7.94 (ddd, J = 12.1, 8.9, 5.9 Hz, 3H), 7.67 (d, J = 2.4 Hz, 1H), 7.55 (qd, J = 5.1, 1.8 Hz, 2H), 7.31 (dd, J = 8.9, 2.4 Hz, 1H), 3.02 (tt, J = 12.8, 3.4 Hz, 1H), 2.06 – 1.87 (m, 2H), 1.62 (t, J = 12.8 Hz, 2H), 1.14 (s, 6H), 1.11 (s, 6H). HRMS (ESI)<sup>+</sup> m/z calculated for C<sub>20</sub>H<sub>24</sub>NO<sub>3</sub>Na 349.1654 [M+Na]<sup>+</sup> found 349.1658. FTIR (cm<sup>-1</sup>): 2760-3000 (rocking CH<sub>2</sub>), 1744 (C=O ester), 1366 (N-O stretch).

**Synthesis procedure of 4HTs (4-hydroxyTEMPO derivatives):**  
To a stirred solution of 4-hydroxyTEMPO (1 eq.) DMAP (0.3 eq.) and the corresponding acid (0.7 eq) in dichloromethane cooled to 0°C in an ice bath, was added EDC (1 eq), and the mixture was warmed to ambient temperature over several hours and stirred for 48-72 h, following the reaction advancement through thin layer chromatography. The solution was diluted with dichloromethane and was washed with water (2 ×), a saturated aqueous solution of sodium bicarbonate (1 ×), and brine (1 ×). The organic layer was dried over sodium sulfate, concentrated in vacuum and purified by silica gel column chromatography to afford the compound as a reddish powder.

**4HT-Benzene:** (yield 53%) <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 7.98 – 7.91 (m, 2H), 7.75 – 7.61 (m, 1H), 7.59 – 7.49 (m, 2H), 5.30 – 5.00 (m, 1H), 1.98 (ddd, J = 11.3, 4.3, 1.7 Hz, 2H), 1.61 (t, J = 11.7 Hz, 2H), 1.13 (s, 12H). HRMS (ESI)<sup>+</sup> m/z calculated for C<sub>16</sub>H<sub>22</sub>NO<sub>3</sub> 276.1600 [M]<sup>+</sup> found 276.1603. FTIR (cm<sup>-1</sup>): 2834-3016 (rocking CH<sub>2</sub>), 1715 (C=O ester), 1363 (N-O stretch).

**4HT-Napthalene:** (yield 46%) <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.67 – 8.55 (m, 1H), 8.18 – 8.09 (m, 1H), 8.08 – 7.91 (m, 3H), 7.74 – 7.58 (m, 2H), 5.26 (tt, J = 11.2, 4.2 Hz, 1H), 2.08 – 1.99 (m, 2H), 1.67 (t, J = 11.7 Hz, 2H), 1.15 (s, 12H). HRMS (ESI)<sup>+</sup> m/z calculated for C<sub>20</sub>H<sub>26</sub>NO<sub>3</sub> 328.1913 [M+2H]<sup>2+</sup> found 328.1916. FTIR (cm<sup>-1</sup>): 2838-3016 (rocking CH<sub>2</sub>), 1712 (C=O ester), 1365 (N-O stretch).

## Chemical oxidation of 4CT-Napthalene:

The chemical oxidation of 4CT-Napthalene has been performed following the already reported procedure<sup>16</sup>. Glacial acetic acid (2 mL) and hydrogen peroxide (5 mL, 29.0–32.0%) was added to 1 M NaClO<sub>4</sub> aqueous solution (100 mL). The solution was added dropwise to the stirred 4CT-Napthalene suspension in 1 M NaClO<sub>4</sub> at ambient temperature. The mixture was left under vigorous stirring during 24 hours.

## Characterization techniques:

<sup>1</sup>H-NMR spectra were acquired using a Bruker ARX 300 MHz spectrometer. Spectra were recorded at room temperature, chemical shifts are written in ppm, and coupling constant in Hz. Multiplicity is presented in the following way: s = singlet, d = doublet, t = triplet, q = quintuplet, m = multiplet. High resolution mass spectrometry (HRMS) has been recorded on a XEVO G2-XS QTOF (Waters) operating on ESI<sup>+</sup>. Thermal analyses were performed with a NETZSCH STA 449F3 device under Argon atmosphere. Fourier transform infrared spectroscopy (FTIR) spectra were collected with a Bruker Vertex 70 device employing KBr pellets, using a DTGS detector at a resolution of 4 cm<sup>-1</sup>. Powder XRD diagrams have been recorded with the Bruker D8 Advanced system. X-ray single crystal diffraction data of 4CT-Naphthalene were collected on an Agilent SuperNova diffractometer equipped with Atlas CCD detector and mirror monochromated microfocus Cu-K<sub>α</sub> radiation ( $\lambda = 1.54184 \text{ \AA}$ ). The structure was solved and refined on F2 by full-matrix least-squares techniques using SHELX programs. All non-hydrogen atoms were refined anisotropically and hydrogen atoms were placed in calculated positions. Multiscan empirical absorption was corrected using CrysAlisPro program (CrysAlisPro, Agilent Technologies, V1.171.37.35 g, 2014). A summary of the crystallographic data and the structure refinement parameters is provided in Table S1. Operando X-ray diffraction patterns were recorded using a PANalytical X'Pert Pro diffractometer operated in Bragg-Brentano reflection geometry with a Long Line Focus Cu-anode X-ray source, and a X'Celerator RTMS detector. Data were collected in a (10-30)<sup>o</sup> 2 $\theta$  range with a step of 0.033<sup>o</sup> and an acquisition time of 5'16'' per diagram. Measurements were performed with a Multi Purpose Sample Stage (MPSS) able to accommodate the in situ Swagelok-type cell. Note that the common beryllium window was removed because it was unstable in the tested electrochemical conditions. Therefore, we replaced the beryllium window by a thin Kapton<sup>®</sup> foil (X-ray film polyimide Kapton<sup>®</sup>, PANalytical B-V); thickness: 10  $\mu\text{m}$ , diameter: 6.4 mm. A threefold-size (in mAh) carbon electrode (80 wt.% activated carbon Norit 1600, 10 wt.% Ketjenblack (KB) EC-600JD (AkzoNobel), and 10 wt.% polytetrafluoroethylene (PTFE)) was used as negative and reference electrode whereas the positive electrode composition was 70 wt.% of active material, 25 wt.% KB, and 5 wt.% PTFE. This mixture was pressed at 10 tons on a stainless steel (AISI 316L) grid current collector. Thick electrodes of 4HT-benzene and DNVBr were obtained pressing at 5 tons a 70 wt.% active material, 25 wt.% KB, 5 wt.% PTFE mixture into a pellet. The latter was then inserted in a specially designed mold and pressed at 2 tons between two stainless steel grids. Peak fitting was performed using Winplotr using a Gaussian function with constant FWHM defined by  $U=0.07^o$ . UV-Vis spectroelectrochemistry was conducted by inserting a home-made three-electrode quartz cell into a Shimadzu UV-2501 PC spectrometer. Each scan was recorded from 250 to 800 nm within approx. 4 min. Calibration curves were performed by dissolution of 4HT-Bz in acetonitrile and 4CT-Napht in ethylacetate. Both these solvents show an absorbance cut-off below 270 nm which allows to follow the signal of the two compounds. Solutions of these compounds show the same color as in the solid state although being more pale as dissolution increases.

## Electrochemical study:

The various electrochemical measurements were performed using SP 300, MPG-2 and VMP3 potentiostats from Bio-logic SAS (Seysinnet-Pariset, France). All the TEMPO derivatives have been characterized by galvanostatic cycling in aqueous NaClO<sub>4</sub> 1 M electrolyte. 4HT-Benzene was also tested in highly concentrated aqueous electrolyte (8 M). Unless otherwise specified, the electrode composition is 70 wt.% active material, 25 wt.% KB, 5 wt.% PTFE. The electrode mixtures have been prepared vigorously mixing the three components in the dry state with the use of a mortar. A co-precipitated mixture has been also prepared. The 4CT-Naphthalene (70 wt.%) has been dissolved in the minimum volume of dimethylsulfoxide (DMSO) then 25 wt.% KB has been added to the solution. The suspension has been vigorously stirred during one hour. Subsequently, it has been added dropwise into a stirred water solution. The precipitate has been finally filtered, washed and dried at 60°C under vacuum overnight. 5 wt.% PTFE has been finally added mixing the powders in a mortar. Caution was taken in preparing thin electrodes with at 7 to 10 mg of active material/cm<sup>2</sup> (0.7-1 mAh/cm<sup>2</sup>) pressed at 5 tons on a stainless steel (AISI 316L) grid current collector. Thicker electrodes of 80 mg/cm<sup>2</sup> were made in two steps. First a pellet was obtained by pressing at 5 tons the active material mixture. This pellet was then introduced between two stainless steel grids and pressed at 2 tons in order to favor the current collector-electrode mixture contacts. A threefold-oversized counter-electrode (80 wt.% activated carbon Norit 1600, 10 wt.% KB, and 10 wt.% PTFE) and a Saturated Calomel Electrode from ALS Co., Ltd Japan (ref: RE-2BP) (SCE) as reference were used in all aqueous tests. Thick electrode full cells have been prepared based on DNVBr 7.8 mAh/cm<sup>2</sup> (80 mg/cm<sup>2</sup>, 1.09 mm for 4HT-benzene) and 8 mAh/cm<sup>2</sup> (80 mg/cm<sup>2</sup>, 0.86 mm for DNVBr). Such cell balancing was found to result in the highest energy density. All full cells tests were performed within an argon-filled glovebox with less than 1 ppm O<sub>2</sub>. An example of thick electrodes full cell has been also made using a pouch cell three electrodes configuration. Two microfiber glass separators were used, the first to isolate the positive (81 mg/cm<sup>2</sup>, 7.9 mAh/cm<sup>2</sup> for 4HT-benzene) and the negative electrode (86 mg/cm<sup>2</sup>, 9 mAh/cm<sup>2</sup> for DNVBr) and another between the negative and reference electrode. This cell balancing was chosen to maximize the energy density. not screened for pouch cells. Pouch cells reference electrodes were composed of Li<sub>0.5</sub>FePO<sub>4</sub>. The cell used was an aluminum bag (8 cm × 5.5 cm) from Sidec (model CA 40) sealed at 190°C, where 1.5 mL of electrolyte (8 M NaClO<sub>4</sub>) have been added. The pouch cell has been prepared in an argon-filled glovebox and tested outside. Regular cyclic voltammetry experiments in solution have been conducted by dissolving 1 mM of the four TEMPO derivatives in acetonitrile/NaClO<sub>4</sub> 2.5 M as the electrolyte and recorded with a Pt micro disc electrode ( $\varnothing = 1.5 \text{ mm}$ ). A large Pt wire has been used as counter electrode whereas the reference electrode consisted in AgNO<sub>3</sub>/Ag from ALS Co., Ltd Japan (ref: RE-7, TBAClO<sub>4</sub> 10 mM in acetonitrile). Potential values in Figure S1 are reported versus an aqueous SCE electrode (Ag/AgNO<sub>3</sub> = +0.36 V vs SCE). Temperature of the electrochemical tests is 22°C. Electrochemical impedance measurements were performed from 180 kHz to 100 mHz with a perturbation voltage of 10 mV. Power tests have been carried out on discharge using the single discharge technique consisting of

relaxation periods until  $\Delta E/\Delta t = 1$  mV/h between each current pulse from  $I_{\max}$  to  $I_{\min}$ . All experiments were conducted twice in order to ensure reproducibility. Maximum experimental errors related to capacity values were kept below 5% using our protocols and set-up.

## Results and discussion

Four TEMPO bearing redox active molecules have been investigated (Figure 2): the commercially available 4-hydroxy TEMPO benzoate (4HT-Benzene), the isomer 4-carboxy TEMPO benzoate and their naphthalene derivatives (4HT-Naphthalene and 4CT-Naphthalene). The last three compounds have been produced upon esterification reaction in order to obtain the benzoate or naphthalate ester in the 4-position of the piperidine ring. Importantly in regard to the targeted application, these compounds can readily be obtained in gram scales by a one pot reaction. The materials have been fully characterized by means of Electrospray ionization high-resolution mass spectrometry (ESI-HRMS), liquid nuclear magnetic resonance (NMR) and Fourier transform infrared (FTIR) spectroscopies as well as thermal analyses (for further details see in supplementary information).

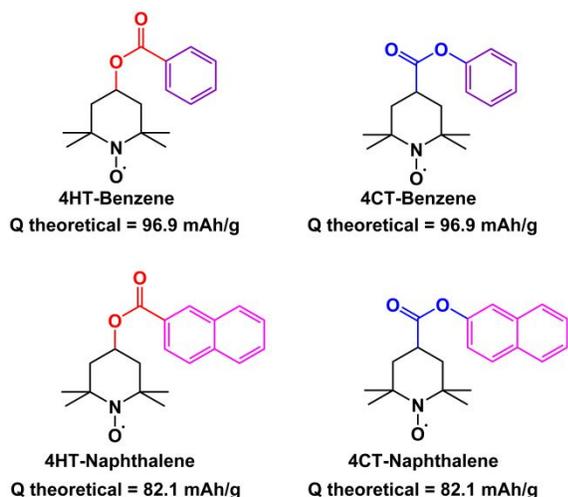


Figure 2. Molecular structure of the four TEMPO derivatives and their theoretical specific capacity.

Single crystals of the 4CT-Naphthalene compound could be obtained and its crystal structure compared to that of one of the two benzene derivative, the 4HT-Benzene. X-ray single crystal structure data study reveals that 4CT-Naphthalene crystallizes in the non-centrosymmetric  $P21$  space group and consists of 1D supramolecular arrangements of molecules (Figure 3a). The  $d_{\text{N-O}}$  distance of 1.286(3) Å found in 4CT-Naphthalene is typical of a free radical single bond N-O<sup>17</sup>. Moreover, the chair conformation of the piperidine ring clearly indicates that the nitrogen is in  $sp^3$  hybridization state confirming N-O is a single bond. The organization of 4CT-Naphthalene radicals into 1D chains is achieved via weak C-H $\cdots$ O interactions between oxygen and hydrogen of the radical rings with a N-O $\cdots$ O distance of 5.766(2) Å (Table 1 and Figure 3b). In the  $bc$  plane, N-O $\cdots$ O distances are shorter than in 1D chains (5.259(3) Å). Comparing structural features of 4CT-Naphthalene with that of 4HT-Benzene<sup>18</sup> brings valuable information: N-O $\cdots$ O distances in 4HT-Benzene are not only longer, but an inversion also occurs as the shortest N-O $\cdots$ O distances run along the 1D chains (6.128(7) Å) whereas the longest distances are found in the  $bc$  plane (6.264(7)-6.287(6) Å).

Table 1. Weak C-H $\cdots$ O interactions in 4CT-Naphthalene

C—H $\cdots$ O	C—H (Å)	H $\cdots$ O (Å)	C $\cdots$ O (Å)	C—H $\cdots$ O (°)
C <sub>2</sub> —H <sub>2</sub> C $\cdots$ O <sub>1</sub> <sup>i</sup>	0.96	2.62	3.553 (3)	163
C <sub>5</sub> —H <sub>5</sub> $\cdots$ O <sub>1</sub> <sup>i</sup>	0.98	2.64	3.623 (3)	178

Symmetry code: (i)  $x+1, y, z$ .

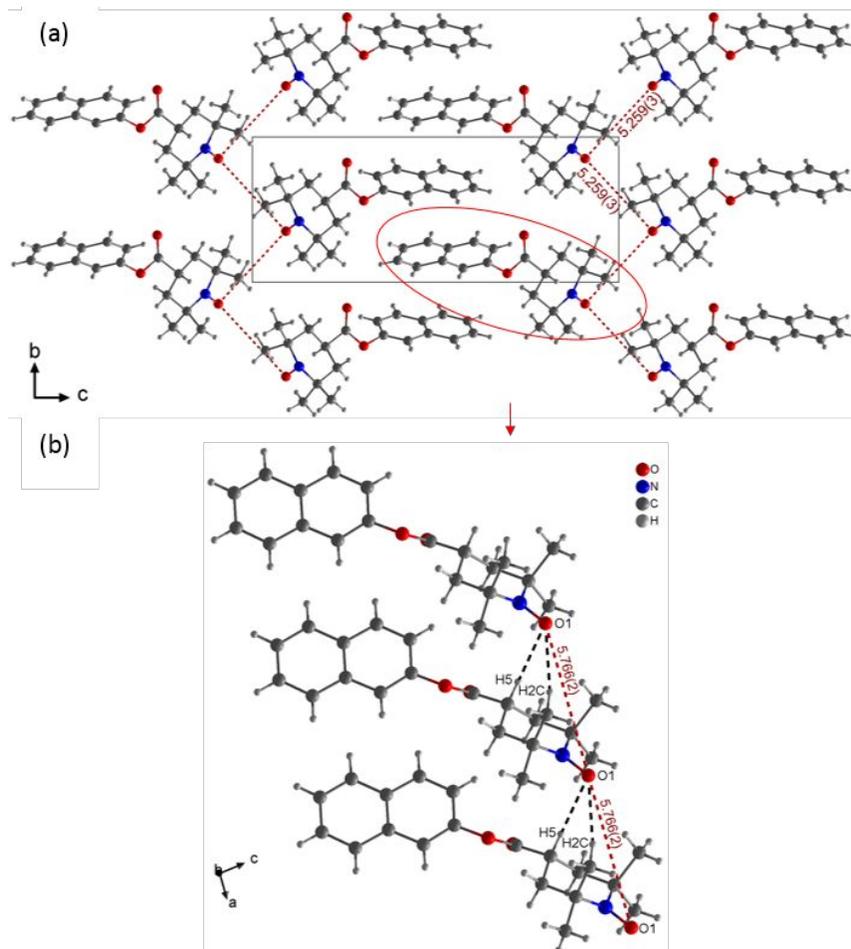


Figure 3. Structural features of 4CT-Naphthalene a) Crystal packing and N-O...O-N distance in the *bc* plane and b) N-O...O-N distance into 1D chains and weak C-H...O interactions represented in dashed black line in 4CT-Naphthalene.

The electrochemical activity of the four materials has been firstly characterized by cyclic voltammetry in the solubilized state using acetonitrile/NaClO<sub>4</sub> 2.5 M as the electrolyte (Figure S1). All derivatives exhibit a single reversible process with  $\Delta E_p$  approaching the reversible one electron value of 58 mV. The characteristic peak potentials on oxidation was not found to vary significantly indicating the substituent group has a weak inductive effect if any. Lastly, derivation of diffusion coefficients (Figure S2 and Table S2, see details in supplementary information) are all similar and comparable to the value reported for the 2,2,6,6-tetramethylpiperidiny-*N*-oxyl (TEMPO) radical<sup>13</sup>.

As shown by galvanostatic experiments, the electrochemical profile is strongly modified in the solid state with the occurrence of two electrochemical steps (Figure 4). This point is reiterated using cyclic voltammetry for the two naphthalene compounds (Figure S3). The reaction starts on oxidation by a smooth increase of the incremental capacity (I) followed by a

plateau type process (II). We note the capacitive contribution of the carbon additive (25 wt.%) to the overall electrochemical response is negligible (Figure S4). The two benzene derivatives appear quite unstable in 1 M NaClO<sub>4</sub> electrolyte showing an important capacity fade during cycling (Figure 5). 4HT-Benzene in particular shows a continuous fading of 70% in 100 cycles (123 hours). Accordingly, UV-Vis analysis of the corresponding electrolyte upon 100 cycles clearly highlights the dissolution of the material (Figure S5). Interestingly, the introduction of more hydrophobic naphthalene moiety significantly improves the cyclability at the expense of a lower specific capacity. In particular, the initial capacity of 4CT-Naphthalene isomer increases from 35 mAh/g up to 44 mAh/g in 60 cycles followed by a capacity loss restricted to 4.5% during the following 40 cycles. Consequently, no color change could be detected in the electrolyte, while chemical oxidation with hydrogen peroxide did not result in any apparent dissolution of the compound (see Experimental part).

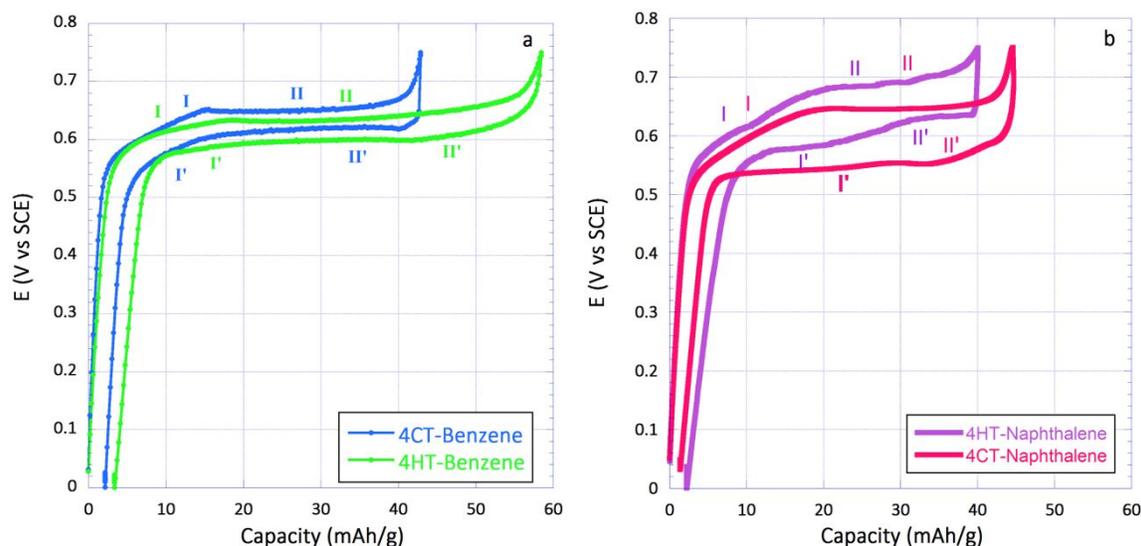


Figure 4. Galvanostatic curves for 4CT-Benzene (blue) on cycle 19, 4HT-Benzene (green) on cycle 1, 4CT-Naphthalene (pink) on cycle 60 and 4HT-Naphthalene (violet) on cycle 22 at 0.075 A/g in 1 M NaClO<sub>4</sub> aqueous electrolyte. Selected cycles correspond for each compound to the maximum capacity obtained during the first 100 cycles as shown in Figure 5. The areal capacity of electrodes are about 0.7 mAh/cm<sup>2</sup>.

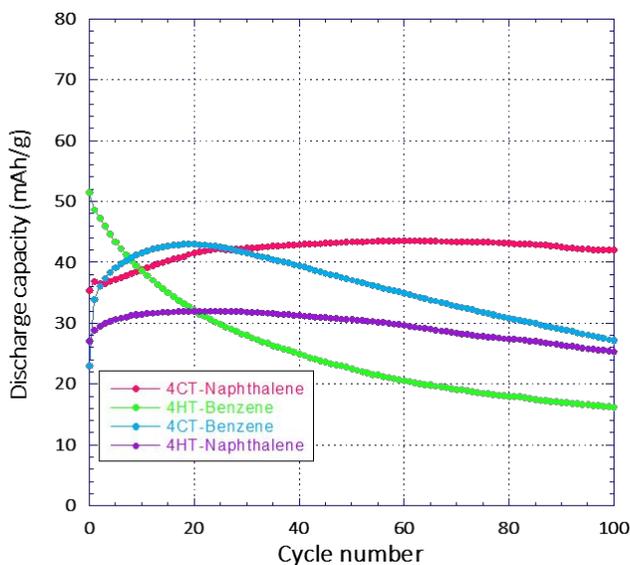


Figure 5. Cyclability curves for the four compounds in 1 M NaClO<sub>4</sub> at 0.075A/g (4CT-Benzene (blue), 4HT-Benzene (green), 4CT-Naphthalene (pink) and 4HT-Naphthalene (violet)). The areal capacity of electrodes are about 0.7 mAh/cm<sup>2</sup>.

To get a better insight, UV-Vis spectroelectrochemistry has been performed during 158 hours while the sample was cycled at 0.075 A/g (194 cycles) within the usual potential window of (0; 0.75) V vs SCE. Simultaneously with the initial increase of capacity during the first 60 hours, a UV band appears at 300 nm (Figure 6). The latter rises during 80 hours and stabilizes throughout the remaining cycles to 0.07mM. This value corresponds in fact to a negligible part of the initial compound (5 wt.%) (Figure S6). The question is now whether, this concentration corresponds to a solubility limit or alternatively,

is limited because part of the active material does not participate in the electrochemical reaction. A first hint is given by the fact that irrespective of the electrolyte to active material mass ratio, 4CT-Naphthalene always gives the same order of maximum specific capacity ( $40 \pm 5$  mAh/g, 52% of the theoretical capacity). Besides, there is no clue so far regarding the source of capacity limitation. In this goal the first issue to be addressed consist in determining if, as suggested above, some grains are poorly electrically connected, or alternatively if all grains react but the electrochemical reaction is thermodynamically limited

because of overwhelming charge repulsion and/or lack of space to host ions within the oxidized host material.

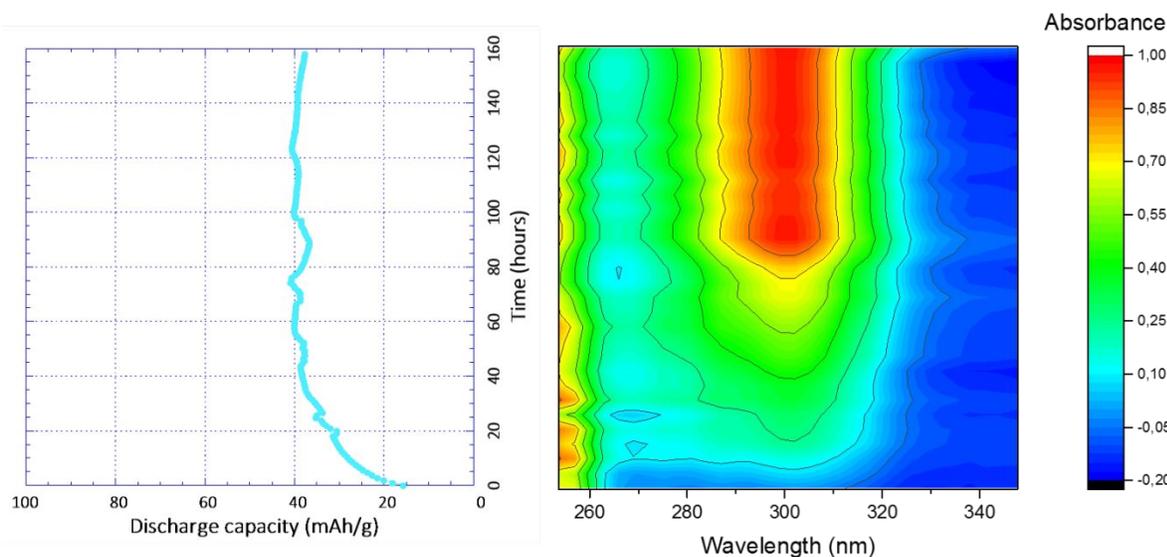


Figure 6. Operando UV-Vis spectroelectrochemical spectra (right) and corresponding evolution of the discharge capacity of 4CT-Naphthalene upon cycling (left) measured at 0.075A/g in 1 M NaClO<sub>4</sub> aqueous electrolyte and within (0; 0.75) V vs SCE potential window. The areal capacity of electrodes are about 0.7 mAh/cm<sup>2</sup>.

To address this point, *Operando* XRD measurements of 4CT-Naphthalene have been performed during galvanostatic cycling. Noteworthy, each scan (10-30°) was 5'16'' therefore enabling a reliable description of the structural behavior by approx. 24 scans for each charge or discharge. The compound has been first cycled within (-0.1; 0.95) V vs SCE. The corresponding overall structural evolution is given in Figures 7a for the first cycle (the first four cycles are shown in Figure S7). The initial phase (P0) is characterized in Figure 7a by three main peaks at 12.26°, 13.61° and 14.62°. During the first part of the charge (0 < Q/Q<sub>0</sub> < 15% or 0 < Scan < 10) the galvanostatic profile is characterized by a smooth variation of the voltage that corresponds to that observed for the potential in figure 4b (0 < Q < 12 mAh/g). Surprisingly however, no angular shift was detected for the initial set of peaks during the first 10<sup>th</sup> scans (55 minutes, Figure S8). Instead, intensities were found to decrease at the expense of a new phase (P1) characterized by a set of growing reflections at constant angle (15.22°, 15.77°, 19.49° and 24.17°) (Figure 7a and S7). Peak fittings allowed a quantitative description of the phase transformation progress referred to as  $\xi_i$  (Figure 7b, pink curve). The latter was derived according to the ratio  $\xi_i = 1 - [I(P0)_i / I(P0)_0]$  where  $I(P0)_0$  and  $I(P0)_i$  are the sum of the integrated intensity of the three main peaks of (P0) listed above, at the beginning and, during the electrochemical cycling respectively. The evolution of (P1) (Figure 7b, violet curve) was tracked by the ratio between the integrated intensity  $I(P1)_i$  of the (P1) main peak at 15.22° on cycle  $i$ , over that obtained by extrapolation of  $I(P1)_i$  at 100% of the theoretical capacity, referred to as  $I(P1)_{MAX}$  (Figure S9). Finally, the evolution of the specific capacity  $Q_n$  normalized by the theoretical one  $Q_{theoretical}$  is also shown in Figure 7b (comparison of these curves vs the voltage is reported in Figure S10). The trends of the structural and electrochemical

evolutions probed by  $\xi_i$  and  $Q_n / Q_{theoretical}$  generally fits well for the whole cycle and the similar slopes of  $1 - [I(P0)_i / I(P0)_0]$  and  $I(P1)_i / I(P1)_{MAX}$  in shows (P0) and (P1) transforms into one another during the reaction. However, at the beginning of the charge and discharge the extent of the phase transformation is up to 10% below that expected by referring to the relative capacity  $Q/Q_0$ . This type of "structural delay" has been observed for the phase transformation between LiFePO<sub>4</sub>/FePO<sub>4</sub><sup>19-22</sup> due to macroscopic heterogeneity associated with the active material grain connectivity to ionic and electronic percolating networks<sup>20</sup>. We note that on the first cycle the capacity on charge is 28.8% higher than on discharge while it is only 12.7% higher on the second cycle (Figure S11). However, the amounts of P1 at the end of the 1<sup>st</sup> and 2<sup>nd</sup> charges are virtually unchanged (Figure S8) while a marked inflection appears on first charge above 0.8V (Figure S12) suggesting side reaction, most presumably oxygen evolution, contributes to the coulombic inefficiency. Importantly, results clearly show that a significant fraction of (P0) (48%) does not react upon charge (Figure 7b). When the current is reversed, 88% of (P1) transforms back to (P0) which fits well with the recovered specific capacity of 41 mAh/g. We note that the residual trace at the end of the reduction (6%) of (P1) suggests either kinetic limitation and/or oxidation of (P0) by dissolved oxygen which was produced on charge.

In conclusion, based on a one electron reaction as proposed in Figure 1, operando XRD shows that the specific capacity of 4CT-Naphthalene scales with the mass ratio of matter that is involved in the electrochemical reaction. Accordingly, either some grains are not connected to the electrical circuit and/or the electrode highly suffers from kinetic hindrance causing some grains to partially react. This last hypothesis can primarily stem either from the efficiency of electronic percolation network

within the composite electrode or, from the electrical conductivity of the compound itself (electronic and/or ionic).

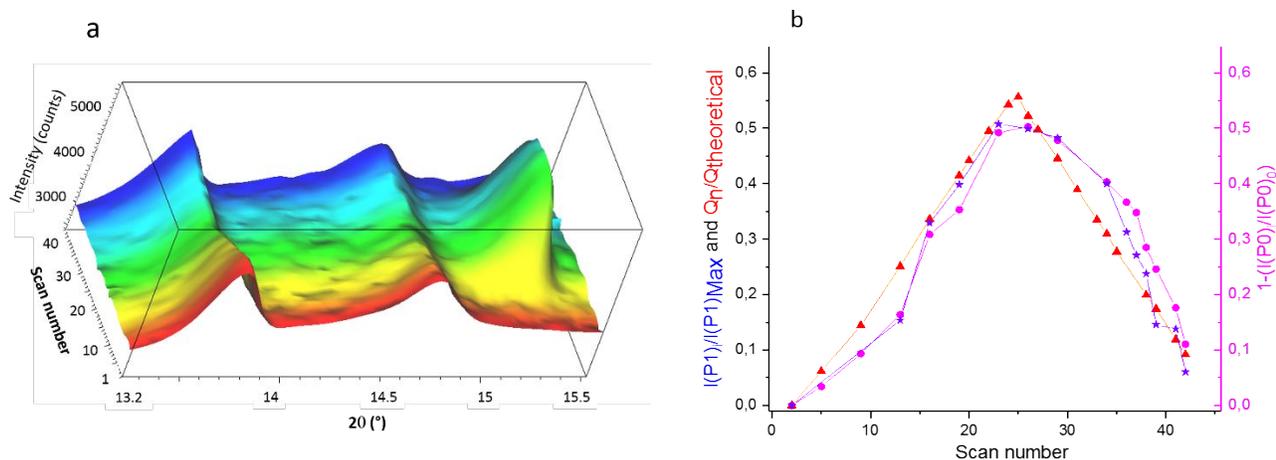


Figure 7. a) XRD diagrams collected for 4CT-Naphtalene during the first cycle within (-0.1; 0.95) V vs SCE showing the phase transformation between the initial phase P0 and the new phase P1 characterized by a peak at 15.22°. b) Evolution of the corresponding structural phase transformation derived by  $\xi_i = 1 - [(P0)_i / (P0)_0]$ , (P0) being the initial phase, compared to the evolution of the fraction of the new phase (P1) and the relative specific capacity ( $Q_n / Q_{\text{theoretical}}$ ).

In this context, the 4CT-Naphthalene has been tested doubling the amount of conductive carbon (40 wt.% of KB, 10 wt.% of CNT) everything else being the same. Surprisingly enough, the discharged capacity in 1M NaClO<sub>4</sub> aqueous electrolyte is virtually unchanged (38 mAh/g) (Figure 8). SEM images of this sample are compared in Figure 9 to that containing 25 wt.% of carbon additive. The two samples distinctively show the three components of the composite electrodes. More particularly, the two mixtures show 4CT-Naphthalene particles are nicely wrapped by nano-grains (and nano-tubes in the case of the 50 wt.

mixture) of the carbon additive which suggests the main cause for the relatively low specific capacity of 4CT-Naphthalene is not related to insulated grains but rather to kinetic hindrance. The size of 4CT-Naphthalene grains and quality of the carbon additive/active grain mixture could be modified by using an alternative protocol to prepare the electrode. The latter consists in co-precipitating a DMSO solution of 4CT-Naphthalene containing 25 wt.% of carbon additive in water (see Experimental part for more details). SEM images of the corresponding electrode mixture show that carbon particles leave this time some un-covered active material grains (Figure 9). As a result, only 29 mAh/g could be attained with this type of electrode which is approx. 25% lower than with the two other electrode mixtures (Figure 8). Complementary images taken at lower magnifications (Figure S13) reiterate these observations and highlight that in all cases most 4CT-Naphthalene grains are a few microns large. Accordingly, in an attempt to unify these results, we believe the slow kinetics of 4CT-Naphthalene electrodes is primarily ascribed to the low electrical conductivity of the compound. Its effect is all the more detrimental to the performance that grains are generally quite large. Therefore, the quality of the mixture with the carbon

additive does improve the performance up to a certain point where the entire surface of the grains is covered (as observed with 25 wt.% of carbon – hand mixed). It would therefore not be surprising that decreasing the grain size while achieving a kind of conducting coating would most likely greatly enhance the electrochemical properties of this compound. In this regard, Ketjenblack is presumably not the most appropriate carbon additive. To nail our point, we provide in the supplementary information the test of the well-known PTAm polymer using the present electrode chemistry. Indeed, results reported in Figure S15 show that the maximum specific capacity that could be measured does not exceed 44 mAh/g (49% of the theoretical capacity). This value significantly departs from that obtained recently by Oyaizu and Nishide's group using an optimized electrode design with only 5 wt.% of single wall carbon nanotubes and an optimized current collector<sup>7</sup>.

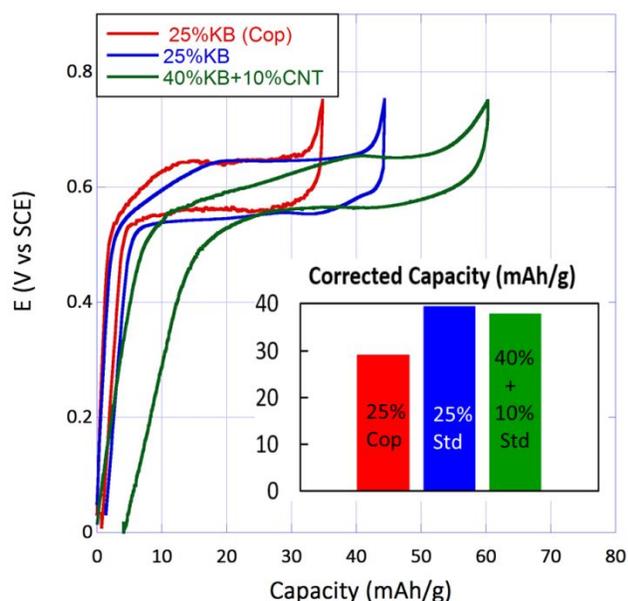


Figure 8. Galvanostatic curves of 4CT-Naphthalene mixed with 25 wt.% KB (blue), 40 wt.% KB + 10% CNT (green) and co-precipitated with 25 wt.% KB (red) obtained in 1 M NaClO<sub>4</sub> aqueous electrolyte (0.075 A/g). Inset shows the specific capacity recovered on discharge after correction of that associated with the carbon (Figure S14). The areal capacity of electrodes are about 0.7 mAh/cm<sup>2</sup>.

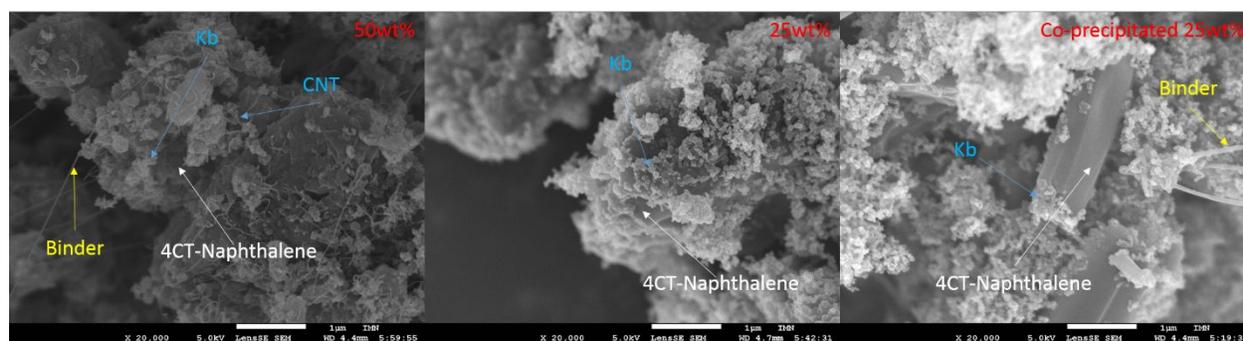


Figure 9. SEM images of 4CT-Naphthalene electrode mixtures with 50 wt.% (left) and 25 wt.% (middle) of conductive carbon (KB). The co-precipitated electrode mixture with 25 wt.% of conductive carbon is also shown (right).

As mentioned above, the oxidized form of the 4HT-Benzene dissolves in 1 M NaClO<sub>4</sub>. Accordingly, a “water-in-salt” approach was evaluated using a more concentrated NaClO<sub>4</sub> electrolyte (8 M) without any other additives<sup>23</sup>. It seems important to remind that the price of NaClO<sub>4</sub>, H<sub>2</sub>O is only 1-2\$/kg<sup>24</sup>. The material has been cycled at 0.075A/g in a (0; 0.9) V potential window vs SCE. Figure 10a,b highlights the effect of the electrolyte concentration. First, an estimation of the equilibrium potentials (defined as  $E_{eq} \approx \langle E \rangle = (E_{pa} + E_{pc})/2$ ) shows the latter is ~50 mV lower in the concentrated electrolyte (8 M) than in the 1 M electrolyte solution which compares well with the potential difference expected from the Nernst equation (i.e.,  $\Delta E = 53$  mV). Second, the cyclability and Coulombic efficiency of the material are significantly improved in the concentrated electrolyte. Indeed, the initial capacity is nearly the same than in the 1M electrolyte but remarkably, it increases a bit from 52 mAh/g to nearly 60 mAh/g (62% of the theoretical capacity) during the first 30 cycles and is then constant for the next 70 cycles (123 hours in total). Notwithstanding, as shown by the Ragone plot in Figure 10c only 64% of the theoretical capacity could be achieved even at a C/10 rate. In conclusion, despite a higher concentration of the electrolyte solves the cyclability issue, the specific capacity of 4HT-Benzene is still hindered.

To gain a better understanding of the 4HT-Benzene redox behavior in concentrated NaClO<sub>4</sub> electrolyte (8M), electrochemical impedance spectroscopy (EIS) measurements have been performed at different states of charge using a sample containing 10 wt.% of carbon additive (Figure 11a,b). This relatively low carbon content is used to exacerbate the electrical response associated with the material itself. Except for the spectrum corresponding to the initial state, all EIS spectra measured during the electrochemical process show a semicircle followed by a diffusion and capacitive part (Figures 11 and S16). The significant change between the partially charged and initial states indicates the resistance associated with the semicircle can be ascribed to a charge transfer process. Interestingly, results show the charge transfer resistance (36 Ohms) is almost stable all along the electrochemical reaction suggesting that the oxidized phase shows similar electrical properties to the reduced one. The decrease of the carbon content to 10% does significantly reduce the capacity to only 11 mAh/g (82% lower than with 25 wt.%, Figure 11 c,d), demonstrating a strong impact of the percolating network. EIS results therefore leave us to believe that the specific capacity of 4HT-Benzene is mitigated by its poor electron conductivity which would hamper the electron transport across the electrode when particles are poorly connected to the carbon network. This

hypothesis is supported by the trend of the charge transfer resistance that scales almost linearly with the inverse of the specific capacity (Figure 11c) and therefore with the number of particles connected to the carbon network. Accordingly, the full capacity (97 mAh/g) could be achieved with 40 wt.% of KB and

10 wt.% of CNT hand mixed with the material (Figure 11d). As a result, both TEMPO derivatives (4HT-Benzene and 4CT-Naphthalene) suffer from kinetic hindrance that inhibits the performance.

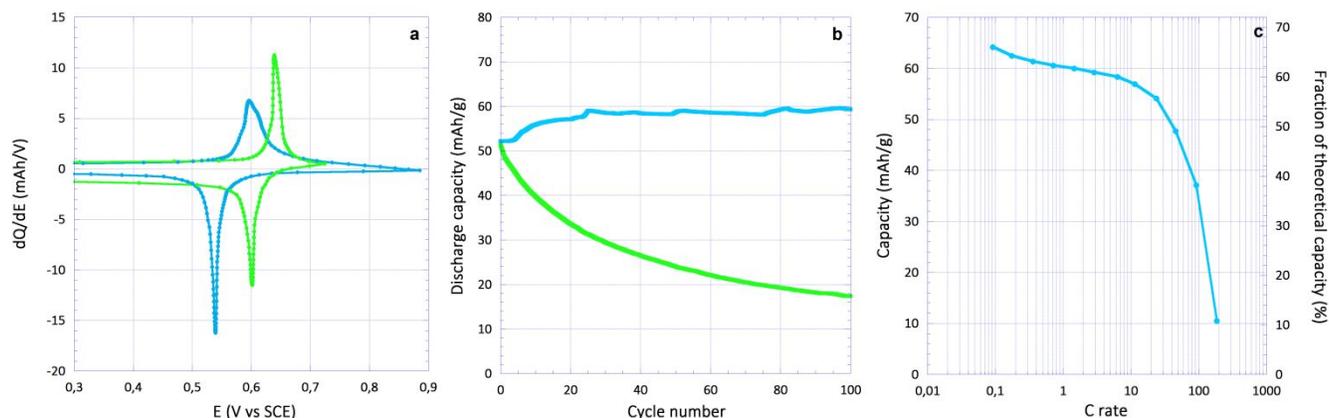


Figure 10. a) Incremental capacity curve on cycle 4 derived from galvanostatic cycling of 4HT-Benzene at 0.075 A/g in highly concentrated 8 M (blue) and 1 M (green) NaClO<sub>4</sub>; b) cyclability of 4HT-Benzene at 0.075 A/g in 8 M (blue) and 1 M (green) NaClO<sub>4</sub>; c) Ragone plot of 4HT-Benzene in 8 M NaClO<sub>4</sub>. Electrodes contain 25 wt.% of carbon additive and the corresponding areal capacity are about 0.7 mAh/cm<sup>2</sup>.

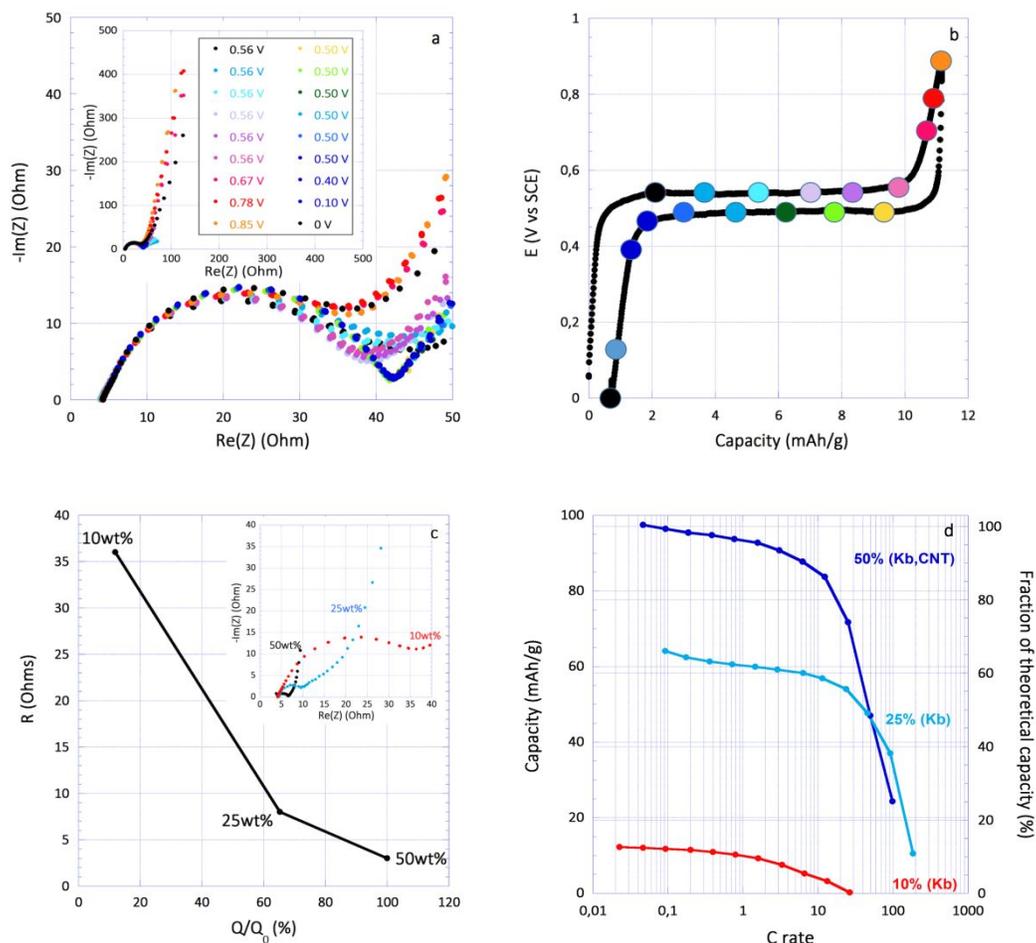


Figure 11. a) EIS measurements performed at different states of charge using a sample of 4HT-Benzene containing 10 wt.% of carbon additive in 8M NaClO<sub>4</sub>; b) Galvanostatic curve of the 4HT-Benzene (0.7 mAh/cm<sup>2</sup>) electrode containing 10 wt.% of carbon additive in 8M NaClO<sub>4</sub>; c) Dependence of charge transfer resistance towards the specific capacity (the capacity has been normalized to the theoretical capacity (Q<sub>0</sub>); inset of c) EIS measurements performed at the end of charge of samples of 4HT-Benzene containing 50 wt.%, 25 wt.% and 10 wt.% of conductive carbon; d) Ragone plot of 4HT-Benzene electrodes containing 50 wt.%, 25 wt.% and 10 wt.% of conductive carbon.

To curtail the impact (mass and cost) of the concentrated NaClO<sub>4</sub> electrolyte at the cell level, a 4HT-Benzene thick electrode of 7.8 mAh/cm<sup>2</sup> (80 mg/cm<sup>2</sup>; 1.10 mm) areal capacity was designed and evaluated (Figure 12). During the first 68 cycles, the capacity sharply rises from 22 to 40 mAh/g which most likely mirrors an activation process associated with the wetting of the electrode porosity. The latter can be mitigated however, by applying a floating period at the end of the charge (0.9 V) until the current drops to 0.015 A/g (C/5) (Figures 12a and S17). With that condition, the discharged capacity slightly increases up to 66 mAh/g<sub>4HT-Benzene</sub> or 5.3 mAh/cm<sup>2</sup> (68% of the theoretical capacity) after 535 cycles (38 days). The coulombic efficiency also increases during the first 30 cycles to stabilize at 99.8%, suggesting an important contribution of the oxygen evolution in the initial stages. A blank electrode containing only the conductive additive indicates that half of the irreversibility actually stems from side reactions going on at the surface of the

carbon (Figure S18). The latter sensibly attenuate with cycling presumably due to the increasing acidity of the electrolyte from neutral pH to pH=4 after 25 cycles (Figure 12b). Remarkably, the power properties of the thick electrode are not strongly impacted up to C-rate as it merely behaves like the thin ones (Figure 12c). This therefore also serves to reiterate that performance are limited by the efficiency of the electrode chemistry in coping for the low conductivity of the material. Results are therefore hardly comparable to those of Oyaizu and Nishide' group<sup>7</sup> who demonstrated the only other thick (>3 mAh/cm<sup>2</sup>) organic based electrode. Indeed, in this case a high utilization of the electrode can only be reached thanks to the use of specific SWNTC conducting additive and 4- $\mu$ m-thick conductive SWNT layer deposited at the surface of the current collector.

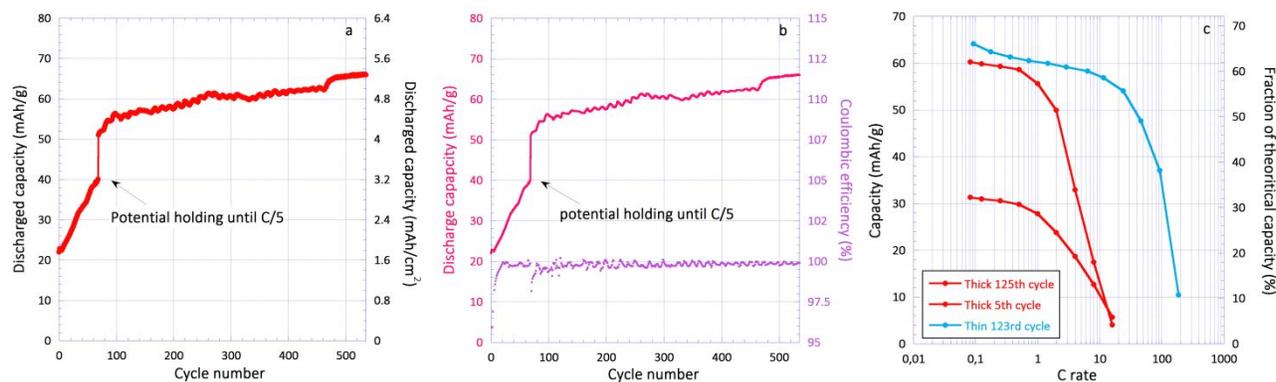


Figure 12. a) Cyclability of 4HT-Benzene in 8 M NaClO<sub>4</sub> aqueous electrolyte (80 mg/cm<sup>2</sup>); b) corresponding cyclability and coulombic efficiency together with c) Ragone plots of both thin (10 mg/cm<sup>2</sup>) and thick electrodes (80 mg/cm<sup>2</sup>).

Although no further attempts have been done to improve the overall kinetics of the electrodes, a full cell has been evaluated with thick electrodes (mixed with 25 wt.% of KB) by coupling the 4HT-Benzene positive electrode to a novel bipyridinium-naphthalene diimide oligomer (referred to as DNVBr) as the negative one. The formula and synthesis of DNVBr are provided in the supplementary information. As demonstrated in our previous work for a molecule of the same family<sup>25</sup> and illustrated in scheme S1, DNVBr is a p and n type active material that releases anions and uptakes cation simultaneously during its electrochemical reduction (charge of the full cell). The present 4HT-Benzene/DNVBr full cell is therefore a mixed, rocking-chair anionic<sup>12,26</sup> cell (that corresponds on charge to the transfer of anions from DNVBr to 4HT-Benzene) and a dual-ion<sup>5</sup> one simultaneously (with the transfer on charge of cations from the electrolyte to DNVBr and corresponding anions to the 4HT-Benzene). Although a detailed study of the electrochemical performance of the DNVBr compound is out of the scope of the present paper, its charge-discharge profile and specific capacity retention are provided in Figure S19 for an 80 mg/cm<sup>2</sup> electrode. The capacity ratio 4HT-Benzene:DNVBr leading to the highest energy density was found to be 0.99 and the areal capacity of both electrodes was approx. 8 mAh/cm<sup>2</sup> (see Experimental part for more details regarding the electrode preparation). For the sake of comparison, an identical full cell but using thin electrodes of 0.7 mAh/cm<sup>2</sup> (10 mg/cm<sup>2</sup> electrodes) has also been evaluated. The cell has been cycled at 0.075 A/g (1C) within a controlled voltage of (0; 1.8) V vs. SCE (Figure 13a, b) by holding the end-of-charge potential until 0.015 A/g (C/5). Under these conditions, an average voltage of

0.96 V was obtained with a remarkably flat specific capacity of 30 mAh/g<sub>active materials</sub> associated with an excellent coulombic efficiency (99.6%) over 285 hours (~12 days, 190 cycles) (Figure 13b,c). These results compare well with that achieved with the thin electrode full cell while allowing much higher areal capacity of 4.6 mAh/cm<sup>2</sup> (vs 0.7 mAh/cm<sup>2</sup> for the thin one). Remarkably such a high areal capacity competes well up to 8C (Figure 13d). We note the slightly reduced discharge voltage arises from a higher polarization associated with the thick electrodes (Figure 13a). In addition it is worth noting that although a high maximum charge voltage of 1.8 V is applied, self-pH-buffering of the electrolyte occurs to around pH=7, thus mitigating current collector corrosion. The specific energy density of this non-optimized thick electrode cell is 28 Wh/kg per mass of both active materials at C-rate and 35 Wh/kg at C/8 rate, the latter rate being still acceptable for stationary storage. This 4HT-Benzene//DNVBr cell is the first full organic cell with close to millimeter thick electrodes to the best of our knowledge. In comparison, the specific energy density obtained for the thin electrode cell saturates to approx. the same value of 36 Wh/kg at C and C/2 rates. However, extrapolating these results to a whole stack cell with electrodes of more than 700 cm<sup>2</sup>, shows that the lower fraction of inactive components inherent to the use of millimeter thick electrodes enables 13.6 Wh/kg<sub>stack</sub> at 1C and 17.4 Wh/kg<sub>stack</sub> at C/8, which represents gains of nearly 13% and 37% respectively, vs. the same cell but having 0.7 mAh/cm<sup>2</sup> electrodes (12.7 Wh/kg<sub>stack</sub>) (Figure 13c, d) (see supplementary information for details regarding calculations).

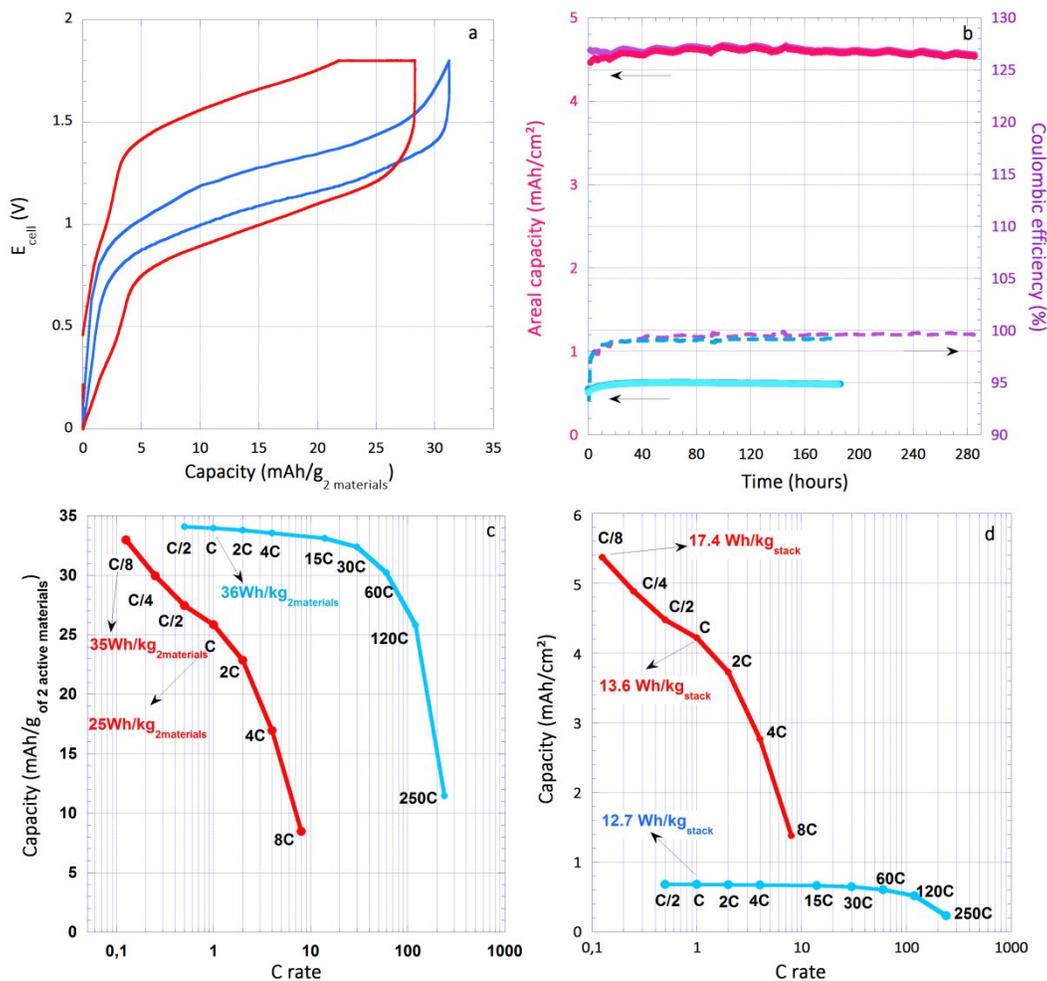


Figure 13. a) Cell voltage ( $E_{\text{cell}}$ )-capacity profile of full cells of 0.7 (blue) and 8  $\text{mAh/cm}^2$  (red) electrodes at 0.075 A/g, b) Cyclability of the full cells (violet thick electrodes full cell; blue thin electrodes full cell) at 0.075 A/g along with Coulombic efficiency; c, d) corresponding Ragone plots in 8M  $\text{NaClO}_4$ .

In the light of these results, pouch cell configurations with similar thick electrodes have also been evaluated to further demonstrate the practicability of the present full organic aqueous battery (see Experimental part for further details). The capacity ratio 4HT-Benzene: DNVBr was set to 1.01. In these conditions, an average output voltage of 0.78 V was obtained

with similar areal capacity (4.5  $\text{mAh/cm}^2$  corresponding to 30  $\text{mAh/g}_{\text{active materials}}$ ), high cyclability (98% of capacity retention after 500 hours), coulombic efficiency (99.6%) at 0.075 A/g (Figure 14a) and rate capability (Figure 14b). More interestingly, an even higher areal capacity (5  $\text{mAh/cm}^2$ ) has been further attained at 0.009 A/g (C/8).

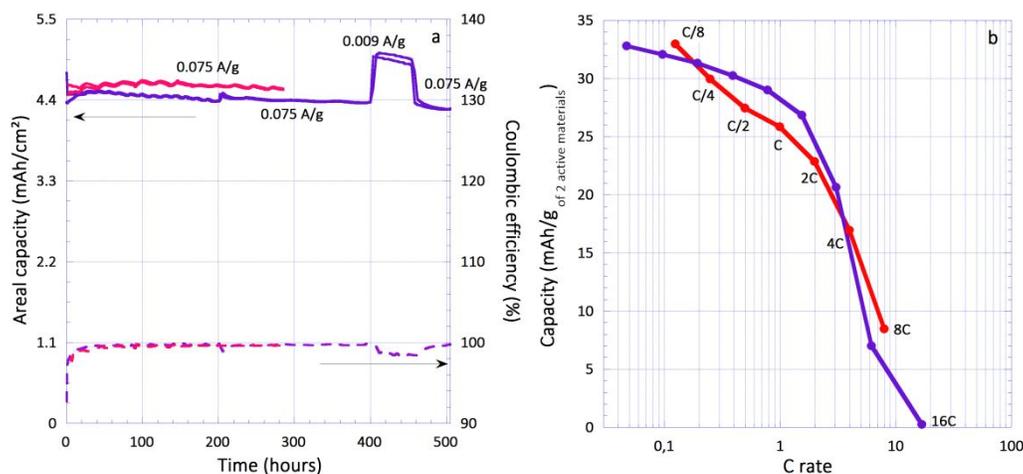


Figure 14. a) Cyclability of the full cells (violet pouch cell; red glass cell) along with coulombic efficiency in 8M NaClO<sub>4</sub>; b) corresponding Ragone plots.

### Conclusion

All the above results serve to demonstrate and rationalize the potentialities and limitations of two small TEMPO molecules in neutral molar aqueous electrolyte. For the first the commercially available 4-hydroxy TEMPO benzoate molecule has been proved to be highly efficient active material in millimeter thick electrodes delivering up to 5.3 mAh/cm<sup>2</sup> at C rate for over 500

cycles in neutral and cheap 8 M NaClO<sub>4</sub> electrolyte. Paired with a new bipyridinium-naphthalene diimide oligomer a novel type of aqueous battery that operates simultaneously in anion-rocking chair and dual-cation/anion mode is demonstrated. The full cell shows excellent cyclability and coulombic efficiency offering 35 Wh/kg<sub>materials</sub> on discharge at C/8. We believe these results represent an important advance towards cheaper and more sustainable energy storage systems.

## ASSOCIATED CONTENT

**Supporting Information.** Further electrochemical details and all the materials' characterizations could be found in the Supporting Information. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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### Author Contributions

All authors have given approval to the final version of the manuscript.

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