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Biodegradable Zn-ion battery with a lignin composite electrode and bio-ionic liquid based electrolyte: possible *in situ* energy generation by lignin electrocatalysis†

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Electrochemical energy storage with lignin-based composite electrodes offers a cheap and renewable source for batteries. Lignin composites have mainly been used in supercapacitors and little has been investigated related to applying these composites in batteries. Here we show a biodegradable hybrid Zn-ion battery with a polymer/lignin composite electrode as a cathode, a Zn anode and bio-ionic liquid electrolytes. Interestingly, the polymer/lignin composite cathode led to a higher discharge capacity compared to the charge capacity over the entire 100 charge/discharge cycling process. It was observed that the lignin concentration in the composite polymer electrode was crucial in obtaining such a phenomenon. Using SEM, electrochemical analysis and NMR, we showed that lignin electrocatalytically oxidizes during battery charging, which produced the extra energy. This results in obtaining a higher discharge capacity compared to the charge capacity. It is anticipated that such an electrocatalytic technique can be extended to other battery systems to harness clean electrochemical energy from biomass.

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Introduction

Development of batteries with sustainable materials is important to reduce the battery cost as well as reduce the complexity of the end-of-life battery recycling process. Biomass composite electrodes are potential alternatives to metal oxide electrodes due to its abundance and biodegradable properties. Besides electrodes, current batteries use toxic and flammable electrolytes, which makes battery recycling a challenging aspect. Therefore, new environmentally friendly electrolytes are equally necessary to develop safe batteries.

Among different biomass, lignin is a naturally occurring biopolymer present in plant cells. It has a complex polymeric structure comprising mainly hydroxybenzene, guaiacyl (4-alkyl-2-methoxyphenol) and syringyl (4-alkyl-2,5-dimethoxyphenol).^{1–3} Due to its vast availability, effective valorization of lignin is of significant interest. Until now, main attention has been given towards lignin fractionation and/or depolymerisation using green chemistry routes to form value-added aromatics as

products.^{2,4–6} However, the feasibility beyond the lab-scale has been questioned mainly due to the complex structure of lignin and its heterogeneity.⁶ In comparison to depolymerisation, some recent studies have shown the prospect of directly catalyzing lignin to electricity using a biomass fuel cell which works at elevated temperatures between 80 and 550 °C.^{7–9} However, the low conversion efficiency, costly electrodes (usually noble metals), elevated temperatures and low power density hinder further exploitation of the process.

Lignin derivatives such as lignosulfonates and its composites with conducting polymers have shown some promise in energy storage.^{10–13} The quinone groups formed by electrooxidation of guaiacyl and syringyl in lignin are excellent moieties for reversible redox reactions, which has been exploited for sensing molecules and for charge storage.^{12–14} Lignin based composite electrodes have been tested as capacitors in acidic or alkaline solutions.^{10–13} For batteries, lignin has been used as binders and electrodes^{15,16} wherein the electrodes were made by carbonizing lignin.^{16–19} Casado *et al.* showed the possibility of Na ion storage in a lignin/poly(3,4-ethylenedioxythiophene) composite electrode in ionic liquids, which showed a capacity of 70 mA h g^{−1} at a low current density of 25 mA g^{−1}.²⁰

Here, we demonstrate a new phenomenon in which lignin present in a polypyrrole composite electrode not only acts as a charge storage medium but also acts as an energy generating source during running of a Zn-ion battery in bio-ionic liquid

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electrolytes. This *in situ* energy generation in effect leads to >100% coulombic efficiency.

Results and discussion

The composite electrode of polypyrrole (PPy) and lignin was obtained by electropolymerisation. The microstructures of the PPy and PPy/lignin composite electrodes are shown in Fig. 1. PPy shows the formation of coagulated large particles (Fig. 1a). The PPy/lignin composite shows a slightly different morphology as the concentration of lignin in the electrolyte is changed. As higher concentrations of lignin is incorporated in PPy, a much smoother morphology (Fig. 1d and e) is obtained with the presence of small particles of around 1 μm in size. An increase in particle size is again observed with further increase in lignin concentration (Fig. 1f). Comparing the microstructures of various composites in Fig. 1, it is evident that lignin significantly changes the PPy morphology.

The Raman and X-ray photoelectron spectroscopy (XPS) of PPy and the PPy/lignin composites are shown in Fig. 2 and 3, respectively. The Raman spectra in Fig. 2 show characteristic peaks of polypyrrole.²¹ From Raman, it can be observed that, on addition of lignin to polypyrrole, almost insignificant shifts are observed. Most of the lignin peaks get merged within the polypyrrole structure. However, changes in the Raman intensity are evident. With the addition of 0.5 mg mL^{-1} lignin in the electrolyte, a slight increase in the Raman intensity is observed, which increases on addition of lignin to 2.5 mg mL^{-1} to the electrolyte and decreases on further addition of lignin. This indicates that the number of active functional groups/cross-linkages in the composite is the highest with the electrolyte containing 2.5 mg mL^{-1} of lignin. Due to the overlap of lignin modes and polypyrrole modes, it is not possible to distinguish the concentration of lignin in the polymer.

The XPS spectra of the PPy and PPy/lignin composite electrodes are compared in Fig. 3. From the XPS survey spectra in Fig. 3a,

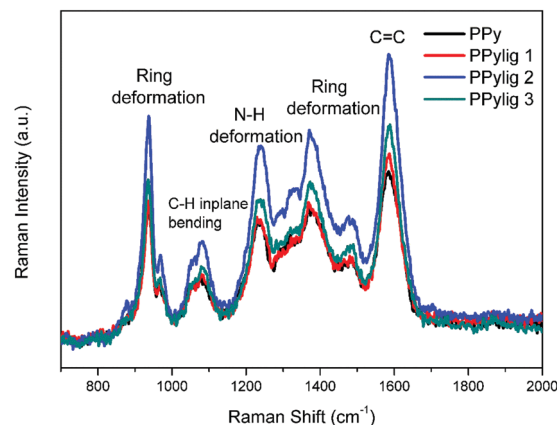


Fig. 2 Raman spectra of polypyrrole and polypyrrole with different lignin concentrations. PPyLig1 was synthesized from an aqueous solution containing 0.05 M LiClO_4 , 0.1 M pyrrole and 0.5 mg mL^{-1} of liginosulfonate. PPyLig2 was synthesized from an aqueous solution containing 0.05 M LiClO_4 , 0.1 M pyrrole and 2.5 mg mL^{-1} of liginosulfonate. PPyLig3 was synthesized from an aqueous solution containing 0.05 M LiClO_4 , 0.1 M pyrrole and 5 mg mL^{-1} of liginosulfonate.

peaks related to C 1s, N 1s, O 1s, Cl 1s and S 2p are observed. The S 2p and Cl 1s come from liginosulfonate and LiClO_4 present in the electrolyte, respectively. The presence of S 2p confirms the formation of the PPy/lignin composite electrode. The C 1s detailed spectra are shown in Fig. 3b. PPy shows three peaks at 284.8 eV (C–C bond), 286.3 eV (C–OH, C–N) and 288.1 eV (C=O).^{22,23} The presence of lignin in PPy broadens the C 1s spectra and an increase in the intensity at 286.3 eV is evident. This peak increase is due to the presence of *p*-hydroxyphenyl, guaiacyl, and syringyl moieties of lignin. The N 1s detailed spectra also show some changes. On addition of lignin in PPy, the N 1s spectra in Fig. 3c change from a symmetric peak to an asymmetric peak. Furthermore, a shoulder at 401 eV is observed, which clearly indicates additional lignin moieties present in the PPy matrix.

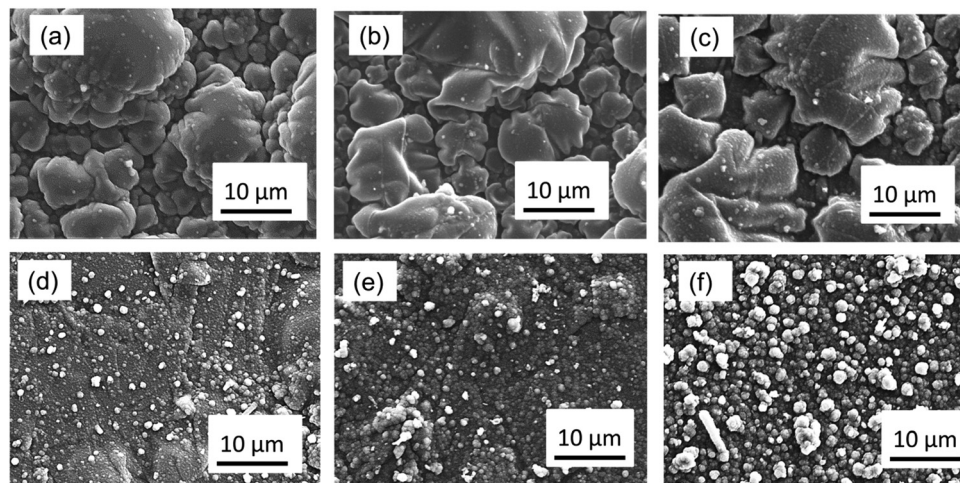


Fig. 1 Microstructure of polypyrrole and the PPy/lignin composite electrode deposited on carbon paper from (a) 0.05 M LiClO_4 , 0.1 M pyrrole, (b) 0.05 M LiClO_4 , 0.1 M pyrrole, 0.5 mg mL^{-1} liginosulfonate, (c) 0.05 M LiClO_4 , 0.1 M pyrrole, 1.25 mg mL^{-1} liginosulfonate, (d) 0.05 M LiClO_4 , 0.1 M pyrrole, 2.5 mg mL^{-1} liginosulfonate, (e) 0.05 M LiClO_4 , 0.1 M pyrrole, 3.75 mg mL^{-1} liginosulfonate and (f) 0.05 M LiClO_4 , 0.1 M pyrrole, 5 mg mL^{-1} liginosulfonate.



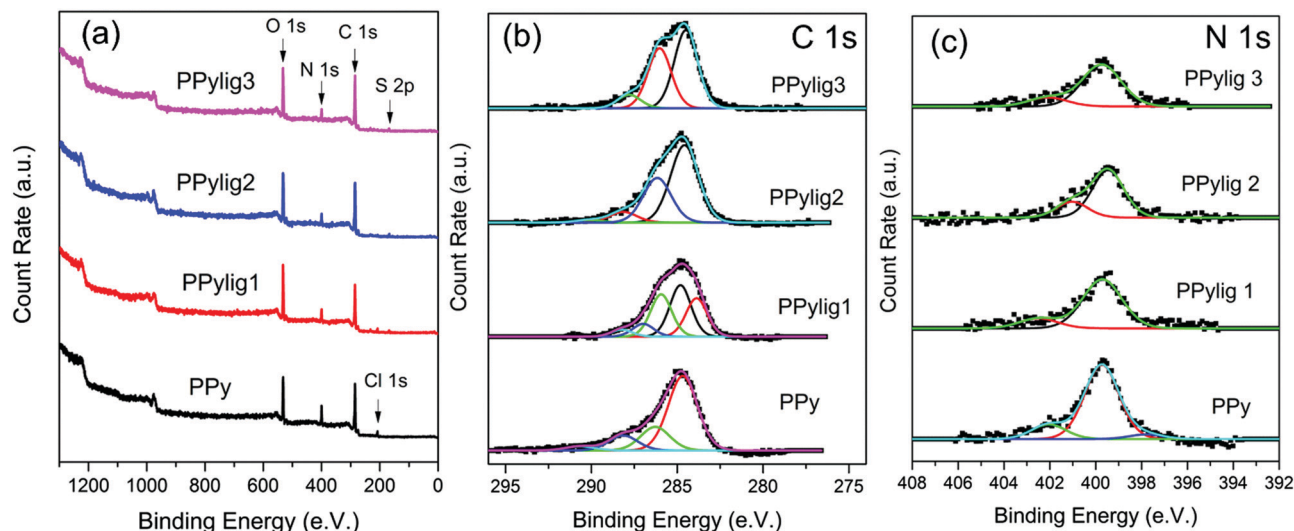


Fig. 3 (a) XPS survey spectra of PPY and PPY with different lignin concentrations, (b) C 1s detailed spectra of PPY and the PPY/lignin composites and (c) N 1s spectra of PPY and the PPY/lignin composites.

Fig. 4a shows the cyclic voltammetry curves of polypyrrole (PPY) and the PPY/lignin composite in 50 wt% choline acetate (ChAc) + 50 wt% water and 1 mol kg⁻¹ of Zn acetate (ZnAc₂) in (50 wt% ChAc + 50 wt% water). PPY shows almost a capacitive

behavior in 50 wt% ChAc + 50 wt% water in the presence and absence of ZnAc₂, whereas in the PPY/lignin composite electrode an oxidation peak at 1.04 V and a large reduction wave centered at 0.95 V are observed in 50 wt% ChAc + 50 wt%

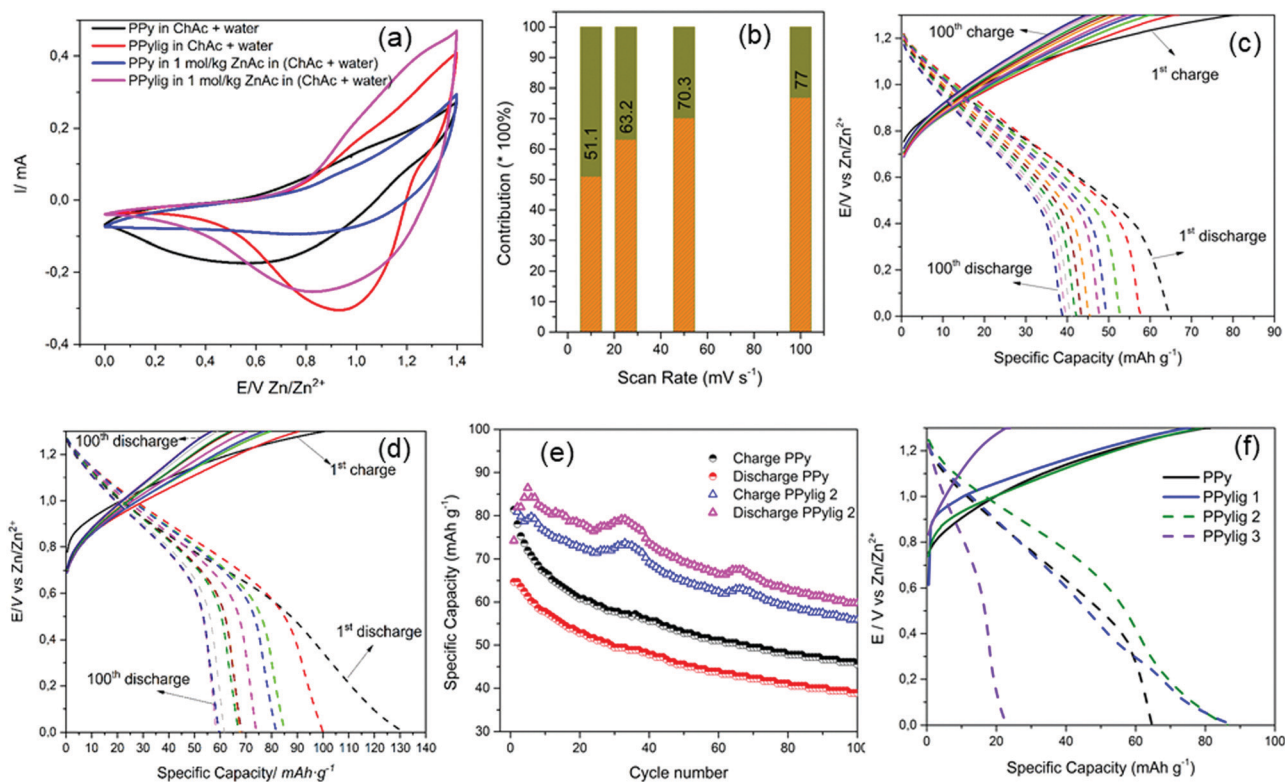


Fig. 4 (a) CV of the PPY and PPY/lignin composite electrodes in different electrolytes at a scan rate of 10 mV s⁻¹. (b) Capacitive contribution to the total capacity at different scan rates (orange represents the capacitive process and the olive color represents the faradaic process). (c) The charge–discharge profile of PPY in 1 mol kg⁻¹ of ZnAc₂ in (50 wt% ChAc + 50 wt% water) at a current density of 200 mA g⁻¹. (d) The charge–discharge profile of the PPY/lignin composite (2.5 mg mL⁻¹ lignin concentration in the electrolyte for deposition) in 1 mol kg⁻¹ of ZnAc₂ in (50 wt% ChAc + 50 wt% water) at a current density of 200 mA g⁻¹. (e) Galvanostatic charge–discharge with the cycle number at a current density of 200 mA g⁻¹. (f) The charge–discharge profile of different PPY/lignin composites in 1 mol kg⁻¹ of ZnAc₂ in (50 wt% ChAc + 50 wt% water).

water. The oxidation/reduction peak is attributed to quinone moieties formed by guaiacyl and syringyl oxidation in lignin.¹⁴ On addition of ZnAc₂ in the electrolyte, a shift in the oxidation peak to 1 V and reduction peak to 0.81 V is observed. The shift in the oxidation/reduction peaks confirms that Zn ions are being stored in the biopolymer matrix. The contribution of the capacitive process in the PPy/lignin composite electrode was determined by plotting log of the current (at a particular voltage) versus log of the scan rate according to the equation $i = av^b$ where a is a constant and b is a value ranging from 0.5 (diffusion controlled contribution) to 1 (capacitive contribution).^{24–26} The current was chosen at the peak potentials at 0.95 V and 0.76 V in the anodic and cathodic processes (from Fig. S1a, ESI†), respectively. From Fig. S1b (ESI†), the value of b was found to be 0.73 and 0.65 for the cathodic and anodic processes, respectively, which suggests the capacitive process to be the main contribution to charge storage. The contribution of capacitive and faradaic processes is shown in Fig. 4b, from which it is evident that with an increase in the scan rate the capacitive process becomes more prominent. The hybridization of both faradaic and capacitive processes is an added benefit in high energy density battery systems²⁷ for fast charging of the battery.

Fig. 4c and d compare the charge discharge of the PPy and PPy/lignin composite electrodes in Zn-ion batteries. In PPy the charge discharge curve shows that an initial discharge capacity of 65 mA h g^{−1} is achieved for Zn storage and the charging process takes a considerably longer time. In comparison, on addition of lignin to PPy, a remarkable change in the charge-discharge behavior is observed in Fig. 4d. A synergic effect is seen with a higher initial discharge capacity of almost 130 mA h g^{−1}, which can be related to the presence of lignin which acts as additional Zn storage sites and leads to an initial higher discharge capacity. However, strangely the discharge capacity is higher than the charge capacity, which can occur due to irreversibility or a secondary reaction that generates *in situ* energy during the charge process. Irreversibility would be detrimental to the battery performance as Zn would be consumed with cycling, leading to lower battery performance; however, *in situ* energy generation might lead to new battery chemistry.

The galvanostatic charge-discharge curves with the number of cycles of PPy and the PPy/lignin composite are compared in Fig. 4e. Compared to Fig. 4d, it is evident that in the first cycle the charge capacity is higher than the discharge capacity for the PPy/lignin composite. However, from the second cycle a higher discharge capacity compared to the charge capacity is observed. Repeated experiments with different PPy/lignin composites showed that such a behavior is observed sometimes in the first cycle but reverses from the second cycle. The instability in the charge/discharge behavior in Fig. 4e can be ascribed to the heterogeneous and complex structure of lignin. With a change in the lignin concentration in PPy, a similar behavior of an increased discharge capacity compared to the charge capacity is seen (Fig. 4f and Fig. S2, ESI†) until a certain concentration. With a higher concentration of lignin in PPy, not only is a significant decrease in capacity storage observed but also the

charge capacity becomes higher than the discharge capacity (PPy/lign3, Fig. 4f). This might be due to a phase change in the PPy/lignin composite (see Fig. 1 wherein a significant change in the morphology is observed with varying lignin in the biopolymer matrix), which inhibits the secondary reaction from taking place. Also, a higher concentration of lignin leads to lowering of the active sites in the polymer as seen from Fig. 2. The galvanostatic charge-discharge curves of different PPy/lignin composites are shown in Fig. S2 (ESI†). Except for the case of PPy/lign3, it is evident that in all the cases the discharge capacity is higher than the charge capacity (Fig. S2, ESI†).

To understand the influence of lignin in generating energy during charging, the electrolyte and electrode were analyzed. The CV of PPy and the PPy/lignin composite in the Zn ion containing electrolyte with and without lignin is shown in Fig. 5a. PPy in the electrolyte shows a capacitive behavior. However, on addition of lignosulfonate in the electrolyte, a clear increase in current from 0.95 V is observed (red line, Fig. 5a), which can be associated with electrocatalytic oxidation/dissolution of lignin.²⁵ In the PPy/lignin composite electrode the CV shows an oxidation/reduction behavior, which is related to the quinone redox process. However, with lignosulfonate in the electrolyte, an increase in current as well as a shift in the oxidation potential by less than 100 mV (pink line, Fig. 5a) is evident.

It has been shown that Zn can accelerate the depolymerisation of lignin in formic acid² and therefore the increase in current in Fig. 5a along with the reduced potential can be associated with the increase in electrocatalytic oxidation/dissolution of lignin. Therefore, on running the battery, the increase in discharge capacity compared to charge capacity can be associated with the electrons generated by the electrocatalytic oxidation/dissolution of lignin during battery charging. To prove the electrocatalytic oxidation/dissolution behavior from the PPy/lignin composite electrode, CV cycles were run in the 50 wt% ChAc + 50 wt% water electrolyte (Fig. S3, ESI†). It is evident from the CV curves (Fig. S3b, ESI†) that with cycling the current related to the redox reaction increases and reaches almost saturation after about 40 cycles. This is in contrast with that observed in PPy (Fig. S3a, ESI†) in the same electrolyte, wherein a decrease in the redox reaction took place. This indicates that lignin is electrocatalytically oxidized/dissolved and generates energy during this process. Furthermore, the microstructure of the PPy/lignin composite observed after CV cycles in Fig. 5b shows formation of a number of particles of about 1 μm in size compared to fewer particles observed in Fig. 1d. Furthermore, on plotting the difference between the discharge capacity and charge capacity with the cycle number for different PPy/lignin composite electrodes (Fig. 5c), it is evident that the capacity difference fades with cycling, which confirms the irreversible lignin oxidation/dissolution during the charge process. However, with different lignin concentration in PPy, an additional average energy of about 5 mA h g^{−1} is still present even after 50 cycles. This suggests that the electro-oxidation/dissolution of lignin in this electrolyte is slow. It has been shown that lignin dissolution is difficult in ionic liquids with low yields of 6% at 110 °C.²⁸ Therefore, it is anticipated that the



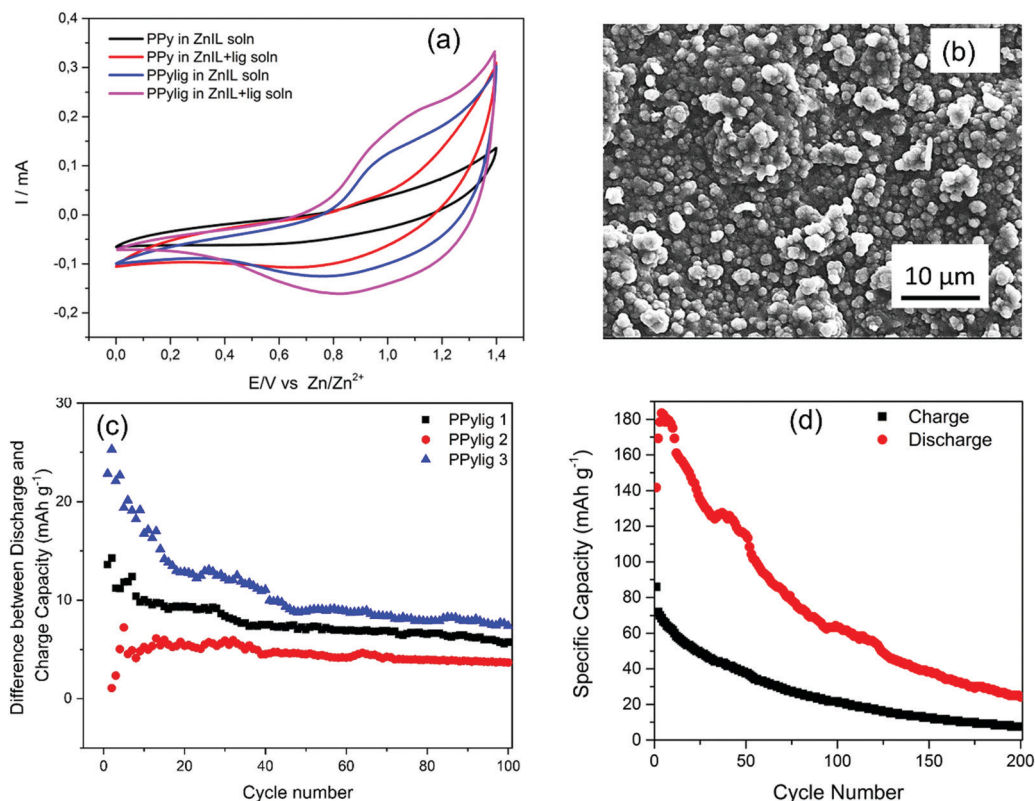


Fig. 5 (a) CV of PPY and the PPY/lignin composite in 1 mol kg⁻¹ ZnAc₂ in (50 wt% ChAc + 50 wt% water) and the same electrolyte containing liginosulfonate (2.5 mg mL⁻¹) performed at 10 mV s⁻¹. (b) SEM of the PPY/lignin composite after 50 CV cycles in 1 mol kg⁻¹ ZnAc₂ in (50 wt% ChAc + 50 wt% water). (c) Specific capacity difference between discharge and charge. (d) Charge/discharge cycles of the PPY/lignin composite electrodeposited on indium tin oxide at 300 mA g⁻¹.

lignin powered battery would result in producing additional energy for a long time and less energy will be needed to charge the battery. In order to relegate the effect of the carbon current collector in producing extra energy, the PPY/lignin composite was deposited on ITO and the battery was tested in the same electrolyte. Fig. 5d again shows a higher discharge compared to charge,

which decreases with the cycle number. Although the specific capacity decreases significantly with the cycle number, the discharge is more than the charge even after 200 cycles, which clearly indicates that the current collector has no role in producing extra energy during charging. However, repeated experiments showed that the battery capacity is significantly affected

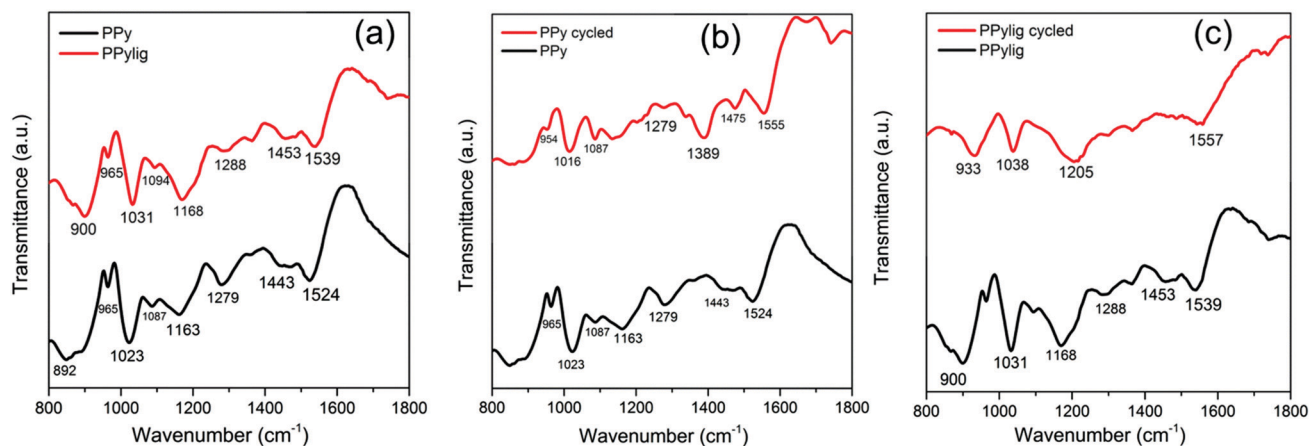


Fig. 6 (a) IR spectra of the electrodeposited polypyrrole (black line) and polypyrrole–lignin composite (red line), (b) comparison of polypyrrole (black line) and polypyrrole after cycling in a Zn-ion battery (red line) and (c) comparison of the polypyrrole–lignin composite (black line) and polypyrrole–lignin composite after cycling in a Zn-ion battery (red line).



The electrode was further investigated with XPS to evaluate the structural changes in the electrodes. XPS of the as synthesized PPy and PPy/lignin and after cycling in the Zn-ion battery are compared in Fig. S9 (ESI[†]). A stark difference in the cycled PPy

The ionic liquid, choline acetate (ChAc), was purchased from IO-LI-TEC, Germany (99%). Zinc acetate and lithium perchlorate (99.99%) were obtained from Sigma-Aldrich (99%) and pyrrole (>98%) was purchased from Alfa Aesar. The pyrrole was distilled prior to use and was stored in a refrigerator.

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