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PAPER

In situ nano-assembly of bacterial cellulose–polyaniline composites

Zhijun Shi,^a Shanshan Zang,^a Fan Jiang,^a Long Huang,^b Dan Lu,^b Yuguang Ma^b and Guang Yang^{*a}

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Bacterial cellulose (BC), produced by *Gluconacetobacter xylinum*, consists of ribbon-shaped nano-fibers in web-like structures. Polyaniline (PAni) is a conductive polymer, the conductivity of which is related to the synthesis and doping process. In this paper, we report on the *in situ* nano-assembly of BC nanofibers and PAni to enhance the electronic conductivity. PAni could be synthesized on the surface of BC nano-fibers and assembled into a novel 3D network. The reaction time of polymerization, the types and concentration of doping protonic acids play a major role on the electroconductivity properties of the composites. The electroconductivity of composite hydrogels was enhanced from 10^{-8} to 10^{-2} S cm⁻¹, and can be further improved by doping with various protonic acids. The BC–PAni nanofiber composite is an electro-conductive hydrogel that combines the properties of hydrogels and conductive systems, and it may potentially be used for flexible displays, biosensors, and platform substrates to study the effect of electrical signals on cell activity, and to direct desirable cell function for tissue engineering applications.

Introduction

As a conductive electroactive polymer, polyaniline is well-known for its ease of synthesis and environmental stability (Fig. 1). Aniline can be polymerized to polyaniline through chemical¹ or electrochemical polymerization.^{2,3} The conductivity of polyaniline can be changed through doping/dedoping and oxidation/reduction, from 10^{-8} S m⁻¹ (an insulator) to 10^2 S m⁻¹ (a conductor).⁴ Conductive electroactive polymers, including polyaniline, find applications as electrical conductors, batteries, chemical and biological sensors, actuators, electromagnetic shielding, antistatic coatings, corrosion protection, and electro-optic and electrochromic devices.^{3,5–12} Their unique electrical, electrochemical and optical properties also could be used as health care materials, from tissue engineering to drug delivery, from neural regeneration to smart implants. All these several special sections provide new opportunities and challenges to these polymers. Many researchers want to use conductive electroactive polymers in biomedical applications, such as implantable electrochemical biosensors, electro-stimulated drug release devices and neural prosthetics. But there are several obvious challenges, in which questionable biocompatibility is the most serious problem.¹³

Bacterial cellulose (BC) is a kind of cellulose, and it can be produced by many microorganisms,^{14,15} one of which is *Gluconacetobacter xylinum*.¹⁶ Sharing the same structural unit (Fig. 1), BC does not only have the properties of common

cellulose products produced by plants, but it also has its own special properties. The special properties of BC are its high chemical purity and crystallinity,¹⁷ high degree of polymerization, excellent biocompatibility and biodegradability.^{18,19} These properties have already led to successful commercialization in a variety of niche markets, including health foods, high-end audio components, speciality paper, and wound care,^{19–21} or even for gene detection.²² Most importantly, BC is a natural polymer hydrogel, which has three-dimensional polymeric networks and imbibe large amounts of water. BC hydrogels are flexible and have good mechanical properties, and they can accordingly easily change their size and shape in response to the environmental stimuli. Moreover, BC can also imbibe other monomeric, reactive and potentially polymerizable monomers into its networks, essentially occupying its void volume and interacting with BC fiber chains. Many researchers have used these

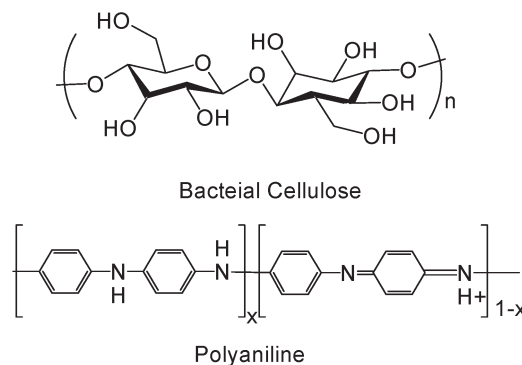


Fig. 1 Chemical structure of BC and PAni ($x = 1$, leucoemeraldine base; $x = 0.5$, emeraldine salt; $x = 0$, pernigraniline).

^aNational Engineering Research Center for Nano-Medicine, College of Life Science and Technology, Huazhong University of Science and Technology, Wuhan, 430074, China. E-mail: yang_sunny@yahoo.com; Fax: 86 027 8779 2265; Tel: 86 027 8779 3523

^bState Key Laboratory of Supramolecular Structure and Materials, Jilin University, Changchun, 130012, China



Fig. 2 Photographs of (a) BC hydrogel and (b) BC-PAni hydrogel and (c) schematic diagram showing the experiment process, the chemical structure and synthesis of PAni in the BC hydrogel.

properties to change or improve the characteristics of BC to achieve optical transparency,^{23–26} electrical conductivity,²⁷ hydrophobicity,²⁸ etc.

The use of plant cellulose, regenerated cellulose and cellulose derivatives in combination with conductive polymers for the preparation of conductive materials has been reported previously.^{29,30} Additionally, there are multiple reports involving BC that document their biocompatibility, biodegradability, and bioadhesion.^{18–20,31} In this paper, we incorporated polyaniline, with BC hydrogels, to synthesize an electroconductive hydrogel that combines the properties of hydrogels and conductive systems. The reaction time of polymerization, the type and concentration of doping protonic acids play a major role in the electroconductivity properties of the composites. The combination of BC hydrogels and inherently conductive electroactive polymers forms an excellent biocompatible and electroconductive hydrogel material, which may engender new applications in



Fig. 3 ESEM images of the surface of a bacterial cellulose membrane cultured for 10 days. (The BC membrane was soaked in water for 2 days and then freeze-dried.)

flexible electrodes, flexible displays, or even be used as healthcare materials, such as tissue engineering scaffolds, deep brain stimulation electrodes, implantable biosensors, and implantable neural prosthetic devices.

Experimental section

Preparation of BC membranes

Gluconacetobacter xylinum (ATCC53582) was used for the biosynthesis of BC. The bacterium was cultured in a Hestrin and Schramm (HS) medium, which was composed of 2% (wt) glucose, 0.5% (wt) yeast extract, 0.5% (wt) peptone, 0.27% (wt) disodium phosphate, and 0.15% (wt) citric acid. After incubating statically for 6–14 days at 26 °C, the BC membranes were dipped into distilled water for 2 days, and then steamed by boiling in a 1 wt% NaOH solution for 30 min to eliminate bacteria and proteins.¹⁸ Afterwards, the BC membranes were purified by being washed in distilled water several times, and were then stored in distilled water at 4 °C.

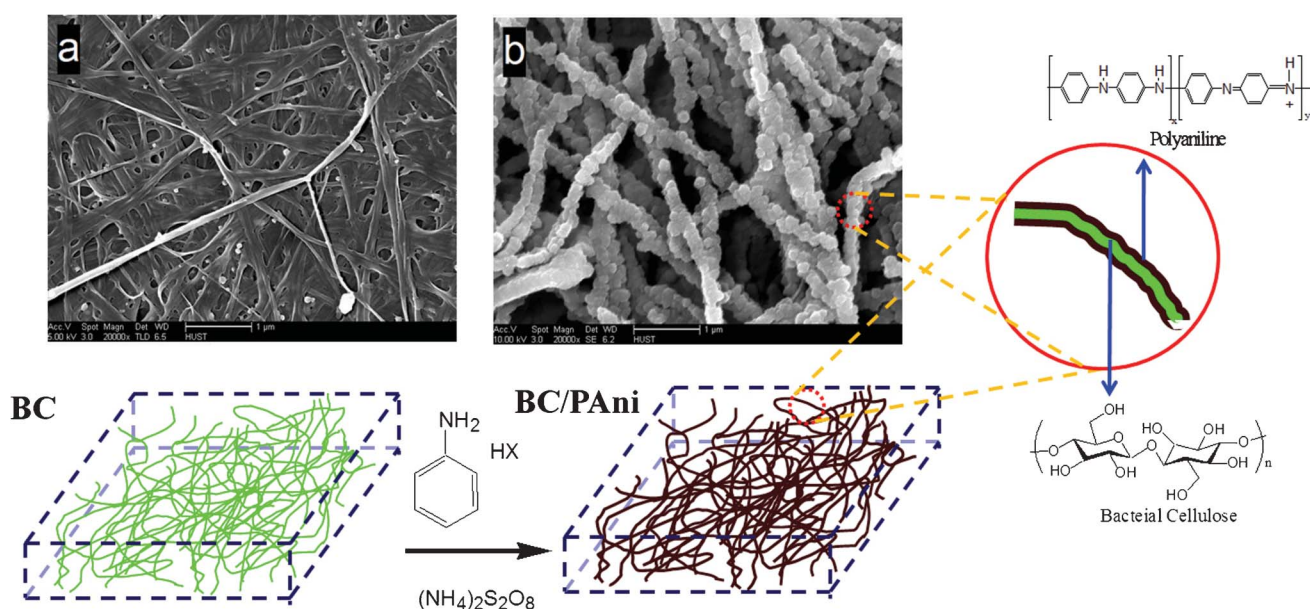


Fig. 4 Schematic diagram of the process of aniline polymerization in the BC hydrogel. (Fig. a is the FESEM image of the hot-pressed BC membrane; and Fig. b is the FESEM image of hot-pressed BC-PAni membrane.)



Fig. 5 XPS of BC, PANi and BC-PAni-2.

Preparation of BC-PAni composite membranes

The BC membrane was dipped into a 100 ml aniline hydrochloride solution (0.5 mol L^{-1}) for about 48 h with stirring, at room temperature, enabling the aniline hydrochloride solution to fully infiltrate through the network of BC fibers. The BC was immersed into 200 ml of an ammonium peroxydisulfate solution (0.25 mol L^{-1}) with stirring. Samples were taken out of the ammonium peroxydisulfate solution after 5, 10, 20, 30, 60 and 120 min. The schematic diagram of the process of aniline polymerization is shown in Fig. 2. The composite samples obtained at 10, 30 and 60 min were named BC-PAni-1, BC-PAni-2 and BC-PAni-3. Finally, dry BC or BC-PAni hybrid sheets were obtained by pressing the BC or BC-PAni hydrogels at 120°C and 1–2 MPa.

Characterization

The surface morphology of all the samples was observed using environment scanning electron microscopy (ESEM, Quanta 200, Holand) and field emission scanning electron microscopy (FESEM, Sirion 200, FEI, Holand). All samples were sputtered with gold, before observation and photography.

X-Ray photoelectron spectroscopy (XPS ESCALAB 250, UK) was used to characterize the surface composition of the BC and the BC-PAni. Monochromatic Al K α X-rays were employed. The X-ray source was 0.5 mm nominal X-ray spot size utilized for both survey and high-resolution spectra. Survey spectra, from 0 to 1200 eV binding energy, were recorded at 50 eV pass energy with an energy step of 1.0 eV. High-resolution spectra were recorded at 20 eV pass energy with an energy step of 0.05 eV, with a typical average of three scans.

The samples were analyzed in KBr discs by Fourier transform infrared spectroscopy (FTIR, VERTEX 70, Germany) in the region $400\text{--}4000 \text{ cm}^{-1}$.

^{13}C cross-polarization magic-angle spinning (CPMAS) and ^{13}C single-pulse excitation magic-angle spinning (SPEMAS) NMR spectra were recorded on a Varian InfinityPlus-300 NMR spectrometer equipped with a standard Chemagnetic magic-angle spinning probe head, operating at 299.98 MHz for ^1H and 75.12 MHz for ^{13}C , respectively. Typically, about 50 mg sample was packed into a 4 mm zirconium oxide rotor and spun at about 6 kHz ($\pm 1 \text{ Hz}$).

Raman microscopy (Renishaw inVia, U.K.) was performed using 325 nm laser excitation.

Table 1 XPS results of the BC, PANi and BC-PAni-2 composites

	XPS ^a			
	C 1s	O 1s	N 1s	S
BC	39.8	60.2	0	0
BC-PAni-2	51.1	40.2	8.7	0
PAni	25.0	46.3	12.2	16.5

^a Atom% was based on the survey scan of C 1s (285 eV), O 1s (532 eV), N 1s (400 eV), S (169 eV, 232 eV).

The conducting properties of samples were measured by using Potentiostats-Electrochemistry Workstation (Par 2273, USA). The samples were placed between two $\text{SnO}_2\text{:F}$ electrodes, by measuring the current in a voltage range $0\sim 1 \text{ V}$ or $-4\sim 4 \text{ V}$.

Results and discussion

In the process of cellulose formation, sub-elementary fibrils are first assembled to form microfibrils ($d = 3\sim 4 \text{ nm}$) and further polymerized to form ribbons ($d = 30\sim 50 \text{ nm}$), which finally developed into a hyperfine network structure through interactive interweaving. For subsequent analysis, the BC membranes were soaked in water for 2 days and then freeze-dried. The ESEM images of the dried BC membranes showed that they formed dense and finely branched nanolevel networks in the planar

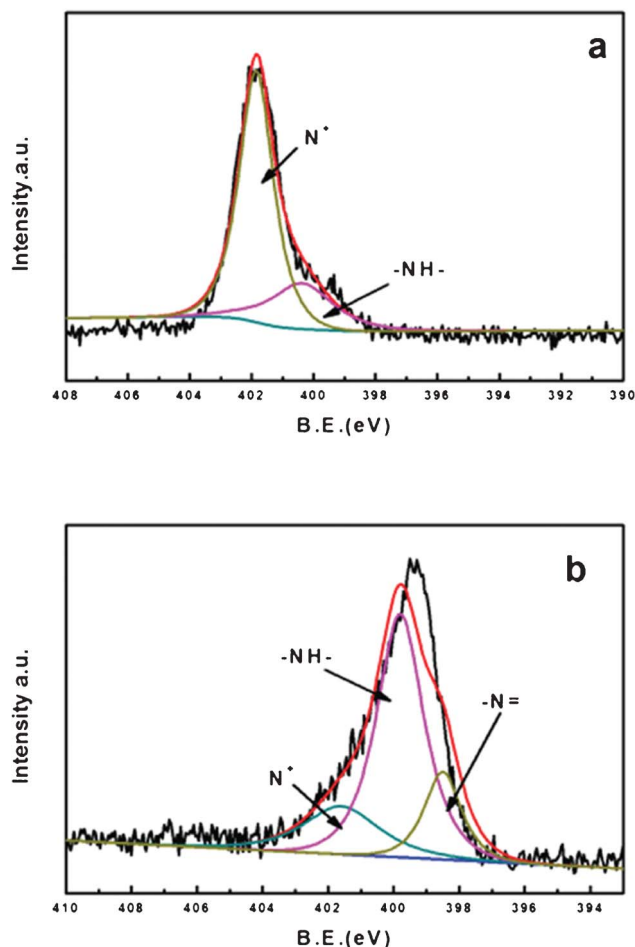


Fig. 6 XPS spectra (N 1s) for (a) PANi and (b) BC-PAni-2.

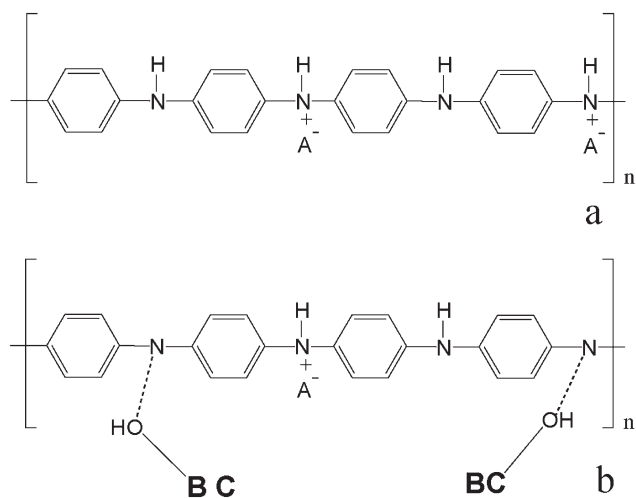


Fig. 7 Scheme of PANi (emeraldine salt form) (a) and BC-PANI (b).

direction (Fig. 3), and the microfibrils are 40 nm in diameter. To obtain purified BC membranes, bacterial cell debris and protein should be totally removed by boiling in 1% wt NaOH solution for about 30 mins.

Dry BC membranes were obtained by pressing the wet BC mat at 120 °C and 1–2 MPa. After the hot press process, the wet BC membrane turned into a highly toughened sheet, with the fracture strength reaching 180 MPa. The FESEM images show the network structure of the purified BC membrane (Fig. 4a).

PANI is currently prepared by the oxidation of aniline with ammonium persulfate as the oxidant in an acidic aqueous medium.^{32,33} The BC membrane was added to a conical flask filled with an aniline salt solution, and then the flask was placed on a shaking table for 48 h to let the aniline salt solution replace the water in the membrane. When the BC membrane was soaked in a solution of ammonium persulfate, polymerization started in the BC membrane substrate to form PANi. The aniline salt lost electrons as it was converted to PANi, and ammonium persulfate accepted these electrons and was reduced to sulfate.³⁴ Based on the section images of samples with different reaction times, it was concluded that the PANi was formed from the surface to the

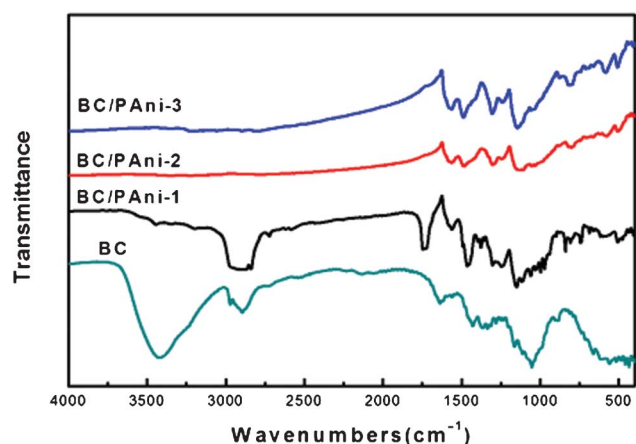


Fig. 8 FT-IR spectra of BC and BC-PANI composites.

Table 2 XPS spectra (N 1s) for (a) PANi and (b) BC-PANI-2

	XPS		
	N ⁺ /N	-NH-/N	-N=N
PANI	75.4	24.6	0
BC-PANI	19.8	62.6	17.6

middle layer of the BC hydrogel; contrast is provided by the color of the BC-PANI composites which are dark green.

The results of FESEM show that BC-PANI composites (Fig. 4b) contain a random assembly of microfibrils. This morphology of uniform microfibrils is very different from that of microfibrils of purified BC. From several images the width of BC microfibrils was measured to be 40–60 nm, and the width of BC-PANI composites microfibrils was 80–120 nm. We propose that the PANi polymerized on the surface of the BC nanofiber.

X-Ray photoelectron spectroscopy (XPS) was used to characterize the surface composition of BC, PANi and BC-PANI. The appearance of S (169 eV, 232 eV) in the PANi sample was attributed to the ammonium persulfate which accepted electrons and was reduced to sulfate. Survey scans of BC showed the C 1s peak at 285 eV and the O 1s peak at 532 eV, respectively. The N 1s peak appears at 400 eV; no nitrogen peak is found in BC, but it is seen in PANi and BC-PANI (Fig. 5). From Table 1 we know that the atom% of O in BC-PANI was lower than that in BC, while the atom% for C in BC-PANI was higher than that of BC, and the difference was attributed to the polymerization of PANi on the BC nanofiber.

Fig. 6a represents the XPS spectra for the nitrogen of PANi. The curve deconvolution resulted in the presence of three peaks attributed to -N= (398.5 eV), -N- (399.8 eV) and N⁺ (401.6 eV).^{35,36} PANi is prepared by the oxidation of aniline with ammonium persulfate as the oxidant in an acidic aqueous medium. The percentage of -N=, -N- and N⁺ was of 0%, 24.6% and 75.4% respectively (Table 2). The PANi sample we obtained is a protonated emeraldine base form with a high conductivity (~3 S cm⁻¹), which is called the emeraldine salt form (ES).³⁷ Fig. 7a shows the molecular structure of ES; from this we know that the nitrogens of ES occurred as protonated nitrogens (N⁺) and as amine nitrogens (-N-). The XPS spectrum for BC-PANI-2 is shown on Fig. 6b. In this case, the

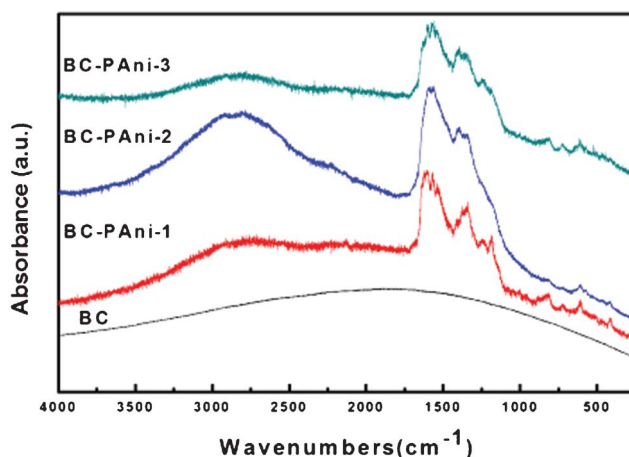


Fig. 9 Laser Raman spectra of BC and BC-PANI.

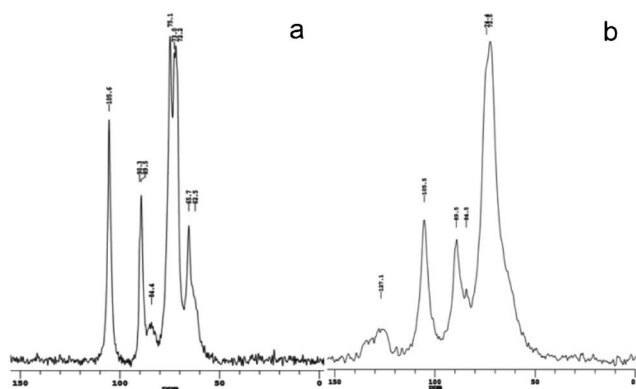


Fig. 10 ^{13}C CPMAS NMR spectra of (a) BC and (b) BC-PAni-1

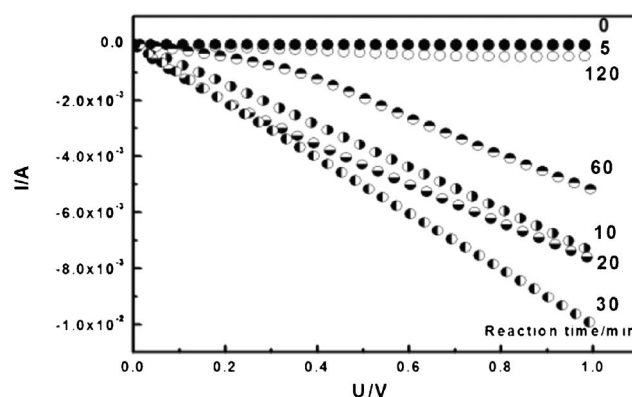


Fig. 11 C-V characteristic curve of BC-PAni with different reaction times. (Linear sweep voltammogram of the samples between two SnO_2/F electrodes from 0 ~ 1 V.)

Table 3 Conductivity variance of BC-PAni with different reaction times

Reaction time (min)	0	5	10	20	30	60	120
Electrical conductivity (S cm^{-1})	6.3×10^{-8}	7.2×10^{-4}	7.8×10^{-3}	5.8×10^{-3}	1.3×10^{-2}	4.5×10^{-3}	6.7×10^{-5}

percentages of $-\text{N}=\text{}$, $-\text{N}-$ and N^+ were of 17.6%, 62.6% and 19.8%. The decrease of the percentage of protonated nitrogen (N^+) could indicate that BC interacts with the PAni through the formation of hydrogen bonds, as suggested in Fig. 7.

BC and BC-PAni composites were also characterized by Fourier transform infrared spectroscopy (Fig. 8). In the spectrum of BC, the peak at 3400 cm^{-1} was assigned to the hydroxyl group, and the peak at $2800\sim 3000 \text{ cm}^{-1}$ was attributed to C-H stretching. The spectrum of all three samples of composites showed two peaks at 1560 cm^{-1} and 1460 cm^{-1} , which were assigned to $\text{N}=\text{Q}=\text{N}$ and $\text{N}-\text{B}-\text{N}$, respectively (Q represents quinonoid and B represents benzenoid moieties). The conductive form of PAni, which consists of about equal amounts of benzenoid and quinonoid units, has approximately equal intensities for the 1560 cm^{-1} and 1460 cm^{-1} absorption bands.³⁸ The sample of BC-PAni-1, in which the intensity for the 1460 cm^{-1} absorption band was much stronger than for the 1560 cm^{-1} band, demonstrated that the amounts of benzenoid units were greater than those of the quinonoid units. Moreover, the sample of BC-PAni-2, in which intensity of the 1460 cm^{-1} absorption band was about equal to the 1560 cm^{-1} absorption band, suggested that with increased progressing reaction time, more benzenoid units would transform to quinonoid units. The BC-PAni-2 sample consists of about equal amounts of benzenoid and quinonoid units, that is to say, the structure of this sample was similar to that of the conductive form of PAni. Therefore, the sample of BC-PAni-2 should have good electrical conductivity.

It is important to inquire how PAni was incorporated into the BC. We observed that the intensity for the 3400 cm^{-1} absorption band, which is attributed to the hydroxyl ($-\text{OH}$), was reduced in BC-PAni-1, and even disappeared in BC-PAni-2 and BC-PAni-3. In addition, the absorption spectra of BC-PAni composites did not show any N-H stretching absorption. Thus, we can conclude that BC may have reacted *via* dehydration-condensation with PAni, leading to cause the disappearance of O-H and N-H bonds.

The laser Raman spectra of BC and BC-PAni samples are shown in Fig. 9; they were obtained with a 325 nm excitation line. As it can be observed, the spectrum of BC indicates dispersion of fluorescence, but the spectra of BC-PAni

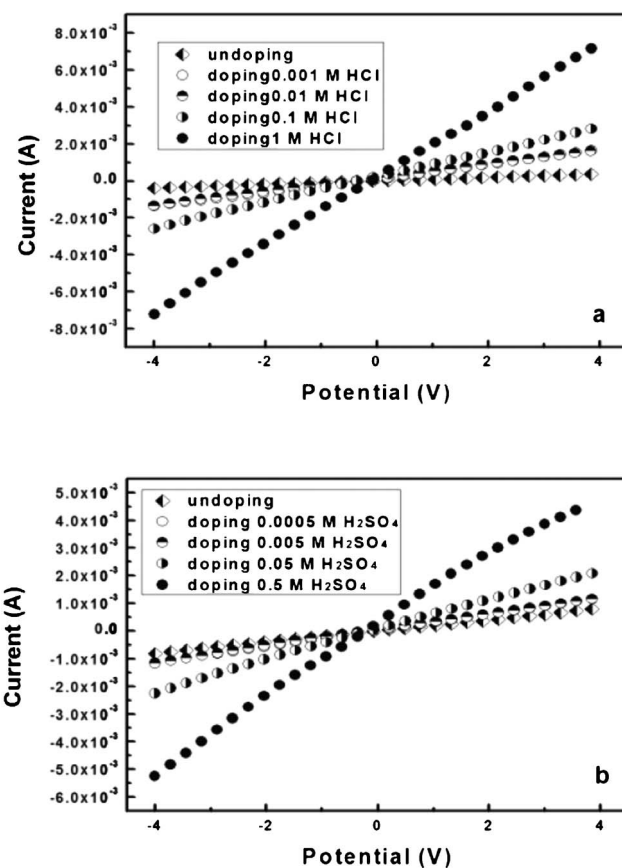


Fig. 12 Linear sweep voltammograms recorded for hybrid BC-PAni films doped by (a) HCl solution and (b) H_2SO_4 solution between -4 and $+4$ V.

Table 4 Conductivity variance of BC–PAni doping by HCl solution and H₂SO₄ solution

C _{HCl} (Mol/L)	1	0.1	0.01	0.001	Undoped
pH	0.01	0.99	1.99	2.73	—
Electrical conductivity (S cm ⁻¹)	1.8×10^{-1}	6.8×10^{-2}	3.7×10^{-2}	3.8×10^{-2}	1.0×10^{-2}
C _{H₂SO₄} (Mol/L)	0.5	0.05	0.005	0.0005	Undoped
pH	0.46	1.54	2.30	3.17	—
Electrical conductivity (S cm ⁻¹)	1.2×10^{-1}	5.5×10^{-2}	2.9×10^{-2}	2.9×10^{-2}	2.0×10^{-2}

composites could be analysed. The relative intensity of the band at 1486 cm⁻¹, assigned to the C=N stretching of the quinoid diimine units, was enhanced when the BC–PAni reaction was terminated at 30 min and 60 min. The bands at 1600 cm⁻¹ (ν (C–C) benzenoid units) and 1148 cm⁻¹ (ν (C–N) benzenediamine units)³⁹ appear in the spectrum of BC–PAni-1, but the bands at 1148 cm⁻¹ (ν (C–N) benzenediamine units) diminished in the samples of BC–PAni-2 and BC–PAni-3. All these changes were consistent with the transformation of benzenoid units into quinoid ones, corresponding to the analysis by FT-IR.

The structures of BC and BC–PAni-1 were confirmed by NMR spectroscopy, as shown in Fig. 10. The spectrum of BC, the region between 60 and 70 ppm is assigned to C6 of the primary alcohol group. The next cluster of resonances, between 70 and 80 ppm, is attributed to C2, C3, C5, and the ring carbons other than those anchoring the glycosidic linkage. The region between 80 and 95 ppm is associated with C4 and that between 100 and 110 ppm with C1, the anomeric carbon.^{40,41} The spectrum of sample BC–PAni-1, compared with that of BC, shows that the peak of C6 is merged with the peaks of C2, C3 and C5, the hydroxyl (–OH) of C6 may react *via* dehydration–condensation with PAni, leading to the C6 peak merging with the other peaks. Finally, the region between 110 and 140 ppm is assigned to the aromatic ring of PAni.^{42,43}

The conductivities of all samples were measured on a Potentiostats-Electrochemistry Workstation. To measure the conductance, the samples were placed between two SnO₂:F electrodes; by measuring the current in a voltage range 0~1 V or –4~4 V, the results are shown in Fig. 11. The reaction times for the polymerizations of aniline were 5, 10, 20, 30, 60, and 120 min. The pure BC membrane is an insulator with a conductivity of 6.3×10^{-8} S cm⁻¹. The conductivity of the BC–PAni composites rose at first and then declined with reaction time; it reached a maximum of 1.3×10^{-2} S cm⁻¹ when the reaction time was 30 min (Table 3). The result is corresponds to the analysis of FT-IR and laser Raman spectroscopy. The PAni was formed gradually in the network of the BC membrane; in the beginning of the reaction, aniline hydrochloride polymerized to leucomeraldine, the fully reduced state of PAni. When the reaction time was about 30 min, the BC–PAni mixtures have a structure similar to that of the emeraldine salt form, the electrical conductive form of PAni, so its conductivity reached the maximum. Afterwards, emeraldine is oxidized to pernigraniline, which made the conductivity decline.

It is well-known that PAni doped by protonic acids displays better conductivity than the undoped ones. The influence of the concentration of the protonic acids can also change and control the BC–PAni hydrogel's conductivity. The samples were given reaction times of 30 min, then doped by 1, 0.1, 0.01, 0.001 Mol L⁻¹ HCl solution, and doped by 0.5, 0.05, 0.005, 0.0005 Mol L⁻¹ H₂SO₄ solution, respectively. Recorded linear

sweep voltammograms are shown in Fig. 12, and electrical conductivity results are shown in Table 4. The sample which was undoped showed an electrical conductivity of 1.0×10^{-2} S cm⁻¹, when doped by 0.001 Mol L⁻¹ HCl solution, the electrical conductivity was improved at 3.8×10^{-2} S cm⁻¹, and when doped by 0.0005 Mol L⁻¹ H₂SO₄ solution, the electrical conductivity was improved from 2.0×10^{-2} S cm⁻¹ to 2.9×10^{-2} S cm⁻¹. Moreover, the electrical conductivity increased when the concentration of HCl/H₂SO₄ solution is promoted. These results suggested that the concentration of hydrogen cations also improved the conductivity of the BC–PAni hydrogel. From Table 4 it can be seen that, in the same concentration of protonic acids, the electrical conductivity of the prepared composites is: HCl > H₂SO₄.

Conclusions

The BC–PAni composite was successfully fabricated by *in situ* nano-assembly with polyaniline. PAni was polymerized and deposited as an adhering layer on the surface of the cellulose fiber. The mixture can form a relatively uniform thin-film conductive material, and its electrical conductivity can reach 10⁻¹ S cm⁻¹. By varying the reaction time, we can get materials with a range of conductivities, and the conductivity of polyaniline can be further improved by controlling the proton acid doping/dedoping. These favorable characteristics can be expected to make the material useful in many fields, such as flexible electrodes, flexible display devices, bio-sensors, as a platform substrate to study the effect of electrical signals on cell activities, to direct desirable cell function for tissue engineering applications, *etc.*

Acknowledgements

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References

- Z. Zhang, Z. Wei and M. Wan, Nanostructures of polyaniline doped with inorganic acids, *Macromolecules*, 2002, **35**(15), 5937–5942.

- 2 J. Liu, Y. Lin, L. Liang, J. A. Voigt, D. L. Huber, Z. R. Tian, E. Coker, B. Mckenzie and M. J. Mcdermott, Templateless assembly of molecularly aligned conductive polymer nanowires: a new approach for oriented nanostructures, *Chem.–Eur. J.*, 2003, **9**(3), 604–611.
- 3 A. G. MacDiarmid, J. C. Chiang, A. F. Richter and A. J. Epstein, Polyaniline: a new concept in conducting polymers, *Synth. Met.*, 1987, **18**(1–3), 285–290.
- 4 D. Li, J. Huang and R. B. Kaner, Polyaniline Nanofibers: A Unique Polymer Nanostructure for Versatile Applications, *Acc. Chem. Res.*, 2009, **42**(1), 135–145.
- 5 P. Chandrasekhar, *Conducting polymers, fundamentals and applications: a practical approach*, Springer, Netherlands, 1999.
- 6 L. Hu, G. Gruner, D. Li, R. B. Kaner and J. Cech, Patternable transparent carbon nanotube films for electrochromic devices, *J. Appl. Phys.*, 2007, **101**, 016102.
- 7 E. M. Genies, A. Boyle, M. Lapkowski and C. Tsintavis, Polyaniline: a historical survey, *Synth. Met.*, 1990, **36**(2), 139–182.
- 8 M. M. Alam, J. Wang, Y. Guo, S. P. Lee and H. R. Tseng, Electrolyte-gated transistors based on conducting polymer nanowire junction arrays, *J. Phys. Chem. B*, 2005, **109**(26), 12777–12784.
- 9 J. Huang, S. Virji, B. H. Weiller and R. B. Kaner, Polyaniline nanofibers: facile synthesis and chemical sensors, *J. Am. Chem. Soc.*, 2003, **125**(2), 314–315.
- 10 Y. Qiao, S. J. Bao, C. M. Li, X. Q. Cui, Z. S. Lu and J. Guo, Nanostructured polyaniline/titanium dioxide composite anode for microbial fuel cells, *ACS Nano*, 2007, **2**(1), 113–119.
- 11 J. Janata and M. Josowicz, Conducting polymers in electronic chemical sensors, *Nat. Mater.*, 2003, **2**(1), 19–24.
- 12 D. W. Hatchett and M. Josowicz, Composites of intrinsically conducting polymers as sensing nanomaterials, *Chem. Rev.*, 2008, **108**(2), 746–769.
- 13 A. Guiseppi-Elie, Electroconductive hydrogels: Synthesis, characterization and biomedical applications, *Biomaterials*, 2010, **31**(10), 2701–2716.
- 14 R. Jonas and L. F. Farah, Production application of microbial cellulose, *Polym. Degrad. Stab.*, 1998, **59**(1), 101–106.
- 15 D. Klemm, F. Kramer, S. Moritz, M. T. Lindstr, M. Ankerfors, D. Gray and A. Dorris, Nanocelluloses: A New Family of Nature-Based Materials, *Angew. Chem., Int. Ed.*, 2011, **50**(24), 5438–5466.
- 16 R. M. Brown and D. Montezinos, Cellulose microfibrils: visualization of biosynthetic and orienting complexes in association with the plasma membrane, *Proc. Natl. Acad. Sci. U. S. A.*, 1976, **73**(1), 143.
- 17 J. Shah and R. Malcolm Brown, Towards electronic paper displays made from microbial cellulose, *Appl. Microbiol. Biotechnol.*, 2005, **66**(4), 352–355.
- 18 D. Klemm, D. Schumann, U. Udhardt and S. Marsch, Bacterial synthesized cellulose–artificial blood vessels for microsurgery, *Prog. Polym. Sci.*, 2001, **26**(9), 1561–1603.
- 19 M. Iguchi, S. Yamanaka and A. Budhiono, Bacterial cellulose—a masterpiece of nature's arts, *J. Mater. Sci.*, 2000, **35**(2), 261–270.
- 20 W. K. Czaja, D. J. Young, M. Kawecki and R. M. Brown Jr, The future prospects of microbial cellulose in biomedical applications, *Biomacromolecules*, 2007, **8**(1), 1–12.
- 21 L. Fu, Y. Zhang, J. Zhang and G. Yang, Bacterial Cellulose for Skin Repair Materials. In *Biomedical Engineering—Frontiers and Challenges*, Fazel-Rezaei, R., Ed *InTech*, 2011, 249–274.
- 22 M. Tabuchi, Nanobiotech versus synthetic nanotech?, *Nat. Biotechnol.*, 2007, **25**(4), 389–390.
- 23 S. Ifuku, M. Nogi, K. Abe, K. Handa, F. Nakatsubo and H. Yano, Surface modification of bacterial cellulose nanofibers for property enhancement of optically transparent composites: Dependence on acetyl-group DS, *Biomacromolecules*, 2007, **8**(6), 1973–1978.
- 24 M. Nogi, K. Handa, A. N. Nakagaito and H. Yano, Optically transparent bionanofiber composites with low sensitivity to refractive index of the polymer matrix, *Appl. Phys. Lett.*, 2005, **87**, 243110.
- 25 M. Nogi and H. Yano, Transparent nanocomposites based on cellulose produced by bacteria offer potential innovation in the electronics device industry, *Adv. Mater.*, 2008, **20**(10), 1849–1852.
- 26 R. T. Olsson, M. A. S. A. Samir, G. Salazar-Alvarez, L. Belova, V. Ström, L. A. Berglund, O. Ikkala, J. Nogués and U. W. Gedde, Making flexible magnetic aerogels and stiff magnetic nanopaper using cellulose nanofibrils as templates, *Nature nanotechnology*, 2010, **5**, 584–588.
- 27 S. H. Yoon, H. J. Jin, M. C. Kook and Y. R. Pyun, Electrically conductive bacterial cellulose by incorporation of carbon nanotubes, *Biomacromolecules*, 2006, **7**(4), 1280–1284.
- 28 A. G. Cunha, C. Freire, A. Silvestre, C. P. Neto, A. Gandini, E. Orblin and P. Fardim, Highly hydrophobic biopolymers prepared by the surface pentafluorobenzoylation of cellulose substrates, *Biomacromolecules*, 2007, **8**(4), 1347–1352.
- 29 Z. Mo, Z. Zhao, H. Chen, G. Niu and H. Shi, Heterogeneous preparation of cellulose-polyaniline conductive composites with cellulose activated by acids and its electrical properties, *Carbohydr. Polym.*, 2009, **75**(4), 660–664.
- 30 S. M. Ebrahim, A. B. Kashyout and M. M. Soliman, Electrical and structural properties of polyaniline/cellulose triacetate blend films, *J. Polym. Res.*, 2007, **14**(5), 423–429.
- 31 D. A. Schumann, J. Wippermann, D. O. Klemm, F. Kramer, D. Koth, H. Kosmehl, T. Wahlers and S. Salehi-Gelani, Artificial vascular implants from bacterial cellulose: preliminary results of small arterial substitutes, *Cellulose*, 2009, **16**(5), 877–885.
- 32 J. Stejskal and R. G. Gilbert, Polyaniline. Preparation of a conducting polymer (IUPAC technical report), *Pure Appl. Chem.*, 2002, **74**(5), 857–868.
- 33 P. Anilkumar and M. Jayakannan, Divergent Nanostructures from Identical Ingredients: Unique Amphiphilic Micelle Template for Polyaniline Nanofibers, Tubes, Rods, and Spheres, *Macromolecules*, 2008, **41**(20), 7706–7715.
- 34 N. V. Blinova, J. Stejskal, M. Trchova, G. Ciric-Marjanovic and I. Sapurina, Polymerization of aniline on polyaniline membranes, *Journal of Physical Chemistry B-Condensed Phase*, 2007, **111**(10), 2440–2448.
- 35 J. Yue and A. J. Epstein, XPS study of self-doped conducting polyaniline and parent systems, *Macromolecules*, 1991, **24**(15), 4441–4445.
- 36 X. L. Wei, M. Fahlman and A. J. Epstein, XPS study of highly sulfonated polyaniline, *Macromolecules*, 1999, **32**(9), 3114–3117.
- 37 J. C. Chiang and A. G. MacDiarmid, Polyaniline: Protonic acid doping of the emeraldine form to the metallic regime, *Synth. Met.*, 1986, **13**(1–3), 193–205.
- 38 E. T. Kang, K. G. Neoh and K. L. Tan, Polyaniline: A polymer with many interesting intrinsic redox states, *Prog. Polym. Sci.*, 1998, **23**(2), 277–324.
- 39 J. Da Silva, D. De Faria, S. de Torresi and M. Temperini, Influence of thermal treatment on doped polyaniline studied by resonance Raman spectroscopy, *Macromolecules*, 2000, **33**(8), 3077–3083.
- 40 R. H. Atalla and D. L. VanderHart, The role of solid state ¹³C NMR spectroscopy in studies of the nature of native celluloses, *Solid State Nucl. Magn. Reson.*, 1999, **15**(1), 1–19.
- 41 K. Masuda, M. Adachi, A. Hirai, H. Yamamoto, H. Kaji and F. Horii, Solid-state ¹³C and ¹H spin diffusion NMR analyses of the microfibril structure for bacterial cellulose, *Solid State Nucl. Magn. Reson.*, 2003, **23**(4), 198–212.
- 42 S. Kaplan, E. M. Conwell, A. F. Richter and A. G. MacDiarmid, A solid-state NMR investigation of the structure and dynamics of polyanilines, *Synth. Met.*, 1989, **29**(1), 235–242.
- 43 S. Kababya, M. Appel, Y. Haba, G. I. Titelman and A. Schmidt, Polyaniline-Dodecylbenzene Sulfonic Acid Polymerized from Aqueous Medium: A Solid State NMR Characterization, *Macromolecules*, 1999, **32**(16), 5357–5364.