


Cite this: *RSC Adv.*, 2018, 8, 5212

New highly hydrated cellulose microfibrils with a tendril helical morphology extracted from agro-waste material: application to removal of dyes from waste water

M. El Achaby,^a N. Fayoud,^a M. C. Figueroa-Espinoza,^b H. Ben youcef^a and A. Aboulkas^c

Cocoa bean shells (CBS) are a by-product of the cocoa bean processing industry. They represent 12–20 wt% of dry cocoa beans, after having been separated from these by a roasting process. CBS often end up as a waste product which contains around 34 wt% of cellulose. The transformation of this waste into valuable and marketable products would help to improve waste disposal. Indeed, the large annual production of this waste makes it a sustainable and renewable bio-source for the production of chemicals and fibers for advanced applications. In this work, new cellulose microfibrils (CMFs) with a tendril helical morphology and highly hydrated gel-like behavior were successfully extracted from CBS waste using a controlled chemical extraction process. During this study, several physico-chemical characterizations were carried out in order to identify the properties of each of the products at different stages of treatment. Microscopic observations show that the extracted CMFs have a tendril helical shape like climbing plant tendrils. Due to this special morphology, the extracted CMFs show a highly hydrated state forming a gel network without additional modifications. The as-extracted CMFs were used as adsorbent material for the removal of methylene blue from concentrated aqueous solution, as an application to wastewater treatment for the removal of basic dyes. Swelling properties, adsorption kinetics and isotherms were carried out in batch experiments. The results indicated that the CMFs have a high swelling capacity (190%). The pseudo second order model can be effectively used to evaluate the adsorption kinetics and the adsorption isotherms can also be described well by the Langmuir isotherm model with a maximum adsorption capacity of 381.68 mg g⁻¹. Thus, the as-extracted CMFs with unique characteristics have the potential to be used as efficient adsorbent material for the removal of different cationic dyes from industrial wastewater.

Received 14th September 2017
Accepted 19th January 2018

DOI: 10.1039/c7ra10239a

rsc.li/rsc-advances

Introduction

Cellulose is considered the most abundant renewable organic polymer on earth having unique properties such as renewability, biodegradability, light weight and good mechanical properties.¹ The world yearly biomass production of cellulose has been estimated to be over 7.5×10^{10} tons, making it an inexhaustible source of raw material for environmentally friendly and biocompatible products.² The major sources of cellulose are wood and cotton, but it can also be derived from a variety of sources, such as annual plants, marine animals,

marine biomass, fungi, bacteria, invertebrates and other agro-industrial wastes.³ Depending on the extraction process, various cellulose forms named fibers, microfibers, microfibrils, nanofibrils or nanocrystals can be extracted for numerous applications.^{4,5}

In nature, depending on the degree of polymerization, the number of chains, and the way these chains pack together, the physiochemical properties of cellulose material can be enormously complicated and highly variable.⁶ Cellulose is considered to be water-insoluble compound and plays an important role in maintaining the structure of cell walls.^{2,7} However, the hierarchical arrangement of cellulose microfibrils (CMFs) in the cell wall varies depending on the origin and nature of plants. In the majority of plants, the MFCs are arranged as linear fibrils surrounded by others cell wall constituents.⁶ But in some cases, CMFs can adopt a defined tendril helical structure embedded in a soft matrix of polysaccharides, aromatic compounds and structural proteins,^{8,9} in which the sugar units and cellulose

^aMaterials Science and Nanoengineering Department, Mohamed 6 Polytechnic University, Lot 660 – Hay Moulay Rachid, Benguerir, 43150, Morocco. E-mail: mounir.elachaby@um6p.ma; Tel: +212 6 620 10 620

^bMontpellier SupAgro, UMR IATE, Montpellier F-34060, France

^cLaboratoire des procédés chimiques et matériaux appliqués (LPCMA), Faculté polydisciplinaire de Béni-Mellal, Université Sultan Moulay Slimane, BP 592, 23000 Béni-Mellal, Morocco



molecules take the form of asymmetric and chiral configurations, at the molecular level. In recent years, considerable efforts have been devoted to extract CMFs from cellulose-rich bio-sourced materials.^{3,7} However, the research in this field is oriented toward of using new bio-sourced materials, new isolation processes, and new treatments for obtaining of CMFs with special characteristics to extend the application of cellulose.

Cocoa bean shells (CBS) are the seed coats covering the cocoa cotyledons and they constitute 12–20 wt% of the dry cocoa beans, and they are separated from cocoa beans after roasting process.¹⁰ The world annual production of CBS is estimated at 700 000 tons,¹⁰ in which the Mexico country produced annually around 27 000 tons, it is the 13th and the 6th world's and Latin America's largest cocoa producer, respectively.¹¹ CBS is thus an abundant and often underexploited by-product from the cocoa industry, which can represent serious disposal and environmental problems, if badly managed.¹² CBS is devoid of any marketable value and is mainly used as raw biomass material (as food additive, garden mulch and soil conditioner) or as fuel for boilers.^{10,13,14} CBS is composed mainly of polysaccharides, hemicellulose, and lignin and other components.^{10,14} making it a sustainable and renewable biosource for production of cellulose and others biomolecules for advanced applications. Indeed, the transformation of this waste into valuable and marketable products, such as cellulose derivatives, would help to valorize this waste and improve its disposal.

Owing its excellent properties, cellulose has generated much attention and interest during these last decades in both academic and industrial fields for various application.^{4,15} Recently, the research focused on the use of cellulose and its modified forms as a new class of versatile adsorbents for wastewater treatment.¹⁶ Although many papers have been published so far discussing the importance of cellulose-based adsorbents in water pollution treatment.^{17–21} Indeed, cellulose-based adsorbents offer significant advantages over commercially available expensive materials/adsorbents for water pollution treatment.²² Recently, cellulose derivatives have been extracted from bio-sourced materials and used as potential adsorbents for waste water treatment. Hussin *et al.* have reported the possible utilization of unmodified microcrystalline cellulose (MCC) extracted from oil palm fronds as adsorbent for removal of methylene blue (MB) from aqueous solution.²³ They found that the extracted MCC exhibited a maximum adsorption capacity of 51.81 mg g^{−1}. Batmaz *et al.* reported that cellulose nanocrystals extracted from cellulosic fibers can also be used as adsorbents for removal of cationic dyes (MB) from concentrated aqueous solutions, and a relatively high adsorption capacity was measured (118 mg g^{−1}).²⁴ Douissa *et al.* have reported the efficiency of extracted cellulose from *Posidonia oceanica* to remove methylene blue from water. The appropriate adsorption capacity (305.45 mg g^{−1}) was found higher than that obtained by raw material (*Posidonia oceanica*), showing the very good potentiality of extracted cellulose as no expensive adsorbent.²⁵

Nevertheless, little research is focused on the extraction of cellulose derivatives from bio-sourced materials and their use directly without further modifications as potential adsorbents

for wastewater treatment. In the current investigation, new CMFs with tendril helical morphology and highly hydrated gel-like behavior were extracted from CBS waste, using the same chemical treatments that are largely used for cellulose extraction from typical terrestrial lignocellulosic materials, *e.g.* alkali and bleaching treatments.^{26,27} After its successful extraction, the as-obtained CMFs were successfully characterized in terms of their physico-chemical properties, and used as potential adsorbents for adsorption of cationic dyes from concentrated aqueous solution, as application for wastewater treatment for the removal of basic dyes. MB was chosen as adsorbate in this study, due to its strong adsorption on solid surfaces and its increased use in textile and paper industry. Swelling properties, adsorption kinetics and isotherms were carried out at batch experiments, and various theoretical models were used to evaluate the maximum adsorption capacity and to understand the adsorption mechanism.

Materials and experimental details

Materials

The dried CBS used in this study were obtained from fermented and roasted cocoa beans collected from Tabasco (Mexico) in 2016. Firstly, the CBS waste was chopped, then dried in a fluidized bed dryer (GRACO, TGA101) for 90 min at 60 °C. The dried CBS was milled using a Knife Shredder (RETSCH SM100, Haan, Germany) equipped with 5 mm sieve. Finally, it was stored in the dark at 4 °C until the extraction was performed. The moisture content of the CBS was about 4.59 wt%. All the analytical grade chemicals used for purification and bleaching treatments were purchased from Sigma-Aldrich and used without further purification. The cationic Methylene Blue (MB) was purchased from Sigma Aldrich (M9140-25G) and used as received.

Treatment of CBS and CMFs extraction

CMFs in highly hydrated gel form were extracted from raw CBS using alkali and bleaching treatments. Firstly, a prewashing of ground CBS was carried out by distilled water for 1 hour at 60 °C under mechanical stirring. Then, the prewashed CBS were treated 3 times with 4 wt% NaOH solution at 80 °C for 2 hours under mechanical stirring. Then, the obtained alkali treated-CBS was subjected to a bleaching treatment with a solution made up of equal parts (v/v) of acetate buffer (27 g NaOH and 75 mL glacial acetic acid, diluted to 1 L of distilled water) and aqueous sodium chlorite (1.7 wt% NaClO₂ in water). The bleaching treatment was performed 3 times for 2 h at 80 °C, under magnetic stirring. The obtained cellulose product was washed several times with distilled water, resulting in highly hydrated white gel composed of CMFs, as confirmed by microscopic observations. The overall steps these purification/extraction and digital images of each obtained product are presented in Fig. 1.

Characterization techniques

Elementary analysis was performed mainly to determine the total carbon, oxygen, hydrogen and nitrogen contents before



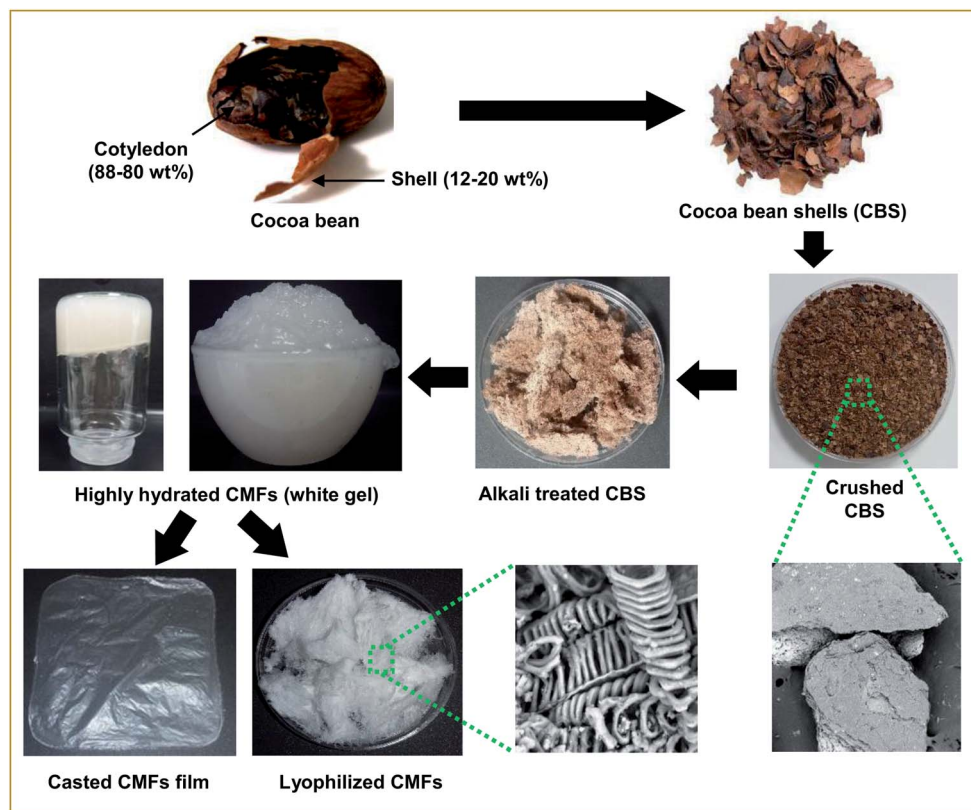


Fig. 1 Schematic representation of raw CBS treatment and digital images of the obtained products.

and after the extraction of cellulose microfibrils. It was carried out on elemental analyzer system GmbH; analyzer vario MICRO V4.0.2. Morphology of samples was evaluated using a scanning electron microscope (SEM) (Phenom ProX) operated at 10 kV. Fourier Transform Infrared Spectroscopy (FTIR) of all studied samples were measured on a Tensor27 apparatus, the experiments were recorded in transmittance mode in the range of 4000–400 cm^{-1} with a resolution of 4 cm^{-1} and an accumulation of 16 scans. Thermogravimetric analysis (TGA) (Mettler Toledo) for all samples was conducted under a nitrogen atmosphere between 25 and 700 $^{\circ}\text{C}$, at a heating rate of 10 $^{\circ}\text{C min}^{-1}$. X-ray diffraction (XRD) characterizations of samples were performed on a Bruker diffractometer D8 Advance using $\text{Cu-K}\alpha$ radiation. The diffraction patterns were obtained at diffraction angles between 5 and 50 $^{\circ}$, at room temperature. The crystallinity index (CrI%) of the samples was estimated using Segal's equation: $\text{CrI} = ((I_{002} - I_{\text{am}})/I_{002}) \times 100$,²⁸ where I_{002} and I_{am} are the peak intensities of crystalline and amorphous cellulose, respectively.

Swelling ratio

The equilibrium swelling ratio (SR) of CMFs was measured in distilled water at room temperature. For that, structural stable film with thickness of 180 μm was successfully casted from the obtained CMFs gel on Petri dish and air dried (Fig. 1). To determine the SR of the film sample, square dried film with an initial weight (W_0) was placed in a glass crystallizer with excess

water. The swollen film was removed after an interval of time. The excess water on the surface of film was removed with filter paper, and the weight of each sample (W_t) was measured. The percentage of SR (%) was calculated using the following eqn (1):

$$\text{SR}(\%) = \frac{(W_t - W_0)}{W_0} \times 100 \quad (1)$$

Adsorption studies

All adsorption studies were performed at batch experiments. A stock solution of MB at a concentration of 1000 mg L^{-1} was prepared in distilled water and suitably diluted to the required initial concentrations. The concentration of the MB dye for all samples during the experimental tests were measured using a UV-visible spectrophotometer (PerkinElmer Lambda 25 spectrophotometer) at a wavelength of 664 nm which correspond to the maximum absorbance for MB.²⁴ For each experiment, the concentration of MB aqueous solution was obtained from linear standard calibration curves ($R^2 = 0.999$).

Adsorption kinetics

The adsorption experiments were conducted at room temperature using an initial concentration of MB of 50 mg L^{-1} . For all experiments, the pH of the initial MB solution was kept at 7. After that, a pre-weighted amount of adsorbent (10 mg) was added to 20 mL of MB solution in glass bottles, and the mixture was slowly stirred (100 rpm) for various contact time intervals (5–150 min). At a predetermined time interval, sample aliquots



(~3 mL) were withdrawn from the bottle, using 25 mm disposable syringe filter (0.45 μm cellulose acetate membrane), and were either introduced directly into plastic cell or diluted with a suitable volume of distilled water to allow for Beer's law concentration analysis on the UV-vis spectrophotometer. The adsorbed amount of MB q_t (mg g^{-1}), at a particular time interval t , was calculated using the following expression (2):

$$q_t = \frac{(C_0 - C_t)V}{W} \quad (2)$$

where C_0 (mg L^{-1}) is the initial concentration of MB, C_t (mg L^{-1}) is the dye concentration at time t , V (L) is the volume of the solution and W (g) is the weight of used adsorbent.

Adsorption at equilibrium state

The adsorption isotherm at the equilibrium state was carried out in the same conditions by varying the initial MB concentration from 25 to 500 mg L^{-1} . The adsorption capacity q_e (mg g^{-1}) at the equilibrium was measured using the following eqn (3):

$$q_e = \frac{(C_0 - C_e)V}{W} \quad (3)$$

where C_0 (mg L^{-1}) is the initial concentration of MB, C_e (mg L^{-1}) is the concentration of MB at equilibrium (150 min of contact time), V (mL) is the volume of the solution and W (mg) is the weight of adsorbent.

Results and discussion

Extraction of CMFs

CMFs were extracted from CBS by means of controlled chemical treatments. Obviously, the initial ground CBS grains have brown color (Fig. 1), due to its diversified constituents (lignin, hemicellulose, cellulose, protein, pectin and others). Starting with the alkali treatment of CBS, the main objective of this treatment was to eliminate the whole non-cellulosic cementing components (lignin, hemicelluloses, pectin, ash, and other impurities), resulting in the alkali-treated CBS. During this alkali treatment, some alkali-labile linkages (ether and ester linkages) between lignin monomers or between lignin and polysaccharides may have been broken, resulting in a partial breakdown of CBS grains into interconnected fibers, with yellowish brown color (Fig. 1). The yield of this alkali-treated CBS was about 39% (in dry mass). The following treatment was the bleaching treatment, with the main purpose to obtain pure cellulosic fibers. This whiteness and purification is to be achieved simply by removing the residual lignin and the other unwanted non-cellulosic components adhering to the cellulose fibers, with the help of bleaching agents. During this treatment, the residual lignin molecules react with NaClO_2 and an oxidative fragmentation takes place resulting into soluble lignin molecules in the bleaching solution, which lead into total removal of lignin, and leave the CMFs unaltered.²⁹ The yield of the extracted CMFs was determined at 20% (in regard to the initial weight of raw CBS). Very interestingly, the CMFs were

extracted in highly hydrated white gel form without any further chemical treatment (Fig. 1).

It is noteworthy that the extracted CMFs gel can easily be transformed into powdered form *via* freeze drying process (Fig. 1) or in form of stable film using the casting process followed by air drying (Fig. 1). This was surprising, because only the bacterial cellulose or modified cellulose can exhibit this characteristic.^{30,31} It have been reported that the natural extracted cellulose could form a thick gel or highly hygroscopic material after a suitable additional modification such as treatments in an alkaline aqueous solution with cooling by physical cross-linking.³² Herein, we report that a highly hydrated CMFs gel can be directly extracted from raw CBS *via* simple bleaching treatment. Morphological analysis by SEM showed that the extracted cellulose exhibits a mixture of linear long microfibrils and tendril helical microfibrils (see below). May be this special morphology is responsible for obtaining of gel-like cellulose with highly hydrate state.

Morphology of CBS and CMFs

As a widely employed technique, SEM was used to investigate and identify the morphological features of CBS and extracted CMFs, examples of the obtained SEM results are shown in Fig. 2. According to the above discussion, lignin molecules, hemicelluloses and other extractive substances were removed from CBS by means of bleaching treatment, leaving only the cellulose product. From the SEM results, it is clearly found that

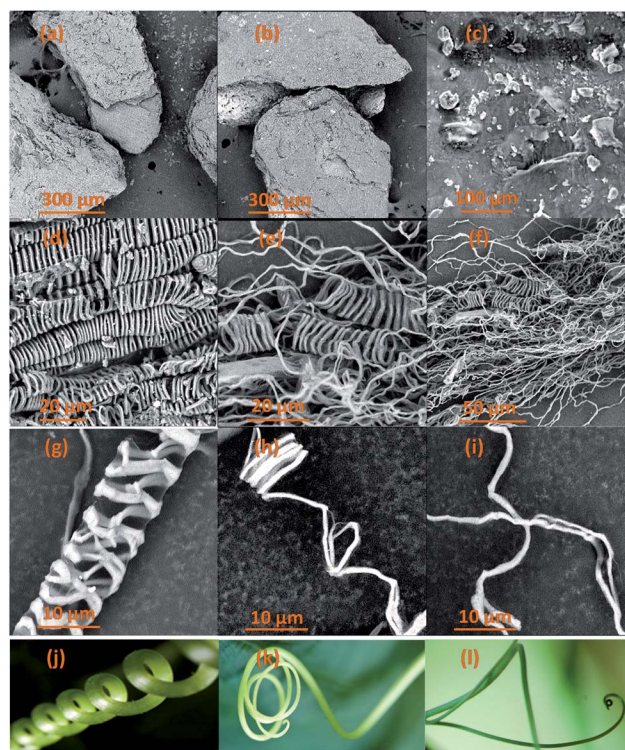


Fig. 2 SEM images of raw CBS (a–c), extracted CMFs (d–i) samples and examples of plant tendrils (j–l) comparable to the morphology of the extracted CMFs.



Table 1 Elementary analysis of CBS, alkali treated CBS and extracted CMFs

	C (%)	H (%)	O (%)	N (%)
CBS	41.64 ± 0.33	5.80 ± 0.02	50.27	2.29 ± 0.03
Alkali treated CBS	35.86 ± 0.12	5.34 ± 0.01	58.39	0.41 ± 0.01
Extracted CMFs	39.00 ± 0.10	6.06 ± 0.18	54.92	0.00 ± 0.00

the morphologies of CBS and the extracted CMFs are extremely different. The crushed CBS exhibit a layer of surface deposits mostly composed of lignin, hemicellulose, and other non-cellulosic substances that cover the CMFs inside (Fig. 2a–c). For the extracted CMFs, a remarkable finding of this new extraction can be seen in Fig. 2d–i. It is found that the morphology of CMFs consisted of a mixture of linear long microfibrils and tendril helical microfibrils. The helical shape of the obtained CMFs was comparable to that of a climbing plant tendril (Fig. 2j–l). The average diameter of individual microfibril helices was determined at 10 μm , whereas the average diameter of microfibril forming helices was measured at about 1 μm . Nonetheless it was not possible to determine the average length of CMFs by SEM microscopy analysis, because the CMFs are much entangled forming a spaghetti-like structure (Fig. 2f).

Elementary analysis

Table 1 shows the result obtained from elemental analysis for all studied samples (CBS, alkali treated CBS and extracted CMFs). From this result, it should be noted that all samples present high carbon and oxygen contents, indicating the organic nature of these samples. The determination of nitrogen (N) in CBS sample indicates that the presence of proteins in the original sample. Notably, a total removal of N (%) content in extracted CMFs indicates a total elimination of proteins after

bleaching treatment. It must also be noticed that the ratio O/C of the extracted CMFs was calculated at about 1.40, wherein 0.83 is the theoretical value of pure cellulose.³³ The relatively high O/C ratio in our case is probably due to the excess of hydroxyl groups in the surface of microfibrils, which can explain in first time the formation of highly hydrated CMFs gel directly after the extraction process.

Structure and crystallinity

Direct information about changes in chemical functionality can be obtained by FTIR spectroscopy which has been extensively used for structural analysis of the material before and after chemical treatments. FTIR spectra of raw CBS, alkali treated CBS and extracted CMFs are shown in the Fig. 3. Additionally, the FTIR spectrum of commercial MCC is also shown in this Fig. 3 for eventual comparison. Obviously, all samples spectra exhibit the characteristic bands of cellulose molecule. The absorption bands located at 3300 cm^{-1} and 2980 cm^{-1} are attributed to the –OH stretching vibration of hydrogen bonded hydroxyl groups, showing the hydrophilic tendency of all samples,³⁴ and the –CH groups of cellulose molecule,²⁹ respectively. In addition, the absorption band at 1610 cm^{-1} is attributed to the adsorbed water.³⁵ Beside these absorption bands, other specific absorption bands can also be seen in all spectra. The band centered at 1160 cm^{-1} is assigned to the C–O–C stretching of the β -1,4-glycosidic ring linkages between the D-glucose units in cellulose.²⁹ Noticeably, this band was gradually going intense from raw CBS to extracted CMFs, indicating that the cellulose content was increased during different chemical treatments starting from raw CBS and ending up with extracted CMFs. In addition, the two bands observed at 1035 and 894 cm^{-1} in all spectra are attributed to the C–O stretching and C–H rocking vibrations of cellulose of the carbohydrates.³⁶

The most distinct spectral changes that can be detected in the alkali treated CBS and the extracted CMFs comparing to the raw CBS are the absorption band located at 1742 cm^{-1} , which is

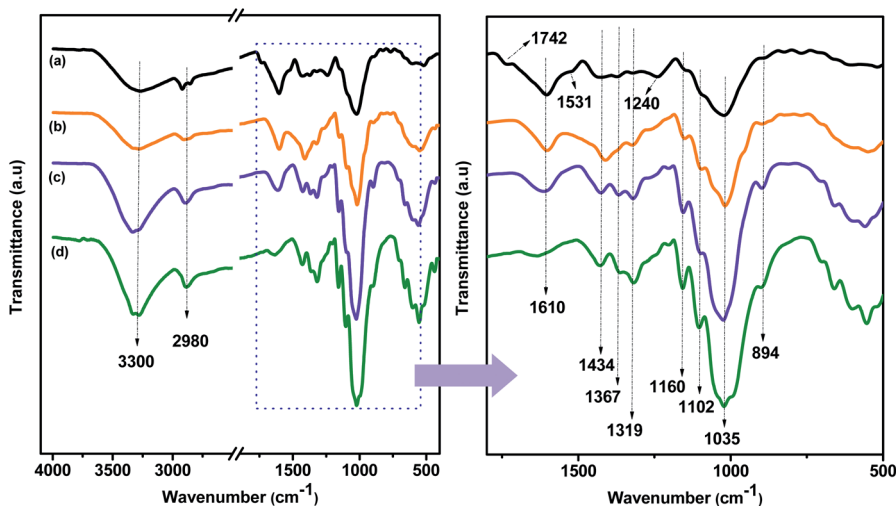


Fig. 3 FTIR spectra of CBS (a), alkali treated CBS (b) and extracted CMFs (c). The FTIR spectrum of commercial MCC (d) is included for comparison.



assigned to the uronic ester and acetyl groups of hemicelluloses and the ester linkage of carboxylic group of the ferulic and *p*-coumaric acids of lignin,³⁷ and the absorption band located at 1531 cm^{-1} which is due to the stretching vibration of N–H band of the protein amide II structure,³⁸ and the C=C stretching from aromatic hydrocarbons of lignin.³⁹ Additionally, the band observed at 1240 cm^{-1} is attributed to the C–O out of plane stretching vibration of the aryl group in the lignin molecules.³⁶ These three bands are clearly present in raw CBS, as it can be seen from the Fig. 3b. However, after the alkali treatment, in the alkali treated SBC spectrum, these bands were drastically reduced confirming that the lignin and hemicelluloses were not completely removed.

The disappearance of these bands in the spectrum of the extracted CMFs was observed, which is an indication that all non-cellulosic components were completely removed after the bleaching treatment. These results were already confirmed by the elemental analysis data (Table 1) in which the % N content was totally removed after bleaching treatment.

The FTIR spectra of all samples show another absorption band located at 1102 cm^{-1} and it is attributed to presence of β -glucosidic ether linkages (C–O–C) resulted in the vibration of wagging, deformation and twisting modes of anhydroglucopyranose ring skeleton.⁴⁰ In addition, the absorbance bands at 1434 cm^{-1} , 1367 cm^{-1} and 1319 cm^{-1} in alkali treated CBS and extracted CMFs samples were mainly associated to the presence of cellulose chain.²⁸ It is interesting to observe that raw CBS sample did not exhibit bands at 1319 cm^{-1} and 1367 cm^{-1} , as shown in Fig. 3b. This is an indication that the CMFs were embedded in the matrix of amorphous non-cellulosic components in the raw CBS, as observed by SEM analysis (Fig. 2). Besides that, the extracted CMFs from raw CBS was comparable with the commercial available MCC in terms of FTIR analysis (Fig. 3), suggesting that pure cellulose has been successfully produced from raw CBS.

XRD analysis was performed to investigate the crystalline structure and the crystallinity index (CrI) of all studied

samples (raw CBS, alkali treated CBS and extracted CMFs). Herein, it should be noted that the presence of a crystallinity feature in all samples is generated from the ordered structure of cellulose, which is generated from intra- and intermolecular hydrogen bonding occur in cellulose *via* hydroxyl groups, resulting in various ordered crystalline arrangements.²⁹ Fig. 4 shows the XRD patterns for raw CBS, alkali treated CBS and extracted CMFs. From this result, all samples show three major peaks located at $2\theta = 14.9^\circ$, 16.8° , and 22.6° , which are characteristic of the crystal polymorphs of cellulose. These peaks correspond to the $1\bar{1}0$, 110 and 200 crystalline planes, respectively.²⁷

The crystallinity index (CrI) determined was found to be about 40.46%, 71.3%, and 75.57% for raw CBS, alkali treated CBS and extracted CMFs, respectively, demonstrating that the crystallinity of the material progressively increases during the chemical extraction. In the case of raw CBS, the cellulose is embedded in the matrix of amorphous non-cellulosic components thus a low crystallinity is determined.²⁹ In the case of CMFs, the increased crystallinity was ascribed to the total removal of amorphous components, leading only cellulose structure that contains amorphous and ordered regions, which was also confirmed by FTIR analysis of all studied samples at different stages of treatment (Fig. 3).

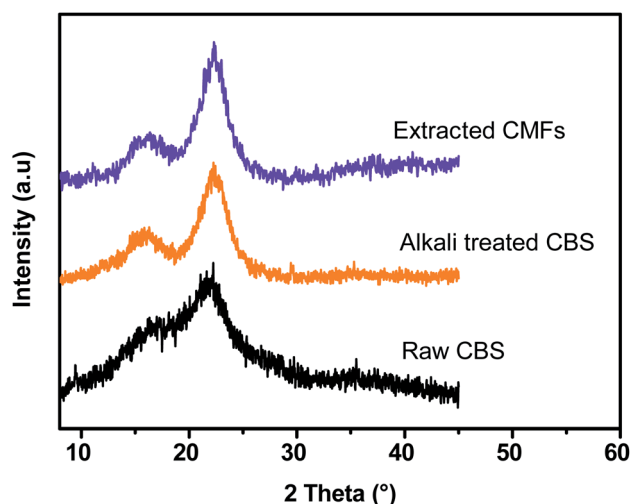


Fig. 4 XRD patterns of CBS, alkali treated CBS and extracted CMFs.

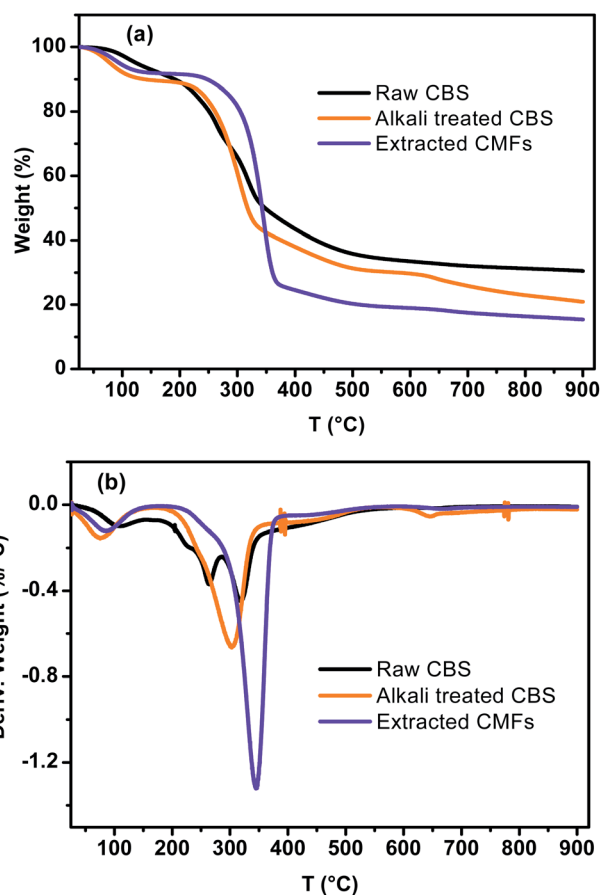


Fig. 5 TGA (a) DTG (b) curves of CBS, alkali treated CBS and extracted CMFs.



Thermal stability

Fig. 5 shows the TGA–DTG curves of CBS, alkali treated CBS and extracted CMFs. From these results, it can be viewed that all samples demonstrated a weight loss at around 100 °C, which was caused by evaporation of loosely bound moisture on the surface and the intermolecular hydrogen bonded chemisorbed water.²¹ The degradation of raw CBS occurs at three different stages. The first and second stages start forward at around 184 °C and 235 °C respectively, which are likely due to the decomposition of non-cellulosic components, such as hemicelluloses and lignin.⁴¹ The corresponding maximum temperatures from DTG curves were observed at 224 °C and 262 °C. The third degradation stage of raw CBS starts at 286 °C with corresponding maximum temperature of 323 °C, which is due to degradation of cellulose molecules.⁴² For alkali treated CBS, a single decomposition stage is observed, which occurs between 184 and 360 °C, indicated by a shoulder peak at DTG curve with a maximum temperature of 304 °C. This degradation is due to decomposition of small amounts of lignin and hemicelluloses that are still presented in this sample as well as of cellulose molecules. This finding likely reflect the partial elimination of lignin, hemicelluloses and other non-cellulosic segments,⁴³ which are started to decompose at low temperatures (<230 °C), this is confirmed by the absence of small shoulder DTG peaks observed in raw CBS curve.

Concerning the extracted CMFs, the degradation starts at 269 °C and the corresponding maximum temperature is reached at 345 °C. It can be noted that the onset thermal decomposition shifted to higher temperature, in regard to raw CBS and alkali treated CBS. This is due to the purification process which lead to the complete elimination of hemicelluloses and lignin residues and other impurities.^{27,28} This finding is comparable to that observed for cellulose microfibrils extracted from sugarcane bagasse,^{26,27} and red algal waste.²⁸

Generally, cellulose thermal degradation involves dehydration, depolymerization and decomposition of glycosyl-units and then formation of a charred residue.³⁴

Additionally, the TGA analysis showed minimum residual mass for extracted CMFs in regard to raw CBS and alkali treated CBS samples, which probably due to the absence of the charred residue that can be generated from the degradation of lignin and hemicelluloses.³⁷

Swelling ratio of CMFs

With regard to the use of CMFs as adsorbent for removal of MB from concentrated aqueous solution, the swelling capacity of the as-extracted CMFs was investigated. It is noteworthy that the CMFs were extracted in highly hydrated gel form (Fig. 1), and they were easily transformed into powdered form *via* freeze drying process or in form of stable film using the casting process followed by air drying. Indeed, the swelling ratio (SR) of CMFs based film was measured by immersing of such film in distilled water for various contact time intervals. The obtained results are shown in Fig. 6. From these results, the SR increased rapidly with the prolongation of immersing time and reached swelling equilibrium within 40 min. The high obtained SR was about 190% and reached after 120 min, suggesting that the CMFs have a high capacity to absorb water. This behavior is due to the special morphology of the extracted microfibrils (mixture of linear long microfibrils and tendril helical microfibrils) as well as the entanglement network (spaghetti-like structure) formed by the microfibrils at their solid state, as observed by SEM observations (Fig. 2). This network may have a high capacity to absorb water throughout its void regions. Additionally, the extracted CMFs may have an excess of hydroxyl groups on their surface, as deduced from the high O/C ratio obtained by elemental analysis (Table 1). The hydroxyl groups on the surface of microfibrils are able to interact with water molecules, thus improving the swelling capacity.

Application of the extracted CMFs for MB adsorption

Freeze dried CMFs were tested as bioadsorbent for removal of MB from concentrated aqueous solution. Adsorption kinetic

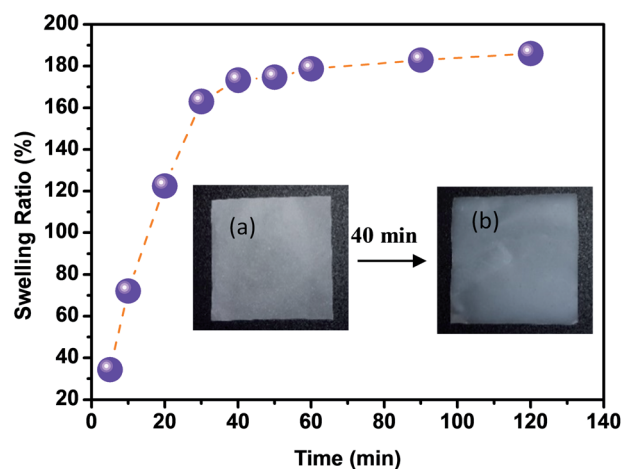


Fig. 6 Swelling ratio of extracted CMFs in form of casted film (inset: CMFs film before (a) and after (b) its swelling at 40 min).

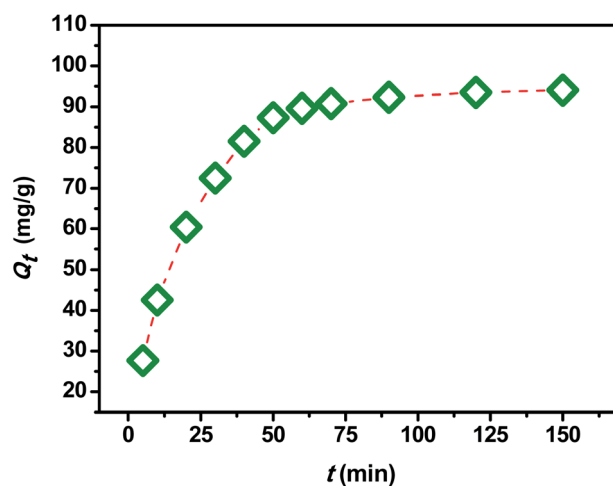


Fig. 7 Adsorption capacity of the extracted CMFs as function of the contact time (*t*).



and isotherm were evaluated at controlled conditions and the obtained results were analyzed and simulated using kinetic and isotherm models. The as-extracted CMFs exhibit native cellulose-like structure and well-defined morphology. It was observed that CMFs is microscale material consisting of linear long microfibrils and tendril helical microfibrils, forming an entangled network (spaghetti-like structure). As such, CMFs in aqueous solution contain micropores in the case of helical microfibrils and contain naturally active hydroxyl groups on their surfaces. In conventional theory of adsorption, there are two adsorption processes, firstly, adsorbate must first be in contact with adsorbent (mass transfer of adsorbate) then the filling of pores due to capillary forces (diffusion of adsorbate through pores).¹⁶ In our case, the removal of MB by CMFs can be ensured by both adsorption processes.

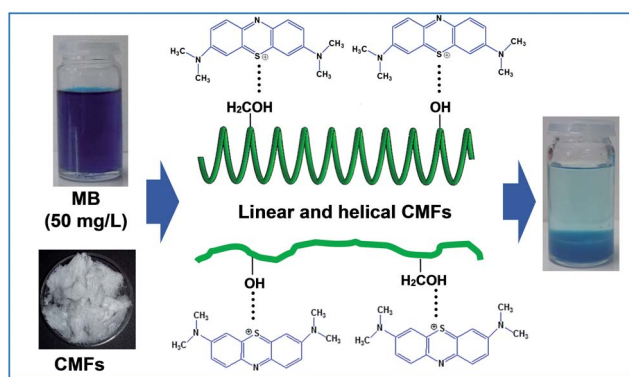


Fig. 8 Adsorption mechanism of MB molecule onto CMFs.

Effect of contact time and adsorption kinetics

In order to investigate the influence of the contact time on the removal of MB by the extracted CMFs, the adsorption capacity was studied at various times ranging from 0 to 150 min. The obtained results are shown in Fig. 7. From this, it can be observed that the MB adsorption process on CMFs presents generally two phases; an initial rapid phase, where adsorption capacity sharply increased and reached an equilibrium state within 60 min, due to the rapid surface adsorption, and a second slow phase associated with the internal surface created by helical morphology of CMFs. This can be explained by the fact that initially the adsorption sites of extracted CMFs were vacant and therefore easily accessible to the MB dye molecules, considering the associative interactions that take place between the MB cationic dye and the CMFs' surface,⁴⁴ as shown in the representative scheme in Fig. 8. However after this initial stage, the remaining active sites of extracted CMFs are difficult to be occupied due to repulsive force towards free MB molecules over times which also resulted in the lower adsorption rate. When repulsive and driving force reached the equilibrium, the reaction equilibrium achieved and the MB adsorption would not be influenced by time of the reaction any more.

In order to evaluate the kinetic mechanism that controls the MB biosorption process, the obtained adsorption data were analyzed and simulated using four kinetic models including the pseudo first order, pseudo second order, Elovich and intra-particle diffusion models. It should be noted that the conformity between the experimental data and the predicted model is based on the values of the determination coefficients (R^2); thus, the value of R^2 closest to the unit will indicate the appropriate model to describe the adsorption kinetic of MB.

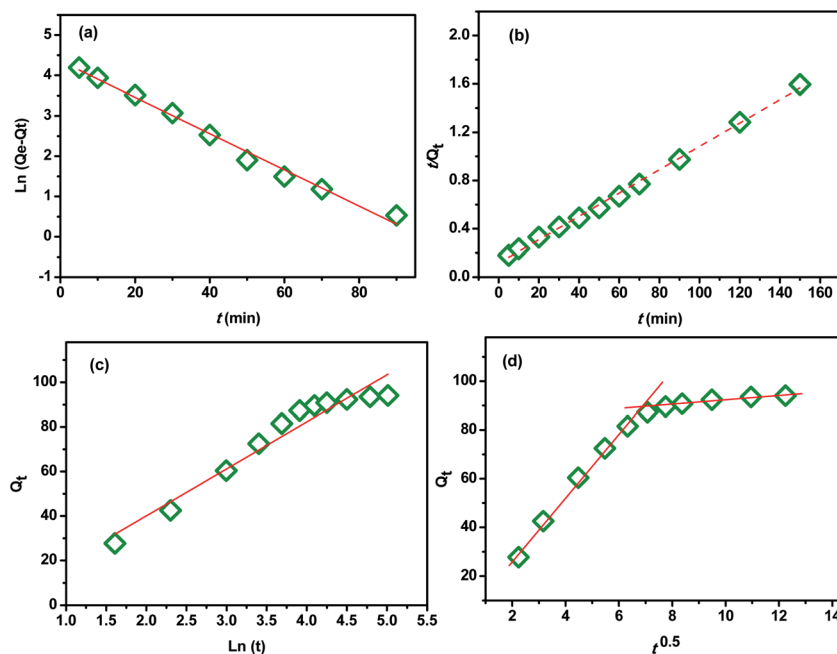


Fig. 9 (a) Pseudo-first order equation, (b) pseudo-second order equation, (d) Elovich and (c) intraparticle diffusion.



Pseudo first order and pseudo second order models. The pseudo first order (eqn (4)) and the pseudo second order (eqn (5)) models were most commonly used to determine the rate constants and to analyze the adsorption process. The linear forms of these kinetic models can be expressed as:

$$\ln(q_e - q_t) = \ln q_e - k_1 t \quad (4)$$

$$t/q_t = 1/k_2 q_e^2 + (1/q_e)t \quad (5)$$

where q_e (mg g^{-1}) is the adsorption capacity at equilibrium state, q_t (mg g^{-1}) is the adsorption capacity at contact time t (min), k_1 (min^{-1}) and k_2 ($\text{g mg}^{-1} \text{min}^{-1}$) are the equilibrium rate constants of pseudo-first-order and pseudo second order models, respectively. For the pseudo first order model, q_e and k_1 were determined from the slope and intercept of the plot of eqn (4). While for the pseudo second order, q_e and k_2 can be calculated from the slope and intercept of the plot of eqn (5). The experimental data were fitted linearly by using eqn (4) and (5), and the obtained results are shown in Fig. 9a and b, respectively. According to the obtained kinetic parameters summarized in Table 2, the R^2 value of the pseudo-second-order ($R^2 = 0.997$) is higher compared with that obtained from the pseudo-first-order ($R^2 = 0.988$). Moreover, the experimentally determined ($q_{e,\text{exp}}$) value, obtained from the linear plot, for the pseudo-first-order model did not agree with the calculated value ($q_{e,\text{calcul}}$) as mentioned in Table 2. Indicating that the adsorption process of MB onto extracted CMFs did not comply with the latter model. However, the linear plot of t/q_t versus t showed a good agreement between experimental and calculated values of q_e . These findings indicate that the pseudo-second-order mechanism is the predominant in the MB adsorption process for the used cellulose.

Therefore, this evidence indicated that the adsorption of MB onto CMFs follows pseudo-second-order kinetic model. A

similar trend was also reported for the adsorption of MB on cellulose extracted from *Posidonia oceanica*,²⁵ and on cellulose nanocrystal filled polyacrylamide nanocomposite hydrogels,⁴⁵ which the adsorption rate was determined by the free adsorption site on the surface of the adsorbent.

Elovich model. The Elovich model (eqn (6)) was used to describe the chemical adsorption mechanism of MB onto extracted CMFs, because it is based on a general second-order reaction mechanism for heterogeneous adsorption processes.⁴⁵ The Elovich equation is expressed in linear form as:

$$q_t = \frac{1}{\beta} \ln(\alpha\beta) + \frac{1}{\beta} \ln t \quad (6)$$

where α ($\text{mg g}^{-1} \text{min}^{-2}$) is the initial adsorption rate and β ($\text{g mg}^{-1} \text{min}^{-1}$) is the desorption constant, which is related to the activation energy and the extent of surface coverage for chemisorption.⁴⁵ The experimental data in Fig. 9d were fitted linearly by using eqn (6) (q_t vs. $\ln t$). The values of α , β , and R^2 are calculated and listed in Table 2. From the obtained data, it can be seen that the Elovich model don't fit to the experimental data based on its R^2 value (0.93). This further indicates that the adsorption of MB onto the extracted CMFs is not a chemical process.⁴⁵

Intra-particle diffusion model. The intra-particle diffusion model (eqn (7)) was used to further determine the mechanism of MB removal in terms of dye molecules diffusion. The intra-particle diffusion equation is expressed as:

$$q_t = k_i t^{1/2} + C_i \quad (7)$$

where q_t (mg g^{-1}) is the dye fraction uptake at time t , k_i ($\text{mg g}^{-1} \text{min}^{-0.5}$) is the intra-particle diffusion constant, and C_i (mg g^{-1}) is a constant that gives an idea about the thickness of boundary layer. If $C = 0$, the intra-particle diffusion is the only rate-limiting step, conversely if $C \neq 0$, the adsorption process is controlled by various adsorption mechanisms.⁴⁸ The plot of q_t vs. $t^{0.5}$ is shown in Fig. 9c and the obtained parameters are listed in Table 2. It's assumed that the transport of adsorbate from the solution phase to the surface of the adsorbent particles may be controlled by one or more steps.⁴⁶ In our study, it can be observed from Fig. 9c that there are two steps during the biosorption process of MB onto extracted CMFs. The initial linear portion with higher slope ($k_{i1} = 12.4$) indicates the transport of dye molecules from the solution to the external surface of extracted CMFs by diffusion through boundary layer.⁴⁵ The second linear portion with lower ($k_{i2} = 0.97$) slope is the gradual adsorption stage with intra-particle diffusion dominating.⁴⁶ The MB is first adsorbed by the outer surface of used CMFs, for that the adsorption rate is very high. Once the outer surface is completely saturated, the dye molecule diffuses into the internal pores within the microfibrils network, and is finally adsorbed by the inner surface of the extracted cellulose. When the dye molecules diffuse through the internal pores, the diffusion resistance increases, which results in a decrease in the diffusion rate. As expected, the values of C (Table 2) show that the constants C are not zero in any case. This showed that pore diffusion was not the rate-limiting step. Consequently, the adsorption process may be of a complex nature consisting of both,

Table 2 Parameters of different kinetic models for MB adsorption on the extracted CMFs

Kinetic model	Parameter	Values
Pseudo-first order model	k_1 (min^{-1})	0.1036
	$q_{e,\text{calcul}}$ (mg g^{-1})	78.10
	R^2	0.988
Pseudo-second order model	k_2 ($\text{g mg}^{-1} \text{min}^{-1}$)	0.0007
	$q_{e,\text{calcul}}$ (mg g^{-1})	104.16
	R^2	0.997
	$q_{e(\text{exp})}$ (mg g^{-1})	95.05
Elovich model	α ($\text{g g}^{-1} \text{min}^{-1}$)	0.127
	β (g mg^{-1})	0.047
	R^2	0.939
Intraparticle diffusion model	First stage	
	k_{i1} ($\text{mg g}^{-1} \text{min}^{-0.5}$)	12.40
	C_{i1} (mg g^{-1})	2.55
	R^2	0.987
	Second stage	
	k_{i2} ($\text{mg g}^{-1} \text{min}^{-0.5}$)	0.97
	C_{i2} (mg g^{-1})	82.52
	R^2	0.918



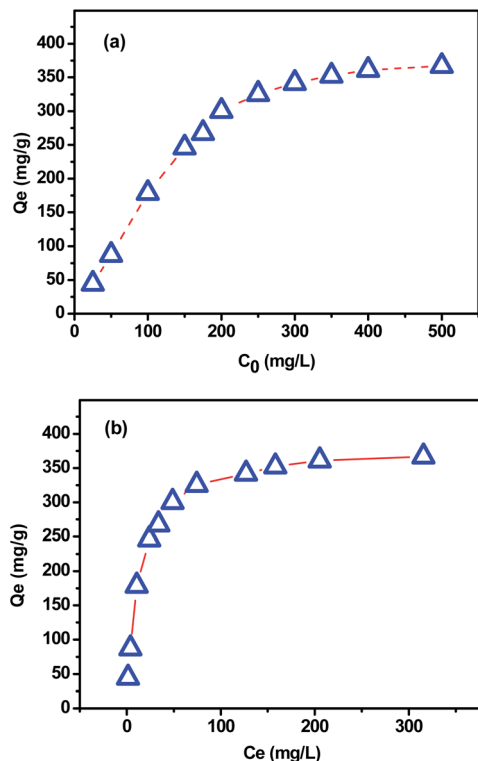


Fig. 10 Adsorption capacity as function of (a) initial and (b) equilibrium MB concentration.

surface adsorption and intra-particle diffusion.⁴⁶ Furthermore, as can be seen in Table 2, the value of R^2 in first adsorption step is larger than that in second step while C_1 was smaller than C_2 . This shows that the intra-particle diffusion can predict the kinetic process at different stages.⁴⁶

Effect of dye concentration and adsorption isotherms

The dye concentration in solution can strongly affect the sorption process. The effect of this parameter on MB removal by using the extracted CMFs was investigated. Fig. 10 represents the adsorption capacity (q_e) as function of initial (C_0) and equilibrium (C_e) MB concentration measured at the equilibrium contact time of 150 min. From these obtained results, it was found that when the initial dye concentration increased from 25 to 500 mg L⁻¹, the adsorbent amount of MB increased from 60 to 380 mg g⁻¹. This behavior is due to the fact that the initial dye concentration provides a powerful driving force to overcome the mass transfer resistance between the aqueous and solid phases.⁴⁶ Beyond the initial concentration of about 300 mg L⁻¹, the adsorption kept nearly constant owing to the saturation of the active sites of extracted CMFs.⁴⁷

In order to determine the parameters associated with MB adsorption, the experimental data were modeled with the three most commonly used isotherm models namely Langmuir, Freundlich and Temkin.

The Langmuir isotherm model (eqn (8)) assumes that the adsorption is monolayer onto identical active sites of adsorbent with uniform adsorption energies and without lateral

interaction and steric hindrance between the adsorbed molecules.⁴⁸ It can be expressed in the linear form as:

$$\frac{C_e}{q_e} = \frac{1}{k_L q_m} + \frac{1}{q_m} C_e \quad (8)$$

where C_e (mg L⁻¹) is the equilibrium concentration of MB, q_e (mg g⁻¹) and q_{\max} (mg g⁻¹) the equilibrium and maximum

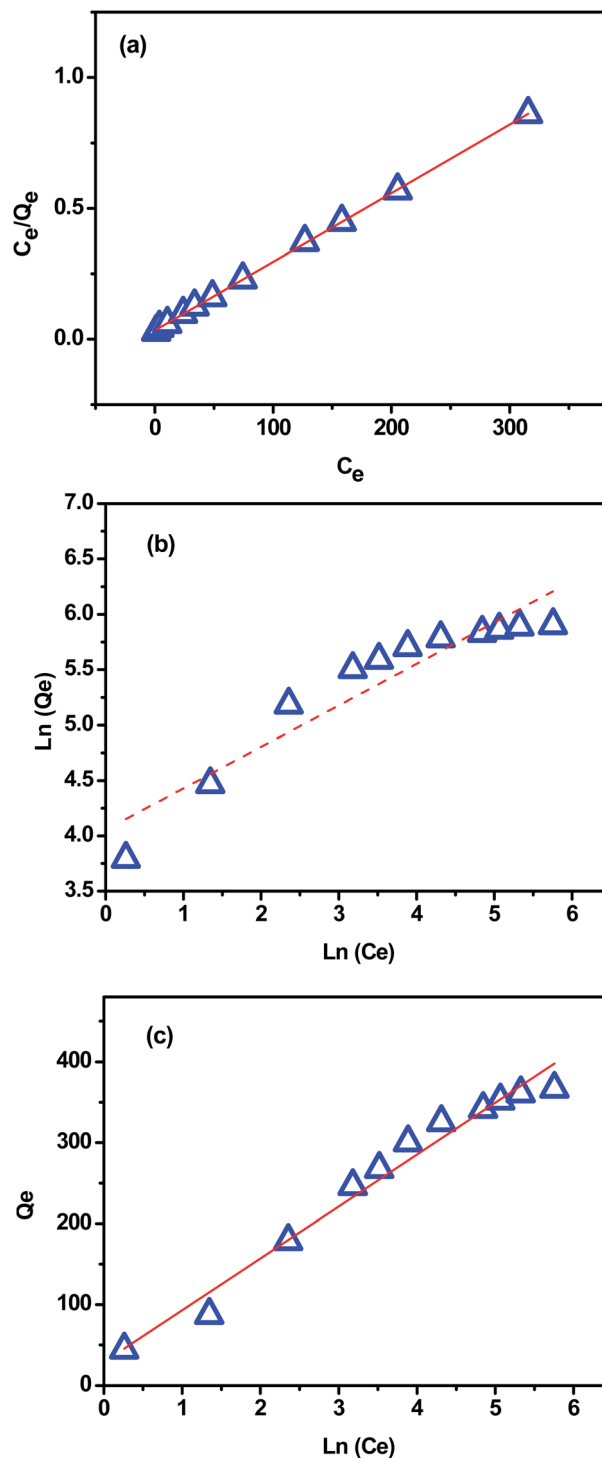


Fig. 11 (a) Langmuir isotherm, (b) Freundlich isotherm and (c) Temkin isotherm.



Table 3 Parameters of different isotherms for adsorption of MB on CCBC sample

Langmuir				Freundlich			Temkin		
q_m (mg g ⁻¹)	k_L (L mg ⁻¹)	R^2	$q_{e(\text{exp})}$ (mg g ⁻¹)	k_F (mg g ⁻¹)	n	R^2	k_T (L mg ⁻¹)	b (J mol ⁻¹)	R^2
381.68	0.079	0.999	375.15	57.68	2.67	0.882	1.56	38.62	0.972

adsorption capacities, respectively, and k_L is the Langmuir equilibrium constant.

The Freundlich isotherm model (eqn (9)) assumes that the adsorption is multi-layered and that the adsorption on surface is non-ideal and reversible, which is not restricted to the formation of monolayer coverage.⁴⁸ It can be expressed in the linear form as:

$$\ln q_e = \ln k_F + \frac{1}{n} \ln C_e \quad (9)$$

where, k_F (L g⁻¹) is the adsorption coefficient, $1/n$ is a constant related to the surface heterogeneity, and q_e (mg g⁻¹) and C_e (mg L⁻¹) are the equilibrium adsorption capacity and the equilibrium MB concentration, respectively.

Temkin model (eqn (10)) considers the effects of some indirect adsorbate/adsorbate interactions on adsorption isotherms and suggests that because of these interactions, the heat of adsorption of all molecules in the layer decreases linearly with coverage.⁴⁸ Its linear form can be expressed as:

$$q_e = \beta \ln k_T + \beta \ln C_e \quad \left[\text{where, } \beta = \frac{RT}{b} \right] \quad (10)$$

where k_T (mL mg⁻¹) and b (J mol⁻¹) are the isotherm constant and Temkin–Pyzhev constant, respectively.

Fig. 11a–c present the plots of Langmuir, Freundlich and Temkin isotherm models, respectively. The theoretical parameters obtained from Fig. 11 are listed in Table 3. According to the obtained data, Langmuir model fits better for the MB

adsorption as its correlation coefficient value was higher and closer to the unity ($R^2 = 0.997$) compared with that from Freundlich and Temkin models (Table 3), indicating that the MB adsorption onto extracted CMFs is limited with monolayer coverage. In addition, the active sites are homogeneously distributed on the surface of the CMFs adsorbent. The maximum adsorption capacity of extracted CMFs calculated using Langmuir model was 381.68 mg g⁻¹, which is close to the experimental results (375.15 mg g⁻¹). Subsequently, it can be concluded that the MB adsorption occurred through monolayer adsorption.

Moreover, the Temkin constant b_T was about 38.62 J mol⁻¹ which indicates that the interactions between the sorbate and the used CMFs were weak (physisorption), because the typical range of bonding energies for an ion-exchange mechanism (chemisorption) is 8000–16 000 J mol⁻¹.¹⁷ This behavior is the case of the majority of cellulose-based adsorbents, where the adsorption is controlled by physical forces (with some exception of chemisorption), and the main physical forces controlling the adsorption are van der Waals forces, hydrophobicity, hydrogen bonds, polarity and steric interaction, dipole induced dipole interaction *etc.*⁵³ Consequentially, the adsorbate gets accumulated on the surface of the adsorbent by the above mentioned interactions.⁵³

Basing on the obtained adsorption results, the extracted CMFs showed a relatively high adsorption capacity. It is worth mentioning that the unique physical aspect (tendrill helical shape) of the extracted CMFs is crucial for the adsorption process. The special morphology of CMFs, mixture of linear long and helical microfibrils as well as their spaghetti-like structure, play an important role in the MB adsorption mechanism. Additionally, the abundant hydroxyl groups on the surface of CMFs is another important factor that contributing to its excellent adsorption performance because the –OH groups could facilitate dye diffusion and adsorption process *via* the formation of hydrogen bonds or van der Waals forces. For CMFs, the adsorption process involves monolayer adsorption, assuming that the adsorption takes place at specific homogeneous sites on the surface of the adsorbent and that all of the adsorption sites are energetically identical.

To demonstrate the significance of this study and the efficiency of the extracted CMFs as bioadsorbent, a comparison of adsorption capacity of the used CMFs with other adsorbent was performed. As shown in Table 4, the maximum adsorption capacity of as-extracted CMFs was found to be greater than other cellulose derivatives extracted from various bio-sourced materials as well as some cellulose based hydrogels, indicating that the extracted CMFs could be considered as

Table 4 Comparison of maximum adsorption capacities of various sorbent for MB removal

Cellulose-based adsorbent	q_{max} (mg g ⁻¹)	References
Cellulose nanocrystals from <i>Carex meyeriana</i> Kunth	217.4	49
Nanocrystalline cellulose–montmorillonite composite	183.8	50
Nanocrystalline cellulose	101.1	51
Hybrid cellulose–cotton fiber	165.4	52
Cellulose nanofibrils	122.2	16
Polyacrylamide/cellulose nanocrystals	326.0	45
Hydrogel nanocomposites of CMC	137.0	20
Regenerated porous CMC-based hydrogels beads	259.7–357.0	47
Microcrystalline cellulose extracted from oil palm	51.8	23
Extracted cellulose from <i>P. oceanica</i>	305.4	25
<i>P. oceanica</i> fibers	5.4	25
CMFs	381.6	This study



a promising low-cost adsorbent for efficient recovery of cationic dyes from aqueous solutions.

Conclusion

This study has demonstrated that new cellulose microfibrils (CMFs) with tendril helical morphology and highly hydrated gel-like behavior can be extracted from cocoa bean shells waste, using simple chemical treatments. It was found that the as-extracted CMFs gel can easily be transformed into powdered form *via* freeze drying process or in form of stable film using the casting process followed by air drying. Morphological analysis demonstrated that the as-extracted CMFs exhibit a mixture of linear long microfibrils, with an average diameter of 1 μm , and tendril helical microfibrils with an average diameter of 10 μm , forming an entangled network gel at its hydrated form. Structural analysis confirmed that CMFs have the same cellulose crystalline form and its structure is comparable to the commercial available microcrystalline cellulose. This study reveals that the as-extracted CMFs have a potential to be used as efficient adsorbent for the removal of different cationic dyes from industrial wastewater. The high swelling capacity of CMFs (190%) and its special morphology were found to be effective for enhancing its adsorption capacity. By using methylene blue as cationic dye, batch studies illustrated that the kinetics of the dye adsorption process was well fitted with the pseudo-second-order kinetic model, and the adsorption isotherms can be also described well by the Langmuir isotherm model with a maximum adsorption capacity of 381 mg g^{-1} . The extraction of CMFs, with specific morphology and properties, from cocoa bean shells waste would not only help to reduce the costs of the production of cocoa products but would also manage the disposal of this waste in an environmental friendly manner through the production of new added value materials. This study will allow us to continue valorizing this waste in order to produce new value-added materials for advanced application, such as biomaterials for food packaging.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The financial assistance of the Office Chérifien des Phosphates (OCP S.A.) in the Moroccan Kingdom toward this research is hereby acknowledged. This work was performed as part of collaboration between the INRA (UMR-IATE, Montpellier, France) and the Materials Science and Nanoengineering Department (MSN) of the Mohamed 6 Polytechnic University (UM6P) in Morocco. The authors would like to thank Dr A. Barakat and B. M. Ruesgas Ramón from INRA-UMR-IATE for their help to improve this work, and Dr Mirna Leonor Suárez Quiroz and Dr Oscar González Ríos from the Instituto Tecnológico de Veracruz (Veracruz, Mexico) for supplying the original CBS.

Notes and references

- 1 S. Kalia, A. Dufresne, B. M. Cherian, B. S. Kaith, L. Avérous, J. Njuguna and E. Nassiopoulou, *Int. J. Polym. Sci.*, 2011, **2011**, 1–35.
- 2 Y. Habibi, L. A. Lucia and O. J. Rojas, *Chem. Rev.*, 2010, **110**, 3479–3500.
- 3 O. Nechyporchuk, M. N. Belgacem and J. Bras, *Ind. Crops Prod.*, 2016, **93**, 2–25.
- 4 R. J. Moon, A. Martini, J. Nairn, J. Simonsen and J. Youngblood, *Chem. Soc. Rev.*, 2011, **40**, 3941–3994.
- 5 M. Prakash Menon, R. Selvakumar, P. Suresh kumar and S. Ramakrishna, *RSC Adv.*, 2017, **7**, 42750–42773.
- 6 S. Y. Ding, S. Zhao and Y. Zeng, *Cellulose*, 2014, **21**, 863–871.
- 7 D. Trache, M. H. Hussin, C. T. Hui Chuin, S. Sabar, M. R. N. Fazita, O. F. A. Taiwo, T. M. Hassan and M. K. M. Haafiz, *Int. J. Biol. Macromol.*, 2016, **93**, 789–804.
- 8 D. R. Smyth, *Development*, 2016, **143**, 3272–3282.
- 9 J.-S. Wang, G. Wang, X.-Q. Feng, T. Kitamura, Y.-L. Kang, S.-W. Yu and Q.-H. Qin, *Sci. Rep.*, 2013, **3**, 3102–3108.
- 10 D. C. G. Okiyama, S. L. B. Navarro and C. E. C. Rodrigues, *Trends Food Sci. Technol.*, 2017, **63**, 103–112.
- 11 FAO, FAOSTAT, <http://www.fao.org/faostat>.
- 12 I. A. Emiola, O. O. Ojebiyi and T. O. Akande, *Int. J. Poult. Sci.*, 2011, **10**, 987–990.
- 13 G. Mancini, S. Papirio, P. N. L. Lens and G. Esposito, *Environ. Eng. Sci.*, 2016, **33**, 843–850.
- 14 F. Fiorelli, J. Vieillard, R. Bargougui, N. Bouazizi, P. N. Fotsing, E. D. Woumfo, N. Brun, N. Mofaddel and F. Le Derf, *J. Colloid Interface Sci.*, 2017, **494**, 92–97.
- 15 A. W. Carpenter, C. F. De Lannoy and M. R. Wiesner, *Environ. Sci. Technol.*, 2015, **49**, 5277–5287.
- 16 C. H. Chan, C. H. Chia, S. Zakaria, M. S. Sajab and S. X. Chin, *RSC Adv.*, 2015, **5**, 18204–18212.
- 17 Y. Zhou, M. Zhang, X. Wang, Q. Huang, Y. Min, T. Ma and J. Niu, *Ind. Eng. Chem. Res.*, 2014, **53**, 5498–5506.
- 18 H. Shi, W. Li, L. Zhong and C. Xu, *Ind. Eng. Chem. Res.*, 2014, **53**, 1108–1118.
- 19 L. Jin, W. Li, Q. Xu and Q. Sun, *Cellulose*, 2015, **22**, 2443–2456.
- 20 H. Qiao, Y. Zhou, F. Yu, E. Wang, Y. Min, Q. Huang, L. Pang and T. Ma, *Chemosphere*, 2015, **141**, 297–303.
- 21 D. Trache, A. Donnot, K. Khimeche, R. Benelmir and N. Brosse, *Carbohydr. Polym.*, 2014, **104**, 223–230.
- 22 H. Y. Yu, D. Z. Zhang, F. F. Lu and J. Yao, *ACS Sustainable Chem. Eng.*, 2016, **4**, 2632–2643.
- 23 M. H. Hussin, N. A. Pohan, Z. N. Garba, M. J. Kassim, A. A. Rahim, N. Brosse, M. Yemloul, M. R. N. Fazita and M. K. M. Haafiz, *Int. J. Biol. Macromol.*, 2016, **92**, 11–19.
- 24 R. Batmaz, N. Mohammed, M. Zaman, G. Minhas, R. M. Berry and K. C. Tam, *Cellulose*, 2014, **21**, 1655–1665.
- 25 N. Ben Douissa, L. Bergaoui, S. Mansouri, R. Khiari and M. F. Mhenni, *Ind. Crops Prod.*, 2013, **45**, 106–113.
- 26 N. El Miri, K. Abdelouahdi, M. Zahouily, A. Fihri, A. Barakat, A. Solhy and M. El Achaby, *J. Appl. Polym. Sci.*, 2015, **132**, 42004, DOI: 10.1002/app.42004.



- 27 M. El Achaby, N. El Miri, A. Aboulkas, M. Zahouily, E. Bilal, A. Barakat and A. Solhy, *Int. J. Biol. Macromol.*, 2017, **96**, 340–352.
- 28 M. El Achaby, Z. Kassab, A. Aboulkas, C. Gaillard and A. Barakat, *Int. J. Biol. Macromol.*, 2018, **106**, 681–691.
- 29 C. S. Julie Chandra, N. George and S. K. Narayanankutty, *Carbohydr. Polym.*, 2016, **142**, 158–166.
- 30 S. Gea, C. T. Reynolds, N. Roohpour, B. Wirjosentono, N. Soykeabkaew, E. Bilotti and T. Peijs, *Bioresour. Technol.*, 2011, **102**, 9105–9110.
- 31 P. Lopez-Sanchez, M. Rincon, D. Wang, S. Brulhart, J. R. Stokes and M. J. Gidley, *Biomacromolecules*, 2014, **15**, 2274–2284.
- 32 Q. Wang, J. Cai, L. Zhang, M. Xu, H. Cheng, C. C. Han, S. Kuga, J. Xiao, R. Xiao, S. Renneckar, C. Han and S. Kuga, *J. Mater. Chem. A*, 2013, **1**, 6678–6686.
- 33 J. O. Zoppe, V. Ruottinen, J. Ruotsalainen, S. Rönkkö, L. S. Johansson, A. Hinkkanen, K. Järvinen and J. Seppälä, *Biomacromolecules*, 2014, **15**, 1534–1542.
- 34 C. J. Chirayil, J. Joy, L. Mathew, M. Mozetic, J. Koetz and S. Thomas, *Ind. Crops Prod.*, 2014, **59**, 27–34.
- 35 M. Jonoobi, J. Harun, P. M. Tahir, A. Shakeri, S. Saifulazry and M. D. Makinejad, *Mater. Lett.*, 2011, **65**, 1098–1100.
- 36 R. M. Sheltami, I. Abdullah, I. Ahmad, A. Dufresne and H. Kargarzadeh, *Carbohydr. Polym.*, 2012, **88**, 772–779.
- 37 W. P. Flauzino Neto, H. A. Silvério, N. O. Dantas and D. Pasquini, *Ind. Crops Prod.*, 2013, **42**, 480–488.
- 38 A. Dmytryk, A. Saeid and K. Chojnacka, *Sci. World J.*, 2014, **2014**, 356328, DOI: 10.1155/2014/356328.
- 39 Y. C. Ching and T. S. Ng, *BioResources*, 2014, **9**, 6373–6385.
- 40 X. Y. Tan, S. B. Abd Hamid and C. W. Lai, *Biomass Bioenergy*, 2015, **81**, 584–591.
- 41 N. Kasyapi, V. Chaudhary and A. K. Bhowmick, *Carbohydr. Polym.*, 2013, **92**, 1116–1123.
- 42 S. M. Luz, J. Del Tio, G. J. M. Rocha, A. R. Gonçalves and A. P. Del'Arco, *Composites, Part A*, 2008, **39**, 1362–1369.
- 43 W. H. Chen, Y. J. Tu and H. K. Sheen, *Appl. Energy*, 2011, **88**, 2726–2734.
- 44 C. Kannan, N. Buvanewari and T. Palvannan, *Desalination*, 2009, **249**, 1132–1138.
- 45 C. Zhou, Q. Wu, T. Lei and I. I. Negulescu, *Chem. Eng. J.*, 2014, **251**, 17–24.
- 46 N. Fayoud, S. Tahiri, S. Alami Younssi, A. Albizane, D. Gallart-Mateu, M. L. Cervera and M. de la Guardia, *Desalin. Water Treat.*, 2016, **57**, 16611–16625.
- 47 T. Benhalima, H. Ferfera-Harrar and D. Lerari, *Int. J. Biol. Macromol.*, 2017, **105**, 1025–1042.
- 48 M. Baghdadi, A. Jafari and A. Pardakhti, *RSC Adv.*, 2016, **6**, 61423–61433.
- 49 X. Yang, H. Liu, F. Han, S. Jiang, L. Liu and Z. Xia, *Carbohydr. Polym.*, 2017, **175**, 464–472.
- 50 S. P. Santoso, L. Laysandra, J. N. Putro, J. Lie, F. E. Soetaredjo, S. Ismadji, A. Ayucitra and Y. H. Ju, *J. Mol. Liq.*, 2017, **233**, 29–37.
- 51 X. Y. He, K. B. Male, P. N. Nesterenko, D. Brabazon, B. Paull and J. H. T. Luong, *ACS Appl. Mater. Interfaces*, 2013, **5**, 8796–8804.
- 52 J. Tao, J. Xiong, C. Jiao, D. Zhang, H. Lin and Y. Chen, *Carbohydr. Polym.*, 2017, **166**, 271–280.
- 53 S. Hokkanen, A. Bhatnagar and M. Sillanpää, *Water Res.*, 2016, **91**, 156–173.

