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Isolation of whiskers from natural sources and their dispersed in a non-aqueous medium

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Abstract

Whiskers have been used as a nanomaterial dispersed in polymer matrices to modify the microscopic and macroscopic properties of the polymer. These nanomaterials can be isolated from cellulose, one of the most abundant natural renewable sources of biodegradable polymer. In this study, whiskers were isolated from sugarcane bagasse and corn cob straw fibers. Initially, the cellulose fiber was treated through an alkaline/oxidative process followed by acid hydrolysis. Dimethylformamide and dimethyl sulfoxide were used to replace the aqueous medium for the dispersion of the whiskers. For the solvent exchange, dimethylformamide or dimethyl sulfoxide was added to the aqueous dispersion and the water was then removed by fractional distillation. FTIR, TGA, XRD, TEM, Zeta and DLS techniques were used to evaluate the efficiency of the isolation process as well as the morphology and dimensions of the whiskers. The dimensions of the whiskers are comparable with values reported in the literature, maintaining the uniformity and homogeneity in both aqueous and non-aqueous solvents.

Keywords: *whiskers, cellulose bleaching, agricultural waste.*

1. Introduction

Agricultural production has expanded in recent years and the lignocellulosic byproducts generated have great potential for exploitation as new raw materials. Agricultural byproducts, such as sugarcane bagasse (SCB) and corn cob straw (CCS), are residues produced on a large scale which, after treatment, can be added to a polymeric matrix in order to alter the physicochemical characteristics of the nanocomposites. Lignocellulosic materials consist mainly of cellulose, hemicellulose and lignin, the contents of which vary depending on the raw material and the physiological characteristics of the plant. SCB is comprised of 42 to 46% cellulose, 21 to 28% lignin and 27 to 29% of hemicellulose^[1-3]. On the other hand, CCS contains 42 to 44% cellulose, 22 to 28% lignin and 27 to 28% hemicellulose^[4-6]. Minor amounts of other polysaccharides (found in the cell walls) and minerals may also be present in these raw materials.

Cellulose is an abundant biopolymer which is biodegradable and renewable and it can be obtained in an environmentally sustainable manner. It has a microfibrillar structure composed of β -1,4-linked anhydro-D-glucose ($C_6H_{10}O_5$) units, and the repeat segment is a dimer of glucose, known as cellobiose. The spatial conformation and arrangement associated with intra- and intermolecular hydrogen bonds lead to the rigid characteristics of cellulose^[7,8]. Six interconvertible cellulose polymorphs, namely I, II, III₁, III₂, IV₁ and IV₂, have been identified, each with its particular set of network parameter characteristics of unit cells, type I being referred to as native cellulose^[7]. In order to isolate the cellulose from lignocellulosic

plant materials, the non-cellulosic components (hemicellulose, lignin, etc.) need to be removed from the fiber through a process known as bleaching. Different treatments have been applied in the bleaching of natural fibers, such as the use of an alkali, followed by an oxidizing/alkaline environment with the use of an oxidant based on chlorinated compounds^[9,10]. Studies have been conducted using oxidants which generate very low levels of waste, such as hydrogen peroxide, thereby reducing the environmental impact^[1,11]. Rosa et al.^[12] has isolated cellulose from rice husk by cellulose extraction in two-step chlorine-free bleaching processes. Other processes are performed without the alkali treatment step, using systems such as nitric/acetic acid^[11] and peracetic acid^[13] and mixtures of hydrogen peroxide with manganese(II) sulfate and zinc oxide or manganese(II) sulfate and titanium dioxide^[14]. In the bleaching process the control variables, such as temperature, time and concentration of reagents, play an extremely important role.

After bleaching, the cellulose isolated by mechanical, chemical or enzymatic treatment can result in microcrystalline cellulose, microfibrillated cellulose or whiskers. The microfibrils which make up the cellulose fiber consist of amorphous and crystalline domains and their acid hydrolysis allows the isolation of the crystalline domains, denoted as whiskers or cellulose nanocrystals, due to the dimensional characteristics of the material. The aspect ratio of the whiskers is dependent on the variables involved in the process used to isolate the whiskers, such as the raw material, temperature, reaction

time, reagent concentration and reagent/raw materials ratio. Habibi et al.^[17] and Moon et al.^[15] reported that whiskers have an acicular shape and length/diameter ratio of 10 to 30, with lengths of 35 to 500 nm and a diameter of 3 to 15 nm, depending on the raw material and treatment used to isolate the whiskers.

The large surface area and crystallinity of the whiskers make these promising materials for incorporation into polymeric matrixes, where they increase the mechanical strength and modify the gas barrier properties^[7,15-17]. On the other hand, the whiskers can be dispersed homogeneously in polymers which are soluble in aqueous solution due to their hydrophilic surface^[18]. However, this restricts their use in the matrices of hydrophobic polymers due to difficulties related to their dispersion in these systems. Studies have been directed toward obtaining an adequate dispersion of whiskers in hydrophobic matrices, using approaches such as the addition of anionic surfactants or *tert*-butanol to the dispersing medium^[19], the use of a cationic surfactant^[20], the use of partially hydrolyzed poly(vinyl alcohol)^[21], surface functionalization of the whiskers^[22-24], the use of lyophilization and redispersion in a non-aqueous solvent. Of these proposed methods, the redispersion of the whiskers in non-aqueous solvents has proved to be a promising technique which allows homogeneous dispersion in hydrophobic polymers, it being necessary to assess the process conditions in relation to the polymer matrix. Yu et al.^[25] isolated whiskers in an aqueous medium and then replaced the water with acetone and subsequently replaced the acetone with chloroform in several successive centrifugation steps and finally the whiskers were incorporated into the poly(3-hydroxybutyrate-co-3-hydroxyvalerate) matrix. Pracella et al.^[26] used an analogous procedure for the solvent exchange of the whiskers suspension and subsequent incorporation of the whiskers into poly(lactic acid).

The whiskers yield from the isolation process, as well as the whiskers suspension in non-aqueous solvents and subsequent incorporation into the hydrophobic polymer matrix, are dependent on the characteristics of the raw material, the bleaching process and the procedure used for the isolation of the whiskers, in addition to the characteristics of the solvent used in the suspension and the polymer matrix employed to produce the nanocomposite.

In this context, the aim of this study was to evaluate the processes used for the isolation of whiskers from sugarcane bagasse and corn cob straw, to promote the exchange of the whiskers dispersing medium and keep the whiskers homogeneously dispersed in non-aqueous solvents to facilitate the obtainment of a nanocomposite polymeric matrix containing whiskers.

2. Materials and Methods

2.1 Materials

The sugarcane bagasse (SCB) was provided by regional ethanol producers (São Paulo, SP Brazil) and the corn cob straw (CCS) by regional organic producers (Pato Branco, PR Brazil). Poly(lactic acid) (Code 3251D) was purchased from NatureWorks, Cargill (Minnetonka, MN, USA). Sulfuric acid, sodium hydroxide and hydrogen peroxide were purchased

from Lafan Química Fina, and dimethylformamide (DMF) and dimethyl sulfoxide (DMSO) were supplied by Nuclear (Brazil). Cellulose was purchased from Merck and lignin obtained by isolation from the raw materials (SCB and CCS) according to regulation TAPPI T222 om-88^[24]. All of the chemical reagents were used without further purification.

2.2 Bleaching of fibers from raw materials

The SCB and CCS raw materials were washed with distilled water, dried and milled and then passed through a 30-mesh sieve. In the next step, the fibers were dried in an oven with air circulation at 105 °C for 4 h prior to the bleaching process. For the bleaching process, three different procedures were applied: (i) treatment with NaOH aqueous solution; (ii) treatment with H₂O₂; and (iii) treatment with Ca(ClO)₂ aqueous solution.

- (i) Treatment with NaOH aqueous solution - The fibers were maintained in a 5% aqueous solution of NaOH (1:20 w/v fiber/solution) at 85 °C for 90 min, under mechanical stirring. In the next step, the dispersion was filtered and the fibers washed until neutral pH and dried at 105 °C for 4 h. The fibers obtained through this treatment are denoted as SCB/NaOH or CCS/NaOH, according to the raw material.
- (ii) Treatment with H₂O₂ - The fibers subjected to treatment (i) were then maintained in an aqueous solution of 5% NaOH and 11% H₂O₂ (1:20 w/v) at 55 °C for 90 min and the dispersion was filtered. The fibers were then washed with distilled water until neutral pH and dried at 105 °C in an oven with air circulation for 4 h. These treated fibers were denoted by SCB/H₂O₂ or CCS/H₂O₂, according to the raw material.
- (iii) Treatment with Ca(ClO)₂ aqueous solution - Fibers subjected to treatment (i) were maintained in an aqueous solution of 2.5% Ca(ClO)₂ and 5% NaOH (w/v), at 45 °C for 240 min and the dispersion was filtered. The fibers were then washed until neutral pH and dried at 105 °C for 4 h. These treated fibers were denoted by SCB/Ca(ClO)₂ or CCS/Ca(ClO)₂, according to the starting raw material.

2.3 Characterization of raw materials and bleached fibers

To evaluate the effect of bleaching on the composition and crystallinity of the lignocellulosic materials, SCB and CCS samples before and after the bleaching processes were characterized by Fourier transformed infrared spectroscopy (FTIR), thermogravimetry analysis (TG) and X-ray diffraction (XRD). The cellulose, lignin and hemicellulose contents of the untreated SCB and CCS were determined in accordance with the respective recommendations of the Technical Association of the Pulp and Paper Industry (TAPPI standards).

2.3.1 Content of cellulose, lignin and hemicellulose in SCB and CCS raw materials

The lignin, cellulose and hemicellulose of the raw materials SCB and CCS were determined according to the TAPPI Standard T222 om-88^[27], the ashes according to the norm TAPPI T211 OM 93^[28] and the extractive was

determined according to the standard TAPPI T 204 om-88^[29]. The holocellulose content (cellulose and hemicellulose) and the pulp were determined according to standard TAPPI T13 m-54^[30], allowing the determination of the hemicellulose content. All measurements are expressed on a dry basis.

2.3.2 Fourier transform infrared spectroscopy (FTIR)

The FTIR spectra for the cellulose, lignin, SCB and CCS samples before and after the bleaching process were obtained on a Shimadzu IR Prestige system, performing 20 scans, with a resolution of 4 cm⁻¹, in the range of 4000 to 600 cm⁻¹ at room temperature. The samples were prepared in the form of compressed KBr disks.

2.3.3 Thermogravimetric analysis (TGA)

Thermal degradation measurements were taken on a Shimadzu TGA-50 thermogravimetric analyzer, from room temperature to 600 °C at a heating rate of 10 °C min⁻¹. The flow rate of the nitrogen purge gas was 50 mL min⁻¹.

2.3.4 X-ray diffraction (XRD)

X-ray patterns of the specimens were obtained on a Philips X'Pert diffractometer (Netherlands), with Cu (K_α) radiation (λ = 1.5418 Å), operating at room temperature, 30 mA and 40 kV. The scanned region of 2 to 40° (2θ) and a pitch of 0.05° s⁻¹ were applied to evaluate the crystallinity index of the SCB and CCS samples, before and after the bleaching processes. The samples were milled to yield material between 120 and 450 mesh sizes. The crystallinity index (C_I) was determined using Equation 1, by comparing the corresponding areas of the crystalline and amorphous fractions, obtained by deconvolution of the diffractogram, where A_a is the area of the amorphous phase and A_t the total area of

$$C_I = \left(1 - \frac{A_a}{A_t} \right) \times 100 \quad (1)$$

2.4 Extraction of whiskers

To extract the whiskers bleached fibers obtained from CCS and SCB were selected, through treatments (ii) with H₂O₂ and (iii) with Ca(ClO)₂, as described in section 2.2. Acid hydrolysis was carried out at 55 °C for 75 min with 60 wt % H₂SO₄ (1:20 w/v) under mechanical stirring. The fiber suspension was diluted with water (five times the volume of the reaction mixture) at room temperature to stop the reaction and after centrifugation the supernatant was removed. Distilled water was then added to the sediment, which was suspended under mechanical stirring, and the suspension was centrifuged until the appearance of the cloud point. This suspension of cellulose whiskers was collected and treated using dialysis bags in a cellulose acetate membrane with a cut off of 10,000 g mol⁻¹ at neutral pH and stored at 5 °C.

2.5 Replacement of dispersive medium of whiskers

The dispersive medium containing the cellulose whiskers was changed from water to DMF or DMSO, in order to solubilize the hydrophobic polymer matrix. The DMF or

DMSO was added to the aqueous suspension of cellulose whiskers and the water eliminated through vacuum distillation at 60 °C, since the boiling point of a non-aqueous solvent is greater than the boiling point of water.

In the exchange of the aqueous dispersion medium containing the whiskers the volumes of solvent before and after fractional distillation were gravimetrically monitored.

2.6 Characterization of whiskers

2.6.1 Whiskers in aqueous and non-aqueous media

In order to estimate the yield of the isolation process of the whiskers and efficiency in the solvent exchange the quantity of whiskers dispersed in the aqueous and non-aqueous medium was determined using the gravimetric procedure. A sample of 2 mL of the whiskers suspension in aqueous medium was placed in Petri dishes and kept in an oven with air circulation at 80 °C to evaporate the solvent. The sample was then maintained at 105 °C until constant weight, and the quantity of whiskers was calculated and expressed as weight per volume of suspension. The yield of the hydrolysis reaction was estimated from the weight of the cellulose content of the bleached fibers. An analogous procedure was used to determine the efficiency of the solvent exchange from water to DMF or DMSO, maintaining the suspension at 130 °C until complete evaporation of the solvent.

2.6.2 Zeta potential analysis (ξ) and dynamic light scattering (DLS)

The surface charges were measured by zeta potential analysis (ξ) from an aliquot of the neutral aqueous suspension of the whiskers with the concentration equalized at 0.005% (w/w). In order to evaluate the stability of the suspensions they were kept under refrigeration at 7 °C and analyzed at 6-month intervals, after sonication (UltraCleaner 1600A) for 30 min and centrifugation under a force of 1370 x g.

Dynamic light scattering was carried out using a Zetasizer Nano ZS system (Malvern, UK), by detecting back-scattered laser-light (θ = 173°) and comparing the coherence of scattering patterns as a function of time, with 0.002 g mL⁻¹ of whiskers in aqueous, dimethylformamide or dimethyl sulfoxide suspensions, at 25 °C. Measurements of the particles in suspension were carried out in triplicate, and each measurement was composed of 15 runs every 10 s. Analysis based on the diffusion rate of the particles in the fluid was conducted to calculate the autocorrelation function and deduce the particle size information.

2.6.3 Transmission electron microscopy (TEM)

Transmission electron microscopy analysis of the whiskers was carried out using a JEOL-1011 TEM electron microscope operating at 80 kV. One drop (0.002 g mL⁻¹) of the suspension of whiskers was diluted in 2 mL of isopropyl alcohol and deposited on a grid coated with copper film (FORMVAR®) and after being almost dried it was stained by adding one drop (~ 4 μL) of 3% uranyl acetate (w/v). The excess liquid was removed and this was followed by drying at room temperature.

3. Results and Discussions

3.1 Characterization of raw materials

Table 1 shows the composition of cellulose, lignin, hemicellulose, ash and extractive for the sugarcane bagasse (SCB) and corn cob straw (CCS). As expected, for both raw materials cellulose predominates in relation to the other components. The percentages of each component are consistent with values reported in the literature^[1-6]. SCB fibers have a higher content of lignin and lower content of hemicellulose than CCS fibers, which affects the physiological characteristics of these fibers.

3.2 Evaluation of the effect of bleaching process

Figure 1 shows the FTIR spectra, in the range of 1400-1850 cm^{-1} , for cellulose and lignin obtained by isolation from the raw materials straw corn cob straw (CCS) and sugar cane bagasse (SCB) before and after the bleaching treatments. In the range of 1400 to 1850 cm^{-1} the cellulose shows absorption bands at 1645 and 1430 cm^{-1} related to the symmetrical angular deformations of the OH and CH_2 (constituents of the chemical structure of the cellulose repeating unit), respectively^[31]. Due to the phenolic groups present in the lignin, the infrared spectrum shows absorption bands at 1600 and 1510 cm^{-1} , related to axial deformations of the aromatic rings^[32]. Besides the characteristic absorption bands of cellulose and lignin, the untreated SCB and CCS present absorption bands at 1730 cm^{-1} , characteristic of the axial stretching of the carbonyl and associated with hemicellulose^[10,14]. The infrared spectra of the untreated SCB and CCS indicate the presence of cellulose, lignin and

hemicellulose, in agreement with the results of the quantitative analysis shown in Table 1. After the alkaline treatment (i) of the raw materials (SCB and CCS) there was a reduction in the intensity of the absorption bands as expected, which is associated with the functional groups present in the polymer chain of the lignin and hemicellulose. Although the NaOH treatment promoted the removal of lignin, the presence of shoulders at 1510 and 1600 cm^{-1} indicates that complete removal of lignin did not occur. In order to reduce the lignin content in the samples, we used the treatments (ii) or (iii), after which the infrared spectra suggest greater efficiency of the bleaching processes, with a reduction in the intensity of the absorption bands related to lignin.

Figure 2 shows the TG curves obtained for the pure cellulose, the lignin obtained by isolation from the raw materials, the raw material and the fibers after the bleaching processes for (a) SCB and (b) CCS. The TG curves of the raw materials exhibited similar behavior with weight loss at 100 °C associated with the removal of water molecules and in temperature range of 250 °C to 400 °C related to the thermal degradation of the constituents of the sample, while at temperatures above 400 °C only the solid waste residue remained. The residue of the lignin isolated from SCB and CCS is higher than the residue of cellulose and other studied samples. At 450 °C there was 23% of residue for the raw material samples and 63% and 71% of residue for lignin from SCB and CCS, respectively. The percentages of residue for the fibers after treatment with hydrogen peroxide were 19% and 13% for SCB and CCS, respectively. These values of residue percentage is related to higher quantity of lignin in SCB than in CCS samples after bleaching

Table 1. Composition mass percentage of sugarcane bagasse (SCB) and corn cob straw (CCS).

Raw material	Cellulose	Lignin	Hemicellulose	Ash	Extractive
SCB	44.6 ± 1.2	24.0 ± 1.3	26.0 ± 2.1	2.8 ± 0.3	2.5 ± 0.3
CCS	47.3 ± 2.1	14.4 ± 0.9	34.8 ± 2.3	1.5 ± 0.1	2.0 ± 0.2

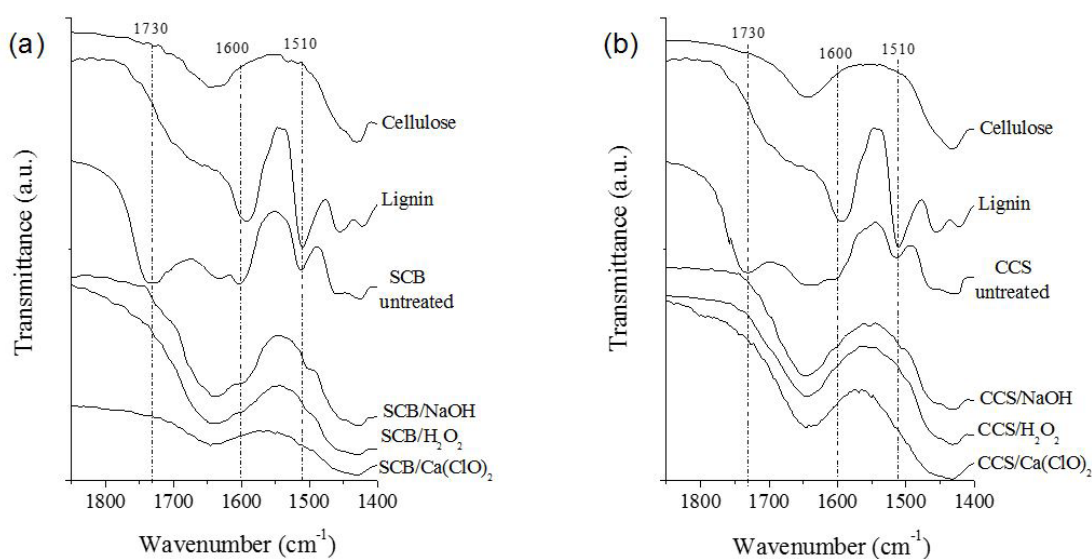


Figure 1. Infrared spectra for pure cellulose, lignin and the samples after the bleaching process (indicated in the figure) using (a) SCB and (b) CCS as raw materials.

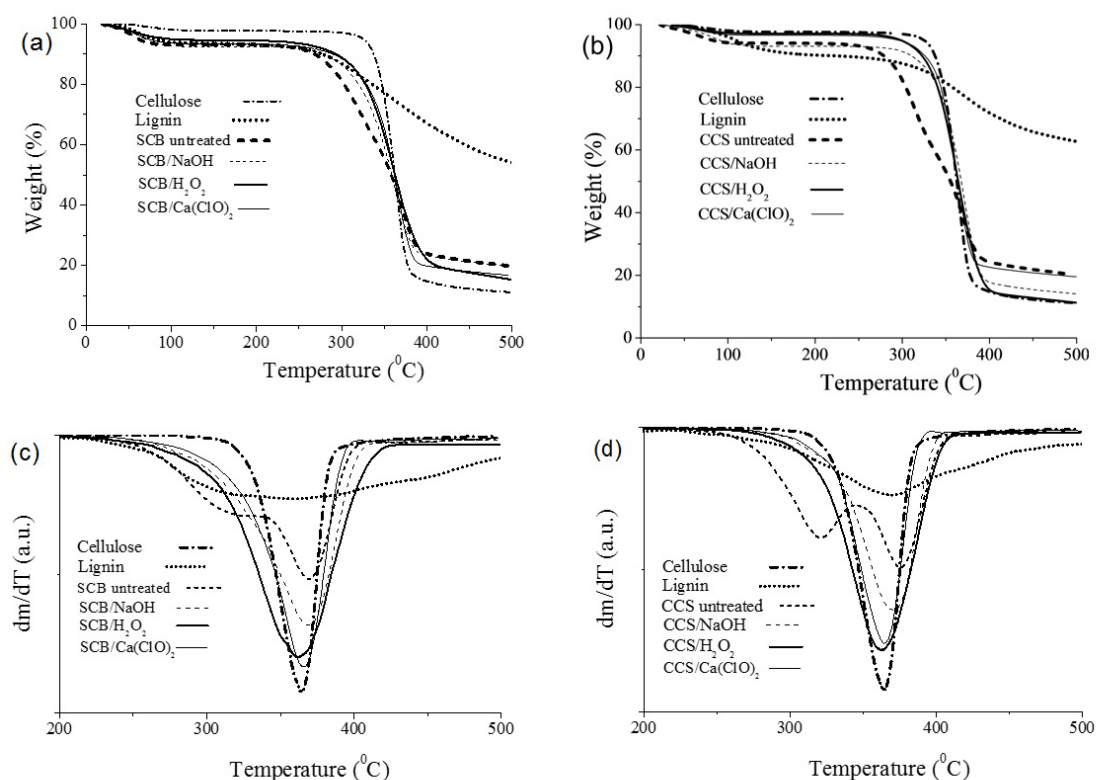


Figure 2. TG curves and DTG curves for (a, c) SCB and (b, d) CCS related to pure cellulose, lignin and the samples after the bleaching processes (indicated in the figure).

processes, as can be seen by relative intensity at 1510 cm^{-1} and 1600 cm^{-1} in the FTIR spectra, since the lignin was not totally removed. The amount of residue observed after the calcium hypochlorite process can be included the presence of calcium compound.

The DTG curves enable the visualization of the different stages of thermal degradation of the components in the samples, as shown in Figure 2c, d for samples derived from SCB and CCS, respectively.

Pure cellulose presents a single-stage thermal degradation event with the maximum degradation rate at $365\text{ }^{\circ}\text{C}$, as described in the literature^[10,33]. Besides the degradation stage associated with cellulose, the raw materials presented other degradation stages related to lignocellulosic material, such as lignin and hemicellulose^[33]. After treatment (i) with NaOH, the behavior of the DTG curve with a single stage of thermal degradation tends to be similar to that of pure cellulose, indicating the removal of the lignocellulosic components. This tendency was noted for the sequence of treatments with hydrogen peroxide and calcium hypochlorite, indicating greater efficiency for these bleaching processes. This behavior was observed for CCS and SCB, in agreement with the residue levels observed in the thermal analysis and infrared spectroscopy.

Figure 3a, b show the diffractograms for the SCB and CCS raw materials after treatments (ii) and (iii), respectively, with peaks at 14.7° , 16.6° , 22.5° and 34.7° , which relate to the families of crystalline planes: (101), (10 $\bar{1}$), (002) and (004),

respectively, characteristic of the cellulose I^[34]. The peaks associated with the crystal structure suggest that the alkali treatment is not aggressive in relation to changes in the cellulose crystal lattice. According to Borysiak and Doczekalska^[35], the crystalline structure of cellulose can change from cellulose I to cellulose II in an aqueous 15% NaOH (w/v) solution.

The crystallinity index was 30% for both of the raw materials studied. After the bleaching process the fibers obtained from SCB showed values for the crystallinity index of 70% and 68%, after treatment with hydrogen peroxide and calcium hypochlorite, respectively. For the fibers obtained from CCS the crystallinity index was 67% after both treatments. For comparison, the crystallinity index determined by Rosa et al.^[12] for rice husk cellulose and microcrystalline cellulose were 67% and 79%, respectively. An increase in the C_1 values indicates a greater quantity of crystalline cellulose in the samples compared to amorphous material, which is related to the effectiveness in the removal of hemicelluloses and lignin, in agreement with the results obtained in the infrared spectroscopy and thermogravimetric analysis.

3.3 Characterization of whiskers

Table 2 shows the yields for the whiskers isolated from the bleached raw materials by acid hydrolysis. A value of 60% (w/w) was obtained for the bleached cellulose, suggesting that the process used for the isolation of the nanocrystals

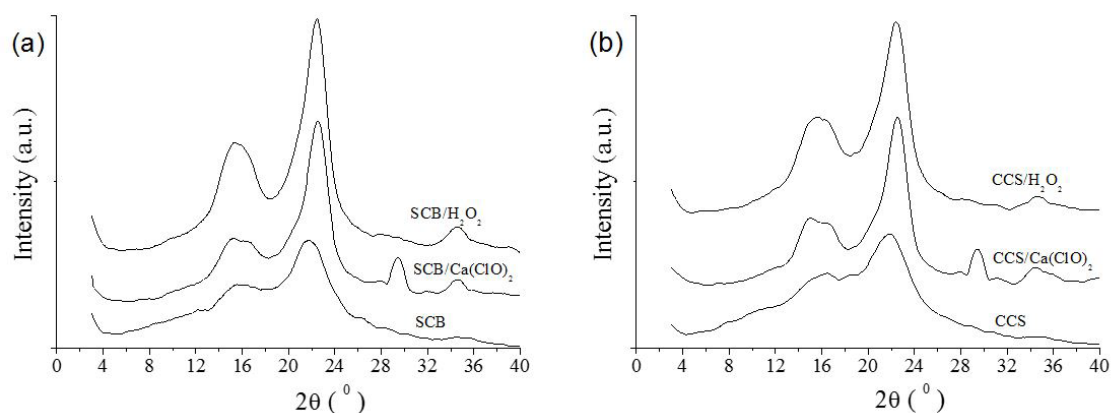


Figure 3. XRD spectra for (a) SCB and (b) CCS raw materials and after treatments with $\text{H}_2\text{O}_2/\text{NaOH}$ or $\text{Ca}(\text{ClO})_2/\text{NaOH}$ (as indicated in the figure).

had a good efficiency, regardless of the bleaching process and raw material used. The literature reports different yields for the whiskers according to the hydrolysis conditions applied for the bleaching of the fibers. Correa^[36] studied the hydrolysis of curauá fibers with 60% H_2SO_4 (v/v) at 45 °C for 75 min and obtained a whiskers yield of 70% while Taipina^[37] using the same kind of fiber obtained a yield of 65%. On the other hand, Oksman and Bondeson et al.^[38] carried out hydrolysis with H_2SO_4 (63.5% w/w), using a 10:1 (v/v) solution/raw material ratio and a reaction time of 2 h and obtained whiskers with a yield of 30%. Dong et al.^[39] reported yields of between 35% and 48% when varying the hydrolysis conditions (64% H_2SO_4 w/w) applying temperatures of 26 °C to 65 °C and reaction times of between 0.25 h and 18 h. Mathew et al.^[40] studied wood waste from the production of ethanol, after bleaching and acid hydrolysis and recorded 46% yield for the whiskers. In this regard, we also conducted preliminary experiments under more severe conditions, with 65% H_2SO_4 (w/w), a temperature of 55 °C and 75 min of reaction and with 60% H_2SO_4 (w/w) at 55 °C for 300 min, and in both cases the yield of whiskers obtained was 30%. However, the suspensions had a dark color, indicating the carbonization of the whiskers.

The stability of aqueous suspensions of whiskers was analyzed via the zeta potential (ξ). The values obtained at 6-month intervals are shown in Table 3. The potential of around -64 mV in neutral aqueous medium (pH between 6.0 and 7.0) is indicative of negative charges on the surface of the whiskers, due to the grafting of sulfate groups in the cellulose hydrolysis process^[41,42]. These groups enable the formation of a colloidal suspension and its stability is due to the electrostatic repulsion of these surface charges^[40]. According to Greenwood^[43], potentials above 30 mV are sufficient for obtaining a suspension with good stability.

The morphology and size of the whiskers were evaluated by transmission electron microscopy (TEM) and dynamic light scattering (DLS). Figure 4a, b show the TEM micrographs for the whiskers dispersed in water. For both raw materials (SCB and CCS) the whiskers were rod shaped (acicular), which is characteristic of structural rigidity. For SCB the average length (L) was 230 ± 35 nm and the average diameter (D) determined by TEM was 16 ± 3 nm, with an

Table 2. Yields obtained for the whiskers after the hydrolysis processes.

Raw material	Bleaching process	Yield (%)
SCB	SCB/ H_2O	60
	SCB/ $\text{Ca}(\text{ClO})_2$	55
CCS	CCS/ H_2O_2	65
	CCS/ $\text{Ca}(\text{ClO})_2$	60

Table 3. Zeta potential (mV) of whiskers.

Storage time (months)	SCB	CCS
0	-64.3 ± 6.9	-65.2 ± 7.8
6	-64.1 ± 1.5	-64.1 ± 0.7
12	-63.2 ± 1.2	-65.6 ± 4.5

aspect ratio (L/D) of 15. For the CCS the values determined by TEM were $L = 288 \pm 62$ nm and $D = 16 \pm 4$ nm, with an L/D ratio of 18. The acicular shape and dimensions are similar to those described in the literature for whiskers, with lengths of around 35 to 500 nm and diameters of between 3 and 15 nm^[7,15]. More specifically, for sugarcane bagasse, Teixeira et al.^[1] obtained average values for the length and diameter of 255 ± 55 nm and 8 ± 3 nm, respectively, while Mandal et al.^[10] reported a shorter length (170 nm) and larger diameter (35 nm).

Lima et al.^[44] and Braun et al.^[45] considered the Broersma relationships using the rotation and translational diffusion coefficients to obtain the dimensions of the whiskers. According to these authors, the spheroidal form factor contains more than one adjustable parameter, allowing the simultaneous determination of the diameter and length, which are related to the size of the whiskers. Thus, using the DLS technique the values obtained for the dimensions of the whiskers were 232 ± 5 nm and 242 ± 15 nm for SCB and CCS, respectively, which are in agreement with the values determined by TEM.

On the other hand, for the whiskers obtained from SCB under more severe hydrolysis conditions, with 65% H_2SO_4 (w/w) at 55 °C for 75 min, the size obtained by DLS was 167 ± 2 nm, which indicates that the hydrolysis conditions

can alter^[39] the dimensions and surface characteristics of the whiskers, as well as their yield. Thus, it is possible to control the aspect ratio (L/D) and surface area, which are important characteristics of nanocrystals, making them of interest for different applications.

The incorporation of whiskers, homogeneously distributed in matrices of water-insoluble polymers presents a great challenge. Approaches commonly applied to overcome this difficulty include the use of surfactants, the surface functionalization of the whiskers and/or the lyophilization and solvent dispersion of hydrophobic polymer matrices. However, a large quantity of surfactants is required to maintain the stability of the suspension due to the high specific surface area of the nanocrystals. Furthermore, as observed by Bondeson and Oksman^[21] the surfactants may cause degradation of the polymer matrix. On the other hand, Peterson et al.^[19] reports that when lyophilizing a suspension of whiskers in aqueous medium and after their dispersion in chloroform, undesirable flakes are formed in the solvent. Viet et al.^[46] studied a suspension of lyophilized whiskers in aqueous medium and their subsequent dispersion in dimethylformamide and dimethyl sulfoxide, while maintaining the aggregates in the suspension. Kamal et al.^[47] used the spray dried, freeze drying and spray freeze drying of whiskers

while maintaining agglomerates in suspension in molten polymer (PLA) at a minimum size of 1 μm .

In this study, we replaced the aqueous dispersion medium with dimethylformamide or dimethyl sulfoxide, as described in the experimental section, and evaluated the morphology and dispersion characteristics of the whiskers in these non-aqueous media.

Figure 5a, b show the TEM micrographs of the whiskers obtained from SCB and CCS as raw materials, respectively, dispersed in dimethylformamide. A homogenous dispersion of the whiskers was maintained, and the lengths (L) were 240 ± 61 nm and 245 ± 80 nm and the diameters (D) were 16 ± 5 nm and 18 ± 6 nm for SCB and the CCS, respectively. These dimensions are in agreement with the values obtained through DLS of $240 (\pm 26)$ for SCB and $250 (\pm 8)$ for CCS in dimethylformamide and $224 (\pm 5)$ for SCB and $244 (\pm 8)$ for CCS in dimethyl sulfoxide.

The solvent exchange procedure was effective and appropriate to maintain the dispersion of the whiskers. As discussed above, the method is simple and inexpensive, indicating that the solvent mixture is not azeotropic and since there was a considerable difference between the boiling temperatures of the solvents efficient water removal could be carried out by fractional distillation.

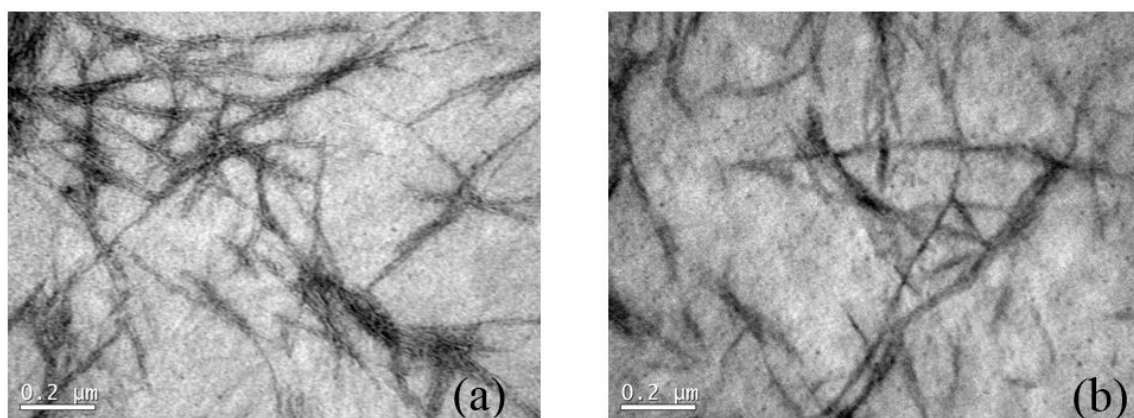


Figure 4. TEM micrographs for whiskers dispersed in aqueous medium obtained from (a) SCB and (b) CCS as raw materials.

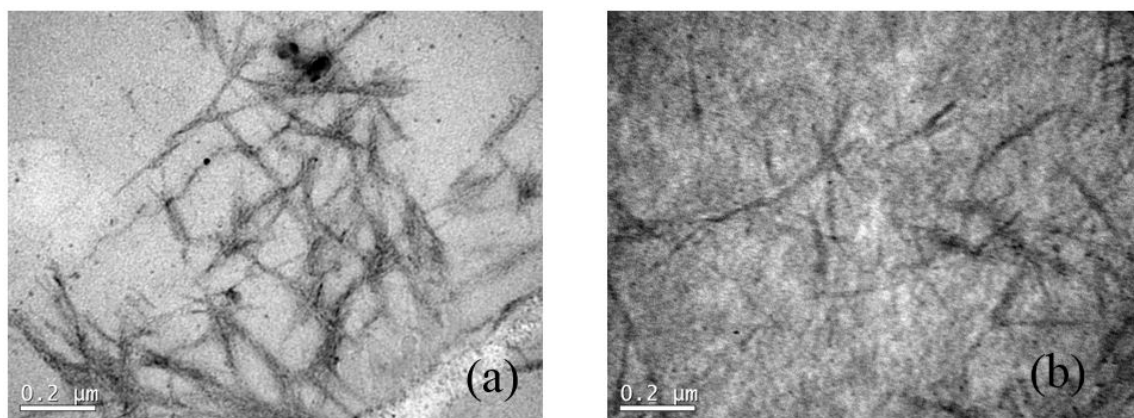


Figure 5. TEM micrographs for whiskers obtained from (a) SCB and (b) CCS as raw materials dispersed in DMF.

4. Conclusions

The procedures used for the isolation of whiskers and the oxidant/alkali bleaching processes followed by acid hydrolysis were found to be effective, providing good yields and dimensional characteristics as well as dispersity in a solvent medium. The isolated whiskers had nanometric dimensions and aspect ratio (L/D) values comparable with those reported in the literature. When the aqueous medium was replaced with a non-aqueous medium using a simple low cost technique, without the need for mechanical work, the whiskers remained dispersed and the morphological characteristics of the whiskers were maintained. The dispersion of the whiskers in a non-aqueous solvent facilitates future studies on the development of techniques to obtain nanocomposite whiskers homogeneously dispersed in matrices of polymers by dissolving them in a common solvent followed by solvent evaporation.

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