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Cellulose nanofiber nanocomposites with aligned silver nanoparticles

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ABSTRACT

Celluloses have attracted much attention as sustainable and abundant materials. Herein, we focus on nanocomposites based on the oxidation-treated nano-sized fibrillated celluloses, namely TOCNs. The silver nanoparticles (AgNPs) were prepared in TOCN aqueous dispersion. Generally, the AgNPs are quickly agglomerated after preparation. For the inhibition of the agglomeration of AgNPs, it is required that AgNPs were prepared under the chelation of TOCN, followed by reduction therein. Therefore, AgNPs possessed the nano-scaled radii and aligned along the TOCN from the atomic force microscopic measurements. The thermal stabilities and mechanical properties were increased. The anisotropic thermal conductivities originated from the orientation of TOCN in nanocomposites were observed. The loading of the large amounts of AgNP fillers led to the drastic increase of the thermal and electrical conductivities. The conductive paths of heat and electron were formed by the contact of AgNP with each other. We functionalized the TOCN papers through the loading of AgNPs and the obtained nanocomposites sheets served as conductors.

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Cellulose nanofiber; TEMPO-mediated oxidation; silver nanoparticle; mechanical property; thermal conductivity

Introduction

In our daily life, petroleum-resource-based synthetic polymers, such as polyolefins, polystyrene and polyvinylchloride have widely spread. However, for these decades, the environmental problems such as global warming and exhaustible resources have been arisen. Cellulose is the most abundant, sustainable and environmental-friendly natural product in the world and attracts much attentions as an alternative material to synthetic polymers in the field of not only agriculture but also polymer science [1]. In addition, it has been reported that isotropic sheet composed of cellulose nanofibers (CNFs) possesses much higher elastic modulus than 10 GPa [2–5]. Therefore, CNFs are well known as remarkable materials with high mechanical properties, light weight, low linear thermal expansion coefficient, biocompatibility and high specific surface [6–11]. CNFs are accepted as fillers in nanocomposites [12, 13], substrates of electrical devices [14–18],...
wrapping papers with high performance [19, 20] and scaffolds for tissue engineering [21, 22]. CNFs are prepared in the two different typical methods; one is the mechanical method such as grinding processes [23, 24] and the other is the chemical method such as 2,2,6,6-tetramethylpiperidine-1-oxyl (TEMPO)-mediated oxidation [25–27]. In particular, the TEMPO treatment selectively and densely introduces carboxyl groups to the 6-positions of a half of glucopyranose rings at the surface of cellulose nanofibers. The nanofibers of TEMPO-oxidized CNFs (TOCNs) show high aspect ratios with radii of about 3–5 nm and lengths of several micro meters. These low radius and high aspect ratio lead to the increasing of entanglements and the decreasing defects of CNFs. Therefore, they improve mechanical properties relative to micro-fibrillated CNF.

In organic–inorganic nanocomposites, where organic matrices are reinforced and functionalized by the nanosized inorganic fillers, the higher performance and functionality have been achieved; for example, mechanical properties [28–30], gas permeabilities [31, 32], higher thermal and electrical conductivities [33–36] and the control of refractive indexes [37–40]. Various inorganic nanofillers for the nanocomposites have been reported. Silver nanoparticles (AgNPs) exhibit catalytic activities [41–43], surface-enhanced Raman scattering activities [44–46], high electrical and thermal conductivities [47, 48] and antimicrobial activities [49, 50]. In the case of nanocomposites, the dispersibility of nanofillers in polymer matrix is a key factor for the achievements of their high performance because of rapid and easy agglomeration of nanofillers in dispersion. For the well-dispersion of AgNPs, it has been reported on the usage of surfactant and in situ preparation methods. Among them, the well-dispersed AgNPs are often prepared in the in situ synthesis through the reductive reaction of metal ions. Moreover, the in situ synthesis by chelation to metal ions leads to the inhibition from random agglomeration as well as the 1D and 2D arrangements of AgNPs along polymer chains and sheets [51–53]. These arrangements induce the percolation of the contacts of nanofillers, the anisotropy of the functional properties and the drastic increase of electrical conductivity with the low loading of nanofillers.

Herein, we focused on TOCN/AgNP nanocomposites and investigated their mechanical and functional properties. The AgNPs were prepared from Ag cation in TOCN aqueous dispersion. In the TOCN dispersion, the Ag ions were chelated with the carboxylate groups at the 6-position of cellulose, and then they are reduced to AgNPs on the TOCN [54, 55]. AgNPs are agglomerated rapidly without any dispersion stabilizers, whereas the chelation effects of TOCN made the high dispersity of AgNPs possible. We observed well-dispersed and aligned AgNPs with atomic force microscope (AFM). In addition, we evaluated the reinforcement effects of AgNPs fillers and anisotropic functional properties through investigation on their mechanical properties as well as functional properties such as Young’s modulus, thermal conductivities and electrical conductivities.

Experimental

Materials

Kenaf bast fibers (Hibiscus cannabinus, Indonesia, 2011) with 2 mm of the average length were provided from Toyota Boshoku co. (Aichi, Japan). All the chemical reagents were purchased with chemical company.

Purification of kenaf bast fibers

Kenaf bast fibers, 20 g, were degreased via Soxhlet extraction using a 1:2 (v/v) mixture of ethanol/toluene for 20 h at 130 °C. Kenaf bast fibers then were dried for 2 days under ambient atmosphere before a half of the degreased biomass was suspended into 1 L of distilled water. For removing the lignin of the sample with Wise method, sodium chlorite, 6.7 g, and acetic acid, 1.2 mL, were directly added to the biomass suspension. The biomass suspension mixture was heated at 80 °C for 1 h. The addition of sodium chlorite, 6.7 g, and acetic acid, 1.2 mL, was repeated four times in 1 h intervals. The mixture was filtered and suspended into distilled water five times. In order to remove the hemicellulose, the biomass samples were suspended in 0.5 L of 4 wt% KOH aq. and stirred for 2 h at 80 °C. Following that, the refined biomass was washed through 5 times filtration and resuspension. Three times of these alkali treatments provided the refined pulp.

TEMPO oxidation treatment

The above refined pulp 10 g, TEMPO 156 mg and NaBr 1.0 g were added into 1 L distilled water. 25 mL sodium hypochlorite solution was added slowly keeping below pH 10 by the automatic titrator. When the constant increase of pH of the solution with addition of 0.5 M NaOH aqueous solution was achieved, the oxidation was stopped. Following that, the oxidized biomass was washed through the five times filtration and redispersion.

The semi-wetted biomass samples were suspended in 1 L acetate buffer solution with pH 4.8. The sodium chlorite was added with the ratio of
1.14 g for 1 g biomass and stirred for 48 h. Following that, the oxidized biomass were washed through the five times filtration and redispersion. These two reactions led to the oxidation of cellulose.

The oxidized biomass pulps dispersed into 700 mL water were fibrillated by the blending mixer with 24,000 rpm for 10 min. The TOCNs were obtained.

Preparation of silver nanoparticles in TOCN suspension

For 215 g of the 0.25 wt% TOCN aqueous dispersion, the silver nitrate aqueous solution was added, which amounts were equal to the 0.3–30 vol% silver against TOCN after reduction. The suspensions were stirred for 1 day under darkness and the ion exchanges were proceeded. 3.2 wt% sodium borohydride aqueous solution with the equivalent to the silver nitrate was added and stirred for 1 day. The reduction of silver ions led to the formation of AgNPs in the TOCN dispersion.

Preparation of TOCN and TOCN/AgNP nanocomposite sheets

After defoaming of TOCN or TOCN/AgNP suspension, the suspensions containing 0.65 g of TOCN were filtered and the sheets were obtained. The sheets were dried under vacuum at 40°C. All the procedures were shown in Figure S1 in the Supporting Information.

Characterization

Infrared absorption (FT-IR) spectroscopy was carried out on Spectrum GX FT-IR System I-KS (Perkin Elmer co. ltd.). The samples were dispersed into KBr and were dried for 24 h at 40°C.

In order to estimate the amount of carboxyl groups in TOCN, the electrical conductivity and pH curves were measured by acid–base titration with an automatic titrator (AUT-501, TOA-DKK Co. Ltd.) and an electrical conductometer (CM-60V, TOA-DKK Co. Ltd.).

The radii of TOCN and AgNPs were measured by atomic force microscopy Nano Navi II E-sweep (SII nanotechnology Co. Ltd.) with cantilever SI-DF20 (SII nanotechnology Co. Ltd., spring modulus: 15 N/m, radii of tip: 10 nm, scanning area: 2 μm × 2 μm).

X-ray diffraction measurements were carried out with RINT2000 (Rigaku Co. Ltd., 40 kV, 20 mA, X-ray source: CuKα (λ = 1.5418 Å) scanning rate: 1/min, sampling steps: 0.02°, scanning region: 2θ = 5°–55°). The crystallite sizes were estimated from Scherrer’s equation (1)

\[ D = \frac{\lambda}{\beta \cos \theta} \]  

\[ \beta^2 = B^2 - b^2 \]  

where \( D \) is the crystallite size, \( \lambda \) is the wavelength of X-ray beam, \( \theta \) is the Bragg angle, \( B \) is the integral width of samples and \( b \) is that of silicon. In addition, the crystallinity of cellulose was calculated through the equation (3)
where $X_c$ is the crystallinity, $A_c$ is the area of the peaks originated from crystalline, and $A_a$ is that of amorphous. For the measurements of the transmittance of the cellulose nanocomposite sheets (thickness: 50 μm), we measured the transmit spectra from 200 to 1000 nm with HITACHI U-2000. In UV–vis absorption measurement, we measured the aqueous dispersion including 0.25 wt% TOCN and AgNP.

For the investigation of thermal properties of TOCN/AgNP nanocomposites, we measured melting points of AgNP in nanocomposites under nitrogen with RIGAKU DSC8230. The scan rate was 5 °C/min. From the Gibbs–Thomson equation (4), the size of AgNP were estimated.

$$ T = T_{\text{bulk}} - \frac{4\sigma M T_{\text{bulk}}}{\Delta H_m r \rho} $$

where $T$ and $T_{\text{bulk}}$ (1223 K) are melting points of silver nanoparticles in nanocomposites and bulk silver, $\sigma$ (1.02 J/m²) is the solid-liquid interface energy, $M$ (107.9 g/mol) is molecular weight, $\Delta H_m$ (11.3 kJ/mol) is the latent heat of fusion, $r$ is the radii of silver nanoparticles and $\rho$ (10.5 g/cm³) is the density.

For thermogravimetric analysis of TOCN and TOCN/AgNP nanocomposites, we carried with RIGAKU Thermo plus EVO II TG8121 under nitrogen. The heating rate was 10 °C/min. The starting temperature was 150 °C for exclusion of effects of water.

The length and width of the trimmed samples for tensile tests were 40 and 5 mm, and they were dried at 40 °C for 24 h before tensile tests. We carried out with Autograph AGS-1kND (SHIMAZU Co. Ltd.). The strain-stress curves were obtained with 20 mm as initial lengths and 1 mm/min of the tensile speed. The density of TOCN was calculated by the floatation method. The initial inclination of the curves was regarded as Young’s modulus.

For the measurement of the volume electric resistance of TOCN sheets and TOCN/AgNP nanocomposites, we measured with Electrometer/High Resistance Meter B2980A and Resistivity Cell 16008B (Keysight technologies Co. Ltd.) in the two ring electrode method. The diameter of the electrode was 26 mm and the applied voltage was 10 V.

For the measurement of the thermal diffusivity by the periodic heating and infrared radiation thermometer method, we carried with Thermowave Analyzer TA3 (Bethel Co. Ltd.). The thermal diffusivity for the in-plane and thickness directions was measured; in-plane means the parallel direction to the sheet surface and thickness means perpendicular one.
We prepared a refined pulp from Kenaf bast fiber through degreasing, Wise treatment and alkaline treatment for removing grease, lignin and hemicellulose, respectively. The refined pulps were fibrillated to nanofibers by TEMPO-catalyst oxidation and fibrillation with Blender. The radii of TOCN were less than 5 nm from the AFM topological images in Figure 1. The amount of the carboxyl group of TOCN were 1.28 mmol/g from the measurements of conductometric titration and the degree of oxidation per anhydroglucose unit of cellulose (DO) was 0.21 [25, 26, 56, 57]. (Supplementary Figure S2) The silver nitrate aqueous solution was added into the obtained 0.25 wt% TOCN dispersion and stirred until ion exchange from sodium ions to silver ions. Following that, the equal amounts of sodium borohydride solution was added and silver ions were reduced to AgNPs. The reduced AgNPs possessed radii less than 5 nm and were aligned along the TOCN as shown in Figure 1. These alignments suggested that Ag ions would be chelated to the carboxyl groups of TOCN and reduced on the TOCN selectively rather than randomly [58]. In the case of the TOCN/20 vol%AgNP composites, the free AgNPs were observed as shown in Supplementary Figure S3. This means that there were too much non-chelated silver ions to form the nuclei of nanoparticle along only the TOCN. In the FT-IR spectra, all the TOCN and TOCN/AgNP nanocomposites showed larger carbonyl stretching bands at around 1600 cm\(^{-1}\) than Kenaf microfibers, while there was no changed in the vibration bands of glucopyranose rings (1060 cm\(^{-1}\)). The latter indicated that polysaccharide structures of cellulose were sustained even after the TEMPO-oxidation and the reduction of Ag ions, as shown in Supplementary Figure S4.

The TOCN/AgNP nanocomposite sheets were prepared by filtration and dried over at 12 h under vacuum. Optical photographs and transmittance spectra of TOCN/AgNP composites and absorption spectra of the TOCN/AgNP aqueous dispersion were shown in Figure 2. TOCN sheets without AgNPs showed high transparency. With the increase of AgNP loading, the sheets turned brownish because of the surface plasmon resonance (SPR) of AgNPs [59]. In the absorption spectra of the

![Figure 3.](image-url)
TOCN/AgNP suspensions, the absorption bands originated from SPR of AgNPs were observed at around 400 nm. These absorption bands were attributed to the AgNPs with less radii than 10 nm. This result corresponded to the size of AgNPs in the AFM images. In addition, this means that the AgNPs were highly dispersed in the TOCN aqueous dispersion without agglomeration [55, 60–62].

The X-ray diffraction measurements were performed for the TOCN and TOCN/AgNP nanocomposites. In their X-ray diffraction profiles in Figure 3a, all the samples showed the peaks originated from 110, 110, and 200 reflections of the cellulose Iβ crystallites. The crystallinity of TOCN in composites was calculated in the area method, as shown in Figure 3b. The effects of the reductive reaction for the preparation of AgNPs were observed and their structure of cellulose Iβ were maintained. The 111 and 200 reflections of AgNPs were appeared at 2θ = 38° and 45° respectively with the loading of AgNPs in TOCN/AgNP nanocomposites. There is no diffraction peak assigned as silver oxide, such as (111) plane (2θ = 33°) [63]. In addition, the crystallite sizes of AgNPs in TOCN/AgNP nanocomposites were estimated from the integral width and the Bragg angle using Scherrer’s equation. As shown in Figure 3c, with the increase of loading amounts of AgNPs, the crystallite size of AgNPs was increased. In particular, AgNPs size in less than 5 vol% AgNPs composites were below 10 nm. These results also supported the nanosized silver particles in the composites.

The thermogravimetric curves of TOCN and TOCN/AgNP composites were shown in Figure 4a. Above around 220°C, the thermal weight loss drastically increased in all the samples. These thermal decomposition were attributed to the decomposition of TOCN. In this study, temperature at 5 wt% loss of TOCN/AgNP nanocomposites was defined as a decomposition temperature $T_{d5}$. $T_{d5}$ were increased with the amounts of AgNP in the composites, as shown in Figure 4b. The loading of AgNP led to the higher thermal stability of TOCN matrix in their nanocomposites. The amounts of residues after thermal decomposition were increased as the loading of AgNPs increased. We calculated the amounts of AgNPs from the residues as shown in Supplementary Table S2.

For investigation of thermal behavior of AgNPs in nanocomposites, we measured their differential scanning calorimetry (DSC). It is known that the metallic nanoparticles possess lower melting points than bulk metals due to their higher specific surface areas [64]. In our nanocomposites, the melting point of only the 2 vol% AgNP nanocomposite was observed at 168°C, which were lower than that (962°C) of bulk silver. The radius of AgNP, which was estimated from the Gibbs-Thomson equation [65, 66], was 5.8 nm. From all the various measurements such as AFM images, UV–vis absorption spectra, X-ray diffraction profiles and DSC thermograms, the AgNPs in the TOCN/AgNP nanocomposites were sustained as nanoparticles even through reduction reaction.

Figure 5a shows the results of tensile tests of Kenaf microfiber, TOCN sheets and TOCN/AgNP nanocomposites. Compared with the Kenaf microfiber sheet, downsizing of fiber diameter of TOCN led to the increase of the Young’s modulus. The reason was that the mechanical defects of the nanofibers were decreased relative to microfibers. For the TOCN/AgNP composites, the low content ratios of the AgNPs provided the effective increase of mechanical modulus, whereas the modulus of the composites with larger content of AgNPs were decreased due to the agglomeration of AgNPs. In Figure 5b, the relationship between the AgNP loading and Young’s modulus was represented as a solid line.
and Young’s modulus predicted by parallel model equation (6) as an upper bound was shown as a dashed line.

\[ E_c = E_f V_f + E_m (1 - V_f) \]  

(6)

where \( E_c \), \( E_f \) and \( E_m \) are the modulus of the composites, fillers and matrix, respectively, and \( V_f \) is the volume fraction of fillers. The modulus of bulk silver (83 GPa) [67] and TOCN were applied as their modulus \( E_f \) and \( E_m \), respectively. At the lower loading of AgNPs, the larger Young’s modulus was exhibited than the estimated value. Moreover, the increase of amount of AgNPs led to higher densities of the composites, which suggested that the void would not emerge. These larger Young’s modulus would be induced by the effect that the nano-sized AgNP possessed the higher modulus than the bulk silver [68, 69]. In contrast, the relation between the tensile strength or elongation at break of those composites and the AgNP fraction was not observed clearly. The reason is that their mechanical parameters received large effects of the reinforcement as well as mechanical defects in the composites. In only the case of the composites with 30 vol% AgNP, it is obviously revealed that the excess AgNP additive led to the decrease of tensile strength or elongation at break.

It is well-known that silver possesses the highest thermal conductivity in single metals [67]. Therefore, we investigated the thermal conductivities of TOCN/AgNP nanocomposites and declared the loading effects of AgNPs. From the DSC measurements and thermal diffusion measurements, their specific heat \( (C_p) \) and the thermal diffusivities to the in-plane \( (\alpha_{in-plane}) \) and thickness \( (\alpha_{thick}) \) direction were evaluated. The thermal conductivities \( (\lambda_{in-plane} \) and \( \lambda_{thick} \)) were calculated from their parameters, as shown in Figure 6. In the TOCN sheets without AgNPs, the anisotropic thermal conductivity in the in-plane and thickness directions were emerged. In

![Figure 5](image)

*Figure 5.* (a) Stress-strain curves of kenaf microfiber, TOCN and TOCN/AgNP nanocomposites with different silver contents. (b) Relationship between observed Young’s modulus and silver content of TOCN/AgNP nanocomposites.

![Figure 6](image)

*Figure 6.* Thermal conductivity (circles: observed, lines: calculated) of TOCN/AgNP nanocomposites with different silver contents in (a) in-plane and (b) thickness direction.
the TOCN sheets, the TOCNs were oriented in the direction parallel to the sheet plane because of the vacuum filtration of TOCN aqueous dispersion. The AgNP loading increased the thermal conductivities to the in-plane direction ($\lambda_{\text{in-plane}}$). From the theory on the thermal conductivities of composite materials, namely the Maxwell equation (7), we calculated the theoretical thermal conductivities of the TOCN/AgNP nanocomposites.

$$\lambda_c = \frac{\lambda_m 2\lambda_m + \lambda_f + 2V_f(\lambda_f + \lambda_m)}{2\lambda_m + \lambda_f - V_f(\lambda_f + \lambda_m)}$$  

(7)

where $\lambda_c$, $\lambda_f$, and $\lambda_m$ are the thermal conductivities of the composite, filler and matrix, respectively, and $V_f$ is the volume fraction of filler. The reported value of the thermal conductivity of AgNP filler ($\lambda_f$) is 430 W/m$\cdot$K [70] and the experimental value of TOCN sheet is employed as the thermal conductivity of the matrix. In the Maxwell theory, it is assumed that the fillers were well-dispersed in the matrix and there was no interaction between the fillers [71]. In Figure 6, the theoretical and experimental thermal conductivities of the TOCN/AgNP nanocomposites were represented. At the low content ratios of the AgNPs, their thermal conductivities were increased in agreement with the Maxwell theory. These results indicated that the AgNPs were distributed homogeneously in the composites without any mutual interaction. In contrast, the nanocomposites with high content ratios of the AgNPs provided the higher thermal conductivities than the calculated values. This suggests that larger amounts of the AgNPs caused the contact of each other in the TOCN matrix and the formation of the thermal conductive paths. In the case of the thermal conductivities in the thickness direction, the nanocomposites including the high amounts of the AgNPs also demonstrated the higher thermal conductivities, whereas the values were much lower than those in the in-plane direction. The difference between the conductivities in the thickness and in-plane directions were attributed to the in-plane alignment of the TOCN and the AgNP in the nanocomposites.

The electrical resistance of the TOCN/AgNP nanocomposites was shown in Figure 7. In the region of the lower content ratios of the AgNP than 15 vol%, the high resistance of the nanocomposites kept constants regardless of the amount of the AgNP fillers. These high resistances were originated from electric insulation of the TOCN matrix. On the other hand, the drastically lower resistances were achieved in the nanocomposites with higher content ratios of the AgNP than 15 vol%. It is suspected that the AgNP fillers were contacted with one another and the conductive paths were formed by the percolation. The volume of the AgNP where the percolation in electrical resistance corresponded to those of thermal conductive percolation. Even in the larger amounts of AgNPs, the decrease of the electrical resistivity were incredibly slight. The reason was that the conductive paths have been formed and the additional AgNP fillers had no large effect on their electrical resistance. The gained electrical resistance of the TOCN/AgNP nanocomposites were compared with those of general semiconductors [72–74].

**Conclusions**

We demonstrated the preparation of the TOCN/AgNP nanocomposites and achieved the functionalization of TOCN nanopapers. The AgNPs were formed through the chelation of Ag ions to carbonyl groups of TOCN, then reduction in aqueous dispersion. The sizes of the AgNPs were less than 10 nm and the nanoparticles were well dispersed in nanocomposites. The AgNPs in the nanocomposites were aligned along the TOCN. The loading of AgNPs increased thermal stabilities of nanocomposites and their mechanical properties. In addition, their thermal conductivities emerged anisotropy between in-plane and thickness direction. With high loading of AgNPs, the higher thermal and electrical conductivities were accessed by their percolation.

**Disclosure statement**

The authors declare no competing financial interest.

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