An Efficient and Sustainable Approach to Prepare Carboxylated Cellulose Nanocrystals for Rubber Reinforcement Featuring Dual Crosslinking Networks

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Abstract

The surface functionalization of CNCs and the construction of strong interfacial adhesion between CNCs and rubber matrix are effective way to achieve high performance rubber/CNCs nanocomposites. Herein, carboxylation of sulphated cellulose nanocrystals (CNC-OSO$_3$H) was conducted in aqueous medium by using citric acid as modifier. Large amount of carboxyl groups was successfully grafted on the surface of CNC-OSO$_3$H, which endows the carboxylated CNC-OSO$_3$H (abbreviate as CNC-CA) with higher chemical reactivity and thermal stability. Subsequently, carboxylated styrene butadiene rubber (XSBR)/CNC-CA nanocomposites with dual crosslinking design were prepared by using polyethylene glycol diglycidyl ether (PEGDE) as the crosslinking agent and CNC-CA as the reinforcing fillers. FTIR investigation found that in the obtained nanocomposites, the carboxyl groups on CNC-CA and XSBR formed hydrogen bonds (physical crosslinking) with each other, and the carboxyl groups formed covalent bond with the epoxy group on PEGDE simultaneously. The coexistence of physical and chemical crosslinking improved the interface compatibility between CNC-CA and XSBR matrix, accelerated the homogenous dispersion of CNC-CA and realized the crosslinking of the matrix itself. As expected, XSBR/CNC-CA nanocomposites with dual crosslinking network showed remarkable enhancement in tensile strength (up to 500%), modulus (up to 151%), work of fracture (up to 348%). This work provides both a facile and green approach to obtain carboxylated CNCs and a convenient method for the preparation of high-performance rubber nanocomposites with multiple interactions.

Introduction

By loading reinforcing fillers, the mechanical properties of composites can be improved largely compared to the neat matrix (Bitinis et al. 2011; Chen et al. 2008; Ramorino et al. 2009; Rooj et al. 2010). With the development of nanoscience, the application of various nano-reinforcing fillers in the field of composite materials has attracted increasing attention (Ding et al. 2019; Iwatake et al. 2008; Potts et al. 2012; Sengupta et al. 2011; Tang et al. 2012; Wu et al. 2018). Cellulose is the most abundant biopolymer resource on the planet and has some nanosized and hierarchical structure. As one kind of nanocellulose, cellulose nanocrystals (CNCs) are considered as a promising “green” filler for composite materials owing to the various advantages, such as large specific surface area, high tensile strength, high aspect ratio, high Young’s modulus, plenty of surface hydroxyl groups, biocompatibility and biodegradability (Kargarzadeh et al. 2018; Klemm et al. 2018; Thomas et al. 2018; Dufresne et al. 2018; Habibi et al. 2010). In 2011, the preparation of CNCs was industrialized via sulfuric acid hydrolysis, and this further promotes the application of CNCs in the field of composite materials. The CNCs prepared by this method have sulfonate groups on the surface and is abbreviated as CNC-OSO$_3$H. Due to the merits of fast reaction speed, simple process, high yield and good dispersion of CNC-OSO$_3$H in aqueous solution, sulfuric acid hydrolysis is the only industrialized preparation method at present. CNCs were first adopted to reinforce rubber in 1995 by Favier et. al, since then plenty of reports about rubber/CNCs composites emerge (Eslami et al. 2020; Flauzino Neto et al. 2016; Jiang and Gu 2020; Kargarzadeh et al. 2017; Nunes 2017; Parambath Kanoth et al. 2015; Peng et al. 2018; Peng et al. 2019; Tian et al. 2017; Favier et al. 1995).
Rachel Blanchard et. al reported that the incorporation of CNC-OSO$_3$H effectively improved the crosslinking density, tensile strength and modulus of the rubber latex for the application in dipped goods (Blanchard et al. 2020). However, the poor heat resistance of CNC-OSO$_3$H caused by sulfonate group and relatively low chemical reactivity remain a problem which cannot be ignored (Jiang et al. 2010; Wang et al. 2007; Roman et al. 2004). Compared with CNC-OSO$_3$H, CNCs prepared by citric acid have higher thermal stability, and the surface carboxyl groups endow CNCs higher reactivity. Yet, due to the weak acidity of citric acid, the yield of CNCs prepared by citric acid alone is quite low (Ji et al. 2019). A large amount of citric acid is needed when mixed with other inorganic acids to prepare CNCs, and it is not only the preparation process is time-consuming but also difficult to realize the recycling of citric acid (Yu et al. 2016).

Carboxylated styrene butadiene rubber (XSBR), a copolymer of styrene, butadiene and a small amount of acrylic acid, is widely used in paper and board coating industry. Cao et. al utilized CNC-OSO$_3$H to improve the mechanical properties of XSBR matrix by XSBR latex/CNC-OSO$_3$H co-sedimentation and two-roll mill blending (Cao et al. 2013). Even though the addition of CNC-OSO$_3$H improved the mechanical properties of XSBR/CNC-OSO$_3$H nanocomposites to a certain extent, there are still some unsolved problems: the poor thermal stability of CNC-OSO$_3$H, the agglomeration of CNC-OSO$_3$H in matrix and inadequate interaction between fillers-rubber (Chen et al. 2015). In order to improve the compatibility and interfacial strength between CNCs and matrix, a lot of efforts have been made. Cao et al. used maleic anhydride grafted tunicate cellulose nanocrystals (t-CNs) to reinforce the epoxidized natural rubber by the construction of dual cross-linked network. The interfacial chemical reaction between carboxyl groups on modified t-CNs and epoxy groups on ENR chains formed covalent crosslinking network, and the hydrogen bonds between hydroxyl groups on t-CNs and epoxy groups on ENR chains served as physical crosslinking network. It is worth mentioning that, even though the covalent bond formed between modified t-CNs and ENR chains effectively improve the interface compatibility and adhesion between filler and matrix, the uncrosslinking matrix still leads to the limited improvement of the mechanical properties of the composites (Cao et al. 2018). Tao et al. designed a novel double-network composite system of reducing end-modified CNCs (mCNC) enhanced thermoplastic elastomer with click reaction and bulk crosslinking. Though, the combination of filler-matrix, filler-filler and matrix-matrix interaction effectively realized the mechanical enhancement of the composites, the modification process of CNCs and the preparation of composites were quite tedious (Tao et al. 2019). The above works demonstrate that the construction of dual crosslinking network is an effective way to improve the properties of rubber composites. As the reactive functional carboxyl group of XSBR provides the possibility for the formation of hydrogen bond and covalent bond between XSBR and reinforcing fillers. The key point is to activate and desulfonate the surface of CNC-OSO$_3$H, subsequently design a strong XSBR-CNCs and XSBR-XSBR interaction network.

In this work, CNC-OSO$_3$H were firstly carboxylated by reaction with citric acid in aqueous phase under a mild condition, the reaction solvent and ethanol used for centrifugal washing could be recycled. Then XSBR/CNC-CA nanocomposites with dual crosslinking network were constructed by using polyethylene
glycol diglycidyl ether (PEGDE) as crosslink agent and 2-methylimidazole (2-mIm) as catalyst. FTIR investigation confirmed that the hydrogen bonds (physical crosslinking) formed among the carboxyl groups on CNC-CA and XSBR, the covalent bonds (chemical crosslinking) formed between the carboxyl groups and the epoxy group of PEGDE. Benefiting from the dual crosslinking structure, the interface compatibility between CNC-CA and XSBR matrix is improved, the homogenous dispersion of CNC-CA is accelerated and the crosslinking of the matrix itself is realized. Compared with XSBR/CNC-CA nanocomposites without PEGDE crosslinking, the tensile strength, modulus and work of fracture of XSBR/CNC-CA nanocomposites with PEGDE crosslinking are significantly improved.

**Experimental**

**Materials**

XSBR latex with 50 wt % solid content was obtained from Shandong Xing Yu gloves Co., Ltd. (Gaomi, China). Cotton linter pulps (CLP) were provided by Silver Hawk Co. Ltd. (Gaomi, China). Citric acid anhydrous (CA, 99.5%) was purchased from Sinopharm Chemical Reagent Co., Ltd. (Shanghai, China). Polyethylene glycol diglycidyl ether (PEGDE, 99%) was purchased from Macklin reagent company (Shanghai, China). 2-methylimidazole (2-mIm, 98%) was purchased from Aladdin Industrial Corporation (Shanghai, China). Sodium hydroxide, sulfuric acid, toluene and dimethylformamide are general commercial products. All regents are analytical grade.

**Preparation of sulfonic cellulose nanocrystals (CNC-OSO$_3$H)**

CNC-OSO$_3$H was extracted from CLP via sulfuric acid hydrolysis as described in our previous work (Wang et al. 2020). In brief, raw CLP was first immersed in 4 wt% NaOH solution for 24 h at room temperature. Then, the obtained slurry was thoroughly washed and dried in a vacuum oven. After that, the pretreated CLP was hydrolyzed in 64 wt% sulfuric acid at 50°C with continuous vigorous stirring, a large amount of water was used to dilute the mixture and terminate the acid hydrolysis process. Finally, the suspension was centrifuged repeatedly and washed until the pH of the centrifugal solution was close to 3, followed by dialyzing against deionized water until neutral pH was achieved. The resultant CNC-OSO$_3$H suspension was concentrated to ca. 2 wt% by rotary evaporation at 40°C, and carefully collected for the following experiment.

**Preparation of citric acid grafting CNCs (CNC-CA)**

First, 20 ml of CNC-OSO$_3$H aqueous solution (20 mg/ml) was added to a flask and ultrasonic treated for 10 min, then 10 g anhydrous CA was added to the flask and ultrasonic treated for another 10 min. Subsequently, the flask was assembled to a distillation device, and the oil bath temperature was set at 120°C. The water in the mixture was gradually evaporated with oil temperature increases. After the oil bath temperature reached 120°C, the CNC-OSO$_3$H surface esterification was performed for 15 h. After the reaction, the obtained CNC-CA was purified from unreacted CA by 4 times dispersion-centrifugation with a large excess of ethanol at room temperature. Then CNC-CA was dispersed in deionized water and dialysis
against deionized water for one week. The CA dissolved in the washing ethanol were recycled by distillation of the solvent in a rotavapor. The content of carboxyl group in CNC-CA was conducted by potentiometric titration and the value is 0.679 mmol/g, the test method is in accordance with our previous work (Zhou et al. 2018).

**Preparation of XSBR/CNC-CA nanocomposites**

The carboxyl content of XSBR was 2.59 mmol/g, the specific test methods are presented in the supporting information. XSBR/CNC-CA nanocomposites films were prepared by a solution casting method. For the preparation of XSBR/CNC-CA nanocomposites with dual crosslinking networks, XSBR (3 g) and the desired amount of CNC-CA, PEGDE, 2-methylimidazole were mixed together, ultrasonically treated for half an hour and then stirred for one hour. The obtained mixture was cast on a glass petri dish and dried by solvent evaporation at 60°C temperature for 12 h, the samples were then further treated at 110°C for six hours to ensure complete crosslinking. The dual crosslinked XSBR/CNC-CA nanocomposites with 1%, 3%, 5%, 7%, 9% CNC-CA were prepared, the resulting sample is marked as X/mC-D (m represents the content of CNC-CA in nanocomposites, X, C, and D are the abbreviations of XSBR, CNC-CA and dual crosslinking network, respectively). The mole of PEGDE is equal to the carboxyl content of CNC-CA in XSBR/CNC-CA nanocomposites. The mass ratio of 2-methylimidazole to PEGDE was 1:1. For comparison, the XSBR/CNC-CA nanocomposites without PEGDE and 2-methylimidazole were also prepared according to the above method, and the resulting sample is marked as X/mC-H (H is the abbreviation of hydrogen bonded crosslinking network).

**Characterizations**

FTIR spectra were conducted on a Bruker tensor 70 spectrometer equipped with a DTG detector under the transmission mode in the wavenumber range 4000 to 400 cm$^{-1}$ or under the ATR mode in the wavenumber range 4000 to 600 cm$^{-1}$ with the resolution of 2 cm$^{-1}$ and 32 scans.

The wide-angle X-ray diffraction (WAXD) patterns were recorded on a Rigaku mart Lab diffractometer with Cu K$\alpha$ radiation ($\lambda = 0.154$ nm) in the range $2\theta = 5$–40° with a scan rate of 5°/min.

Thermogravimetric analysis (TGA) measurements were carried out using a thermogravimetric analyzer (NETZSCH TG 209F1 Libra) under nitrogen gas flow at a rate of 100 mL/min. Samples were heated from 30 to 600°C at 10°C/min.

Dynamic light scattering (DLS) was used to characterize the average particle size of CNC-CA and CNC-OSO$_3$H in suspension. DLS was performed on a Malvern Nano ZS90 light scattering instrument.

The transmission electron microscopy (TEM) was conducted to observe the morphological differences of CNC-CA and CNC-OSO$_3$H. The aqueous solution of CNC-CA and CNC-OSO$_3$H was treated by ultrasonic to obtain a uniform suspension. A small drop of 0.1 wt% solution was deposited on a carbon-coated copper grid. TEM observations were carried out on a JEM-2100 electron microscope operated at an acceleration voltage of 300 kV.
SEM images were recorded with a Quanta FEG 250 scanning electron microscope. The cryogenically fractured surfaces of neat XSBR and XSBR/CNC-CA nanocomposites specimens were sputtered with gold before observations.

The tensile test was performed by Instron 5943 with a tensile rate of 50 mm/min. Before testing, neat XSBR and XSBR/CNC-CA nanocomposite films were cut into rectangle with dimensions of 30×5×0.2 mm³. At least 5 specimens were measured for each sample.

The dynamic mechanical analyses (DMA) of neat XSBR and X/mC-D nanocomposites were determined with a TA Q800 at the oscillation frequency of 1.0 Hz and the heating rate of 3°C/min. The experiment was conducted in the tensile mode at temperature range of -100 to 100°C.

The swelling behaviors in water and toluene were measured according to the following procedure: three preweighed (m₀) specimens of X/mC-D nanocomposites were immersed in deionized water or toluene at 60°C for 72 h, respectively. Then the weights of the samples after swelling were recoded as m₁. The swelling ratios were calculated according to the following formula.

\[
\text{Swelling ratio} = \frac{m_1 - m_0}{m_0} \times 100\%
\]

The solubility of neat XSBR and XSBR/CNC-CA nanocomposites was conducted by immersing a certain weight of specimens into toluene/DMF mixed solution at 60°C for 72 h, then ultrasound for an hour.

**Results And Discussion**

**Preparation and characterization of CNC-CA**

The preparation route and reaction mechanism of CA modified CNC-OSO₃H are presented in Scheme 1. Large amount of hydroxyl group on the surface of CNC-OSO₃H reacted with carboxyl group of CA and sulfonate of CNC-OSO₃H was replaced by carboxyl group during the reaction. The FTIR spectra of CNC-CA and CNC-OSO₃H were provided in Fig. 1a. It is worth noting that all CNC-CA were washed thoroughly to eliminate all residual reactants before analysis. As shown in Fig. 1a, in the FTIR spectra of CNC-CA, there is a new band appeared at 1725 cm⁻¹ which corresponds to the C = O vibration from the ester groups of the carboxylic acids (Yu et al. 2014). Compared with the spectrum of CNC-OSO₃H, the band at 812 cm⁻¹ which corresponds to C-O-S vibration disappeared in CNC-CA (Wang et al. 2020). Those evidences suggest that the CA have been grafted successfully on CNC-OSO₃H and most of the sulfonate on the surface of CNC-OSO₃H was removed during the modification process. The removal of sulfonate was also verified by EDS elemental analysis, as shown in table S1, after the surface modification, no sulfur element can be detected.
The crystalline structure of CNC-CA and CNC-OSO$_3$H were analyzed by X-ray diffractions. Both CNC-CA and CNC-OSO$_3$H show diffraction peaks at $2\theta$ of 14.8°, 16.4° and 22.7°, corresponding to the typical X-ray diffraction pattern of form-I crystal of cellulose. Based on the above investigation, it is confirmed that CNC-CA has the same crystalline structure as CNC-OSO$_3$H. The application of CNC-OSO$_3$H in polymer compounds is limited due to the poor heat resistance. Here the thermal degradation of CNC-CA and CNC-OSO$_3$H was compared by thermogravimetric analysis (TGA). As shown in Fig. 1c. and d, CNC-CA displays a markedly enhanced thermal stability compared with CNC-OSO$_3$H. The maximum weight-loss rate temperature of CNC-CA is 354°C, which is 143°C higher than that of CNC-OSO$_3$H. According to the FTIR results of the CNC-CA and CNC-OSO$_3$H, it is deduced that removal of the sulfate groups is a crucial factor in improving the thermal stability of CNCs.

The morphology of CNC-OSO$_3$H and CNC-CA were observed by using TEM (shown in Fig. 2a and b). The individual cellulose nanocrystals of both CNC-OSO$_3$H and CNC-CA can be clearly distinguished. Compared with CNC-OSO$_3$H, the surface of CNC-CA is much more blurry because the surface of CNC-CA is covered with small molecules of citric acid. Because of the strong hydrogen bond interaction between the carboxyl groups of citric acid, CNC-CA is more likely to agglomerate after drying. The average particle sizes of CNC-OSO$_3$H and CNC-CA were estimated by dynamic mechanical analyses (DLS), the values are around 187 nm and 161 nm, respectively. The decrease of the size of cellulose nanocrystals after modification may be due to the further etching of the amorphous region on the surface of CNC-OSO$_3$H by CA.

As mentioned in the experimental section, after surface modification of CNC-OSO$_3$H, un-grafted CA can be separated and reused. To examine the effectiveness of recycled CA, CNC-CA are prepared by using pristine CA, first and second recycled CA as the modifiers. The results are shown in Fig. S1a and b. It can be observed that all these three samples can be stably dispersed in deionized water. In addition, the similar FTIR spectra of these three kinds of CNC-CA indicates that recycled CA can also modify CNC-OSO$_3$H effectively.

**Mechanical Properties of XSBR/CNC-CA Nanocomposites**

In order to study the effect of dual crosslinking network on the mechanical properties of XSBR nanocomposites, the tensile tests were performed. The tensile curves and mechanical property parameters of neat XSBR, X/mC-H and X/mC-D nanocomposites containing 1, 3, 5, 7 and 9 wt% CNC-CA are shown in Fig. 3. For neat XSBR, the stress increased slowly until break, and achieved tensile strength and strain at failure of 2.56 MPa and 389%, respectively. For the X/mC-H nanocomposites, even though there are some improvement by adding CNC-CA into the XSBR matrix, the improvement is quite limited (shown in Fig. 3a). XSBR is not crosslinked by covalent bonds in the X/mC-H nanocomposites, and the role of hydrogen bond is very limited. The tensile strength of X/mC-H nanocomposites is far lower than the requirement of industrial application. Therefore, covalent crosslinking of the nanocomposites is an essential step to further improve its properties. Here, end-epoxy group PEGDE was used as crosslinking
agent to realize chemical crosslinking of nanocomposites through the reaction between epoxy group of PEGDE and carboxyl group of XSBR and CNC-CA. As expected, the mechanical properties of X/mC-D nanocomposites show a significant improvement compared to the X/mC-H ones (as shown in Fig. 3b). For example, with only 5 wt% CNC-CA addition, tensile strength and strain at failure of X/5C-D nanocomposites increase to 8.59 MPa and 456%, respectively, which are even superior to that of X/9C-H nanocomposites. When the content of CNC-CA increases to 9 wt%, the X/9C-D nanocomposites show tensile strength of 12.65 MPa, which are nearly five times that of the neat XSBR, and it's valuable that X/9C-D keeps same elongation at break with pure XSBR. Moreover, from the charts of Fig. 3c. and d., it can be seen that the tensile stresses of the nanocomposites with dual crosslinking networks are nearly double of that with only hydrogen bond, moreover, the fracture energy of the samples with dual crosslinking networks is also significantly improved.

**Dynamic mechanical performance of XSBR/CNC-CA Nanocomposites**

The temperature dependence of the storage modulus (E') and the loss tangent (tan δ) for neat XSBR and X/mC-D nanocomposites with different CNC-CA contents are shown in Fig. 4a and Fig. 4b. A successive increase of the E' values can be observed by increasing the amount of CNC-CA (as shown in Fig. 4a.). For example, the storage modulus at -70°C increases from ∼1049 to 3083 MPa with the addition of CNC-CA from 0 to 9 wt% in the XSBR matrix. The enhancement in modulus below glass transition temperature is a good evidence for the strong reinforced tendency of CNC-CA in the XSBR matrix. Meanwhile, the improvement of crosslinking degree brought by the dual crosslinking network (the covalent bond networks and the H-bond networks) also contributes to the improvement of the modulus.

Figure 4b. shows the variation of tanδ as a function of temperature for the neat XSBR and X/mC-D nanocomposites. XSBR exhibits a peak tanδ value at a temperature around -3°C, which corresponds to the glass transition temperature (T_g) of XSBR. While the T_g of X/mC-D nanocomposites gradually decreases to lower temperature (to -6°C) with the increase of CNC-CA content. Two effects might decrease the T_g value: one is, as reported by Cao et. al, the hydrogen bond formed between filler particles and matrix may act as an internal plasticization resulting in the shifting of T_g to a lower temperature with increasing filler content (Cao et al. 2013); another is, excessive 2-methylimidazole might act as plasticizer in the X/ mC-D nanocomposites because the amount of 2-methylimidazole used is proportional to the CNC-CA content in the nanocomposite. However, further research is needed to clarify this question. Besides, the intensity of tan δ, which indicates the damping properties, decreases with the increase of CNC-CA content, which is mainly due to the decrease of matrix content and the enhancement of filler's restriction effect on the chain mobility of rubber matrix.

**The dispersion of CNC-CA in XSBR/CNC-CA Nanocomposites**
The uniform dispersion of fillers in the matrix plays a key role on improving the properties of nanocomposites. The aggregation of fillers in the matrix forms stress concentration points and has adverse effects on the properties of the nanocomposites.

Figure 5a-f. illustrate the SEM images of cryogenically fractured surfaces of the neat XSBR and X/mC-D nanocomposites containing 1, 3, 5, 7 and 9 wt% CNC-CA, respectively. Compared to the micrograph of neat XSBR, the morphology of CNC-CA in the XSBR matrix can be easily identified. The CNC-CA appears as white dots and their concentration on the fracture surface of the nanocomposites increases directly with the increase of the CNC-CA loading. A uniform distribution of CNC-CA in the XSBR matrix can be observed in all X/mC-D nanocomposites (as shown in Fig. 5b-f). It is obvious that those CNC-CA dots are partial embedded in XSBR matrix, demonstrating good compatibility between the fillers and matrix. Furthermore, no gaps, voids or pull-out cracks are observed on the surface, indicating excellent interfacial adhesion between the CNC-CA and XSBR matrix. This excellent compatibility and interface adhesion can be ascribed to the hydrogen bond between XSBR and CNC-CA and the formation of ester group among PEGDE, XSBR and CNC-CA. This uniform distribution and excellent interfacial adhesion are anticipated to play an important role on the improvement of the mechanical performance of the resulting X/mC-D nanocomposites.

**Swelling properties and solubility of XSBR/CNC-CA Nanocomposites**

The water and toluene uptake ratios of X/mC-D nanocomposites containing 1, 3, 5, 7 and 9 wt% CNC-CA were conducted to investigate the hydrophilicity and crosslinking degree. The X/mC-D nanocomposites were immersed in deionized water or toluene for 72 h at 60°C and weighed. Figure 6. shows the water uptake ratios of neat XSBR and X/mC-D nanocomposites. The water uptake ratios of X/mC-D nanocomposites with CNC-CA content ≤ 7 wt% was much lower than that of neat XSBR. This may be attributed to the dual crosslinking network which makes the components of nanocomposites compact more densely and restricts the chain mobility of the rubber matrix, subsequently inhibits the diffusion of water in the nanocomposites effectively. But, the hydrophilicity of X/mC-D nanocomposites increases with the increase of hydrophilic CNC-CA content, especially when the CNC-CA content > 7 wt%, there is a dramatic increase of water uptake ratios with increase of CNC-CA content. As can be seen from Fig. 6, the water uptake ratios of X/9C-D is almost twice that of neat XSBR. This phenomenon is due to the dual role of CNC-CA in X/mC-D nanocomposites. On one hand, the formation of hydrogen bond between CNC-CA and XSBR molecular chains inhibits the movement of XSBR molecular chains and reduces the possibility of hydrogen bond formation between XSBR molecular chains and water, resulting in the decrease of hydrophilicity of X/mC-D nanocomposites. But, due to hydrogen bonds have saturability, when CNC-CA content reaches a certain level, the number of hydrogen bond between CNC-CA and XSBR molecular chains remains unchanged with increase of CNC-CA. So, the effect of hydrogen bond on the hydrophilicity of X/mC-D nanocomposites doesn't increase when CNC-CA content reaches a certain level. On the other hand, CNC-CA has the inherent hydrophilicity, more CNC-CA, more hydrophilicity is introduced in the X/mC-D nanocomposites. As a result of the above two aspects, water uptake ratios of X/mC-D
nanocomposites increase with increase of CNC-CA content and has an interrupt increase when the content of CNC-CA > 7%. On the contrary, the toluene absorption ratios decrease with the increasement of CNC-CA content, which should be attributed to the increased hydrophilicity caused by the introduction of CNC-CA and the dual crosslinking network restricts the chain mobility of XSBR.

In order to confirm the formation of dual crosslinking network, the X/mC-D and X/mC-H nanocomposites were immersed in DMF/toluene mixed solution for 72 h at 60°C and ultrasound treating for 1h to check the swelling and dissolution behavior. As shown in Fig. S2, all the X/mC-H nanocomposites sheets (upper) and neat XSBR disintegrated and dissolved in the DMF/toluene mixed solvent. In sharp contrast, the shape of all the X/mC-D nanocomposites sheets (below) keep well. The above experimental phenomena further prove that PEGDE has an effective crosslinking effect on the XSBR/CNC-CA nanocomposite.

Structural Characterization and proposed dual crosslinking networks of XSBR/CNC-CA nanocomposites

Structural Characterization of XSBR/CNC-CA nanocomposites

FTIR is a powerful tool for studying the intermolecular interactions, hydrogen bonds and chemical reactions. The FTIR spectra of neat XSBR, PEGDE, XSBR/PEGDE, CNC-CA, and CNC-CA/PEGDE are shown in Fig. 7. In order to make the results clear, here, the mass ratios of XSBR to PEGDE and CNC-CA to PEGDE were set to be 1:1, and the samples were treated at 110°C for six hours. As shown in Fig. 7, the three peaks of 3055, 1252, 854 cm\(^{-1}\) in the FTIR spectrum of PEGDE are the characteristic peaks of epoxy group. Compared to the FTIR spectra of single XSBR and PEGDE, a new band at 1725 cm\(^{-1}\), which corresponds to the C = O vibrations of ester group, appeared in the XSBR/PEGDE blends as shown in Fig. 7a. This reveals that the carboxyl groups of XSBR reacted with the epoxy groups of PEGDE and formed \(\beta\)-hydroxyl esters. The spectra of CNC-CA/PEGDE blends, is shown in Fig. 7b. The three characteristic peaks of PEGDE disappeared in the FTIR spectrum of CNC-CA/PEGDE blends, which means that the carboxyl group of CNC-CA and the epoxy group of PEGDE are esterified. From the above analysis results, we confirmed that the epoxy group on PEGDE can react with carboxyl group of both XSBR and CNC-CA, therefore, PEGDE can effectively crosslink XSBR/CNC-CA nanocomposites.

The existence of hydrogen bonds in X/mC-D nanocomposites was also confirmed by FTIR. As shown in Fig. S3, the intensity of the band between 3500 – 3250 cm\(^{-1}\) corresponding to the hydrogen bond became stronger after compounding the CNC-CA with XSBR, indicating there is a strong hydrogen bond interaction between XSBR and CNC-CA. According to the above results, it can be concluded that dual crosslinking networks of the covalent bond networks and the hydrogen bond networks have been formed in XSBR/CNC-CA nanocomposites, both contribute to improve the mechanical properties of XSBR/CNC-CA nanocomposites.
The proposed model of dual crosslinking networks for crosslinked XSBR/CNC-CA nanocomposites

Based on the above discussion, it can be concluded that dual crosslinking networks featured of hydrogen bonds network and covalent bonds network have been formed in X/mC-D nanocomposite. The esterification of epoxy group of PEGDE with carboxyl group of XSBR and CNC-CA is shown in Scheme S1, the illustration of hydrogen bonds network and dual crosslinking network is shown in Scheme 2. Compared with the ionic bond and covalent bond, hydrogen bond is much weaker so that its ability on the improvement of the mechanical properties of XSBR/mC-H is very limited. By further constructing covalent bonds in the X/mC-D nanocomposites using PEGDE as crosslinking agent, the rubber chains and CNC-CA are held together tightly. The combination of the two networks can effectively improve the compatibility and interface adhesion between CNC-CA and XSBR, as well as the interactions between XSBR-XSBR, which is beneficial to the improvement of the physical and mechanical properties of XSBR/CNC-CA nanocomposites.

Conclusions

In this work, a green and environment friendly method of CA grafting CNC-OSO$_3$H was provided. As-prepared CNC-CA has higher thermal stability and excellent compatibility with XSBR. Meanwhile the dual crosslinking networks in XSBR/CNC-CA nanocomposites were constructed. By using PEGDE as the crosslinker, covalent bonds which serves as the chemical crosslinking network were formed between the XSBR and CNC-CA and PEGDE. The hydrogen bond formed between XSBR and CNC-CA serves as the physical crosslinking network. Besides, 2-methylimidazole the catalyst for esterification acts as plasticizer to endow the XSBR/CNC-CA nanocomposites with better ductility. The combination of dual crosslinking networks and excessive 2-methylimidazole is beneficial for the improvement of the strength and ductility of XSBR/CNC-CA nanocomposites. As CNC-CA content reaches to 9 wt%, the tensile strength of the X/mC-D nanocomposite is five times that of the neat XSBR with the same elongation at break. Beyond that, the addition of CNC-CA improves the solvent resistance of XSBR nanocomposites. This research provides a facile way to the surface modification of CNC-OSO$_3$H and the fabrication of dual crosslinking XSBR nanocomposites with superior properties.

Declarations

Supporting Information Available

Element analysis data of CNC-OSO$_3$H and CNC-CA; Photograph and FTIR spectra of CNC-CA-1st, CNC-CA-2nd and CNC-CA-3rd; The dissolution and swelling pictures of neat XSBR, X/mC-H and X/mC-D nanocomposites in DMF/toluene solution; FTIR spectra of neat XSBR, CNC-CA and X/mC-D
nanocomposites; The esterification involved in the preparation of X/mC-D nanocomposites; The
determination of carboxyl content in XSBR.

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Ethical standards  Compliance with ethical standards

Conflict of interest  The authors declare no competing financial interest.

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**Figures**
**Figure 1**

(a) FTIR spectra, (b) XRD patterns of CNC-CA and CNC-OSO3H, (c) TGA thermograms, (d) Derivative thermogravimetry of CNC-CA and CNC-OSO3H.

![Figure 1](image)

**Figure 2**

TEM micrographs of (a) CNC-OSO3H and (b) CNC-CA.

![Figure 2](image)
Figure 3

The tensile stress-strain curves of (a) neat XSBR and X/mC-H nanocomposites, (b) neat XSBR and X/mC-D nanocomposites, (c) the tensile strength and (d) fracture energy as a function of filler contents.
Figure 4

The temperature dependence of the storage modulus ($E'$) (a) and the loss tangent ($\tan \delta$) (b) for neat XSBR and X/mC-D nanocomposites.

Figure 5

SEM images of the cryogenically fractured surfaces of neat XSBR and X/mC-D nanocomposites: (a) XSBR, (b) X/1C-D, (c) X/3C-D, (d) X/5C-D, (e) X/7C-D, (f) X/9C-D.
Figure 6

Toluene and water uptake ratios of neat XSBR and X/mC-D nanocomposites.
Figure 7

FTIR spectrum of (a) neat XSBR, PEGDE, XSBR/PEGDE and (b) CNC-CA, PEGDE, CNC-CA/PEGDE.

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