Adding 40% DMSO and 2 wt % CaCl₂ increased the tensile strength of the [Amim]Cl for 4 h at 90°C dissolving, regenerating, and pressurized drying. When 4 wt % waste corrugated cardboard was dissolved in [Amim]Cl/CaCl₂, different dissolution systems were selected to construct dissolution systems to produce value-added products from pretreatment of waste corrugated cardboards (P-WCCs). The dissolution behaviors of P-WCCs before and after ball milling were studied in different dissolution systems. The regenerated cellulose films were quickly and efficiently prepared via dissolving, regenerating, and pressurized drying. When 4 wt % waste corrugated cardboard was dissolved in [Amim]Cl for 4 h at 90°C, the regenerated cellulose films featured tensile strengths as high as 59.00 MPa. Adding 40% DMSO and 2 wt % CaCl₂ increased the tensile strength of the film to a maximum value of 85.86 MPa. This demonstrates that DMSO improves the ability of WCC to dissolve in ionic liquids; Ca²⁺ improves the tensile strength and thermal stability of the regenerated cellulose film but reduces its transparency. This work provides a new, simple, and highly efficient way to use WCCs for packaging and wrapping.

1. INTRODUCTION

In 2018, China’s express industry generated about 50.7 billion packages,¹ 80% of which used corrugated boxes as packaging materials but the recovery rate of these packaging materials was less than one-tenth. This resulted in a large number of paper resources being wasted and the main component of the corrugated board being cellulose. Cellulose is a type of polymer with high strength, degradability, and renewability, making it a promising material in biomass reusing. However, there are hydrogen bond network supramolecular structures in the macromolecular structure of cellulose, which makes cellulose insoluble in water and most common organic solvents, severely limiting its applications.² A specific range of the DP of cellulose can be dissolved in a NaOH aqueous solution, as the NMNO solvent system has high energy consumption, poor thermal stability, and may be accompanied by side reactions because of oxidation.³⁴ At present, the LiCl/DMAC solvent system is mainly limited to the laboratory because of LiCl is expensive, the cellulose solution is not stable, and cellulose molecular chain easily gather.⁵ Ionic liquids have proven to be excellent green solvents for cellulose and have great potential in the dissolution and processing of lignocellulose.⁶ 1-Butyl-3-methylimidazolium chloride ([Bmim]Cl), 1-ethyl-3-methylimidazolium acetate ([Emim]OAc), and 1-allyl-3-methylimidazolium chloride ([Amim]Cl) have been used for homogeneous esterification of cellulose as well as for the formation of films and fibers.⁷ Ionic liquids can destroy the hydrogen bonding between spiral cellulose molecular chains, due to the high concentration of anionic intrusions in the cellulose intramolecular and intermolecular hydrogen bonds, forming new hydrogen bonds to divide cellulose.⁸ The main factor affecting the solubility in ionic liquids is the structure of ionic liquids. It has been found that the stronger the hydrogen bond acceptance capacity of anions in the ionic liquid structure, or the short alkyl chain, smaller central group, or strong electron-withdrawing groups such as carbon—carbon double bond and hydroxyl group in the cationic group structure, the stronger the solubility.⁹ It was reported that [Bmim]Cl can dissolve 10% cellulose,¹⁰ [Amim]Cl can dissolve 14.5% cellulose,¹¹ and [Emim]OAc can dissolve 20% of untreated cellulose.¹² But both [Bmim]Cl and [Emim]OAc have a lower melting point and viscosity and greater solubility with regard to cellulose.¹³¹⁴ [Emim]OAc has a complex synthesis procedure than those of [Amim]Cl and [Bmim]Cl; [Bmim]Cl and [Emim]OAc are so hygroscopic that the viscosity determination measurements with the air contact are not reliable.¹⁵ The solubility of imidazolium ionic liquids is because of the strong hydrogen bonding that arises from their alkali and low solvent viscosity. Cl⁻, [OAC]⁻, and other strong basic anions are strong electron donors, as they can easily form a strong hydrogen bond with H in a natural polysaccharide molecule —OH and cooperate with the hydrogen bond between the imidazole cation and O in the natural polysaccharide molecule...

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–OH to form a electron donor–receptor complex, so as to dissolve the natural polysaccharide. Recently, it has been found that cationic structures also play an important role in this behavior during cellulose dissolution. The regenerated cellulose films can be prepared by dissolving, settling, and precipitating lignocellulose (cellulose, hemicellulose, and lignin) with ionic liquids. Moreover, it is easy to prepare these films and their chemical properties are fairly robust. The dissolution of cellulose in ionic liquids involves not only interactions between the solute and the solvent but also mass transfer in the ionic liquid. Ionic liquids are relatively viscous, so it is often necessary to add organic cosolvents such as dimethyl sulfoxide (DMSO) to improve the dissolution rate and reduce the viscosity of the solution system and save time. Xu studied how Ca²⁺ promotes interactions between cellulose chains, which can improve the tensile strength of the regenerated cellulose films. The regenerated cellulose films have been widely used in various fields due to their high performance, such as in wastewater prevention, groundwater pollution treatment, medical equipment, industrial pervaporation, and bio-reactors. Cellulose can also adsorb heavy metals, adsorb organic pollutants, perform antibacterial functions, and act as drug carriers after grafting functional groups onto cellulose or performing other physical modifications. In recent years, the regenerated cellulose films have also received extensive attention in green packaging, coating, screen display devices, optoelectronic materials, and electronic flexible screens. Sixta presented a novel regenerated cellulose fiber process of the Lyocell type using [DBNH]-[OAc] as a solvent, and the resultant cellulose solution has a high dissolution power and low viscosity. Its excellent properties make the Ioncell-F fibers suitable as a reinforcing material in composite structures and other technical applications.

In this work, we seek to achieve high-value utilization of waste corrugated cardboard using a [Amin]Cl/DMSO/CaCl₂ dissolution system. DMSO can improve the dissolution of cellulose in [Amin]Cl and CaCl₂, can improve the mechanical property of the films. In addition, the recovery of [Amin]Cl is pretty high and the recovered [Amin]Cl still has a good ability to dissolve fiber, and the performance of the cellulose film (R-WCC film) regenerated from the recovered [Amin]Cl is also very good.

2. RESULTS AND DISCUSSION

2.1. Composition of Fibers. Waste corrugated cardboard is mainly composed of cellulose, hemicellulose, and lignin, accounting for 52.02, 6.79, and 10.43% of the total composition, respectively. From Table 1, it can be inferred that the cellulose ratio of P-WCC is increased to 82.19%, the amounts of ash and impurities are greatly reduced, the ratios of lignin and hemicellulose are reduced by the NaOH solution treatment, and the pulp can be bleached by H₂O₂. Finally, after the KOH solution treatment, the WCC became fluffy, which facilitated the dissolution.

Moreover, the DP of cellulose in WCC and P-WCC is 1635 and 600, respectively, and the DP of cellulose in the P-WCC film decreased to 485, according to GB/T 1548-2016. This may be due to the protonation of Cl⁻ with water molecules to form HCl, resulting in the acid drop of polysaccharide molecules in the dissociation reaction.

2.2. Viscosity Analysis. The addition of a cosolvent can have a strong influence on the viscosity, polarity, solvation, and other physical properties of an ionic liquid. Measuring the viscosity of cellulose in an ionic liquid system can help to clarify the dissolution mechanism of such a mixture.

As shown in Figure 1, DMSO can greatly improve the rheological properties of the solvent, and adding just 10% DMSO at 70 °C can reduce the solution viscosity by 58.94%. Therefore, adding DMSO at a lower temperature has a greater overall impact on the viscosity of [Amin]Cl. The viscosity of [Amin]Cl decreases with the increase of DMSO content over 40%, but the change is negligible. With the addition of the cosolvent DMSO, the viscosity of the dissolving system decreases, the internal friction of the ionic liquid decreases, and the hydrocarbon binding force decreases. This makes the cellulose molecular chains disperse better in [Amin]Cl, thereby improving the solubility of cellulose.

2.3. Solubility Properties. The polarizing micrographs of P-WCC in [Amin]Cl at 90 °C is shown in Figure S1. Figure 2a shows the solubility and dissolution time of P-WCC and BM-P-WCC in [Amin]Cl at 90 °C. The solubility of the fiber after ball milling increases from 10.79 to 19.01 wt %. This increase in solubility is due to the breaking of hydrogen bonds between cellulose molecules under mechanical force, which destroys the crystalline regions. Ball milling treatment reduces the fiber size and increases the specific surface area, thus improving the solubility as observed by Bodvik. Cellulose absorbs the mechanical energy during the ball milling process as shown in the X-ray diffraction (XRD) data in Figure S2; the crystallinity of the P-WCC fiber was 72.19%, while that of the BM-P-WCC fiber was reduced to 55.42%, such that the crystalline structure of cellulose changes from a metastable cellulose I type structure to a stable cellulose II type structure. Further, the cross-linking between cellulose molecules in the II type structure is closer, decreasing the dissolution time.

Figure 2b shows that as the temperature increases, the solubility and dissolution time increase continuously. In this

Table 1. Composition of WCC, P-WCC, and P-WCC Film

|        | cellulose (%) | hemicellulose (%) | lignin (%) | ash (%) | other additives (%) |
|--------|---------------|-------------------|------------|---------|---------------------|
| WCC    | 52.02         | 6.79              | 10.43      | 15.71   | 15.05               |
| P-WCC  | 82.19         | 7.11              | 7.65       | 0.92    | 2.13                |
| P-WCC film | 81.62      | 6.22              | 7.03       | 1.24    | 3.89                |
experiment, 4 wt % BM-P-WCC was added to [Amim]Cl to examine the effect of the dissolution temperature on its solubility and dissolution time. The solubility is 16.73 wt %; when the temperature increases to 150 °C, the solubility increases to 38.31 wt % and the dissolution time decreases from 195 to 54 min.

Adding DMSO can reduce the viscosity of the cellulose solution and effectively improve the solubility of cellulose in an ionic liquid.25 BM-P-WCC (4 wt %) was added to 5 g of [Amim]Cl and dissolved at 90 °C for 4 h. Figure 2c shows that as the amount of DMSO increases, the solubility of BM-P-WCC in the ionic liquid first increases and then decreases. When the DMSO content is more than 20%, the solubility begins to increase slowly; the solubility reaches a maximum of 25.47 wt % when the DMSO content is 40% and then begins to decrease. In other words, the time required for 5 g of [Amim]Cl to dissolve 0.2 g of BM-P-WCC decreases from 163 to 25 min. These results show that the addition of DMSO reduces the viscosity of the solvent and effectively improves the solubility of WCC fibers in [Amim]Cl. Under the action of a certain temperature and proper cosolvent, the internal friction force of the ionic liquid decreases and the binding force of the hydrocarbon also decreases, which manifests as a decrease in viscosity. The diagram of the dissolution mechanism of WCC fibers in ionic liquids is shown in Figure 3. The cellulose molecular chains can be better dispersed in [Amim]Cl, and thus the solubility of WCC fibers in [Amim]Cl improves under such conditions, a conclusion also reached by Andanson26 and Lv.52

2.4. Mechanical Characterization. The regenerated cellulose films (named as P-WCC film and BM-P-WCC film) were prepared by adding 4 wt % P-WCC or BM-P-WCC to [Amim]Cl and dissolving for 4 h at variable dissolution temperatures. As shown in Figure 4a, the tensile strength of the film first increases and then decreases as the dissolution temperature increases, and the tensile strength of the BM-P-
WCC film is slightly higher than that of the P-WCC film. The tensile strengths of the P-WCC and BM-P-WCC films had maximum values of 59.00 and 62.10 MPa, respectively at 90 °C. When the temperature increased above 110 °C, the tensile strengths of these films became poor. At temperatures higher than 150 °C, the film formation became noticeably difficult and the tensile strengths were recorded as 0 MPa. The WCC fiber dissolves in [Amim]Cl through a nonderivative form, which can better retain the original mechanical strength of lignocellulose in the fiber. When the temperature is too high, the cellulosic macromolecules oxidize and degrade in ionic liquids, resulting in poor tensile strength. According to Figure 4b, when the dissolution time increases from 3 to 4 h, the tensile strengths of the P-WCC and BM-P-WCC films increase from 48.69 and 51.16 MPa to 59.00 and 62.10 MPa, respectively. The tensile strengths of the films begin to decrease when the dissolution time continues to increase beyond this point, denoting an optimal dissolution time of 4 h; this is consistent with the research results of Zhang.53 Because lignocellulose is easy to oxidize and degrade in ionic liquids,54−57 when the dissolution time is too long, the tensile strength of the regenerated cellulose film decreases.

When the amount of the added fiber is close to the maximum solubility of [Amim]Cl, the viscosity of the ionic liquid will be very high and the viscosity of the film-forming solution has a great influence on the film-forming effect. Therefore, we studied the influence of different added WCC ratios on the tensile strength of the P-WCC and BM-P-WCC films. In Figure 4c, the regenerated cellulose films were prepared by dissolving BM-P-WCC or P-WCC at variable ratios in [Amim]Cl at 90 °C for 4 h. When 4 wt % WCC was added, the tensile strength of the resulting P-WCC and BM-P-WCC films reached their maximum values of 59.00 and 62.10 MPa, respectively, and then decreased as more WCC fibers were added. When the fiber ratio is greater than 8 wt %, a cellulose hydrogel forms due to the high viscosity of the film-forming solution. In addition, the tensile strength of the BM-P-WCC film is slightly higher than that of the P-WCC film; crystallinity may affect the tensile strengths of the films, as shown in Figure S2, which is consistent with the results of Zheng.14

As BM-P-WCC was proved to be too difficult to be regenerated into a film using the [Amim]Cl/DMSO system, only the tensile strength of the regenerated cellulose film prepared using P-WCC was measured here. DMSO film was prepared by adding 4 wt % P-WCC, different DMSO ratios, and dissolving at 90 °C for 4 h. As shown in Figure 4d, the tensile strength of the film decreases as the DMSO ratio increases and at a ratio of 40%, the tensile strength of the P-WCC film is 28.65 MPa, after which the tensile strength sharply decreases. DMSO can prevent the oxidation and decomposition of fibers in ionic liquids and improve the
dissolution efficiency of fibers in [Amim]Cl. However, too much DMSO will promote the acid-catalyzed water depolymerization of cellulose in [Amim]Cl and will reduce the tensile strength of the film. It shows the change in the tensile strength of the regenerated cellulose film (named as CaCl₂ film) in Figure 4d prepared by adding 4 wt % P-WCC and 40% DMSO at 90°C at 4 h of dissolution, using varying CaCl₂ additions of 0−4 wt %. The tensile strength of the CaCl₂ film first increases and then decreases as the amount of CaCl₂ increases. At a CaCl₂ addition of 2 wt %, the tensile strength of the film reaches a maximum value of 85.86 MPa, which is 45.53% higher than that of the P-WCC film (59.00 MPa) and 199.86% higher than that of the DMSO film (28.65 MPa). The tensile strength of common commercial polyolefin films (PE, PP) is typically 20−40 MPa, and the tensile strength of the cellulose film regenerated from ionic liquids ([EMIMO]Ac) and ([Bmim]Cl) is, respectively, about 70% and 34 MPa. The regenerated porous cellulose films were prepared by dissolving in the LiCl/DMAc solvent; its tensile strength reached 29.22 MPa and the tensile strength of all-cellulose films regenerated from 7 wt % NaOH/12 wt % urea aqueous solution change from 30 to 135 MPa. Thus, this CaCl₂ film has the potential to replace petroleum-based plastic bags as packaging materials. The same anion of CaCl₂ and [Amim]Cl improves the tensile strength of the regenerated cellulose film, which can promote the dissolution and regeneration of cellulose in ionic liquids. Meanwhile, Ca²⁺ is cross-linked with the cellulose molecular chains to enhance the tensile strength of the film via the mechanism shown in Figure 5. The excess CaCl₂ will also adhere to the surface of cellulose and reduce the tensile strength of the CaCl₂ film.

2.5. Micromorphology Analysis. Figure 6 shows the scanning electron microscopy (SEM) images of the surface and cross section of the films prepared under different conditions. There are a few particles on the surface (Figure 6a-1) and cross section (Figure 6a-2) of the P-WCC film and a few protrusions and holes on the cross section, which may be caused by the removal or fracturing of larger fibers when the film is quenched. In contrast, the surface (Figure 6b-1) of the BM-P-WCC film is more smooth, compact, and bright owing to ball milling, and the cross-sectional structure (Figure 6b-2) is even and smooth without a grainy appearance.
the DMSO film is rough but the fiber structures are exposed and are interlaced and stacked to form a layered structure (Figure 6c-2). The surface of the CaCl₂ film is still relatively rough with obvious fiber structures (Figure 6d-1) and the cross section features some pores (Figure 6d-2). The inner fibers are closely interwoven into a thin network structure but this structure is slightly flat and dense compared with that of the DMSO film (Figure 6c-1). The addition of CaCl₂ inhibited the
production of protons from the active hydrogen at the second position on [Amim]+ in the ionic liquid, thus weakening the acid-catalyzed water depolymerization of cellulose. This allows Ca2+ to cross-link on the cellulose molecular chain, thus enhancing the tensile strength of the film. Xu28 demonstrated that such nonrigid fiber structures can be effectively cross-linked by Ca2+.

2.6. Roughness Analysis. Figure 7 shows the three-dimensional (3D) topographic maps of the P-WCC, BM-P-WCC, DMSO, and CaCl2 films using atomic force microscopy.
The tensile strength, surface roughness (Figure S5), and transmittance (Figure S6) of the BM-P-WCC and R-WCC films are relatively close, indicating that the recovered ionic liquid still has a good ability to dissolve the fiber and that this change has little effect on the physical properties of the film.

### 3. CONCLUSIONS

Four kinds of regenerated cellulose films with good mechanical properties were successfully prepared using low-cost waste corrugated cardboard (WCC) instead of high-grade pulp as a raw material. Adding 4 wt % fiber in [Amim]Cl and dissolving for 4 h at 90 °C yielded the best results. The regenerated cellulose films were smooth, compact, and transparent or semi-transparent, depending on the DMSO ratio, CaCl₂ ratio, and whether or not ball milling was used. The films exhibited good mechanical properties, such as a maximum tensile strength of 85.86 MPa for the CaCl₂ film. These films also had good thermal stability and could be used in fields with low transparency requirements. The BM-P-WCC film had a smooth and uniform surface and high transmittance and could be applied in fields requiring higher requirements. The transmittance of the P-WCC film was slightly lower than that of the BM-P-WCC film. With the addition of the cosolvent DMSO, the tensile strength of the DMSO film decreased, whereas the DMSO film prepared with 40% DMSO had low tensile strength, low transmittance, and poor transparency. [Amim]Cl was a nonderivatizing solvent for the WCC fiber. Moreover, a deeper analysis has led to a greater understanding of this route. The XRD patterns indicate a transformation in the crystal structure (from cellulose I to cellulose II) that occurred during the dissolution and regeneration processes. TGA results demonstrated that the DMSO and CaCl₂ films possessed high thermal stability similar to that of the P-WCC film. Therefore, the present work provides a simple and effective method to convert low-grade cellulose resources into high-value cellulose-based products.

### 4. MATERIALS AND METHODS

#### 4.1. Materials.

- Waste corrugated cardboard (C-type corrugated box), [Amim]Cl (C₇H₁₁N₂Cl, ≥99%, Lanzhou Institute of Chemical Technology, Chinese Academy of Sciences).
- DMSO (AR, ≥99.8%, Tianjin Fuyi Fine Chemical Co., Ltd.), NaOH (AR, ≥96.0%, Chengdu Kelong Reagent Plant), KOH (AR, ≥96.0%, Chengdu Kelong Chemical Reagent Plant), K₂CO₃ (AR, ≥96.0%, Chengdu Kelong Chemical Reagent Plant), Na₃PO₄·12H₂O, Shanghai Aladdin Biochemical Technology Co., Ltd.), and phosphotungstic acid (AR, H₃O₃PW₁₂·3H₂O, Shanghai Aladdin Biochemical Technology Co., Ltd.) were used in this study.

#### 4.2. Pretreatment of Waste Corrugated Cardboard.

As shown in Figure 10, waste corrugated cardboard was cut into small pieces (1–2 cm), crushed and passed through a 700 μm sieve, treated with a 2 wt % NaOH solution in the ratio 1:10 (g/mL), and then stirred at 90 °C for 2 h. Subsequently, the samples were treated with a 20% H₂O₂ solution in the ratio 1:20 (g/mL) and stirred at 70 °C for 1 h. Finally, this sample was treated with a 4 wt % KOH solution in the ratio 1:10 (g/mL) and stirred at 90 °C for 2 h. The precipitate was washed with deionized water to a neutral pH and dried under vacuum at 40 °C for 48 h to obtain the waste corrugated board (named as P-WCC), which was kept sealed when not in use. One-half of the above-mentioned P-WCC was used to prepare BM-P-WCC via ball milling with a rotation speed of 4000 rpm for 10 h and kept sealed when not in use.

#### 4.3. Preparation of the Regenerated Cellulose Films.

The preparation procedure of the regenerated cellulose films is shown in Figure S1. After pretreatment, 0.8 g of P-WCC, 0.2 g...
of CaCl₂, and 40% DMSO (to 20 g of [Amim]Cl) were dispersed into 20 g of [Amim]Cl. The mixture was then heated at 90 °C for 4 h under magnetic stirring, so that a homogeneous 4 wt % P-WCC/[Amim]Cl solution was obtained. The resulting solution was cast onto a poly-(tetrafluoroethylene) plate with a spreader to obtain a 0.5 mm thick layer after 30 min setting time and immediately immersed into a coagulation bath (deionized water) to form a cellulose-based hydrogel, which was labeled as Hydrogel. The cellulose-based hydrogel was then washed at least five times with distilled water until the residual [Amim]Cl was completely removed (the scrubbing solution was reserved for later use). Finally, applying 900 MPa at 50 °C for 20 min using a Kaiser rapid sheet former (2DA, Estanit GmbH, Germany), the regenerated cellulose films were obtained. The films were dried at 60 °C for 24 h for characterization.

4.4. Chemical Compositions of WCC and P-WCC. The mass fraction of cellulose, hemicelluloses, lignin, and ash content in WCC and P-WCC were characterized by the Fan method, as shown in Table 1.64

4.5. Viscosity of [Amim]Cl/DMSO. The viscosity of [Amim]Cl/DMSO was measured for 0.5 min using a modular advanced rheometer (HAAKE MARS 40, HAAKE Technology Co., Ltd., Germany), with an FL22 4B/ss Vane rotor and CCB25 measuring cup. For each sample, at least three specimens were tested, and the average value was reported.

4.6. Solubility Properties of Fibers in [Amim]Cl. The dissolving property of P-WCC and BM-P-WCC in [Amim]Cl was observed using a polarizing microscope (Nikon E200, Nikon Co., Ltd., Japan). The solubility was calculated following Liu,65 where the dissolution time refers to the time required to completely dissolve 0.2 g of P-WCC or BM-P-WCC in 5 g of [Amim]Cl.

4.7. Mechanical Characterization. The tensile strength of the regenerated cellulose film was measured on an electronic universal material testing machine (3367, Instron, U.K.) at a crosshead speed of 5 mm/min. Specimens 50 mm in length and 10 mm in width were used for these tests, and a gauge length of 25 mm was maintained. For each sample, at least five specimens were tested, and the average value was reported. The mechanical tests were done by following the ASTM D-882 standard.

4.8. Micromorphology Analysis. Samples (cut into squares of 10 mm × 10 mm) were lifted with tweezers and attached to the stage with a conductive paste. Then, they were sputter-coated with a thin layer of platinum before being observed by scanning electron microscopy (SEM) (Phenom-World Pro, FEI). The film was placed in liquid nitrogen for 10 s and quenched to obtain the fracture surface; the cross-sectional structures of the films were observed at 10 V and the surface structures of the films were observed at 5 V.

4.9. AFM Analysis. Films were cut into squares of 10 mm × 10 mm for observation, and the samples were measured by atomic force microscopy (AFM) (S1000N, Hitachi High-Tech Co., Ltd.) at 8.5 V, 299 kHz, tapping mode, C = 32 N/m, tip diameter = 7 nm, and probe type = SI-DFP2.

4.10. TGA Analysis. The thermal decomposition behavior of the samples was investigated using a synchronous thermal analyzer (STA 449 F5, Netzsch Instruments Manufacturing Co., Ltd.) under a nitrogen atmosphere at a heating rate of 10 °C/min, gas flow of 20 mL/min, and scanning temperature of 20−550 °C.

4.11. Optical Performance. Films were cut into specimens of 40 mm × 9 mm before testing, and the transmittance of a film laid on the inner wall of the colorimetric vessel was measured by UV spectrophotometry (Lambda950, PerkinElmer Instruments) over the wavelength range of 380−780 nm.

4.12. Recovery Rate of Ionic Liquids. The scrubbing solution reserved after preparing the regenerated cellulose films was filtered with a Nylon Millipore filter (Φ = 0.45 μm), and the water solvent was removed by rotary evaporation (RE-3000 B, Shanghai Xiande Experimental Instrument Co., Ltd.) at 105 °C under a 0.1 MPa vacuum. 66 3A zeolite was added to remove impurities, and the material was vacuum-dried at 80 °C for 24 h to obtain the recovered ionic liquid.

ASSOCIATED CONTENT

Supporting Information
The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acsomega.0c02713.
Data of the polarizing micrographs of P-WCC in [Amim][Cl]; XRD spectrum of P-WCC and BM-P-WCC; FTIR and $^1$H NMR of [Amim][Cl] and recovered [Amim][Cl]; AFM graphic and optical performance figure of the R-WCC film (PDF)

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**Notes**

The authors declare no competing financial interest.

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