High Oxygen Nanocomposite Barrier Films Based on Xylan and Nanocrystalline Cellulose

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The goal of this work is to produce nanocomposite film with low oxygen permeability by casting an aqueous solution containing xylan, sorbitol and nanocrystalline cellulose. The morphology of the resulting nanocomposite films was examined by scanning electron microscopy and atomic force microscopy which showed that control films containing xylan and sorbitol had a more open structure as compared to xylan-sorbitol films containing sulfonated nanocrystalline cellulose. The average pore diameter, bulk density, porosity and tortuosity factor measurements of control xylan films and nanocomposite xylan films were examined by mercury intrusion porosimetry techniques. Xylan films reinforced with nanocrystalline cellulose were denser and exhibited higher tortuosity factor than the control xylan films. Control xylan films had average pore diameter, bulk density, porosity and tortuosity factor of 0.1730 μm, 0.6165 g/ml, 53.0161% and 1.258, respectively as compared to xylan films reinforced with 50% nanocrystalline cellulose with average pore diameter of 0.0581 μm, bulk density of 1.1513 g/ml, porosity of 22.8906% and tortuosity factor of 2.005. Oxygen transmission rate tests demonstrated that films prepared with xylan, sorbitol and 5%, 10%, 25% and 50% sulfonated nanocrystalline cellulose exhibited a significantly reduced oxygen permeability of 1.1387, 1.0933, 0.8986 and 0.1799 cm³·μm/m²·d·kPa respectively with respect to films prepared solely from xylan and sorbitol with a oxygen permeability of 189.1665 cm³·μm/m²·d·kPa. These properties suggested these nanocomposite films have promising barrier properties.

Keywords: Nanocomposites; Xylan; Nanocrystalline cellulose; Oxygen barrier

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Biobased plastics have generated increasing interest to replace petrochemical based polymers in light of their ready availability, sustainability, limited environmental impact and simplified end-of-life disposal issues [1-4]. High barrier packaging materials require low oxygen permeability with good mechanical strength. Replacing existing oxygen barriers such as aluminum and synthetic polymers with biopolymers is a research topic of increased interest. Early studies examined the use of chitosan, starch and cellulose derivatives to address these challenges and were shown to have good film forming properties [5].

Acetylated galactoglucomannan (AcGGM) hemicellulose was found to be an excellent candidate for making new renewable barrier materials [6]. The oxygen barrier permeability of the AcGGM films were found to be similar to, or lower than, the values reported on oxygen barrier films made from glucuronoxylan [7] and other polysaccharides, such as starch [8], chitosan [9] and mixtures of various polysaccharides [10]. Hartman reported oxygen permeability of 2.0 cm³·μm/m²·d·kPa for GGM-sorbitol film [6]. The oxygen barrier properties of films obtained from a mixture of O-acetyl-galactoglucomannan and either alginate or carboxymethylcellulose were also studied. The reported values of oxygen permeability of GGM/alginate/
glycerol blend were 4.6 cm$^3$/μm$^2$·d·kPa [6]. The oxygen permeability of the GGM films was lower than that of glycerol-plasticized amylose and amylopectin films [11], but not as low as that of sorbitol-plasticized aspen glucuronoxylan films [7,12]. The oxygen permeability of oat spelt arabinoxylan films plasticized with 40% sorbitol was 4.7 cm$^3$/μm$^2$·d·kPa, which is slightly lower than that of GGM films [13]. Biobased free-standing films and coatings with low oxygen permeability of 1 cm$^3$/μm$^2$·d·kPa have been also prepared from a wood hydrolysate [14]. Films made from these polysaccharides are brittle and therefore to form cohesive films requires plasticizers such as sorbitol and xylitol. However, even with the plasticizers, the mechanical properties of these films have been considered to be less than desirable. Nanocrystalline cellulose has been studied as reinforcements of various synthetic and some natural polymer matrices [15-17] to improve the strength properties owing to their high bending strength of 10 GPa, and elastic modulus of 143 GPa [18-19]. Lagaron et al. [20] discussed the role of crystalline structure of polymers and emphasized that high crystallinity improves barrier properties. Nanocrystalline cellulose is greater than 60% crystalline [21-22] and this property together with the resulting rigid hydrogen-bonded network of nanocrystalline cellulose can cause an increase in tortuosity and smaller pore size for nanocomposites which may be utilized to create high barrier materials. In a recent study, spruce galactoglucomannans (GGM) and konjac glucomannan (KGM) were mixed with nanocrystalline cellulose (NCC) to study the mechanical and barrier properties of the films [23,24]. The tensile strength of unplasticized KGM films increased by 30% but the mechanical properties of the plasticized films were not affected with increased in nanocrystalline cellulose. The presence of 5% of nanocrystalline cellulose did not significantly affect the oxygen permeability of the films.

Xylan is one of the most common hemicelluloses, is the most abundant polysaccharide in nature after cellulose, and is an attractive resource for film production [25-27]. Prior studies by Saxena et al. [28] have shown that the addition of 7% nanocrystalline cellulose to xylan-sorbitol films increased the tensile energy absorption of the resulting films by 445% and the tensile strength of the film by 141%. Recent studies demonstrated that films with 10% nanocrystalline cellulose exhibited a 74% reduction in specific water transmission properties with respect to films prepared solely from xylan/sorbitol, and a 362% reduction in water transmission rate with respect to xylan films reinforced with 10% softwood kraft fibers [29]. The objective of the current study is to evaluate the oxygen barrier properties of xylan-nanocrystalline cellulose composite films.

Materials and Methods

Materials

Oat spelt xylan was obtained from Aldrich and was determined to contain 81.0% xylose, 9.8% arabinose, 7.6% glucose, 1.4% galactose and 0.2% of mannose. A commercial elemental chlorine-free (ECF) bleached softwood (SW) kraft pulp was used as received. Dialysis tubes were purchased from Spectrum Labs. All other reagents and solvents were purchased from Aldrich and used as received.

Preparation of Nanocrystalline Cellulose

Nanocrystalline cellulose was prepared following the procedure outlined by Pu [30]. In brief, softwood kraft pulp was ground in a Wiley mill to pass through a 20-mesh screen. The acid hydrolysis of the milled pulp was accomplished with 64 wt % sulfuric acid at 45°C for 45 min with stirring. The reaction was halted by adding a 10-fold excess of deionized (DI) water. After standing overnight, the water was decanted and the settled celluloses were collected with a minimal amount of water. Excess acid was removed with a water wash (200 ml), and centrifuged for 20 min at 11,000 rpm. The aqueous phase was discarded and the wash cycle was repeated an additional two times. The solids suspension was then dialyzed against DI water for 3 days using regenerated cellulose dialysis tubing with a 12000-14000 molecular weight cut off until the pH of the solution reached 7. Sonication was performed on the neutral nanocrystalline cellulose solution for 35 min with an Ultrasonic Processor GEX-500 sonicator with standard titanium alloy probe (tip diameter 1.27 cm) at 28 Watts power input while immersed in an ice bath. The average yield of nanocrystalline cellulose was 35%. Conductometric titration was used to measure the bulk charge on nanocrystalline cellulose following the method described by Katz (1984) [31]. A known amount of nanocrystalline cellulose (0.31% of suspension containing 0.50 g of whisker) was taken and mixed with 5.00 ml of 0.01 M NaCl before measurement. The solution was titrated with 0.05 M NaOH.

The change in conductivity was recorded with a conductometer. The bulk charge for the nanocrystalline cellulose was found to be 9.73 mmol acid groups/100 g. The bulk carboxylic acid group content of the fibers was measured by
conductometric titration methodology. Air-dried fibers equivalent to 1.50 g of oven-dried fibers were added to 300.00 ml of 0.10 M HCl and stirred for 1 h. The pulp was filtered using a vacuum flask and then washed with at least 400 ml of deionized water. The washed pulp was treated with 0.001 M NaCl and 1.50 mL of 0.10 M HCl, stirring and bubbling nitrogen through the solution. The treated pulp was finally titrated with 0.05 M NaOH in 0.25 ml aliquots. The change in conductivity was recorded with conductometer. The bulk charge for the starting cellulose fibers was found to be 3.82 mmol carboxylate groups/100 g.

Preparation of xylan- sulfonated nanocrystalline cellulose nanocomposite film

Xylan composite films were formed by adding an aqueous suspension (35.00 ml) of sulfonated nanocrystalline cellulose to xylan (0.25 g). The solid nanocrystalline cellulose content was 0.0, 5.0, 10.0, 25.0, and 50.0 wt% of the total mixture of sorbitol, xylan, and nanocrystalline cellulose. Sorbitol (0.25 g, 1.37 mmol) was then added to the mixture with stirring and this solution was heated to 95ºC for 15 min. The whole solution was then poured into polystyrene petri dishes (4.60 cm in radius) and allowed to dry at room temperature for three days. The thickness of resulting composite films for xylan-sulfonated nanocrystalline cellulose films with 10%, 25% and 50% sulfonated nanocrystalline cellulose was 0.0812 mm, 0.0877 mm, 0.1312 mm, respectively.

Characterization

The cross-section and surface morphology of the composite films were analyzed by Hitachi S800, thermally assisted field emission (TFE) scanning electron microscope (SEM) with an accelerating voltage of 12 kV. The samples were sputter-coated with gold prior to examination. Nanocrystalline cellulose and the composite films were analyzed using Dimension 3100 scanning probe microscope and Nanoscope III controller. The images were acquired in tapping mode in air using 1–10 Ω cm phosphorus (n) doped silicon tip with the cantilever resonance frequency of 150 kHz. Scans were done at 5 microns.

Oxygen permeability Analysis

The oxygen transmission of the films was measured using a Mocon Ox-Tran 2/211 apparatus (Modern Controls Inc., Minneapolis, USA) with a coulometric sensor in accordance with ASTM method D 3985-95 [32].

Mercury Intrusion Porosimetry Analysis

Micromeritics’ AutoPore IV 9500 Series was used to measure the porosity, bulk density, and average pore diameter and tortuosity factor of the control and nanocomposite films.

Results and Discussion

This study examines the oxygen barrier properties of xylan-nanocrystalline cellulose composite films. By AFM analysis, the sulfonated nanocrystalline celluloseic were observed to have rod like structure with an average length of 150-200 nm and a width of less than 20 nm (Fig. 1). AFM images, acquired using tapping mode, of the xylan/sorbitol films reinforced with nanocrystalline cellulose show well dispersed sulfonated nanocrystalline cellulose on xylan surface in comparison to more open structure of xylan/sorbitol control films (Fig. 2).

Table 1 Effect of sulfonated nanocrystalline cellulose dosage on oxygen transmission rate of xylan film.

| Samples                                      | Specific oxygen transmission rate (cm³/m²·day) |
|----------------------------------------------|-----------------------------------------------|
| Control (Xylan)                              | 354.950                                       |
| Xylan + 5% sulfonated nanocrystalline cellulose | 1.442                                         |
| Xylan + 10% sulfonated nanocrystalline cellulose | 1.364                                         |
| Xylan + 25% sulfonated nanocrystalline cellulose | 1.038                                         |
| Xylan + 50% sulfonated nanocrystalline cellulose | 0.139                                         |
| EVOH                                          | 3.5                                           |
| Polyester, oriented, Polyvinylidene chloride (PVdC) coated | 9.15                                          |
| MFC film                                      | 17.0-18.5                                     |

FIG. 1. AFM image of sulfonated nanocrystalline cellulose with a concentration of 0.7% by weight.
The measured oxygen permeability is lower or comparable to the often used barrier plastic ethylene vinyl alcohol (EVOH) [33] and the films made from microfibrillar cellulose [34], see Table 1.

Oxygen permeability values were calculated by dividing the oxygen transmission rates by the differential partial pressure of oxygen across the film (1 atm or 101.3 kPa) and multiplying by the film thickness in microns [5]. Table 2 summarized the oxygen permeability of some of the literature work and current work. The oxygen transmission rates as summarized in Table 1 at 25% and 50% dosage of nanocrystalline cellulose decreased drastically with respect to control xylan films and are the two lowest values that we obtained in this study. It will be an interesting subject to explore the porosity, bulk density and tortuosity factor at these two levels and the control xylan films.

As summarized in Table 3, the density and tortuosity factor of the composite film increased while the pore diameter and porosity decreased as the loading of sulfonated nanocrystalline cellulose increased in the xylan-based films.

Table 2 Oxygen Permeability of biopolymer films.

| Sample                                      | Oxygen permeability (cm³·μm²·d·kPa) |
|---------------------------------------------|-------------------------------------|
| GGM-sorbitol                                | 2.0[6]                              |
| GGM/alginate/glycerol blend                 | 4.6[6]                              |
| Oat spelt arabinoxylan films plasticized with 40% sorbitol | 4.7 [13]                      |
| Biobased free-standing films and coatings from a wood hydrolysate | 1.0 [14]                        |
| Xylan + 50% sulfonated cellulose whiskers   | 0.1799 (current study)              |

SEM images of the control xylan film surface showed agglomerated structures on the surface in comparison to a more uniform surface for the nanocrystalline cellulose-xylan films (Fig. 3 (a) and 3 (b)).

Oxygen transmission rate at 5% and 10% charge of nanocrystalline cellulose doesn’t differ much but a significant drop of transmission rate as compared to control. We studied xylan-10% nanocrystalline cellulose film under SEM (Fig. 3) and AFM (Fig. 2b) and found that control xylan film surface in Fig. 3 (a) shows agglomeration in comparison to well dispersed sulfonated nanocrystalline cellulose on xylan surface in Fig. 3 (b). The uneven structure and agglomeration of the xylan can be the cause of higher oxygen transmission rate of control xylan film in comparison to xylan reinforced with 10% sulfonated nanocrystalline cellulose.

Table 3. Mercury intrusion porosimetry measurements of control xylan and nano-composite films.

| Sample                                      | Average pore diameter (μm) | Bulk density (g/ml) | Porosity (%) | Tortuosity factor |
|---------------------------------------------|---------------------------|---------------------|--------------|-------------------|
| Control (Xylan)                             | 0.1730                    | 0.6165              | 53.0161      | 1.258             |
| Xylan+25% sulfonated nanocrystalline cellulose | 0.1301                    | 0.8705              | 40.8501      | 1.700             |
| Xylan+25% sulfonated nanocrystalline cellulose | 0.0581                    | 1.1513              | 22.8906      | 2.005             |

SEM cross-section images of freeze fracture of control xylan films showed a rough texture with small cracks in the film as summarized in Fig. 4(a) and 4(b). The same analysis for the xylan film reinforced with nanocrystalline cellulose exhibited smooth fractured surface and less porous structure (see Fig. 4(c) and Fig. 4(d)).
The dense structure of composite films with reduced surface porosity is due to the combination of the nanocrystalline cellulose and the resulting rigid hydrogen-bonded network within the nanocrystalline cellulose and with the matrix [2]. This causes an increase in the tortuosity factor and bulk density with decrease in porosity and average pore diameter of the composite films resulting in lower oxygen transmission rates of composite films in comparison to the control.

Nanocrystalline cellulose have been reported to have crystallinity values greater than 60% [21] which would further reduce the transmission of oxygen molecules through these nanocomposite films. The tortuosity factor measurements from mercury porosimetry techniques showed that the control xylan films have a tortuosity factor of 1.258 whereas the xylan films containing 50% nanocrystalline cellulose had a tortuosity factor of 2.005. The combination of 50% nanocrystalline cellulose and xylan showed the lowest oxygen transmission rate of 0.139 cm³/m²-day). Figures 5 and 6 show the rate of oxygen transmission rate for the control xylan and xylan reinforced with 50% nanocrystalline cellulose respectively. Control xylan films have much higher rate of oxygen transmission than xylan reinforced with nanocrystalline cellulose films.
Conclusions

Nanocrystalline cellulose was incorporated into xylan slurry to prepare high oxygen barrier biodegradable films. It appears that the high degree of crystallinity of nanocrystalline cellulose and the dense composite structure of the formed films with nanocrystalline cellulose and xylan leads to a film that has reduced transmission properties. These effects were attributed to a rigid hydrogen-bonded network of hydrogen bonded nanocrystalline cellulose to form an integrated matrix as described by percolation theory.

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References

1. L. Shen, E. Worrell and M. Patel, Biofuels, Bioprod. Biorefin. 4, 25 (2010). doi:10.1002/bbb.189
2. A. Samir, F. Alloin and A. Dufresne, Biomacromolecules 6, 612 (2005).
3. A. J. Ragauskas, C. K. Williams, B. H. Davison, G. Britovsek, J. Cairney, C. A. Eckert, W. J. Frederick, J. P. Hallett, D. J. Leak, C. L. Liotta, J. R. Mielenz, R. Murphy, R. Templer and T. Tschaplinski, Science 311, 484 (2006). doi:10.1126/science.1114736
4. L. Petersson and K. Oksman, Compos. Sci. Technol. 66, 2187 (2006). doi:10.1016/j.compscitech.2005.12.010
5. J. M. Krochta, E. A. Baldwin and M. O. Nisperos-Carriedo, Lancaster: Technomic (1994).
6. J. Hartman, A. C. Albertsson, M. S. Lindblad and J. Sjöberg, J. Appl. Polym. Sci. 100, 2985 (2006). doi:10.1002/app.22958
7. M. Gröndahl, E. Eriksson and P. Gatenholm, Biomacromolecules 5, 1528 (2004). doi:10.1021/bm04925n
8. P. Dole, C. Joly, E. Espuche, I. Alric and N. Gontard, Carbohydr. Polym. 58, 335 (2004). doi:10.1016/j.carbpol.2004.08.002
9. B. L. Butler, P. J. Vergano, R. F. Testin, J. M. Bunn and J. L. Wiles, J. Food Sci. 61, 953 (1996). doi:10.1111/j.1365-2621.1996.tb10909.x
10. I. Arvanitoyannis and C. G. Biliaderis, Carbohydr. Polym. 38, 47 (1999). doi:10.1016/S0144-8617(98)00087-3
11. A. W. Rindlav, M. Stading, A. M. Hermansson and P. Gatenholm, Carbohydr. Polym. 36, 217 (1998). doi:10.1016/S0144-8617(98)00025-3
12. P. Linder, R. Bergman, A. Bodin and P. Gatenholm, Langmuir 19, 5072 (2003). doi:10.1021/la0341355
13. K. S. Mikkonen, S. Heikkinen, A. Soovre, M. Peura, R. Serimaa, R. A. Talja, J. Appl. Polym. Sci. 114, 457 (2009). doi:10.1002/app.30513
14. U. Edlund, Y. Z. Ryberg and A. C. Albertsson, Biomacromolecules 11, 2532 (2010). doi:10.1021/bm100767g
15. M. A. S. A. Samir, F. Alloin and A. Dufresne, Biomacromolecules 6, 612 (2005).
16. Kvien, J. Sugiyama, M. Votruba and K. Oksman, J Mater. Sci. 42, 8163 (2007). doi:10.1007/s10853-007-1699-2
17. L. Petersson, I. Kvien and K. Oksman, Composites Sci. Technol. 67, 2535 (2007). doi:10.1016/j.compscitech.2006.12.012
18. X. D. Cao, H. Dong and C. M. Li, Biomacromolecules 8, 899 (2007). doi:10.1021/bm0610368
19. S. Harbaugh, N. K. Loughnane, M. Davidson, L. Narayanan, S. Trott, Y. G. Chushak and M. O. Stone, Biomacromolecules 6, 1055(2005). doi:10.1021/bm049291k
20. M. Lagaron, R. Catala and R. Gavaa, Mater. Sci. Technol. 20, 1 (2004). doi:10.1179/026708304225010442
21. M. M. De Souza Lima and R. Boström, Carbohydr. Polym. 58, 335 (2004). doi:10.1016/j.carbpol.200300268
22. O. Mielenz, R. F. Testin, J. M. Bunn and J. Hallett, A. Ebringerova and T. Heinze, Macromol. Rapid Commun. 25, 771 (2004). doi:10.1002/marc.200300268
23. S. Mikkonen, A. P. Mathew, K. Pirkkalainen, R. Serimaa, C. Xu, S. Willför, K. Oksman and M. Tenkanen, Carbohydr. Polym. 17, 69 (2009). doi:10.1007/s10570-009-9380-3
24. S. Mikkonen, M. I. Heikkila, H. Helén, L. Hyvönen and M. Tenkanen, Carbohydr. Polym. 79, 1107 (2010). doi:10.1016/j.carbpol.2009.10.049
25. P. Coughlan and G. P. Hazlewood, Hemicellulose and Hemicelluloses (Eds). Portland Press Ltd, NC, U.S.A 1993.
26. A. Ebingerova and T. Heinze, Macromol. Rapid Commun. 21, 542 (2000). doi:10.1002/1521-3972(20000601)21:9<542::AID-MARC542>3.0.CO;2-7
27. I. Gabrielli and P. Gatenholm, J. Appl. Polym. Sci. 69, 1661 (1998). doi:10.1002/(SICI)1097-4628(19980822)69:8<1661::AID-APP19>3.0.CO;2-X
28. A. Saxena, T. Elder, P. Shaobo and A. J. Ragauskas, Composites Part B: 40, 8 (2009).
29. A. Saxena and A. J. Ragauskas, Carbohydr. Polym. 78, 357 (2009). doi:10.1016/j.carbpol.2009.03.039
30. Y. Pu, J. Zhang, T. Elder, Y. Deng, P. Gatenholm and A. J. Ragauskas, Composites Part B: Eng. 38, 360 (2007). doi:10.1016/j.compositesb.2006.07.008
31. S. Katz, R. P. Beatson and A. M. Scallan, Sven. Papperstidn 87, 48 (1984).
32. ASTM, Standard test method for oxygen transmission rate through plastic film and sheeting using a coulometric sensor, designation D 3985-9, in: Annual Book of ASTM Standards, American Society for Testing and Materials, 1995.
33. R. T. Parry, Principles and applications of modified atmosphere packaging of foods, Blackie Academic & Professional, England, 1 (1993).
34. Syverud and P.Stenius, Cellulose 16, 75 (2009). doi:10.1007/s10570-008-9244-2
35. A. Ebringerova and T. Heinze, Macromol. Rapid Commun. 21, 542 (2000). doi:10.1002/1521-3927(20000601)21:9<542::AID-MARC542>3.0.CO;2-7
36. I. Gabrielli and P. Gatenholm, J. Appl. Polym. Sci. 69, 1661 (1998). doi:10.1002/(SICI)1097-4628(19980822)69:8<1661::AID-APP19>3.0.CO;2-X
37. A. Saxena, T. Elder, P. Shaobo and A. J. Ragauskas, Composites Part B: 40, 8 (2009).
38. A. Saxena and A. J. Ragauskas, Carbohydr. Polym. 78, 357 (2009). doi:10.1016/j.carbpol.2009.03.039
39. Y. Pu, J. Zhang, T. Elder, Y. Deng, P. Gatenholm and A. J. Ragauskas, Composites Part B: Engineering 38, 360 (2007). doi:10.1016/j.compositesb.2006.07.008
40. S. Katz, R. P. Beatson and A. M. Scallan, Sven. Papperstidn 87, 48 (1984).
41. ASTM, Standard test method for oxygen transmission rate through plastic film and sheeting using a coulometric sensor, designation D 3985-9, in: Annual Book of ASTM Standards, American Society for Testing and Materials, 1995.
42. R. T. Parry, Principles and applications of modified atmosphere packaging of foods. Blackie Academic & Professional, England, 1 (1993).
43. Syverud and P.Stenius, Cellulose 16, 75 (2009). doi:10.1007/s10570-008-9244-2