

§ Supplementary Material §

Cooperative effects of cellulose nanocrystals and sepiolite when combined on ionic liquid plasticised chitosan materials

Pei Chen^{a,b}, Fengwei Xie^{b,*†}, Fengzai Tang^c, Tony McNally^{b,**}

^a College of Food Science, South China Agricultural University, Guangzhou, Guangdong 510642, China

^b International Institute for Nanocomposites Manufacturing (IINM), WMG, University of Warwick, Coventry CV4 7AL, United Kingdom

^c WMG, University of Warwick, Coventry CV4 7AL, United Kingdom

* Corresponding author. Email addresses: d.xie.2@warwick.ac.uk, fwhsieh@gmail.com (F. Xie)

** Corresponding author. Email address: t.mcnnally@warwick.ac.uk (T. McNally)

† This author leads the research.

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1 Figures

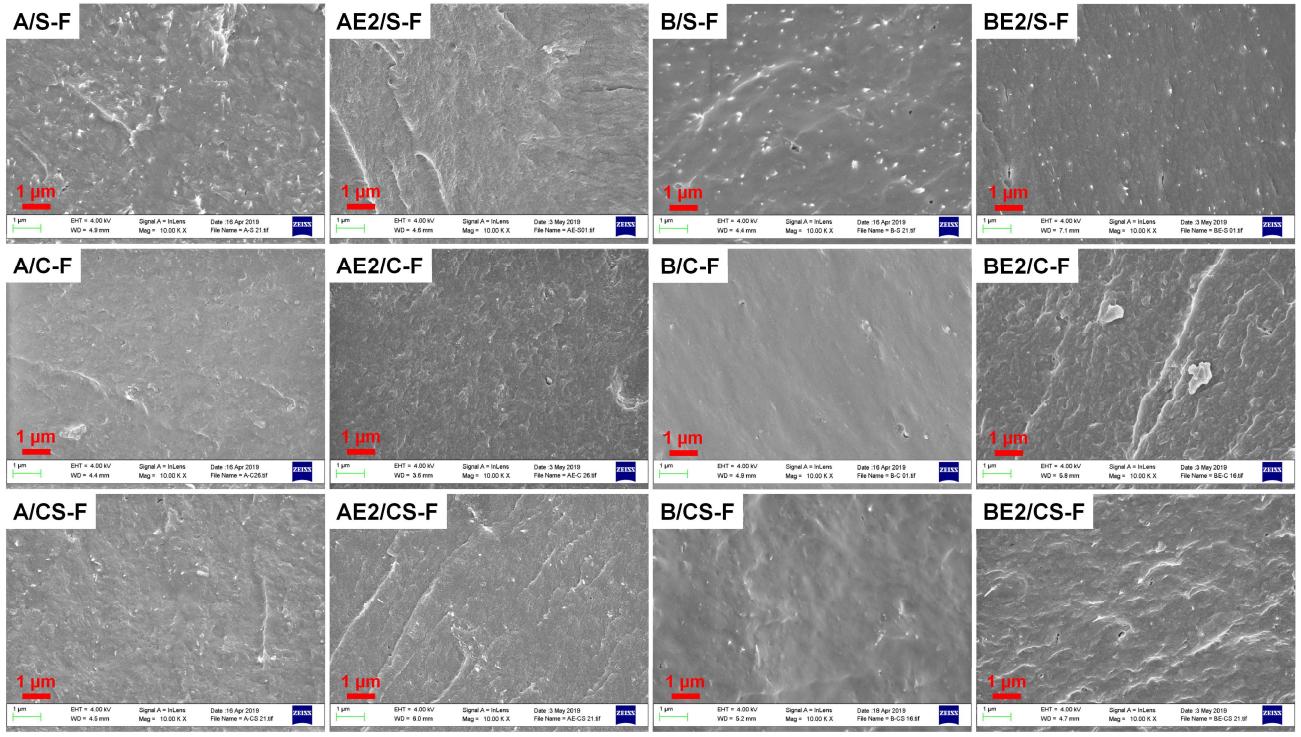


Figure S1. Scanning electron microscopy (SEM) images of the different bionanocomposite films.

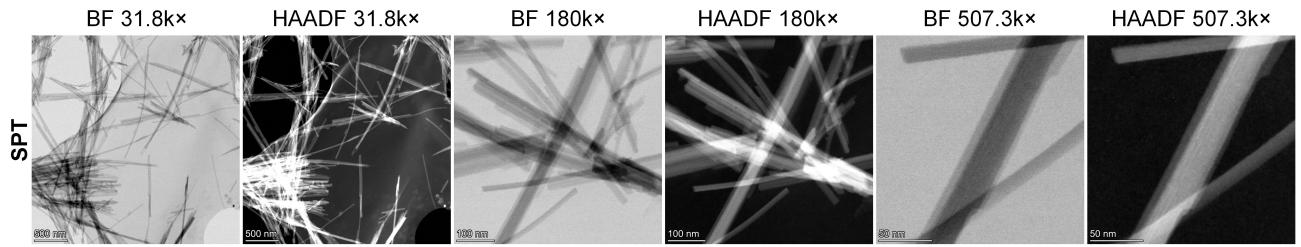


Figure S2. Scanning transmission electron microscopy (STEM) images of sepiolite (SPT). BF, bright field; HAADF, High-angle annular dark-field.

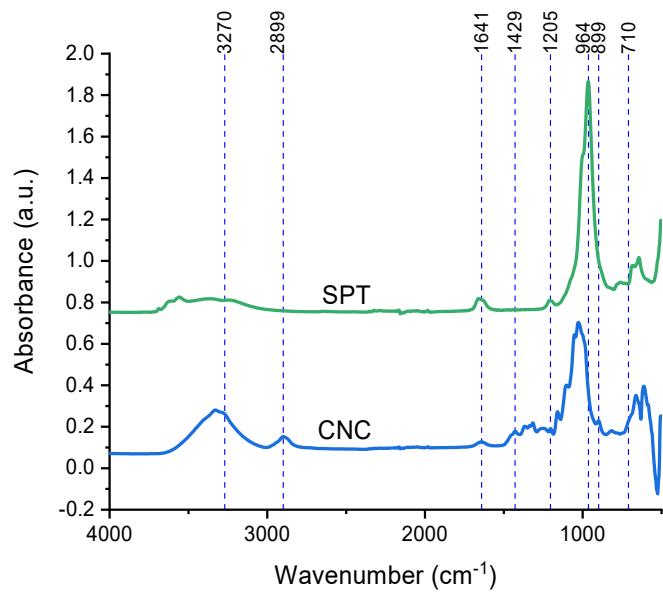


Figure S3. Fourier-transform infrared (FTIR) spectrum of cellulose nanocrystals (CNCs) and sepiolite (SPT).

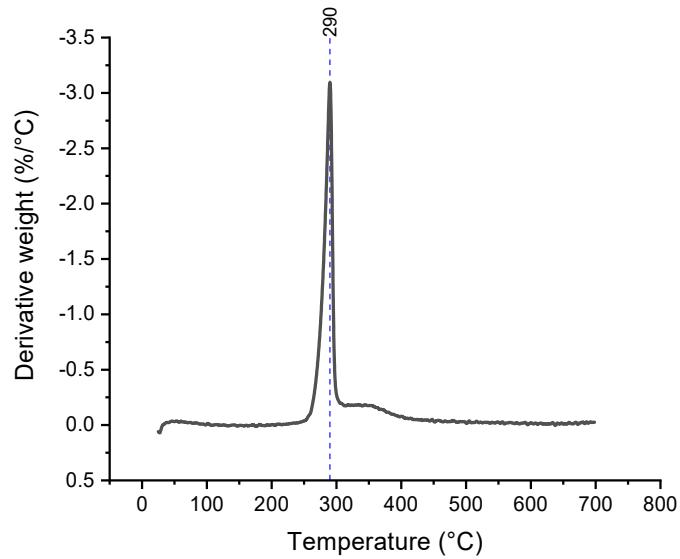


Figure S4. Derivative weight vs. temperature curve measured by thermogravimetric analysis (TGA) for cellulose nanocrystals (CNCs).

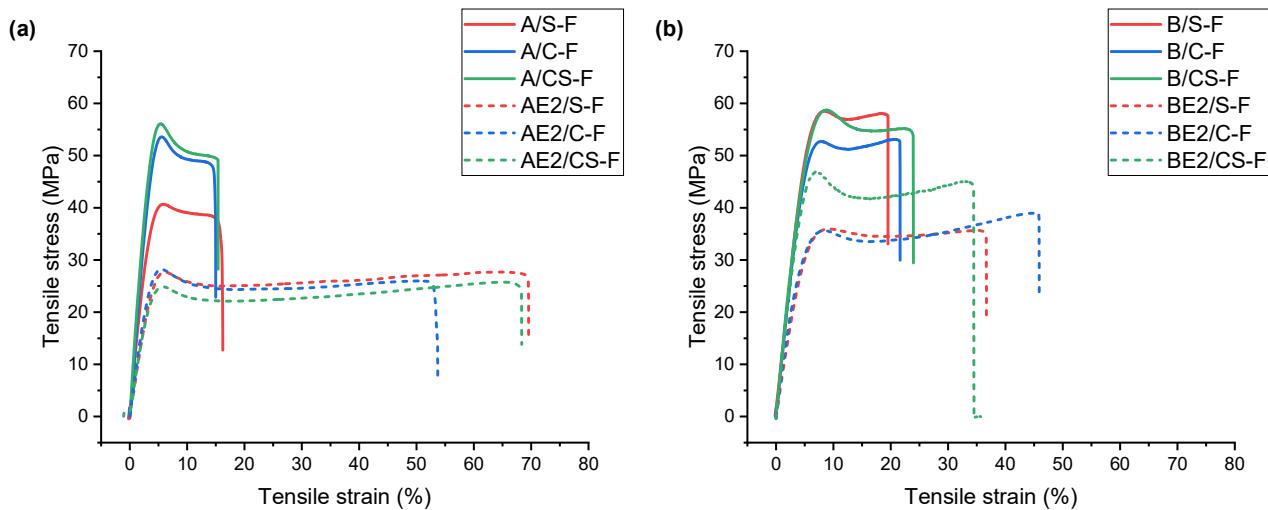


Figure S5. Representative stress–strain curves under tensile testing for different biopolymer composite films: a) chitosan matrix; and b) chitosan/CMC matrix.

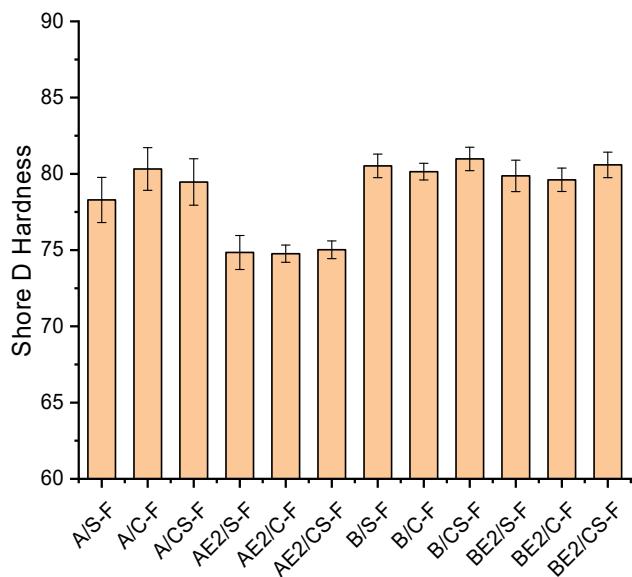


Figure S6. Shore D hardness values of the different biocomposite films.

2 Notes to figures

Figure S3 shows that for cellulose nanocrystals (CNCs), the broad bands in the 3680–3000 cm⁻¹ region are due to O—H stretching vibrations and the peak at 2899 cm⁻¹ corresponds to C—H stretching vibrations. The 1430 cm⁻¹ band is assigned to C6—CH₂ bending [1]. The band at 899 cm⁻¹ is attributed to C—O stretching and C—H vibration in cellulose [2]. There is a sulphate peak at 1205 cm⁻¹ resulting from the esterification reaction, suggesting this CNCs was obtained by acid hydrolysis. The O—H stretching at 3270 cm⁻¹ and the out-of-plane bending at 710 cm⁻¹ indicates this CNCs is of the cellulose I β type [1]. For sepiolite (SPT), the band at 964 cm⁻¹ can be assigned to Si—O stretching [3].

References

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3. McKeown, D.A.; Post, J.E.; Etz, E.S. Vibrational analysis of palygorskite and sepiolite. *Clays Clay Miner.* **2002**, *50*, 667–680, doi:10.1346/000986002320679549.