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Supporting Information

RAFT-mediated Pickering Emulsion Polymerization with Cellulose Nanocrystals Graft Random Copolymer as Stabilizer

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Figure S1. TEM image of CNCs.

Sample	Composition (wt%)			[N]/[O]	[C]/[O]
	С	0	N	_ [1,],[0]	[_],[_]
CNCs	50.69	40.31			1.68
Poly(AN-r-BA)	69.70	19.10	11.20	0.67	4.88
Poly(VT-r-BA)	62.44	11.30	26.06	2.63	7.37
Poly(VT-r-BA)-g-	56 95	20.15	22 90	1 30	3 76
CNCs	50.75	20.13	22.70	1.50	5.70

Table S1. EDS data of CNCs, Poly(AN-*r*-BA), Poly(VT-*r*-BA) and Poly(VT-*r*-BA)-*g*-CNCs.















Figure S2. EDS picture of CNCs (a), Poly(AN-*r*-BA) (b), Poly(VT-*r*-BA) (c) and Poly(VT-*r*-BA)-*g*-CNCs (d).

FT-IR spectra of poly(AN-*r*-BA), poly(VT-*r*-BA), CNCs and poly(VT-*r*-BA)-*g*-CNCs. Poly(AN-*r*-BA), poly(VT-*r*-BA), CNCs and poly(VT-*r*-nBA)-*g*-CNCs was further characterized by IR measurement. The click chemistry of poly(AN-*r*-BA) with sodium azide lead to poly(VT-*co*-BA) containing tetrazolium group. As shown in figure 5, the FTIR spectrums of poly(AN-*r*-BA) and 5 poly(VT-*r*-BA)had their obvious IR characters. In spectrogram line of poly(AN-*r*-BA), the peak absorption at 2240 cm⁻¹ was in accordance with C≡N stretching vibration, absent in the IR spectrum of poly(VT-*r*-BA) due to C≡N groups converted into tetrazolium group. Moreover, there was additional peak at 1545 cm⁻¹ and 1610 cm⁻¹ associated with tetrazolium group, appears with the IR spectrum of poly(VT-*r*-BA). Therefore, these phenomenon manifest successfully preparation of poly(VT-*r*-BA) and CNCs. In IR spectra of CNCs, all characteristic was found. Meanwhile, the peak at 1723 cm⁻¹ attributed to C=O stretching vibration derived from poly(VT-*r*-BA) and 1059 cm⁻¹ from CNCs were observed in IR spectrum line poly(VT-*r*-BA)-CNCs at the same time. On the basis of analying these character, it clearly demonstrated that the successful synthesis of poly(VT-*r*-nBA)-*g*-CNCs.



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Figure S3. The FT-IR spectrogram of poly(AN-*r*-BA), poly(VT-*r*-BA), CNCs and poly(VT-*r*-BA)-*g*-CNCs.