Supporting Information

Glow discharge electrolysis plasma induced synthesis of cellulose-based ionic

hydrogels and their multiple response behaviors

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Figure S1. Photographs of wet hydrogel (a), dried hydrogels (b), and hydrogel after swelling equilibrium (c).

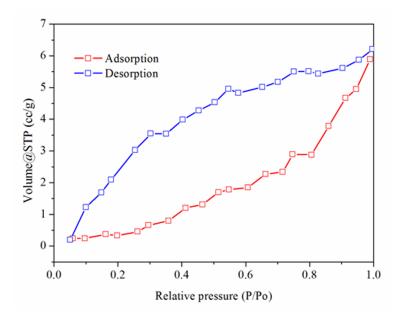


Figure S2. Nitrogen sorption isotherm of cellulose-based ionic hydrogel (the discharge voltage and discharge time is 570 V and 90 s, respectively).

Figure S2 shows the N_2 sorption isotherm of the freezed dried cellulose-based ionic hydrogel after absorbing water sufficiently. The sample was measured using automated surface area and pore size analyzer (Quantachrome Autosorb-1 MP). The surface area was 2.131 m²/g.

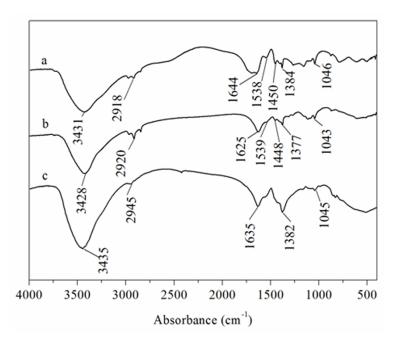


Figure S3. FT-IR of cellulose-based ionic hydrogels after adsorbing heavy metal ions (a) Na^+ , (b) Zn^{2+} , and (c) Fe^{3+} .

Figure S3 shows the infrared spectrogram of cellulose-based ionic hydrogels after adsorbing heavy metal ions. As the Na⁺, Zn²⁺, Fe³⁺ ions were adsorbed by cellulose-based ionic hydrogels, the peak at 1722 cm⁻¹ (C=O stretching vibration) (Fig. 7b) was disappeared completely implying that complexation was possible and new peaks appeared at 1644, 1625, 1635 cm⁻¹ on curve (a), (b) and (c), respectively.¹ In addition, the peaks at 1543 and 1456 cm⁻¹ (the stretching vibration of COO⁻) (Fig. 7b) were shifted to lower wavenumbers after adsorbing Na⁺, Zn²⁺ metal ions, or even disappeared after adsorbing Fe³⁺. Besides, the intensity of the peaks of Zn²⁺ was weaker than Na⁺. All results indicated that oxygen atoms in cellulose-based ionic hydrogels donated unshared electron pairs to the metal ions to form coordinate-covalent bonds and the coordination between Fe³⁺/Zn²⁺ and carboxyl group was obvious stronger than that Na⁺.²⁻⁴

References:

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