

Electrochemically induced reversible formation of carboxymethyl chitin hydrogel and tunable protein release

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1. The amount of iron in the hydrogel

The total amount of iron in the hydrogel was determined by method of phenanthroline spectrophotometric^{1, 2}. Briefly, the electrodeposited hydrogel on the iron electrode was dissolved in 10 mL 0.1 M Na₂SO₄ containing ascorbic acid (10 mM). Then, 10 mL of the resulted solution, 2.5 mL of hydroxylamine hydrochloride (1%), 5.0 mL of HAc/NaAc (pH=4.6) and 5.0 mL of phenanthroline (0.1%) were added into a 50 mL volumetric flask. Ultrapure water was added to the scale. The volumetric flask was shaken to mix the solution. The final solution was put still for 10 min before test. The Fe³⁺ concentration in the solution was recorded on a UNICO UV-2000 Spectrophotometer at 510 nm. The solution without Fe³⁺ was used as a control. Ferrous ammonium sulfate was used for calibrating the Fe³⁺ analysis.

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