

Electronic Supporting Information

Fabrication and characterization of ferritin-chitosan-lutein shell-core nanocomposites and lutein stability and release evaluation *in vitro*

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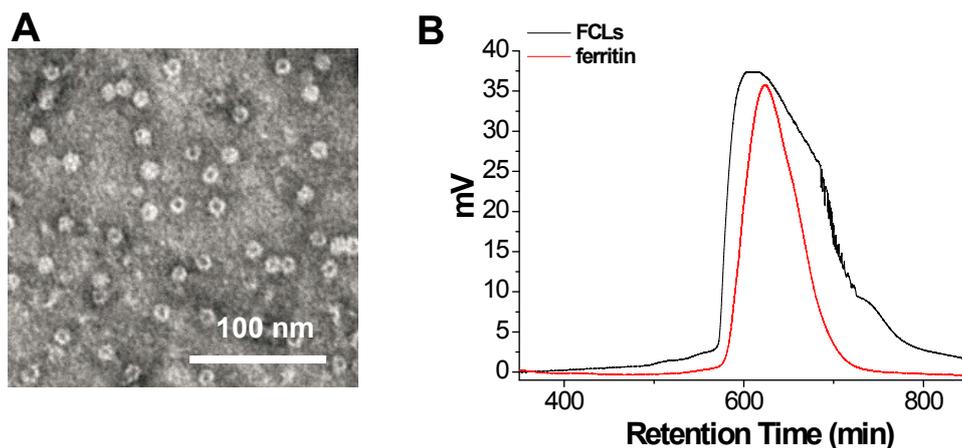


Fig. S1 (A) TEM of the apoSSF (2.0 μ M) simply mixed with lutein (300 μ M) followed by dialysis by Mops buffer (20 mM, pH 6.0) with two buffer changes (every 8.0 h intervals); (B) Size exclusion chromatography of ferritin and FCLs at pH 6.0.

Procedure and condition of Size exclusion chromatography experiment

Size exclusion chromatography was performed by a purificating column (1.5 \times 120 cm) filling with Sephacryl S-300 resin (GE. Inc., USA) consisting of an UV detector (280 nm) (4.6 \times 250mm, 5 μ m). Samples were eluted by the use of a mobile phase consisting of 50 mM Mops, 0.15 M NaCl. The injection volume was 5 mL, and the flow rate of the mobile phase was 0.4 mL/min. The flowed samples were collected by 10 mL tubes with a rate of 10 min/tubes.

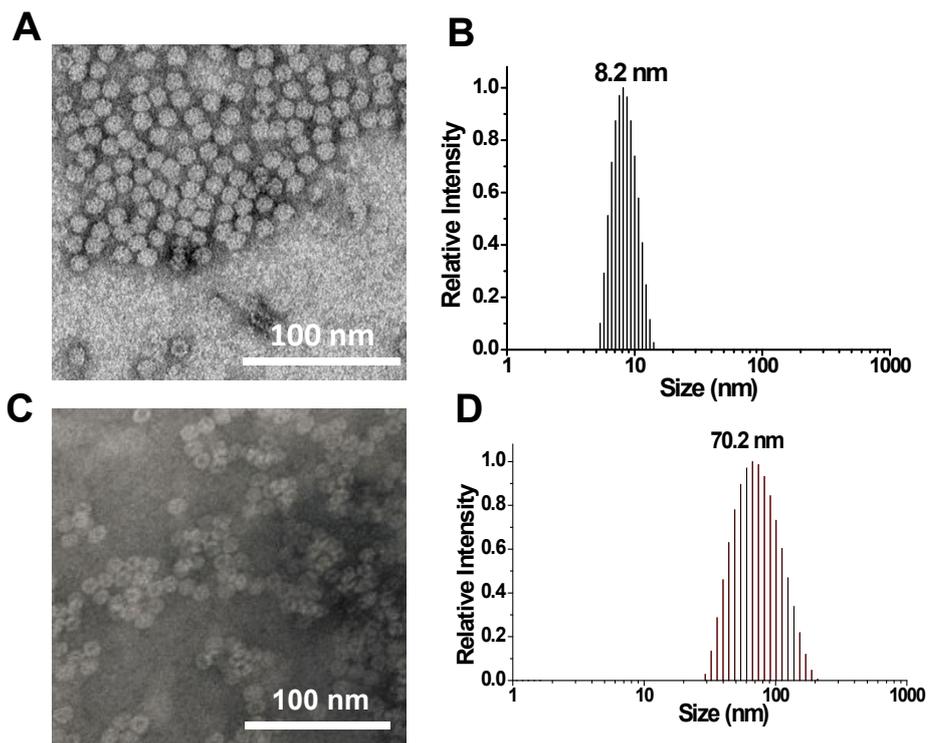


Fig. S2 (A) TEM of ferritin-lutein complex in the presence of 17% ethanol; (B) DLS of ferritin-lutein complex in the presence of 17% ethanol; (C) TEM of ferritin-lutein complex in the presence of 30% ethanol; (D) DLS of ferritin-lutein complex in the presence of 30% ethanol.

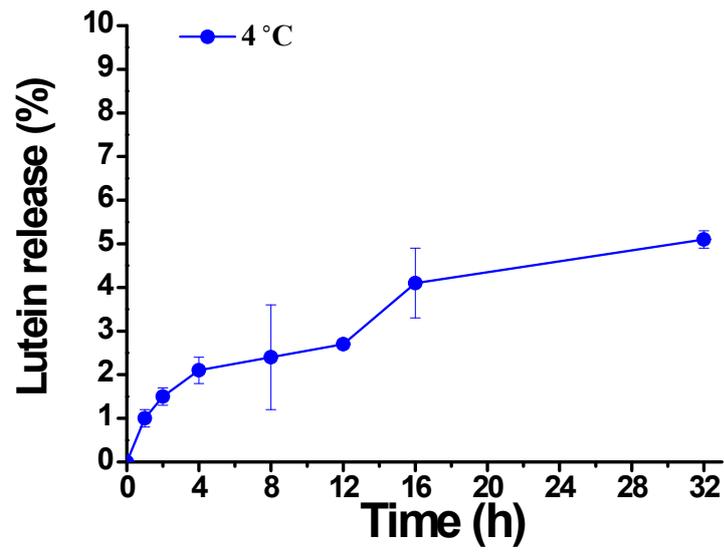


Fig. S3 Kinetics of lutein release from FCLs at different dialysis time.