

Facile Strategy to Enable Nanoparticles for Simultaneous Phase Transfer, Folate Receptor Targeting, and Cisplatin Delivery

Mochamad Zakki Fahmi and Jia-Yaw Chang*

Department of Chemical Engineering, National Taiwan University of Science and Technology, Taipei 106, Taiwan, Republic of China

*Corresponding Author: jychang@mail.ntust.edu.tw

Supporting Information:

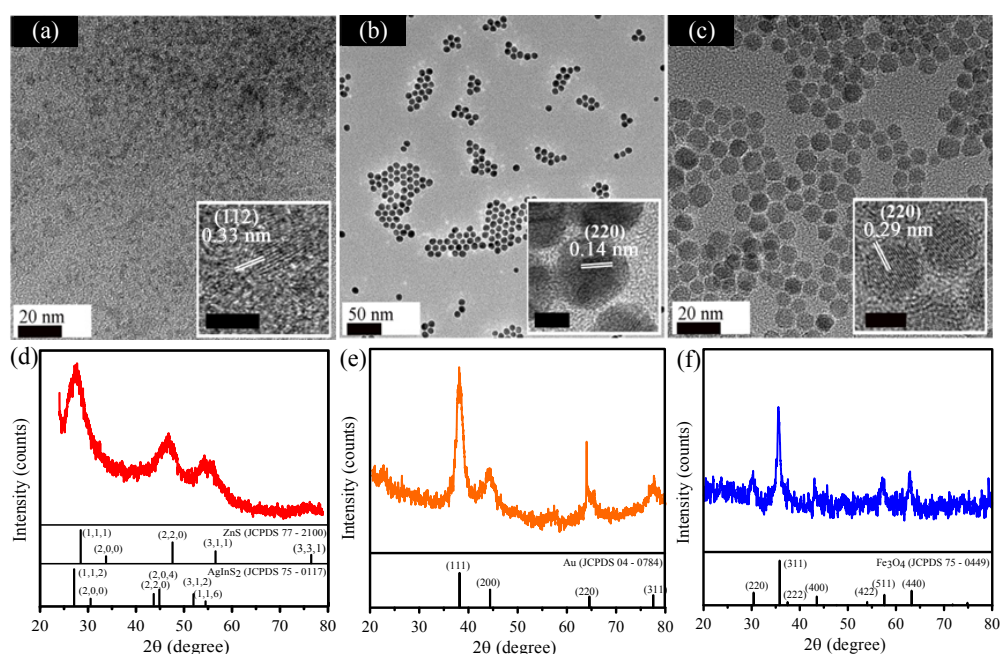


Figure S1. TEM images of (a) AQD, (b) GNP, and (c) FNP. The insets with lattice fringes show high-resolution TEM images of the corresponding nanoparticles; it can be seen that they are crystalline. The scale bar is 2 nm. XRD patterns of as-prepared AQD (d), GNP (e), and FNP (f). XRD patterns of (d) AgInS₂ (JCPDS 75-0117) and ZnS (JCPDS 77-2100); (e) Au (JCPDS 04-0784); (f) Fe₃O₄ (JCPDS 75-0449) are also shown as reference.

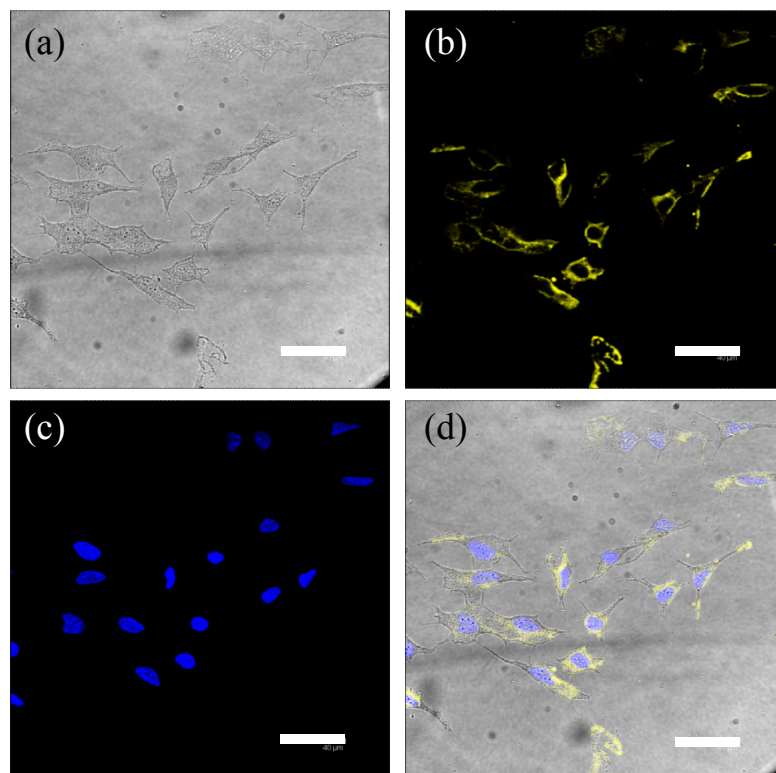


Figure S2. Confocal laser images of HeLa cells treated with AQD@folate: (a) transmission image and luminescent images containing, (b) yellow emission of AQD@folate, (c) blue emission of DAPI, and (d) an overlay of (a)–(c). Scale bars represent 40 μm .

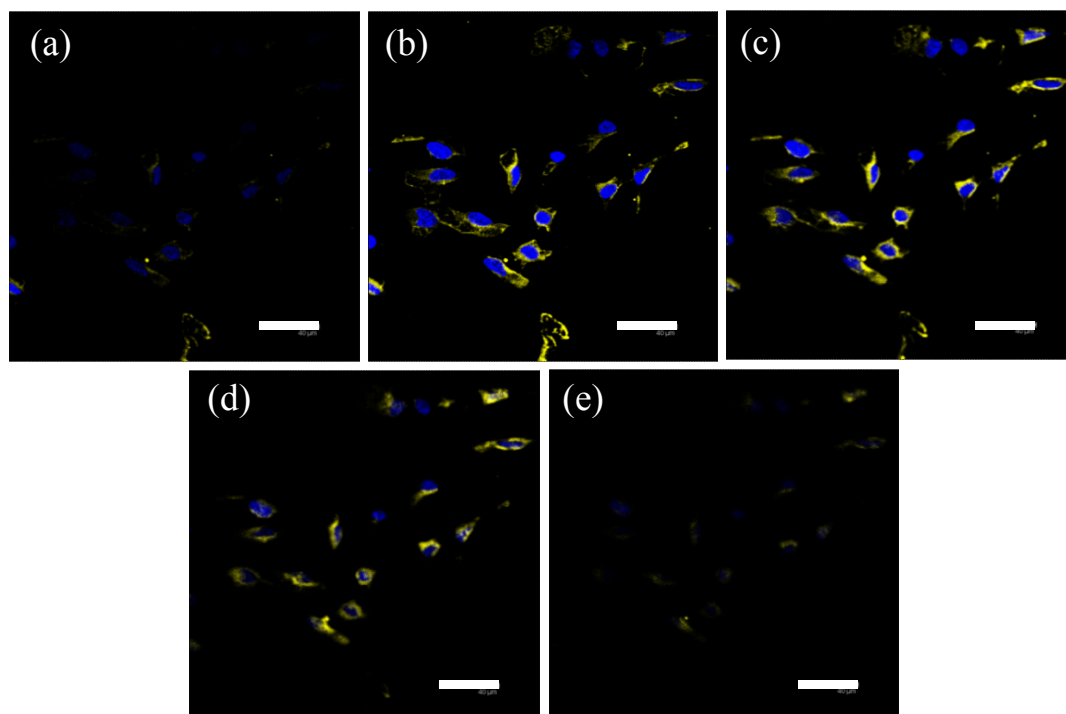


Figure S3. Confocal microscopy Z-stack sectioning of HeLa cancer cells treated with AQD@folate for 2 h, fixed with alcohol, and subsequently stained with DAPI. Slices were taken from the top (a) to the bottom (e) of the HeLa cells; consecutive images in each column are separated by 2 μm . Yellow fluorescence originates AQD@folate and blue emission at 460 nm, showing the location of the HeLa cell nuclei. Scale bars represent 40 μm .

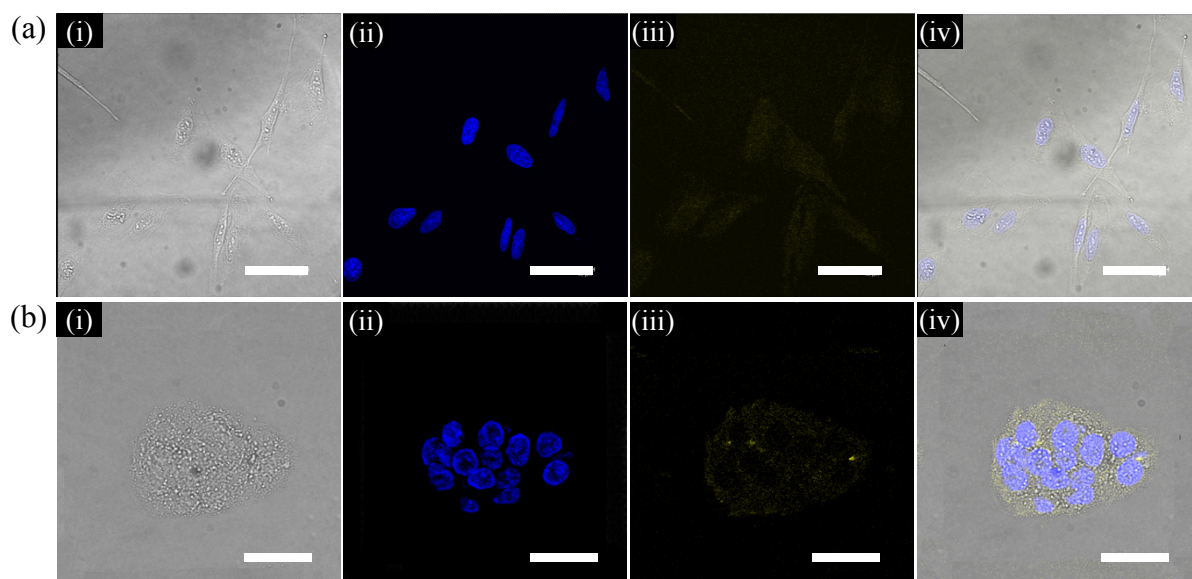


Figure S4. Confocal laser images of HeLa cells treated with AQD@OA (a) and HepG2 cell treated with AQD@folate/CDDP (b). Series of images consists : (i) transmission image and luminescent images containing (ii) blue emission of DAPI as well as (iii) yellow emission of AQD@OA, and (iv) an overlay of (i)–(iii). Scale bars represent 40 μm .

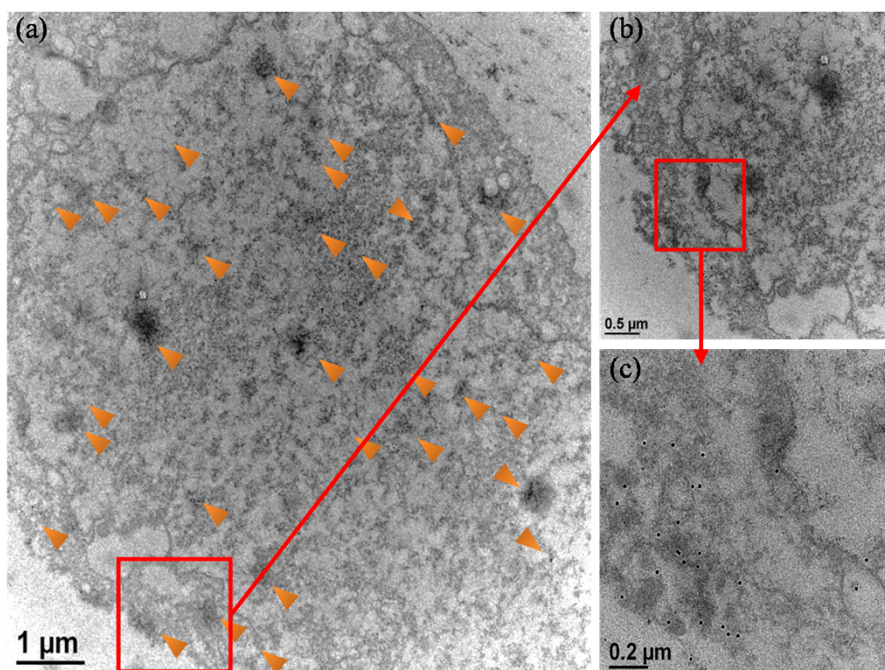


Figure S5. (a) TEM images showing the internalization of GNP@folate into HeLa cells. (b) and (c) High-magnification images of the area delimited by the square in panels (a) and (b), respectively. Brown arrows indicate the presence of GNP@folate nanoparticles.

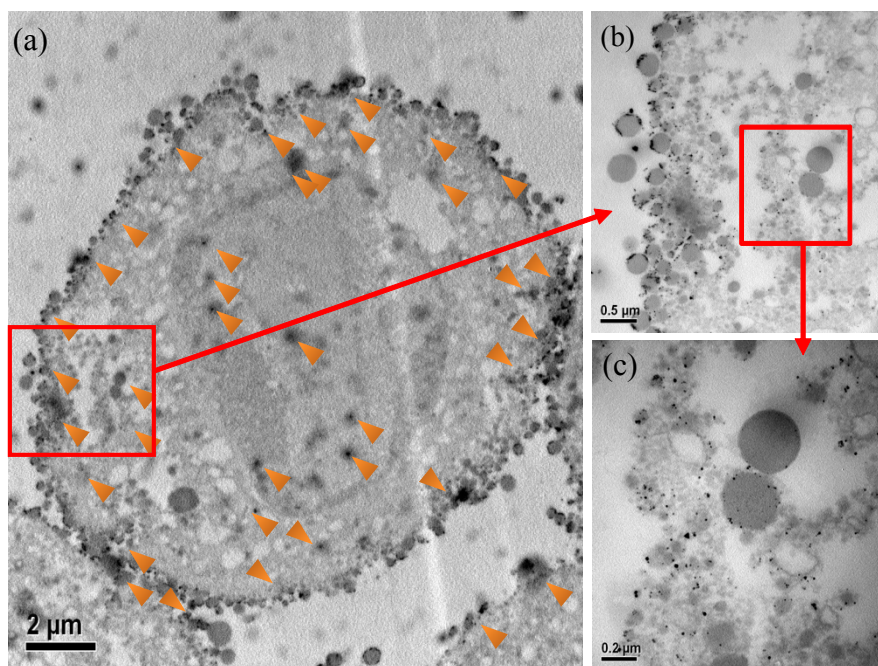


Figure S6. (a) TEM images showing the internalization of FNP@folate into HeLa cells. (b) and (c) High-magnification images of the area delimited by the square in panels (a) and (b), respectively. Brown arrows indicate the presence of FNP@folate nanoparticles.

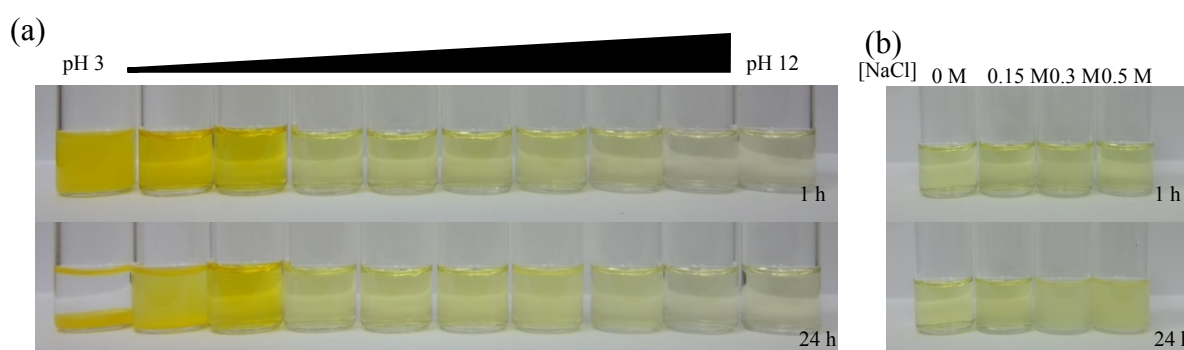


Figure S7. (a) The photograph images of AQD@folate in aqueous solutions of varying pH (from 3 to 12) at room temperature after storing for 24 h. (b) The photograph images of AQD@folate dispersed in pH 7 solutions containing different NaCl concentrations (from left to right: 0 M, 0.15 M, 0.3 M, and 0.5 M) after storing for 24 h.

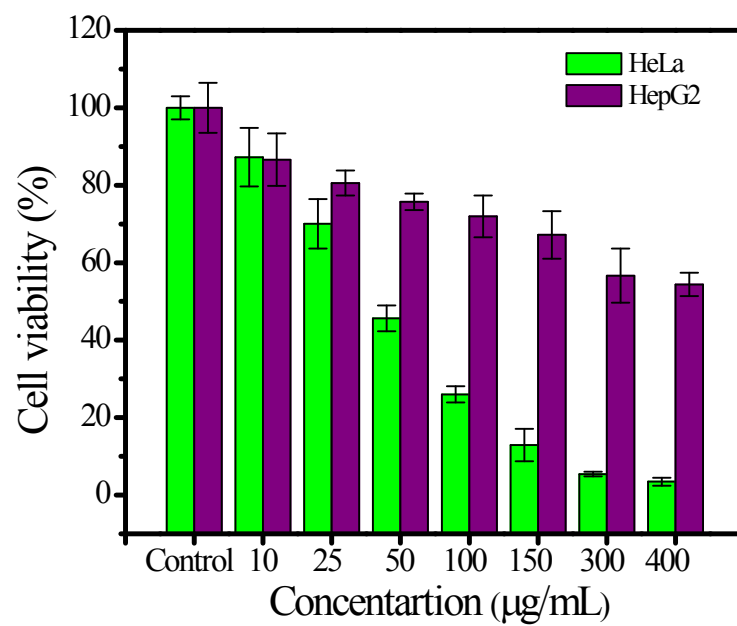


Figure S8. Comparison cell viability graphs of HeLa and HepG2 cancer cells after 24 h incubation with ACD@folate/CDDP

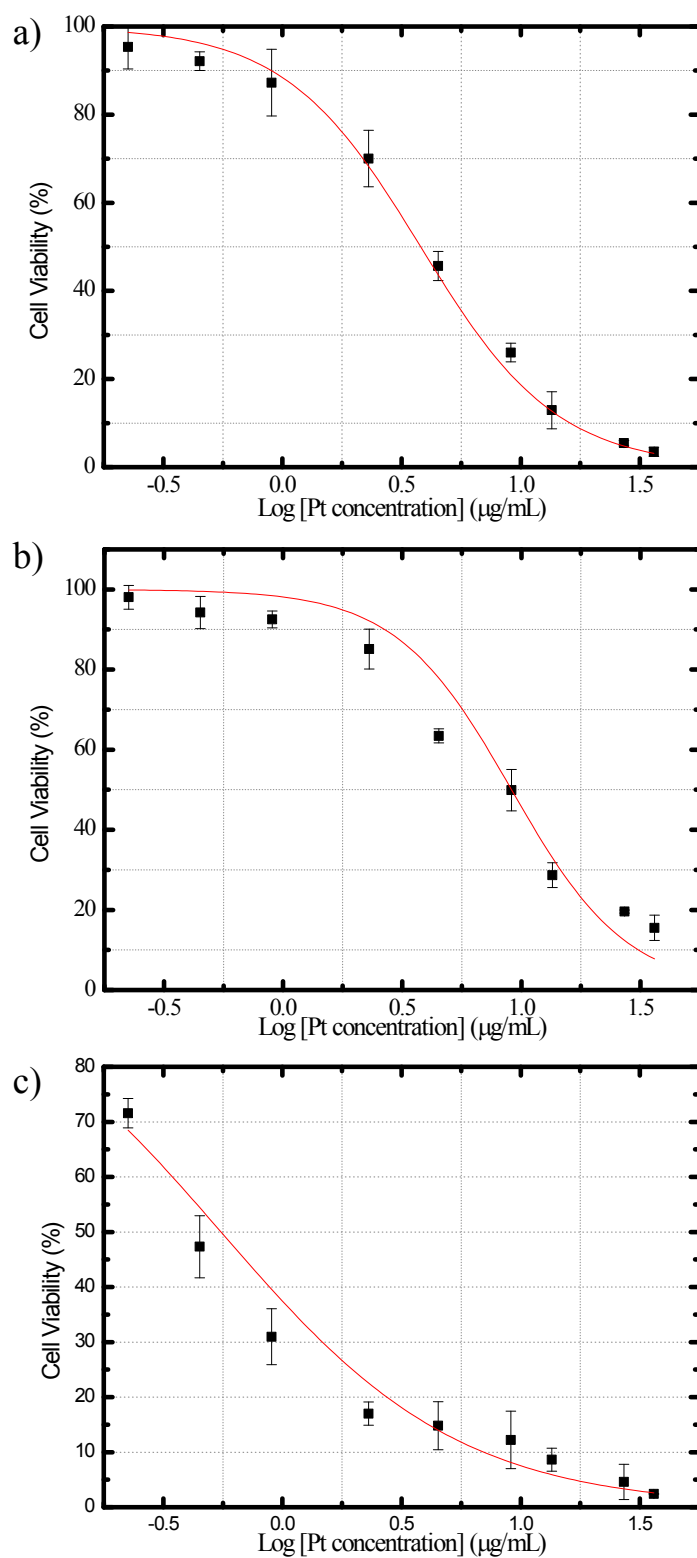


Figure S9. Cell viability graphs of HeLa cancer cells after 24 h incubation with (a) AQD@folate/CDDP, (b) AQD@CDDP, and (c) free CDDP. IC₅₀ values can be determined on red fitted curve, which showed at 0.169 µg/mL, 0.455 µg/mL, and -0.565 µg/mL, respectively, for each log [Pt concentration] or 1.477 µg/mL, 2.856 µg/mL, and 0.272 µg/mL, respectively, for each [Pt concentration].