

Supporting Information

Preparation and folic acid conjugation of fluorescent polymer nanoparticles for cancer cell targeting

Xinxin Qiang,^a Tong Wu,^a Jiangli Fan,^a Jingyun Wang,^b Fengling Song,^a Shiguo Sun,^a
Xiaojun Peng*

^a State Key Laboratory of Fine Chemicals, ^b School of Life Science and
Biotechnology, Dalian University of Technology,
2 Linggong Road, 116024 Dalian, China

* Corresponding author: pengxj@dlut.edu.cn

Contents

NMR spectral of synthesized dyes

Optical spectral of dye 4

Images of the fNPs emulsion

FT-IR spectra of NPs

Zeta potential of NPs

Cytotoxicity of FANPs

Confocal images and flow cytometer measurement

NMR spectral of synthesized dyes.

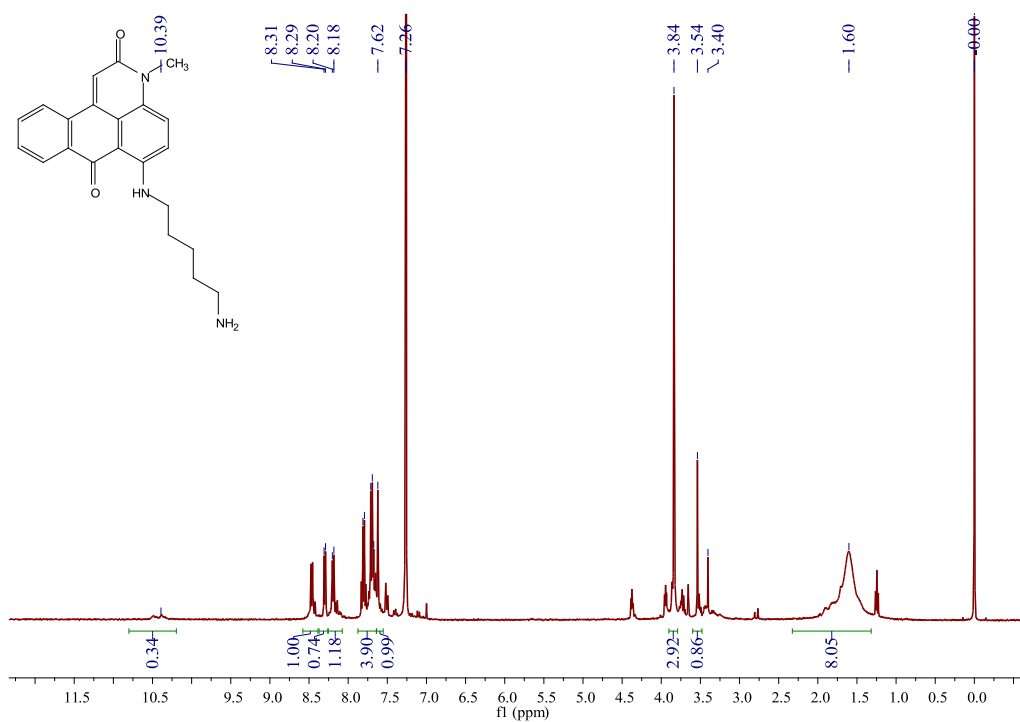
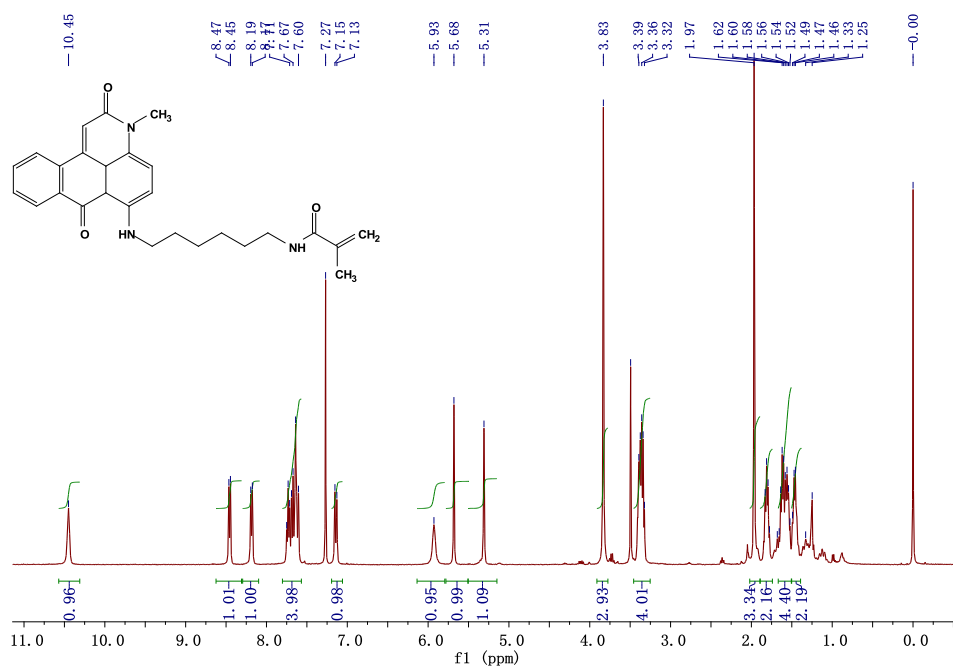


Fig. S1 ¹H-NMR spectra of dye 2.

a)



b)

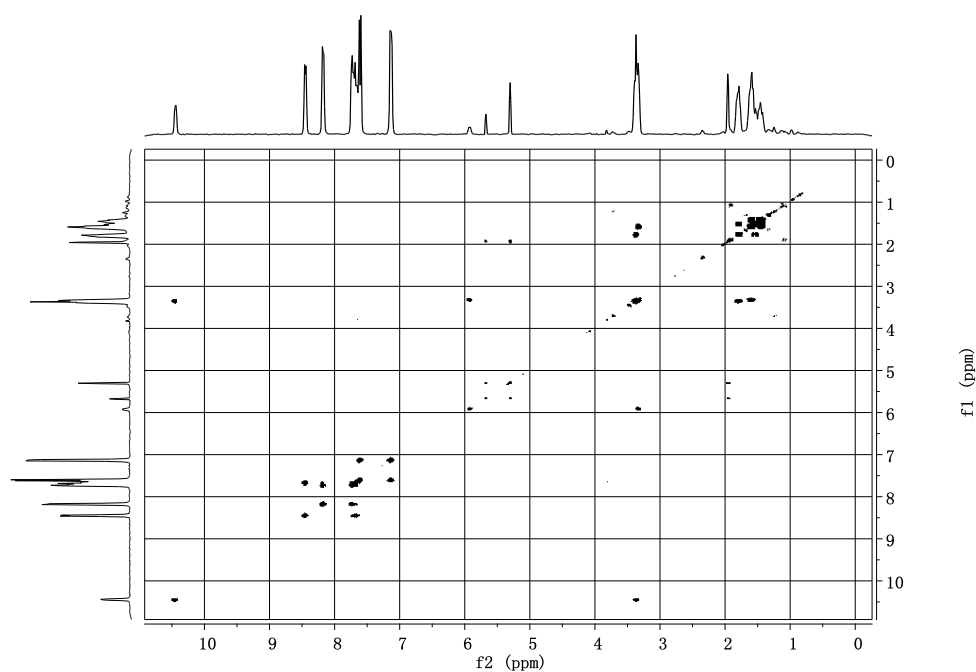
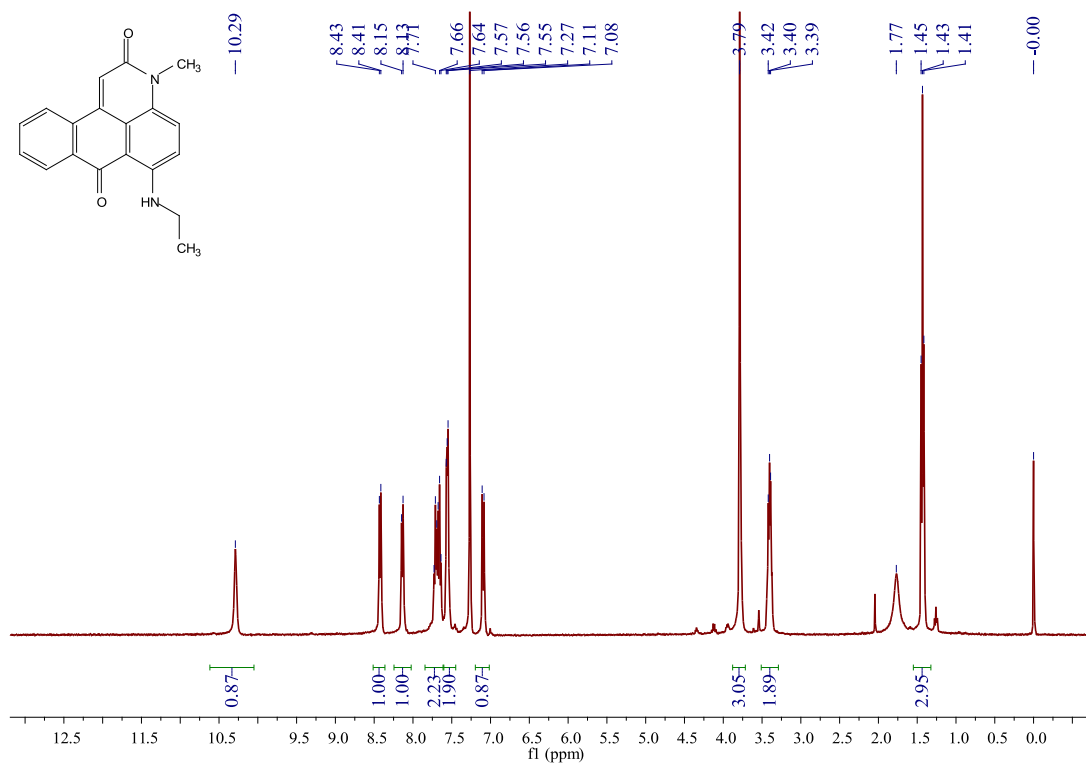


Fig. S2 NMR spectra of dye 3 a) ^1H -NMR; b) ^1H - ^1H COSY.

a)



b)

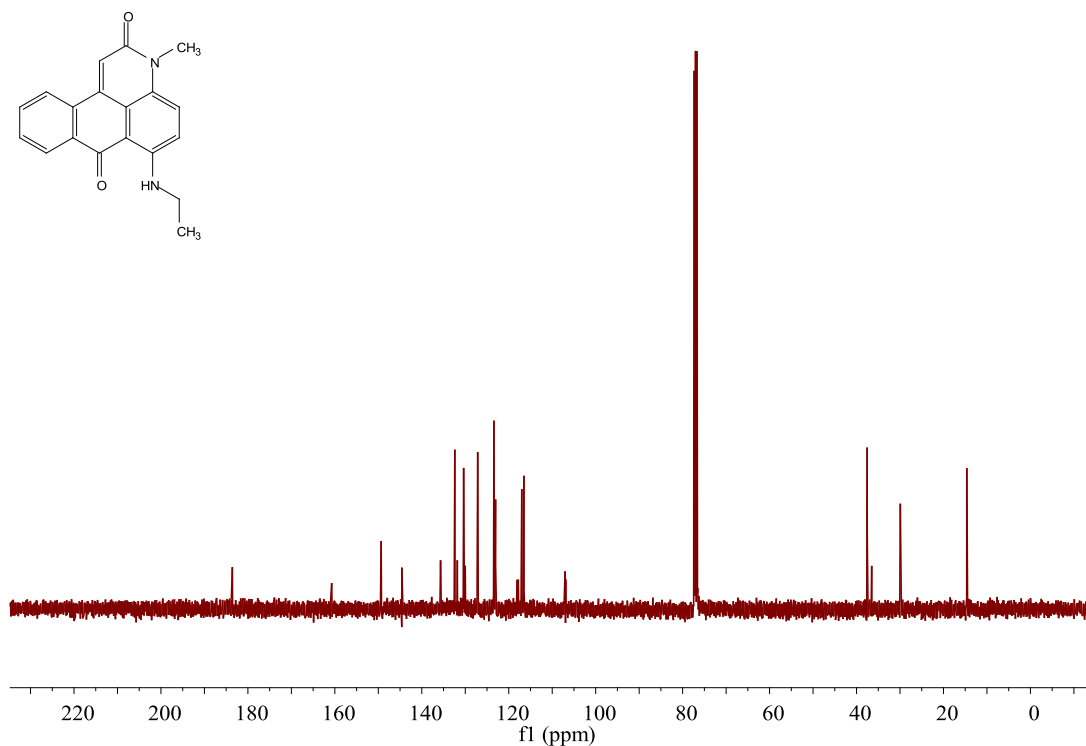


Fig. S3 NMR spectra of dye 4 a) ¹H-NMR; b) ¹³C-NMR.

Optical spectral of dye 4

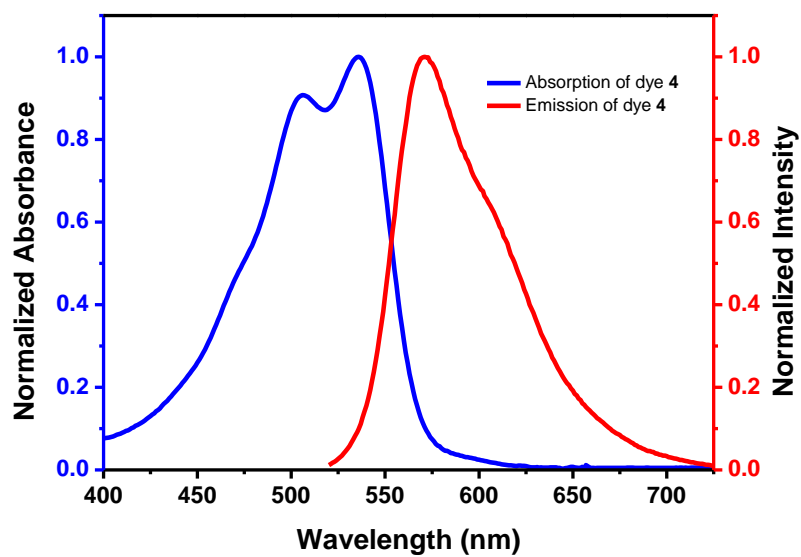


Fig. S4 Normalized absorption and emission of dye 4 in ethanol.

Images of the fNPs emulsion



Fig. S5 Images of the fNPs emulsion illuminated by a) sunlight; b) 365 nm of UV light.

FT-IR spectra of NPs

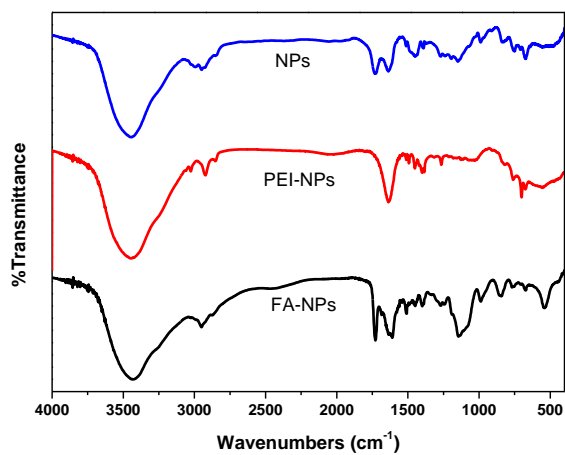


Fig. S6 FT-IR spectra of NPs.

Zeta potential of NPs

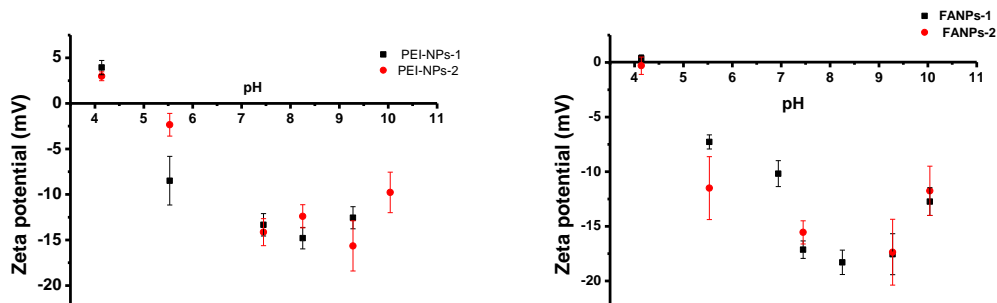


Fig. S7 Zeta potential test of PEI-NPs (right) and FANPs (left) towards pH.

Cytotoxicity of FANPs

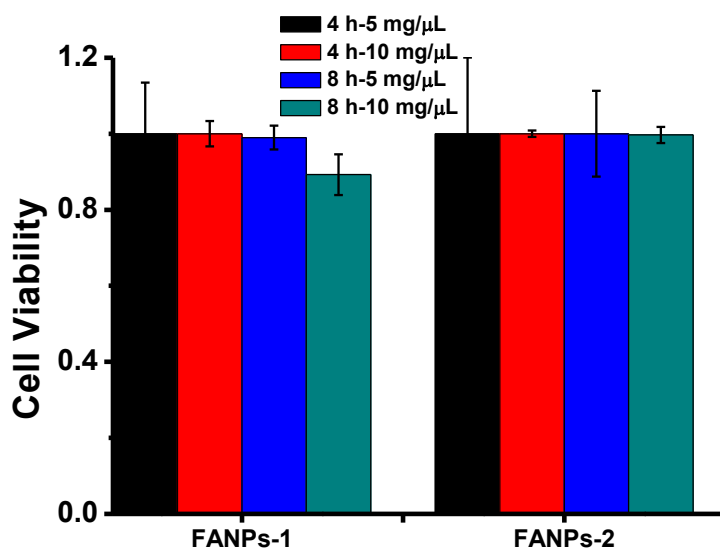


Fig. S8 Cell viability of FANPs towards HeLa cells.

Confocal images and flow cytometer measurement

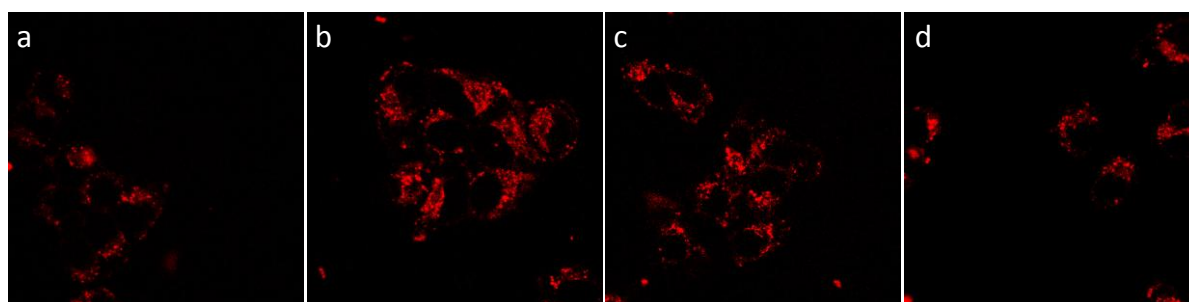


Fig. S9 Confocal images of HeLa cells incubated with FANPs (FANPs-1, FANPs-2, FANPs-3, which prepared using 2% PEI solution of a) 0.1 mL; b) 0.2 mL; c) 0.5 mL respectively) and d) large NPs.

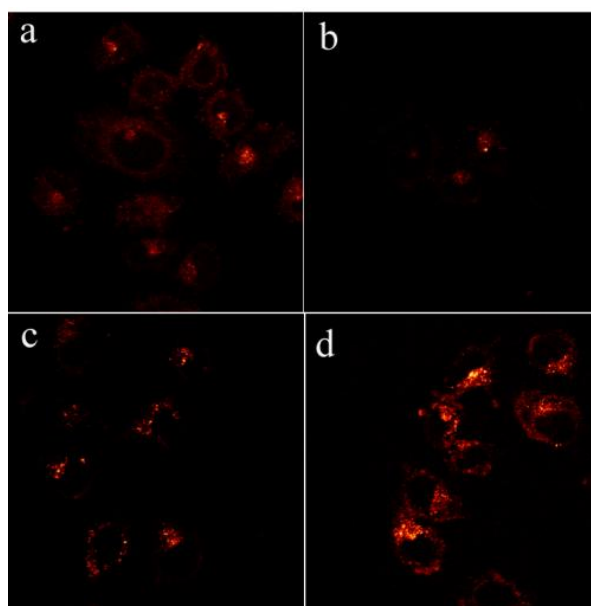


Fig. **S10** Confocal fluorescent microscopy of cancer cell lines following incubation with NPs. MCF-7 Cells were incubated with a) PEI-NPs and b) FANPs-2 for 4 h. HeLa cells were incubated with c) PEI-NPs and d) FANPs-2 for 4 h.

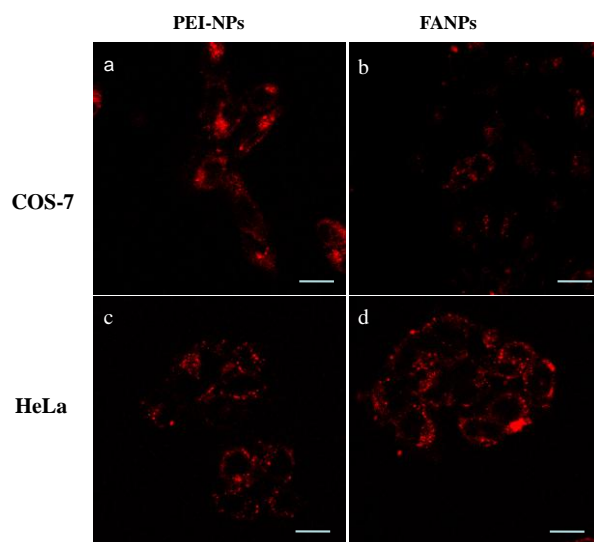


Fig. **S11** Confocal fluorescent microscopy of cancer cell lines following incubation with NPs. COS-7 Cells were incubated with a) PEI-NPs and b) FANPs-2 for 4 h. HeLa cells were incubated with c) PEI-NPs and d) FANPs-2 for 4 h. Scale bar is 20 μm .

Table **S1** Uptake ratio of HeLa and HepG2 cells towards NPs measured by flow cytometer (incubated with NPs at concentration of 10 $\mu\text{g}/\text{mL}$ for 4 h).

NPs	Cell lines	Uptake Ratio (%)
-----	------------	------------------

Control	HeLa	0.37
(no NPs)	HepG2	0.4
FANPs-1	HeLa	1.7
FANPs-2	HeLa	3.6
FANPs-2	HepG2	0.87
large NPs	HeLa	2.07

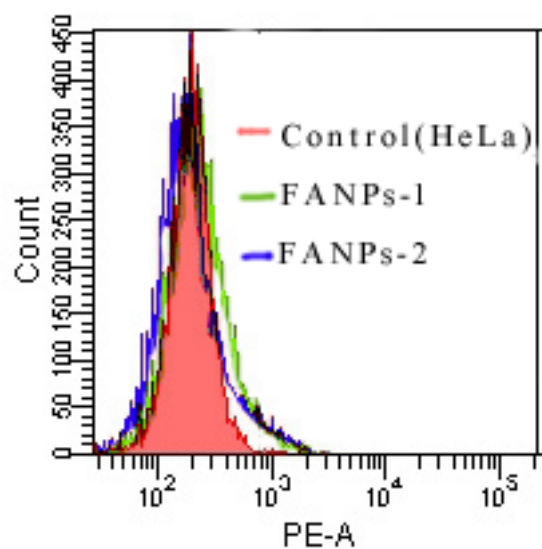


Fig. S12 Flow cytometry data showed the relative cellular uptake of HeLa cells incubated with Control (no NPs), FANPs-1 and FANPs-2 (at concentration of 10 $\mu\text{g}/\text{mL}$ for 4 h).

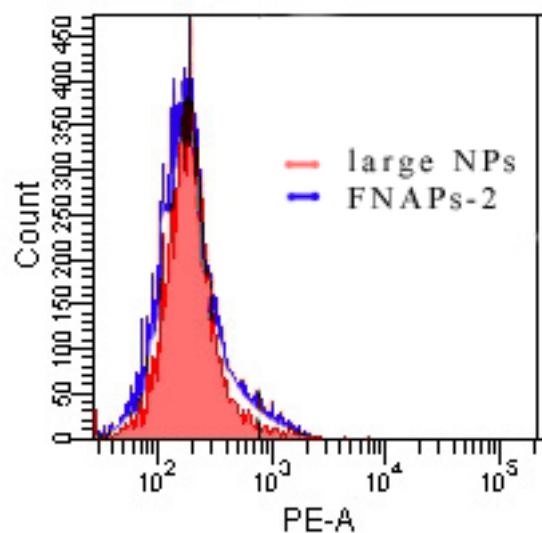


Fig. S13 Flow cytometry data showed the relative cellular uptake of HeLa cells incubated with large NPs and FANPs-2 (at concentration of 10 $\mu\text{g}/\text{mL}$ for 4 h).

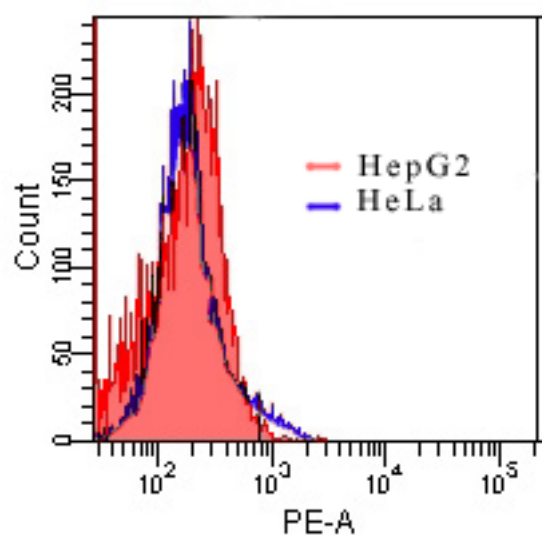


Fig. S14 Flow cytometry data showed the relative cellular uptake of HeLa and HepG2 cells incubated with FANPs-2 (at concentration of 10 $\mu\text{g}/\text{mL}$ for 4 h).