Electronic Supplementary Material (ESI) for Journal of Materials Chemistry B. This journal is © The Royal Society of Chemistry 2016

## Construction of AuNSs-AA@Fe<sub>3</sub>O<sub>4</sub>-PEI/PEG

In Figure S1-A, the TEM result showed that the form of Fe<sub>3</sub>O<sub>4</sub>-PEI@AuNSs-AA/PEG complexes was almost same as Fe<sub>3</sub>O<sub>4</sub>-PEI@AuNSs-EGCG/PEG. As shown in Figure S1-B, the diffraction peaks (311, 511, 400) correspond to the face-centered cubic structure of magnetite according to JCPDS card (No. 88-0315), and the peaks related to Au (red color) appeared which can be attributed to the crystal planes of 111, 200, 220 and 311 of the face-centered cubic structure (FCC) of Au (JCPDS No. 65-8601). This XRD results were further proved that the construction of Fe<sub>3</sub>O<sub>4</sub>-PEI@AuNSs-EGCG/PEG NPs was success.

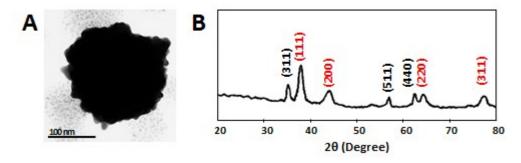


Figure S1. A) TEM of AuNSs-AA@Fe<sub>3</sub>O<sub>4</sub>-PEI/PEG. B) XRD pattern of AuNSs@Fe<sub>3</sub>O<sub>4</sub>-PEI/PEG (black: EGCG method; gray: ascorbic acid method)

## Normal cell culture

Human normal prostate epithelial RWPE-1 cells were maintained in keratinocyte serum-free medium (GIBCO, NY, USA) supplemented with 50 mg/l bovine pituitary extract, 5% l-glutamine and 5 μg/l epidermal growth factor. Cells were maintained in a humidified incubator (5% CO<sub>2</sub>) at 37 °C. RWPE-1 cells were seeded into 96-well plates at a density of 1 × 10<sup>4</sup> cells/well. After 24 h incubation, cells were treated with different concentration of Fe<sub>3</sub>O<sub>4</sub>-PEI@AuNSs-EGCG/PEG and Fe<sub>3</sub>O<sub>4</sub>-PEI@AuNSs-AA/PEG, respectively. After incubation for another 24 h, cytotoxicity was

measured by MTT method.

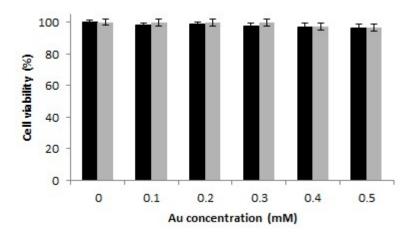


Figure S2. Cytotoxicity of RWPE-1 cells after Fe<sub>3</sub>O<sub>4</sub>-PEI@AuNSs-EGCG/PEG (black) and Fe<sub>3</sub>O<sub>4</sub>-PEI@AuNSs-AA/PEG (grey) treatment with different concentrations, respectively.