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A charge-conversional intracellular-activated polymeric prodrug for tumor therapy

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Abstract

Stimuli-responsive drug-delivery systems have attracted great attention for controlled drug release. The pH value is relatively low in the tumor tissues and the concentration of glutathione is significantly high in the cells. According to these features, a charge-conversional polymeric prodrug that is sensitive to pH and redox is designed. 3,3′-Dithiodipropionic acid modified paclitaxel (DTPA-PTX) and 2,3-dimethylmaleic anhydride (DMMA) are conjugated to the amino groups of poly(ethylene glycol)-*b*-poly(L-lysine) (mPEG-*b*-PLL). The surface charge of the obtained mPEG-*b*-(PLL-*co*-(PLL-DMMA)-*co*-(PLL-DTPA-PTX)) nanoparticles (DMMA-NPs) can change from negative at blood pH to positive at tumor extracellular pH, implying that the DMMA-NPs will have a prolonged blood circulation time through avoiding rapid clearance by reticuloendothelial system and an enhanced cell uptake at the tumor site. Compared to the control group, the *in vitro* studies show that the DMMA-NPs exhibit effective cellular uptake, rapid intracellular drug release, and significantly enhanced cytotoxicity against MCF-7 tumor cells, owing to the tumor-relevant pH and reductive conditions. These indicate that the charge-conversional DMMA-NPs provide an excellent platform for the potential tumor therapy.

Key Words：Chemotherapy, nanoparticles, paclitaxel, stimuli-responsive

1. Introduction

Within the last decade, polymeric drug delivery systems have emerged as a promising platform for cancer therapy.[1-6] Compared to the conventional chemotherapeutic agents, polymeric drug delivery systems for cancer therapy have the following advantages: 1) improving drug solubility; 2) altering the pharmacokinetics and tissue distribution; 3) prolonging circulation time through avoiding rapid clearance by reticuloendothelial system; 4) enhancing the drug accumulation in the tumor tissue *via* the Enhanced Permeability and Retention (EPR) effect; 5) reducing the side effects.[7-9]

An effective nanoparticular drug delivery system should possess multiple functions, such as prolonged circulation time, enhanced cellular internalization, and rapid intracellular drug release. In recent years, stimuli-responsive drug-delivery systems have attracted great attention in cancer therapy. Typical biological stimuli include pH, temperature, redox and enzymes.[10-17] Among all of the stimuli, pH and redox are the two most attracting one.[18] pH value varies in normal tissues and tumor tissue, the normal physiological pH value in the blood and normal tissues is about 7.4, but the pH value in cancerous tissues (pH 6.5 - 7.2), endosomes and lysosomes (pH 4.5 - 6.5) are slightly acidic.[19, 20] Thus pH-responsive nanoparticles can be designed to control drug release in specific tumor site. For instance, Wang et al. reported a dual pH-sensitive polymer-drug conjugate nanoparticulate system for efficient anticancer drug delivery [21]. In addition, the concentration of glutathione (GSH) is different between the extracellular (~2.0 to 20.0 μM) and intracellular (~10 mM) environments, this redox-responsiveness can be used for rapid intracellular drug release.[17, 22, 23] For example, Zhang et al. reported a paclitaxel-loaded redox-sensitive micelles system which accomplished rapid drug release under reducing condition.[24] Chen and co-workers reported redox and pH-responsive DOX-loaded micelles (PEG-SS-PTMBPEC), which showed significant release of

protein at pH 6.0 or under a reductive condition containing 10 mM dithiothreitol (DTT) at pH 7.4 compared to that at normal physiological pH 7.4.[25]

Even rapid progresses have been made in the development of multi-stimuli-responsive nanoparticles, there are still some insufficient aspects should be considered. Some nanoparticles were designed to achieve high drug loading capacity and rapid intracellular drug release, but other aspects such as enhanced cellular internalization and prolonged circulation time, had not been considered.[26] In particular, drug delivery system which can take advantages of pH and redox dual responsiveness of carriers for prolonged blood circulation, enhanced cellular internalization and triggered intracellular release of payload, was seldom reported.[27]

Therefore, in spite of the great efforts have been made in the stimuli-responsive field, two intractable problems are still remained. One is the contradiction between long circulation time and the efficient tumor cellular uptake, as prolonged circulation time needs slightly negative charge on the polymeric surface while enhanced cellular internalization needs positive charge on its surface. In the past decades, persistent efforts have been made in the charge-conversional nanoparticles. For example, Wang and co-workers designed zwitterionic polymer-based nanoparticles which exhibited prolonged circulation time and promoted the tumoral cell internalization *in vivo*, owing to the charge switching.[28] The other problem is the contrary requirement between stability of nanoparticles during blood circulation and intracellular drug release. Cai and co-workers developed redox-responsive doxorubicin prodrug (DEX-PEI(-SS-DOX)) micelles which reduced drug leaking during blood circulation and improved the drug release rate through the stability of the drug conjugating and clearly redox-responsive behavior.[29] Compared to encapsulated drugs through electrostatic or hydrophobic interactions, the drugs conjugated to the nanocarriers through chemical bonds is

relatively stable. So we decided to develop a polymer-drug conjugate which is stable in blood circulation and has rapid drug release profile under intracellular condition.

As previously described, we should solve all the problems in one system to achieve high therapeutic efficiency. Therefore, a charge-conversional polymer-drug conjugate engineered with disulfide bonds was designed (Scheme 1). The conjugate is sensitive to both pH and redox stimuli. DTPA-PTX was conjugated to the amino groups of mPEG-*b*-PLL to produce the disulfide-contained polymer-PTX conjugate mPEG-*b*-(PLL-*co*-(PLL-DTPA-PTX)), where the disulfide bond would be cleaved under the high concentration of GSH (intracellular level), thus drug could release rapidly in cells. The polymer-PTX conjugate was then modified with DMMA to produce mPEG-*b*-(PLL-*co*-(PLL-DMMA)-*co*-(PLL-DTPA-PTX)) nanoparticles (DMMA-NPs). DMMA was reacted with the amino group of polymer-PTX conjugate, which generated a pH-sensitive amide bond and a carboxylic acid group. The amide bond remains stable in neutral and alkali pH values, but is cleaved quickly under slightly acidic conditions. The nanoparticles should therefore ideally remain negatively charged at the blood pH, while change to positively charged at the tumor tissue pH. In addition, the disulfide-contained nanoparticles have rapid intracellular drug release because of the reductive conditions in tumor cells.[30] Thus the drug delivery system has great potential to achieve the goals including prolonged circulation time, enhanced cellular internalization and rapid intracellular drug release. In the present study, preparation and characterization of DMMA-NPs were presented as well as *in vitro* tests.

Scheme 1

2 Experimental

2.1 Materials

Methoxyl poly(ethylene glycol) (mPEG, M_n = 5000) and 3,3′-dithiodipropionic acid (DTPA)

were purchased from Aldrich and used as received. N^ε-Benzyloxycarbony-L-lysine (H-Lys(Z)-OH)

was purchased from GL Biochem Co.Ltd. (Shanghai, China). Amino-terminated methoxyl poly

(ethylene glycol) (mPEG-NH$_2$, M_n = 5000) was prepared according to our previous work.[31]

N^ε-benzyloxycarbonyl-L-lysine-N-carboxyanhydride (Lys(Z)-NCA) was synthesized according to

the literature.[32] N,N-Dimethylformamide (DMF) was stored over calcium hydride (CaH$_2$) and

purified by vacuum distillation with CaH$_2$. Triethylamine (TEA) was dried and distilled with CaH$_2$.

Paclitaxel (PTX) was purchased from Beijing Huafeng United Technology Company, China.

3-(4,5-Dimethyl-thiazol-2-yl)-2,5-diphenyl   tetrazolium   bromide   (MTT)   and

4′,6-diamidino-2-phenyl-indole dihydrochloride (DAPI) were purchased from Sigma and used as

received. Fluorescein isothiocyanate (FITC), DMMA and glutathione (GSH) were purchased from

Aladdin reagent company, China. Succinic anhydride was purchased from Sinopharm Chemical

Reagent Co. Ltd. and used without further purification.

2.2 Synthesis of DTPA-PTX

In a 250 mL flame-dry flask, PTX (0.500 g, 0.585 mmol) was dissolved in 80 mL of dry THF.

DTPA (0.185 g, 0.88 mmol) and EDC•HCl (0.340 g, 1.78 mmol) were then added and the mixture

was stirred at 30 °C for 2 h. Subsequently, DMAP (0.073 g, 0.598 mmol) was added. The

esterification reaction was maintained at 30 °C for 48 h. The solution was precipitated with 800 mL

0.1 M diluted hydrochloric acid. After filtration and drying under vacuum for 48 h, white solid was

obtained.

2.3 Synthesis of mPEG-*b*-PLL block copolymer

Methoxyl   poly(ethylene   glycol)-*b*-poly(L-lysine)   (mPEG-*b*-PLL)   were   synthesized   by

ring-opening polymerization of Lys-NCA using mPEG-NH$_2$ as a macroinitiator and subsequent

deprotection of benzyloxycarbonyl groups. Typically, mPEG-NH$_2$ (3.00 g, 0.60 mmol) was dissolved

in dry DMF (40 mL) after an azeotropic dehydration process with toluene. Then, Lys(Z)-NCA (3.43

g, 10.8 mmol) was dissolved in dry DMF (40 mL) and added to the mPEG-NH$_2$ solution in dry DMF

via a syringe under nitrogen. The reaction mixture was stirred for 3 days at 30 °C under a dry

nitrogen atmosphere and then precipitated into excess ice cold diethyl ether to obtain methoxy

polyethylene glycol-block-poly-ε-(benzyloxycarbonyl)-L-lysine [mPEG-*b*-PLL(Z)].

mPEG-*b*-PLL was obtained through the deprotection of benzyloxycarbonyl groups on the

PEG-*b*-PLL(Z). In brief, mPEG-*b*-PLL(Z) (4.70 g) was dissolved in trifluoroacetic acid (47 mL).

HBr/acetic acid (33 wt %) (6.15 mL) was then added at 0 °C. The reaction mixture was stirred for 45

min at 25 °C, and then precipitated into excessive ice cold diethyl ether. The product was further

purified by dialyzed (MWCO 3500 Da) against distilled water. White solid was obtained by

lyophilization.

2.4 Synthesis of mPEG-*b*-(PLL-*co*-(PLL-DTPA-PTX))

3,3′-dithiodipropionic acid modified PTX (DTPA-PTX) was conjugated to the amino groups of

mPEG-*b*-PLL to obtain a disulfide-contained polymer-PTX conjugate

mPEG-*b*-(PLL-*co*-(PLL-DTPA-PTX)). In brief, in a 50 mL flame-dry flask, DTPA-PTX (0.072 g,

0.069 mmol), EDC·HCl (0.039 g, 0.20 mmol) and NHS (0.024 g, 0.21 mmol) were dissolved in 10

mL of dry DMF and the reaction mixture was maintained stirring for 12 h at room temperature.

Subsequently, mPEG-*b*-PLL (0.50 g, 0.069 mmol) and TEA (0.015 g, 0.14 mmol) were dissolved in

dry DMF, and then added into the above-mentioned solution drop by drop. The condensation reaction

was maintained at 25 °C for 48 h. The solution was precipitated with excess amount of ice cold

diethyl ether. The product was dissolved in DMF and dialyzed (MWCO 3500 Da) against DMF to remove unreacted small molecules, then the solution was dialyzed (MWCO 3500 Da) against distilled water to remove the solvent DMF and the product was obtained by lyophilization.

2.5 Synthesis of mPEG-*b*-(PLL-*co*-(PLL-DMMA)-*co*-(PLL-DTPA-PTX))

Charge-conversional conjugate polymer mPEG-*b*-(PLL-*co*-(PLL-DMMA)-*co*-(PLL-DTPA-PTX)) (DMMA-NPs) was prepared as follows: mPEG-*b*-(PLL-*co*-(PLL-DTPA-PTX)) (0.10 g, 0.013 mmol) was dissolved in 20 mL of distilled water. pH of the solution was adjusted to 8 - 9 by adding 0.1 M NaOH, and stirred for 2 h at room temperature. 2,3-dimethyl malefic anhydride (DMMA) (0.475 g, 0.380 mmol) was then added, in the process the pH of the solution was kept 8 - 9 using 0.1 M NaOH. When the pH of solution was constant, the reaction mixture was stirring for another 12 h. The solution was dialyzed (MWCO 3500 Da) against distilled water and white solid was obtained by lyophilization.

As a control, mPEG-*b*-(PLL-*co*-(PLL-SA)-*co*-(PLL-DTPA-PTX)) (SA-NPs) was synthesized through a similar procedure as above. Details were described as follows: mPEG-*b*-(PLL-*co*-(PLL-DTPA-PTX)) (0.060 g, 0.0075 mmol) was dissolved in 15 mL of distilled water. pH of the solution was adjusted to 8 - 9 by adding 0.1 M NaOH, and then stirred for 2 h at room temperature. Succinic anhydride (SA) (0.023 g, 0.23 mmol) was added in several times to keep the pH $\approx$ 8 - 9 using 0.1 M NaOH. The subsequent steps were identical with the synthesis of the mPEG-*b*-(PLL-*co*-(PLL-DMMA)-*co*-(PLL-DTPA- PTX)).

2.6 Synthesis of FITC-labeled NPs

mPEG-*b*-(PLL-*co*-(PLL-DTPA-PTX)) (50 mg) was dissolved in 2 mL of distilled water, and then FITC (2.5 mg) in DMF (3 mL) was added. After stirring for 24 h at room temperature under

dark, the solution was dialyzed (MWCO 3500 Da) against distilled water and then yellow solid were

obtained by lyophilization.

2.7 Characterizations

The ^{1}H NMR, zeta potential, dynamic laser scattering (DLS) were performed as in our previous

studies [33, 34].

2.8 Calculation of PTX loading content by elemental analysis

The drug loading content (DLC) of PTX in the DMMA-NPs was calculated according to

equations:

PTX wt.% = (Mass of PTX in DMMA-NPs)/(Mass of DMMA-NPs)×100%   (Eq. 1)

PTX wt.%/M(PTX) = S wt.%/2M(S)   (Eq. 2)

where M(PTX) is the molecular weight of PTX, S wt.% is the mass percent content of sulphur

in the DMMA-NPs, M(S) is the atomic weight of sulphur.

2.9 Cell culture

Breast carcinoma MCF-7 cell lines were obtained from the Cell Bank of the Shanghai Institute

of Biochemistry & Cell Biology in the Chinese Academy of Sciences and cultured at 37 °C in a 5%

CO_2 atmosphere in Dulbecco's modified Eagle's medium (DMEM, Gibco) supplemented with 10%

fetal bovine serum (FBS), penicillin (50 U mL^{-1}), and streptomycin (50 U mL^{-1}).

2.10 MTT assays

The cytotoxicities of DMMA-NPs and SA-NPs were evaluated against MCF-7 cells by MTT

assay.[35] MCF-7 cells were seeded in 96-well plates (6000 cells per well) in 100 μL of DMEM

medium and incubated at 37 °C in a 5% CO_2 atmosphere for 24 h. The culture medium was replaced

with 200 μL of fresh medium without fetal bovine serum (FBS) containing DMMA-NPs or SA-NPs

at pH 7.4 or 6.5. After 3 h incubation, the medium were replaced with 200 μL of fresh medium at pH 7.4. After another 21 h or 45 h incubation, cell viability was analyzed using MTT and measured on a Bio-Rad 680 microplate reader at a wavelength of 490 nm.

2.11 Confocal laser scanning microscopy (CLSM) observation

MCF-7 cells were seeded in six-well plates at a density of 2×10^5 cells per well in 2 mL DMEM medium and incubated at 37 °C in a 5% CO_2 atmosphere for 24 h. The culture medium was replaced with 2 mL of fresh medium without FBS containing FITC-labeled DMMA-NPs or SA-NPs at pH 7.4 or 6.5. After 1 and 2 h incubation, the cells were washed with PBS and fixed with 4% formaldehyde for 20 min at room temperature. The cells nuclei were stained by DAPI. At last, the cells were monitored by confocal laser scanning microscopy (Carl Zeiss LSM 780).

3 Results and discussion

3.1 Preparation and characterization of DMMA-NPs

The preparation of DMMA-NPs was shown in Scheme 2. First, DTPA-PTX and mPEG-*b*-PLL were synthesized; then, DTPA-PTX was conjugated to the amino groups of the mPEG-*b*-PLL through condensation reaction, finally, DMMA or SA was conjugated to the mPEG-*b*-(PLL-*co*-(PLL-DTPA-PTX)) through amide formation.

Scheme 2

^{1}H NMR shows that DTPA was successfully conjugated to PTX, the signals at δ 2.87 ppm were contributed to the methylene peaks of DTPA ($-CH_2-CH_2-S-S-CH_2-CH_2-$). Furthermore, through the comparison of the ^{1}H NMR spectra of free PTX and DTPA-PTX, the 2′-CH proton peak shifts from 4.76 ppm (free PTX) to 5.61 ppm (conjugated PTX). However, the 7-CH proton peak remains where it was before (Figure 1). Therefore, the esterification reaction of PTX and DTPA took place

preferentially at the 2′-hydroxyl of PTX instead of the 7-CH, owing to the 2′-hydroxyl of PTX has

less steric hindrance than the 7-hydroxyl [36, 37]. The ESI-MS of DTPA-PTX shows peak at m/z =

1044.4 corresponding to the (M-H)⁻, which is entirely consistent with the theoretical calculation

value (Figure 2).

Figure 1

Figure 2

In this study, mPEG-*b*-PLL was prepared as before. The ^{1}H NMR spectra of methoxyl

poly(ethylene glycol)-*b*-poly(N^ε-benzyloxycarbonyl-L-lysine) [mPEG-*b*-P(LL(Z)] and mPEG-*b*-PLL

(Figure 3) confirmed that mPEG-*b*-PLL was synthesized successfully. All the peaks in the spectra

were well assigned. The degree of polymerization (DP) of Lys(Z) units in mPEG-*b*-P(LL(Z)) was

calculated by comparing the integration of methylene peaks of Lys(Z) groups (–CH$_2$–CH$_2$–CH$_2$–

CH$_2$–NH) with the integration of methylene peaks of poly(ethylene glycol) (-**CH$_2$-CH$_2$**-O-). Finally,

the DP of mPEG-*b*-P(LL(Z)) was calculated to be 17. Through a deprotection process of

mPEG-*b*-P(LL(Z)), the typical peaks of carbobenzoxy groups at δ 5.13 and 7.25 ppm disappeared,

indicating that the deprotection was complete. The DP of mPEG-*b*-PLL was also calculated to be 17

through the same method for the mPEG-*b*-P(LL(Z)).

Figure 2

Figure 3

Then, DTPA-PTX was conjugated to mPEG-*b*-PLL through a typical condensation reaction to

obtain disulfide containing polymer (mPEG-*b*-(PLL-*co*-(PLL-DTPA-PTX))). As shown in Figure 4,

comparing the ^{1}H NMR spectrum of mPEG-*b*-(PLL-*co*-(PLL- DTPA-PTX)) with that of DTPA-PTX,

typical signals of phenyl proton (7.2 - 8.0 ppm) of PTX appear, demonstrating that DTPA-PTX was

successfully conjugated to the mPEG-*b*-PLL. The DLC of PTX was calculated through two methods: one method was calculated by comparing the peak integration of phenyl proton of PTX with that of PEG methylene proton signal (δ 3.5 ppm), the result was 10.3 wt.% (Figure 4); the other one was the elemental analysis (EA), the DLC of PTX was calculated to be 7.9 wt.% by analysis the EA data (Table 1).

Table 1

Figure 4

Finally, mPEG-*b*-(PLL-*co*-(PLL-DTPA-PTX)) was reacted with 2,3-dimethyl malefic anhydride (DMMA) to obtain the charge-conversional nanoparticles DMMA-NPs. The characteristic resonance of methyl (-C(**CH₃**)=C(**CH₃**)-) of DMMA at δ 1.64 - 1.87 ppm was displayed in the ¹H NMR spectrum of mPEG-*b*-(PLL-*co*-(PLL-DMMA)-*co*-(PLL-DTPA-PTX)), indicating that DMMA was successfully conjugated (Figure 5). The degree of substitution of DMMA on PLL is 76%. mPEG-*b*-(PLL-*co*-(PLL-DTPA-PTX)) was reacted with succinic anhydride (SA) to obtain the non-charge reversal nanoparticles SA-NPs, where the degree of substitution of SA on PLL is 53%.

Figure 5

3.2 Dynamic laser scattering (DLS) and zeta potentials of DMMA-NPs

The sizes of nanoparticles in sub-100 nm range are regarded as more effective in the solid tumor penetration.[38, 39] As shown in Figure 6, the hydrodynamic radii (R_h) of DMMA-NPs and SA-NPs determined by DLS were 39.4 ± 10.9 and 35.7 ± 8.37 nm, respectively. This suggests that the nanoparticles should have good penetration ability in solid tumors.

It is well known that cell membranes are negatively charged, so that nanoparticles surface charge plays an important role in cellular uptake. Neutral and slightly negatively charged

nanoparticles have prolonged circulation time than positively charged nanoparticles, but they show

poorer internalization as compared to the positively charged nanoparticles.[40] The DMMA-NPs have

pH-sensitive DMMA groups that would be cleaved in slightly acidic condition and resulted in the

change of zeta potential of nanoparticles. The zeta potentials of DMMA-NPs at pH 7.4 (blood pH)

and pH 6.5 (tumor tissue pH) were shown in Figure 7. The zeta potential of DMMA-NPs increased

quickly at pH 6.5, it reached 0 mV in 6 min and became positive in 10 min. The zeta potential of

DMMA-NPs at pH 7.4 also increased, but it increased in a much slower rate than that at pH 6.5. The

increase of zeta potential of DMMA-NPs stopped after 30 minutes at both pH 7.4 and pH 6.5. These

indicated that the DMMA-NPs would be negatively charged in the normal physiological

environment and have long blood circulation time. After the DMMA-NPs reached tumor tissue sites,

the slightly acidic pH would cause the break of the pH-sensitive DMMA amide bond and change the

surface charge of the nanoparticles to positive, which would contribute to the internalization of the

nanoparticles. In contrast, no significant increase of zeta potential was observed for the SA-NPs, it

maintained at negative charge around -22 mV at both pH 7.4 and pH 6.5. Therefore the SA-NPs were

not charge-conversional nanoparticles and quite different with the DMMA-NPs.

<center>Figure 6 , Figure 7</center>

3.3 Cellular uptake of DMMA-NPs

Confocal laser scanning microscopy (CLSM) was used to investigate the cellular uptake

behaviors of DMMA-NPs in MCF-7 breast carcinoma cancer cell at pH 7.4 and 6.5 (Figure 8).

DMMA-NPs were labeled by FITC for internalization investigation. After 1 h incubation of

DMMA-NPs at pH 6.5, some green fluorescence was observed in the cell membrane and cytoplasm,

but little green fluorescence was observed at pH 7.4. After incubation for 2 h, the quantity of green

fluorescence increased at both pH 7.4 and 6.5, the green fluorescence was distributed mostly in the cytoplasm at pH 6.5, but little green fluorescence was observed in the cell membrane at pH 7.4. Thus, the charge-conversional nanoparticles enhanced cell internalization significantly, which was in consistent with the results of zeta potential analyses at pH 7.4 and 6.5.

Figure 8

3.4 *In vitro* cytotoxicity of copolymers

The *in vitro* cytotoxicities of charge-reversal DMMA-NPs and non-charge-reversal SA-NPs nanoparticles were evaluated using MTT assay against MCF-7 cell line in fresh serum-free DMEM at pH 7.4 and 6.5. As shown in Figure 9, after incubation for 24 and 48 h, the cell viabilities of MCF-7 cells treated with DMMA-NPs were all about 80% at pH 7.4 even if the highest concentration reached 50 μg/mL. In contrast, the cell viabilities incubated with DMMA-NPs could be less than 10% under the tumor extracellular pH 6.5, which indicated that the charge-reversal DMMA-NPs nanoparticles more effectively inhibited the proliferation of MCF-7 cells at pH 6.5 than at pH 7.4 (Figure 9A, 9B). Compared to the charge-reversal DMMA-NPs nanoparticles, after incubation of 24 h and 48 h against MCF-7 cell, the cell viabilities treated with non-charge-reversal SA-NPs were all above 70% at pH 7.4 and 6.5 (Figure 9C, 9D) at all the incubation concentrations. These demonstrated that pH-sensitive charge-reversal DMMA-NPs nanoparticles exhibited a more effective inhibition proliferation against tumor cells at tumor extracellular pH_e 6.5 than non-charge-reversal SA-NPs nanoparticles. This phenomenon was also observed in Wang and co-workers' work upon $^\oplus$NP/Pt@PPC-DA delivery system for overcome cisplatin resistance in cancer therapy [36]. In addition, owing to the existence of redox-responsive disulfide bonds in the nanoparticles, the charge-reversal DMMA-NPs nanoparticles at pH 7.4 and non-charge-reversal

SA-NPs nanoparticles at pH 7.4 and 6.5 showed slightly toxic against MCF-7 tumor cells. This was

demonstrated through *in vitro* cytotoxicity analyses of non-charge-reversal SA-NPs nanoparticles

against MCF-7 cell line in fresh DMEM at pH 7.4. MCF-7 cells were incubated for 1 h in fresh

DMEM containing 1 mM GSH or not, and then replaced with fresh DMEM containing

non-charge-reversal SA-NPs nanoparticles for another 47 h. As shown in Figure 9E, the half

maximal inhibitory concentration (IC_{50}) of SA-NPs with non-GSH was 28 times higher than that was

treated with 1 mM GSH. The phenomenon was contributed to the effect of disulfide linkages in the

nanoparticles. Similar phenomenon was also observed in Chen and his co-workers' study upon redox

and pH-sensitive polymer-drug conjugates for efficient delivery of paclitaxel.[30] In summary, the

charge-conversional DMMA-NPs nanoparticles have enhanced cellular uptake and inhibition against

the proliferation of tumor cells.

Figure 9

4 Conclusions

In conclusion, we have successfully designed a pH and redox dual responsive polymer-drug

conjugate for PTX delivery. The charge-reversal DMMA-NPs nanoparticles were well-synthesized

through conjugation of DTPA-PTX and DMMA to the amino groups of mPEG-*b*-PLL. The obtained

DMMA-NPs' surface charge changed from negative at blood pH to positive at tumor extracellular

pH. The charge reversal feature enhanced cellular uptake of nanoparticles in tumor extracellular

environment. Subsequently the disulfide linkages in the nanoparticles were cleaved due to the redox

reaction intracellular, leading to rapid drug release. Thus these demonstrate that the polymer-drug

conjugate have the advantages including prolonged circulation time, enhanced cellular

internalization and timely intracellular drug release, which enables the DMMA-NPs to be a promising alternative for tumor therapy.

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Captions to Figures

Scheme 1. Illustration of charge-conversional polymer-drug conjugate nanoparticles (DMMA-NPs) for long circulation, enhanced cellular uptake as well as intracellular redox-responsive release of PTX.

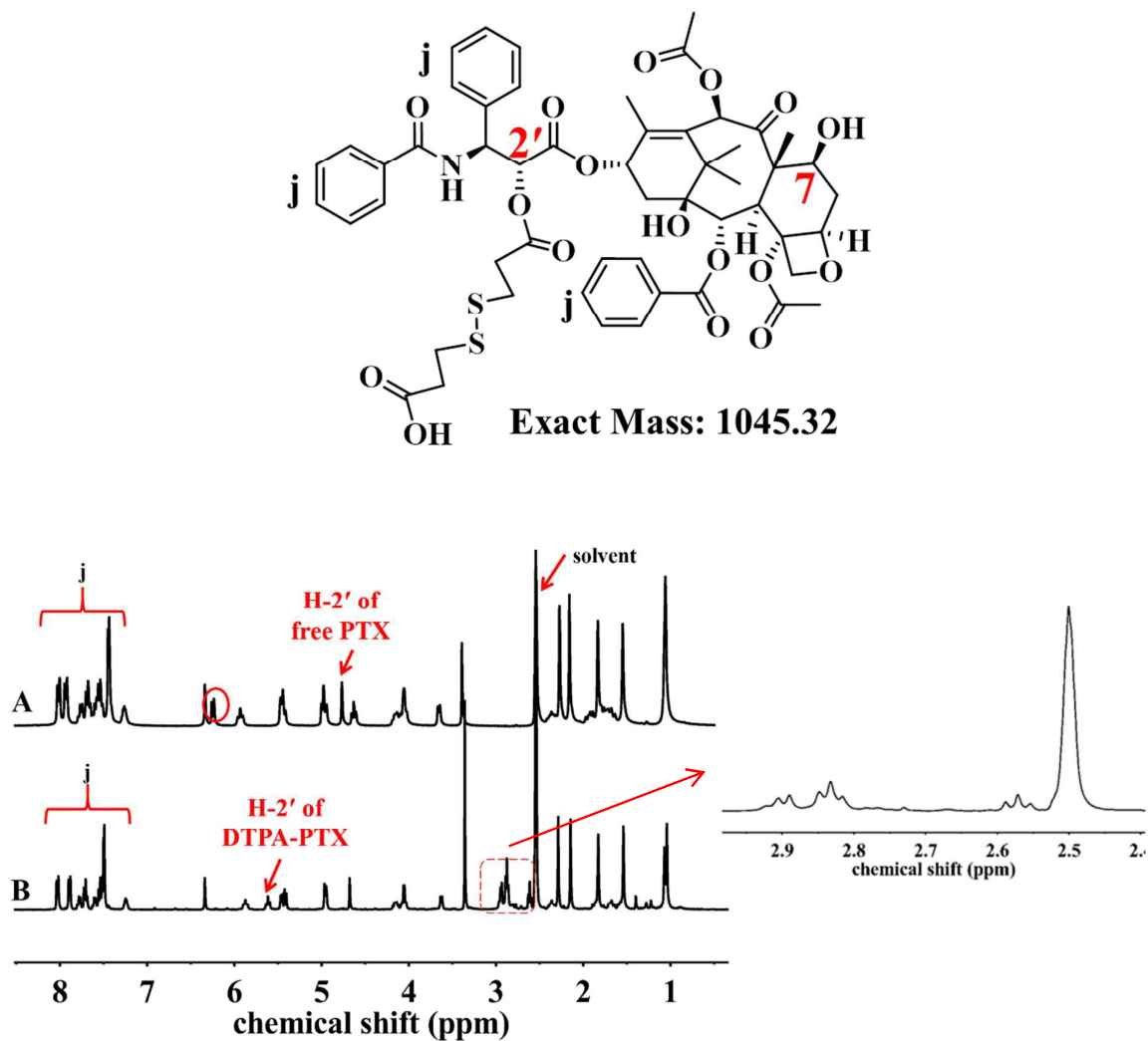Scheme 2. Synthesis of mPEG-*b*-(PLL-*co*-(PLL-DMMA)-*co*-(PLL-DTPA-PTX)).

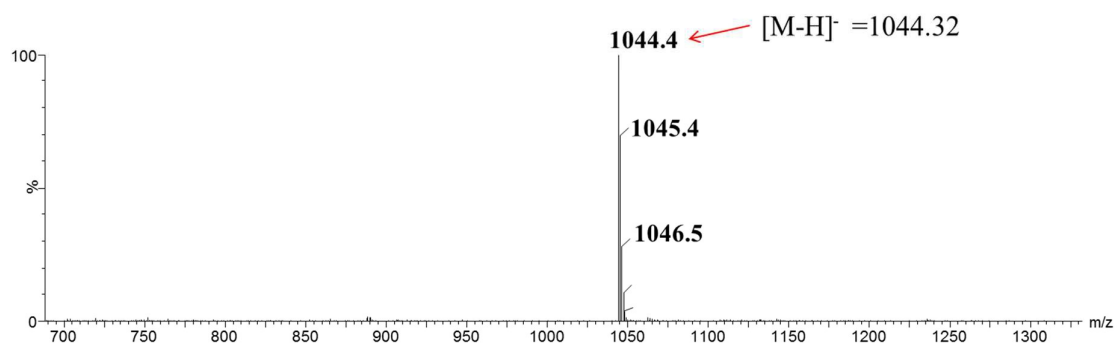Figure 1. ^{1}H NMR spectra of PTX (A), DTPA-PTX (B) in DMSO-d_6.
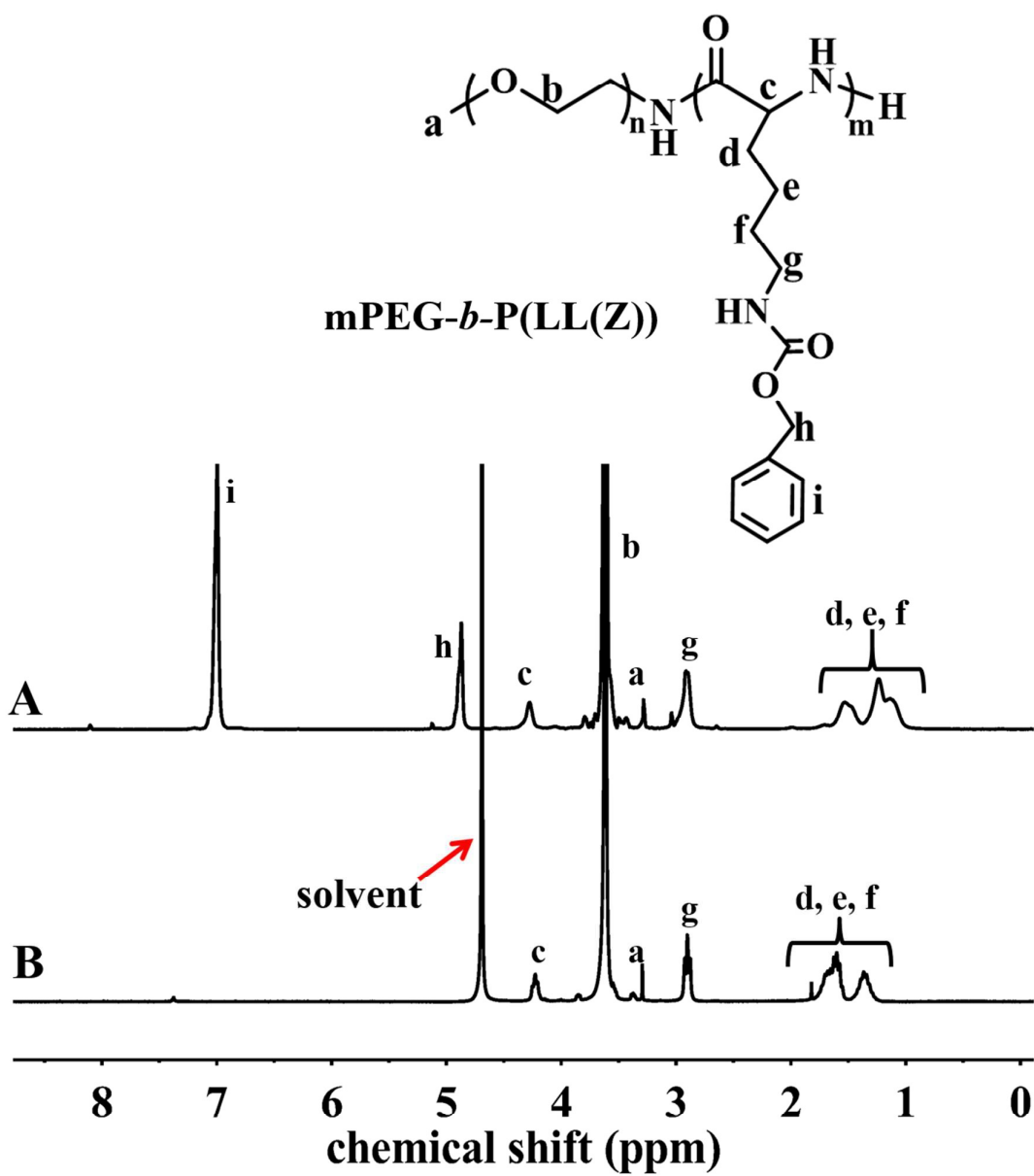
Figure 2. ESI mass spectrum of DTPA-PTX.

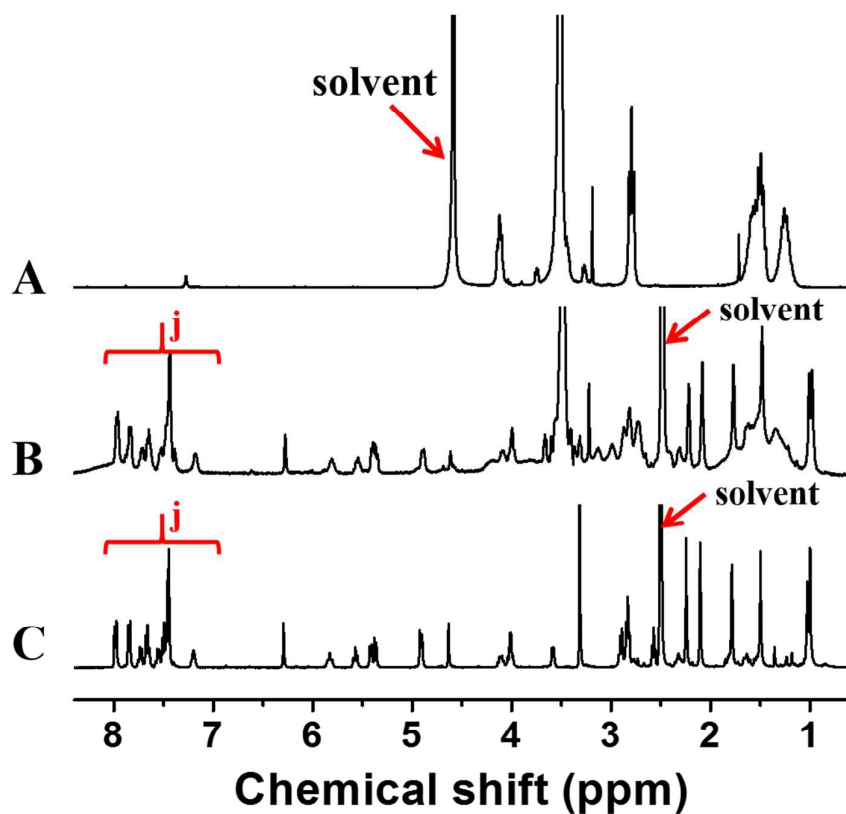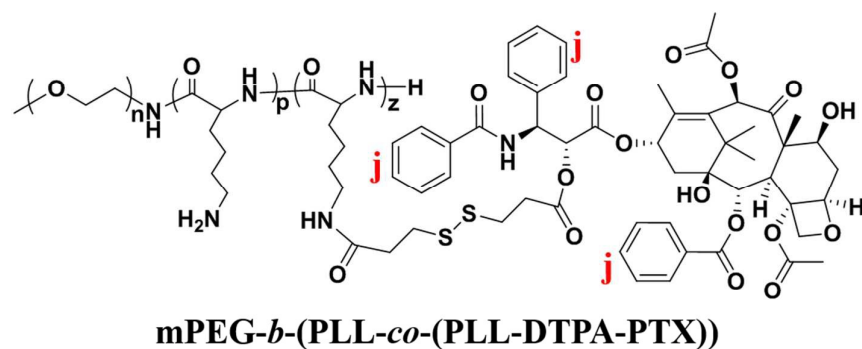Figure 3. ^{1}H NMR spectra of mPEG-*b*-P(LL(Z)) in CF$_3$COOD (A) and mPEG-*b*-PLL in D$_2$O (B).

Figure 4. ^{1}H NMR spectra of mPEG-*b*-PLL (A) in D$_2$O, mPEG-*b*-(PLL-*co*-(PLL-DTPA-PTX)) (B) and PTX (C) in DMSO-d_6.
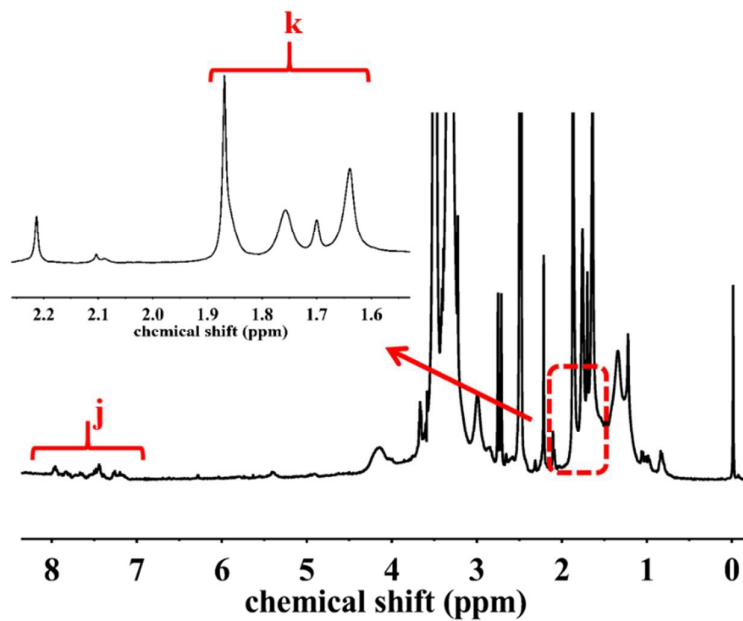
Figure 5. ^{1}H NMR spectrum of mPEG-*b*-(PLL-*co*-(PLL-DMMA)-*co*-(PLL-DTPA-PTX)) in DMSO-d_6.
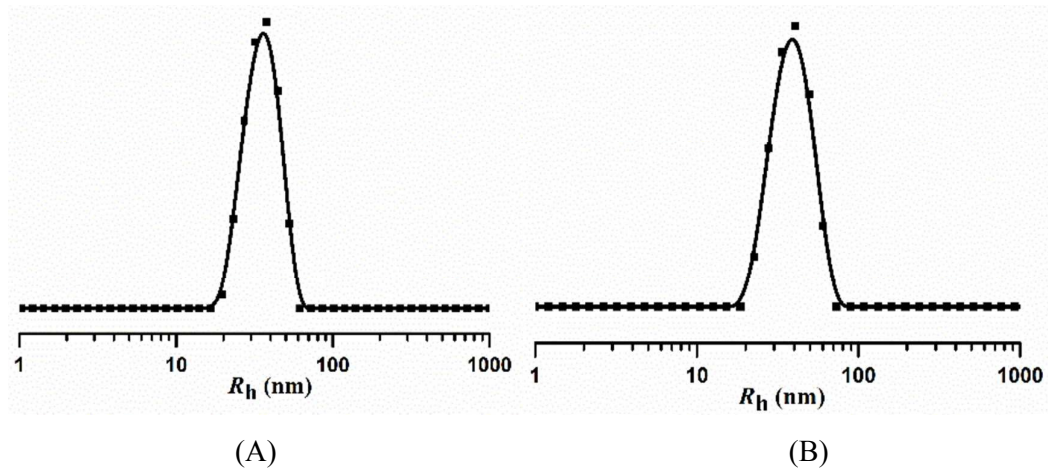
Figure 6. Size of DMMA-NPs (A), SA-NPs (B) determined by DLS.

Figure 7. Zeta potentials of DMMA-NPs (A) and SA-NPs (B) at pH = 6.5 and pH = 7.4 with different incubation time.

Figure 8. Confocal laser scanning microscopy images of MCF-7 cells after incubation with FITC-labeled DMMA-NPs for 1 and 2 h at different pH.

Figure 9. *In vitro* cytotoxicity of DMMA-NPs at 24 h (A), 48 h (B); SA-NPs at 24 h (C) ,48 h (D) to MCF-7 at different pH. *In vitro* cytotoxicity of SA-NPs incubated with free or 1 mM GSH to MCF-7 for 48 h (E).

Table 1. Elemental analysis of DMMA-NPs.

	N (%)	C (%)	H (%)	S (%)
DMMA-NPs	3.79	50.12	7.251	0.596

mPEG-*b*-(PLL-*co*-(PLL-DMMA)-*co*-(PLL-DTPA-PTX))

Scheme 1. Illustration of charge-conversional polymer-drug conjugate nanoparticles (DMMA-NPs) for long circulation, enhanced cellular uptake as well as intracellular redox-responsive release of PTX.

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mPEG-*b*-P(LL(Z))

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mPEG-*b*-(PLL-*co*-(PLL-DTPA-PTX))

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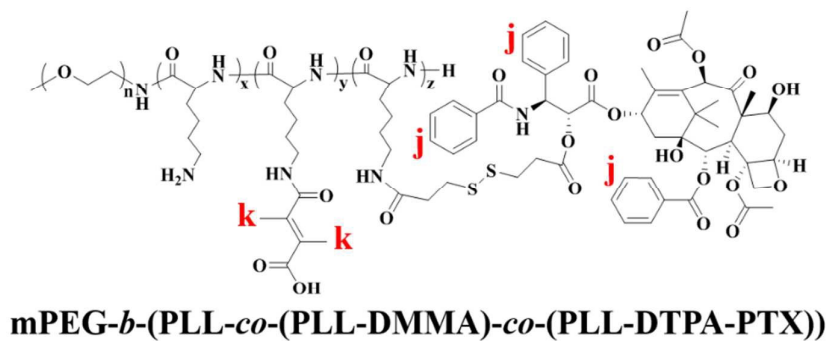
mPEG-*b*-(PLL-*co*-(PLL-DMMA)-*co*-(PLL-DTPA-PTX))

Figure 5. ^{1}H NMR spectrum of mPEG-*b*-(PLL-*co*-(PLL-DMMA)-*co*-(PLL-DTPA-PTX)) in

DMSO-d_6.

(A)   (B)

Figure 6. Size of DMMA-NPs (A), SA-NPs (B) determined by DLS.
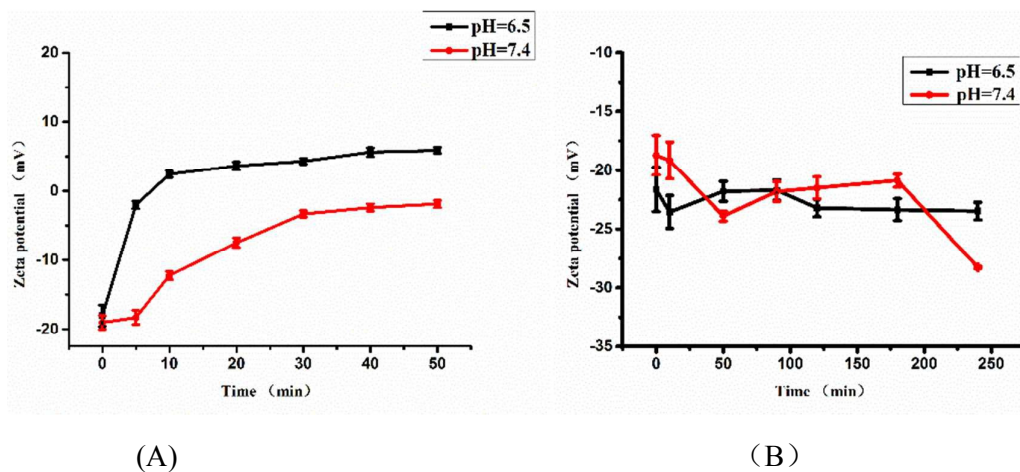
(A)   （B）

Figure 7. Zeta potentials of DMMA-NPs (A) and SA-NPs (B) at pH = 6.5 and pH = 7.4 with
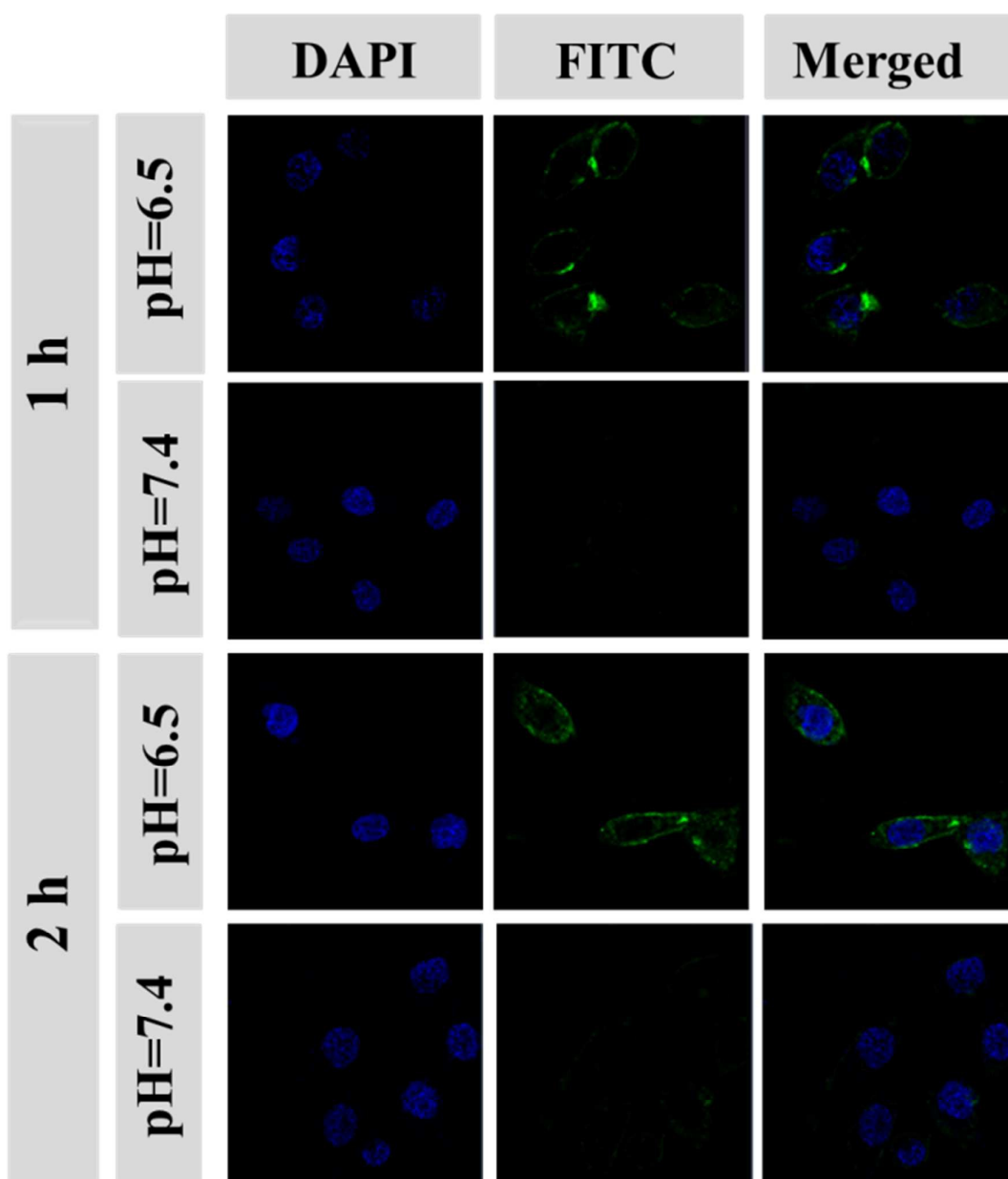
different incubation time.

Figure 8. Confocal laser scanning microscopy images of MCF-7 cells after incubation with FITC-labeled DMMA-NPs for 1 and 2 h at different pH.
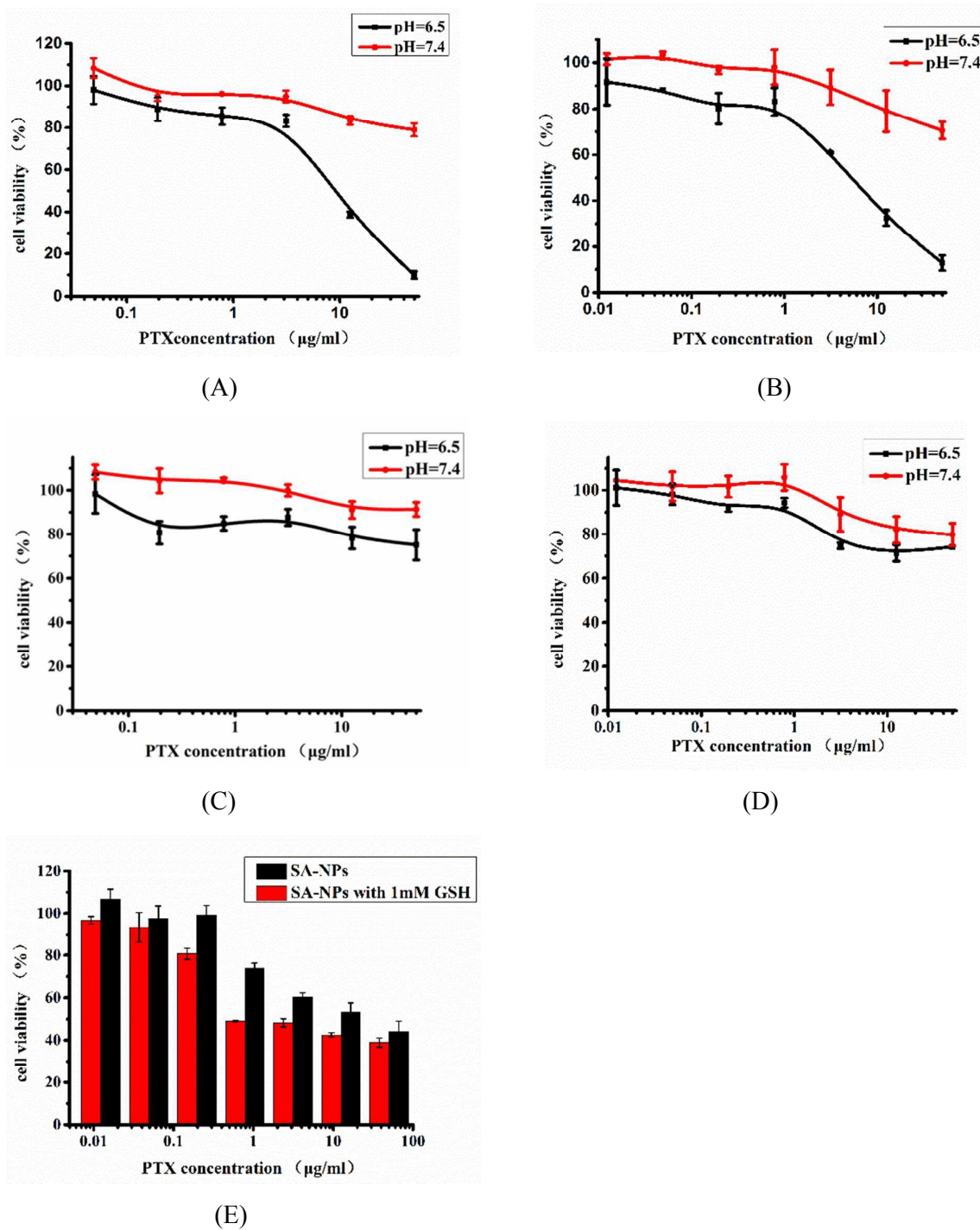
(A)

(B)

(C)

(D)

(E)

Figure 9. *In vitro* cytotoxicity of DMMA-NPs at 24 h (A), 48 h (B); SA-NPs at 24 h (C) ,48 h (D) to

MCF-7 at different pH. *In vitro* cytotoxicity of SA-NPs incubated with free or 1 mM GSH to MCF-7

for 48 h (E).

Table of contents entry

A pH and redox dual responsive polymer-PTX conjugate with prolonged circulation time, enhanced cellular internalization and timely intracellular drug release, is reported.