


 Cite this: *RSC Adv.*, 2020, **10**, 28891

Polymers prepared through an “ATRP polymerization–esterification” strategy for dual temperature- and reduction-induced paclitaxel delivery

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Clinically, the nanotherapy of tumors has been limited by the drug content, efficiency of targeted release, and bioavailability. In this study, we fabricated an amphiphilic block polymer, poly(2-methacryloyloxyethyl thiocicciccarboxylate)-block-poly(*N*-isopropylacrylamide) (PMAOETC-*b*-PNIPAM), using an “ATRP polymerization–esterification” strategy for paclitaxel (PTX) delivery. The hydrophobic drug paclitaxel was encapsulated based on hydrogen bond interactions between PTX and the PMAOETC and PNIPAM blocks, together with hydrophobic interactions between PTX and PMAOETC segments, affording PTX-laden polymer micelles with ~30% drug loading content. The critical micelle concentration of the PTX-loaded polymeric micellar aggregates was 34.53 mg l⁻¹, as determined through fluorescence spectroscopy, which indicated favorable stability during infinite dilution by body fluids. The phase transition temperature of the micelles was tunable (36.10–39.48 °C) via adjusting the lengths of the blocks. The PTX-laden micelles showed the release of a significant amount of PTX in cancerous tissue, while negligible cytotoxicity was shown against HCT-116 cells in PBS at pH 7.4 and 37 °C. Further *in vivo* anticancer studies revealed that antitumor treatment using the PTX-laden micelles caused a significant suppression in tumor volume compared with a free-PTX-treated group. This study provides a reference for improving drug content levels and optimizing the therapeutic effects of drug delivery systems from the perspective of polymer preparation.

Received 21st June 2020

Accepted 21st July 2020

DOI: 10.1039/d0ra05422d

rsc.li/rsc-advances

1. Introduction

Recently, as a fatal disease, the study of malignant tumors has become a vast emerging research field due to the wide variety of tumors and their complicated pathogenesis. However, efficacious therapeutic approaches, including gene therapy, drug therapy, surgical treatment, and chemoradiotherapy, face unprecedented challenges due to several reasons. In terms of antitumor drugs, some drug characteristics, such as high molecular weight and polyanionicity, make drugs more cumbersome to deliver,¹ and non-selectivity, low stability, poor solubility, and heterogenous biological dispersion can arise. Additionally, side effects from these drugs can be devastating, and the serious toxicity of anticancer drugs puts them at risk of dissociation in the reticuloendothelial system (RES).² To alleviate these dilemmas, self-assembled drug nanovehicles based on amphiphilic block polymers have evoked great interest as an

eye-catching platform due to their ability to load, solubilize, stabilize, and conserve drugs.^{3–5} Generally, amphiphilic polymers hold infinite promise for cancer therapy due to the coexistence of hydrophobic and hydrophilic shells. Furthermore, hydrophobic drugs can be encapsulated in polymers and assembled into nanocarriers as part of the inner core in aqueous solutions. Park *et al.* found that PEG-PDLLA had an encapsulation efficiency even lower than 2%.⁶ Considering that high drug content and drug bioavailability in carriers is crucial for cancer treatment, some drug delivery systems (DDSs) with satisfactory drug loading capacities have been fabricated *via* adjusting the hydrophobic/hydrophilic balance of the polymers and the polymer/drug ratio.⁷ Even so, regulatory strategies are difficult to control, and improvements in DDS performance are still limited.⁸ Only drugs that enter the nuclei work efficiently.⁹ However, lower drug content levels in polymeric-based micelles and subsequent torpid and partial release may delay the process of drug accumulation within nuclei. Above all, drugs that are released from nanocarriers are apt to decompose and can leak out before reaching the lesion location, causing serious harm to normal tissue and having a feeble effect on cancerous tissue.^{10–12}

The assembly of hydrophobic drugs and amphiphilic polymers that can respond to specific redox, glucose, pH, enzyme, and temperature stimuli ensures that drugs can exist in the blood

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circulation stably under the protection of the hydrophilic segment of the assembly.^{13–16} Among the various types of polymers, disulfide-bond-containing polymers can respond rapidly to glutathione (GSH) concentrations that differ from those in normal tissue; the GSH concentration is 2–20 μM in the blood circulation, 2–10 mM in mammalian cell cytoplasm, and *ca.* 20 mM intracellularly.¹⁷ Besides, 1,4-dithiothreitol (DTT) also can be utilized to cleave disulfide bonds into mercaptan and achieve targeted drug release.^{18–20} In 2017, the redox and pH dual-responsive polymer PEG-SS-COOH was synthesized for DOX delivery; DOX-laden micelles formed at a 1 : 1 molar feeding ratio of DOX to COOH and displayed a loading content (LC) of 24.1%.²¹ Poly(*N*-isopropylacrylamide) (PNIPAM), a frequently used thermo-sensitive component, was introduced into the structure and adjusted to obtain a smart polymer that could respond to the specific temperature of tumor lesions. Additionally, its obvious phase transition around a low critical solution temperature (LCST) made it possible to use for hydrophobic drug encapsulation.²²

To our knowledge, only a few studies have focused on improving the drug content and optimizing the therapeutic efficacy from the angle of polymer preparation. In this work, we designed an amphiphilic polymer, poly(2-methacryloyloxyethyl thiocticcarboxylate)-*block*-poly(*N*-isopropylacrylamide) (PMAOETC-*b*-PNIPAM), with dual temperature- and reduction-responsiveness based on the ATRP polymerization of both hydroxyethyl methacrylate (HEMA) and NIPAM, followed by esterification with thioctic acid (TA). PTX-laden polymers were prepared through synergistic loading from PMAOETC and PNIPAM, and they were able to self-assemble into globular nanomicelles at a critical micelle concentration (CMC) of 34.53 mg l^{−1}. This, combined with a zeta potential (ζ) of around 30 mV, reflects the favorable stability of the DDS during blood circulation. The obtained PTX-loaded nanomicelles have an adequate LC of 29.36% and EE of 78.33%, which exceeds the encapsulation capacity of PNIPAM by over 10%.²² In view of the significantly higher GSH concentration and temperature at lesion sites than in normal human tissue, the dually sensitive polymer contains appreciable levels of pendant disulfides and thermo-sensitive PNIPAM blocks. The disassembly behavior of the micelles was investigated in aqueous solution containing 10 mM GSH and DTT, and an *in vitro* PTX release study was performed in the presence of corresponding stimuli mimicking the physiological cancerous environment. Finally, the cytotoxicity of the PTX-laden micelles was evaluated *via* MTT assays against HCT116 cells. The *in vivo* anticancer efficacy was assessed using mice bearing 4T1 tumors, and the results suggested that the PTX-laden micelles were safe and efficient for tumor suppression. In light of these characteristics of PMAOETC-*b*-PNIPAM, this strategy of “ATRP polymerization-esterification” will hopefully promote drug loading into carriers and lead to satisfactory therapeutic effects.

2. Experimental section

2.1 Materials and reagents

All chemicals were of analytical grade and purchased from commercial suppliers. NIPAM and cuprous chloride (CuCl) were

purified, and the solvents used for reactions needed to have any water removed prior to use. Unless specified, reagents were used without further purification.

2.2 Synthetic procedure

PMAOETC-*b*-PNIPAM was synthesized *via* a three-step process, including the ATRP polymerization of hydroxyethyl methacrylate (HEMA) and *N*-isopropylacrylamide (NIPAM), and esterification with thioctic acid (TA). The typical synthesis procedure is shown in Scheme 1.

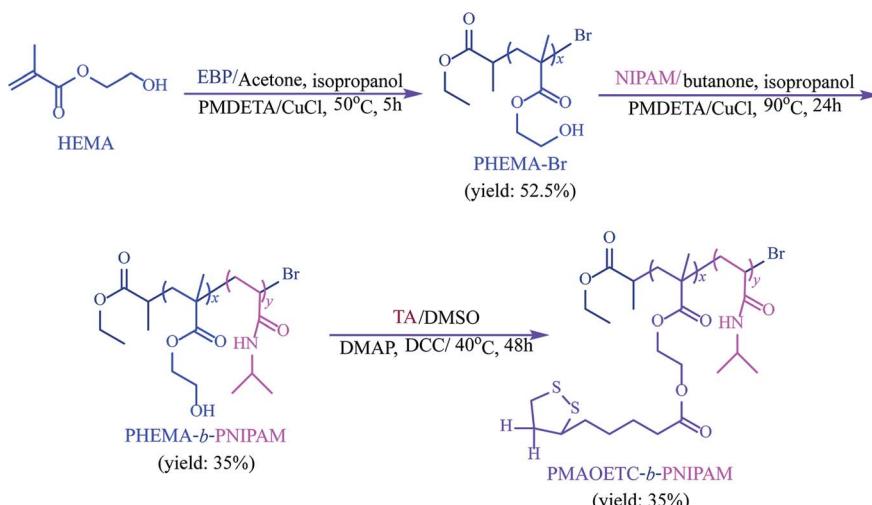
2.2.1 Synthesis of the macromolecular initiator PHEMA-Br.

The ATRP polymerization of HEMA was implemented employing ethyl 2-bromopropionate (EBP) as an initiator at EBP : HEMA molar ratios of 1 : 30 and 1 : 50. Representative, HEMA (3.65 ml, 30 mmol) and EBP (128 μl , 1 mmol) were dissolved into acetone-isopropanol mixed solvent (6 ml, v/v = 2/1) under stirring. Next, *N,N,N',N',N'*-pentamethyldiethylenetriamine (PMDETA, 633 μl , 3 mmol) and CuCl (0.199 g, 2 mmol) were added promptly, and “freeze-pump-thaw” circulation was used to thrice remove air and moisture from the system. Then, polymerization was carried out at 50 °C for 5 h. The polymerization was quenched *via* air-cooling, and the cyan mixture was collected after precipitation with excess ether. This was followed by dissolution in the mixed solvent and then the solution was passed through a neutral aluminum column to eliminate the catalyst and unreacted ligands. The residue achieved after rotary evaporation was precipitated in ether thrice to eliminate unpolymerized HEMA, and the white precipitate was vacuum dried at 30 °C overnight (yield: 52.5%). ¹H NMR (400 MHz, DMSO-*d*₆) δ (ppm): 1.12–1.21 (m, 3H of $\text{CH}_3\text{CH}_2\text{OCCO}$ – and 3H of $-\text{CH}(\text{CH}_3)\text{COO}$ – in EBP residue), 1.50–1.60 (m, 1H, $-\text{CH}(\text{CH}_3)\text{COO}$ – of the EBP structure), 0.77–1.04 (m, 3H, $-\text{CH}_2\text{C}(\text{CH}_3)(\text{COO})$ – of main chains), 1.78 (m, 2H, $-\text{CH}_2\text{C}(\text{CH}_3)(\text{COO})$ – of main chains), 3.90 (m, 2H, $-\text{COOCH}_2\text{CH}_2\text{OH}$), 3.58 (m, 2H, $-\text{COOCH}_2\text{CH}_2\text{OH}$), 4.81 (s, 1H, $-\text{COOCH}_2\text{CH}_2\text{OH}$). FTIR (film) ν (cm^{−1}): 1166 (C–O stretching vibration), 1726 (−C=O stretching vibration), 2848–2981 (−C–H stretching vibration), 3080–3755 (−OH).

2.2.2 Synthesis of PHEMA-*b*-PNIPAM.

PHEMA-*b*-PNIPAM was prepared using PHEMA-Br as a macromolecular initiator through ATRP polymerization at PHEMA-Br : NIPAM : PMDETA : CuCl molar ratios of 1 : 100 : 3 : 2 and 1 : 200 : 3 : 2. Typically, PHEMA-Br (0.131 mmol, 0.50 g) and NIPAM (13.1 mmol, 1.48 g) were completely dissolved in anhydrous isopropanol–butanone mixed solvent (4 ml, v/v = 1/1). The ligand PMDETA (0.393 mmol, 81.7 μl) and catalyst CuCl (0.262 mmol, 25.9 mg) were blended into the system. “Freeze-pump-thaw” circulation was carried out thrice. After the polymerization process was carried out at 90 °C for 24 h, the reaction mixture was purified *via* passing it through a neutral aluminum column, followed by rotary evaporation. The obtained liquor was precipitated in excess ether thrice. Then, the precipitant was vacuum-dried at 25 °C for 24 h (yield: 35%). ¹H NMR (400 MHz, DMSO-*d*₆) δ (ppm): 0.70–1.22 (m, 6H of EBP residue, 3H of $-(\text{CH}_2)\text{C}(\text{CH}_3)\text{COO}$ –, and 6H belonging to $-\text{CONHCH}(\text{CH}_3)_2$ of PNIPAM), 1.61–2.06 (m, 2H from $-\text{CH}_2\text{CH}(\text{CH}_3)\text{COO}$ – of PHEMA, 2H of $-\text{CH}_2\text{CH}(\text{CONH})$ –, and 1H of $-\text{CH}_2\text{CH}(\text{CONH})$ – in PNIPAM), 3.71–4.05 (m, 2H from $-\text{COOCH}_2\text{CH}_2\text{OH}$ of PHEMA, and 1H from





Scheme 1 A schematic diagram of the synthesis of the polymer PMAOETC-*b*-PNIPAM.

-CONHCH(CH₃)₂ of PNIPAM), 6.93–7.53 (s, 1H, -CONHCH(CH₃)₂ of PNIPAM). FTIR (film) ν (cm⁻¹): 1542 (–N–H bending vibration), 1645 (amide –C=O stretching vibration), 2848–2981 (–C–H stretching vibration), 3136–3500 (N–H stretching vibration).

2.2.3 Synthesis of PMAOETC-*b*-PNIPAM. PMAOETC-*b*-PNIPAM was produced through the esterification of PHEMA-*b*-PNIPAM and TA in the presence of 4-dimethylaminopyridine (DMAP) and the dehydrant *N,N*'-dicyclohexylcarbodiimide (DCC). Representative, PHEMA-*b*-PNIPAM (0.40 g, 8.07 \times 10⁻² mmol, -OH content: 2.26 mmol), TA (3.39 mmol, 0.699 g) and DMAP (2.26 mmol, 0.276 g) were dissolved in anhydrous DMSO (25 ml) in a 100 ml Schleck flask with a stirring bar and kept in an ice-water bath for 3 h under a nitrogen atmosphere to ensure the adequate activation of carboxyl. Next, a solution of DCC (2.712 mmol, 0.560 g) in refined DMSO (15 ml) was added dropwise into the above system over a 1 h period, and the reaction proceeded at 40 °C for 48 h. The mixture was filtrated twice to remove the *N,N*'-dicyclohexylurea (DCU) precipitate. Then, the solution was dialyzed against deionized water for 48 h, during which the water was renewed every 3 h. The solution was lyophilized and freeze-dried afterwards (yield: 35%). ¹H NMR (400 MHz, CDCl₃) δ (ppm): 1.63 (m, 4H, -OCOCH₂CH₂CH₂-CH₂- from side chains of PMAOETC), 2.36 (m, 2H, -OCO(CH₂)₄CH(S–S)- and -SSCH₂CH–H- of PMAOETC), 1.42 (m, 2H, -OCOCH₂CH₂CH₂CH₂- of PMAOETC), 3.06 (m, 2H, -SSCH₂CH–H- of five-membered rings belonging to PMAOETC), 3.53 (2H, -OCOCH₂CH₂CH₂CH₂- from side chains of PMAOETC), 1.72 (m, 1H, -SSCH₂CH–H- of five-membered rings ascribed to PMAOETC side chains). FTIR (film) ν (cm⁻¹): 2855–2928 (PMAOETC –C–H stretching vibration), 1634 (amide –C=O stretching vibration), 1539 (amide –N–H bending vibration), 1157 (–C–O stretching vibration), 3129–3561 (–N–H stretching vibration).

2.3 The generation of PMAOETC-*b*-PNIPAM polymeric micelles in aqueous solution

A dialysis method was applied to form the PMAOETC-*b*-PNIPAM block polymer micellar solution. Polymer powder (30 mg) at

different reaction feed ratios was dissolved completely in DMSO (10 ml) and transferred into a dialysis tube (MWCO: 7000) for dialysis against 2000 ml of deionized water for 72 h. Deionized water was renewed within predetermined time intervals. Once DMSO was completely removed, the translucent solution was gathered into a 25 ml volumetric flask for measurements. The maximum concentration of the micellar solution was 1200 mg l⁻¹.

2.4 Characterization and determination

¹H NMR spectra were recorded using a 400 MHz Bruker Avance spectrometer (Bruker, Germany) at 25 °C with DMSO-*d*₆ and CDCl₃ as solvents, and tetramethylsilane was used as the internal standard. Fourier transformation infrared (FTIR) analysis was performed using an EQUINX55 FTIR spectrophotometer (Bruker Corp., Germany) through the KBr pellet method. A gel permeation chromatography system (GPC, Waters Corp., USA) with a Waters 515 pump and Waters 2410 differential refractive index detector was used to determine the number-averaged molecular weight (M_n), weight-averaged molecular weight (M_w) and polydispersity index (PDI) data from the polymers; linear polystyrene and chromatographically pure THF were selected as the calibrated standard and eluent, respectively.

A FluoroMax-4 spectrofluorophotometer (HORIBA Scientific, USA) was adopted to investigate the micellization behavior and CMC using a pyrene probe. In brief, an acetone solution of pyrene (4.8 μ l, 5 \times 10⁻⁴ M) was added into the sample cell and dried for 2 h to ensure that the acetone fully evaporated. Then, micellar solutions (4 ml) with different concentration gradients (1 \times 10⁻⁴ to 5 \times 10⁻¹ mg ml⁻¹) were added into the corresponding cells to allow for assembly with pyrene while stirring overnight. The fluorescence spectra were collected at an excitation wavelength of 331 nm, the emission wavelength ranged from 350–550 nm, and the slit width for both excitation and emission was 2 nm.

A dynamic light scattering (DLS, BI-90Plus, America) instrument equipped with an argon laser was selected to detect



hydrodynamic diameter (D_h), polydispersity index (PDI), and ζ data from the micelles (100 mg l⁻¹). DLS analysis was carried out under the following conditions: λ , 660 nm; deflection angle, 90°; power output, 15 mW; 25 °C. For reduced micelles, reduced GSH (12.3 mg) was blended into the polymer micellar solution (4 ml, 1 × 10⁻¹ mg ml⁻¹) under stirring at 25 °C. Afterwards, the D_h values of the reduced micelles were detected at 0, 0.5, 4, and 24 h. The morphologies and size dispersions of micelles with/without GSH treatment were observed *via* JEM-2100 transmission electron microscopy (TEM, Electron Corp., Japan) at an accelerating voltage of 200 keV. Prior to observations, blank, GSH-reduced, PTX-laden, and released (under thermal and reductive conditions consistent with tumor sites) micellar solutions (10 μ l, 1 × 10⁻¹ mg ml⁻¹) were transferred onto carbon film, followed by air-drying.

A UV-vis spectrophotometer (UV-6100S, Mapada, Shanghai) was used to record the transmittance of micellar solutions (1 × 10⁻¹ mg ml⁻¹) at temperatures ranging from 20–70 °C, as this can serve as a critical parameter for evaluating thermal responsiveness.

2.5 The reduction-induced assembly behavior of polymer micelles

The assembly behavior of PMAOETC-*b*-PNIPAM micelles treated with 10 mM GSH in PBS at pH 7.4 was detected using DLS and TEM at different time intervals. Specifically, 4 ml of prepared micellar solution containing 0 or 10 mM GSH was subjected to gentle stirring at 37 °C for 0, 0.5 h, 4 h, and 24 h.

2.6 The loading of PTX into micelles *via* self-assembly

As a model anticancer drug, PTX was loaded into the polymer micelles *via* dialysis. PMAOETC-*b*-PNIPAM (40 mg) and PTX (10 mg) were dissolved entirely in DMSO (6 ml). Next, a small amount of deionized water was added dropwise until the solution became turbid. The mixture was then encased in a dialysis bag (MWCO: 7000) and dialyzed against deionized water for an additional 72 h. The water was replaced every 8 h. Dissociated PTX was removed *via* low-speed centrifugation, and PTX-laden micelles were obtained *via* lyophilization and then collected and stored at 4 °C for further measurements. LC and EE values for PTX were calculated on the basis of eqn (1) and (2):

$$\text{LC (\%)} = (\text{amount of PTX in polymer/amount of PTX-laden polymer}) \times 100\% \quad (1)$$

$$\text{EE (\%)} = (\text{amount of PTX in polymer/amount of PTX added originally}) \times 100\% \quad (2)$$

Prior to measurements, the linear relationship between the PTX concentration and UV absorbance at 210 nm was obtained *via* a calibration curve to achieve the equation:

$$C (\text{mg l}^{-1}) = (A - 0.0128)/0.39082 \quad (R^2 = 0.9982) \quad (3)$$

where C represents the concentration of PTX in standard solution, and A denotes the absorbance at 210 nm. Additionally,

drug-encapsulated micelles were dissolved in methanol to detect the LC of the synthesized PTX-loaded micelles.

2.7 The temperature- and reduction-induced release behavior of PTX-laden micelles

To assess the responsiveness of the PTX-laden nanoparticles to thermal and reduction stimuli, PBS solution of lyophilized PTX encapsulating polymer powder (4 mg, 1 mg ml⁻¹) was encased in a dialysis bag (MWCO: 3500) and dialyzed against 200 ml of PBS (pH: 7.4 and 5.6) containing different reductants (GSH and DTT) at 10 mM under gentle stirring at 30, 37, and 42 °C for 85 h. At predetermined time intervals, 4 ml aliquots were extracted from the dialysis medium and supplemented with isometric fresh medium. The cumulative PTX release amount was assessed *via* determining the medium absorbance at 210 nm of samples taken out after different time intervals. The cumulative PTX release was calculated as follows:

$$\text{Cumulative PTX release (\%)} = (M_t/M_o) \times 100 \quad (4)$$

where M_t and M_o denote the mass of PTX at time t and the amount of PTX encapsulated in the micelles, respectively.

2.8 The *in vitro* cytotoxicity of PTX encapsulating micelles

The human colon cancer cell line HCT116 was purchased from Mingjin Biology (Shanghai, China) and cultured in DMEM supplemented with 10% fetal calf serum (Gibco, America) under fully humidified conditions containing 5% CO₂ at 37 °C. The cytotoxicities of prepared blank and PTX-laden polymer micelles against HCT116 cells were measured using MTT assays. HCT116 cells were seeded on 96-well plates at a density of 1 × 10⁵ per well and cultured in 200 μ l of DMEM with 10% Hyclone fetal bovine serum for 24 h at 37 °C in 5% CO₂. Next, the medium was removed, cells were rinsed with PBS thrice, and 200 μ l of free PTX, blank and PTX-encapsulated micellar solutions with different concentration gradients from 1 to 500 mg l⁻¹ were suspended in pure DMEM and incubated for 72 h. After that, the cells were eluted with PBS (pH 7.4) thrice and supplemented with another 180 μ l of fresh DMEM and 20 μ l of MTT solution (5 mg ml⁻¹). After the final 4 h, 150 μ l of DMSO was added into the sediment and the mixture was shaken for 10 min. HCT116 cells were seeded into fresh culture medium and incubated under the same conditions to act as a negative control.

The absorbance values were recorded using a Model 680 universal microplate reader (Bio-Rad laboratories (UK) Ltd) at 490 nm. The cell viability was estimated *via* the following formula:

$$\text{Cell viability (\%)} = \text{OD}_{\text{Sample}}/\text{OD}_{\text{Control}} \times 100\% \quad (5)$$

where OD_{Sample} and OD_{Control} signify the optical densities of the sample and negative control wells, respectively. Statistical significance was evaluated using Student's *t*-test. All quantitative data are presented as mean value ± standard deviation.



2.9 *In vivo* animal studies

2.9.1 *In vivo* anticancer efficacy. The murine breast cancer cell line 4T1 was supplied by Mingjin Biology (Shanghai, China) and cultured in RPMI 1640 supplemented with 10% fetal calf serum (Gibco, America) under humidified conditions containing 5% CO₂ at 37 °C. The conventional digestion and selection of cells in the exponential growth phase were performed for subsequent experiments.

Male Balb/c mice (20 ± 2 g), purchased from the Laboratory Animal Center of Xi'an Jiaotong University, were selected to investigate the anticancer efficacy. Animals were fed standard lab chow and tap water. All animal procedures were performed in accordance with the Guidelines for Care and Use of Laboratory Animals of Xi'an Jiaotong University, and experiments were approved by the Animal Ethics Committee of Xi'an Jiaotong University Health Center. The mice were divided into 5 groups randomly ($n = 3$). Then, 0.1 ml of PBS containing 1×10^6 4T1 murine breast cancer cells was inoculated subcutaneously below the mouse shoulder. PTX, and blank and PTX-entrapped micelles were injected into each group when the tumor volume reached 100 mm³ (PTX dose: 5 mg kg⁻¹; tumor volume = $(L \times W^2)/2$, herein L and W denote the length and width of the tumor, respectively). The remaining injections were performed on days 3, 6, 9, 12, and 15. The control groups were injected with isometric normal saline and PBS. The tumor sizes and mice body weights were monitored at predetermined time intervals.

2.9.2 Histological testing. On day 15 of treatment, the five mice groups were sacrificed; their major organs, including hearts, livers, spleens, lungs, and kidneys, were removed, fixed in 4% formaldehyde, processed with paraffin, and sliced. The sections obtained were dyed with hematoxylin and eosin (H&E) for histological analysis.

3. Results and discussion

3.1 Synthesis and characterization of PMAOETC-*b*-PNIPAM block polymers

Temperature- and reduction-sensitive PMAOETC-*b*-PNIPAM block polymers were synthesized *via* the ATRP polymerization of HEMA and NIPAM, using EBP and PHEMA-Br as initiators, respectively. This was followed by esterification with TA. The chemical structure of each product was characterized *via* ¹H NMR and FTIR.

Fig. 1 shows the ¹H NMR spectra of PHEMA, PHEMA-*b*-PNIPAM, and PMAOETC-*b*-PNIPAM. The characteristic peak from hydroxyl protons belonging to PHEMA appears at 4.81 ppm (Fig. 1A), and the signal from protons of amide units is located at 7.16 ppm (Fig. 1B). The representative peaks from protons attributed to the five-membered ring of TA emerge at 1.72, 2.36, and 3.06 ppm (Fig. 1C). This indicates that PHEMA-Br, PHEMA-*b*-PNIPAM, and PMAOETC-*b*-PNIPAM were successfully prepared. Furthermore, the chemical signals from both double bond protons of NIPAM (Fig. 1B) and hydroxyl protons attributable to PHEMA units (Fig. 1C) disappear, which further confirms that the ATRP polymerization of HEMA and the esterification of TA were successfully carried out.

Additionally, FTIR analysis was also performed to verify the components of PMAOETC-*b*-PNIPAM and its precursors. Due to the existence of hydrogen bonds interacting with the amide structures, the stretching vibration of -NH- was split into two signals at 3294 and 3419 cm⁻¹.

The -C-H (2855–2928 cm⁻¹), -C=O (1634 cm⁻¹ from amide carbonyl, 1724 cm⁻¹ from ester carbonyl of PMAOETC), -C-O (1157 cm⁻¹) and -NH- (3129–3561 cm⁻¹) stretching vibrations, and -NH- (1539 cm⁻¹) bending vibration also validate the chemical structure of PMAOETC-*b*-PNIPAM (Fig. 1D(c)).

Chemical composition and molecular weight (M_n) values were estimated using the integration area ratio of the terminal methyl peak of EBP at 1.21 ppm to the peak attributed to hydroxyl from HEMA at 4.81 ppm. Similarly, the integration area ratio of the hydroxyl peak from PHEMA at 4.81 ppm to the peak area of methyne from PNIPAM at 3.91 ppm was identified as illustrating the repeat unit number of NIPAM. When the hydroxyl peak from PHEMA at 4.81 ppm disappeared after esterification, the M_n value of PMAOETC-*b*-PNIPAM was assessed according to the following formula:

$$M_n(\text{PMAOETC-}b\text{-PNIPAM}) = M_n(\text{PMAOETC-}b\text{-PNIPAM}) + xM_n(\text{TA}) - xM_n(\text{H}_2\text{O})$$

where x signifies the repeat unit number of TA, and $M_n(\text{TA})$ and $M_n(\text{H}_2\text{O})$ are 206.3182 and 18 g mol⁻¹, respectively. There are about 28 and 40 HEMA repeat units in PHEMA-Br, and 10 and 21 NIPAM repeat units polymerized in PHEMA-*b*-PNIPAM. Therefore, it can be deduced that the maximum conversion ratios of HEMA and NIPAM are 93.3% and 10.0%, respectively. The polymers with the composition formulae of PMAOETC₂₈-*b*-PNIPAM₁₀, PMAOETC₂₈-*b*-PNIPAM₂₁, and PMAOETC₄₀-*b*-PNIPAM₁₀ are hereafter denoted as P1, P2, and P3, respectively.

In addition, the GPC curves of the polymers are shown in Fig. 2; it is clear that symmetric elution curves with single peaks and, moreover, without tailing and impurity peaks are exhibited over the whole retention region, manifesting the good control of the polymer synthesis process. The molecular weight and PDI data for all polymers are presented in Table 1.

3.2 The self-assembly of PMAOETC-*b*-PNIPAM polymers in aqueous solution

PNIPAM and PMAOETC served as the hydrophilic and hydrophobic blocks in the polymers and are expected to self-assemble into micelles with an appropriate hydrophilic/hydrophobic balance (Scheme 2). As a fluorescent probe, pyrene was introduced into the micelles for encapsulation into the micellar cores to analyze the CMC by virtue of changes in the fluorescence intensity ratio. Excitation spectra at 330 nm, emission spectra ranging from 350–550 nm, and the intensity ratio (I_{383}/I_{372}) *versus* the logarithm of micellar concentration were employed to determine CMC. I_{383}/I_{372} *vs.* log C plots of the polymeric micelles are shown in Fig. 3. The corresponding concentration at the intersection of two tangents of the fitting curve is regarded as the CMC. As such, 19.49, 26.74, and 8.14 mg l⁻¹ are the CMC values corresponding to P1, P2, and P3, respectively (Table 2). Micelles form easily and are dynamically



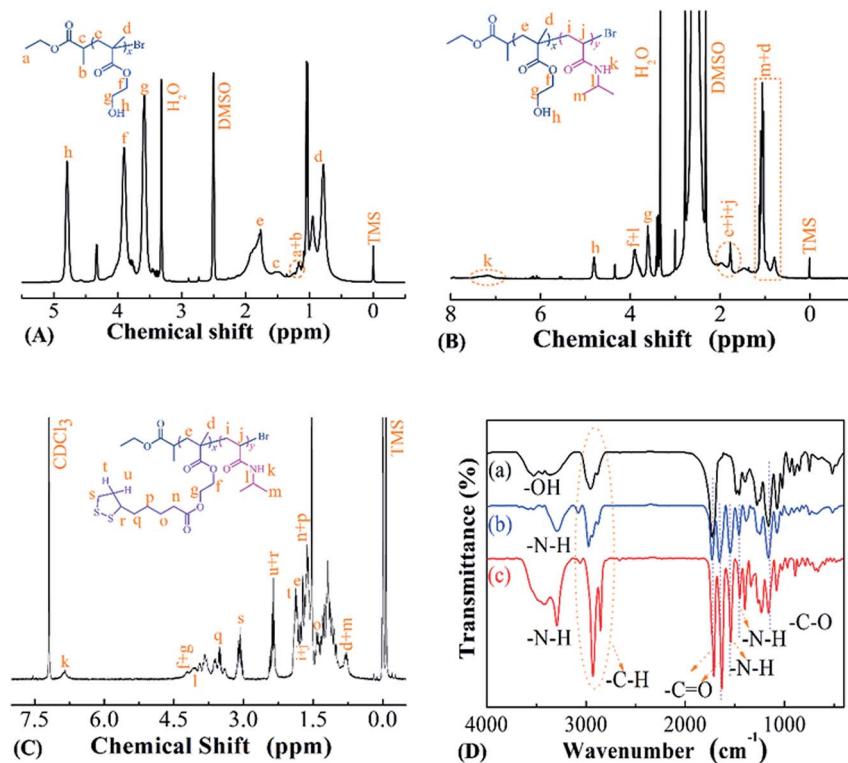


Fig. 1 ¹H NMR spectra of (A) PHEMA-Br, (B) PHEMA-*b*-PNIPAM, and (C) PMAOETC-*b*-PNIPAM. (D) FTIR spectra of (a) PHEMA-Br, (b) PHEMA-*b*-PNIPAM, and (c) PMAOETC-*b*-PNIPAM.

stable throughout multi-dilution from bodily fluids. It can be concluded that the decreased CMC value is due to a decrease in the PNIPAM block concentration and an increase in the PMAOETC segment concentration. Comparing the CMC values of blank micelles and their PTX-loaded counterparts, the slight increases may be due to the participation of hydrophobic PTX and the larger micellar aggregates that are generated. As part of the hydrophobic core, the PTX-loading content directly affects the CMC value, as shown by the PTX-loaded P1 and P2 micelle values of 69.83 and 34.53 mg l⁻¹, respectively. The enhanced hydrophobicity of the P2 micelles originates from the higher

level of PTX encapsulation, leading to a lower CMC value than that of the PTX-laden P1 micelles.

The zeta potential values (ζ , Table 2) illustrate that the polymer micelles are positively charged, which is probably due to the fact that both PMAOETC and PNIPAM are ionized in aqueous solution. The absolute ζ values of around 30 mV suggest that the polymer micelles have favorable stability. Specifically, the absolute value of ζ was enhanced from 29.24 to 36.19 mV upon increasing the length of the NIPAM block, while there was a negligible effect from PMAOETC on ζ in the presence of identical PNIPAM block lengths. Stronger hydration originating from intense inter- and intra-molecular hydrogen bond interactions between NIPAM and H₂O can improve the micellar solution stability. In general, positively charged micelles tend to interact with negatively charged cancer cell membranes, facilitating subsequent PTX release.

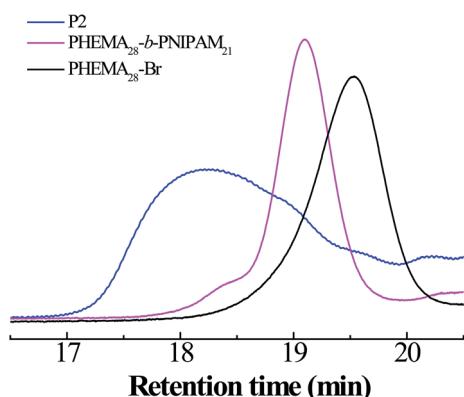
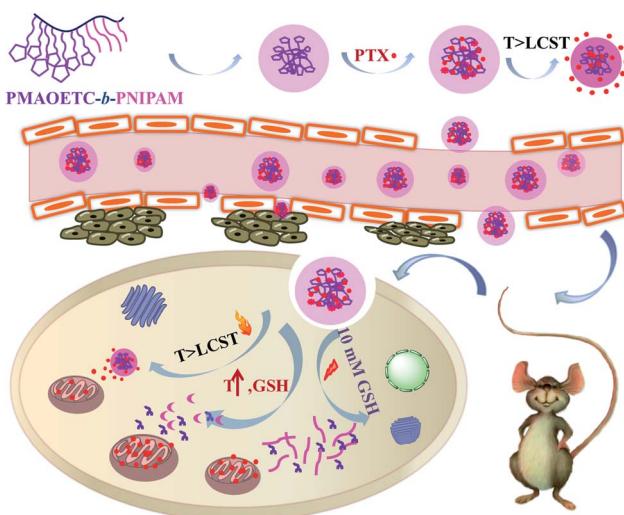


Fig. 2 The GPC curves of PHEMA₂₈-Br, PHEMA₂₈-*b*-PNIPAM₂₁, and PMAOETC₂₈-*b*-PNIPAM₂₁ (P2).



Table 1 The molecular weight parameters of the polymers

Feed molar ratio	Experimental molecular weight from ^1H NMR	Conversion molar ratio (%)	Sample	Molecular weight from GPC		
				M_n	M_w	PDI (M_w/M_n)
$n_{\text{EBP}}/n_{\text{HEMA}} = 1 : 30$	3825	93.3	$\text{PHEMA}_{28}\text{-Br}$	7287	9480	1.30
$n_{\text{EBP}}/n_{\text{HEMA}} = 1 : 50$	5400	80.0	$\text{PHEMA}_{40}\text{-Br}$	10 820	13 525	1.25
$n_{\text{PHEMA}_{30}\text{-Br}}/n_{\text{NIPAM}} = 1 : 100$	4965	10.0	$\text{PHEMA}_{28}\text{-b-PNIPAM}_{10}$	9630	13 675	1.42
$n_{\text{PHEMA}_{30}\text{-Br}}/n_{\text{NIPAM}} = 1 : 200$	6219	10.0	$\text{PHEMA}_{28}\text{-b-PNIPAM}_{21}$	17 909	23 819	1.33
$n_{\text{PHEMA}_{50}\text{-Br}}/n_{\text{NIPAM}} = 1 : 100$	6540	10.0	$\text{PHEMA}_{40}\text{-b-PNIPAM}_{10}$	19 091	23 291	1.22
$n_{\text{PHEMA}_{30}\text{-b-PNIPAM}_{100}} n_{\text{-OH}}/n_{\text{NIPAM}} = 1 : 1.5$	10 761	66.7	$\text{PMAOETC}_{28}\text{-b-PNIPAM}_{10}$	22 024	37 000	1.68
$n_{\text{PHEMA}_{30}\text{-b-PNIPAM}_{200}} n_{\text{-OH}}/n_{\text{NIPAM}} = 1 : 1.5$	12 015	66.7	$\text{PMAOETC}_{28}\text{-b-PNIPAM}_{21}$	29 655	45 075	1.52
$n_{\text{PHEMA}_{50}\text{-b-PNIPAM}_{100}} n_{\text{-OH}}/n_{\text{NIPAM}} = 1 : 1.5$	14 820	66.7	$\text{PMAOETC}_{40}\text{-b-PNIPAM}_{10}$	30 258	50 833	1.68

Scheme 2 A schematic illustration of the self-assembly, PTX loading, accumulation, and targeted release of PMAOETC-*b*-PNIPAM.

solution was treated with 10 mM GSH and analyzed at predetermined time intervals. It is noteworthy that the more thorough the reduction by GSH, the larger the D_h value of the micelles. As seen from the DLS data (Fig. 4A(a-d)), the D_h values of the P2 micelles reduced by GSH at 0, 0.5 h, 4 h, and 24 h are 151.1, 184.3, 233.7, and 521.8 nm, respectively. A globular P2 micelle structure with a PMAOETC core and PNIPAM shell is observed, showing that D_h varies from 65 to 122 nm, centered at

95 nm (Fig. 4B), to 80 to 217 nm, averaging about 210 nm, after GSH reduction for 24 h (Fig. 4C). This change is consistent with DLS analysis. The homogenously dispersed micelles become big micellar aggregates composed of micelles of various sizes after reduction. In fact, disulfide bonds are liable to break into mercaptan in the presence of reduced GSH. The stronger hydrophilicity of mercaptan compared to that of disulfide bonds leads to hydrogen bonds and other interactions and subsequent micellar aggregate formation. In addition, the enhanced hydrophilicity of mercaptan results in stronger swelling, thereby increasing the shell layer thickness and micellar size.²³⁻²⁵ By contrast, the micellar size from DLS is much larger than that captured *via* TEM, which is mainly due to the dehydration of micelle solution that occurs during the process of TEM sample preparation.^{26,27}

After loading PTX onto the polymers, the distinguished core-shell structure of the resulting PTX-laden micelles is remarkable (Fig. 4D); the D_h values of PTX-loaded P2 micelles are distributed within the range of 111–133 nm and focused at 122 nm. The hydrophobic anticancer drug PTX can be encapsulated by means of hydrophobic stacking with PMAOETC segments as the inner core with the PNIPAM layer as an outer shell to stabilize the PTX-laden micelles and further prevent PTX leakage before reaching the cancerous site. After PTX loading into the PMAOETC block, the micellar size increased to 122 nm compared to a size of 95 nm for blank P2 micelles. Hence, it was validated that PTX-laden micelles were successfully prepared.

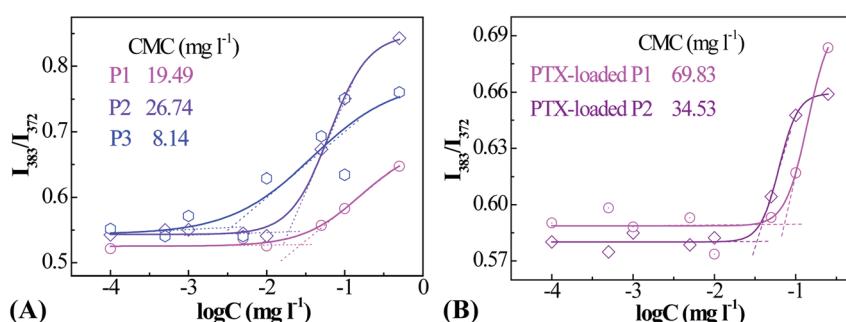
Fig. 3 Fluorescence intensity ratio (I_{383}/I_{372}) vs. logarithm of concentration plots for blank (A) and PTX-laden (B) PMAOETC-*b*-PNIPAM block polymer micelles.

Table 2 The physicochemical parameters of the PMAOETC-*b*-PNIPAM block copolymer micelles

		P1	P2	P3
CMC (mg l ⁻¹)	Blank micelles	19.49	26.74	8.14
	PTX-entrapped micelles	69.83	34.53	—
Zeta potential (ζ , mV)		29.24 \pm 0.90	36.19 \pm 1.38	30.17 \pm 0.69
LCST (°C)		39.48	36.10	—
LC (%)		12.70	22.53	29.36
EE (%)		46.83	69.60	78.33

The morphology of the PTX-laden micelles after release in PBS at pH 5.6 with 10 mM GSH at 42 °C after 83 h is displayed in Fig. 4E. D_h ranged from 81 to 311 nm, concentrated at 194 nm, for the PTX-laden P2 micelles after release under thermal and GSH dual stimulation. Significantly, despite the many holes and alveoli that emerged in the core region of the micelles, the core-shell structure remained. One possible explanation is that the initial hydrophobicity changed when the disulfide bonds were reduced to mercaptan by GSH, followed by the reassembly of the micelles. The uneven holes may be attributed to various degrees of reduction. After 83 h of PTX release, the PTX-laden micelle

size can be up to 194 nm, which is ascribed to two reasons. First, the GSH-induced reduction of disulfide bonds into mercaptan led to enhanced hydrophilicity and swelling, and micellar sizes larger than those of the blank micelles after reduction. Second, at the end of the release process, 40% of PTX still remained in the PMAOETC segments, causing the micellar size to increase to some extent. Thus, the aforementioned results reveal that the PMAOETC-*b*-PNIPAM polymeric micelles can be reduced by GSH in cancerous tissue.

As the thermally sensitive PNIPAM was introduced into polymer, the thermo-sensitivity of the prepared micelles was

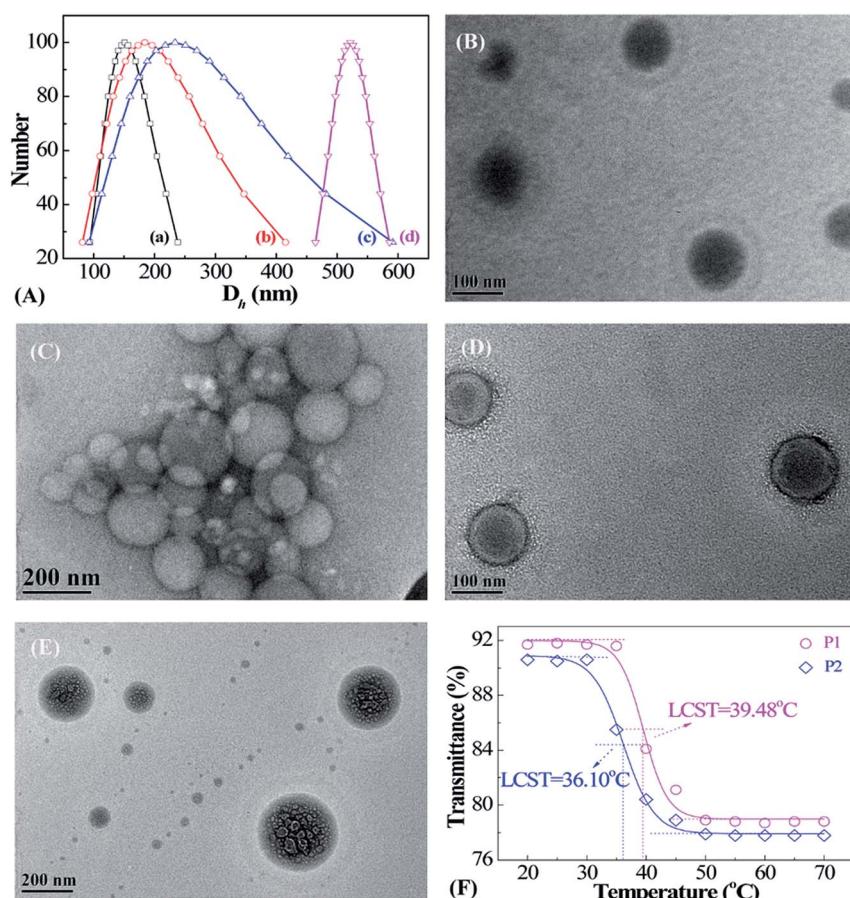


Fig. 4 (A) Dynamic diameter (D_h) values of P2 micelles in aqueous solution containing (a) 0 mM GSH, and 10 mM GSH for (b) 0.5 h, (c) 4 h, and (d) 24 h. TEM photographs of blank P2 micelles treated with (B) 0 mM and (C) 10 mM GSH after 24 h. TEM photographs of (D) PTX-laden P2 micelles and (E) PTX-laden P2 micelles after release in PBS at pH 5.6 containing 10 mM GSH at 42 °C for 85 h. (F) Transmittance–temperature plots of polymeric micelles obtained via UV-vis spectroscopy.



investigated using a UV-vis spectrometer (Fig. 4F). The results indicate that a definite phase transition appears as the temperature increases, and the LCSTs of the P1 and P2 micelles are 39.48 and 36.10 °C, respectively. Obviously, P2 contains more NIPAM units than P1, leading to the higher molecular weight of P2, which results in a lower LCST than P1. In cases in which there are the identical amounts of PMAOETC, the molecular weight of PNIPAM determines the LCST directly, which can be explained using Flory–Huggins theory.²⁸ This theory states that the molar volume ratio, r , of a polymer to a solvent decreases with decreasing molecular weight, and χ_c increases, thus causing P1 to have a higher LCST. This agrees with conclusions drawn by Furyk, Laukkonen, and Xia *et al.*^{29–31} However, P3 does not exhibit obvious thermo-responsive behavior in the temperature range of 20–70 °C, which is mainly attributed to the flexible chains of PMAOETC shielding the PNIPAM blocks. Based on these results, we demonstrated that the polymers containing 28 MAOETC repeat units possess both reduction- and thermo-responsiveness.

3.4 PTX encapsulation and *in vitro* temperature- and reduction-induced PTX release

PTX, as a template anticancer drug, is widely applied using drug delivery systems so that it can have the most optimal effects in cancer therapy. PTX is solubilized for interactions with the hydrophobic PMAOETC core. The D_h value of the homo-dispersed PTX-loaded micelles becomes larger when PTX is introduced into the micelles. Only a limited amount of PTX can be loaded into the PNIPAM section in view of the hydrogen bond interactions between them.²² The LC values of the P1, P2, and P3 micelles are 12.70%, 22.53%, 29.36%, respectively, and the corresponding EE values are 46.82, 69.60%, and 78.33%, respectively (see Table 2). Additionally, a comparison between PTX-laden P1 and P3 micelles shows that the longer the PMAOETC block, the more PTX can be encapsulated into the micelles. Meanwhile, PNIPAM₁₀ and PNIPAM₂₀ possess LC values of 10.31 and 12.64%, respectively, and EE values of 38.43 and 40.62%, respectively, lower than the loading capacities of the polymeric micelles. Both PTX-encapsulating P1 and P2 micelles have sufficient LC and EE values, especially since the longer PNIPAM segment can be conducive to stabilizing the micellar structure and preserving PTX in the micellar core. The results are consistent with the CMC values of the PTX-encapsulated micelles. Taking the PTX content and synergistic responsiveness of the micelles into consideration, the lyophilized PTX-laden P2 micellar powder is suitable for further *in vitro* release studies.

Furthermore, *in vitro* temperature- and reduction-triggered PTX release experiments involving PTX-encapsulating micelles were performed under the following conditions: in PBS (pH 7.4) at 37 °C (almost equivalent to the LCST of P2) or 30 °C (lower than the LCST of P2), and in PBS (pH 5.6) containing 10 mM DTT or 10 mM GSH at 42 °C, corresponding to the tumor intracellular environment. Only approximately 18.47% PTX is isolated from the PTX-loaded micelles in PBS at pH 7.4 and 30 °C over the whole release process, while the accumulated

PTX release amount reached 28.86% in PBS at pH 7.4 and 37 °C (Fig. 5(a and b)). It is indicated that the PNIPAM blocks respond to temperature changes and a phase transition happens afterwards, leading to micellar shrinkage and the subsequent diffuse release of PTX. These results show that PTX is steadily attached to the micelles in normal tissue. Furthermore, the PTX release amount sharply increases to 37.21% at 48 h and 61.03% at 83 h upon the addition of 10 mM DTT at 42 °C (Fig. 5(c)). For the release results under GSH stimulation at 42 °C, the cumulative release of PTX reaches 33.27% within 34 h and tends to even out thereafter, reaching 40.87% in the end (Fig. 5(d)). The higher amount of PTX release from PTX-laden P2 micelles in 10 mM DTT compared to isometric GSH is due to the stronger reduction abilities of DTT. In our study, the PTX-laden P2 micelles were shown to respond satisfactorily to characteristic reductants and increased temperatures at tumor sites.

3.5 The cytotoxicity of PTX-laden polymeric micelles

As a drug delivery vehicle, it is essential to evaluate the cytotoxicity of PTX-laden P2 micelles against HCT116 cells. Free PTX presents dose-dependent cytotoxicity in the concentration range from 0.1 to 20 mg l⁻¹, and the IC₅₀ value was deduced to be approximately 3.29 µg ml⁻¹ (Fig. 6). However, the empty micelles exhibit relatively lower cytotoxicity than that of free PTX, achieving almost 90% cell viability, and some surpass 100% when the micellar concentration is over 300 mg l⁻¹. This may be ascribed to the function of TA in facilitating cell growth. For PTX-loaded micelles, the slight decrease in cell viability is not only caused by a tiny amount of PTX being released under the stimulation of reduced GSH in cancer cells, but it is also due to a trace amount of PTX leaking from the micelles under the culture conditions (PBS at pH 7.4 and 37 °C), which is in accordance with the results illustrated in Fig. 5(a). We demonstrate that the empty micelles are nontoxic, and the PTX-laden polymeric micelles can be used as PTX nanocarriers.

3.6 *In vivo* studies

3.6.1 Anticancer efficacy. The anticancer efficacy of free PTX and PTX-laden micelles were estimated in 4T1-tumor-

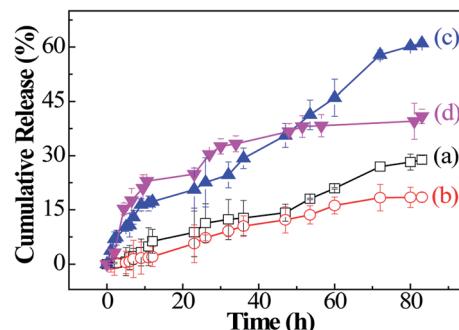


Fig. 5 *In vitro* PTX release plots of PTX-laden P2 micelles in PBS (pH 7.4) at 37 °C (a), PBS (pH 7.4) at 30 °C (b), PBS (pH 5.6) containing 10 mM DTT at 42 °C (c), and PBS (pH 5.6) containing 10 mM GSH at 42 °C (d).



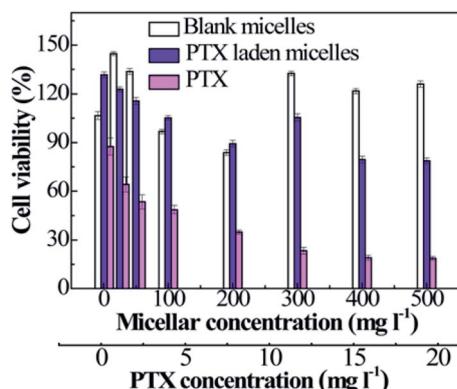


Fig. 6 The viabilities of HCT116 cells treated with free PTX, blank micelles, and PTX-loaded micelles as a function of the micellar or PTX concentration ($n = 3$).

bearing Balb/c mice were studied to show the relationship between tumor volume/body weight and treatment time. In this study, normal saline and PBS (control), free PTX, blank micelles, and PTX-entrapped P2 micellar solutions were injected intravenously. The relative tumor volumes (V_0 and V denote the volume initially and after the measurement period, respectively) and mice body weights (W_0 and W signify the body weight initially and after the measurement period, respectively)

were calculated every three days (Fig. 7A and B). As expected, treatment using PTX-loaded micelles caused a distinguishable suppression in tumor volume, 1.34 times higher than the free-PTX-treated group (Fig. 7A). From the results displayed in Fig. 4E and 5, the synergistic effect of reduced GSH, DTT, and temperatures higher than the LCST led to the rupturing of the micelles and simultaneous PTX release, thereby causing tumor cell apoptosis. Free PTX did not achieve an improved anticancer effect because of its strong hydrophobicity. Moreover, the body weights of mice treated with the PTX-entrapped micelles exhibited a promising trend (Fig. 7B), reflecting that the dual-responsive micelles can circumvent the side effects of therapeutic agents such as PTX.

3.6.2 Histological assays. Haematoxylin and eosin (H&E) staining was employed to study the histological characteristics of excised major organs (hearts, livers, spleens, lungs, and kidneys) from 4T1-tumor-bearing mice treated with PTX-laden nanoformulations. The typical histological features are presented in Fig. 7C. The results indicate the presence of distinct heart damage caused by free PTX,³² however, no pathological abnormalities appear in the mice treated with the PTX-laden micelles in comparison with the control groups. Thus, the PTX-loaded micelles can potentially be applied as an effective anticancer formulation *in vivo* due to the circumvention of cardiotoxicity from PTX. Apart from enhanced PTX

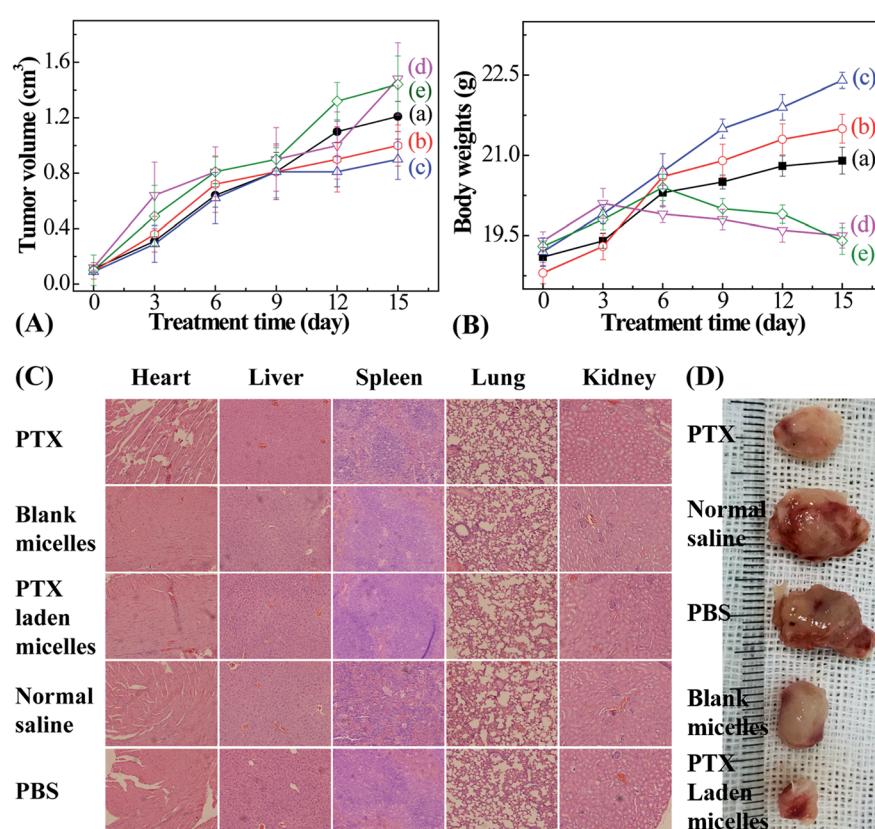


Fig. 7 Tumor volume (A) and body weight (B) trends of 4T1-tumor-bearing Balb/c mice after the intravenous injection of free PTX (a), blank micelles (b), PTX-entrapped micelles (c), normal saline (d), and PBS (e) at a PTX dose of 5 mg kg^{-1} . Histological images of sliced heart, liver, spleen, lung, and kidney samples from mice on day 15 (200 \times) (C), and the excised tumor sizes after 15 days of treatment (D).



accumulation at cancerous sites under thermal and reduction stimuli, the negligible pre-leakage and low retention of drugs in healthy organs and tissue mean that the carrier can circumvent the severe side effects of free PTX. These results highlight the biosafety and anticancer efficacy of the PTX-laden nanoformulations. The volumes of the excised solid tumors seen in Fig. 7D are consistent with the tumor size and mice body weight trends seen during the overall treatment process (Fig. 7A and B).

4. Conclusions

In summary, a type of temperature- and reduction-responsive amphiphilic block polymer, PMAOETC-*b*-PNIPAM, was prepared *via* an “ATRP polymerization–esterification” strategy and utilized for PTX delivery. Through synergistic loading using the PMAOETC and PNIPAM blocks, the PTX-laden nanomicelles displayed an improved drug loading capacity (LC of 29.36% and EE of 78.33%) and favorable stability (ζ of approximately 30 mV). Additionally, under the trigger of 10 mM DTT at 42 °C, about 61.03% of the total PTX was released from the carriers in a sustainable manner within 83 h. Furthermore, the PTX-laden nanomicelles circumvented the systematic toxicity of free PTX and even caused a remarkable suppression in tumor volume, with tumors 1.34 times smaller than the free-PTX-treated group during *in vivo* studies. Given these results, this “ATRP polymerization–esterification” method is feasible for the formation of a DDS with an appropriate drug content that can achieve satisfactory therapeutic effects.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work is supported by the Fundamental Research Funds for the Central Universities (No. GK201601003), the National Natural Science Foundation of China (No. 21572123), and the Natural Science Research Fund of Shaanxi University of Science and Technology (2019BJ-43).

References

- 1 S. M. Elbashir, J. Harborth, W. Lendeckel, A. Yalcin, W. Klaus and T. Tuschl, *Nature*, 2001, **411**, 494–498.
- 2 M. I. Yatoo, A. Saxena, M. H. Malik, M. K. Sharma and U. Dimri, *J. Anim. Sci. Adv.*, 2014, **4**, 705–709.
- 3 D. Mei, Z. Lin, J. Fu, B. He, W. Gao, L. Ma, W. Dai, H. Zhang, X. Wang, J. Wang, X. Zhang, W. Lu, D. Zhou and Q. Zhang, *Biomaterials*, 2015, **42**, 52–65.
- 4 F. Danhier, P. Danhier, C. J. De Saedeleer, A.-C. Fruytier, N. Scheleiche, A. Rieux, P. Sonveaux, B. Gallez and V. Préat, *Int. J. Pharm.*, 2015, **479**, 399–407.
- 5 Y. Li, J. Lin, H. Wu, Y. Chang, C. Yuan, C. Liu, S. Wang, Z. Hou and L. Dai, *Mol. Pharm.*, 2015, **12**, 769–782.
- 6 L. Palanikumar, E. S. Choi, J. Y. Oh, S. A. Park, H. Choi, K. Kim, C. Kim and J.-H. Ryu, *Biomacromolecules*, 2018, **19**, 3030–3039.
- 7 G. Verma and P. A. Hassan, *Phys. Chem. Chem. Phys.*, 2013, **15**, 17016–17028.
- 8 V. C. Yang, *J. Controlled Release*, 2019, **311–312**, 322–323.
- 9 B. Yang, Y. Lv, J. Zhu, Y. Han, H. Jia, W. Chen, J. Feng and X. Zhang, *Acta Biomater.*, 2014, **10**, 3686–3695.
- 10 C. T. Nguyen, T. H. Tran, M. Amiji, X. Lu and R. M. Kasi, *Nanomedicine*, 2015, **11**, 2071–2082.
- 11 H. Chen, S. Kim, L. Li, S. Wang, K. Park and J.-X. Cheng, *Proc. Natl. Acad. Sci. U. S. A.*, 2008, **105**, 6596–6601.
- 12 Y. Zhang, T. Ren, J. Gou, L. Zhang, X. Tao, B. Tian, P. Tian, D. Yu, J. Song, X. Liu, Y. Chao, W. Xiao and X. Tang, *J. Controlled Release*, 2017, **261**, 352–366.
- 13 W. Yuan, L. Li and H. Zou, *RSC Adv.*, 2015, **5**, 80264–80268.
- 14 T. Yin, J. Wang, L. Yin, L. Shen, J. Zhou and M. Huo, *Polym. Chem.*, 2015, **6**, 8047–8059.
- 15 Q. Zhou, Y. Hou, L. Zhang, J. Wang, Y. Qiao, S. Guo, L. Fan, T. Yang, L. Zhu and H. Wu, *Theranostics*, 2017, **7**, 1806–1819.
- 16 M. Simona, N. Julien and C. Patrick, *Nat. Mater.*, 2013, **12**, 991–1003.
- 17 F. Q. Schafer and G. R. Buettner, *Free Radical Biol. Med.*, 2001, **30**, 1191–1212.
- 18 Z. Zhou, Y. Shen, J. Tang, M. Fan, E. A. V. Kirk, W. J. Murdoch and M. Radosz, *Adv. Funct. Mater.*, 2009, **19**, 3580–3589.
- 19 P. Sun, H. Du, L. Gao, W. Zhu, X. Li and Z. Shen, *Acta Polym. Sin.*, 2012, **8**, 789–793.
- 20 Y. Wang, H. Du, L. Gao, H. Ni, X. Li, W. Zhu and Z. Shen, *Polym. Chem.*, 2013, **4**, 1657–1663.
- 21 J. Y. Teo, W. Chin, X. Ke, S. Gao, S. Liu, W. Cheng, J. L. Hedrick and Y. Y. Yang, *Nanomedicine*, 2017, **13**, 431–442.
- 22 J.-W. Xu, X. Ge, L.-H. Lv, F. Xu and Y.-L. Luo, *Macromol. Rapid Commun.*, 2018, **39**, 1800628.
- 23 C. Wu, J. Yang, X. Xu, C. Gao, S. Lü and M. Liu, *Eur. Polym. J.*, 2016, **83**, 230–243.
- 24 L. Wu, Y. Zou, C. Deng, R. Cheng, F. Meng and Z. Zhong, *Biomaterials*, 2013, **34**, 5262–5272.
- 25 R. Wei, L. Cheng, M. Zheng, R. Cheng, F. Meng, C. Deng and Z. Zhong, *Biomacromolecules*, 2012, **13**, 2429–2438.
- 26 J. Ding, J. Chen, D. Li, C. Xiao, J. Zhang, C. He, X. Zhuang and X. Chen, *J. Mater. Chem. B*, 2013, **1**, 69–81.
- 27 J. Chang, Y. Li, G. Wang, B. He and Z. Gu, *Nanoscale*, 2013, **5**, 813–820.
- 28 K. C. Tam and S. Dai, *Langmuir*, 2003, **19**, 5175–5177.
- 29 S. Furyk, Y. Zhang, D. Ortiz-Acosta, P. S. Cremer and D. E. Bergbreiter, *J. Polym. Sci., Part A: Polym. Chem.*, 2006, **44**, 1492–1501.
- 30 A. Laukkonen, L. Valtola, F. Winnik and H. Tenhu, *Macromolecules*, 2004, **37**, 2268–2274.
- 31 Y. Xia, A. D. N. Burke and H. Stöver, *Macromolecules*, 2006, **39**, 2275–2283.
- 32 T. Yin, L. Wang, L. Yin, J. Zhou and M. Cao, *Biomaterials*, 2015, **61**, 10–25.

