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Dual acid/glutathione-responsive core-degradable/shell-sheddable block copolymer nanoassemblies bearing benzoic imines for enhanced drug release†

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The development of amphiphilic block copolymer (ABP)-based nanoassemblies that degrade in response to dual stimuli at dual locations (e.g. hydrophobic cores and core/corona interfaces) offers a promising platform for controlled drug delivery. This work harnesses the features of an acid-labile benzoic imine (BzIm) bond and a glutathione (GSH)-cleavable disulfide linkage. We synthesized a poly(ethylene glycol) (PEG)-based dual location dual acid/GSH-degradable ABP with BzIm pendants in a hydrophobic poly-methacrylate block and a disulfide at the block junction. The acid-catalyzed hydrolysis rate of BzIm depends on substituents attached to its *para*-position. Hydrolysis is faster with electron-donating substituents (methoxy), and slower with electron-withdrawing ones (bromo and nitro). Well-defined ABP synthesized by reversible addition–fragmentation chain transfer (RAFT) polymerization of a methoxy-substituted BzIm methacrylate in the presence of a disulfide-labeled PEG-based chain transfer agent, enables self-assembly to form colloidal-stable, monomodal, and spherical nanoassemblies. These nanoassemblies are capable of encapsulating cancer drug doxorubicin (Dox) and exhibit enhanced release of Dox through core degradation upon the cleavage of BzIm bonds in acidic pH and shell detachment upon the cleavage of disulfide bonds in the presence of GSH. Moreover, Dox-loaded nanoassemblies show excellent uptake by HeLa cell multi-tumor spheroids, demonstrating their potential as drug delivery nano-carriers. This study highlights the importance of substituent effects on the hydrolysis of BzIm and the dual acid/GSH-responsive strategy for developing a promising drug delivery system with precise control over drug release.

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Introduction

Amphiphilic block copolymers (ABPs) that exhibit stimuli-responsive degradation (SRD) have emerged as a promising platform for controlled drug delivery.^{1–3} The general synthetic approach for SRD-exhibiting ABPs involves polymerization in the presence of cleavable bond-functionalized initiators or monomers *via* reversible-deactivation radical polymerization (RDRP), typically by atom transfer radical polymerization (ATRP) and reversible addition–fragmentation chain transfer (RAFT).^{4–8} This enables the incorporation of cleavable bonds into polymer precursors at various positions, such as the backbone, junction, and side chain.⁹ The intrinsic amphiphilicity of ABPs allows them to self-assemble to form stable aqueous nanoassemblies.^{10–12} In response to specific stimuli, the cleavage of SRD linkages causes the disassembly of nanoassemblies through changes in hydrophobic/hydrophilic balance, main chain degradation, or corona detachment, leading to site-specific and enhanced release of encapsulated drugs.^{13–15}

When SRD-based polymeric nanoassemblies are designed for intracellular delivery, the selection of cleavable linkages is critical. Acidic pH is a stimulus found in subcellular and tissue microenvironments, such as endosomes and lysosomes (pH = 4–5.5), and tumors (pH = 6.5–6.8), respectively.^{16–18} Various covalent bonds that are cleavable in acidic pH ranges have been explored, including ketal, acetal, 2,3-dialkylmaleamic amide, β -thiopropionate, oxime, hydrazone, and benzoic imine (BzIm) linkages.¹⁹ Among these pH-sensitive linkers, BzIm (C=N) bonds have attracted particular interest due to their facile synthesis *via* a catalyst-free click-type reaction between carbonyl and amine groups.² In an acidic environment, the C=N bond is cleaved into the corresponding carbonyl and amine precursors. As acid-responsive degradation directly regulates drug release process *via* degradation-induced disassembly, it is critical to control their cleavage kinetics by

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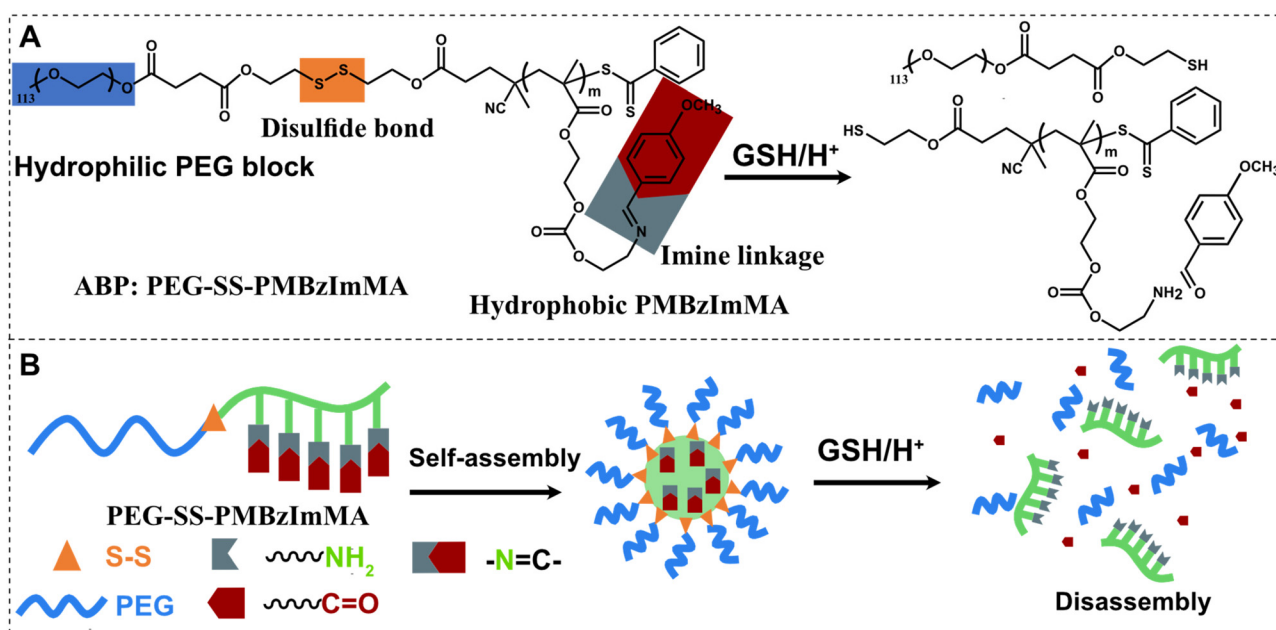
chemical design. Mechanistic studies show that C=N hydrolysis is initiated by protonation of the N atom and followed by nucleophilic attack of H₂O. Consequently, the rate of acid catalyzed imine cleavage depends on electrophilicity, which can be tuned by substituent effects on the C=N bond and phenyl ring for BzIm *via* inductive, resonance, and steric effects.²⁰

Built on these promising acid-responsive properties, the development of BzIm-based (co)polymers has been extensively explored as acid-responsive drug delivery nanocarrier in various forms including self-assembled nanoassemblies,^{21,22} core-crosslinked nanogels,^{23,24} and polymer-drug conjugates.^{2,25} These systems are generally fabricated by post-polymerization modification of either amine or aldehyde-labelled (co)polymer precursors. Recently, our group demonstrated direct polymerization²² and post-polymerization modification²¹ approaches as effective means to synthesize a well-defined poly(ethylene glycol)-based ABP containing hydrophobic poly(methacrylate) with pendant BzIm groups. Despite these advances, BzIm-bearing nanoassemblies suffered from slow release and aggregation. More precise control over degradation behavior of imine and resulting drug release is highly demanded. Inspired by the structure-dependent hydrolysis of BzIm bonds,²⁶ we hypothesized that *para*-substituents on BzIm groups with varying electronic properties (electro-donating or withdrawing) could tune their acid-catalyzed cleavage kinetics and thus drug release profiles.

Further studies have explored polymers with dual or multiple stimuli-responsive degradation mechanisms.^{27,28} Glutathione (GSH), a tripeptide with a cysteine residue and a

pendant thiol group, is a reducing agent that is found at millimolar concentrations in the cytosol of cells, and can be elevated in cancer cells.^{29–32} GSH-responsive disulfide (SS) group has been explored together with a BzIm linkage for the construction of dual acid/GSH-responsive degradable ABP nanoassemblies.^{33–38} However, these nanoassemblies were designed with both BzIm and disulfide linkages in a single location, either at core/corona interface or in hydrophobic core. We postulated that the integration of SS bonds into the block junction of PEG-based ABP-bearing pendant BzIm bonds could facilitate synergistic effects for controlled drug release. This strategy known as dual location dual acid/glutathione-responsive degradation causes dual responses at dual locations as in the cores and at core/corona interfaces, thus enabling to not only independently regulate the release of encapsulated molecules but also facilitate synergistic/accelerated release at dual locations.^{39–41}

In this work, we explored tunable imine hydrolysis and dual location dual acid/GSH response to construct dual location dual acid/GSH-degradable ABPs and their nanoassemblies for enhanced control over drug release (Scheme 1). A series of BzIm-bearing methacrylate monomers with various *para* substituents (RBzImMA) (R = MeO, H, NO₂, and Br) were synthesized to investigate the effect of their electronic properties on acid-catalyzed hydrolysis rates. Using a new imine methacrylate with MeO substituent (*e.g.* MBzImMA) with enhanced acid-responsive hydrolysis, well-defined PEG-based ABPs with pendant BzIm groups in the hydrophobic block and a SS positioned at the block junction was synthesized by a controlled radical polymerization technique. The copolymer was charac-



Scheme 1 Illustration of dual location dual acid/GSH-degradable ABP with BzIm pendants in the hydrophobic block and a disulfide bond at the block junction (A) and its nanoassemblies exhibiting synergistic/enhanced release of encapsulated drug molecules in dual acid/GSH responses at dual locations as in cores and at core/corona interfaces (B).



terized for aqueous self-assembly, dual acid/GSH-responsive degradation and drug release, as well as cellular uptake with a HeLa multi-cellular tumor spheroid model. Our results show that the dual acid/GSH-responsive nanoassemblies enabled synergistic control over drug release and minimized degradation products. Drug-loaded nanoassemblies demonstrated superior uptake by HeLa spheroids compared to free drug.

Experimental

Instrumentation and materials

NMR spectra were recorded using 500 MHz Varian spectrometer. The CDCl_3 singlet at 7.26 ppm was selected as the reference standard. Molecular weight and molecular weight distribution were determined using gel permeation chromatography (GPC) with Malvern equipped with a Viscotek VE1122 pump and a refractive index detector. Tetrahydrofuran (THF) was used as eluent at room temperature at a flow rate of 1.0 mL min^{-1} . Molecular weight and molecular weight distribution were calculated relative to linear poly(methyl methacrylate) standards from Fluka. Aliquots of the polymer samples dissolved in THF were filtered through a 0.45 μm PTFE filter to remove any THF-insoluble species and injected into the GPC with a drop of anisole as a flow rate marker.

Dynamic light scattering (DLS) with a Malvern Instruments Nano S ZEN1600, transition electron microscope (TEM) using a Talos L120C TEM operated at 80 kV electrons, fluorescence spectroscopy with Varian Cary Eclipse fluorescence spectrometer, and UV/vis spectroscopy with Agilent Cary 60 UV/vis spectrometer were used as described in our previous publications.^{42,43}

Most reagents including triethylamine (Et_3N , 99.5%), carbonyl diimidazole (CDI, $\geq 90\%$), 2-hydroxyethyl methacrylate (HEMA, $\geq 99\%$), ethanolamine (EA, $\geq 98\%$), 4-methoxybenzaldehyde (98%), 4-nitrobenzaldehyde (98%), 4-bromobenzaldehyde (99%), 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU, 98%), succinic anhydride (SA, $\geq 99\%$), 4-cyano-4-(phenylcarbo-*nothioylthio*) pentanoic acid (CTA, 97%), 4-(methylamino)pyridine (DMAP, 99%), 1-ethyl-3-(3 dimethylaminopropyl) carbodiimide-HCl (EDC), acetyl chloride ($\geq 98\%$), ethylene glycol vinyl ether ($\geq 97\%$), pyridinium *p*-toluenesulfonate (PPTS, 98%), doxorubicin (Dox, $-\text{NH}_3^+\text{Cl}^-$ form, $>98\%$), and deuterium chloride solution (DCl, 35 wt% in D_2O) from Sigma-Aldrich as well as 2,2'-azodi(2-methylbutyronitrile) (AMBN) from Wako Chemicals were purchased and used as received. Poly(ethylene glycol) monomethyl ether (PEG, MW = 5000 g mol^{-1} , EO# = 113) purchased from Sigma-Aldrich was dried by azeotropic distillation with toluene to remove residual moisture. A carbonylimidazole (CI)-activated 2-hydroxyethyl-methacrylate (HEMA-CI) were synthesized as described in our previous publication.^{22,44}

Synthesis of RBzImOH precursors

RBzImOH intermediates were synthesized based on our previous report with R-substituted benzaldehyde.²² A solution of

EA (1.22 g, 0.02 mol) dissolved in DCM (5 mL) was mixed with a solution containing R-benzaldehyde dissolved in DCM (50 mL) in presence of sodium sulfate (5 g) under stirring at room temperature for 13 h. Solvent was removed by rotary evaporation and residue was dried in a vacuum oven at room temperature for 13 h.

MBzImOH. 4-Methoxybenzaldehyde (2.72 g, 0.02 mol). Yield = 3.4 g (96%). $^1\text{H-NMR}$ (CDCl_3 , ppm): 8.15 (s, 1H, $-\text{NCHC}_6\text{H}_4-$), 7.6 and 6.9 (d, 4H, $-\text{NCHC}_6\text{H}_4\text{OCH}_3$), 3.85 (t, 2H, $\text{HOCH}_2\text{CH}_2\text{NCH}-$), 3.8 (s, 3H, $-\text{C}_6\text{H}_4\text{OCH}_3$), 3.66 (t, 2H, $\text{HOCH}_2\text{CH}_2\text{NCH}-$).

NBzImOH. 4-Nitrobenzaldehyde (3.02 g, 0.02 mol). Yield = 3.8 g (98%). $^1\text{H-NMR}$ (CDCl_3 , ppm): 8.4 (s, 1H, $-\text{NCHC}_6\text{H}_4\text{NO}_2$), 8.22 and 7.88 (d, 4H, $-\text{C}_6\text{H}_4\text{NO}_2$), 3.93 (t, 2H, $\text{HOCH}_2\text{CH}_2-$), 3.81 (t, 2H, $\text{HOCH}_2\text{CH}_2-$).

BrBzImOH. 4-Bromobenzaldehyde (3.70 g, 0.02 mol). Yield = 4.3 g (94%). $^1\text{H-NMR}$ (CDCl_3 , ppm): 8.21 (s, 1H, $-\text{NCHC}_6\text{H}_4\text{Br}$), 7.54 (q, 4H, $-\text{C}_6\text{H}_4\text{Br}$), 3.89 (t, 2H, $\text{HOCH}_2\text{CH}_2-$), 3.71 (t, 2H, $\text{HOCH}_2\text{CH}_2-$).

Synthesis of RBzImMA

Substituted benzoic imine methacrylates (**RBzImMAs**) were synthesized based on our previous report using **RBzImOH**.²² The dried **RBzImOH** precursor was mixed with a solution consisting of HEMA-CI and DBU dissolved in DCM under stirring at room temperature for 13 h. The reaction mixture was washed with saturated NaHCO_3 solution three times and dried over sodium sulfate. After the removal of solvent, the product was dried in a vacuum oven for 13 h.

NBzImMA. **NBzImOH** (1.74 g, 9 mmol), HEMA-CI (2.02, 9 mmol), DBU (0.27 g, 0.2 mmol), and DCM (100 mL). Yield = 3.1 g (98%). $^1\text{H-NMR}$ (CDCl_3 , ppm): 8.4 (s, 1H, $-\text{NCHC}_6\text{H}_4-$), 8.2 and 7.9 (d, 4H, $-\text{NCHC}_6\text{H}_4\text{NO}_2$), 6.1 and 5.5 (s, 2H, $\text{CH}_2\text{C}(\text{CH}_3)-$), 4.5 (t, 2H, $-\text{CH}_2\text{CH}_2\text{OC}(\text{O})\text{O}$), 4.3 (m, 2H, $-\text{CH}_2\text{CH}_2\text{OC}(\text{O})\text{O}$), 4.3 (m, 2H, $-\text{CH}_2\text{CH}_2\text{NCH}-$), 3.9 (t, 2H, $-\text{CH}_2\text{CH}_2\text{NCH}-$), 1.9 (s, 3H, $\text{CH}_2\text{C}(\text{CH}_3)-$).

MBzImMA. **MBzImOH** (0.16 g, 0.9 mmol), HEMA-CI (0.20 g, 0.9 mmol), DBU (0.03 g, 0.18 mmol), and DCM (10 mL). Yield = 0.27 g (89%). $^1\text{H-NMR}$ (CDCl_3 , ppm): 8.2 (s, 1H, $-\text{NCHC}_6\text{H}_4-$), 7.6 and 6.9 (d, 4H, $-\text{NCHC}_6\text{H}_4\text{OCH}_3$), 6.1 and 5.5 (s, 2H, $\text{CH}_2\text{C}(\text{CH}_3)-$), 4.4 (t, 2H, $-\text{CH}_2\text{CH}_2\text{OC}(\text{O})\text{O}$), 4.3 (m, 2H, $-\text{CH}_2\text{CH}_2\text{OC}(\text{O})\text{O}$), 4.3 (m, 2H, $-\text{CH}_2\text{CH}_2\text{NCH}-$), 3.8 (m, 5H, $-\text{CH}_2\text{CH}_2\text{NCHC}_6\text{H}_4\text{OCH}_3$), 1.9 (s, 3H, $\text{CH}_2\text{C}(\text{CH}_3)-$).

BrBzImMA. **BrBzImOH** (2.05 g, 9 mmol), HEMA-CI (2.01 g, 9 mmol), DBU (0.27 g, 1.8 mmol), and DCM (10 mL). Yield = 3.3 g (95%). $^1\text{H-NMR}$ (CDCl_3 , ppm): 8.21 (s, 1H, $-\text{NCHC}_6\text{H}_4-$), 7.54 (q, 4H, $-\text{C}_6\text{H}_4\text{Br}$), 6.1 and 5.6 (s, 2H, $\text{CH}_2\text{C}(\text{CH}_3)-$), 4.5 (t, 2H, $-\text{CH}_2\text{CH}_2\text{OC}(\text{O})\text{O}$), 4.3 (m, 2H, $-\text{CH}_2\text{CH}_2\text{OC}(\text{O})\text{O}$), 4.3 (m, 2H, $-\text{CH}_2\text{CH}_2\text{NCH}-$), 3.85 (m, 5H, $-\text{CH}_2\text{CH}_2\text{NCHC}_6\text{H}_4\text{OCH}_3$), 1.9 (s, 3H, $\text{CH}_2\text{C}(\text{CH}_3)-$).

Synthesis of PEG-COOH

SA (0.22 g, 2.2 mmol) dissolved in anhydrous DCM (5 mL) was mixed with an organic solution containing PEG (2 g, 0.4 mmol) and DMAP (49 mg, 0.4 mmol) in DCM (15 mL) in an ice-bath under stirring for 48 h. The resultant mixture was washed with



brine solution three times, dried over sodium sulfate, and then precipitated from cold diethyl ether to remove excess succinic anhydride. The product, white solid, was isolated by vacuum filtration and dried in a vacuum oven at room temperature for 13 h. Yield = 1.35 g (65%). $^1\text{H-NMR}$ (CDCl_3 , ppm): 4.25 (t, 2H, $-\text{CH}_2\text{OC}(\text{O})-$), 3.5–3.8 (m, $-\text{CH}_2\text{CH}_2\text{O}-$ of PEG main chain), 3.36 (s, 3H, CH_3-), 2.6 (m, 4H, $-\text{CH}_2\text{CH}_2\text{COOH}$).

Synthesis of PEG-SS-CTA

In the first step to synthesize CTA-ss-OH, CTA (0.2 g, 0.72 mmol) dissolved in anhydrous DCM (15 mL) was mixed with an organic solution containing ss-diol (166 mg, 1.1 mmol), EDC (270 mg, 1.4 mmol) and TEA (145 mg, 2 mmol) in DCM (85 mL) in an ice-bath under stirring. After proceeding at room temperature for 48 h, the solvent was evaporated, and the product was purified by silica gel column chromatography. The product, a red oil, was collected as the third band of total five bands. Yield = 0.06 g (30%); R_f = 0.35 (ethyl acetate/hexane = 1/1, v/v). $^1\text{H-NMR}$ (CDCl_3 , ppm): 7.9–7.3 (m, 5H, $-\text{SC}(\text{S})\text{C}_6\text{H}_5$), 4.4 (t, 2H, $-\text{CH}_2\text{OC}(\text{O})-$), 3.9 (t, 2H, $-\text{CH}_2\text{CH}_2\text{OH}$), 2.94 (t, 2H, $-\text{CH}_2\text{SSCH}_2\text{CH}_2\text{OH}$), 2.88 (t, 2H, $-\text{SSCH}_2\text{CH}_2\text{OH}$), 2.7 (t, 2H, $-\text{CH}_2\text{CH}_2(\text{CH}_3)\text{C}(\text{CN})-$), 2.4 (t, 2H, $-\text{CH}_2\text{CH}_2(\text{CH}_3)\text{C}(\text{CN})-$), 1.94 (s, 3H, $-(\text{CH}_3)\text{C}(\text{CN})-$).

In the second step to synthesize PEG-ss-CTA, EDC (79 mg, 0.41 mmol) dissolved in anhydrous DCM (20 mL) was mixed with an organic solution containing PEG-COOH (0.5 g, 98 μmol), DMAP (3.6 g, 29 μmol) and CTA-ss-OH (0.16 g, 0.39 mmol) in DCM (80 mL) in an ice-bath under stirring for 24 h. The resultant mixture was washed with brine solution three times, dried over sodium sulfate, and then precipitated from cold diethyl ether to remove excess CTA. The pink solid product was isolated by vacuum filtration and dried in a vacuum oven at room temperature for 13 h. Yield = 0.46 g (70%). $^1\text{H-NMR}$ (CDCl_3 , ppm): 7.9–7.3 (m, 5H, $-\text{SC}(\text{S})\text{C}_6\text{H}_5$), 4.34 (m, 4H, $-\text{CH}_2\text{OC}(\text{O})-$), 4.24 (t, 2H, $-\text{OCH}_2\text{CH}_2\text{OC}(\text{O})-$), 3.5–3.8 (m, $-\text{CH}_2\text{CH}_2\text{O}-$ of PEG main chain), 3.38 (s, 3H, $\text{CH}_3\text{O}-$), 2.92 (t, 4H, $-\text{CH}_2\text{SSCH}_2$), 2.7 (m, 2H, $-\text{CH}_2\text{CH}_2(\text{CH}_3)\text{C}(\text{CN})-$), 2.69 (m, 4H, $-\text{OC}(\text{O})\text{CH}_2\text{CH}_2\text{C}(\text{O})\text{O}-$), 2.4 (t, 2H, $-\text{CH}_2\text{CH}_2(\text{CH}_3)\text{C}(\text{CN})-$), 1.94 (s, 3H, $-(\text{CH}_3)\text{C}(\text{CN})-$).

RAFT polymerization to synthesize PEG-SS-PMBzImMA (ImSSP)

MBzImMA (0.55 g, 1.64 mmol), PEG-SS-CTA (0.12 g, 22 μmol), AMBN (1.08 mg, 7 μmol) and anhydrous anisole (0.92 mL) were mixed in a 5 mL Schlenk flask. The mixture was deoxygenated by purging with nitrogen for 45 min, and then immersed into an oil bath at 70 $^\circ\text{C}$ to initiate polymerization. For kinetics study, aliquots were taken out periodically to follow monomer conversion by $^1\text{H-NMR}$ and molecular weight by GPC. Polymerization was stopped by cooling the reaction mixture in an ice-bath and exposing to air.

Investigation of acid-catalyzed imine degradation using $^1\text{H-NMR}$ spectroscopy

Hydrolysis in acidic conditions were carried out based on our previous report.²² For the hydrolysis of monomers, **RBzImMA**

(10.0 μmol) dissolved in CD_3CN (0.3 mL) was mixed with the as-prepared PB solutions in D_2O (0.1 mL, 0.2 M, pD = 5.0, 6.8 and 7.4). For PEG-SS-pMBzImMA block copolymer, the purified ImSSP (10 mg) dissolved in CD_3CN (0.3 mL) was mixed with the PB solutions in D_2O (0.1 mL, 0.2 M, pD = 5). The resultant mixtures were subjected to $^1\text{H-NMR}$ analysis at 25 $^\circ\text{C}$ for given periods of time.

Aqueous micellization of ImSSP by nanoprecipitation method

PBS solution (10 mL, pH = 7.4) was dropwise added to an organic solution of ImSSP (10 mg) dissolved in THF (2 mL) at a rate of 1 mL min^{-1} using a syringe pump under stirring. The resultant mixture was kept stirring at room temperature for 40 h to remove THF, yielding an aqueous solution of nanoassemblies at 1 mg mL^{-1} .

Colloidal stability and acid/GSH responsive disassembly of aqueous nanoassemblies

Aliquots of empty nanoassemblies (1 mL, 1 mg mL^{-1}) were incubated with acetate buffer solution (3 mL, 0.2 M, pH = 5) or GSH (10 mM) under stirring at room temperature. In parallel, empty nanoassemblies were kept on bench at room temperature. DLS analysis was followed for given periods of time.

Preparation of Dox-loaded NPs by dialysis

PBS solution (10 mL, pH = 7.4) was added dropwise to an organic solution consisting of Dox (2 mg), Et_3N (5 μL) and ImSSP (20 mg) dissolved in DMF (2 mL) at a rate of 1 mL min^{-1} using a syringe pump under stirring for 1 h. The resulting mixture was transferred to a dialysis tubing (MWCO = 11–13 kDa) for dialysis against PBS solution (1 L, pH = 7.4) at room temperature for 24 h to remove free (not encapsulated) Dox molecules and residual DMF. The resulting dispersion was passed through a 0.45 μm PES filter, yielding an aqueous Dox-loaded nanoparticles (Dox-NPs) at 1.6 mg mL^{-1} .

To determine the loading level of Dox in micelles, aliquots of aqueous Dox-NPs (1 mL) were mixed with DMF (5 mL). The resultant mixture was subjected to filtration using a 0.45 μm PTFE filter to remove insoluble salts and analyzed by UV/vis spectroscopy.

Acid/GSH-responsive release of Dox from Dox-NPs

Aliquots of Dox-NPs (1.7 mg mL^{-1} , 2 mL) were mixed with buffer solutions (2 mL) at different pHs = 5, 6.8 and 7.4 as well as 10 mM GSH. The mixtures were transferred to a dialysis tubing (MWCO = 11–13 kDa) and immersed into corresponding buffer solutions (40 mL) at 25 $^\circ\text{C}$ under stirring. Aliquots of outer solutions (3 mL) were taken out at given interval to record fluorescence spectra excited at $\lambda_{\text{ex}} = 480 \text{ nm}$. Equivalent amounts of fresh buffer solutions were added back to keep sink condition.

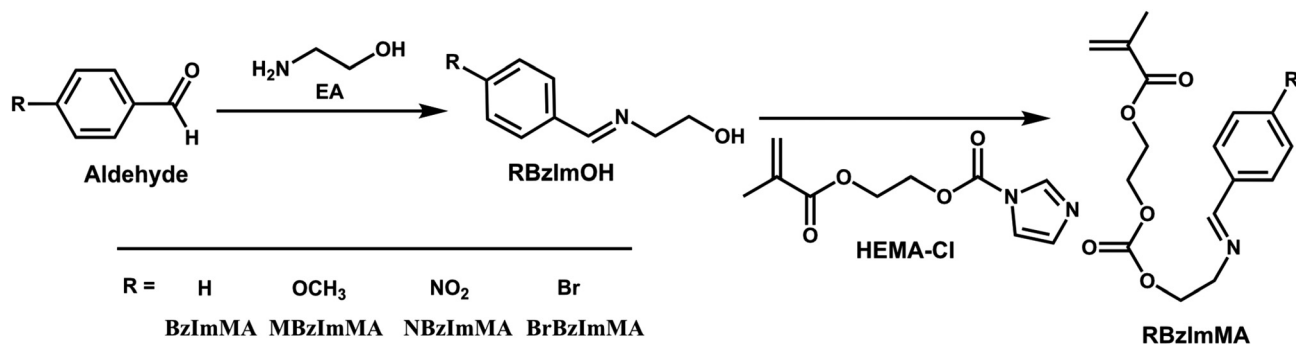
Cellular uptake with HeLa multi-cellular tumor spheroids

HeLa cells were maintained in DMEM (Wisent) supplemented with 10% cosmic calf serum (Hyclone) with 100 U penicillin (Wisent), and 0.1 mg mL^{-1} streptomycin (Wisent). Cells were



maintained in humidified incubators at 37 °C with 5% CO₂. Cells were split using 0.5% trypsin (Wisent) as per standard protocols. Multi-cellular tumor spheroids were generated from

HeLa cells in 96-well plates (BioLite, Thermo Scientific). Wells were coated with 1.5% agarose (Biotechnology Grade, BioShop), then seeded with 500–1000 cells in 150 µl of media,



Scheme 2 Our approach to synthesize a series of methacrylates bearing BzIm group with different *para* substituents (RBzImMAs).

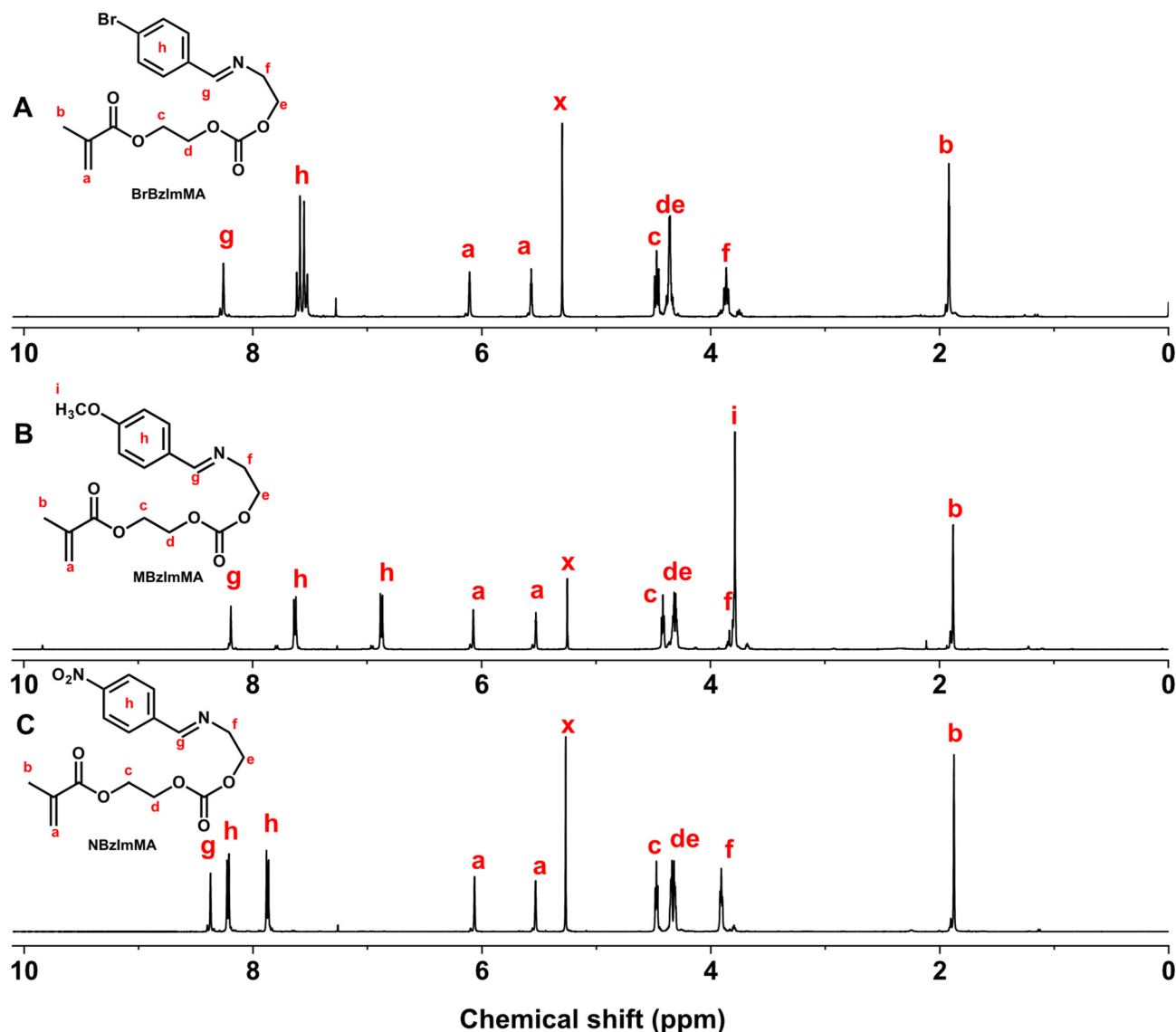


Fig. 1 ¹H-NMR spectra of RBzImMAs, including BrBzImMA (R = Br, A), MBzImMA (R = Me, B), and NBzImMA (R = NO₂, C) in CDCl₃.



which were left to aggregate with gravity at 37 °C and 5% CO₂. Spheroids were grown for 6 to 10 days and were monitored daily using an Invertoskop 40 C light microscope. Once spheroid formation was confirmed, they were transferred into 24-well plates coated with agarose in 2 mL of media. Spheroids were incubated with Dox or nanoassemblies with encapsulated Dox for 6 hours to compare Dox penetration. Spheroids were imaged using a Nikon TiE microscope epifluorescence microscope using the 4× objective, an LED Heliophor and filters for excitation at 488 nm and emission at 561 nm, and a Photometrics Evolve EMCCD camera with NIS Elements acquisition software. The images were exported as TIFFs and analyzed using Image J (NIH). The average intensity of Dox fluorescence in a defined surface area was measured for each spheroid and normalized to their initial values at time = 0.

Results and discussion

Synthesis of substituted BzIm methacrylates

Our synthesis of dual location dual acid/GSH-degradable ABP began with BzIm-bearing methacrylates with varying substituents. Scheme 2 illustrates our approach with two steps to synthesize a series of BzIm-bearing methacrylates with different R substituents on the *para* position of phenyl group (**RBzImMAs**). The R groups were chosen based on their electronic properties: electron-donating groups (methoxyl group) and electron-withdrawing groups (bromo and nitro groups). Note R = H (**HBzImMA**), *p*-methoxy (**MBzImMA**), bromo

(**BrBzImMA**), and nitro group (**NBzImMA**). The synthesis and NMR analysis of **HBzImMA** are described in our previous publication.²²

The first step was the facile coupling reaction of EA with various substituted benzaldehydes to synthesize their **RBzImOH** precursors bearing a BzIm linkage. The mole ratio of EA to aldehydes was kept at 1/1. Sodium sulfate was used to remove water, a biproduct formed during reactions, to promote forward reactions to form BzIm bonds. ¹H-NMR analysis confirms the successful synthesis of **MBzImOH** (Fig. S1†), **BrBzImOH** (Fig. S2†), and **NBzImOH** (Fig. S3†). Next step was the conjugation of the formed **RBzImOH** precursors with HEMA-Cl at a 1/1 mole ratio in the presence of DBU as a base catalyst, yielding **RBzImMAs**. ¹H-NMR spectra of the purified **RBzImMAs** (Fig. 1) show their characteristic peaks, including the peak at ~8.2 ppm (g) corresponding to imine proton as well as two peaks at 5.5–6.1 ppm (a) presenting methacrylate protons. The ¹H-NMR analysis, together with ¹³C-NMR results (Fig. S4–S6†), confirms the successful synthesis of BzIm methacrylates.

Impact of substituents on acid-catalyzed hydrolysis of BzIm bond

Given the synthesis of **RBzImMAs**, the effect of the substituents (R = H, MeO, Br, and NO₂) on acid-catalyzed hydrolysis of BzIm bonds was examined by ¹H-NMR analysis to estimate their acid-catalyzed hydrolysis kinetics in their corresponding polymers. Aliquots were dissolved in a mixture of CD₃CN/PB

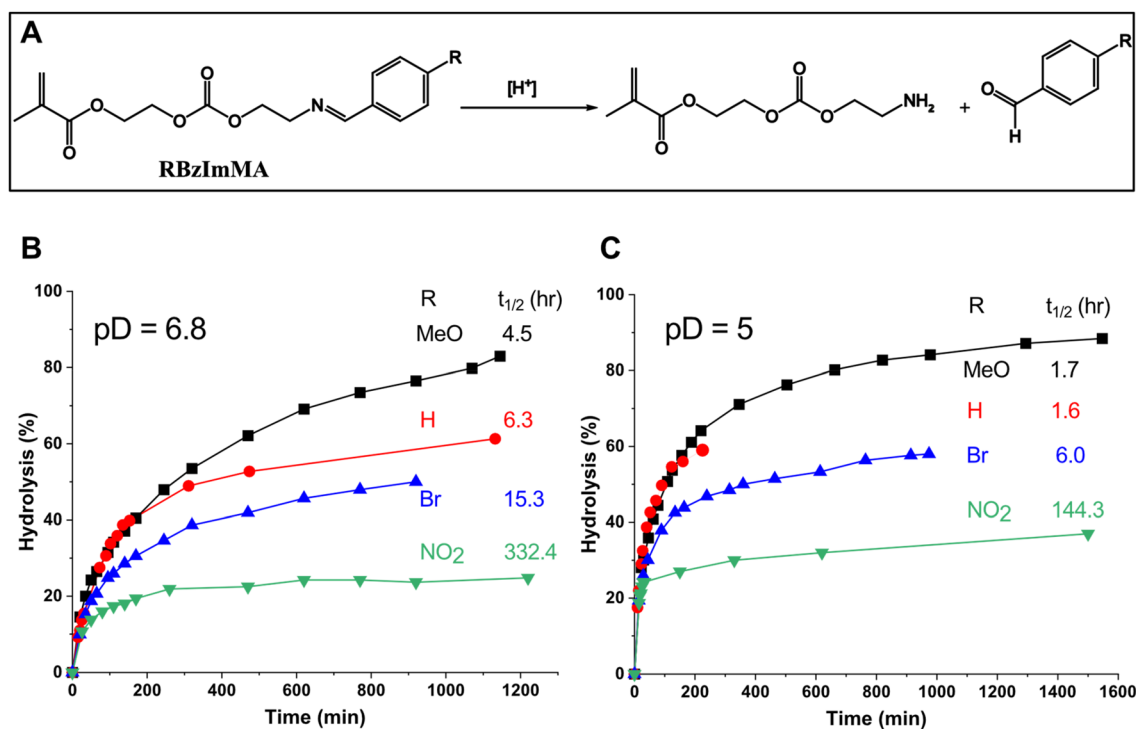


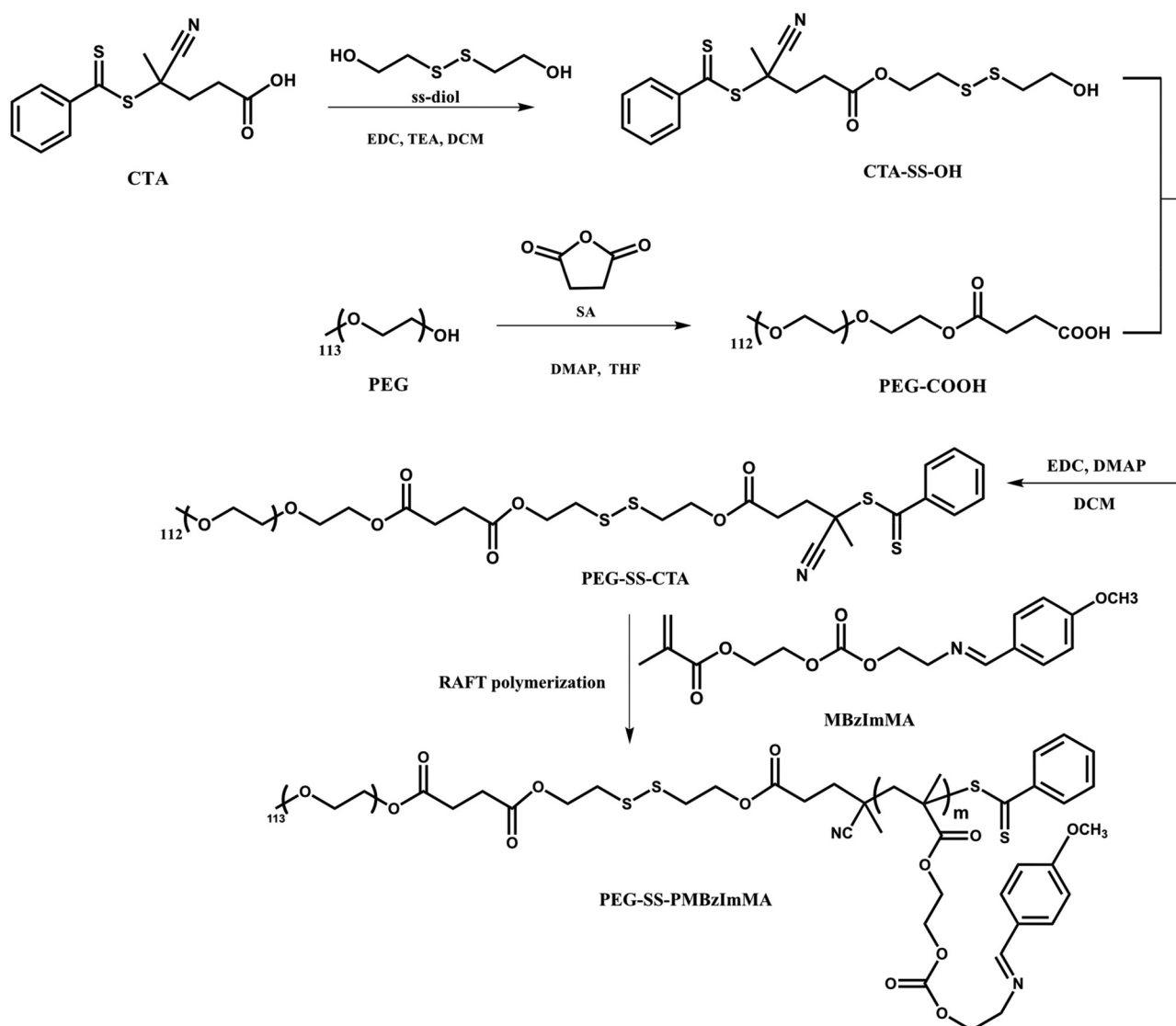
Fig. 2 Schematic illustration of acid-catalyzed hydrolysis of **RBzImMAs** upon the cleavage of BzIm bonds (A) and evolution of their percentage hydrolysis incubated at pD = 6.8 (B) and 5.0 (C) with their $t_{1/2}$ values.



D₂O (3/1, v/v) adjusted to pD 6.8 and 5.0, mimicking tumoral and *endo*/lysosomal pH, respectively. Their ¹H-NMR spectra were recorded over incubation times at given pDs (Fig. S7–S9† for **MBzImMA**, **BrBzImMA**, and **NBzImMA**, and Fig. 2 in our previous publication²² for **HBzImMA**). As illustrated in Fig. 2A, acid-catalyzed hydrolysis of **RBzImMAs** results in the cleavage of BzIm bonds, generating amine-bearing methacrylates and R-aldehydes. ¹H-NMR spectra show that the imine proton peak at ~8.2 ppm decreased, while a new peak at ~10 ppm equivalent to aldehyde proton appeared and increased over incubation time. Their integral ratios as aldehyde/(aldehyde + imine) were used to determine the percentage hydrolysis of BzIm bonds. This quantity, along with the time when a half concentration of BzIm linkages is cleaved ($t_{1/2}$), were used for quantitative evaluation and comparison of acid-catalyzed hydrolysis rate of **RBzImMAs** with different substituents. The results are shown in Fig. 2B at pD = 6.8 and Fig. 2C at pD =

5.0. At both pDs, the percentage hydrolysis of BzIm bonds increased over incubation time for all methacrylates.

At pD = 6.8, the hydrolysis rate turned to increase in the order of R = MeO > H > Br > NO₂ group (Fig. 2B). Compared with unsubstituted **HBzImMA** ($t_{1/2}$ = 6.3 h), the electron-donating MeO group enhanced the hydrolysis rate (**MBzImMA**, $t_{1/2}$ = 4.5 h), while the electron-withdrawing Br (**BrBzImMA**, $t_{1/2}$ = 15.3 h) and NO₂ (**NBzImMA**, $t_{1/2}$ = 332.4 h) groups slowed down the hydrolysis rate. The acid-catalyzed hydrolysis of imine bonds can be initiated with the protonation of the imine nitrogen, generating an electrophilic carbocation intermediate that facilitates nucleophilic attack by water, leading to the formation of an unstable tetrahedral intermediate and ultimately resulting in imine bond dissociation.⁴⁵ The plausible reason of this observed substituents effect could be due to their ability to stabilize generated carbocation intermediates. For example, electron-donating MeO group could stabilize the



Scheme 3 Synthesis of PEG-SS-CTA macromediator and PEG-SS-PMBzImMA (ImSSP) block copolymer by RAFT polymerization.



formed carbocation through resonance in phenyl ring, thus increasing hydrolysis rate; while electron-withdrawing Br and NO₂ groups could destabilize it, impeding acid-catalyzed hydrolysis.

When being incubated at pD = 5, a similar trend of substituent effect on imine hydrolysis rate was observed, following the order of R = MeO \approx H > Br > NO₂ group (Fig. 2C). Interestingly, no significant difference in $t_{1/2}$ is observed for **MBzImMA** and **HBzImMA**, likely due to their rapid hydrolysis at lower pD = 5. As expected, the hydrolysis rates of BzIm bonds were greater at pD = 5.0, compared with pD = 6.8 for all methacrylates. For example, the $t_{1/2}$ was 1.8 h (4.5 h at pD = 6.8) for **MBzImMA** and 6 h (15.9 h at pD = 6.8) for **NBzImMA**.

These results suggest that the acid-catalyzed hydrolysis of BzIm bonds is significantly influenced by the electronic properties of their *para* substituents.

Given its rapid hydrolysis, **MBzImMA** was chosen to explore the synthesis of amphiphilic block copolymer for further experiments, *e.g.* aqueous micellization, acid/GSH-responsive disassembly and Dox release as well cellular uptake.

Synthesis of PEG-SS-PMBzImMA (ImSSP)

As depicted in Scheme 3, RAFT polymerization of **MBzImMA** was explored to synthesize ABP bearing a disulfide junction for hydrophilic PEG block and hydrophobic polymethacrylate block bearing pendant BzIm groups. The approach began with

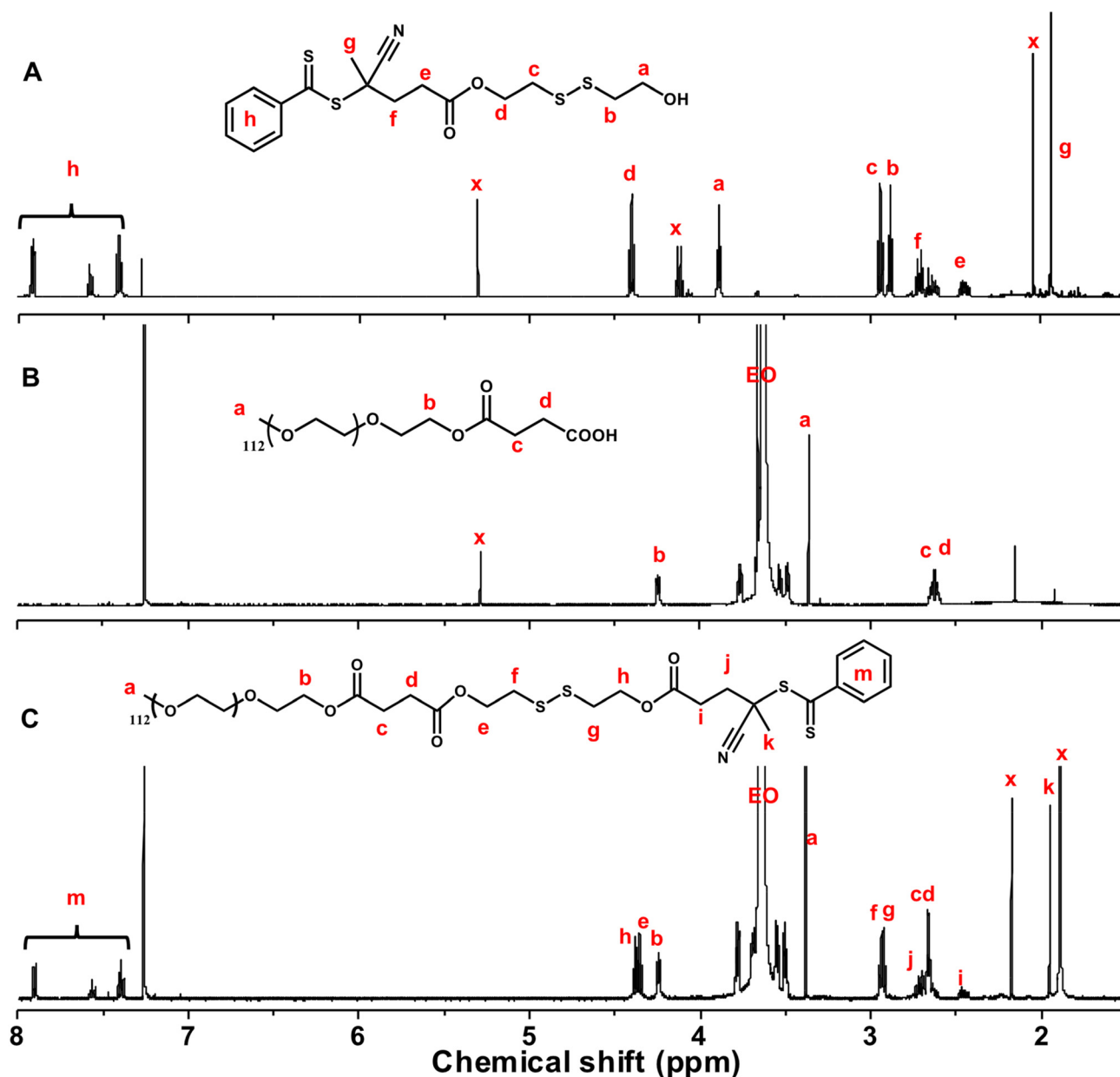


Fig. 3 ¹H-NMR spectra of CTA-SS-OH (A), PEG-COOH (B), and PEG-SS-CTA (C) in CDCl₃. x denotes residual solvents or impurities.



three-step synthesis of a disulfide-labeled PEG-based chain transfer agent (PEG-SS-CTA). The first step involved the reaction of CTA with excess SS-Diol to synthesize a disulfide-labelled dithiocarbonyl CTA with a terminal hydroxyl group (CTA-SS-OH). The desired mono-substituted product was purified and collected as the third band out of total five during silica gel column chromatography with an eluent solution

(ethyl acetate/hexane = 1/1, v/v) at a yield of 30%. The second step involved the reaction of PEG (MW = 5000) with excess succinic anhydride, yielding a PEG functionalized with carboxylic acid (PEG-COOH). The last step was an EDC-mediated coupling reaction between CTA-SS-OH and PEG-COOH to synthesize PEG-SS-CTA. Their chemical structures were confirmed by $^1\text{H-NMR}$ analysis (Fig. 3). Specifically, $^1\text{H-NMR}$ spectrum of

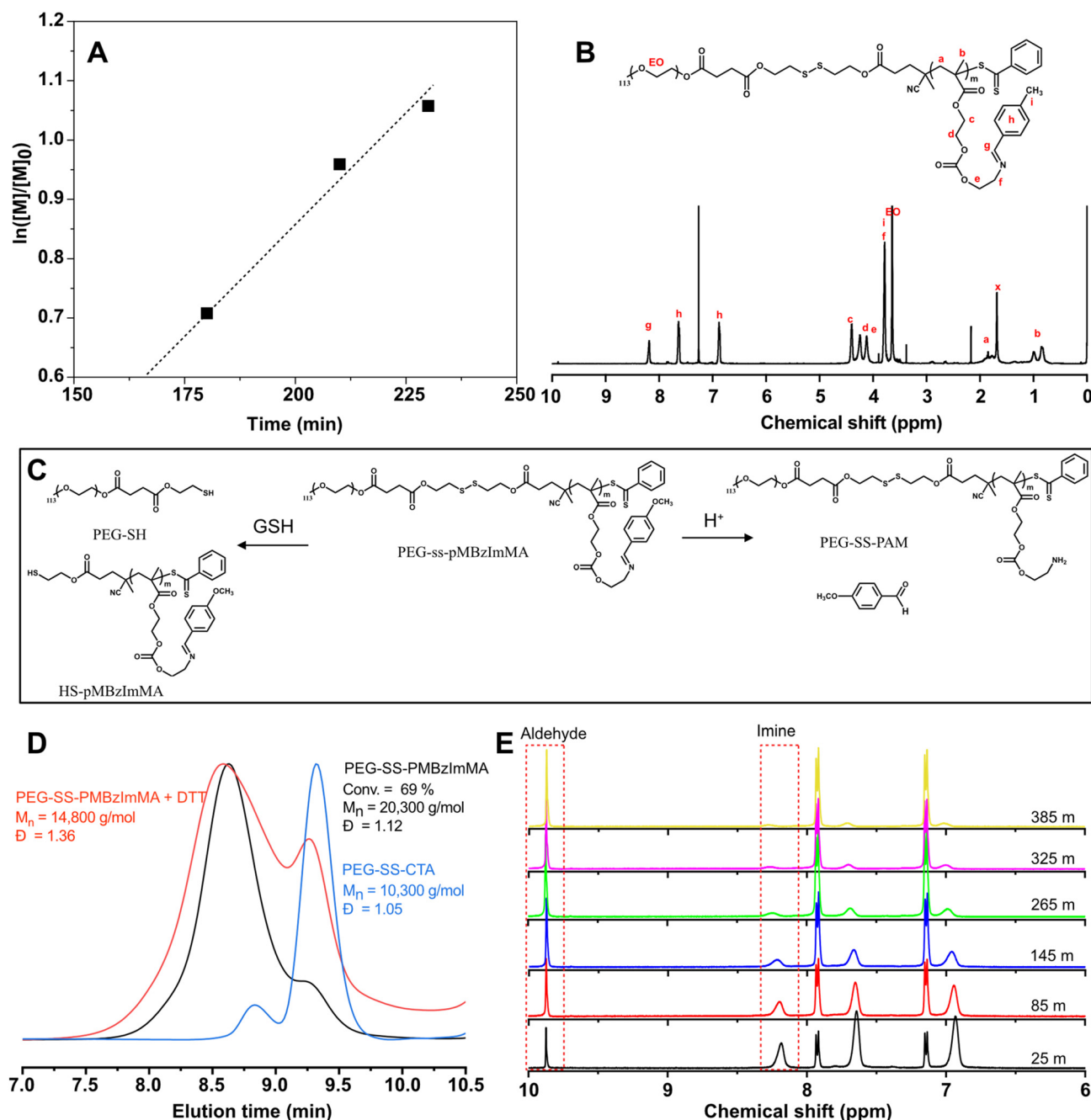


Fig. 4 Synthesis and acid/GSH-responsive cleavage of ImSSP. First-order kinetic plot for RAFT polymerization of MBzImMA in presence of PEG-SS-CTA (A); $^1\text{H-NMR}$ spectrum of purified PEG-SS-PMBzImMA at 65% monomer conversion (B); conditions for RAFT polymerization: $[\text{MBzImMA}]_0/[\text{PEG-SS-CTA}]_0/[\text{AMBn}]_0 = 75/1/0.3$ in anisole at 70°C and $[\text{BzImMA}]_0/\text{anisole} = 0.6$ wt/wt; degradation of ImSSP upon the cleavage of junction disulfide linkage and pendant BzIm bond in the presence of GSH or in acidic pHs (C). Overlaid GPC diagrams before and after GSH-induced cleavage, compared with PEG-ss-CTA (D); and overlaid $^1\text{H-NMR}$ spectra at $\text{pD} = 5.0$ over incubation time (E).



PEG-SS-CTA (Fig. 3C) shows the characteristic peaks at 7.3–8.0 ppm (m) corresponding to phenyl protons in CTA moiety and the peak at 3.2 ppm (a) corresponding to methoxy protons in PEG moiety. The integrals of all peaks were consistent with the numbers of their protons. From the integral ratio of peaks (a and h), the conjugation efficiency was estimated to be >95%.

With the successful synthesis of PEG-SS-CTA and optimal acid-cleavable monomer **MBzImMA**, RAFT polymerization for **MBzImMA** was conducted in the presence of PEG-SS-CTA as a macromediator, initiated with AMBN (an azo-initiator) at 70 °C. This yielded a shell-sheddable/core-degradable PEG-SS-PMBzImMA (ImSSP), consisting of hydrophilic PEG block conjugated *via* a disulfide bond to hydrophobic PMBzImMA block having pendant BzIm bonds. Kinetic analysis revealed that monomer conversion reached 65% within 4 h after an induction period of *ca.* 165 min (Fig. 4A). The crude polymer was purified by precipitation in diethyl ether. The ¹H-NMR spectrum of the purified PEG-SS-PMBzImMA in Fig. 4B shows the characteristic peaks at 8.3 ppm (g) for the imine bond, 7.4–7.7 ppm (h) for the phenyl group, 0.7–1.1 ppm (b) for the backbone methyl group of PMBzImMA block as well as 3.6 ppm (EO) for PEG block. The presence of imine peak in ¹H-NMR spectrum after purification confirms its stability during RAFT polymerization. Based on their integral ratio, the degree of polymerization (DP) was estimated to be 33, forming PEG-SS-PMBzImMA₃₃. Its GPC trace was shifted to higher molecular weight region after polymerization (Fig. 4D), indicating successful chain extension from PEG-SS-CTA with acid-cleavable PMBzImMA block. A small shoulder in low molecular weight region appeared to overlap with PEG-SS-CTA, suggesting that the block copolymer could contain a small portion of unreacted PEG-SS-CTA. The block copolymer had a low dispersity of $D = 1.12$ and a number average molecular weight of $M_n = 20.3 \text{ kg mol}^{-1}$, which was in good agreement with the theoretical $M_n = 20.1 \text{ kg mol}^{-1}$ calculated based on the targeted DP of PMImBzMA blocks and monomer conversion.

Acid/GSH-responsive degradation of ImSSP

The synthesized PEG-SS-PMBzImMA contains a reductively cleavable disulfide linkage at the block junction and acid-labile BzIm pendants in the hydrophobic block. Fig. 4C shows schematic illustration of its degradation in response to reduction and acidic pH. To investigate reductive degradation using GPC, the copolymer was incubated with DTT, a reducing agent (5 equivalent excess to disulfide bonds). As seen in Fig. 4D, the M_n decreased from 20.3 kg mol^{-1} to 14.8 kg mol^{-1} as a consequence of the cleavage of junction disulfide bond to generate PEG-SH and SH-PMBzImMA. The GPC diagram of the degraded products shows two peaks: a pronounced shoulder peak at lower MW region corresponding to PEG-SS-CTA and a peak in higher MW region corresponding to SH-PMBzImMA.

To investigate acid-responsive degradation using ¹H-NMR, the copolymer was incubated at pD = 5.0. Similar to its MBzImMA monomer counterpart, the peak at 8.3 ppm cor-

responding to BzIm proton in the copolymer decreased over incubation time, while the peak at 9.8 ppm corresponding to aldehyde proton in *p*-methoxybenzaldehyde as a degraded product increased (Fig. 4E). The $t_{1/2}$ was determined to be 1.9 h at pD = 5 for the copolymer, which is slightly higher than that (1.77 h) for **MBzImMA** monomer under the similar condition. Further, the degraded products were isolated by centrifugation for their structural analysis. ¹H-NMR spectrum in Fig. S10† confirms the generation of methoxybenzaldehyde and PEG-SS-PAM as degraded products upon the cleavage of pendant BzIm bonds.

Aqueous micellization and acid/GSH-responsive disassembly

The synthesized diblock copolymer consists of a hydrophilic PEG block and a hydrophobic pMBzImMA block with multiple imine pendants, connected *via* a disulfide linkage. Its amphiphilicity was utilized for fabricating polymer micelles at 1 mg mL^{-1} . The formed micellar aggregates are composed of imine-containing hydrophobic PMBzImMA cores, surrounded with sheddable hydrophilic PEG corona. They had a z-average diameter of $94 \pm 0.1 \text{ nm}$ with a monomodal distribution by DLS analysis (Fig. 5A).

Upon incubation with GSH, both z-average diameter and polydispersity (PDI) of the nanoassemblies increased rapidly, with the occurrence of large aggregates (diameter > 1 μm) within 24 h and PDI rising from 0.1 to 0.3. Such change is likely due to the destabilization and aggregation of formed hydrophobic cores after the loss of their hydrophilic coronas upon the cleavage of disulfide linkages at core/corona interfaces. In acidic pH = 5, their diameter significantly decreased to 3.5 nm. This is probably attributed to the cleavage of imine bonds in hydrophobic cores to generate primary amine pendants, which could be further protonated to form highly water-soluble PEG-SS-PAM. A similar decrease in their diameter to 2.2 nm was observed when the nanoassemblies were treated with both H⁺ and GSH simultaneously.

Loading and acid/GSH-responsive release of Dox

To explore the potential of ImSSP-based nanoassemblies as an intracellular delivery nanocarrier for chemotherapy, Dox, a

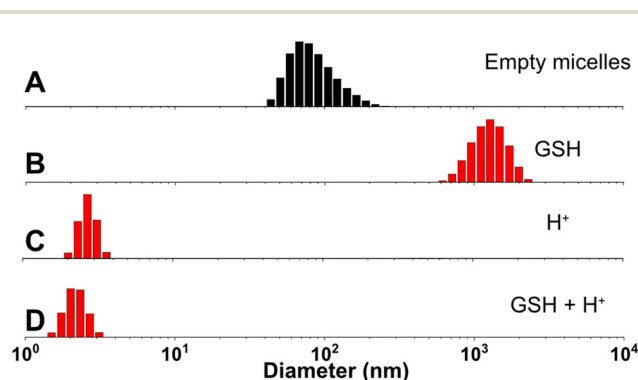


Fig. 5 DLS diagrams of nanoassemblies formed through self-assembly of PEG-SS-PMImBzMA before (A) and after being incubated with 10 mM GSH (B), in acidic pH = 5 (C), and both (D).



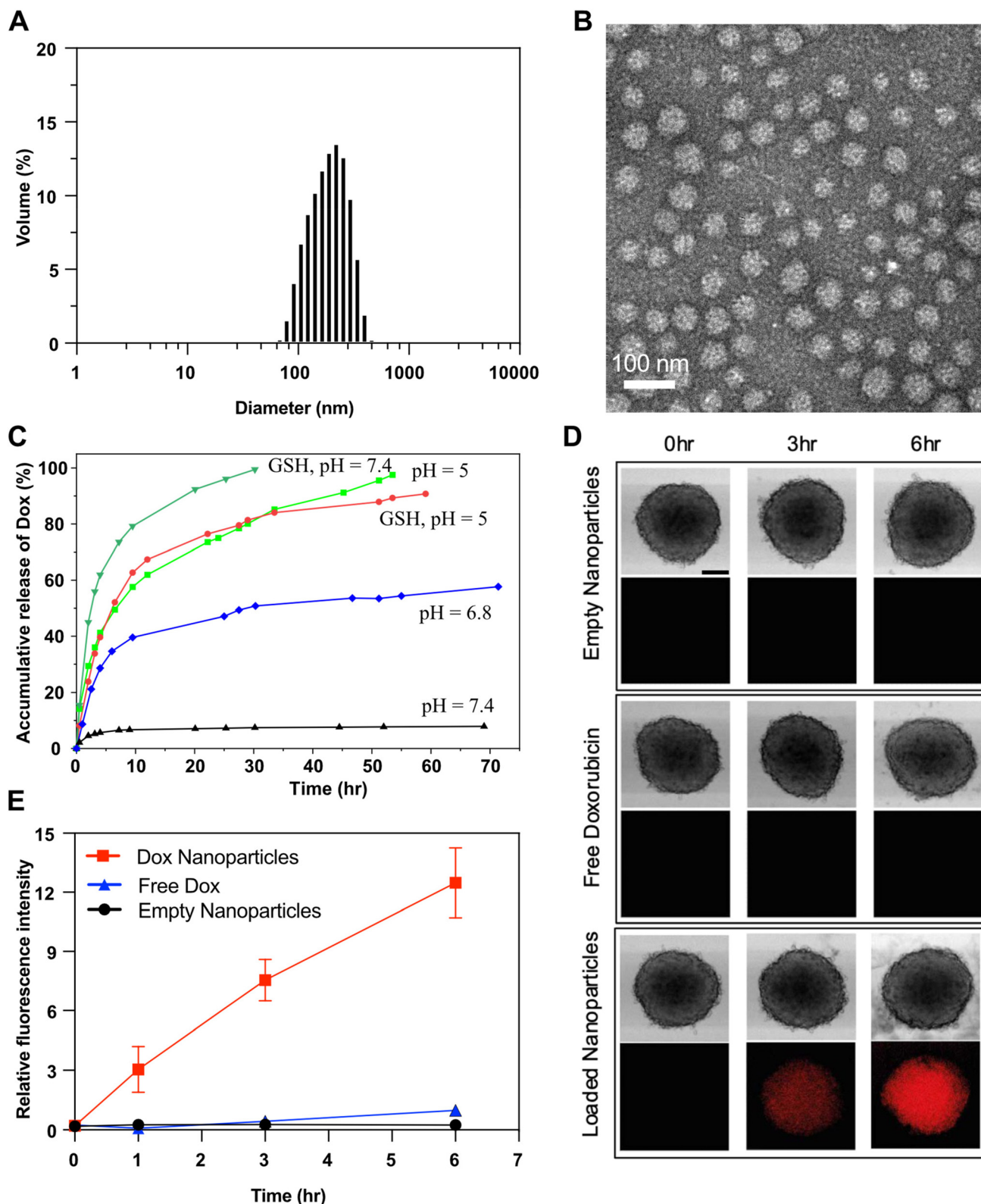


Fig. 6 DLS diagram (A) and TEM image (B, scale bar = 100 nm) of Dox-loaded NPs. %Dox release over incubation time at acidic pHs = 5.0 and 6.8 and GSH, compared with in absence of external stimuli at physiological pH = 7.4 (C). Fluorescence microscope images of HeLa multicellular tumor spheroids incubated for 6 hours with empty nanoparticles, free Dox and Dox-loaded nanoparticles. $n = 4$, scale bar = 200 μm . (D). Quantitative analysis of fluorescence intensity in the spheroids (E).



clinically approved cancer drug, was encapsulated in the nanoassemblies using a dialysis method. The resulting Dox-loaded nanoparticles (Dox-NPs) had a diameter of 160.6 ± 0.1 nm as measured by DLS analysis (Fig. 6A), which is larger than their empty counterparts (94 ± 0.1 nm). The increase in size could be due to the formation of larger hydrophobic cores upon incorporation of hydrophobic Dox. TEM analysis (Fig. 6B) revealed that Dox-NPs had a uniform spherical morphology with an average diameter of 48 ± 7 nm. This smaller size determined by TEM, compared to DLS, is attributed to the shrinkage of nanoparticles in the dried state, which is commonly observed for block copolymer-based nanoassemblies. Using the extinction coefficient of Dox ($12\,400\text{ M}^{-1}\text{ cm}^{-1}$) in DMF/water mixture (1/5, v/v), the loading efficiency of Dox was determined to be 11% with a loading content of $18.6\ \mu\text{g mL}^{-1}$ (Fig. S11†).

Self-assembled Dox-NPs could disintegrate upon the acid-catalyzed hydrolysis of pendant imine linkages in their hydrophobic cores, disrupting hydrophilic/hydrophobic balance. Alternatively, NPs could also disassemble in the presence of GSH due to the loss of the hydrophilic PEG corona upon the cleavage of junction disulfide bonds. As shown in Fig. 6C, Dox release was significantly enhanced in acidic pH or in the presence of GSH, compared to that in physiological pH at 7.4. For instance, only 8% of Dox was released in neutral pH after >80 h, while 50% and 80% of Dox were released at pH = 6.8 and pH = 5, respectively. These enhanced Dox release results are consistent with previously observed pH-dependent hydrolysis kinetics of BzIm bonds. Additionally, Dox release was also higher in the presence of GSH, which is attributed to the disintegration of Dox-NPs upon the cleavage of disulfide bonds at interfaces.

Biological evaluation of the release of Dox by HeLa multi-cellular tumor spheroids

Next, we evaluated the ability of the dual-responsive nanoparticles to penetrate and release Dox in tumors. Multi-cellular tumor spheroids grown from cultured HeLa cells *in vitro* were used to mimic solid tumors *in vivo*. HeLa cells were induced to form spheroids, which were then treated with Dox-NPs or with free Dox and empty NPs as controls. As shown in Fig. 6D, bright field (top) and fluorescence images (bottom) of HeLa spheroids after treatment revealed fluorescence signal with Dox-NPs after 3 h, but not the controls (free Dox and empty NPs). The fluorescence intensity in spheroids treated with Dox-NPs increased further after 6 h. Quantitative analysis of fluorescence intensity (Fig. 6E) indicated that spheroids treated with Dox-NPs had a pronounced increase in fluorescence, whereas there was only small increase for free Dox and no increase for the spheroids incubated with empty nanoparticles. These results suggest that the formed Dox-NPs can effectively penetrate and release Dox in the HeLa spheroids. Combined with their dual-responsive nature in the tumor microenvironment (acidic pH and elevated GSH), these results highlight the potential of Dox-NPs as a promising drug delivery system for cancer therapy *in vivo*.

Conclusion

In this study, we have developed and optimized BzIm-based dual acid/GSH-responsive degradable ABPs and their nanoassemblies for enhanced drug release. The ABPs were designed with acid-cleavable BzIm pendants in hydrophobic blocks and GSH-cleavable disulfides at PEG hydrophilic/hydrophobic block junctions. Various methacrylate monomers bearing BzIm groups (**RBzImMAs**) with different *para*-substituents (R = MeO, H, NO₂, and Br) were synthesized. Their acid-catalyzed hydrolysis rate by ¹H-NMR analysis was revealed in the increasing order of MeO > H > Br > NO₂, which could be attributed to the electronic properties of their *para* substituents. Thus, tunable hydrolysis kinetics of BzIm bonds could be useful for designing distinct pH-responsive delivery systems.

Given that **MBzImMA** exhibited an enhanced acid-catalyzed hydrolysis rate, a dual acid/GSH-responsive ImSSP (PEG-SS-PMBzImMA) was synthesized through direct RAFT polymerization in the presence of PEG-SS-CTA. ¹H-NMR and GPC results confirmed that disulfide linkage and BzIm pendants in synthesized ImSSP were cleaved in response to acid and GSH. The resulting ImSSP ABP self-assembled to form colloiddally stable, monomodal, and spherical nanoassemblies, as confirmed by both TEM and DLS analysis. They were capable of Dox encapsulation and exhibited enhanced release of Dox in response to acidic pH and GSH, mimicking the tumor acidic microenvironment and intracellular cytosol, respectively. This enhanced release could be attributed to the degradation of hydrophobic cores to hydrophilic polymers upon the cleavage of BzIm bonds as well as the detachment of the sheddable PEG corona upon the cleavage of SS bonds at core/corona interfaces. Moreover, biological evaluation using HeLa multi-cellular tumor spheroids confirmed that the penetration of Dox-loaded nanoassemblies and their release is greater compared to free Dox.

These results show that ImSSP-based nanocarriers are robust acidic pH and GSH-responsive drug release systems, leveraging the tunable hydrolysis kinetics of BzIm bonds by varying substituents. Further biological evaluation in cell and animal models of these dual-responsive nanoassemblies could pave the way their application as delivery system for cancer therapies.

Data availability

The data supporting this article have been included as part of the ESI.†

Conflicts of interest

There are no conflicts to declare.

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References

- C. Li, Z. Deng and E. R. Gillies, *Curr. Opin. Biomed. Eng.*, 2023, **25**, 100437.
- X. Hu, A. M. Jazani and J. K. Oh, *Polymer*, 2021, **230**, 124024.
- M. A. Beach, U. Nayanathara, Y. Gao, C. Zhang, Y. Xiong, Y. Wang and G. K. Such, *Chem. Rev.*, 2024, **124**, 5505–5616.
- N. Corrigan, K. Jung, G. Moad, C. J. Hawker, K. Matyjaszewski and C. Boyer, *Prog. Polym. Sci.*, 2020, **111**, 101311.
- X. Hu, G. Szczepaniak, A. Lewandowska-Andralojc, J. Jeong, B. Li, H. Murata, R. Yin, A. M. Jazani, S. R. Das and K. Matyjaszewski, *J. Am. Chem. Soc.*, 2023, **145**, 24315–24327.
- J. Jeong, X. Hu, R. Yin, M. Fantin, S. R. Das and K. Matyjaszewski, *J. Am. Chem. Soc.*, 2024, **146**, 13598–13606.
- X. Hu, R. Yin, J. Jeong and K. Matyjaszewski, *J. Am. Chem. Soc.*, 2024, **146**, 13417–13426.
- A. M. Jazani, H. Murata, M. Cvek, A. Lewandowska-Andralojc, R. Bernat, K. Kapil, X. Hu, F. De Luca Bossa, G. Szczepaniak and K. Matyjaszewski, *Chem. Sci.*, 2024, **15**, 9742–9755.
- Q. Zhang, N. R. Ko and J. K. Oh, *Chem. Commun.*, 2012, **48**, 7542–7552.
- A. Harada and K. Kataoka, *Prog. Polym. Sci.*, 2006, **31**, 949–982.
- A. S. Mikhail and C. Allen, *J. Controlled Release*, 2009, **138**, 214–223.
- X.-B. Xiong, A. Falamarzian, S. M. Garg and A. Lavasanifar, *J. Controlled Release*, 2011, **155**, 248–261.
- A. M. Jazani and J. K. Oh, *Polym. Chem.*, 2020, **11**, 2934–2954.
- K. K. Bawa, A. M. Jazani, Z. Ye and J. K. Oh, *Polymer*, 2020, **194**, 122391.
- J. K. Oh, *Polym. Chem.*, 2019, **10**, 1554–1568.
- I. F. Tannock and D. Rotin, *Cancer Res.*, 1989, **49**, 4373–4384.
- P. Watson, A. T. Jones and D. J. Stephens, *Adv. Drug Delivery Rev.*, 2005, **57**, 43–61.
- S. Bazban-Shotorbani, M. M. Hasani-Sadrabadi, A. Karkhaneh, V. Serpooshan, K. I. Jacob, A. Moshaverinia and M. Mahmoudi, *J. Controlled Release*, 2017, **253**, 46–63.
- S. Binauld and M. H. Stenzel, *Chem. Commun.*, 2013, **49**, 2082–2102.
- K. Neuvonen, F. Fülöp, H. Neuvonen, A. Koch, E. Kleinpeter and K. Pihlaja, *J. Org. Chem.*, 2003, **68**, 2151–2160.
- K. Bairagi, J. T. Liu, A. Thinphang-nga and J. K. Oh, *Macromolecules*, 2023, **56**, 4307–4317.
- X. Hu and J. K. Oh, *Macromol. Rapid Commun.*, 2020, **41**, 2000394.
- Y. Li, Q. N. Bui, L. T. M. Duy, H. Y. Yang and D. S. Lee, *Biomacromolecules*, 2018, **19**, 2062–2070.
- J. Xue and P. Liu, *Eur. Polym. J.*, 2024, **217**, 113313.
- J. Li, Y. Zhou, C. Li, D. Wang, Y. Gao, C. Zhang, L. Zhao, Y. Li, Y. Liu and X. Li, *Bioconjugate Chem.*, 2015, **26**, 110–119.
- N. Kongkatigumjorn and D. Crespy, *Polym. Chem.*, 2024, **15**, 4491–4518.
- N. Chan, B. Khorsand, S. Aleksanian and J. K. Oh, *Chem. Commun.*, 2013, **49**, 7534–7536.
- X. Fu, L. Hosta-Rigau, R. Chandrawati and J. Cui, *Chem*, 2018, **4**, 2084–2107.
- A. Russo, W. DeGraff, N. Friedman and J. B. Mitchell, *Cancer Res.*, 1986, **46**, 2845–2848.
- F. G. Ottaviano, D. E. Handy and J. Loscalzo, *Circ. J.*, 2008, **72**, 1–16.
- C. Hwang, A. J. Sinskey and H. F. Lodish, *Science*, 1992, **257**, 1496–1502.
- G. Wu, Y.-Z. Fang, S. Yang, J. R. Lupton and N. D. Turner, *J. Nutr.*, 2003, **134**, 489–492.
- Y. Wang, Q. Luo, W. Zhu, X. Li and Z. Shen, *Polym. Chem.*, 2016, **7**, 2665–2673.
- J. Zhang, H. Tang, Y. Shen, Q. Yu and Z. Gan, *Macromol. Rapid Commun.*, 2018, **39**, 1800139.
- Y. Zhang, J. Ding, M. Li, X. Chen, C. Xiao, X. Zhuang, Y. Huang and X. Chen, *ACS Appl. Mater. Interfaces*, 2016, **8**, 10673–10682.
- Z. Su, Y. Xu, Y. Wang, W. Shi, S. Han and X. Shuai, *Biomater. Sci.*, 2019, **7**, 3821–3831.
- L. Sun, H. Wei, X. Zhang, C. Meng, G. Kang, W. Ma, L. Ma, B. Wang and C. Yu, *Polym. Chem.*, 2020, **11**, 4469–4476.
- X. Hu, H. Li, S. Luo, T. Liu, Y. Jiang and S. Liu, *Polym. Chem.*, 2013, **4**, 695–706.
- A. M. Jazani, N. Arezi, C. Shetty, S. H. Hong, H. Li, X. Wang and J. K. Oh, *Polym. Chem.*, 2019, **10**, 2840–2853.
- Y. Ding, C. Du, J. Qian, L. Zhou, Y. Su, R. Zhang and C.-M. Dong, *Polym. Chem.*, 2018, **9**, 3488–3498.
- Z. Su, Y. Xu, Y. Wang, W. Shi, S. Han and X. Shuai, *Biomater. Sci.*, 2019, **7**, 3821–3831.
- D. Biswas, S. Y. An, Y. Li, X. Wang and J. K. Oh, *Mol. Pharmaceutics*, 2017, **14**, 2518–2528.
- S. Y. An, S. H. Hong, C. Tang and J. K. Oh, *Polym. Chem.*, 2016, **7**, 4751–4760.
- G. Zhang and J. K. Oh, *ACS Appl. Polym. Mater.*, 2020, **2**, 2319–2326.
- E. H. Cordes and W. P. Jencks, *J. Am. Chem. Soc.*, 1963, **85**, 2843–2848.

