Judging Enzyme-Responsive Micelles by Their Covers: Direct Comparison of Dendritic Amphiphiles with Different Hydrophilic Blocks

Gadi Slor,* Alis R. Olea,* Silvia Pujals, Ali Tigrine, Victor R. De La Rosa, Richard Hoogenboom, Lorenzo Albertazzi,* and Roey J. Amir*

ABSTRACT: Enzymatically degradable polymeric micelles have great potential as drug delivery systems, allowing the selective release of their active cargo at the site of disease. Furthermore, enzymatic degradation of the polymeric nanocarriers facilitates clearance of the delivery system after it has completed its task. While extensive research is dedicated toward the design and study of the enzymatically degradable hydrophobic block, there is limited understanding on how the hydrophilic shell of the micelle can affect the properties of such enzymatically degradable micelles. In this work, we report a systematic head-to-head comparison of well-defined polymeric micelles with different polymeric shells and two types of enzymatically degradable hydrophobic cores. To carry out this direct comparison, we developed a highly modular approach for preparing clickable, spectrally active enzyme-responsive dendrons with adjustable degree of hydrophobicity. The dendrons were linked with three different widely used hydrophilic polymers—poly(ethylene glycol), poly(2-ethyl-2-oxazoline), and poly(acrylic acid) using the CuAAC click reaction. The high modularity and molecular precision of the synthetic methodology enabled us to easily prepare well-defined amphiphiles that differ either in their hydrophilic block composition or in their hydrophobic dendron. The micelles of the different amphiphiles were thoroughly characterized and their sizes, critical micelle concentrations, drug loading, stability, and cell internalization were compared. We found that the micelle diameter was almost solely dependent on the hydrophobicity of the dendritic hydrophobic block, whereas the enzymatic degradation rate was strongly dependent on the composition of both blocks. Drug encapsulation capacity was very sensitive to the type of the hydrophilic block, indicating that, in addition to the hydrophobic core, the micellar shell also has a significant role in drug encapsulation. Incubation of the spectrally active micelles in the presence of cells showed that the hydrophilic shell significantly affects the micellar stability, localization, cell internalization kinetics, and the cargo release mechanism. Overall, the high molecular precision and the ability of these amphiphiles to report their disassembly, even in complex biological media, allowed us to directly compare the different types of micelles, providing striking insights into how the composition of the micelle shells and cores can affect their properties and potential to serve as nanocarriers.

INTRODUCTION

Polymeric nano-assemblies, amongst them polymeric micelles, have shown great potential as drug delivery systems (DDS) as well as in many other biomedical applications.1-3 This is due to the ability to dramatically increase the very low water solubility of lipophilic drug molecules by encapsulating them inside the hydrophobic cavities of the assemblies, simultaneously shielding them from the hostile biological environment. In addition, in many cases, these assemblies have sizes...
that allow passive accumulation in cancerous or inflamed tissues due to the enhanced permeability and retention (EPR) effect.\textsuperscript{4,5} Despite all advantages proven in numerous scientific reports, there are still challenges to overcome to increase the translation of polymeric DDS from academic research into the clinic.\textsuperscript{6} Hence, it is essential to further conduct fundamental research in this important area in order to gain deeper understanding of the parameters that govern the stability and functionality of such carriers and open the way for their broader application in biomedicine.

It is clear that DDS should be, on one hand, extremely stable to withstand the high dilution and interactions with blood components in order to allow their circulation in the body while maintaining their cargo of active drug molecules.\textsuperscript{7} On the other hand, the carriers should be able to release the drugs when the DDS has reached the target site.\textsuperscript{8−11} To address this need, over the last three decades, there has been a great interest in utilizing stimuli-responsive polymeric micelles as DDS to allow selective release of their therapeutic cargo.\textsuperscript{12,13} There are many reported examples of polymeric micelles that disassemble due to changes in pH,\textsuperscript{14−17} temperature,\textsuperscript{18−22} or redox potential,\textsuperscript{23−26} while there are significantly fewer examples of polymeric nanocarriers that can disassemble due to the presence of a designated enzyme.\textsuperscript{27−30} Enzymes are very appealing for triggering the disassembly of drug containing micelles since they are already present in the body, known for their high substrate specificity and in many cases specific enzymes are overexpressed in diseased tissues.\textsuperscript{31−33} Polymeric micelles are typically formed by the self-assembly of amphiphilic diblock copolymers so that the hydrophobic block forms the core and the hydrophilic block forms the micellar corona. It is clear that in the biological environment, most of the interactions between the micelle and its surroundings occur through the micelle’s corona.\textsuperscript{34,35}
interesting to note that although most reported DDS are based on poly(ethylene glycol) (PEG), the use of additional types of promising hydrophilic polymers such as poly(2-oxazoline) and polyacrylates has also been reported, inspired by the increasing human population that carries anti-PEG antibodies leading to an immune-response upon treatment with PEG-based therapeutics. To allow the rational design of DDS, it is critical to compare and study the behavior of different corona forming polymers in order to rationally select the most suited hydrophilic block. For a direct comparison between different hydrophilic polymers as micellar shells, it is essential that the hydrophobic core-forming block will be identical. Using dendrons as the hydrophobic core forming block provides many advantages compared to linear hydrophobic polymer blocks due to their well-defined structure and monodispersity and hence can be ideal for the purpose of the abovementioned comparison.

Here, we report the synthesis of clickable, fluorescently labeled enzyme-responsive dendrons, which can be reacted with a variety of azide-functionalized hydrophilic polymers. In this work, we studied and compared three types of commonly used hydrophilic polymers: PEG, poly(2-ethyl-2-oxazoline) (PEtOx), and poly(acrylic acid) (PAA). PEG and PETOx, which are noncharged polymers, were selected as they are considered "stealth" polymers that minimize interactions with native proteins and other biomolecules and hence elongate circulation time of their conjugates in vivo. PAA is polyionic at physiological pH and is much more hydrophilic than the other two polymers. Because of the comparative nature of this work, we decided to use polymers with similar molecular weights of approximately 5 kDa. Each type of polymer was clicked with two types of dendrons that differ in their hydrophobicity containing either four hexyl or four nonyl ester end-groups (Scheme 1). This allowed both head-to-head comparison between the three hydrophilic polymers and also to understand how altering the hydrophilic/hydrophobic ratio affects the micelles that are formed from each polymer.

**EXPERIMENTAL SECTION**

**Instrumentation.** HPLC: All measurements were recorded on a Waters Alliance e2695 separation module equipped with a Waters 2998 photodiode array detector. All solvents were purchased from Bio-Lab Chemicals and were used as received. All solvents are of high-performance liquid chromatography (HPLC) grade. 

**NMR:** Spectra recorded were on Bruker AVANCE I and AVANCE III 400 MHz spectrometers, as indicated. Chemical shifts are reported in parts per million and referenced to the solvent. SEC: all measurements were recorded on Visotec GPCmax by Malvern using the refractive index detector, and PEG standards (purchased from Sigma-Aldrich) were used for calibration. Absorbance and fluorescence spectra: measurements were recorded on a Tecxan Infinite M200Pro device or Agilent Technologies Cary Eclipse Fluorescence Spectrometer. MALDI-TOF MS: analysis was conducted on a Bruker AutoFlex MALDI-TOF MS (Germany). An α-cyano-4-hydroxycinnamic acid matrix was used. Mass spectra were acquired on a Bruker AutoSpec HRMS (EI) Micromass (UK) or Sympat High Definition MS (ESI), Waters Inc. (USA). Dynamic light scattering (DLS): all measurements were recorded on a Corduran technology VASCOY—particle size analyzer. Confocal microscopy: imaging was performed on a Zeiss LSM 800 confocal microscope, using 63× plan-apochromat oil immersion objective. Flow cytometry: analysis was performed using a spectral analyzer (Sony SA3800) flow cytometer, 96/384w.

**Materials.** Poly(ethylene glycol) methyl ether (5 kDa), allyl bromide (99%), 3-mercaptopropionic acid (98%), 2,2-dimethoxy-2-phenylacetophenone (99%), 4-(dimethylamino)pyridine (99%), ethyl α-bromoisobutyrate (98%), Fmoc-Lys(Boc)-OH (98%), N-hydroxysuccinimide (NHS, 99%), copper(I) bromide (CuBr, 98%), N,N′,N,N′-pentamethyldihexilenediaminetriazine (PMDETA, 99%), 3-bromopropionic acid (97%), bovine serum albumin (BSA), Triton X-100, porcine liver esterase (PLE), and Sephadex LH20 were purchased from Sigma-Aldrich. 1-Hexanol (98%) was purchased from Acros Organics. Propargyl bromide (80% in toluene), chlorotriphenylmethane (Trt–Cl, 98%), 4-nitrophenol (99%), triethylsilane (98%), N,N′-dicyclohexylcarbodiimide (99%), propargyl amine (98%), tert-butyl acrylate (tBA, 99%), and anhydrous K2CO3 (99%) were purchased from Alfa Aesar. 3,5-Dihydroxy benzoic acid was purchased from Apollo scientific. Cy5-NHS was purchased from Lumiprobe. Cystamine hydrochloride (98%), potassium hydroxide, N,N-disopropylpyrrolidylamine, 1-nanonol, and sodium azide (NaN3) were purchased from Merck. Silica gel (60 Å, 0.040–0.063 mm), sodium hydroxide, anhydrous Na2SO4 (granular, 10–60 mesh), piperidine (peptide synthesis), N,N-dimethylformamide (DMF, peptide synthesis), trifluoroacetic acid (HPLC grade), and all solvents were purchased from Bio-Lab and were used as received. Deuterated solvents for NMR were purchased from Cambridge Isotope Laboratories (CIL), Inc. PrestoBlue and Dulbecco’s modified Eagle medium (DMEM) with high glucose and pyruvate, fetal bovine serum, trypsin with 0.25% EDTA, and penicillin–streptomycin were purchased from Thermo Fisher. The dendrons and azide-functionalized hydrophilic polymers were synthesized, as detailed in the Supporting Information.

**General Procedure for CuAAC Click Reaction between Polymer-N3 and Dendron.** CuBr (2 equiv with respect to polymer−N3) was loaded in a 4 mL glass vial, which was sealed with a rubber septum. The vial was deoxygenated with three vacuum-nitrogen cycles and backfilled with nitrogen. In a separate 4 mL vial polymer−N3 (1 equiv), dendron (1.3 equiv) and PMDETA (2 equiv) were dissolved in DMF (100–200 mg polymer/mL) and purged with nitrogen for 2 min. This mixture was added into the CuBr containing vial using the nitrogen flushed syringe and needle. The vial was thoroughly vortexed until clear green solution was obtained (approximately 30 s). The reaction was stirred at room temperature for 1 h, filtered through a syringe filter [0.44 μm, hydrophilic poly(tetrafluoroethylene)], purified using LH20 (Sephadex) size exclusion column, and eluted with MeOH. Fractions that contained the product (identified by bright yellow color) were pooled, MeOH was evaporated to dryness, and the product was dried on high vacuum. All polymers were obtained as bright yellow solids.

**CMC. Preparation of Diluent.** Nile Red stock solution (0.88 mg/mL in ethanol) was diluted into a phosphate buffer (PB) (100 mM, pH 7.4) to afford a final concentration of 1.25 μM.

**Preparation and Measurement of Samples.** The polymer-dendron amphiphile was directly dissolved in the diluent to give a final concentration of 250 μM. Solution was vortexed vigorously until the amphiphile completely dissolved and further sonicated for 15 min in an ultrasonic bath. This solution was consecutively diluted by a factor of 1.5 with the diluent to afford a series of 24 samples. 150 μL of each sample was loaded onto a 96 well plate, and a fluorescence emission scan was performed for each well. In order to determine the amphiphile’s critical micelles’ concentration (CMC)—the maximum emission of Nile Red (at about 630 nm) was plotted versus the amphiphile’s concentration. This procedure was repeated three times for each amphiphile, and mean value is reported as the CMC and the standard deviation as the measurement error.

**Enzymatic Degradation Experiments.** A micellar solution of the tested amphiphile was prepared by directly adding PB (pH 7.4) to solid polymer to a final concentration of 160 μM. The vial was vortexed until full solubility was obtained and then placed in an ultrasonic bath for 15 min. PB stock solution or PB was added (30 μL into 1470 μL or 14 μL into 686 μL for HPLC or fluorescence experiments, respectively, to yield the final PB concentration of 1.4 μM), and degradation was followed at 37 °C either by monitoring the
area under the peak of the parent amphiphile by HPLC or the fluorescence emission at 540 nm. Each experiment was conducted thrice, and the reported values in each time point are the mean values and the standard deviation is the error.

**CPT and PTX Encapsulation Procedure.** The tested amphiphile was dissolved in MeOH (1 mg/mL). 1 mL of amphiphile solution was mixed with 1 mL of drug solution (0.5 mg/mL in DCM). Solvents were removed in vacuum forming a thin film layer, which was further dried on high vacuum for 1 h. Then, 1 mL of PB was added, and the vial content was stirred vigorously and placed in an ultrasonic bath for 30 min. The undissolved drug was filtered off using a syringe filter (0.45 μm hydrophilic Nylon), and the clear solution was analyzed by HPLC. The drug concentration was calculated by calibration curve at 360 nm for camptothecin (CPT) and 224 nm for paclitaxel (PTX).

**Blood Protein Interaction: Micelle Incubation with BSA-Cy5.** The interaction between micelles and Cy5-labeled BSA was measured using Förster resonance energy transfer (FRET). Micelles solution in PBS (pH 7.4) at 145.5 μM was mixed with BSA (10% v/v labeled with cyanine 5) at the 5.5 μM final concentration or with the same volume of PBS. Each sample was 60 μL in final volume inside a 96-well plate (flat-bottom, transparent, NUNC). The samples were excited at the 420 nm wavelength, and a fluorescence spectra was collected between 450 and 750 nm, with a 5 nm step. A reading was performed for each well every 15 min for 5 h, with the initial time point being approximately 15 min after BSA addition (or PBS, respectively).

**Cytotoxicity with PrestoBlue.** The cellular toxicity was assessed using the PrestoBlue assay (ThermoFisher). HeLa cells were seeded at a density of 5000 cells/well in a 96-well plate (Nunc, transparent, flat-bottom plate). After 24 h, micelles were added to the final concentration of 160 μM in full DMEM. The same volume of PBS was added in the negative control, while Triton X-100 0.01% v/v was added in the negative control, while Triton X-100 0.01% v/v was respectively).

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**Micelle Incubation with HeLa Cells (Microscopy).** HeLa cells were seeded at 30000 cells/well density in 8-well LabTek, 200 μL/well, 24 h prior to the experiment. Medium was changed with fresh DMEM (10% FBS), and micelles were diluted 3× to the final concentration of 160 μM on the cells. Samples were imaged in a confocal microscope (Infinite M200 Pro from Tecan) by sampling the emission from the bottom well at 600 nm, while exciting at 550 nm. Each sample was taken in three replicates, distributed randomly on each row (using randomizer.org). The signal was normalized using the negative and positive controls between 0 and 100%, respectively.

**Molecular Design and Synthesis.** We chose 7-(diethylamino)coumarin-3-carboxylic acid (7-DEAC) as the fluorescent tag due to its excimer formation ability that allows to distinguish whether the amphiphiles disassembled into unimers or remained as micelles under various conditions.53,54 Once 7-DEAC dyes are forced to be in close proximity within the micelles, their emission maxima shifts from 480 to ~540 nm, and once the micelles disassemble, the emission shifts back to 480 nm. Our synthetic methodology allows simple preparation of libraries of amphiphilic diblock copolymers that can be examined and compared in many aspects ranging from micellar stability and enzymatic degradability to more complex biological studies that are enabled due to the dendron’s unique fluorescent response.

The synthetic route of the dendron is based on high yielding converging synthesis (Scheme 1). The synthesis started from the thiol–yne reaction between the AB2 branching unit and pre-made thiol-functionalized degradable hydrophobic end-groups, followed by coupling with a propargylated lysine-based taggable unit for fluorescent labeling. The dendron was then deprotected and labeled with the coumarin dye. In parallel, PEtOx-N3 (5 kDa) with narrow molecular weight distribution was synthesized by cationic ring opening polymerization of 2-ethyl-2-oxazoline using methyl tosylate as the initiator and terminated by sodium azide as previously reported.55,56 mPEG-N3 (5 kDa) was synthesized from commercially available mPEG-OH in three high yielding synthetic steps (Figure S21).

**RESULTS AND DISCUSSION**

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**Flow Cytometry on HeLa Cells Incubated with Micelles.** HeLa cells were seeded in a 24-well plate at 120000 cells/well in 1 mL/well 24 h before the experiment. Micelles were added by diluting 3× to the 160 μM final concentration, for either 1 or 6 h of incubation. Cells were washed 2× with warm PBS, then trypsinized with 250 μL/well for 3–4 min at 37 °C, mixed with 750 μL/well full DMEM, centrifuged 3 min at 180 g, and then resuspended in 1 mL/well warm PBS.

The fluorescence spectra with 405 nm excitation was recorded for 8000–10000 cells per sample using a spectral analyzer (Sony SA3800) flow cytometer. For data analysis, the spectral signal was gated in two “channels” representing unimers (420–500 nm) and micelles (550–700 nm).

**Encapsulation Stability in the Presence of BSA.** Micellar solution of the tested amphiphile was prepared in PBS (176 μM).

Hydrophobic Cy5 derivative was added directly (2 μL/mL from 2 mM Cy5 solution in EtOH), and solution was thoroughly vortexed. Then, 50 μL of either BSA solution (55 mg/mL in PBS) or PBS were added into 450 μL of the above solution, and solution was vortexed to obtain final concentrations of 160 and 4 μM for amphiphile and Cy5, respectively, and 5.5 mg/mL for BSA. Absorbance of all final solutions was measured at T0 and the emission spectra was recorded every 30 min for 2 h (λex = 420 nm).

**Imaging of Cy5 Release Experiments on HeLa Cells.** Micelle solution in PBS (480 μM) was mixed with Cy5 solution in ethanol (2 mM) to the final Cy5 concentration 12 μM. Mixture was vortexed and filtered through nylon 0.45 μm syringe filters (PureTech). Then, each solution was diluted with fresh DMEM (10% FBS) by three fold to final micelle and Cy5 concentration of 160 and 4 μM, respectively. Absorbance of each solution was measured in order to verify similarity in concentrations. Imaging was performed similarly to the cell internalization experiment. One field of view was followed for 1 h for each sample. For 405 nm excitation (5 mW diode laser, 1%), the acquisition was split in three channels representing unimer (400–500 nm), micelle (500–617 nm), and possible FRET fluorescence (656–700 nm). For total Cy5 fluorescence, 640 nm excitation (5 mW diode laser, 0.2%) was used, with 656–700 nm acquisition. Also, an electronically switchable illumination and detection module (ESID, bright field-like) signal was acquired using a 488 nm (10 mW) diode laser.

For image analysis (on Fiji ImageJ), regions of interest were drawn manually either outside the cells or containing cell cytoplasm and membrane, without the nucleus. Mean fluorescence inside cells and median outside cells were plotted using GraphPad Prism.
PMDETA catalyst as reported by Matyjaszewski. All polymers reacted neatly, and full conversion was confirmed by $^1$H NMR spectroscopy and size exclusion chromatography (SEC). Purification of the amphiphiles after the click reaction was performed using preparative SEC, and all polymers were obtained in high purity and yield. In the case of the PAA-based amphiphiles, an additional deprotection step of the $t$-butyl esters was performed. All final amphiphiles were characterized by $^1$H and $^13$C NMR, HPLC, SEC (PAA injected in its protected form), MALDI (except PAA), and UV–vis spectroscopy.

**Micellization.** Upon completing the synthesis of the six amphiphiles, we studied their self-assembly in aqueous media (PB, pH 7.4, at 37 °C). At first, the CMC of each polymer was determined using the Nile Red method. The CMC values of the amphiphiles with hexyl end-groups were determined to be $5 \pm 1$, $6 \pm 1$, and $10 \pm 2 \mu M$ for PEtOx-Hex, PEG-Hex, and PAA-Hex, respectively. The CMC values of the amphiphiles with nonyl end-groups were determined to be $3 \pm 1$, $5 \pm 1$, and $9 \pm 1 \mu M$ for PEtOx-Non, PEG-Non, and PAA-Non, respectively. Even though there is a substantial difference in the chemical composition of the micelle coronas, the CMCs of the three amphiphiles in each series are very similar with slightly higher CMC value for the PAA-based amphiphiles, most likely due to the greater hydrophilicity and repulsion of the charged PAA chains. Next, we used DLS to measure the diameters of the different micelles (Figure 1). The hexyl polymers self-assembled into micelles with diameters of $20 \pm 2$, $21 \pm 2$, and $20 \pm 4 \text{ nm}$ for PEtOx-Hex, PEG-Hex, and PAA-Hex, respectively, and the nonyl ones into $28 \pm 5$, $26 \pm 5$, and $32 \pm 5 \text{ nm}$ for PEtOx-Non, PEG-Non, and PAA-Non, respectively. It was fascinating to see the similar sizes of the micelles despite the different hydrophilic shells, demonstrating the key contribution of the hydrophobic block in directing the self-assembly of these polymeric amphiphiles into micelles.

**Enzymatic Degradation Rate.** After characterizing the self-assembly of the amphiphiles, we studied the enzymatic degradation rates of the six amphiphiles with PLE. PLE can selectively cleave the ester bonds between the dendron and the hydrophobic end-groups, exposing highly hydrophilic carboxylic acids on the dendron chain ends. This enzymatically induced modification will turn the polymer from amphiphilic into fully hydrophilic and therefore should cause the disassembly of the micelles (Figure 2A). Three methods were used to monitor the degradation of the amphiphiles and the disassembly of the micelles. Due to the high precision and purity of the polymers, we could directly monitor the degradation of the starting material and the appearance of the degraded polymer using HPLC (Figure 2D,E, solid lines and Figures S33 and S34). Simultaneously, we followed the enzymatically induced disassembly by monitoring 7-DEAC fluorescent response under the same conditions. Once an amphiphile’s end-groups are cleaved, it becomes fully hydrophilic and diffuse away from the micelle. This leads to a decrease in 7-DEAC excimer formation, and this spectral response can be quantified by measuring the decrease in fluorescence emission at 540 nm and the increase at 480 nm (Figure 2B–E, dashed lines). Finally, DLS was used to determine whether micelles are present in solution before and after 24 h incubation at 37 °C with or without PLE (Figure 1, dashed lines). Notably, the nonyl amphiphiles showed high stability in the presence of PLE even at concentration as high as $1.4 \mu M$. PEtOx-Non and PEG-Non showed approximately 10% degradation over 24 h, and around 20% degradation was observed for PAA-Non (Figure 2E).

Strikingly, although the change in the hydrophilic to hydrophobic ratio is relatively small for the nonyl and hexyl series as reflected by the small differences in CMC values, the hexyl-based amphiphiles were significantly more susceptible to enzymatic degradation. While full degradation of PEG-Hex required longer time than PEtOx-Hex, their $t_{1/2}$ was almost identical. PAA-Hex, in contrast, showed ultrafast response to PLE and was fully degraded after less than 30 min (Figure 2D, full lines). The faster degradation of the PAA amphiphile may be related to the higher hydrophilicity of the PAA compared to PEG and PEtOx, while the longer chain length of PEG-5k compared to PEtOx-5k and/or the slightly better anti-fouling behavior of PEG may be responsible for the slower degradation of the PEG amphiphile. Fluorescence measurements showed a decrease in the longer wavelength emission of the micelle and increase in unimer emission, indicating that the enzyme led to disassembly of the micelles. Excellent correlations were observed between the degradation kinetics obtained by HPLC and the decrease in fluorescence intensity at 540 nm (Figure 2D,E), dashed lines), which is indicative of the disassembly of the cleaved polymers into unimers. This clearly demonstrates that, indeed, it is the cleavage of the hydrophobic end-groups by the enzyme that caused the micelles to disassemble. Last, DLS measurements after incubation with PLE confirmed the disassembly of the hexyl-based micelles, while the nonyl-based series stayed intact (Figure 1, dashed lines). In the control experiments conducted in the absence of PLE, all amphiphiles showed micelles with similar sizes to the ones measured at $t_0$ after 24 h incubation (Figure 1, dotted lines, and Figures S48 and S49). The similarities in CMCs and micelle sizes did not hint at the extreme difference in enzymatic degradation kinetics between the hexyl and nonyl polymers. The best example for this difference is seen for the two PAA-based amphiphiles, whereby PAA-Hex degraded in minutes while PAA-Non showed a very limited degree of degradation even after 24 h of incubation. This shows once more the tremendous effect of the hydrophobic block on the micellar dynamics and the importance of molecular precision when designing enzyme-responsive polymeric amphiphiles.
Drug Encapsulation Capacity. Next, we investigated how the different hydrophilic shells affect the micelles encapsulation capacity for hydrophobic cargo. For this purpose, we chose the widely used hydrophobic anticancer drug CPT, which has very poor aqueous solubility. The CPT-loaded micelles were made by the thin-film hydration method, as previously reported. Unencapsulated CPT was removed by filtration through a 0.45 μm filter, and the filtrate was analyzed by HPLC to determine the drug concentration and the drug to polymer weight ratio. The CPT encapsulation results for the hexyl series were 5 ± 1, 18 ± 3, and 8 ± 1 weight percentage for PEtOx-Hex, PEG-Hex, and PAA-Hex, respectively (Figure 3A, black columns and Table 1). Surprisingly, the nonyl series showed lower encapsulation capacities of 1.0 ± 0.1, 14 ± 5, and 4.0 ± 0.3 weight percentages for PEtOx-Non, PEG-Non, and PAA-Non, respectively (Figure 3A, pink columns). These results are in good agreement with previous reports by Luxenhofer and co-workers, who observed a reduction in drug loading capacity upon increasing the hydrophobicity of the hydrophobic block in poly(2-oxazoline)-based micellar nanocarriers. In both, amphiphile series PEG micelles showed a higher encapsulation capacity than PAA micelles while PEtOx micelles showed the lowest encapsulation capacity. To determine whether the encapsulation ability of the different polymers is general or depends on the payload as previously suggested, we also prepared formulations of another, slightly less hydrophobic anti-cancer drug, paclitaxel (PTX). Surprisingly, the trends with PTX were completely different and PEtOx micelles showed significantly higher encapsulation capacity than PEG micelles and PAA micelles (Figure 3B and Table 1). It is important to point out that the chemical composition of the micellar core in each of the hexyl and nonyl series is completely identical and, therefore, the observed differences

Figure 2. (A) Schematic representation of the enzymatic degradation of the hydrophobic end-groups turning the polymeric amphiphile into fully hydrophilic polymers leading to micelle disassembly. Pictures of the fluorescence of PEG-Hex after 30 h incubation without (B) and with (C) PLE demonstrating the system’s spectral response. Enzymatic degradation profiles of hexyl- (D) and nonyl- (E) based micelles as monitored by HPLC (solid lines) and fluorescence spectroscopy (normalized decrease of emission intensity at 540 nm, dashed lines). [amphiphile] = 160 μM, [PLE] = 1.4 μM.

Figure 3. Encapsulation capacities of (A) CPT and (B) PTX in the different micelles (in μM and drug/polymer wt %). [amphiphile] = 160 μM in PB 7.4.
in encapsulation capacities are solely dictated by the different micellar coronas. This may mean that the common way of thinking and illustrating micelles with their hydrophobic cargo trapped only inside the micelle core is not accurate and a significant amount of the cargo might be in the corona, solubilized also between the hydrophilic polymer chains.\textsuperscript{35,44,68} In addition, the hydrophilic corona may influence the core packing and hydration, which can also influence the drug encapsulation behavior for drugs with different hydrophobicity. After comparing the observed trends in encapsulation capacities, it is clear that the type of hydrophilic polymers that are chosen for DDS should be carefully selected to fit with the drug to be encapsulated.

**Interaction with Blood Proteins.** After analyzing the in vitro properties of the six types of micelles, we moved on to study how these structural features influence their biological interactions. One of the first things that occur upon introduction of nanoparticles into the bloodstream, in addition to significant dilution, is the interactions with proteins and their adsorption resulting in the formation of a protein corona,\textsuperscript{69−72} which has been demonstrated to be key for shielding DDS from the immune system.\textsuperscript{3} Therefore, we investigated the interactions between the different micelles and albumin, which is the most abundant serum protein,\textsuperscript{73} as an initial evaluation of micelle—protein interactions. In order to understand the differences in protein adsorption, we incubated the micelles with BSA labeled with Cy5, which serves as a FRET acceptor for the micelle fluorescence (and much less with the shorter wavelength unimer’s fluorescence).\textsuperscript{74} FRET is highly dependent on spatial proximity (1−10 nm)\textsuperscript{75} and therefore the intensity of Cy5 emission qualitatively correlates with the interaction between the micelles and BSA. BSA is known to have hydrophobic regions that can interact with unimers and therefore significantly destabilize the micelles.\textsuperscript{76−78} This type of interaction translates into an increase in the unimer emission (480 nm) and a decrease in micelle emission (540 nm), as illustrated in Figure 4A and B. The increase in the hydrophobicity of the dendron changes the unimer−micelle equilibrium and decreases the interactions between unimers and BSA, indicating an increase in micellar stability (Figure 4C). PEtOx and PEG micelles showed overall similar behavior with slightly weaker interactions of PEtOx micelles with BSA indicated by a smaller peak of the FRET signal. For both PEtOx and PEG hexyl micelles, there was a moderate increase in the unimer emission, which is indicative of their interaction with BSA (Figure 4C). Almost no changes were observed for PEtOx and PEG nonyl micelles, which did not show an increase in the unimer peak upon incubation with BSA. On the other hand, PAA-Hex showed complete disassembly due to interactions with BSA as indicated by the total disappearance of the micelle fluorescence in addition to a significant increase in the unimer signal intensity (Figure 4C and D). The reason for this increase might be the change in the 7-DEAC microenvironment that can influence its quantum yield dramatically. PAA-Non also showed an increase in the unimer signal intensity, but the micelle signal still partially

| Drug/Polymer Molar Ratio and Loading Capacity | Loading Capacity [%] |
|---------------------------------------------|----------------------|
| ![Image](https://example.com/image1.png) | ![Image](https://example.com/image2.png) |
| (drug)/[polymer] | CPT | PTX | CPT | PTX |
| PEtOx-Hex | 0.84 ± 0.16 | 3.09 ± 0.04 | 4.3 ± 0.9 | 29.2 ± 0.3 |
| PEG-Hex | 3.36 ± 0.53 | 0.54 ± 0.13 | 15.5 ± 2.4 | 6.7 ± 1.6 |
| PAA-Hex | 1.40 ± 0.19 | 0.59 ± 0.01 | 7.1 ± 1.0 | 7.3 ± 0.1 |
| PEtOx-Non | 0.24 ± 0.01 | 2.16 ± 0.50 | 1.3 ± 0.1 | 21.9 ± 5.1 |
| PEG-Non | 2.69 ± 1.01 | 1.53 ± 0.35 | 12.5 ± 4.7 | 16.6 ± 3.7 |
| PAA-Non | 0.77 ± 0.06 | 0.32 ± 0.03 | 3.9 ± 0.3 | 4.0 ± 0.3 |

**Figure 4.** (A) Illustration of the two main possible interaction pathways with Cy5-labeled BSA: interaction of BSA either with unimer (left) or micelle (right). (B) Selected fluorescence emission spectrum with arrows highlighting the contribution of the different species to the spectrum. (C) Fluorescence spectra of the micelles with (red lines) and without (blue lines) BSA-Cy5 and (D) zoom out into PAA amphiphiles emission spectra. [amphiphile] = 160 μM, [BSA-Cy5] = 5.5 mg/mL, λ\textsubscript{ex} = 420 nm.
remains, and the Cy5 signal is the most intense amongst all micelles, suggesting the strongest micelle–BSA interactions among the different amphiphiles. To monitor the destabilization of the micelles over time, we calculated the unimer to the micelle fluorescence ratio (Figure S50). All ratios were nearly constant during 5 h, except for PAA-Non, which showed an increase in the unimer/micelle ratio. Overall, this suggests that for all micelles, the interactions with BSA happen almost immediately after addition, leading to various types and degrees of interactions. While the more hydrophobic nonyl micelles generally interact less with BSA, the PAA-Non showed remarkable differences in the degree of internalization of the micelles. To assess whether this difference is due to disassembly of the micelles outside or inside the cells, we analyzed the ratiometric images (Figure S5A bottom row). For all amphiphiles except PAA-Hex, the ratiometric analysis indicated the presence of micelles outside the cells, while inside cells or on the cell membrane, all amphiphiles were mostly in their unimer form (Figure 5C). However, slightly more micelles were observed inside the cells for the more stable PEtOx-Non and PEG-Non (Figure 5B). The micelles of PEtOx-Non and PEG-Non that were shown to be more stable showed a very weak signal within the cells. While PEtOx-Hex and PEG-Hex had similar distributions in intracellular vesicles, PAA-Hex and PAA-Non bound mostly to the cell membrane and had the most intense fluorescence emissions, similar to the trend observed for the incubation with BSA. Thus, we can assume that the hydrophilic block directs the cellular fate of the micelles toward the endo-lysosomal compartments for PEtOx and PEG and membrane-bound for PAA. Interestingly, the relatively small change in hydrophobicity causes a notable decrease in internalization efficiency for the nonyl micelles. To assess whether this difference is due to disassembly of the micelles outside or inside the cells, we analyzed the ratiometric images (Figure 5A bottom row). For all amphiphiles except PAA-Hex, the ratiometric analysis indicated the presence of micelles outside the cells, while inside cells or on the cell membrane, all amphiphiles were mostly in their unimer form (Figure 5C). However, slightly more micelles were observed inside the cells for the more stable PEtOx-Non and PEG-Non (Figure 5B,C). We can assume that the less stable PAA micelles disassembled outside the cells more readily into unimers that could then intercalate into the plasma membrane. For the polymers that were localized in endosomal vesicles, it may be that they internalized as micelles and very rapidly disassembled inside the endosomal vesicles into unimers. To achieve higher degree of internalization and obtain more intense fluorescence signal inside the cells, the experiment was repeated with a longer incubation time of 6 h (Figure S52). The longer incubation time indeed led to a stronger signal within the cells, which was mostly observed in the unimer channel. Although the ratiometric images indicated the presence of unimers inside the cells, it is most likely that the increased internalization cannot be attributed to the disassembly of the micelles outside of the cells over time. This assumption is based on the high micellar stability for PEtOx-
Non and PEG-Non, which remained highly stable when incubated with BSA, PLE, and outside of the cells. The degree of internalization was further validated by spectral flow cytometry after incubation times of 1 and 6 h using both unimer and micelle channels (Figure S53).

**Encapsulation Stability and Cargo Release in the Presence of BSA and Cell Culture.** In addition to the crucial effect of micellar stability on the ability of the micelles to retain their molecular cargo, one of the major drawbacks in physical encapsulation of hydrophobic compounds is the possibility of premature leakage, that is, before reaching the tumor through passive targeting via the EPR effect, due to migration of the cargo into native hydrophobic regions in the surroundings of the carrier such as proteins or membranes.35,79 Hence, after studying the stability of the micelles, we evaluated their ability to retain their cargo of hydrophobic molecules in a biological environment. Therefore, another FRET-based experimental setup was designed in order to study how encapsulated hydrophobic cargo behaves in the presence of BSA. A hydrophobic Cy5 derivative (Figure S17), which served as a model for an encapsulated lipophilic drug, was physically encapsulated within the different micelles, and non-labeled BSA was added to the micellar solution. The encapsulated Cy5-derivative undergoes significant FRET with the fluorescence of the micelles while its migration from the micelles to BSA or its precipitation due to micelle disassembly into unimers should translate into a reduction in FRET efficiency (Figure 6A).

Prior to the fluorescence measurements, the absorbance spectra of all tested solutions containing micelles and BSA, as well as the controls without BSA, were measured to verify that the concentrations of the polymers and Cy5 were similar for all solutions (Figure S54). Next, fluorescence spectra were measured every 30 min over 2 h. Upon addition of BSA, both PEtOx and PEG micelles showed a decrease of ~30% in FRET-related emission, while a slight increase in both the micelle and unimer emission was observed (Figure 6B,C). These results indicate that the majority of encapsulated Cy5 molecules remained entrapped inside the micelles as complete Cy5 release would lead to a complete disappearance of the FRET signal and a substantial increase in micelle fluorescence would be expected. A slight increase in unimer emission is attributed to the interaction of the micelles with the BSA, as described above (Figure 4). Interestingly, the least stable PAA-Hex micelles showed a significantly lower FRET signal in the absence of BSA, which was sustainably reduced by 50% upon the addition of BSA. In addition, a strong increase in the unimer emission was also observed, indicating the low stability of the PAA-Hex micelles and their tendency to disassemble due to interaction with BSA. Unlike the PAA-Hex, the PAA-Non-based micelles showed only a moderate decrease of ~30% of the FRET signal in the presence of BSA, similar to the PEG and PEtOx micelles. The notable increase in the unimer emission of the PAA-Non can again be attributed to the stronger interaction with BSA (Figure 6A). The response to BSA was extremely fast in this experiment as noted before in the Cy5-labeled BSA assay, and no major changes were observed over time, which is indicative of the high encapsulation stability of the PEG, PEtOx, and PAA-Non micelles (Figure S55).

Intrigued by the encapsulation stability of the micelles in the presence of BSA, we decided to study the release of the Cy5 dyes in HeLa cell culture. Two possible mechanisms for the release of physically encapsulated cargo from polymeric micelles can be envisioned: (i) spontaneous leakage or (ii) disassembly and release. Spontaneous leakage would leave the micelles intact, while the Cy5 molecules would exit and accumulate inside cells. On the other hand, disassembly based...
release could occur outside the cells followed by internalization of the unimers together with the released Cy5 dyes. Alternatively, the Cy5 containing micelles could be internalized followed by disassembly and cargo releasing in the endosomal vesicles. Based on the inferior stability noted in all previous experiments for the PAA micelles, we expected that the release of encapsulated cargo would be significantly faster in the presence of cells as a large fraction of PAA micelles disassembled in the presence of HeLa cell culture. To study the effect of micellar corona composition on the release kinetics of hydrophobic cargo, we decided to prepare micelles only from the more stable nonyl amphiphiles. The three types of micelles were loaded with Cy5 and incubated with HeLa cells. The same cells were followed for 1 h, measuring the fluorescence of the unimer and micelle (excitation at 405 nm) as well as directly following the Cy5 (excited at 640 nm, Figures 7A and S57). For every time point, the fluorescence was measured inside the cytoplasm of 10 cells and the mean values were compared to the signal outside the cells (Figure 7B). This unique setup provided important new insights regarding the behavior of the different micelles. By monitoring the ratio of unimer/micelle fluorescence, the assembly state of the amphiphiles in each pixel is revealed (Figure 7C). In addition, the Cy5 channel enabled direct monitoring of the cell internalization kinetics of the encapsulated cargo (Figure 7B).

We hypothesized that the combination of the ratiometric data together with the direct tracking of Cy5 release would shed light on the release mechanism of the different micelles. First, we examined the assembly state of the micelles. As seen in the previous internalization experiments (Figure S5), PAA-Non micelles showed significant disassembly both outside and inside the cells, which increased over time (Figure 7C).

As expected, the PEtOx-Non and PEG-Non micelles were much more stable than the PAA-Non micelles and almost no disassembly was observed, as indicated by the nearly constant unimer/micelle fluorescence ratio (Figure S57). Therefore, we

Figure 7. Release of encapsulated Cy5 onto HeLa cells over time. (A) Fluorescence with 640 nm excitation is shown inside HeLa cells at different time points for hydrophobic Cy5 encapsulated in micelles or free, in full DMEM (10% FBS), scale bar is 20 μm. The quantification of fluorescence from confocal images with (B) as mean fluorescence intensity in the Cy5 channel, with 640 nm excitation or (C) 405 nm excitation is shown as ratio of unimer/micelle signal. Regions of interest were manually drawn around cell cytoplasm, including the cell membrane and excluding the nucleus; n = 10 cells. [amphiphile] = 160 μM, [Cy5] = 4 μM.
were rather surprised to discover that the release and internalization of Cy5 was significantly slower from PAA-Non micelles compared with PEG-Non and PEtOx-Non. While both PEtOx-Non and PEG-Non samples showed dramatic increase in the Cy5 signal inside the cells, PAA-Non showed delayed and more gradual increase, even though its tendency to undergo micellar disassembly is much higher. The results of these experiments demonstrate that lower micellar stability does not always correlate with faster release kinetics, and that disassembly of the carrier is not essential for cargo release (Figure 8). Furthermore, these findings clearly emphasize the importance of tracking both the carrier and the cargo when studying the internalization mechanism and kinetics of DDS.

![Figure 8](image)

**Figure 8.** Schematic illustration of two mechanisms of cargo release by either leakage from stable micelles or disassembly of the polymeric carrier.

### CONCLUSIONS

In summary, we designed a highly modular approach for preparing clickable spectrally active enzyme-responsive dendrons with adjustable degree of hydrophobicity. The dendrons were synthesized using a thiol–yne reaction and then conjugated by the CuAAC click reaction with three different hydrophilic polymers, namely PEG, PEtOx, and PAA, with similar molecular weights, enabling us to prepare six amphiphiles in different compositions. The dendrons, when conjugated with either hydrophilic or hydrophobic polymers, self-assembled into micelles with different molecular weights, enabling us to prepare six polymeric micelles with different compositions.

The drug loading results clearly indicate the importance of the nature of the hydrophilic shell for the encapsulation of hydrophobic drugs. In addition, the composition of the hydrophilic block had a strong effect on the interactions of both unimers and micelles with BSA, which in the case of PAA-Non led to the complete disassembly of the micelles. Cell internalization experiments revealed a substantial difference in the membrane binding and internalization rate between PAA-Non and PEtOx-Non micelles. PAA-based amphiphiles localized more on the cell membrane and internalized to a greater extent than the PEG- and PEtOx-based amphiphiles. In all cases, the more hydrophobic nonyl amphiphiles internalized significantly more slowly than the hexyl ones. Using a hydrophobically modified Cy5 dye as a model for encapsulated drug molecules, it was interesting to see its slower release and cell internalization in the case of the less stable PAA-Non-based micelles, while much faster Cy5 cell internalization was observed for the PEG-Non and PEtOx-Non micelles, which seemed to self-assemble into significantly more stable micelles. These encapsulation and release experiments in cell culture revealed the complexity of studying release mechanisms and the importance of directly tracking both the carrier and the cargo. Overall, the ability to directly compare micelles with different shells and the resulting comparative results provide important fundamental insights into how the composition of the shell and core of such polymeric micelles can affect their properties and potential to serve as nanocarriers for DDS.

### ASSOCIATED CONTENT

* Supporting Information
  
  The Supporting Information is available free of charge at [https://pubs.acs.org/doi/10.1021/acs.biomac.0c01708](https://pubs.acs.org/doi/10.1021/acs.biomac.0c01708).

  Synthetic procedures, amphiphiles and micelles characterization data, detailed experimental protocols, control experiments, cytotoxicity data, and additional confocal imaging (PDF)

### AUTHOR INFORMATION

**Corresponding Authors**

Lorenzo Albertazzi — Institute for Bioengineering of Catalonia (IBEC), The Barcelona Institute of Science and Technology, 08028 Barcelona, Spain; Department of Biomedical Engineering, Institute of Complex Molecular Systems (ICMS), Eindhoven University of Technology (TUE), Eindhoven 5612 AZ, The Netherlands; orcid.org/0000-0002-6837-0812; Email: L.Albertazzi@tue.nl

Roey J. Amir — Department of Organic Chemistry, School of Chemistry, Faculty of Exact Sciences, BLAVATNIK Center for Drug Discovery, ADAMA Center for Novel Delivery Systems in Crop Protection, and The Center for Physics and Chemistry of Living Systems, Tel-Aviv University, Tel-Aviv 6997801, Israel; Tel Aviv University Center for Nanoscience and Nanotechnology, Tel-Aviv University, Tel-Aviv 6997801, Israel; orcid.org/0000-0002-8502-3302; Email: amirroey@tauex.tau.ac.il

**Authors**

Gadi Slor — Department of Organic Chemistry, School of Chemistry, Faculty of Exact Sciences, Tel-Aviv University, Tel-Aviv 6997801, Israel; Tel Aviv University Center for...
Complete contact information is available at:

https://pubs.acs.org/10.1021/acs.biomac.0c01708

Author Contributions

G.S. and A.R.O. authors contributed equally.

Funding

R.J.A. thanks the Israel Science Foundation (grant no. 1553/18) for the support of this research. This project has received funding from the European Union’s Horizon 2020 research and innovation program under the Marie Skłodowska-Curie [grant agreement no. 713673 (THERACAT)].

Notes

The authors declare the following competing financial interest(s): R.H. and V.R.D.L.R. are co-founders of Avroxa BVBA that commercializes poly(2-oxazoline)s as Ultroxa. The other authors have no conflicts of interest to declare.

ACKNOWLEDGMENTS

G.S. thanks the Marian Gertner Institute for Medical Nanosystems in Tel Aviv University for their financial support. R.H. and V.R.D.L.R. are grateful to Ghent University and FWO for continuous financial support.

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