Cite this: *Polym. Chem.*, 2020, **11**, 2775

Received 20th February 2020,
Accepted 30th March 2020

DOI: 10.1039/d0py00280a

rsc.li/polymers

Influence of surface charge on the formulation of elongated PEG-*b*-PDLLA nanoparticles†

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Polymeric vesicles (polymersomes) are an important class of nanoparticles for various biomedical applications. Their interaction with biological systems is strongly dependent on their topological features such as size, shape and surface charge. We have recently developed a versatile method that enables the formation of tubes or bowl-shaped vesicles (stomatocytes) out of spherical vesicles composed of the biodegradable block copolymer poly(ethylene glycol)-poly(D,L-lactic acid) (PEG-*b*-PDLLA), via a dialysis process. Applying this method for particles with different surface charge is however a far from trivial task, as the shape change process is affected by spontaneous membrane curvature, which is strongly influenced by the presence of charged polymer end groups. Here we describe an optimized procedure to attain an effective shape transformation toward tubes of PEG-*b*-PDLLA polymersomes containing PEG blocks with amine (A) or carboxylic acid (CA) end groups. The salt concentration employed during the dialysis process turns out to be key, with the CA-polymersomes requiring substantially lower concentrations than unmodified or A-ones. The ability to control now reliably shape and surface charge in these polymer vesicles, allows a future systematic analysis of the effect of these topological parameters on the biological response of the nanoparticles.

Introduction

Nanoparticle morphology is recognised to be a key determinant of performance in the field of nanomedicine.^{1–6} Contrary to other particle features such as size and surface charge, shape only recently received attention as an important factor determining the behaviour of particles in a biological context.

High-aspect ratio nanoparticles have been shown to have improved biophysical properties with regard to for example flow characteristics (influencing circulation and distribution) and interactions with cells/tissues.^{6–8}

An important class of nanoparticles that have been used in a diverse range of applications including drug delivery,⁹ radiotherapy,¹⁰ cancer cell targeting,¹¹ and nanoreactors¹² are polymeric vesicles, or polymersomes.^{13,14} The compartmentalized nature of polymersomes facilitates encapsulation of various types of cargo within their inner lumen, protecting them from unwanted degradation whilst facilitating their delivery to cells through passive or active uptake (using homing motifs such as peptides or antibodies).^{4,15} The shape effect of course also holds for polymersomes, and this has inspired researchers to develop effective methods for the construction of non-spherical vesicular structures.

For this purpose different processes have been reported such as polymerisation-induced self-assembly (PISA),^{16–18} film re-hydration,^{19–21} or co-solvent/evaporation.⁷ The PRINT technology²² and moulding^{23,24} have also proven to be very useful to create nanoparticles with different shapes, albeit that these techniques predominantly have been used for solid systems. In our group we have constructed non-spherical polymersomes by employing a dialysis procedure; block copolymers were dissolved in an organic solvent, after which water was added to induce polymer assembly. The organic solvent was subsequently removed *via* dialysis. By careful control over the latter step a shape change could be realized. We were able to demonstrate this both for non-degradable block copolymers such as poly(ethylene glycol)-poly(styrene)^{25,26} and, more recently, biodegradable block copolymers.^{1,27} In this way, we have generated tubular polymersomes (nanotubes) and bowl-shaped vesicles known as stomatocytes comprising poly(ethylene glycol)-poly(D,L-lactide) (PEG-*b*-PDLLA) block copolymers.

Surface charge is another important feature of nanoparticles that can have a major effect on their performance as drug delivery systems. Particle charge has been shown to influence phagocytosis, cellular uptake and biodistribution.^{28–30} Reports on various polymeric nanoparticles indicate better cel-

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†Electronic supplementary information (ESI) available. See DOI: 10.1039/d0py00280a

lular uptake and biodistribution for positively charged particles.^{31–33} Negatively charged particles were usually less toxic^{34,35} and more prone to permeate tissues for targeted delivery.^{36,37} Internalization pathways have also been linked to surface charge, with positively charged particles tending towards internalization *via* clathrin receptors, while negatively charged counterparts are prone for caveolae mediated uptake.³³ With this in mind, methods should be available in which both shape and surface charge can be effectively engineered to tailor the performance of nanoparticles *in vivo*.

Various strategies exist for the incorporation of charge onto polymeric nanoparticles including synthetic approaches and post-modification.^{30,33,35,38–42} Herein, we present a facile approach whereby the surface charge of non-spherical polymersomes was modified by blending uncharged methoxy-terminated PEG-*b*-PDLLA block copolymers with either amino (A) or carboxyl (CA) terminated variants. Assembly was conducted using the solvent switch methodology, where water was slowly added to a solution of polymer in organic solvent. By systematically varying the salt concentration during the subsequent dialysis procedure, we were able to transform the differently functionalized spherical polymersomes into nanotubes and, under certain conditions, stomatocytes. Having attained a robust route to the fabrication of functionalized polymersomes (either spherical or tubular) with different surface groups, their toxicity was tested against human retinal (ARPE-19) cells – where they yielded no evidence of harmful effects to cells.

Research and discussion

Using either monomethoxy, carboxyl- or NHBoc-terminated poly(ethylene glycol) macroinitiators of narrow polydispersity,

block copolymers (BCPs) were synthesized through the organo-catalyzed ring-opening polymerization (ROP) of *D,L*-lactide monomers using 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) as a catalyst (S1).²⁷ This yielded BCPs with a well-defined (45 repeats, 6.5 kDa), hydrophobic PDLLA block (after reaction for 2 h at room temperature) with a hydrophilic PEG block of either 1 or 2 kDa for uncharged and charged BCPs, respectively (Fig. 1A and B). This difference in length was chosen to ensure that the incorporated charged groups were accessible at the particle surface. All BCPs had polydispersity values of approximately 1.1, which highlights the excellent control that was achieved using this method (Fig. 1A/S2–S3). Subsequent deprotection of the PEG-terminal amine was performed using trifluoroacetic acid (TFA) in DCM to yield the primary amine, with no detrimental effect on the reaction or polydispersity. The use of acid-terminated PEG did not appear to interfere with the polymerisation. The composition of all BCPs was assessed using ¹H NMR spectroscopy, which confirmed that precise polymerization control was achieved (Fig. 1A/S4–S6). In this way, a range of well-defined polymers were prepared as a first step in formulating biocompatible polymersomes that would be able to display either negative or positive charge at neutral pH through blending with the non-charged variant.

Self-assembly of BCPs was performed using the solvent switch method, by dissolving the polymer in a 4 : 1 mixture of THF : dioxane prior to dropwise addition of water up to 50 vol% over 2 h (Fig. 1C).²⁷ The formation of polymersomal nanostructures was indicated by the increased turbidity of the solution, without the formation of unwanted aggregates. A- or CA-modified polymersomes were formed using the same process, with the introduction of 10 wt% (with respect to total polymer content) of either A or CA terminated BCPs, respectively. From cryo-TEM measurements, polymersomal mor-

Fig. 1 (A) Characterisation of synthesized poly(ethylene glycol)-*block*-poly(*D,L*-lactide) (PEG-*b*-PDLLA) copolymers; (B) molecular structure of PEG-*b*-PDLLA; (C) formulation scheme for the preparation of PEG-*b*-PDLLA vesicles.

phologies were evident in all three samples, with some influence of charge upon the size (Fig. 2A–C). Unmodified polymersomes, comprising neutral PEG-*b*-PDLLA, attained sizes of approximately 500 nm which was also the case for the A-modified samples; however, there was a significant difference in size for the CA-doped samples which were clearly smaller (*ca.* 200–300 nm). This was attributed to the difference in solvent interactions between A- and CA-containing polymers that would influence the kinetics of polymersome formation, thereby altering the average size of the nanostructures. A uniform membrane thickness of 15–20 nm was observed for all formulations.

Inducing shape transformations of PDLLA-based block copolymers from a spherical to elongated tubular morphology was achieved using osmotically induced deflation, as described previously.²⁷ However, the introduction of charged polymers affected the shape transformation process markedly, showing different morphological changes under the same dialysis conditions (overnight dialysis against water for spheres, or salted water for elongated shapes). Unmodified and CA-modified particles formed well-defined elongated tubes, with an average size of 100 nm (width) $\times$ 1 μ m (length), whereas A-modified charged variants were not as elongated and more closely resembled peanut shaped morphologies as opposed to tubes (Fig. 2D/E). During the shape-change process, an internal volume reduction of *ca.* 50% occurred, leading to an irreversible change in morphology owing to the semi-permeability characteristics of the membrane. We

hypothesized that addition of charged groups at the surface of the polymersomes would induce distinct (lateral) electrostatic interactions, influencing the membrane flexibility, its spontaneous curvature and, as a result, the shape change pathway. Spontaneous membrane curvature (influenced by the packing of polymer chains in the membrane, their interactions with each other and surrounding solvent) is a controlling factor in the shape transformation of polymersomes into non-spherical morphologies.^{1,43}

As the ionic strength of the dialysis medium will have an effect on the charge repulsion in the polymersome membrane, we set out to systematically vary the salt concentration during the dialysis process upon the shape transformations of the A- and CA-modified polymersome variants (Fig. 3). The use of 10 mM sodium chloride solution resulted in small bi-layered spheres for A-doped assemblies, but elongated tubes for CA-doped particles. Raising the salt concentration of the solution to 25 mM NaCl resulted in the formation of large peanut-like shapes for assemblies with amine surface, often shorter than the ones obtained with 50 mM NaCl. However, when the same concentration of salt was used for the CA-modified polymersomes, this yielded elongated tubes together with a small population of bowl-shaped structures (stomatocytes). At 100 mM NaCl, mainly stomatocytes were formed from the A-doped polymersomes. Stomatocytes were also seen for the polymersomes with CA surface, in equal amounts with the elongated tubes. These findings highlight the complex nature of polymersomal shape transformations, where oblate and

Fig. 2 Cryo-TEM images of polymersomes with or without shape transformation. No shape transformation was observed when dialysing polymersomes against water (A = PEG-*b*-PDLLA/B = 10 wt% NH₂-PEG-*b*-PDLLA/C = 10 wt% CO₂H-PEG-*b*-PDLLA). Shape transformation was observed when dialysing polymersomes against 50 mM NaCl (D = PEG-*b*-PDLLA/E = 10 wt% NH₂-PEG-*b*-PDLLA) or 10 mM NaCl (F = 10 wt% CO₂H-PEG-*b*-PDLLA). All scale bars = 0.2 μ m.

Fig. 3 Cryo-TEM images of modified polymersomes after shape transformation using different salt concentrations. (–) 10 wt% CO₂H-PEG-*b*-PDLLA and (+) 10 wt% NH₂-PEG-*b*-PDLLA based polymersomes. All scale bars = 0.2 μm.

prolate morphologies can be formed through controlled osmotic deflation and are significantly influenced by salt concentration, composition and the molecular characteristics of the bilayer.⁴³ Based on these results, 10 mM and 50 mM NaCl were chosen for dialysis of CA- or A-modified and unmodified polymersomes, respectively, in order to controllably obtain tubular morphologies.

Further physical characterisation of polymersomes and tubes was performed using DLS (Fig. 4). The average particle size was, in agreement with the cryo-TEM images, in the region of 400–500 nm for unmodified and A-modified poly-

Fig. 4 Physical data for polymersomal formulations. (A) Tabulated results of dynamic light scattering (DLS) measurements and (B) intensity distribution data ($n = 3$ for each sample, standard deviation reported as $\pm x$ in A).

mersomes, whereas their CA-modified counterparts were smaller at around 200–300 nm. It should be noted that DLS is not best suited for non-spherical particles, as it is based on the Stokes–Einstein relationship, and this consideration should thus be taken into account during the analysis of the tubes.⁴⁴ All particles possessed a negative zeta-potential due to the negative character of poly(ethylene glycol)-rich membranes in solution, which was diminished in case of the amine-containing polymersomes.^{45–47} No significant difference in zeta-potential was observed between unmodified and CA-modified polymersomes. Although the zeta-potential values did not yield major differences between the different polymersomal formulations in terms of surface charge, the presence of carboxylic acid and primary amine groups at the surface of polymersomes could still induce different interactions in a biological context.

To evaluate the suitability of the differently shaped and charged particles for application in biology, we investigated their cytotoxicity. In keeping with our assertions regarding the biodegradable, and therefore biocompatible nature of PEG-*b*-PDLLA block copolymers, there was no evident toxicity towards human retinal (ARPE-19) cells at concentrations up to 1.25 mg mL⁻¹ (with 24 h incubation), which was very high compared to other polymeric systems as presented in the literature where studies are usually limited to 0.5 mg mL⁻¹ due to onset of toxicity (Fig. 5).^{48–51} Taken together, these results highlight the suitability of this platform for further studies in biomedical

Fig. 5 Toxicity studies of spherical and tubular polymersomes towards ARPE-19 cells after 24 h incubation (viability determined using AlamarBlue assays).

research to investigate the effect of both charge and shape on biological performance.

Conclusion

In this work, we have explored the influence of formulation factors, such as salt concentration, upon the assembly of PEG-*b*-PDLLA copolymers bearing 10 wt% of either carboxylic acid or amine-modified derivatives. Using optimised conditions we have generated tubular and spherical polymersomes using copolymer blends and characterised their morphology using both cryo-TEM and DLS. Furthermore, neither spherical nor tubular polymersomes (with different surface characteristics) showed any toxicity to cells up to concentrations of 1.25 mg mL⁻¹. This platform constitutes a robust technology that can be readily tuned in terms of morphology and composition to suit future applications in, for example, therapeutic drug delivery.

Abbreviations

A	Amine group
ARPE-19	Retinal pigmented epithelium cell line 19
BCPs	Block copolymers
CA	Carboxylic acid group
CryoTEM	Cryogenic transmission electron microscopy
DLS	Dynamic light scattering
PEG- <i>b</i> -PDLLA	Poly(ethylene glycol)- <i>block</i> -poly(D,L-lactide)

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors would like to thank Imke Welzen-Pijpers and Alexander Mason for CryoTEM measurements. We thank the European Union's Horizon 2020 research and innovation programme Marie Skłodowska-Curie Innovative Training Networks Nanomed (no. 676137) for funding. We thank the Ser Cymru II programme for support of DSW; this project received funding from the European Union's Horizon 2020 research and innovation programme under the Marie Skłodowska-Curie grant agreement no. 663830.

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