Macromolecular design and preparation of polymersomes

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From drug delivery to nanoreactors and protocells, polymersomes have gained considerable interest from researchers due to their novel applications. However, one of the main challenges is the selection of the most appropriate synthetic route. It is crucial to consider factors such as desired application, functionalisation, and environment of the polymersome when designing a synthetic route. This review will explore the current scope of polymersome synthesis and preparation methods, and will conclude by highlighting the most recent reports on polymersome related systems such as virus-like nanoparticles. From the choice of monomers, to polymerisation techniques, and to preparation methods used, we aim for this review to be utilised as a tutorial guide for the synthesis of a range of polymersomes, each with varying characteristics and applications.

1. Introduction

Compartmentalisation is crucial for life to occur.\(^1\) Almost all cells, be it eukaryotic or prokaryotic, consist of an outer protective vesicle containing the vital organelles needed to carry out a specific role in the body. Compartmentalisation provides a secure environment to facilitate essential biological processes such as DNA replication and aerobic respiration.\(^2\) Billions of years of evolution have allowed eukaryotic and prokaryotic cells to perfect the art of cellular compartmentalisation. As a result, such systems have a highly complex and specialised structure, with each organelle existing as its own entity within the cell, e.g. nucleus, mitochondria, lysosomes \( \text{etc} \). In 1960s, artificial vesicular cell mimics consisting of a phospholipid bilayer, known as liposomes (Scheme 1), were first discovered.\(^3,4\) These vesicles contain an aqueous solution core and can be used as a mode of transport for small hydrophilic/hydrophobic molecules. The liposome phospholipid bilayer can fuse with cell membranes and subsequently release its aqueous payload into a cell. This discovery was promising as new methods were being developed for the delivery of drugs and other biological molecules to cells within the body. However, liposomes are prone to stability and permeability issues.\(^5\)

In 1999, the term polymersome was first used to describe a vesicle formed from the self-assembly of an amphiphilic block copolymer by the Discher group.\(^6\) However, polymer vesicles have been known for some time. With pioneering work being conducted by the Eisenberg\(^7\) and Meijer groups\(^8\) in 1995. Polymersomes are similar to liposomes in terms of function and morphology but differ in chemistry. The vesicle bilayer is formed from hydrophilic and hydrophobic blocks and encapsulates an aqueous solution core like liposomes. However, by using synthetic monomer building blocks, characteristics such as stability, biodegradability and type of release mechanism

Scheme 1 2D and 3D representations of a liposome and a polymersome made from their respective subunits.\(^9\) Adapted from E. Rideau, R. Dimova, P. Schwille, F. R. Wurm and K. Landfester, Chem. Soc. Rev., 2018, 47, 40.
can be controlled.\textsuperscript{10--12} Furthermore, surface modification of polymersomes with certain sugars/proteins can direct the vesicle towards specific cells and targeted drug delivery can be achieved.\textsuperscript{13} As mentioned previously, drug delivery is the most widely researched application of polymersomes.\textsuperscript{14} But polymersomes have been synthesised for use in gene delivery in biological applications as well as being used as nanoreactors in enzyme catalysis and cascade reactions.\textsuperscript{15} More recently, polymersomes have found significant importance in the establishment of cellular biomimetics. Here, polymersomes are utilised as artificial organelles and/or cells. Their functions as cellular biomimetics include the uptake and release of specific cargoes, mimicking living cellular function.\textsuperscript{16--18}

It is worth mentioning the theory behind the formation of a polymersome from its corresponding block copolymer. Essentially, the amphiphilic nature of block copolymers enables self-assembly of the linear chains into nanoparticles of varying morphologies. In almost all cases, a macroinitiator consisting of a pre-synthesised hydrophilic polymer chain is polymerised with a hydrophobic monomer. As the hydrophobic block increases in the length, the overall amphiphilicity of the block copolymer changes as does the morphology of the nanoparticle (Scheme 2). Micelles, worm-like, rod-like and vesicles are achieved with an increasing DP of the hydrophobic block. The relative volume fractions of the hydrophilic and hydrophobic block are described by the packing parameter \( P \).\textsuperscript{20}

Many key review articles focusing on the applications of smart polymersomes have been published in the past decade. Notably, those published by Meier and van Hest have done well to highlight the current research of the biological and biomedical applications.\textsuperscript{5,21} However, after conducting an extensive literature search, it was found that there is a lack of review articles highlighting current research in the synthesis and preparation of polymersomes. Therefore, we present a review that aims to provide an overview of the chemistry and processing of polymersomes. This review will introduce the choice and variety of monomers that can be used for the synthesis of degradable and non-degradable polymersomes. Next, the types of polymerisation and preparation techniques are discussed with detailed examples. The review will conclude by exploring other nanoparticle-like systems similar to polymersomes such as virus-like nanoparticles (VLNPs). From the combination of monomers to the polymerisation and preparation methods used (Scheme 3), this review will serve as a comprehensive guide in synthesising a variety of polymersomes for multiple applications.

\section{2. The chemistry of polymersomes}

\subsection{2.1 The choice of monomers}

The main characteristic that defines polymersomes is the amphiphilic block copolymer that forms the bilayer of the vesicle. This is achieved by incorporating, at a minimum, a hydrophilic homopolymer block and a hydrophobic homopolymer block. By controlling properties such as degrees of polymerisation (DP), composition and ratio of the hydrophilic to hydrophobic blocks, the desired vesicle morphology can be obtained.

In general, polymersomes can be split into two categories. Degradable and non-degradable polymersomes. The degree of degradability of a polymersome can be dictated by the choice of monomers used. Non-degradable polymersomes typically include conjugated/aromatic hydrocarbon blocks such as poly(ethylethylene) (PEE),\textsuperscript{6} poly(butadiene) (PBD)\textsuperscript{22} and poly(styrene) (PS)\textsuperscript{22--26} as well as poly(dimethysiloxane) (PDMS).\textsuperscript{27--29} These polymers form the hydrophobic block of the copolymers but lack any bonds susceptible to hydrolytic cleavage. As a result, polymersomes synthesised with any of these monomers are considered non-degradable and are not biocompatible. Research on non-biocompatible polymersomes for biomedical applications has largely been disregarded as biocompatible vesicles are needed.

Polymersomes made from water soluble monomers including poly(ethylene glycol) (PEG),\textsuperscript{30--34} poly(2-methoxazoline)
Minireview

Polymer Chemistry

Table 1  Examples of non-degradable and degradable polymersomes

| Polymers           | Degradable | Polymerisation technique | Preparation method | Encapsulated payload | Ref. |
|--------------------|------------|--------------------------|--------------------|----------------------|------|
| PEG-β-PEE          | No         | Anionic/ hydrogenation    | Film rehydration   | N/A                  | 6    |
| iPSt-β-PBD         | No         | Anionic                  | Self-assembly      | N/A                  | 22   |
| PST-β-PEG          | No         | RAFT                     | Phase inversion    | N/A                  | 23   |
| PST-β-NIPAM-co-SPO | No         | RAFT                     | Double emulsion (w/o/w) | N/A | 24   |
| PVBC-β-PS-PVBC     | No         | RAFT                     | Self-assembly      | N/A                  | 25   |
| PST-β-Dec          | No         | ATRP                     | Self-assembly      | N/A                  | 26   |
| PDMS-b-PMOXA       | No         | Cationic                 | Film rehydration   | Paclitaxel, carboxyfluorescein | 27-29 |
| pPEGMA-S-S-PLA     | Yes        | ATRP                     | Nanoprecipitation  | Doxorubicin          | 30   |
| pPEG-b-PLA         | Yes        | Ring opening             | Film rehydration   | BSA and ASNase       | 31   |
| PEG-b-PCL          | Yes        | Ring opening             | Film rehydration   | Quercetin            | 32   |
| PMOXA-b-PDMS-b-PMOXA| Yes      | Ring opening             | Film rehydration   | Neurotrophins and Curcumin | 33   |
| PLA-b-PEG-b-PLA    | Yes        | Ring opening             | Double emulsion (w/o/w) | Lisonopril and atorvastatin | 34   |
| PEG-b-PLeu-b-PLGA  | Yes        | Ring opening             | Film rehydration   | Doxorubicin          | 41   |
| PLGA-b-PPhe        | Yes        | Ring opening             | Phase inversion    | N/A                  | 42   |

Current research on degradable polymersomes has led to the production of a catalogue of polypeptide-based polymersomes. Advantages in using amino acid-based monomers include complete biocompatibility, almost zero toxicity and easier functionalisation due to the variety of amino acids functional groups in the polymer chain. The review by latrou et al. in 2018, describes the properties and characteristics of polypeptide-based polymersomes and provides an excellent summary of some recent examples of these polymersomes.

Examples of degradable and non-degradable copolymers are summarised in Table 1.

2.2 Polymerisation techniques

Living polymerisation techniques are characterised by the absence of chain termination and chain transfer reactions during polymerisation, as well as having a much larger rate of initiation compared to the rate of propagation. As a result, all chains grow at a near constant rate owing to very little distribution in the molecular weights of the polymer chains. Living polymerisation techniques are mainly employed in the synthesis of block copolymers as the lack of chain termination in these reactions allow polymer blocks to be synthesised in stages by the sequential addition of different monomers. Living polymerisation techniques such as ROP, RAFT and ATRP allow extensive control over parameters such as the composition and block lengths of the hydrophilic and hydrophobic chains. Seen as the assembly of block copolymers into polymersomes vesicles is entirely dictated by these parameters, living polymerisation is paramount in the synthesis of well-defined amphiphilic block copolymers. The advantages and challenges of each technique will be explored in this section.

2.2.1 Ring opening polymerisation. Polymers composed of cyclic monomers make up a large percentage of degradable polymersomes as they contain hydrolytically cleavable ester and carbonate links in the backbone of the polymer chain. ROP provides an easy and effective method of incorporating such functional groups and heteroatoms into the backbone. For the synthesis of polymersomes, cationic ring opening polymerisation (CROP) and anionic ring opening polymerisation (AROP) are the two main techniques used. With CROP being used mainly for oxazoline, epoxide and carbonate polymerisation whilst AROP is used for lactone and lactide polymerisation. Peptide-based polymersomes can also be synthesised by ROP but requires a specialised version of the polymerisation technique. This technique is known as N-carboxyanhydride (NCA) polymerisation. The α-amino acids NCA monomers, to be ring-opened, are synthesised from their respective α-amino acids derivatives via the Leuchs or fuchs-farthing reaction. Ring opening is initiated usually by nucleophilic attack of primary amines. Due to the variety of side-chain functionalities present in amino acids, the polypeptides produced are usually stimuli-responsive and undergo conformational change in response to a change in its environment.

In 2020, Meier et al. reported the synthesis of a new bio-compatible amphiphilic block copolymer poly(ethylene oxide)-block-poly(2-(3-ethylheptyl)-2-oxazoline) (PEO-b-PEHOx) via an optimised microwave-assisted CROP procedure. Meier and co-workers opted for a nosylated PEO macroinitiator as small molecule nosylates (Nos) show faster initiation and better reactivity compared to tosylates. The final optimised procedure used Nos-PEO in chlorobenzene at 140 °C as these conditions gave the lowest dispersity and highest conversion. The two different preparation methods used gave complex nanoparticle morphologies. Film rehydration of PEO-b-PEHOx gave multi-compartment micelles (MCMs) with increasing PEHOx block length and the solvent switch method gave the morphologies of pseudo-vesicles and yolk/shell nanoparticles at large PEHOx block lengths. In 2019, van Hest et al. reported the step-wise
ROP synthesis of poly (ethylene glycol)-b-poly(caprolactone-g-trimethylene carbonate) (PEG-b-PCL-g-PTMC) yielding ionic-
ly-induced nanoworms as asymmetric therapeutic car-
briers.51 This is a perfect example of sequential addition of
different monomers producing a well-defined triblock copoly-
mer. Each step produced an ever growing macroinitiator until
finally reacted with the final monomer, TMC-Q (Scheme 4).

The inclusion of the quaternary ammonium-TMC
(PTMC-Q) chain end allows an ioni
cally-induced morphology change from micelles to nanoworms by varying the ionic
strength of the solution with NaCl. The morphology change is
ultimately driven by the adjustment of the interchain inter-
avtions thus altering the packing parameter of the block copoly-
mer. These nanoworms can be loaded with chemotherapeu-
tics such as doxorubicin (DOX) and can be used for targeted
drug delivery. The cationic nature of the terpolymer facilitates
greater adhesion and membrane penetration with negatively
charged cancer cells.

As mentioned previously, ROP is an effective method in
synthesising block copolymers that incorporate heteroatoms in
the polymer backbone. In 2020, Wang et al. synthesised a dual
stimuli-responsive, polyorganophosphazene (POP) diblock
copolymer poly((mPEG-SSamino)(N,N-diisopropylethylene
diamino)phosphazene) (PPDP) via thermal ring opening poly-
merisation to encapsulate and achieve stimuli-responsive
release of DOX.52 The hydrophobic block of this polymer con-
sists of an entirely heteroatom backbone of nitrogen and phos-
phorus atoms. A PEG-S-S-NH2 macrominitiator was first syn-
thesised in a two-step process. Using this macroinitiator, the
thermal ring opening of Hexachlorocyclotriphosphazene (six
membered nitrogen and phosphorus cyclic ring) proceeds at
250 °C with catalytic amounts of AlCl3. The S–S disulphide
bond is redox-responsive whereas the diisopropyl-
ethylenediamino (DPA) moiety accounts for the pH-responsive-
ness. POPs have gathered a lot of attention recently for drug
delivery due to their excellent biocompatibility and degradabil-
ity into non-toxic nitrogen/phosphorus-based products.53,54

2.2.2 Reversible addition–fragmentation chain transfer
polymerisation (RAFT). Since it was first reported in 1998 by
Rizzardo et al.,55 RAFT polymerisation has been used exten-
sively in polymer chemistry for the synthesis of block copoly-
mers. This living radical polymerisation technique makes use
of a thiocarbonylthio chain transfer agent (CTA) to control the
molecular weight and dispersity of each polymer block. This is
particularly useful when trying to synthesise polymersomes via
Polymerisation-Induced Self-Assembly (detailed explanation
section 3.3). In most cases, increasing the length of the hydro-
phobic block changes the morphology of the assembled block
copolymer. Therefore, it is paramount to control the length of
the hydrophobic block to ensure the correct morphology is
obtained. The main advantage RAFT polymerisation offers is
its tolerance to various functional groups and wide tempera-
tures ranges as well as many solvents including water.56 High
monomer conversion and polymer purity are also character-
istic of RAFT polymerisation. However, RAFT is not completely
infallible. The CTA must be fine-tuned with respect to the
monomers beings used. The main challenge of RAFT is ensur-
the CTA is not only compatible with the first monomer,
but must be compatible with all sequential monomers of the
block copolymer thereafter (Scheme 5).57

Using RAFT dispersion polymerisation, Cheng et al. syn-
thesised a self-assembling diblock copolymer poly(ethylene
oxide)-b-poly(styrene) PEO-b-PSt which produces ultra-thick
membrane, surfactant-resistant polymersomes.58 A PEO-CTA
macrominitiator was used to polymerise the hydrophobic PSt
block. Methanol was chosen as the choice of solvent for the
reaction as it solubilises the PEO-CTA macrominitiator and
Styrene well but solubilises the PST block poorly. The poor
solubility of PST is the driving force for self-assembly. Low PST
block lengths gave micellar and worm-like morphologies
whereas larger block lengths gave the desired vesicle mor-
phology for polymersomes. The large hydrophobic PST block
lengths also gave the polymersomes ultra-thick membranes.
As a result, the resistance to surfactant solubilisation of the
polymersomes was greatly improved, reporting less than 7% leak-
age of the encapsulated cargo after 30 days in a surfactant
solution.

Mantovani and coworkers also used RAFT dispersion poly-
merisation to produce two well define amphiphilic block co-
polymers using a PEG acrylate macrominitiator and 2-(acryl-
oxy)ethyl-3-chloro-4-hydroxybenzoate (ACH) or 2-(3-chloro-4-

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Scheme 4  How molecular programming of the terpolymer achieves
the ioni
cally-induced morphology switch from micelles to nanoworms.
(a) the structure of the (PEG-b-PCL-g-PTMC) terpolymer and the mor-
phology change by increasing the ionic strength of the solution with
NaCl. (b) The electrostatic interaction and membrane insertion between
the terpolymer and cancer cells.54 Adapted from J. C. M. van Hest,
Small, 2019, 15, 8.
hydroxystirbenzamidoethyl acrylate (CHB). Parameters such as choice of solvent, CTA and temperature were carefully selected to ensure the ‘livingness’ of the growing polymer chains. The one-pot RAFT dispersion polymerisation method, developed by Perrier et al., achieved high monomer conversions of ≥94% with vesicle morphology being obtained at ACH and CHB hydrophobic block lengths of DP = 34 (f = 35% ± 10). The vesicles were loaded with terbinafine and cyanocobalamin to determine their potential as nanocarriers of hydrophilic and hydrophobic drugs.

Cortizo et al. synthesised a triblock copolymer poly(vinylbenzoate)-b-poly(ethylene glycol)-b-poly(vinyl benzoate) (PVBz-P-PEG-PBVBz) via RAFT polymerisation. A PEG macroinitiator was synthesised bearing two thiocarbonylthio functional groups at either end of the hydrophilic PEG block. Vinyl benzoate was then polymerised from both ends of the macroinitiator producing a BAB triblock copolymer as seen in Scheme 5. The formation of vesicles via self-assembly in THF/water solution was reported when the triblock copolymer had a hydrophilic PEG weight fraction f of f = 30–40%.

2.2.3 Atom transfer reversible polymerisation (ATRP). ATRP is one of the main living reversible deactivation radical polymerisation techniques used to synthesise amphiphilic block copolymers (along with RAFT). ATRP reactions mainly employ a transition metal halide catalyst, usually copper-based, to form carbon–carbon bonds between vinyl monomers, thus synthesising polymer chains. An alkyl halide initiator and usually an ammine-based ligand is also used in ATRP. Rapid activation/deactivation of the dormant/radical species enables an equal rate of propagation of all polymer chains. As a result, this sequence controlled radical polymerisation technique produces polymers with narrow molecular weight distributions. Other advantages of ATRP also include high tolerance to many functional groups, access to cheap and commercially available reagents as well as ease of use. At the end of the polymerisation reaction, the copper-containing compounds are still present in the reaction mixture. This poses a problem for commercial utilisation of ATRP technique. However, water treatment products such as CupriSorb by Seachem can be used to remove copper, if the reaction is carried out in water.

Theato et al. have developed a unique ‘breathable’ polymer-some from the triblock copolymer poly[(ethylenglycol)methyl ether]-b-poly(N,N-dimethylamino ethyl methacrylate-co-2,2,2-trifluoroethyl methacrylate)-b-poly(4-[4-methoxy-phenylazo]phenoxy methacrylate) (PEG-b-P(DMAEMA-co-TFEMA)-b-PMEPPMA) via step-wise ATRP. A PEGylated ATRP initiator was used to polymerise DMAEMA and TFEMA to form the first statistical polymer block. The resulting diblock was used as a macroinitiator to polymerise MEPPMA to form the triblock copolymer. The amphiphilic nature of the polymer allowed self-assembly into vesicles which exhibited triple stimuli responsive character to CO2, O2 and light (Scheme 6). The membrane permeability of the vesicle can be fine-tuned by changing a single stimulus or a combination of stimuli.

In 2020, Liu et al. reported the synthesis of novel cationic nanoparticles (cNPs) from the coself-assembly of poly(lactic-co-glycolic acid)-b-poly(2-(dimethylamino)ethyl methacrylate) (PLGA-b-PDMA) and poly(lactic-co-glycolic acid)-b-poly(ethylene glycol) (PLGA-b-PEG) block copolymers via ATRP. The use of a PLGA macroinitiator was employed in the homopolymerisation of the PDMA and PEG blocks separately. The two diblock copolymers coself-assemble to form cNPs that bind to cell-free DNA (cfDNA) within inflamed joints and inhibits rheumatoid arthritis. PDMA blocks provides the cationic nature of the polymeric nanoparticles which binds to the negatively charged DNA. Meanwhile, PEG blocks increase accumulation and the retention time of cNPs in the joints thus lowering the administration frequency.

Lee et al. used a disulphide bridge-containing bifunctional initiator to produce a triblock copolymer poly(oligo(ethylene glycol)methyl ether methacrylate)-b-poly(2-(diisopropyl amino)ethyl methacrylate)-b-poly(2(methacryloyloxy)ethyl phosphoryl-
choline) POEGMA-b-PDPA-b-PMPC for DOX encapsulation and release.66 The POEGMA, PDPA and PMPC blocks were synthesised by polymerisation of the respective monomers to full conversion then sequential addition of the next monomer in the sequence. The pH-sensitive nature of PDPA allows pH-responsive release of DOX.

2.3 Block copolymer architectures

To further add diversity to the synthetic route, a vast choice of block copolymer architectures has been employed in the formation of polymersomes. To recap, the main feature of a polymersome is the amphiphilicity of the bilayer membrane. The copolymer assemblies so that the hydrophilic blocks form the exterior and interior layers whilst the hydrophobic blocks form the middle layer. These well-defined hydrophilic and hydrophobic regions are utilised for the encapsulation and transportation of various hydrophilic and hydrophobic drugs. AB, ABA, BAB, ABC block copolymers are the main architectures that are synthesised in literature. With blocks A and C being hydrophilic and block B being hydrophobic. Other alternative architectures such as comb and dendrimersomes exist too but are less studied.

2.3.1 AB diblock copolymers. The most prevalent and well reported block copolymer architecture, the first polymersome described back in 1999 was a PEO-b-PEE diblock copolymer. Much of the early research in polymersomes were diblock copolymers and still to this day, over 20 years later, they are just as significant now as they were back then.67–69 Consisting of a hydrophilic block and a hydrophobic block, the copolymer assemblies so that hydrophobic blocks aggregate forming the middle of the membrane thus reducing there interfacial contact area with the aqueous solution. The hydrophobic blocks then form the exterior and interior layers of the membrane (Scheme 7).

2.3.2 ABA triblock copolymers. Interestingly, ABA triblock copolymers have two possible conformations when forming a vesicle bilayer. The first conformation, like AB diblock copolymers, is cylindrical where the hydrophobic interaction of the B blocks form the middle layer and the two adjoining A blocks form the exterior and interior layers.71–73 The second possible conformation forces the triblock copolymer to curve in on itself. The hydrophobic loop forms the middle layer of the membrane whereas the two hydrophilic blocks, again, form the exterior and interior layers.

2.3.3 BAB triblock copolymers. Similar to AB diblock copolymers, BAB block copolymers only have one possible conformation. Here, the two hydrophobic ends of the copolymer chain aggregate to form the middle layer like the previous example whilst the looped hydrophilic central block forms the exterior and interior layers.74 In the majority of BAB triblock copolymer synthesis methods, a bifunctional hydrophilic polymer macroinitiator is employed. The hydrophobic polymer blocks are then polymerised off both sides of the macroinitiator.25,75

2.3.4 ABC triblock copolymers. ABC block copolymers are possibly the most interesting and unique of all the block copolymers architectures due to the two differing hydrophilic blocks A and C. From Scheme 7, the confirmation of the block copolymer is cylindrical like the first conformation of ABA triblock copolymers. However, due to the difference in the properties of blocks A and C such as hydrophilicity, charge and molecular weight, asymmetric membranes are obtained.76,77 Usually the hydrophilic block with the larger block length forms the exterior layer whilst the other forms the interior layer. However, changing the environmental conditions such as temperature or pH can cause spontaneous rearrangement and even inversion of the hydrophilic blocks affecting the permeability of the membrane in systems where polyelectrolyte complexes are contained within the membrane.78,79 In non-polyelectrolyte complex systems, chemical and physical crosslinking is key in fine-tuning the permeability of the membrane.80 In general, the greater the degree of crosslinking within the membrane, the lower the rate of uptake/release of cargoes.81 As a result, crosslinking has a significant role in mimicking cellular biomimetics as well as finding uses in drug delivery systems and enzymanic nanoreactors.82,83

2.3.5 Other architectures. Post-polymerisation functionalisation of polymer chains with hydrophilic or hydrophobic polymer grafts (depending on the hydrophilic nature of the backbone) result in the formation of comb/bottle brush-like block copolymers. Recent examples of comb/bottle brush-like polymersomes have been synthesised for the transportation and delivery of genetic material. In 2020, Dong and coworkers synthesised comb-like polycarbonates with varying lengths of cationic branches.84 These nanoparticles transport siRNA via encapsulation within the cationic brushes. Xu and coworkers have synthesised heparin based comb-like copolymer for use as a gene delivery vector.85 Cationic side chains of poly(poly (ethylene glycol) methyl ether methacrylate) (PPEGEMA)-b-poly(dimethylaminoethyl methacrylate) PPEGEMA-b-PDMAEMA were grafted onto the negatively charged heparin backbone and successfully encapsulated the plasmid pRL-CMV with low cytotoxicity and high gene transfection efficiency.

Dendrimersomes are a unique class of synthetic vesicles with the term being coined by Percec in 2005.86 They are formed from the self-assembly of low generation (1st or 2nd) amphiphilic Janus dendrimers consisting of a hydrophilic and hydrophobic side (Scheme 8). Percec and co-workers are considered the pioneers of dendrimersomes and have contributed
immensely to the field with a number of seminal papers and reviews being published on them. In one of his more recent publications, Percec et al. have developed amphiphilic dendrimersomes that engulf bacteria (Pseudomonas aeruginosa) via endocytosis mimicking the role of macrophages of the body. The development of these synthetic protocols that are capable of carrying out complex biological processes in collaboration with living matter is another step closer to the development of fully synthetic and artificial cells.

3. Preparation of polymersomes

The next stage in the formation of polymersomes is choosing which preparation method to use. Many have been reported over the years but essentially, they can be classified into two distinct categories, solvent-free and solvent displacement. Solvent free methods include film rehydration, PISA and electroformation where the amphiphilic block copolymers are formed in aqueous solution. Solution displacement methods such as direct injection, emulsion phase transfer and microfluidics involve the use of organic solvent to dissolve the block copolymers. The addition of this solvent-polymer solution in an aqueous solution and subsequent removal of the organic solvent produces polymersomes. Properties such as hydrodynamic radius and size distribution can be controlled using these methods. In recent times, it has been found that the shape of polymersomes can also be controlled very effectively. Polymersomes were long thought of as spherical entities but in the 2019 review by Stenzel and co-workers, they highlight many different examples of non-spherical polymersomes, including preparation methods and applications.

3.1 Rehydration

Rehydration of thin polymer films is the most common preparation technique used due to it being an inexpensive process that is relatively easy to perform. A sample of a block copolymer is dissolved in an organic solvent. The selected solvent must solvate the hydrophilic and hydrophobic blocks of the copolymer well. This polymer-solvent solution is then placed on a solid surface, such as a metal or glass plate, and the solvent is evaporated. Remaining on the surface is a thin dehydrated polymer film. Upon the addition of an aqueous solution, water molecules permeate through small defects in the polymer film. The accumulation of aqueous solution causes bulging and eventual separation of the vesicles from the polymer film is achieved (Scheme 9). This technique produces a fairly broad distribution of vesicle sizes. To narrow the distribution, vesicles are extruded through a filter with set pore diameters.

Direct hydration using low molecular weight PEG can also be a very effective polymersome preparation method. In 2015, Sui et al. reported a modified direct hydration method to produce PEG-b-PCL polymersomes. The PEG-b-PCL block copolymer with an appropriate PEG block ratio was dissolved in PEG 550 at a relatively high concentration. The addition of water to this solution drove the rapid formation of highly concentrated polymersomes. This method yields polymersomes of high quality, sufficient size and good control over membrane thickness. Moreover, this method does not involve the use of organic solvent or any agitation methods that could otherwise effect the quality of the polymersomes.

3.2 Electroformation

Electroformation is another solvent-free preparation method and can be considered as an aided-film rehydration method. Here, block copolymers are deposited onto electrodes (platinum, gold or indium-titanium oxide (ITO) glass) via the same process as film rehydration (Scheme 9). By applying an alternating current to the electrodes, the rate of water diffusion across the polymer film can be controlled. As a result, the degree of bulging and separation of the vesicles can be controlled to give a narrow distribution of polymersome sizes.

3.3 Polymerisation-induced self-assembly (PISA)

One of the more recent methods of preparation that has been extensively used in research is polymerisation-induced self-assembly. This method exploits the differences between the solubility of the monomer and polymer in solution. Like the example in Scheme 2, as a hydrophobic monomer is polymerised from a hydrophilic chain macroinitiator, the resulting
block copolymer becomes increasingly hydrophobic. The amphiphile will self-assemble so that the interfacial tension between the water and hydrophobic block is minimised. Recently, more information regarding the proposed mechanism of self-assembly has been published by Ianaro et al. in 2019. It was found that liquid–liquid phase separation precedes self-assembly of the amphiphile. Liquid–liquid phase separation is responsible for determining the size, membrane thickness and other structural properties of polymersomes. The morphology of the nanoparticle is related to the packing parameter $P$ of the amphiphile which, in turn, is related to the DP of the hydrophobic block. Initially, self-assembly does not occur at very low DP. However, the DP of the hydrophobic block will reach a critical micelle concentration (CMC) upon which micelles start to form. Worm-like, rod-like and vesicle morphologies are achieved at higher DPs. Therefore, a specific morphology can be obtained by fine-tuning the weight fractions of the hydrophilic and hydrophobic blocks or controlling the DP of the hydrophobic block. PISA is a popular method in producing nanoreactors with selective membrane permeability as the permeability is dictated by the $f$ ratios of the differing blocks. PISA has become one of the most popular preparation methods today as polymersome assembly occurs in situ with no post polymerisation processing needed to be done, thus saving time and resources. In addition to PISA, self-assembly of block copolymers into polymersomes can be driven by a change in stimuli such as pH, temperature etc. In 2018, Battaglia et al. reported the self-assembly of amphiphilic copolymers poly(2-(methacryloyloxy)ethyl phosphorylcholine)-b-poly(2-(diisopropylamino)ethyl methacrylate) (PMPC-b-PDPA) into polymersomes driven by a change in pH or temperature. It was found that an increase in temperature (from 5°C–60°C) and pH (from pH 2.0–7.0+) leads to an increase in deprotonation of the PDPA block. Deprotonation leads to an increase in hydrophobicity of the block and therefore the amphiphile self-assembles to form a vesicle. Temperature and pH can then be varied to fine-tune the morphology of the polymersome.

### 3.4 Direct injection

The most common of the solvent displacement methods, direct injection exploits the differences in solubility of the polymer in organic solvent and in aqueous solution. Like film rehydration, direct injection is an easy and inexpensive process to perform hence its popularity in literature. An organic solvent which solvates all blocks of the amphiphile is selected to disolve the block copolymer. Next, the polymer-solvent solution is injected into an aqueous solution where the hydrophobic block becomes insoluble due to an increase of interfacial tension between the block and surrounding water molecules. This interaction triggers the self-assembly of vesicles (Scheme 10). The reverse process can also produce polymersomes where water is injected into the polymer-solvent solution to trigger self-assembly. Direct injection can produce a broad distribution of vesicle sizes. However, the size distribution can be homogenised by extrusion as well as changing the solvent and concentration of solvent too.

### 3.5 Emulsion phase transfer

Emulsion phase transfer takes the direct injection method and adds more control, producing a homogenous sample of uniform vesicle sizes. As highlighted in Scheme 11, an emulsion of water-in-oil droplets are produced by vigorous agitation of a small amount of aqueous solution in a solution of polymer in organic solvent. In a separate vessel, an oil-in water biphasic system is set up, with a layer of amphiphiles sitting on the oil/water interface. The emulsion is then poured into the biphasic system. Due to density differences between the oil and water, the oil-coated water droplets sink to the aqueous phase and are coated in another layer of amphiphiles. The final product is a water-in-oil-in-water (w/o/w) double emulsion. The production of vesicles by w/o/w double emulsion allows the encapsulation of hydrophilic in the aqueous core and/or hydrophobic drugs in the oil layer.

### 3.6 Microfluidics

Emulsion phase transfer can be taken one step further to give almost complete control over the production and size distri-
bution of polymersomes. This method is known as microfluidics and is perhaps the most recent and up-to-date advancement in the polymersomes preparation field. From Scheme 12, a jet of organic solvent containing dissolved amphiphiles (middle fluid) is injected through the 1st junction into an aqueous solution (inner fluid) to produce amphiphile stabilised water-in-oil droplets. These droplets are then injected through a 2nd junction into an amphiphile-coated aqueous phase (outer fluid) to produce w/o/w double emulsion vesicles. By varying the flow rates through each junction, the size of the resulting polymersome can be controlled.

4. Beyond polymersomes

Since the first use of the term polymersome in 1999, research in this area of polymer chemistry has come a very long way since. Starting from the simplest diblock copolymer of PEO-b-PEE, we arrive at multi stimuli-responsive multiblock copolymers and complex architectures in today’s research. This rapid progression driven by intense research in the field has led many academic groups to explore alternative systems related to polymersomes with the same applications and goals in mind. In recent years, researchers have been inspired by some of the oldest organisms in the world to achieve targeted drug delivery. Viruses have long been regarded as excellent vectors for intracellular delivery due to their size and ability to change morphology to traverse the body of its host on a cellular level. By designing polymeric systems that mimic viruses, researchers have managed to synthesise a variety of highly specialised and efficient nanoparticles for intracellular delivery of many cargoes.

Armes et al. have recently reported a Dengue virus-mimicking, pH-responsive, framboidal ABC triblock copolymer vesicles that target breast cancer cells. poly(glycerol monomethacrylate)-b-poly(hydroxypropyl methacrylate)-b-poly(2-(diisopropylamino)ethyl methacrylate) PGMA-b-PHPMA-b-PDPA vesicles were synthesised via RAFT emulsion polymerisation and were prepared by PISA (Scheme 13). A small fraction (3%) of the hydrophilic PGMA chains were functionalised with phosphorylcholine (PMPC). This targeting ligand for the SR-B1 receptor is crucial for intracellular uptake by triple negative breast cancer cells in which this receptor is overexpressed. The pH-responsive nature of the nanoparticles allows for release of a payload (protein, gene or drug) inside these cancer cells.

Wang et al. also reported the synthesis of a pH-responsive virus-like nanoparticle consisting of an acetylated dextran-based diblock copolymer of poly(methyl methacrylate)-b-lauryl methacrylate) (MMA-b-LMA). The MMA-b-LMA block copolymer mimics the capsid protein for endosomal escape and acetylated dextran resembles the viral core for encapsulating payloads. It was found that conjugating the tumour penetrating peptide TT1 to the virus-like nanoparticle increases its cellular uptake in colon carcinoma cells.

As well as synthetic polymeric systems, a variety of peptide-based virus-like nanoparticles have been reported. These nanoparticles take the protein structure of a virus and are functionalised to improve its physiological properties inside the body. Functionalisation strategies include PEGylation for increased biocompatibility and retention in the body. As well as sugar/peptide conjugation for specific receptor targeting. Viruses such as the physalis mottle, cowpea mosaic, tobacco...
mosaic and hepatitis B have all been studied as vectors for drug delivery and vaccinations.  

5. Conclusion and outlook

In this review, we have seen that polymersomes are diverse in synthesis and preparation as there is no standard synthetic route in their formation. The amphiphilic block copolymers are synthesised via one of three main living polymerisation techniques. ROP is used for the polymerisation of cyclic monomers whereas RAFT and ATRP polymerise vinyl monomers with the use of a hydrophilic or hydrophobic macronitiator with mono- or di-functionality. Depending on the chemistry of the amphiphiles, polymersomes can be prepared via solvent-free or solvent dissolution methods. Solvent-dissolution methods such as microfluidics offer finer control over polymersome size and size distribution. However, for quick and easy preparation of polymersomes with less emphasis on size uniformity, solvent-free methods followed by extrusion are perfectly viable. Evidently, the synthetic route is entirely dictated by factors such as the monomers used and the eventual application of the polymersome.

Drug delivery still remains the main application of polymersomes as both hydrophilic and hydrophobic cargoes can be encapsulated within the nanoparticle. Furthermore, stimuli-responsive release of the encapsulated cargoes can be achieved by incorporating stimuli-responsive polymer blocks within the amphiphile. However, there is growing interest in utilising polymersomes as nanoreactors for biological cascade reactions where compartmentalisation is key for these biological processes to occur. Research interest has also grown in synthesising virus-like nanoparticles as vectors for intracellular cargo delivery too. These specialised nanoparticles have the ability to change morphology to pass through cellular membranes and barriers. Sugars and/or peptides can be conjugated to the virus head for targeted delivery to a desired destination.

Considering their significant contribution towards targeted drug delivery, why have we not seen polymersomes reach clinical trials or the commercial market yet? Unfortunately, there are a number of drawbacks that have made the use of these drug delivery vectors not viable yet. Firstly, and quite possibly most importantly, only a handful of polymers have been approved for use in pharmaceuticals by government health and safety agencies such as the MHRA and FDA. These include PEG, PLA, PCL, and PMMA. Any of the more unique, less reported stimuli-responsive polymers will need to be extensively researched, tested and approved by such agencies first. With only a limited number of building blocks to choose from, very few combinations of polymersomes can be produced for actual clinical use. The preparation of polymersomes also presents a few challenges holding back polymersomes for commercial use. During the preparation, any presence of residual cytotoxic organic solvent or monomers in the polymersomes renders them non-biocompatible regardless whether the amphiphiles are biocompatible themselves. Poor drug loading efficiencies, lack of absolute control over the targeting and release of drugs and the issue of scalability for production on a commercial level presents further problems to be addressed. For polymersomes to be incorporated in pharmaceuticals in the future, we believe cytotoxicity, drug loading efficiencies and long-term stability of polymersomes must be extensively researched and understood. Furthermore, new methods to allow the facile and cost-effective production of polymersomes on an industrial scale must also be explored.

Conflicts of interest

There are no conflicts to declare.

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