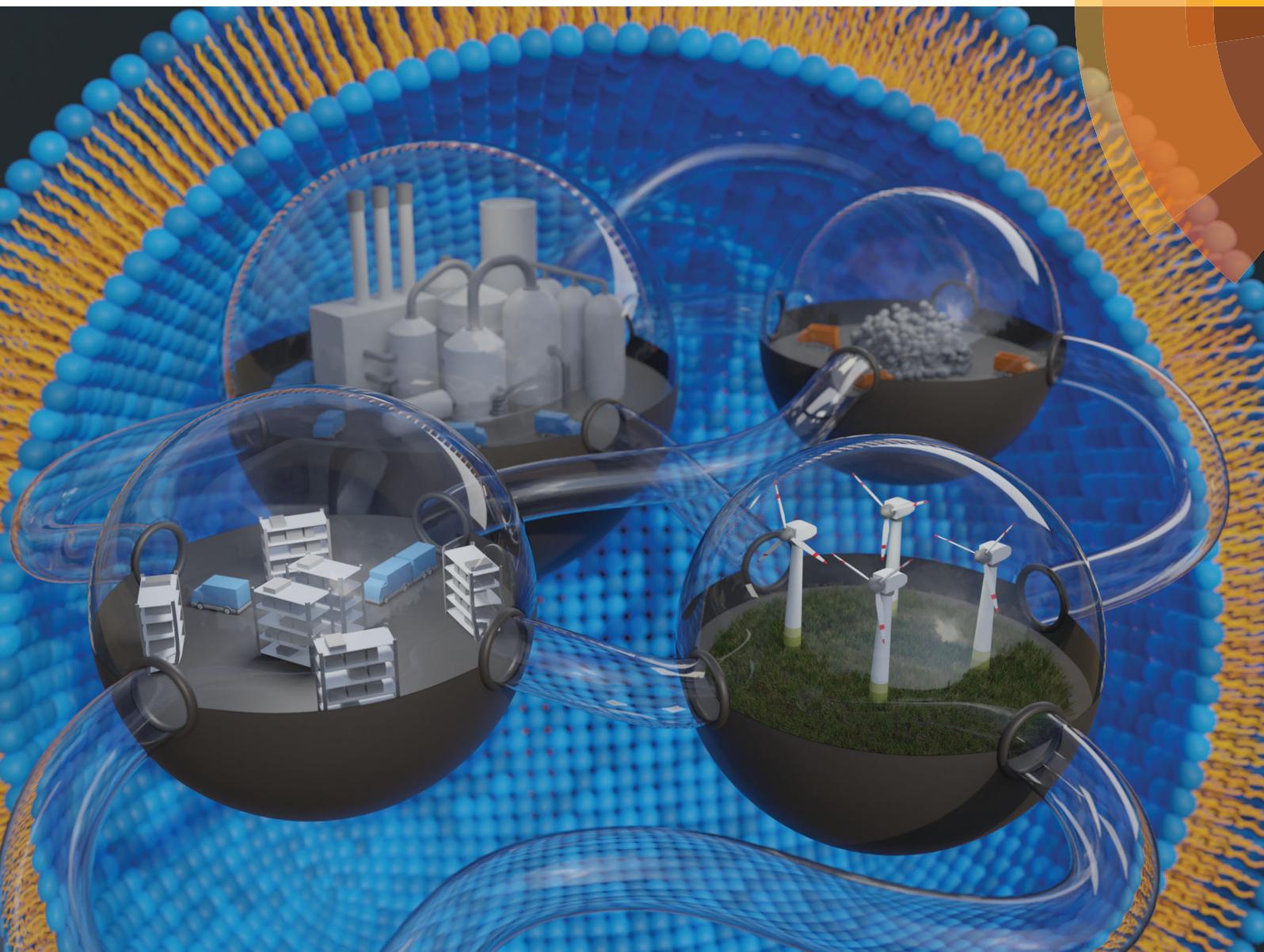


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Liposomes and polymersomes: a comparative review towards cell mimicking



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Liposomes and polymersomes: a comparative review towards cell mimicking

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Cells are integral to all forms of life due to their compartmentalization by the plasma membrane. However, living organisms are immensely complex. Thus there is a need for simplified and controllable models of life for a deeper understanding of fundamental biological processes and man-made applications. This is where the bottom-up approach of synthetic biology comes from: a stepwise assembly of biomimetic functionalities ultimately into a protocell. A fundamental feature of such an endeavor is the generation and control of model membranes such as liposomes and polymersomes. We compare and contrast liposomes and polymersomes for a better *a priori* choice and design of vesicles and try to understand the advantages and shortcomings associated with using one or the other in many different aspects (properties, synthesis, self-assembly, applications) and which aspects have been studied and developed with each type and update the current development in the field.

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1. Introduction

Cells, prokaryotic and eukaryotic, are undeniably integral to all forms of life and even essential for life apparition due to their compartmentalization.^{1–4} Within the compartments, sensitive entities like the genetic material, are protected, processes are regulated and can statistically occur due to concentration effects, as

is the case in protein synthesis. The driving force of reproduction and growth is made possible *via* cell division and energy is generated through the use of proton gradients between cell membranes using adenosine triphosphate (ATP)-synthase or between specialized organelles like the chloroplasts. Moreover, cells only exist because of the barrier that separates the intracellular components from the extracellular media: its membrane.

The fundamental importance of cell membranes has incited scientists not only to study them, but also to create synthetic analogs and models, such as bilayer structures based on natural or synthetic molecules ranging from lipids to surfactants and amphiphilic block copolymers.

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Emeline Rideau is currently a postdoctoral researcher under the supervision of Prof. Dr Katharina Landfester at the Max Planck Institute for Polymer Research, Germany. She obtained her BSc in biomolecular sciences from the University of St Andrews (UK) and her DPhil from the University of Oxford (UK) in organic chemistry in 2016 with Prof. Stephen Fletcher. Her current research interests include synthesis and self-assembly of block copolymers into polymersomes and hybrid vesicles as cellular mimics.



Rumiana Dimova

Rumiana Dimova obtained her PhD from Bordeaux University (France) in 1999. Afterwards she joined the Max Planck Institute of Colloids and Interfaces as a postdoctoral fellow, where she became a group leader in 2000. Since then she has been leading the Biophysics Lab in the Department of Theory and BioSystems. Her main research interests are in the field of membrane biophysics. In 2014, she was awarded the EPS Emmy Noether distinction for women in physics.



But why are synthetic cells necessary, when natural cells are so efficient, well-designed and a product of billions of years of evolution?

From the biological point of view, there has been a lot of effort to elucidate the inherent complexity of life, and biotechnology and bioengineering try to use this knowledge to improve man-made technology. However as life is so immensely complex, the usage and control of organisms in biotechnology are highly difficult.¹ Thus there is a need for simplified and controllable models of life both for a deeper understanding of fundamental biological processes and further man-made applications.

Natural cell membranes consist of a phospholipid bilayer, to which a cornucopia of proteins and small molecules (cholesterol, carbohydrates *etc.*) are bound.⁵ Hence it is not surprising that natural lipids (mostly phospholipids)^{6,7} or their synthetic alternatives can self-assemble into vesicles (spherical bilayer) on their own, termed liposomes (liposomes – from the Greek some = “body of”). Liposomes are a simplified model of a cell

membrane without its ornaments. Liposomes have been extremely well studied since the 1960s, but they are unstable and sensitive and have poor modular chemical functionality as we discuss in Section II of this review.⁸ Thus, despite lipids being highly biocompatible, there has been a great motivation in recent years to use versatile, easily obtainable and tunable entities, which behave in many aspects like lipids: amphiphilic polymers.^{1,9–11} The general physical properties of lipids and amphiphilic polymers are similar: they both are composed of a polar end (hydrophilic or charged head in lipids typically a phosphate moiety and 1 or 2 hydrophilic block(s) in polymers) covalently bound to a hydrophobic tail (1–2 aliphatic chains in lipids and 1 hydrophobic block in polymers).¹² Thus based on these facts amphiphilic polymers can also self-assemble into polymeric vesicles, or polymersomes (Fig. 1).

In this review, we will first explore the differences and similarities in physical properties between liposomes and polymersomes to try to understand their advantages and limitations (Section II). We will also briefly explain the different methods to obtain those vesicles (Section III) and discuss stimuli-responsive vesicles as primordial functionalization of those assemblies towards cell mimicking (Section IV). Finally, we summarize recent progress in creating artificial cells (Section V).

II. Comparison of physical properties

Despite their similar amphiphilic nature, liposomes and polymersomes exhibit important physical differences summarized in Table 1. Those properties have significant repercussion in the application of the vesicles notably from the perspective of generating synthetic protocells.

(a) Size

Phospholipids have a molecular weight of 100–1000 g mol⁻¹ whereas commonly used amphiphilic block copolymers typically have molecular weights M_n of >1000 g mol⁻¹ as polymers are



Petra Schulle

Her research interests range from single molecule biophysics to bottom-up synthetic biology of artificial cells.

Petra Schulle obtained her PhD in 1996 with Manfred Eigen at the MPI for Biophysical Chemistry in Göttingen. After a postdoctoral stay at Cornell University (Ithaca, NY) she established a research group at the MPI Göttingen in 1999 and accepted a call as professor and chair of biophysics at the BIOTEC of the TU Dresden in 2002. In 2011, she was appointed as a scientific member of the Max Planck Society and director at the MPI of Biochemistry, Martinsried.



Frederik R. Wurm

After a two-year stay at EPFL (CH) as a Humboldt fellow, he joined the department “Physical Chemistry of Polymers” at MPIP and finished his habilitation in Macromolecular Chemistry in 2016.

Frederik R. Wurm is currently heading the research group “Functional Polymers” at the Max Planck Institute for Polymer Research (MPIP). In his interdisciplinary research, Frederik designs materials with molecular-defined functions for degradable polymers, nanocarriers with controlled blood interactions, adhesives, and phosphorus flame-retardants. He has published more than 150 research articles to date. He received his PhD in 2009 (JGU



Katharina Landfester

creating functional colloids for new material and biomaterial applications.

Katharina Landfester received her doctoral degree in Physical Chemistry after working in 1995 at the MPI for Polymer Research (MPIP). After a postdoctoral stay at the Lehigh University (Bethlehem, PA), she worked at the MPI of Colloids and Interfaces in Golm leading the mini-emulsion group. From 2003 to 2008, she was a professor at the University of Ulm. She joined the Max Planck Society in 2008 as one of the directors of the MPIP. Her research focuses on



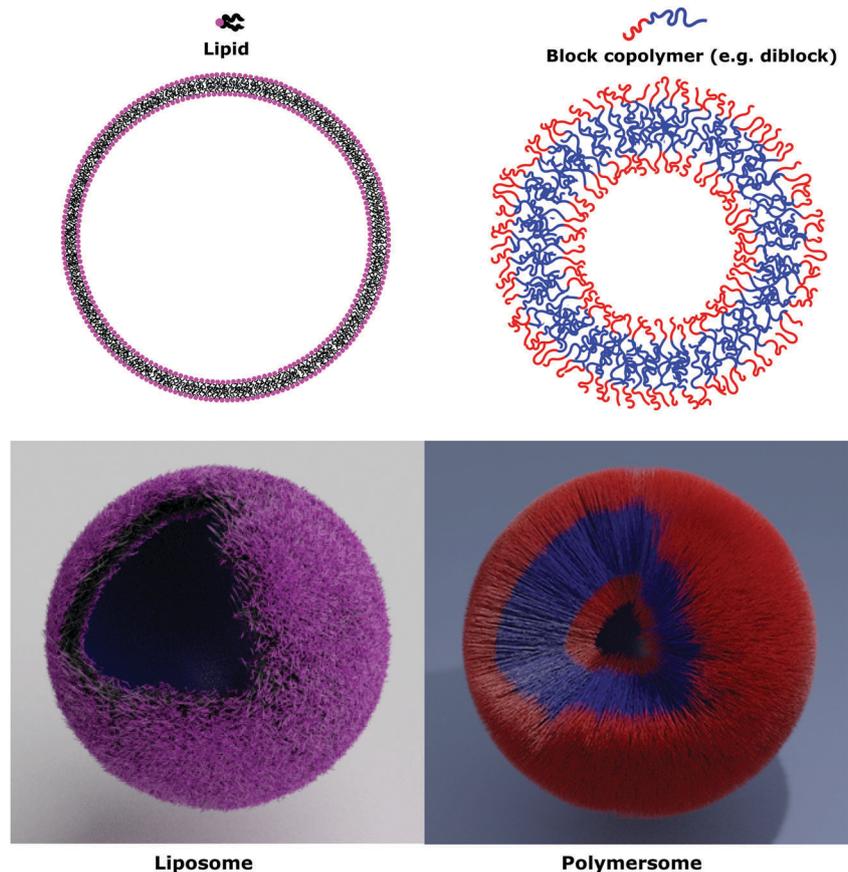


Fig. 1 Subunits, 2D and 3D structures of liposomes and polymersomes.

easier to synthesize controllably and in a low dispersity fashion than oligomers.¹² The molecular weight characteristics of the vesicular subunits result in the bilayer thickness of the self-assembled vesicles to be 3–5 nm for liposomes.^{2,12,13} For polymersomes, reported values are variable, but typically thicker, on average 5–50 nm.^{2,8,14} The membrane thickness of polymersomes was found to be dependent on the polymer M_n or degree of polymerization, especially for the hydrophobic block,^{2,11,12,15,16} hence explaining its much greater range. In comparison the cell membrane is 8–10 nm thick, including the lipid bilayer, membrane proteins and small molecules (cholesterol and carbohydrates).¹⁷ In this respect, the appropriately designed polymersomes compare better to natural cells than liposomes, despite the latter being chemically more similar to a cell. Liposome membranes are thinner than cell membranes due to the absence of membrane proteins.

In terms of diameter, liposomes and polymersomes can both form small vesicles (nm range) or giant vesicles ($>1 \mu\text{m}$) depending on the method used for preparation.¹⁵ The obtained vesicles are usually characterized and classified based on their diameter and lamellar properties (Fig. 2). These classifications are not well defined and vary between authors. The term small unilamellar vesicles (SUVs) is often used for vesicles of 20–100 nm diameter, large unilamellar vesicles (LUVs) for 100 nm–1 μm vesicles and giant unilamellar vesicles (GUVs) for vesicles of $>1 \mu\text{m}$ diameter.⁹ In this review we commonly refer to SUVs for

the smaller vesicles typically with diameters around 100 nm and GUVs for the macroscopic vesicles. In comparison, animal cells are typically 10–30 μm large and plant cells 10–100 μm also depending on their type and thus most model cells will focus on the formation and properties of GUVs. Even cell organelles like mitochondria are macroscopic (typically $1 \times 2 \mu\text{m}$)¹⁸ and though typically represented by SUVs for simplification, proper models should account for their large size. Sometimes vesicles are not unilamellar and can be referred to as multilamellar (MLV) (several bilayers) or multivesicular (MVV) (vesicles in vesicles also called vesosomes) and are usually larger in order to accommodate the volume of internal structures.^{9,19} MLV and UV, even of similar size, have very different physicochemical properties like permeability, stability, elasticity, toughness *etc.* which lead to different applications. MLV are particularly common in the development of drug delivery systems.⁹ Moreover, despite block copolymer membranes being thicker than lipid membranes, it has been shown that small-sized ion-channels can span a triblock copolymer¹ and grafted polymer membranes.¹³ These interesting results prove that polymeric membranes possess a high degree of adaptability to form thinner membrane rafts to incorporate the peptidic channel and thus are also biocompatible. We noticed that even though lipid-GUVs have been relatively well studied, reports on polymeric GUVs have only become increasingly more popular since 2012.



Table 1 Comparison of the measurements of liposomes, polymersomes and eukaryotic cells. SUV/LUV/GUV: small/large/giant unilamellar vesicles

	Liposomes	Polymersomes	Eukaryotic cells
Molecular weight of building compounds	10^2 – 10^3 g mol ⁻¹	10^3 – 10^4 g mol ⁻¹	10^2 – 10^5 g mol ⁻¹
Membrane thickness	3–5 nm	5–50 nm	8–10 nm (including membrane proteins)
Diameter	SUV: 20–100 nm LUV: 100–1000 nm GUV: 1–200 μm	SUV: 20–100 nm LUV: 100–1000 nm GUV: 1–200 μm	Animal cells: 10–30 μm Plant cells: 10–100 μm
Encapsulating volume (calculated)	SUV: 10^{-15} – 10^{-13} μL LUV: 10^{-13} – 10^{-9} μL GUV: 10^{-9} – 10^{-3} μL	SUV: 0 – 10^{-13} μL LUV: 0 – 10^{-9} μL GUV: 10^{-9} – 10^{-3} μL	Animal cells: 10^{-6} – 10^{-5} μL Plant cells: 10^{-6} – 10^{-3} μL

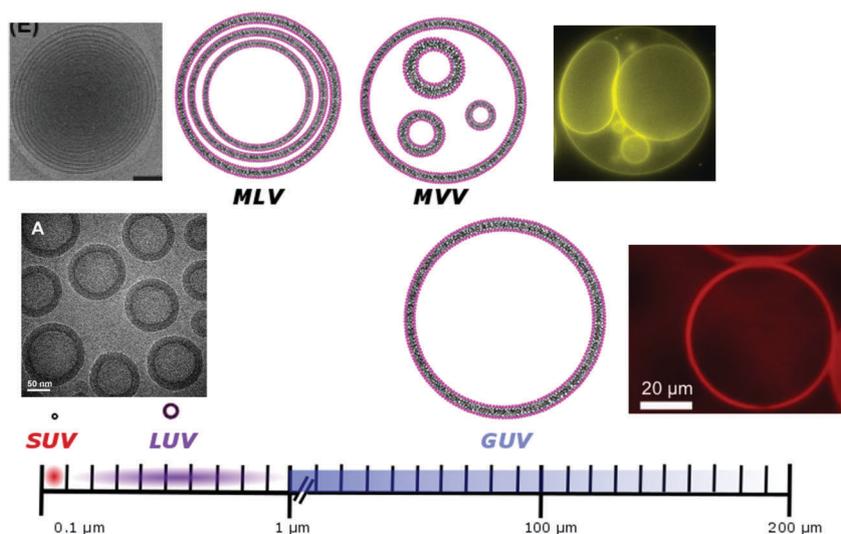


Fig. 2 Different classifications of possible vesicles. SUV/LUV/GUV: small/large/giant unilamellar vesicles; MLV: multilamellar vesicles; MVV: multivesicular vesicles. Microscopy images: SUV/LUV: Adapted from ref. 20 with permission from The Royal Society of Chemistry. GUV: Reprinted from J. Petit, I. Polenz, J. C. Baret, S. Herminghaus and O. Baumchen, *Eur. Phys. J. E: Soft Matter Biol. Phys.*, 2016, **39**, 59. <https://doi.org/10.1140/epje/i2016-16059-8.21>. MLV: reprinted from S. Allen, O. Osorio, Y.-G. Liu and E. Scott, Facile assembly and loading of theranostic polymersomes via multi-impingement flash nanoprecipitation, *J. Controlled Release*, **262**, 91–103. Copyright (2017), with permission from Elsevier.²² MVV: adapted from K. Oglęcka, J. Sanborn, A. N. Parikh and R. S. Kraut, *Front. Physiol.*, 2012, **3**, 120 (<https://www.frontiersin.org/articles/10.3389/fphys.2012.00120/full>).²³

Because membranes are essentially just a compartmentalization, the encapsulating volume of the vesicles is an important property to consider. Based on the membrane thickness and vesicular size, we calculated the estimated internal volume to determine how the differences in membrane thickness affect the encapsulation

efficiency and how these relate to eukaryotic cells (Table 1). For SUVs and LUVs, the encapsulated volume is small ($>10^{-9}$ μL) and the variation in internal volume for polymersomes is high and greatly dependent on the membrane thickness, whereas liposomes are much less affected as the membrane thickness is



insignificant compared to the vesicular size. For GUVs, in the case of both polymersomes and liposomes, the membrane thickness is negligible compared to the vesicle size and a well-designed synthetic vesicle could fit the volume of a eukaryotic cell ($\sim 10^{-4}$ μL) for protocell synthesis purposes.

(b) Permeability and stability

Vesicles, independently of their amphiphilic building blocks, have the significant advantage over other carriers to be able to encapsulate both hydrophilic and hydrophobic cargos within their lumen and membrane respectively (Fig. 3a). However, liposomes are always described as problematically leaky (Fig. 3b), resulting in poor retention efficiency of cargos.²⁴ Their chemical properties, size, saturation levels and method of preparation can modulate their permeability.^{6,9,25} The high permeability of liposomes is deemed a consequence of their high lateral fluidity linked to their low molecular weight and is discussed in more detail in Section Ic (Fig. 3c).^{12,26} The high dynamic mobility of membranes is also found in cells, making liposomes a good choice for artificial cell generation in this respect. In contrast, polymersomes retain their cargos with greater efficiency. Polymersomes are thermodynamically more inert as their lateral diffusivity is very low.^{11,12}

This property often depends on entanglement and varies with the overall M_n of the polymer and the nature of its hydrophobic block and is also determined by the glass-transition temperature (T_g) of the hydrophobic block.^{16,27,28} The polymersomes' low membrane permeability can also be problematic in certain applications (e.g. nanoreactors) as passive diffusion of small molecules through the membrane is low and does not accurately mimic cell fluidity.^{8,29} However, because of the versatility of polymer chemistry, advanced polymersomes can be functionalized for tunable permeability and lateral mobility and for example the grafted polymer PDMS-*g*-PEO generates vesicles of similar thickness (5 nm) and high fluidity to those of liposomes and cells.¹³

As a further consequence of the amphiphiles' chemical nature and lateral fluidity, liposomes are short-lived while polymersomes have a greater life-span (Fig. 3d). Stability can mean the chemical stability of the lipids or the physical stability of the vesicles (number, size, structure) and also is dependent on the storing conditions. Liposomes can be stable for months when handled appropriately and stored under an inert gas at low temperature.⁹ 1,2-Dipalmitoyl-*sn*-glycero-3-phosphocholine (DPPC) SUVs were reported to be stable for 48 h, phosphatidylethanolamine (PE)

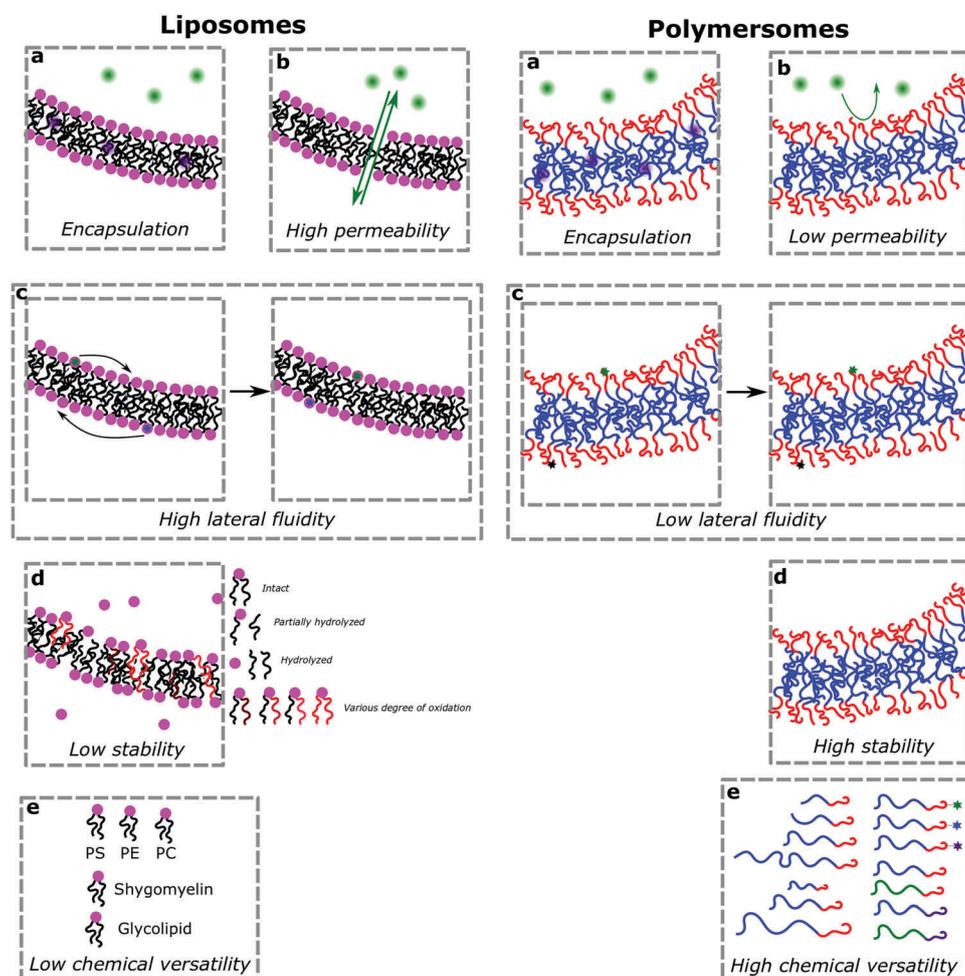
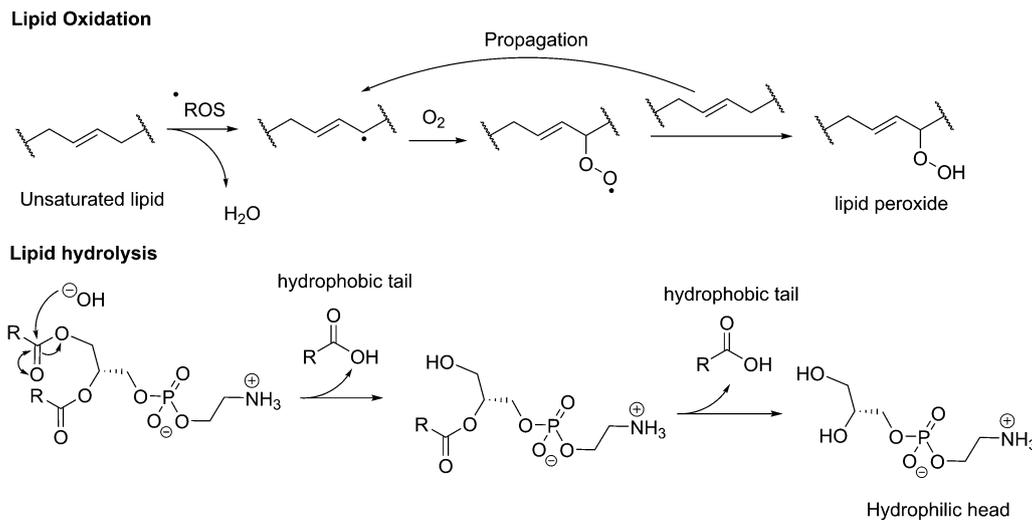


Fig. 3 Comparison of properties between liposomes (left) and polymersomes (right).





Scheme 1 Chemical instability of lipids due to oxidation or hydrolysis. ROS: reactive oxygen species.

SUVs for 12 days, and LUVs for weeks.⁶ The issue of liposome stability arises from the lipids' chemical instability rather than entropically favored disassembly.^{6,9,30} This is why the overall stability of liposomes is dependent on the nature of the lipids used. Their unsaturated fatty acid chains are prone to oxidation by reactive oxygen species (ROS) which causes changes in the overall liposome properties such as its permeability. In addition, the ester moieties can undergo hydrolysis, the covalent bond between the hydrophilic and hydrophobic fractions is cleaved, degrading the liposomes (Scheme 1). Oxidative degradation can be minimized by protecting lipids from light, keeping them under an inert atmosphere, at low temperature ($-20\text{ }^{\circ}\text{C}$), in a highly pure state (no heavy metals, free fatty acids or lyso-phosphatidylcholine (lyso-PC)) and sometimes stabilized by antioxidants while storing them.^{6,30} However, such measures are harder to maintain once they are self-assembled into liposomes in water or injected *in vivo*.

These chemical instability challenges are significantly improved when using block copolymers.^{1,24,28,31} Even though they are still prone to oxidative addition and potential cross-linking of the olefins (if present), the covalent bonds between the blocks are more difficult to break than the ester linkage used in phospholipids (*e.g.* ether bond in poly(butadiene)-*block*-poly(ethylene oxide) (PB-*b*-PEO)). Moreover, degradation of some of the repeating units of the polymers has a lower impact than on the low molecular weight lipids. Hence polymersomes have a greatly increased chemical stability lifetime compared to liposomes and were shown to be stable after at least one month.^{32,33} Even up to 6 months of stability at room temperature has been reported which is probably not the limit.^{34,35} Polymersomes are seemingly more stable than liposomes due to their membrane thickness, entanglement, and lateral diffusivity.² Despite the general trend that polymersomes are more stable and less permeable than liposomes, such properties are entirely dependent on many different aspects most importantly the size of the polymeric vesicles, the nature of the amphiphiles, the method of preparation, and storage conditions. The stability and permeability of polymersomes are consequently highly variable and

unfortunately actual values are rarely mentioned and assumptions on these properties should be used cautiously.

(c) Mechanical and rheological properties

Presumably, the most essential mechanical properties of polymer and lipid membranes are bending and stretching elasticity as they are related to the vesicle resistance to deformation and stretching experienced during drug delivery (resistance in the high shear rate of blood circulation and deformation through tiny vessels such as capillaries) or during cellular processes (*e.g.* division, fusion, osmotic shock). The membrane response to bending and stretching is characterized by the bending rigidity and the area compressibility or stretching elasticity modulus (the shear elastic modulus is zero for fluid membranes). Table 2 shows data about the mechanical, rheological and electrical properties of polymer and lipid membranes, which are easy to assess from measurements on GUVs.^{26,36–38} Depending on the membrane thickness, polymersome membranes can exhibit significant stiffness as demonstrated by the values of the bending rigidity modulus. For comparison, we have included data for lipid membranes in the gel phase as some of their properties appear to be close to those of polymersomes. Even though the bending rigidity of polymersomes might approach that of gel-phase lipid membranes, the response of the former to a deformation will differ significantly from that of a gel-phase vesicle because the latter is not fluid: gel-phase vesicles relax much faster after deformation induced *e.g.* by electric pulses.³⁹

Upon stretching, membrane tension builds up. When it reaches the so-called lysis tension, which is about $5\text{--}10\text{ mN m}^{-1}$ for phosphatidylcholine membranes, the membrane ruptures.^{40,41} The lysis tension of polymersomes is high, and exceeds 20 mN m^{-1} ,⁴² which is why they are referred to as tough vesicles. The membrane lysis tension is directly related to the critical transmembrane potential leading to poration of vesicles. The critical poration potential of gel-phase vesicles and polymersomes is several times higher compared to that of fluid membranes^{39,42} (one of the reasons being the larger membrane thickness). In fluid



Table 2 Typical values (order of magnitude) for the characteristic properties of polymersome bilayers and of pure lipid membranes in fluid and gel phases

Membrane material property	Polymer membranes ^a	Lipid membranes in the fluid phase	Lipid membranes in the gel phase
Bending rigidity ^b	35–400 $k_B T$	20 $k_B T$	350 $k_B T$
Stretching elasticity ^c	120 mN m ⁻¹ ; 470 mN m ⁻¹	200 mN m ⁻¹	850 mN m ⁻¹
Lysis tension ^f	20–30 mN m ⁻¹	5–10 mN m ⁻¹	> 15 mN m ⁻¹
Critical poration potential ^e	4–9 V	1 V	10 V
Pore edge tension ^g	10–50 pN	10–50 pN	—
Shear surface viscosity ^d	2 × 10 ⁻⁶ N s m ⁻¹	3–7 × 10 ⁻⁹ N s m ⁻¹	Diverges

^a The values reported for polymersomes depend on the diblock copolymer used (and the respective membrane thickness) and can vary strongly.

^b Data from ref. 38 and 43–49; the value for gel-phase lipid membranes corresponds to the bending rigidity of DMPC measured at around 5 degrees below the main phase transition temperature of the lipid. ^c Data from ref. 43, 46, 47 and 50. ^d Data from ref. 43, 44 and 51. ^e Data from ref. 39 and 42. ^f Data from ref. 41–43 and 52. ^g Data from ref. 53 and 54.

membranes, pores will reseal to avoid the contact of water with the hydrophobic part of the bilayer and the energy penalty for this is contained in the so-called pore edge tension. The resealing of pores in polymersomes is slowed down, because of the higher membrane viscosity,⁴³ and is arrested in gel-phase membranes³⁹ (upon poration, gel-phase GUVs develop cracks which do not reseal even within minutes as the membrane viscosity diverges⁴⁴). One study estimated the shear viscosity of polymersome membrane to be 500 times higher than that of fluid lipid membranes.⁴³ Consequently, molecular mobility in the polymersome membrane is much less compared to liposomes where diffusion of membrane probes is fast. The latter also leads to faster domain coarsening in phase separated fluid vesicles.

(d) Chemical versatility

Natural and synthetic lipids^{55,56} are already highly specialized molecules that suffer from a large limitation of the available chemical functionalization space (Fig. 3e). Their physical and chemical properties are relatively restricted as small modifications can lead to large repercussions in the properties of their small molecules and thus their self-assembly. The modification of the head of lipids is achievable but is restricted to hydrophilic moieties so that lipids still self-assemble into vesicles. The modification of the lipid tail(s) requires often several synthetic steps as shown for example for lipid **1** in Scheme 8 which is synthesized in 9 steps.⁵⁷

From Table 3, we observe that commonly used lipids forming liposomes are exclusively natural phospholipids and even sometimes used as a naturally occurring mixture such as soy-PC or hydro soy-PC. Their structures and sizes are fairly similar (molecular weight ~ 800 g mol⁻¹) and only vary in aliphatic chain lengths (13–17Cs long; 0–2 degree of saturation and always *cis*) and end groups (primary amine, tertiary amine, diol, or carboxylic acid) which determine the overall charge of the lipids as the phosphate diester moiety is always deprotonated at pH 7 ($pK_a < 7$). Despite their relative low chemical diversity, these lipids have widespread gel-liquid crystal transition temperatures (T_m) even among the same type of phospholipids (–22 to 60 °C). Most importantly, T_m can be higher than room temperature, a property that is significant for liposome self-assembly (see Section III). T_m depends on the nature of the amphiphiles, notably their saturation level, and represents the temperature at which the amphiphile's physical state changes from an ordered gel phase to a disordered liquid crystalline phase.²⁵ A few

groups synthesized non-natural lipids for their self-assembly into vesicles. For example, the group of Zumbühl synthesized modified diamidophospholipids, which self-assembled into interesting d-form vesicles or cuboid liposomes by using hydrogen bonding networks from well-designed lipids.^{58–61}

Block copolymers have the advantage of being adjustable and easily tunable to desired characteristics of the vesicles (Fig. 3e).^{2,28} Primarily, a large pool of blocks (Table 4 lists some typical examples) is available and their overall molecular weight M_n and block ratio can be controlled *via* classical ionic or controlled radical polymerization techniques.^{1,3,24} These designable properties allow for the formation of a variety of polymersomes with adjustable shape, fluidity, entanglement, permeability, stability, and responsiveness,^{11,12} and the polymers can be tailored to appropriate physical properties or biocompatibility.¹ Contrary to lipids, block copolymers are however polydispersed. Even if their distribution is most often low, thanks to modern polymerization, polymers' dispersity can affect their self-assembly and reproducibility can be challenging when using different batches of the same polymers.¹⁵ The most common polymer architecture used in forming polymersomes are di-(AB) or tri-block copolymers (ABA or rarely ABC) (where A (and C) is the hydrophilic block and B the hydrophobic block) (Table 4);¹⁶ however, polymersomes formed from branched, grafted comb or star polymers have also been reported,^{63–65} emphasizing the superior synthetic versatility of polymers over lipids (Fig. 4).

The amphiphilic composition is the driving force for the self-assembly of lipids and block copolymers into various structures.⁶⁶ As the hydrophilic and hydrophobic moieties are chemically incompatible, but covalently coupled, no microphase separation takes place.^{1,15} Their rearrangement into vesicles is entropically favored, as it reduces non-favored interactions in aqueous media by aggregating the hydrophobic groups together away from polar solvent. The hydrophobic and hydrophilic groups are said to be microphase-separated. However, while natural lipids have been designed to self-assemble only into vesicles, as block copolymers can be synthesized with a wide variety of volume ratios of the blocks, they can self-assemble into different morphologies (entropically favored microphase separations).¹⁵ The nature of the polymer and the hydrophilic/hydrophobic ratio are the major parameters which determine whether polymersomes (bilayer), spherical micelles (monolayer), cylindrical micelles, rods, lamellae (bilayer), hexagonal cylinders *etc.* are formed.²⁴



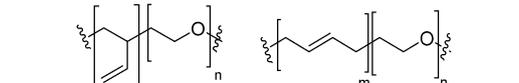
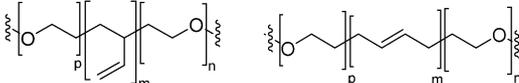
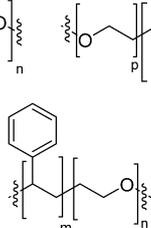
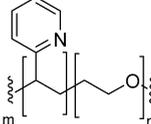
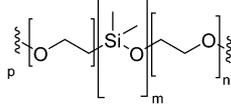
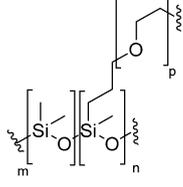
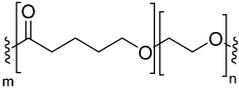
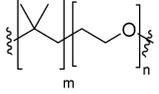
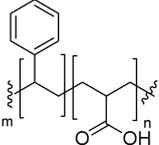
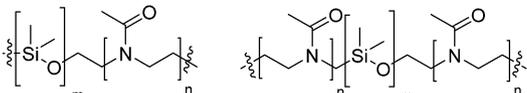
Table 3 List of typical lipids that can self-assemble into liposomes

Lipid	Structure	T_m ($^{\circ}\text{C}$)	Charge at pH 7 pK_a^{62}
Phosphatidylcholine (PC)			
DMPC		23	Zwitterionic ≤ 1 (P(O)(OR) ₂ OH)
DPPC		41	
DSPC		55	
	$n = 12$ (DM), 14 (DP), 16 (DS)		
DOPC		-22	Zwitterionic ≤ 1 (P(O)(OR) ₂ OH)
POPC		-2	
SOPC		6	
	$n = 7$ (DO), 5 (PO), 7 (SO)		
Soy PC (mixture)		-20	Zwitterionic ≤ 1 (P(O)(OR) ₂ OH)
	(63%)		
Hydro Soy PC (mixture)		52	Zwitterionic ≤ 1 (P(O)(OR) ₂ OH)
	(89%)		
Phosphatidylglycerol (PG)			
DMPG		23	Negative ~ 3.0 (P(O)(OR) ₂ OH)
DPPG		41	
DSPG		55	
	$n = 12$ (DM), 14 (DP), 16 (DS)		
DOPG		-18	Negative ~ 3 (P(O)(OR) ₂ OH) ~ 4 (COOH) ≥ 9.7 (NH ₃ ⁺)
Phosphatidylethanolamine (PE)			
DMPE		50	Zwitterionic ≤ 1.7 (P(O)(OR) ₂ OH) ≥ 9.6 (NH ₃ ⁺)
DPPE		60	
	$n = 12$ (DM), 14 (DP)		
Phosphatidylserine (PSE)			
DMPS		38	Negative ~ 3 (P(O)(OR) ₂ OH) ~ 4 (COOH) ≥ 9.7 (NH ₃ ⁺)
DPPS		51	
	$n = 12$ (DM), 14 (DP)		
Sphingomyelin (SM)			
Palmitoyl-SM		41	Zwitterionic ≤ 1 (P(O)(OR) ₂ OH) ~ 17 (OH)

T_m : gel-liquid crystal transition temperature. For lipid abbreviations see the table of abbreviations.



Table 4 List of typical copolymers used to form polymersomes

Copolymer	Structure	Charge at pH 7 pK_a
PB- <i>b</i> -PEO		
PEE- <i>b</i> -PEO		
PS- <i>b</i> -PEO		
P2VP- <i>b</i> -PEO		Neutral
PDMS- <i>b</i> -PEO		
PDMS- <i>g</i> -PEO		
PCL- <i>b</i> -PEO		
PIB- <i>b</i> -PEO		
PS- <i>b</i> -PAA		Negative pK_a PAA: 4.25
PDMS- <i>b</i> -PMOXA		Neutral

PB: poly(butadiene); PEO: poly(ethylene oxide); PEE: poly(ethyl ethylene); PS: polystyrene; P2VP: poly(2-vinylpyridine); PDMS: polydimethylsiloxane; PCL: polycaprolactone; PIB: polyisobutylene; PAA: polyacrylic acid; PMOXA: poly-2-methyl-2-oxazoline; b: block; g: grafted.

The microphase separation depends on 3 parameters: the volume fractions of the blocks (f), the degree of polymerization (N) and the Flory-Huggins parameter χ (degree of block incompatibility).⁶⁶ Theoretical phase diagrams between f and the segregation parameter χN using self-consistent mean-field

theory are used to determine which block ratios are more likely to yield the desired morphology. The critical packing parameter $P_c = v/(a \cdot l)$ is also used to predict the most likely morphology of the self-assembled aggregate (where v is the volume of the hydrophobic block, a is the area of the hydrophilic block and l



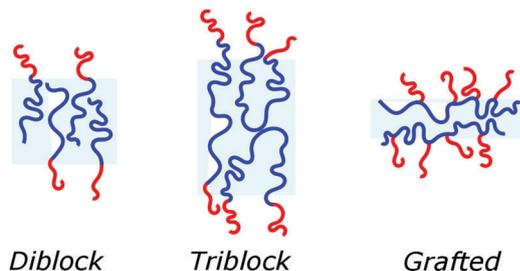


Fig. 4 Different types of block copolymer, which can self-assemble into polymersomes.

is the length of the block copolymer).^{16,24} For $P_c < 1/3$, spherical micelles are formed, for $1/3 < P_c < 1/2$, cylindrical micelles, $1/2 < P_c < 1$ vesicles, for $P_c = 1$, planar bilayers, and for $P_c > 1$, inverted structures. P_c was first established for surfactants and lipids; however, it is generally also accepted for amphiphilic block copolymers. However, determining these parameters is not trivial, especially as a block copolymer head group is more indefinite than those in surfactants or lipids. A less accurate but more straightforward prediction of micro-phase separation of the block copolymer pre-synthesis is the hydrophilic weight fraction $f_w = M_n(\text{hydrophilic block})/M_n(\text{polymer})$.^{16,24,35} Which aggregates are formed greatly depends on the nature of the blocks (hydrophilicity or hydrophobicity, sterics, degrees of freedom, polymer polydispersity, rigidity, etc.) but as a general rule $0.25 < f_w < 0.40$ yields polymersomes.¹² However, predicting self-assembly morphologies is highly complex as thermodynamics and kinetics properties can

lead to the same polymer generating different morphologies depending on the method used. For example, PEO₄₅-*b*-PMMA₅₁ (PMMA: poly(methyl methacrylate)) was shown to form micelles (29 nm in diameter) when using acetone injection in water and polymersomes (550 nm) by the addition of water/benzyl alcohol to the polymeric solution in THF/MeOH.⁶⁷ The determination of self-assembly morphologies by the block copolymer packing parameter has been well exemplified by polymerization-induced self-assembly (PISA) (see Section III Vesicle preparation).⁶⁸

(e) Mixed vesicles and phase separation

Both liposomes and polymersomes have their own advantages and shortcomings. However, the careful design of functional polymersomes or liposomes for customizing the vesicle properties can be synthetically demanding and challenging.

One alternative to obtain vesicles with the advantage of both polymersomes and liposomes is to form capsosomes (liposomes in a polymer shell) (Fig. 5).^{2,3,69,70} This multi-compartmentalized scaffold allows for the desired permeable membrane of the liposomes and the extra stability of the polymeric nanocarriers. These systems are interesting for cell mimicry as the cell uses organelles to further compartmentalize different functions. However, such systems are not trivial to generate and diverge from the simple modelling of a membrane (multiple bilayers rather than a single bilayer).

An increasingly promising alternative for membrane studies and for the pharmaceutical industry is generating lipid-polymer hybrid vesicles, also called lipopolymerosomes, HLP (hybrid liposome-polymerosome) or HUV (hybrid unilamellar vesicles).^{13,14,71,72}

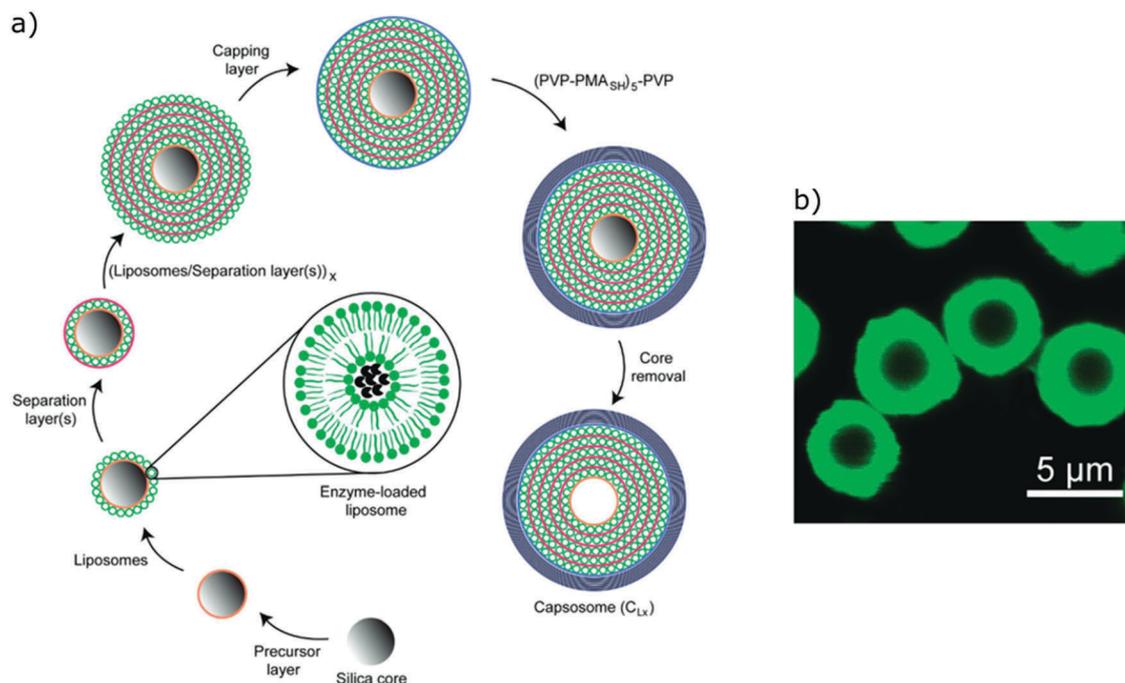


Fig. 5 (a) Schematic representation of capsosomes. (b) Confocal laser scanning microscopy of a capsosome with eight liposome deposition steps. Adapted with permission from R. Chandrawati, L. Hosta-Rigau, D. Vanderstraaten, S. A. Lokuliyana, B. Städler, F. Albericio and F. Caruso, *ACS Nano*, 2010, 4(3), 1351–1361 (<https://pubs.acs.org/doi/abs/10.1021/nn901843j>).⁶⁹



Here the membranes are formed from mixing both lipids and polymers. Hybrid vesicles can offer an easier platform to finely control the physical properties of vesicles with the best characteristic of both types (biocompatibility and biofunctionality and synthetic control of robustness, permeability and responsiveness).⁸ Indeed, a great variety of vesicles with little additional synthetic effort can be efficiently obtained. This variability can be achieved by playing with the nature and functionalization of lipids and polymers and their respective ratio resulting in a library of vesicles with many different properties. Despite the advantages and interests related to hybrid vesicles, only a few systems have been described and almost exclusively GUVs presumably due to the difficulty in distinguishing between phase separation on smaller vesicles, which is normally achieved using selective fluorescent tagging.^{11,73} A key parameter for the formation of stable hybrid vesicles is the compatibility between the block copolymer and the lipid (chemical structure, size, thermodynamic parameters *etc.*) and their ratio.⁸ At first glance, it seems surprising that such different amphiphiles can form hybrid vesicles as lipids form 3–5 nm thick membranes while block copolymers form on average thicker membranes (5–50 nm). Thus, hybrid vesicle self-assembly can suffer from geometrical incompatibility. Nonetheless, hybrid structures have been described for polymers of relatively high molecular weights, for

example with PC/PE/Lissamine rhodamine-PE (Liss Rhod-PE) and PMOXA-*b*-PDMS-*b*-PMOXA (PMOXA: poly(2-methyl-2-oxazoline); PDMS: polydimethylsiloxane) ($M_n = 9000 \text{ g mol}^{-1}$).⁷⁴

The main property to take into account when dealing with hybrid vesicles is their phase separation.^{8,11,14} Homogeneous vesicles (statistically distributed lipid and polymer), vesicles with domains (phase separation within the membrane) or complete phase separation of the lipids and the polymer (coexistence of liposomes and polymersomes) can form (Fig. 6). The nature, size difference, ratio of lipids and polymers and the method of formation determine which type of phase separation is obtained. If the size difference is large, the formation of domains would be entropically favored in order to minimize the exposure of hydrophobic moieties to water or to hydrophilic moieties.⁸ The demixing of polymer and lipids is particularly advantageous for studying the concept of lateral heterogeneity found in plasma membranes and further discussion can be found in Section IV.

III. Vesicle preparation

Over the years, many different preparation methods for liposomes have been developed, most of which have also been applied to the

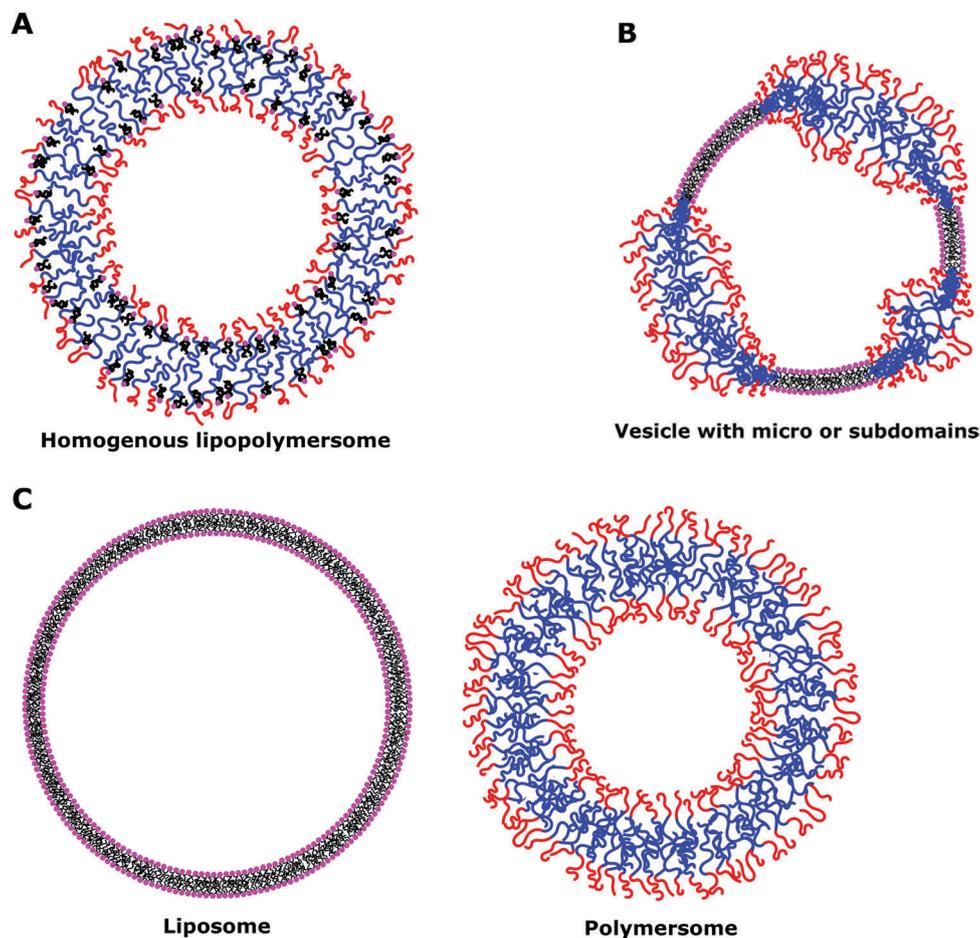


Fig. 6 Possible scenarios of phase separation of lipids and block copolymer vesicles.



Table 5 Preparation methods of lipids and block copolymer vesicles. SUV/GUV: small/giant unilamellar vesicles; MLV: multilamellar vesicles

Methods	Type	Size	Additives	Polydispersity	Ref.
Film rehydration	Solvent free	SUV, MLV	—	Large	6, 15, 76 and 80
Solid rehydration	Solvent free	SUV, MLV	—	Large	81 and 82
Electroformation	Solvent free	GUV	—	Low	46, 67, 77 and 83
Gel-assisted hydration	Solvent free	GUV	Agarose or PVA	Low	83–85
Solvent injection (ethanol, ether)	Solvent displacement	SUV	Solvent	Large	15 and 25
Emulsion phase transfer	Solvent displacement	GUV	Solvent, surfactant	Low	86–88
Microfluidics	Solvent displacement	SUV or GUV	Solvents, surfactant, sugars, polyelectrolytes, polymers <i>etc.</i>	Low	21 and 89–92

formation of polymersomes.^{2,15} Those methods dictate the size, yield, polydispersity, and type (unilamellar, multilamellar, vesosomes) of the resulting vesicles. In 2018, some of the mechanisms behind the formation of polymersomes were elucidated.⁷⁵ Conditions to self-assemble liposomes and polymersomes may vary due to the difference in stability, solubility, and stiffness of the lipid and polymer membranes. The nature of the amphiphile (chemical properties, molecular weight, polydispersity), additives (ions, small molecules, surfactants) or organic solvent used, concentration of the amphiphile, temperature, aqueous medium *etc.* can all affect the size, yield and polydispersity of the vesicles.^{15,76}

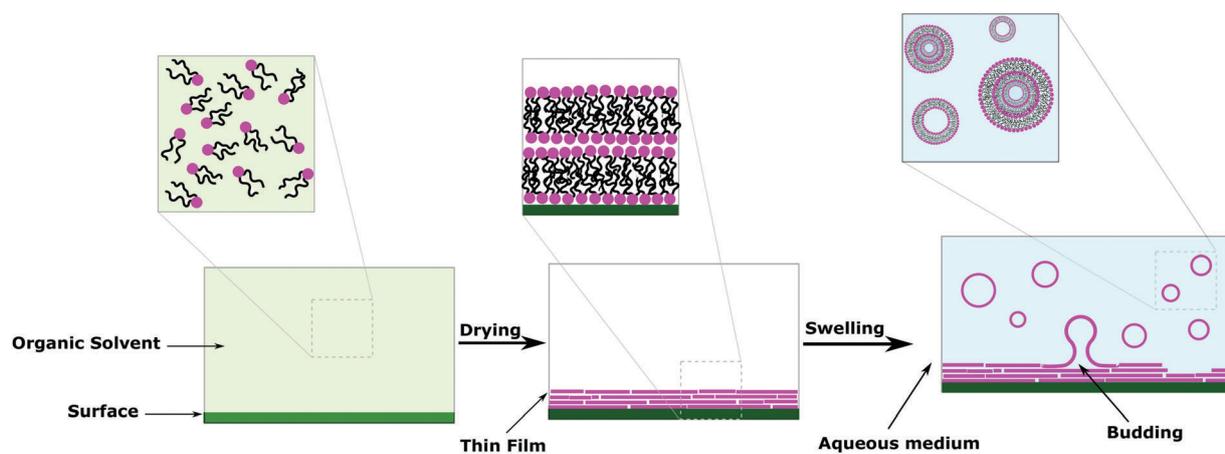
Another important aspect of liposome formation is that vesicles will only be obtained at temperatures above the T_m of the lipid used.^{6,25,77} Having to form liposomes above their T_m can be problematic when encapsulating temperature-sensitive entities (*e.g.* protein, DNA *etc.*).^{6,25,77} For block copolymers, the T_g and T_m are dependent on the degree of polymerization and polymeric architecture and thus much less defined than for lipids. In fact, the T_g of polystyrene (PS) can vary from -10 °C at a $M_n = 500$ g mol⁻¹ up to 100 °C at a $M_n > 10^4$ g mol⁻¹.⁷⁸ In terms of blocks, PB and PDMS have T_g s well below room temperature (PB(1,2) $T_g \approx -21$ °C ($M_n = 105$ k); PB(1,4) $T_g \approx -77$ °C ($M_n = 50$ k); PDMS $T_g \approx -127$ °C),⁷⁸ while polycaprolactone (PCL) and PEO have a low T_g but a T_m above room temperature (PEO, $T_g \approx -54$ °C and $T_m \approx 66$ °C ($M_n = 90$ k); PCL $T_g \approx -72$ °C and $T_m \approx 65$ °C ($M_n = 60$ k))⁷⁸ and others have

only transition temperatures above room temperature (PS ≈ 100 °C ($M_n = 1$ k); polyacrylic acid (PAA) ≈ 70 °C ($M_n = 1.8$ k));⁷⁸ PMOXA ≈ 55 °C ($M_n = 4$ k)⁷⁹). Nonetheless, block copolymer self-assembly is usually carried out at room temperature regardless of the T_g and of the block used. It remains unclear why formation of polymersomes at room temperature using high T_g or T_m blocks is possible in opposition to lipids.

Despite the variety of methods developed, they can be classified into two types: solvent free methods and solvent displacement ones (Table 5).^{6,15,76,80} In solvent free techniques, only the amphiphiles are hydrated in an aqueous medium and thus no organic solvents are present in the vesicle solution. In solvent displacement techniques, the amphiphiles are dissolved in an organic solvent and then placed in aqueous medium followed by removal of the organic solvent.

(a) Solvent free methods

Solvent free methods are all variations of film rehydration (Fig. 7).^{6,15,76,80} Before the formation of the vesicles, the amphiphiles are dissolved in the organic solvent and placed on a solid surface (*e.g.* glass, Teflon, metal wires) before evaporating the solvent entirely to form a thin layer on that surface (Fig. 10a). Then addition of the aqueous medium causes swelling of the film and ultimately forms vesicles once the bilayer leaves the surface. The swelling and budding stage can be facilitated by mechanical stirring or sonication. Film rehydration generates MLVs and SUVs with broad size distributions, but is nonetheless the



Can be aided by sonication, vigorous stirring, electric field, hydrogels.

Fig. 7 Self-assembly of lipids and block copolymer vesicles using solvent-free methods.



most commonly used method to form vesicles as the procedure is straightforward, no special equipment is required and it reproducibly yields a high number of vesicles completely free of organic solvents. Solid rehydration is conceptually similar but uses a bulk powder of the amphiphiles rather than a cast solution. Due to the problematic high polydispersity obtained with film hydration, commonly sequential extrusion is used to achieve better distribution control. This technique consists of repeatedly passing a crude heterogeneous mixture of vesicles (usually MLVs) through a polycarbonate membrane with small pores (typically 100 nm diameter but can be any desired size) to obtain homogeneously distributed SUVs of diameter matching the pore size.^{30,80,93}

Aided film rehydration methods such as electroformation or gel-assisted hydration are also quite popular as they allow the formation of GUVs with much lower polydispersity than simple film hydration.¹⁵ These methods are also desirable as they allow the formation of vesicles for challenging amphiphiles.⁹ Electroformation was pioneered in 1986 on liposomes, and consists of depositing a thin film of amphiphiles on electrodes (Pt or indium titanium oxide (ITO) glass).^{77,83,94} After addition of the aqueous medium, an alternating current is applied facilitating the swelling and release of the vesicles. The electric current is thought to cause fluctuation in the film and inter-layer repulsion resulting in detachment from the electrodes.⁹ Electroformation is the method of choice to obtain liposome GUVs and was reproduced for polymersomes in 1999 on Pt wires.⁴⁶ And has been repeated for a larger pool of block copolymers and extended to ITO glass.^{43,67,95–97} Remarkably, despite the fact that electroformation is well-established for liposomes, special in-house devices are almost exclusively used and poorly detailed. A recent study by Dionzou *et al.* clearly pointed out that the nature of the block copolymer and conditions needed to reproducibly obtain polymersome GUVs are challenging to determine.⁶⁷

In gel-assisted hydration,^{83–85,98,99} the surface is pre-coated with dissolved agarose or poly(vinyl alcohol) (PVA) in warm water, and dried to form a thin gel layer before forming the amphiphile layer. The driving force is the swelling of the hydrogel when the coated surface is placed in aqueous medium. This method has only been described more recently (2009)⁹⁸ and has been gaining in popularity. It also requires more readily available equipment (coverslip rather than an electroformation chamber). However, PVA or agarose could also get embedded in the vesicular membrane modifying its properties and behavior. One major drawback of all film hydration methods is the difficulty in encapsulating water-soluble molecules.⁸⁰ Indeed, any desired entities need to somehow intercalate between the amphiphilic film layers which can be difficult for macromolecules and charged molecules but had nonetheless been reported.⁷⁷

Another approach that has become very popular recently is polymerization-induced self-assembly (PISA), which has been used as a solvent-free polymer-specific methodology to form vesicles.^{68,100–103} This method relies on the solubility differences commonly in water between the monomer and polymer: as the hydrophilic monomer polymerized on a hydrophilic macroinitiator, also called the stabilizer-block; the polymer becomes increasingly

more hydrophobic, entropically driving the amphiphile self-assembly. After reaching an initial critical degree of polymerization, micellar nucleation occurs. Further polymerization leads to spherical nanoparticle assemblies (first spheres then worms, other intermediate structures, and finally vesicles) (Fig. 8). Targeted assemblies can be achieved by adjusting the amphiphile concentration or the degree of polymerization of the stabilizer-block. PISA has been achieved by using diverse polymerizations but by far the most commonly used one is reversible addition fragmentation transfer (RAFT) polymerization to form diblock copolymers but also triblock copolymers and uses organosulfur based chain-transfer agents (CTAs). Most commonly, PISA is limited to self-assembled structures in the nanometer regime (nanoparticles, worms, or SUVs of 10–100 nm). Recently, Albertsen *et al.* described the photo-initiated PISA of 2-hydroxypropyl methacrylate (HPMA) onto PEO, forming GUVs (up to 10 μm).¹⁰¹ The Ru-photosensitizer generates radicals upon exposure to 450 nm radiation, which induce the polymerization of HPMA. These GUVs underwent what the author describes as phoenix behaviors where the change in osmotic pressure during self-assembly leads to the cyclic growth and collapse of the vesicles, catalyzed by exposure to blue light. Over time the GUVs thus gradually shrank, and increased in number demonstrating that SUVs might be more thermodynamically stable than GUVs. PISA perfectly exemplifies the versatility of block copolymers compared to lipids and allows the shortcutting of the different stages of assembly. However, because the method relies on hydrophilicity modulation, PISA is limited to a certain pool of specific blocks (typically HPMA) exhibiting the correct properties but has nonetheless shown incredible diversity.

(b) Solvent displacement methods

Solvent displacement methods define any techniques where organic solvents and the aqueous phase coexist to promote the self-assembly of the amphiphiles before removal of the organic phase. These techniques have the great advantage of facilitating the efficient encapsulation of water-soluble molecules but have the difficulty of organic solvent removal. Organic solvent molecules trapped in the hydrophobic bilayer or dissolved in the aqueous phase would cause exposure of sensitive materials (protein, DNA, small molecule) to an organic solvent, which might sometimes limit their application since the organic solvent can either inactivate the biological macromolecule or be toxic.²⁵ Those methods also often require the use of additives such as surfactants (typically small molecules or block copolymers containing PEG), which get embedded in the membrane as also amphiphilic.^{104–106}

Solvent injection consists of slowly adding a solution of amphiphiles in organic solvent (most of the time ether or ethanol) to an aqueous medium under vigorous stirring (Fig. 9).^{15,25} Such vesicles are SUVs, often MLVs and have a broad size distribution but it is an easy and efficient method to obtain vesicles like film hydration. Vesicle polydispersity and lamellarity can be homogenized by extrusion. Similar methods use reduced pressure, dialysis and/or centrifugation as means to remove organic solvents after mixing of the aqueous and organic phases to obtain similar vesicle populations.



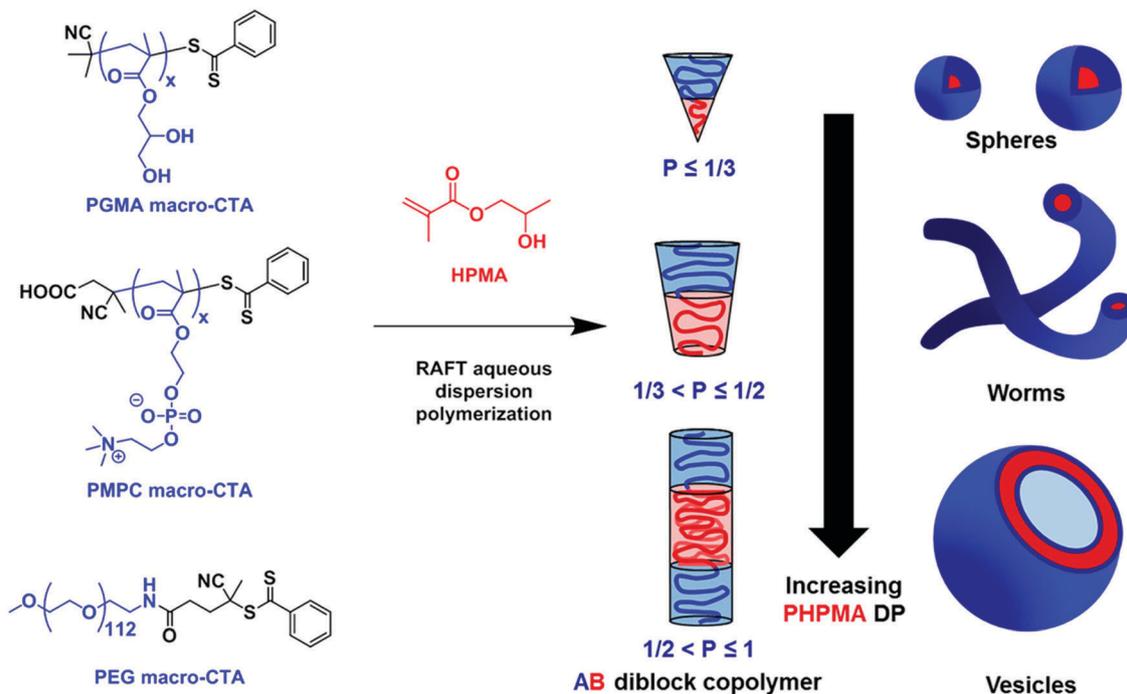


Fig. 8 Polymerization-induced self-assembly (PISA) of 2-hydroxypropyl methacrylate (HPMA). A variety of macromolecular chain transfer agents (CTAs) were used to produce various assemblies depending on the degree of polymerization of HPMA. Reprinted with permission from N. J. Warren and S. P. Armes, *J. Am. Chem. Soc.*, 2014, **136**(29), 10174–10185. Copyright (2014) American Chemical Society⁶⁸ (<https://pubs.acs.org/doi/abs/10.1021/ja502843f>).

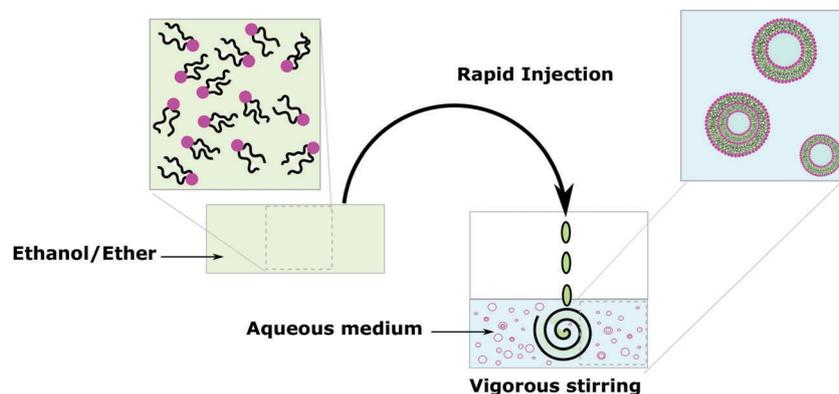


Fig. 9 Self-assembly of lipids and block copolymer vesicles by using solvent injection.

A more controlled manner to form vesicles in defined size distribution is by using emulsion phase transfer (Fig. 10).^{80,86–88} An initial w/o (water-in-oil) emulsion is prepared by vigorously stirring a small quantity of aqueous media to a solution of amphiphiles in an organic solvent, effectively forming water droplets coated with the amphiphile. Then the emulsion is poured on top of an oil-in-water biphasic system where amphiphiles dissolved in the organic solvent should have assembled into a monolayer at the interface. Due to their density difference, the monocoated water droplets from the emulsion sink to the aqueous phase through the amphiphilic saturated interface generating w/o/w double emulsions. Water droplet migration can be facilitated by using sucrose in the inner water droplet and equiosmotic aqueous glucose in the biphasic system due to their density

difference⁸⁰ and/or centrifugation (emulsion-centrifugation),^{91–93} which can also (partially) remove the trapped organic solvent, thus generating low polydispersed vesicles. As the outer and inner layers of the vesicles are formed at separate stages, complete control of the vesicular structure is possible and as such asymmetric vesicles can be obtained (see Section Va Asymmetric membranes for more details).^{86,87} Moreover, encapsulation can be achieved by doping the inner water droplet with desired hydrophilic cargos contrasting with the solvent-free methods.

An improvement to the emulsion phase transfer methods is microfluidics. Microfluidics is probably the most advanced solvent displacement method and has allowed exquisite control and progress in constructing artificial cells.^{21,89,90,107} This water-in-oil-in-water (w/o/w) double emulsion technique selectively



Can be controlled by microfluidics

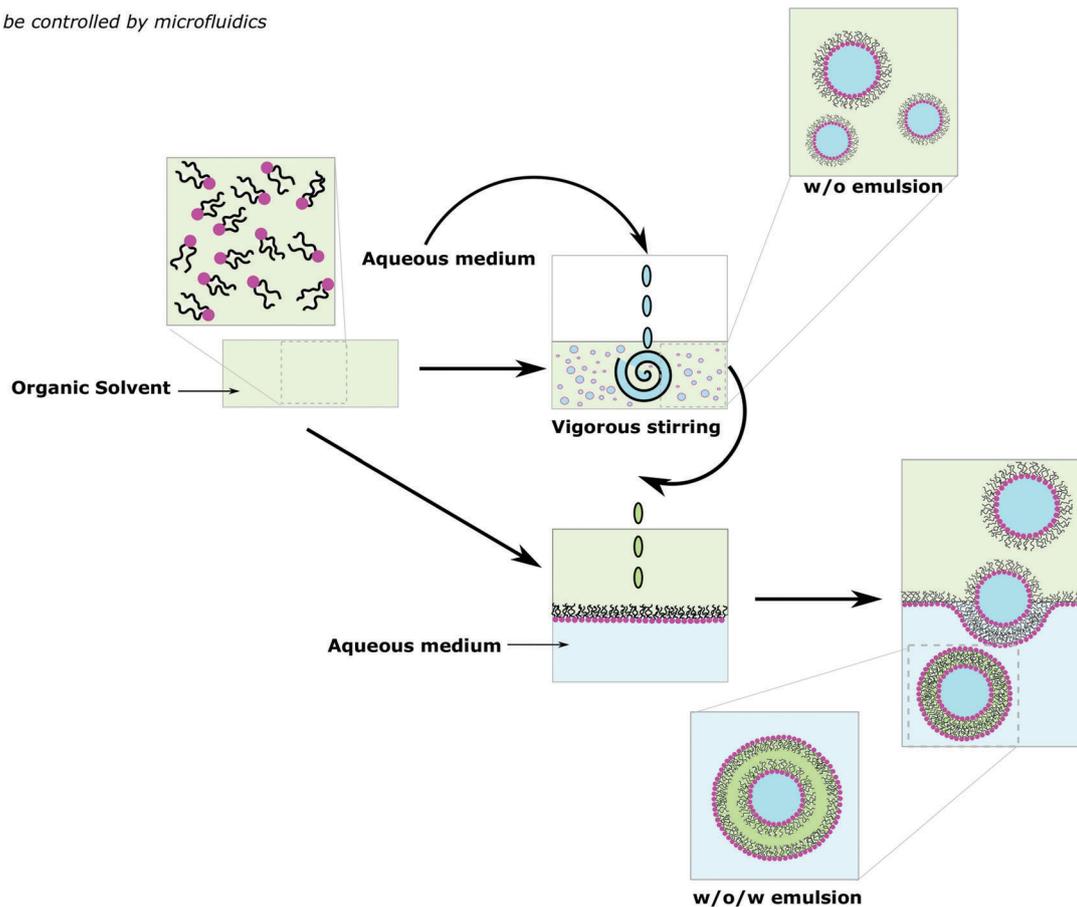


Fig. 10 Self-assembly of lipids and block copolymer vesicles by using emulsion phase transfer.

forms vesicles homogeneous in size from SUVs to GUVs depending on the requirement.^{80,108} Amphiphile-stabilized water-in-oil (w/o) emulsions are first prepared using organic solvents immiscible with water at the first junction. Then these droplets flow to a second junction through an aqueous phase coated with an amphiphile to obtain w/o/w double emulsions (Fig. 11). A monolayer of amphiphile is obtained at both the w/o interphases. Solvent extraction from the double emulsion to form vesicles is a process called dewetting.^{105,109} Dewetting of the double emulsion can be achieved by using ethanol in the outer aqueous phase^{21,110} surfactants,¹⁰⁵ octanol as the oil-phase rather than the traditional hexane or chloroform,¹¹¹ or solvent evaporation.^{109,112} Absolute dewetting is a fundamentally challenging process and most

vesicles formed by solvent-displacement methods still have, in the best cases, trace amounts of solvents embedded within their membrane. Moreover, testing for remaining solvent is rarely done in advanced systems.^{90,113} Highly complex mixtures of components are required for solvent removal, stabilization and formation (surfactant, sugars, polyelectrolytes, non-amphiphilic polymers *etc.*) which are imbedded in the membrane or in the vesicular lumen.^{21,89,105,114,115} These impurities will affect the vesicle properties and impair potential encapsulated bio-components and are often overlooked. In addition, as the inner amphiphilic monolayer is much smaller than the outer monolayer, considerable rearrangement needs to occur which sometimes leads to undesired aggregation within the membrane if the number

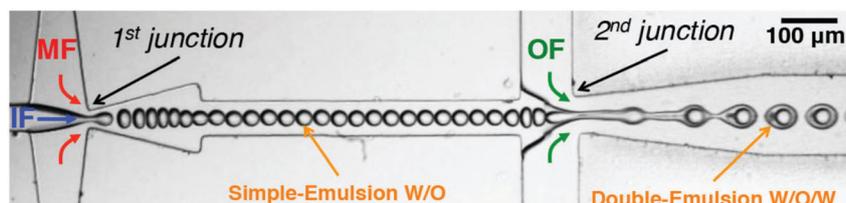


Fig. 11 Production of water-in-oil-in-water (w/o/w) double emulsion by using a microfluidic device. Amphiphile-stabilized water-in-oil (w/o) emulsions are first prepared using organic solvents (middle fluid, MF) immiscible with water (inner fluid, IF) at the first junction. The droplets flow to a second junction through an aqueous phase (outer fluid, OF) coated with an amphiphile to obtain w/o/w double emulsions. Adapted from J. Petit, I. Polenz, J. C. Baret, S. Herminghaus, O. Baumchen, *Eur. Phys. J. E: Soft Matter Biol. Phys.*, 2016, **39**, 59. <https://doi.org/10.1140/epje/i2016-16059-8.21>.



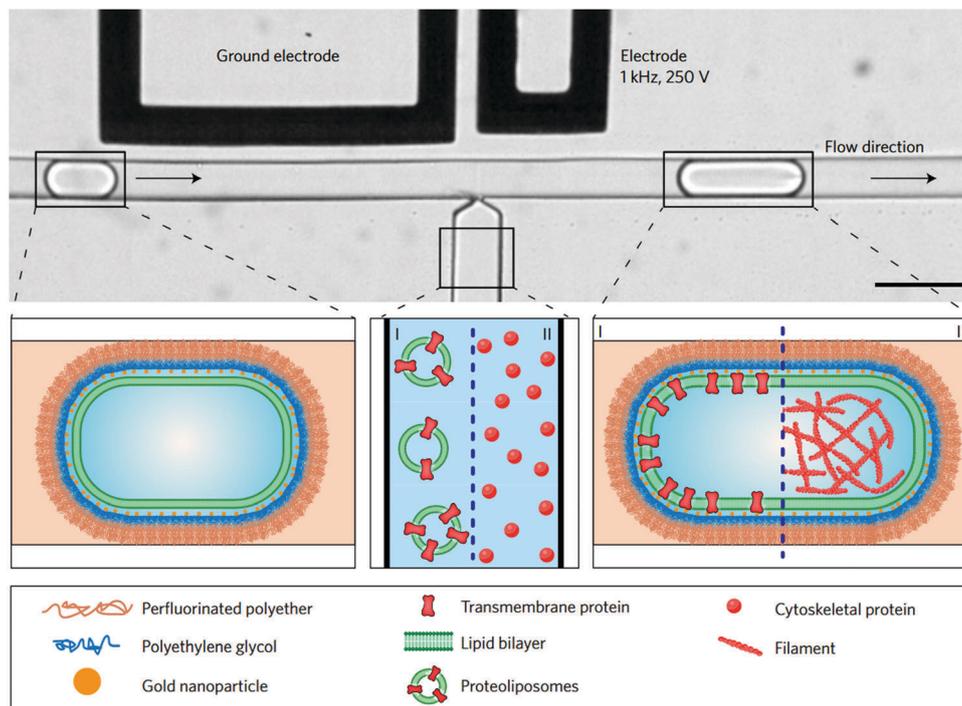


Fig. 12 Sequential bottom-up assembly of transmembrane and cytoskeletal proteins into droplet-stabilized GUVs by high throughput pico-injection. Scale bar 50 μm . Reproduced with permission from M. Weiss, J. P. Frohnmayer, L.T. Benk, B. Haller, J. W. Janiesch, T. Heitkamp, M. Börsch, R. B. Lira, R. Dimova, R. Lipowsky, E. Bodenschatz, J. C. Baret, T. Vidakovic-Koch, K. Sundmacher, I. Platzman and J. P. Spatz, *Nat. Mater.*, 2018, **17**, 89–96.¹⁰³

of amphiphiles is in excess.^{80,105,109,112} The final size of the vesicle is controlled by the initial inner water droplet, achieving very low size distributions. This method also shows a high encapsulation efficiency of a variety of proteins and small molecules and it was also proven that these entities can be co-encapsulated, an important aspect to generate advanced protocells.

Pico-injection microfluidics has been developed as a novel means for post self-assembly encapsulation in droplet-supported vesicles.^{107,113,116} This active encapsulating technique is used for the manipulation of compartments after their formation and the sequential addition of multiple components which would not be otherwise incompatibly added in bulk. By means of electroporation, single drop injection is carried out, precisely controlling the cargo concentration in each vesicle. Recently, Weiss *et al.* described the sequential loading of transmembrane F_0F_1 -ATP synthase and integrin embedded into the lipid bilayer and the encapsulation of cytoskeletal proteins resulting in actin filaments and microtubule reconstitution by high-throughput microfluidics pico-injection (Fig. 12).¹¹³ Pico-injection has proven to be an excellent technique to achieve the controlled encapsulation of complex bio-relevant macromolecules in the membrane or inside droplet-stabilized GUVs. The technique is however still rarely used as it is so far limited to lipid-droplets and requires highly specialized equipments.

(c) Microscopy observation

Depending on the size of the vesicles obtained, different microscopy techniques can be used for the observation and characterization of the vesicles (Fig. 13).¹⁵ For GUVs, optical microscopy is most

frequently used, as it is non-destructive and allows for direct visualization under physiological conditions.¹ Alternatives such as fluorescence microscopy are also popular as the enclosed compartment with encapsulated soluble fluorescent dyes and/or the membrane using lipophilic fluorophores either by covalent or non-covalent linkage can be simultaneously labelled specifically with fluorochromes with high sensitivity (typically 50 fluorescent molecules per mL or 1 fluorescent molecule per 1000 lipids).¹ However, due to the limited resolution of conventional optical microscopy only GUVs can be easily observed.¹ For SUVs and LUVs, transmission electron microscopy (TEM) is favored as it has a resolution 10^5 higher than optical microscopy.¹⁵ Cryo-TEM is preferred as it allows for the observation of a sample in its hydrated state. TEM can also be used to determine the membrane thickness of SUVs and GUVs.^{9,12,15} However, the electrons may affect the state of the specimen and it also has to be considered that most of the experiments are not conducted under wet conditions, but require drying or freezing of the sample. State-of-the-art techniques for super-resolution such as scanning near-field optical microscopy (SNOM), photo-activated localization microscopy (PALM), stimulated emission depletion microscopy (STED), or structured illumination microscopy (SIM) would allow the wet imaging of smaller vesicles but are still very rarely used.^{117–119}

IV. Stimuli-responsive vesicles

Liposomes and polymersomes can be chemically modified (functionalized) to tune their properties, which is particularly important for certain biomedical applications.^{15,76,120–122}



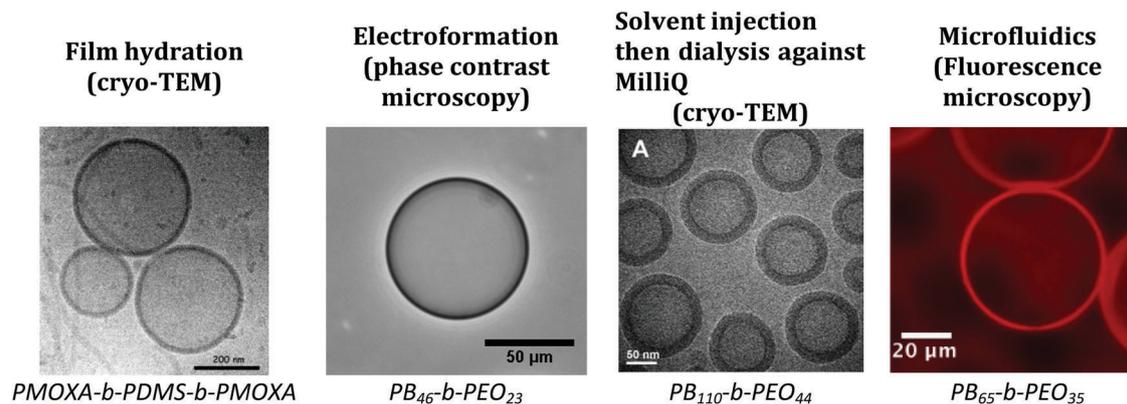


Fig. 13 Microscopy images obtained for different techniques. TEM: transmission electron microscopy. Film hydration: Reprinted from K. Kita-Tokarczyk, J. Grumelard, T. Haefele, W. Meier, Block copolymer vesicles—using concepts from polymer chemistry to mimic biomembranes, *Polymer*, **46**(11), 3540–3563. Copyright (2005), with permission from Elsevier.¹⁵ Electroformation: in-house image. Solvent injection: adapted from ref. 20 with permission from The Royal Society of Chemistry. Microfluidics: Reprinted with permission from J. Petit, I. Polenz, J. C. Baret, S. Herminghaus and O. Baumchen, *Eur. Phys. J. E: Soft Matter Biol. Phys.*, 2016, **39**, 59. <https://doi.org/10.1140/epje/i2016-16059-8.21>.

An important application of small vesicles is drug delivery.²⁴ To be efficient, such vesicles must achieve two characteristics: delivery of the drugs at the desired site and release of the vesicle content only at that location. Frequently, a responsive stimulus is necessary for specific and efficient drug delivery. Most stimuli-responsive or ‘smart’ vesicles release their cargo upon activation either due to (a) full disruption (disassembly) usually caused by making the hydrophobic moiety of the amphiphile more hydrophilic or (b) tuning the vesicles’ permeability for a slow controlled release of encapsulated molecules (‘breathing vesicles’) or (c) changing their supramolecular morphology.²⁸ These changes can be reversible or irreversible and arise from changes to the supramolecular assemblies (e.g. H-bonding network) or chemical changes in the molecular unit.¹²³ Functionalization can also be used in nanoreactors to enhance chemical processes.²⁴ Moreover, stimuli-responsive behaviors are intrinsically the basis of biological processes (cell signaling, immunity, energy and small molecule production *etc.*) and is an important step in biomimetic studies.²⁴ Stimuli can be of biological or chemical nature such as pH changes, response to ROS, enzyme, glucose *etc.* or physical stimuli such as light, temperature, ultra-sound, magnetic fields, pressure *etc.*^{24,124,125} and can be combined for multiple responsivities.

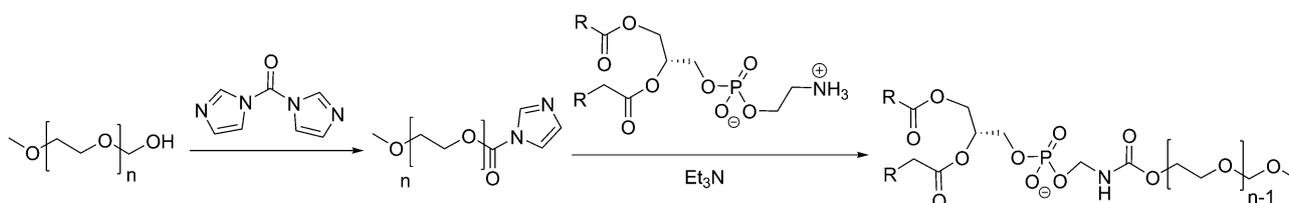
Liposomes are usually functionalized to overcome their low stability, leakiness, and low blood circulation times. PEGylated lipids (PEG: poly(ethylene glycol)) have been shown to be successful in forming liposomes as PEG is hydrophilic and its covalent attachment to the hydrophilic head of the lipid does

not modify its amphiphilicity.^{9,30,126} PEG-stabilized liposomes are thus the most prominent class of functional liposomes and are most commonly formed by covalently grafting a PEG chain to lipids and doping the liposomes with these specialized lipids. The synthesis of PEGylated lipids is achieved by adding a terminal activated PEG chain (for example with imidazole) to an amine terminated lipid (PE) (Scheme 2), which forms MLV liposomes (112–136 nm).¹²⁶ PEGylated liposomes are also called ‘stealth’ liposomes. The addition of biocompatible PEG chains increasingly helped the bioavailability of liposomes *in vivo* by reducing potential interactions with immune cells (e.g. macrophages) by coating the liposome surface and reducing liposome aggregation.^{19,127} These ‘stealth’ liposomes have greatly increased blood circulation time from 1 h to 1–2 days and thus improved drug distribution.^{1,12,19}

(a) Biological or chemical stimuli-responsive vesicles

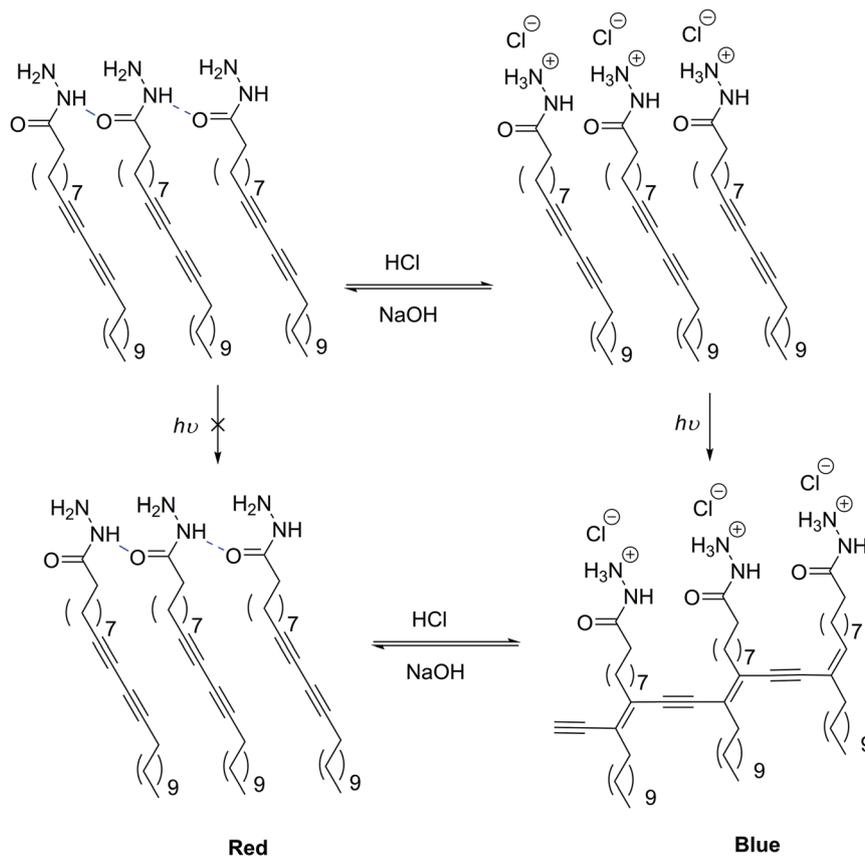
pH-Sensitive LUVs and SUVs are the most studied release systems as one can make use of physiological pH-gradients.^{122,128,129} Extracellularly, inflammatory tissues and tumors are more acidic than healthy tissues, or intracellularly endosomes and lysosomes also show acidic conditions (below pH 7), making pH change an ideal trigger for drug delivery.¹²⁵

pH-Sensitive liposomes are commonly generated using a small amount of anionic lipid derivatives such as *N*-succinyldioleoyl-phosphatidylethanolamine (COPE),¹³⁰ cholesteryl hemicussinate (CHEMS),^{131–133} a cholesterol derivative, or α -tocopherol acid succinate¹³¹ implemented in neutral lipid vesicles. In response



Scheme 2 Example synthesis of PEGylated-lipids by Allen *et al.*¹²⁶



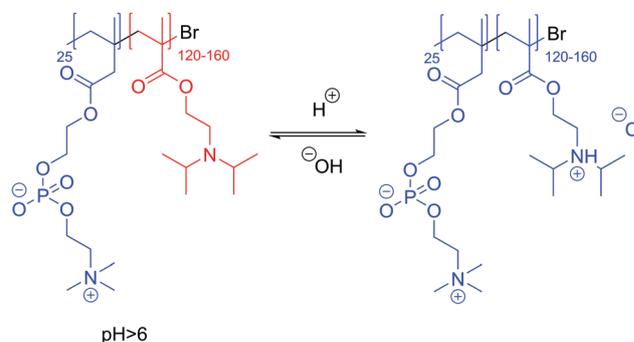


Scheme 3 pH-Responsive SUVs due to a supramolecular change in the head group hydrogen bonding network.¹³⁴

to an acidic environment, for example below 5.8 (pK_a of CHEMS),¹³³ the negative moiety of these lipids gets protonated switching from vesicles to inverted micelle structures. pH-Sensitive colorimetric reversible LUVs (≤ 800 nm) were described using the small amphiphilic molecule polydiacetylene (PDA) and could be used for bio-imaging.¹³⁴ By its nature, PDA's structure is analogous to a lipid as it is composed of a hydrophilic head (acetohydrazide), rather than the much longer repeated hydrophilic moiety found in block copolymers. Upon addition of HCl, the hydrogen bonding network of the amide moiety is disrupted, replaced by an ionic stabilization of the terminal amine by chloride anion and presumably some charge repulsion causing a rearrangement of the macromolecular assembly without vesicular disassembly (Scheme 3). This supramolecular assembly of the vesicles allows them to undergo photo-polymerization visualized as a shift from a red to a blue color. In the neutral form, polymerization of the diacetylene groups is not possible due to geometrical constraints determined by the hydrogen bonding network of the head group. Later Yuan and Hanks also added an amphiphilic fluorophore BO558 in these PDA vesicles (≤ 450 nm) which resulted in a reversible switching of the fluorescence (on when neutral and off when acidic) which could also be used in bio-imaging.¹³⁵

pH-Responsive polymersomes (exclusively LUVs or smaller) can also be formed from hydrolytic, acid cleavable polymers, or are most commonly composed of ionizable blocks such as a polyacid (*e.g.* carboxylic or sulfonic acid) or a polybase

(*e.g.* amines).^{15,24,124,125} In the presence of different pH values, their hydrophobic fraction becomes more water-soluble by ionization modification and thus the polymersomes disassemble. pH-Responsiveness has the advantage of being virtually instantaneous compared to other chemical modifications such as hydrolysis. Armes and coworkers¹³⁶ described a poly(2-methacryloyloxyethyl phosphorylcholine)-*b*-poly(2-(diisopropylamino)ethyl methacrylate) (PMPC-*b*-PDPA) block copolymer where PMPC is a phospholipid-like hydrophilic block and PDPA a polybase (Scheme 4). At $pH > 6$, PDPA (pK_a 6.3) acts as a hydrophobic block and stable polymersomes (100 nm) can be formed.



Scheme 4 Example of pH-responsive polymers bearing the polybase PDPA. Blue: hydrophilic. Red: hydrophobic.¹³⁶



Upon lowering the pH, the tertiary amine is protonated, becoming hydrophilic and causing vesicles' disassembly. The authors further proved that the anti-cancerous drug doxorubicin can be efficiently encapsulated and retained when mixed in the presence of PMPC-*b*-PDPA at pH 2 and then adjusted to pH 7.4. These polymersomes were subsequently used to provide efficient targeted intracellular delivery of drugs, antibodies, DNA, *etc.*¹²⁵ In a more advanced system, Rodriguez-Hernandez and Lecommandoux reported reversible poly(glycolic acid)-*b*-poly-L-lysine (PGA-*b*-PLys) polymersomes (100–200 nm) where the PGA block is hydrophobic and PLys hydrophilic at pH < 4, and the hydrophilicities of the blocks revert at pH > 10 where the PGA block is hydrophilic and PLys hydrophobic.¹³⁷

Eisenberg and coworkers detailed pH-responsive PEO₄₅-*b*-PS₁₃₀-*b*-PDEA₁₂₀ (PDEA: poly(2-diethylaminoethyl methacrylate)) LUVs clearly exemplifying the 'breathing' effect of the membrane.¹³⁸ The pH-responsive block PDEA causes a change in vesicle sizes (from 250 nm to 480 nm), wall thickness (25 nm to 80 nm), structure and appearance (apparent swelling of the membrane at lower pH) (Fig. 14). These transformations undoubtedly affect the overall polymersome permeability creating a notably more porous membrane for protons.

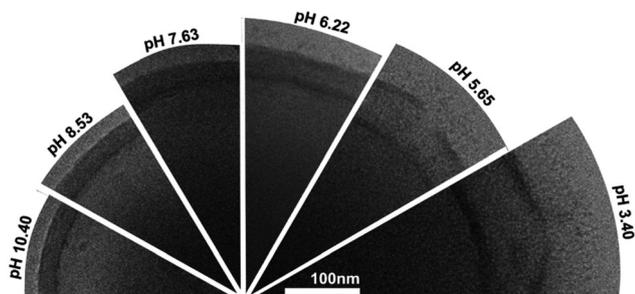
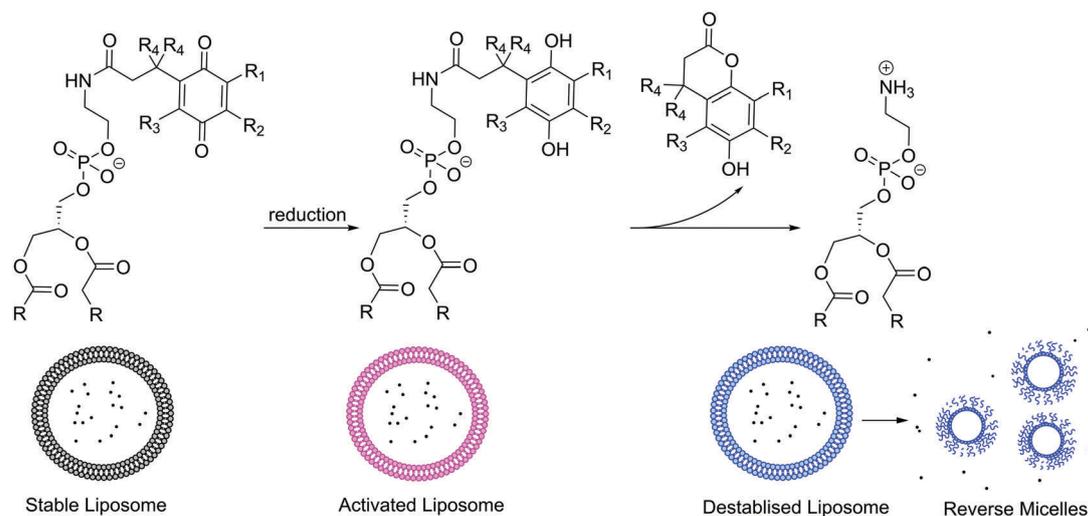


Fig. 14 Cryo-TEM image of a pH-responsive PEO-*b*-PS-*b*-PDEA polymersome at different pH values. Reprinted with permission from S. Yu, T. Azzam, I. Rouiller and A. Eisenberg, *J. Am. Chem. Soc.*, 2009, **131**(30), 10557–10566. Copyright (2009) American Chemical Society.¹³⁸

Many biological processes are triggered by changes of redox potential. Thus, similarly to pH gradients, reduction or oxidation sensitive moieties can conveniently be triggered for controlled release or imaging particularly for intracellular delivery due to the redox potential difference between the different organelles and the cytosol.¹²⁵ In healthy cells, ROS such as peroxides, hydroxyl radicals, singlet oxygen *etc.* are produced as a byproduct of aerobic metabolism and are kept under control by antioxidants, while any damage to cells is repaired. Infection, inflammation, cancer, and many other diseases cause oxidative stress, an overproduction of ROS in cells, which cannot be controlled by the body's natural defenses. Thus, redox-responsive vesicles are attractive for pharmaceutical purposes. Reduction or oxidation sensitive moieties such as ferrocene, disulfide bonds (cleavage under reductive conditions), quinoline *etc.* can all be implemented in the lipid's head or tail or polymer blocks.¹³⁹ For example, quinoline functionalized DOPE forms stable liposomes (100 nm); however, under reducing conditions, quinoline switches to hydroquinone that can rearrange as a lactone, releasing the free PE lipid.¹⁴⁰ This lipid is not capable of maintaining a bilayer and rearranges into reverse micelles, releasing its cargo (Scheme 5).

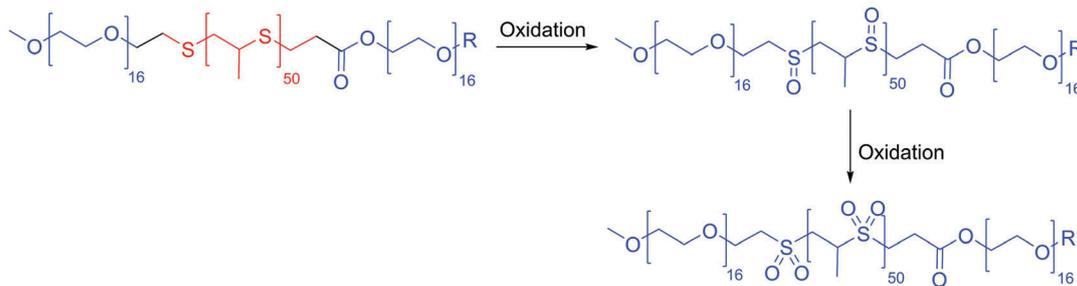
Many redox responsive polymersomes have also been described. The first oxidative responsive polymersomes (100 nm) were described in detail by Napoli *et al.* in 2004 using PEO-*b*-PPS-*b*-PEO (PPS: polyphenylene sulfide) (Scheme 6).²⁷ The thioether in the hydrophobic block becomes oxidized to a sulfoxide initially and ultimately sulfone becoming hydrophilic and thus changing the morphology of the polymersomes to micelles. The ferrocenylsilane block or in general ferrocene containing polymers have been used to form redox active organometallic vesicles.¹⁵ These polymersomes are disrupted in the presence of high concentration of ROS released by the diseased cells causing the oxidation of ferrocene to ferrocenium which destroys the vesicles.²⁴

Enzymes, being highly specialized biological entities, have superior efficiency for selective and sensitive delivery of vesicular cargo. Enzyme-responsive polymersomes were used for drug delivery as certain enzymes are overexpressed in pathological



Scheme 5 Example of redox responsive lipids bearing quinoline groups.¹⁴⁰





Scheme 6 Oxidative responsive polymersomes by Napoli *et al.*²⁷

regions.²⁴ For example, Cathepsin B is found in abundance in tumor tissues and specifically degrades Gly-Phe-Leu-Gly (GFLG) sequences.¹⁴¹ Lee *et al.* used a PEO-*b*-PDLLA (PDLLA: poly(D,L-lactide)) block copolymer with a GFLG linker which self-assembles into SUVs (124 nm). The block copolymer is cleaved in the presence of Cathepsin B (at tumor site) causing disassembly and the release of any drugs for example the anti-epidermal growth factor receptor (anti-EGFR), a chemotherapy agent.¹⁴²

(b) Physical stimuli-responsive vesicles

Temperature is another popular trigger for release of vesicular cargo as temperature modulates cell activity. Thermoresponsive liposomes rely on the transition temperature of natural phospholipids rather than synthetically modified lipids. Typically DPPC and MSPC (1-myristoyl-2-stearoyl-*sn*-glycero-3-phosphocholine) are used as these lipids have a T_m between 41 and 45 °C, a physiologically relevant temperature range especially for targeted drug delivery.^{121,143–146} The thermo-responsiveness of the liposomes can be modulated by using a mixture of lipids. Nonetheless, lipid-only temperature sensitive liposomes remain rare as lipid modification is difficult and the simple doping of liposomes with thermosensitive polymers represents a much more achievable modulative platform for thermoresponsive vesicles. For example Pippa *et al.* formed C₁₂H₂₅-PNIPAM-COOH/DPPC (PNIPAM: poly(*N*-isopropylacrylamide)) hybrid vesicles (100–200 nm) behaving similarly to pure polymeric vesicles.¹⁴⁷

Multiple thermoresponsive polymersomes have been described most frequently also relying on PNIPAM as the thermoresponsive block.^{124,125,128} PNIPAM can undergo a phase transition from hydrophilic to hydrophobic upon heating at its lower critical solution temperature (LCST) of ~32 °C. Thus, at the physiological temperature of ~37 °C, the PNIPAM block is hydrophobic by displacing the hydrogen bond network responsible for its solvation (hydrophilicity) below its LCST. The block copolymer is thus amphiphilic and PEO-*b*-PNIPAM can self-assemble into polymersomes (>1 μm) (Fig. 15). Below the LCST, the thermoresponsive block is solvated in water by favored hydrogen bonding and causes the polymersome to dissociate releasing any encapsulated substances. LCST polymers such as PNIPAM can thus be used to release drugs locally when using hypothermic patches or ice packs, local cryosurgery probes or tissue freezing.¹⁴⁸

Interestingly, almost all thermoresponsive polymersomes use LCST responsive moieties, which intuitively seem more troublesome to use than an upper critical solution temperature

(UCST) block since the polymersomes have to be stored at elevated temperatures and release their cargo at lower temperatures. Tumor tissues or infection sites tend to have more elevated temperature (+2–5 °C) than healthy tissues.¹⁴⁹ Therefore, polymers with an UCST in water would be much more interesting and easier to use than LCST polymers. Only very recently was a poly(phosphonate) block (PPE) shown to have a UCST.¹²³ PPE-*b*-PEO was subsequently proven to self-assemble into polymersomes (100–200 nm) at room temperature and to fully disassemble upon heating up to 60 °C as the PPE block became hydrophilic and to reversibly re-assemble when cooling (Scheme 7). Unfortunately, full disintegration of the polymersomes did not occur at the physiological temperature; however, these block copolymers are a compelling initial proof of concept that polymersomes with UCST behavior are achievable and further work should be done to tune these PPE blocks to a UCST close to 40 °C.

Light can initiate a simple, non-invasive, efficient, and on-demand trigger since time, exposure, wavelength, and intensity can be well controlled, and it is easier to handle than chemical responses.^{24,125} However, most light-responsive moieties used in vesicles are triggered by UV or visible light irradiation which limits the vesicle application to very close to the surface area of the body,¹²⁵ and near-infrared sensitive assemblies would be more appropriate for deep tissue drug-release. Bayer *et al.* developed photo-responsive liposomes (<200 nm) using a synthetic derivative of PC with a 2-nitrobenzyl moiety and showed the release of an hydrophobic cargo (Nile red) upon irradiating the sample at 350 nm (Scheme 8).⁵⁷

Other examples of photo-labile phospholipids were described by Suzuki *et al.*, who used glycerophosphocholines presenting two terminal 2-nitrobenzyl groups.¹⁵⁰ By UV irradiation at 350 nm the vesosomes release their double-stranded DNA (dsDNA) cargo into the outer GU (10 μm diameter). Ces and coworkers described UV-C responsive PC derivatives composed of diacetylene.¹⁵¹ The UV irradiation catalyzes the polymerization of diacetylene effectively cross-linking the phospholipid aliphatic chains creating pores in the membrane, a process already described in Scheme 3.

Photo-responsive polymersomes have been described using azobenzene (reversible *cis/trans* isomerization), spiropyran (ring opening), *O*-nitrobenzyl (photocleavage of block junction), coumarin, 2-diazo-1,2-naphthoquinone *etc.* moieties which also change hydrophobicity (conformational changes) or cause cleavage leading to the dissociation of the vesicles.^{16,125} Not all light-triggered polymeric systems result in irreversible disassembly of polymersomes.



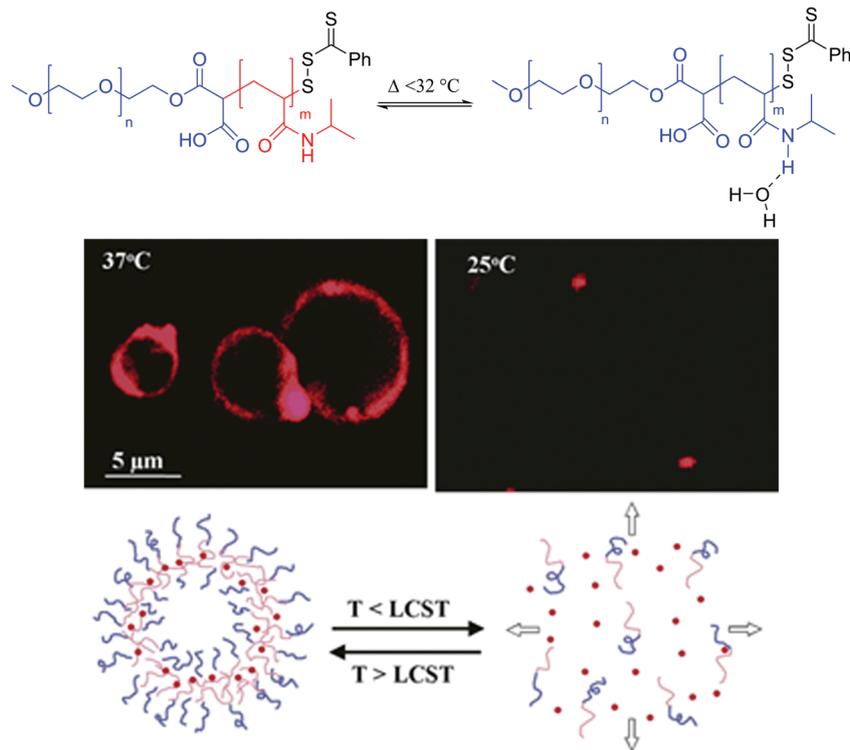
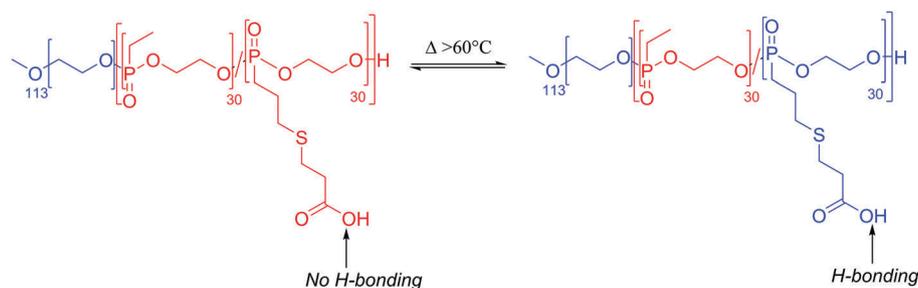
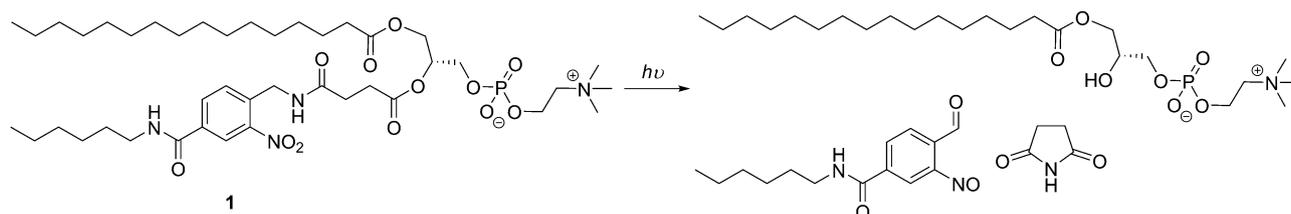


Fig. 15 Thermoresponsive block copolymer bearing the thermosensitive PNIPAM block with LCST behavior. Reprinted with permission from S. Qin, Y. Geng, D. E. Discher and S. Yang, *Adv. Mater.*, 2006, **18**, 2905–2909. Copyright (2006) Wiley.¹⁴⁸



Scheme 7 Thermosensitive PPE block with UCST behavior.¹²³



Scheme 8 Photo-responsive modified PC which can assemble into photo-responsive liposomes.⁵⁷

For example, Su *et al.* described light-responsive PAA-*b*-PAzoM (PAzoM: poly{6-[4-(4-methylphenylazo)phenoxy]hexyl acrylate}) polymersomes (2–4 μm) which can switch between spherical polymersomes and earlike shaped ones depending on the isomerization of the azobenzene moiety.¹⁵²

Vesicles, whether liposomes or polymersomes, can be modified in a variety of ways. Their primary purpose is the encapsulation

and release of drugs. By encapsulating drugs, they are protected against enzymatic degradation and the encapsulation can improve the biodistribution and reduce side-effects.^{19,25,76} SUVs make efficient carriers of small molecules in contrast to most other nanomedicine agents,²⁸ and have been shown to deliver both hydrophilic (encapsulated) and hydrophobic compounds (embedded in the membrane) to target tissues.¹²⁷ We noticed



that by far almost all functionalized or stimuli responsive vesicles are small (LUVs or SUVs) and the few micron-sized vesicles which have been reported are only a few μm ($< 5 \mu\text{m}$). For biomedical purposes, LUVs are more adapted, as they are more malleable, can easily circulate in the body and can even be internalized into cells. Nonetheless there is a clear scarcity of functional and/or stimuli-responsive GUVs (ideally $> 10 \mu\text{m}$) which are essential for characterization of the membrane mechanical properties and for the generation of protocells and biomimetic compartments.

V. Mimicking cell functions in vesicles

Vesicles have been extensively studied and developed from a biomedical perspective because they are biocompatible. Nonetheless, because of their compartmentalization effect, vesicles can also be advantageous to use for enhancing chemical reactions and biosynthesis as reactors or to generate protocells.^{153–155} For example, incompatible reactions could be achieved by simply compartmentalizing them.⁴ Ultimately a cell can also be described as a highly evolved microreactor which encapsulates many nanoreactors (organelles). Constructing simplified synthetic cells is an increasingly expanding area because it contributes to an understanding of fundamental biological processes and provides new and unique prospects for future therapeutic applications, from self-regulating bioreactors to new biomedical devices.^{90,107,156–159} Ultimately, the goal would be to understand cell processes sufficiently using models so that we could replace natural living cells with functional synthetic cells.

(a) Membrane functionality

We have already shown that synthetic compartmentalization can be readily achieved with liposomes and polymersomes. However, cell membranes are very complex: they are asymmetric, have

lipid rafts, use a variety of means to transport ions and small molecules through the membrane in a controlled manner, communicate between cells, act together to form tissues *etc.*

Transversal heterogeneity. In a cell, there is a clear difference between the intracellular and the extracellular environment and thus a demand for physiologically distinguishing the inner and outer sides of the membrane. In terms of lipid compositions, sphingomyelin (SM) and PC are mostly found on the outer layer and PE, phosphatidylserine (PSe) and phosphatidylinositol (PI) in the inner layer (Fig. 16). As such, the membrane is said to be asymmetric. This feature allows proper functioning of many cellular processes: protein localization and orientation, cell recognition, membrane permeability, membrane curvature and shape *etc.*^{87,160–163} Asymmetric vesicles are particularly attractive as cell membrane models for the orientation of membrane proteins. Mimicking the asymmetry of the plasma membrane was found to be challenging firstly for generating them and maintaining them due to transversal diffusion giving a limited lifetime to the asymmetry.^{87,115,161} This flip-flop was found to be variable and in the range of a few hours to 24 h and influenced by contamination from self-assembly and use of bulky fluorescent tags or spin labels used to measure the diffusion rate like (2,2,6,6-tetramethylpiperidin-1-yl)oxyl (TEMPO).

Lipid-only asymmetric liposomes have been described as they naturally entropically favor specific lateral and transversal lipid phase separation due to the intrinsic properties of the lipid such as the angle and charge of hydrocarbon chains, as depicted in Fig. 16.^{161,163–165} However, controlling asymmetry in liposomes to understand the principles behind their distribution or achieve unnatural distributions can mostly only be achieved by tuning the chemical composition of their chemical environment for example through pH modulation.¹⁶⁴ Phosphatidylglycerol (PG) could selectively (80–90%) rearrange to equilibrate on the inner layer of the SUVs (100 nm) when the intravesicular environment was basic and on the outer leaflet when the intravesicular

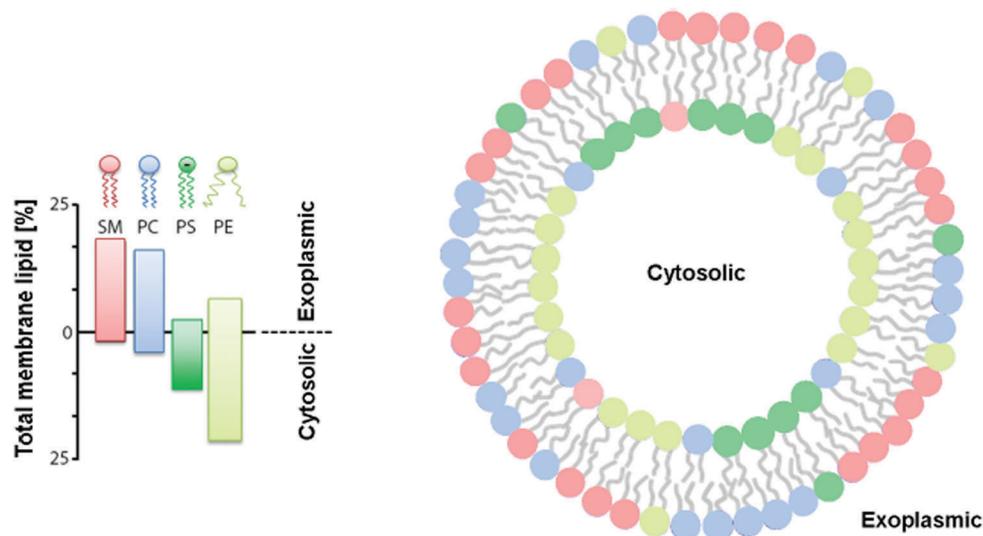


Fig. 16 Distribution of phospholipids in human red blood cells. Reproduced with permission from D. Marquardt, B. Geier and G. Pabst, *Membranes*, 2015, 5(2), 180–196 (<http://www.mdpi.com/2077-0375/5/2/180>).¹⁶¹



environment was acidic while PE remained equally distributed. The basis behind such preferential tuning is that PGs are acidic and thus rearrange on the proton depleted side of the membrane as a means to equilibrate the transmembrane proton gradient.

Inducing asymmetry within the vesicle membrane can also be achieved due to membrane curvature. Tuning the chemical composition of the amphiphiles can dictate a preferred orientation of the amphiphile within the membrane.^{166–169} This is particularly advantageous for polymersomes as the block copolymer length is easily adjustable synthetically in contrast to liposomes as discussed in Section II d. Mixing two diblock copolymers with differently sized hydrophilic blocks could result in a hydrophilic block with the smaller volume in the inner membrane and the larger one on the outer membrane for example with $PS_{295}\text{-}b\text{-}PAA_n$ ($n = 12$ or 74) polymersomes (100 nm).¹⁶⁷ More frequently ABC triblock copolymers are described to form asymmetric membranes presumably as they are easier to design and control than diblock copolymers. Meier and his group investigated two similar ABC polymers $PEO\text{-}b\text{-}PDMS\text{-}b\text{-}PMOXA$ ¹⁷⁰ and $PEO\text{-}b\text{-}PCL\text{-}b\text{-}PMOXA$ ³⁵ which formed asymmetric SUVs (~ 100 nm) and GUVs (~ 5 μm) respectively. Asymmetry is achieved by tuning the block length. $PEO_{45}\text{-}b\text{-}PDMS_{65}\text{-}b\text{-}PMOXA_{346}$ was determined to form vesicles with the PMOXA block oriented on the outside and

$PEO_{40}\text{-}b\text{-}PDMS_{45}\text{-}b\text{-}PMOXA_{97}$ with the reverse arrangement. In analogy to liposomes, Liu and Eisenberg described asymmetric vesicles (120 nm) which can switch orientation depending on pH changes, impeccably exemplifying the influence of the chemical composition on asymmetry.¹⁷¹ $PAA_{26}\text{-}b\text{-}PS_{890}\text{-}b\text{-}P4VP_{40}$ (P4VP: poly-(4-vinyl pyridine)) has two relatively short hydrophilic blocks and thus their orientation in polymersomes cannot be dictated by membrane curvature alone. Both PAA and P4VP are however pH sensitive. At low pH values, the blocks are protonated, thus PAA is neutrally charged, whereas P4VP is positively charged, while at high pH they are deprotonated, PAA is negatively charged and P4VP is neutral. When the blocks carry charges, intra- and intermolecular repulsive interactions are exhibited. Hence, the charged block swells and effectively has a larger volume than the neutral one. Following the membrane curvature restriction, the charged block would be favored on the outer layer of the polymersome membrane (Fig. 17).

Generating asymmetry in lipid and polymer membranes has recently been largely simplified by solvent double emulsion methodologies such as microfluidics or emulsion phase transfer which allows the highly controlled construction of each layer in the vesicular bilayer separately regardless of the nature of the amphiphiles.^{86,87,90,162,172,173} Weitz and coworkers described the self-assembly of asymmetric liposomes (0.5–2 μm) by emulsion phase transfer with POPC (1-palmitoyl-2-oleoyl-*sn*-glycero-3-phosphocholine) and POPS (1-palmitoyl-2-oleoyl-*sn*-glycero-3-phospho-L-serine).⁸⁶ The aliphatic chains of the lipids are the same but POPS has a small anionic head group and POPC a large zwitterionic head group. Both asymmetric vesicles (with POPC or POPS on the inner layer) were generated despite the unfavoured curvature of POPC in the inner layer. In an interesting recent example, Peyret *et al.* developed hybrid POPC/PB-*b*-PEO asymmetric GUVs (~ 10 μm) using emulsion-centrifugation (Fig. 18).⁸⁷ A polymer-stabilized sucrose droplet is initially formed and poured into a lipid-stabilized glucose_(aq)/toluene interface, leading to the formation of GUVs with an outer lipid monolayer and an inner polymer monolayer. The reverse hybrid asymmetric vesicles were more difficult to achieve due to the low stability of the POPC-droplets. Interestingly, polymer only asymmetric membranes have yet to be synthesized by these methods.

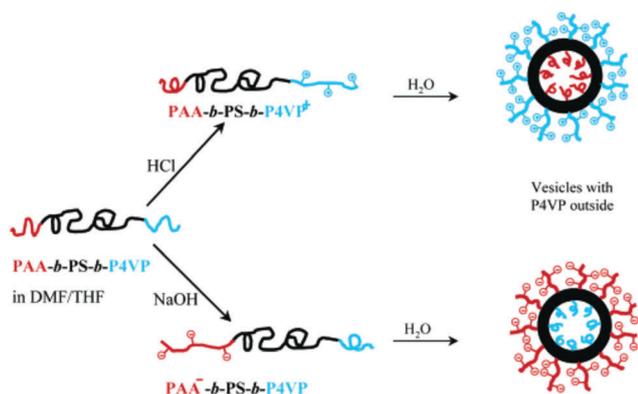


Fig. 17 Reversible asymmetric polymersome from $PAA_{26}\text{-}b\text{-}PS_{890}\text{-}b\text{-}P4VP_{40}$. Reprinted with permission from (F. Liu and A. Eisenberg, *J. Am. Chem. Soc.*, 2003, **125**(49), 15059–15064).¹⁷¹ Copyright (2003) American Chemical Society. <https://pubs.acs.org/doi/abs/10.1021/ja038142r>.

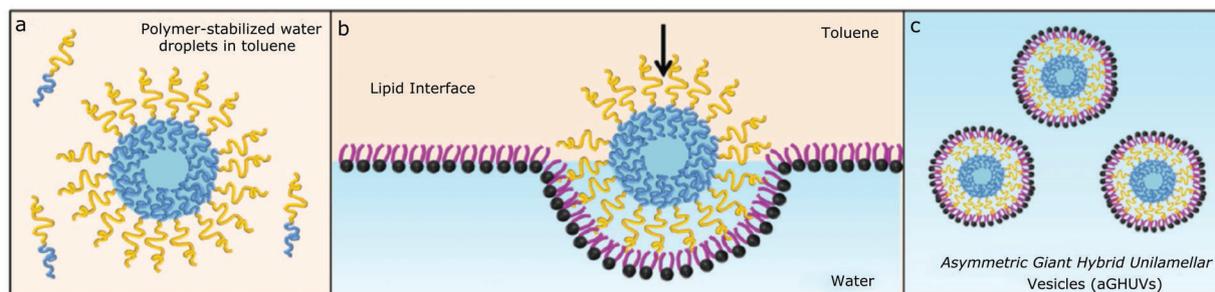


Fig. 18 Preparation of asymmetric hybrid GUVs by emulsion-centrifugation. (a) Preparation of sucrose filled polymer-stabilized droplets in toluene. (b) Crossing the amphiphile-stabilized glucose/toluene interface helped by centrifugation, forming (c) asymmetric hybrid vesicles. Adapted from A. Peyret, E. Ibarboure, J.-F. Le Meins and S. Lecommandoux, *Adv. Sci.*, 2017, **5**(1) with permission from Wiley. (<https://onlinelibrary.wiley.com/doi/abs/10.1002/adv.201700453>).⁸⁷



As previously discussed, a major drawback of microfluidics and droplet emulsion transfer methodologies is the unassessed contamination of the membrane with organic solvents which alter the asymmetric membrane's properties such as in this case the transversal diffusion of the amphiphile.^{89,115,161} With an emphasis on reducing solvent impurities, Takeuchi and coworkers developed lipidic GUVs ($\sim 4 \mu\text{m}$) formed from the non-uniform cleavage of an asymmetric DOPC/DOPS (1,2-dioleoyl-*sn*-glycero-3-phosphocholine/1,2-dioleoyl-*sn*-glycero-3-phosphoserine) planar lipid bilayer by a pulsed microfluidic jet flow.¹¹⁵ Sinusoidal undulations cause deformation in the planar membrane generating two types of GUVs. The larger ones ($150 \mu\text{m}$) contained a large amount of solvent while the smaller ones ($3\text{--}20 \mu\text{m}$) did not. The authors showed that transversal diffusion is inhibited by the presence of *n*-decane in the vesicular bilayer of the larger GUVs while the smaller ones efficiently mimic the transversal diffusion found in apoptotic cells.

Lateral heterogeneity. Asymmetry in membranes commonly refers to the differentiation between the inner and outer layers of the vesicles. Nevertheless, even symmetrical membranes (seemingly identical inner and outer layer composition) can be non-uniform. In cell membranes, small domains ($50\text{--}500 \text{nm}$) rich in glycopingolipids and cholesterol can be found.¹⁷⁵ These so called "lipid rafts" were determined to have many biological purposes such as intracellular signaling, membrane trafficking, uptake and notably to dictate protein function.^{8,176} It is still however not well understood how they are formed and behave.^{175,177} The study of their formation, properties and dynamic rearrangement would contribute to the understanding of domains in the plasma membrane.^{8,176,178} In synthetic membranes, domains can be nano- or micron-sized. Micron-sized domains in GUVs are easily observed by optical microscopy and thus lateral heterogeneity studies typically focus on these domains in GUVs. Nanodomains ($< 300 \text{nm}$) are too small for detection^{8,72} using an optical microscope but can however be detected by Förster resonance energy transfer (FRET) or dual-color fluorescence cross-correlation spectroscopy (DC-FCCS).^{72,177} For example, nanodomain formation in hybrid LUVs (200nm) formed from PDMS-*b*-PMOXA and 1,2-dimyristoyl-*sn*-glycero-3-phosphocholine (DMPC) has been described by Winzen *et al.* and analyzed by DC-FCCS.¹¹

Mixtures of lipids are frequently used to generate liposomes more physiologically relevant to model membranes.^{114,179–182} These mixed lipid membranes can phase separate, due to differences in the melting temperature of the lipids, their aliphatic chains (> 2 carbons) or headgroups.^{8,127} It is not well understood what dictates transversal *versus* lateral heterogeneity in lipid membranes. Moreover, beyond the nature of the lipids themselves, mixing of lipids is also greatly influenced by temperatures, impurities (even in trace amounts), lipid ratios and lipid compositions in each vesicle. An increasing number of model membranes using liposomes have shown that lipid domains can be formed when using the appropriate lipids and ratios.^{176,183–186} Because of its role in lipid rafts, cholesterol phase separation has also been studied extensively. For example, DOPC/DPPC (50/50) and DOPC/DSPC (50/50) (DSPC 1,2-distearoyl-*sn*-glycero-3-phosphocholine) GUVs ($30 \mu\text{m}$) both exhibited

gel-fluid phase separation but by adding 10 mol% of cholesterol, the vesicles became homogeneous (domains disappeared) and with 20 mol% cholesterol, new fluid domains ($1\text{--}20 \mu\text{m}$) were formed (liquid ordered) which differ from the original domains (liquid disordered). For cholesterol ≥ 50 mol%, domains were no longer visualized by optical microscopy. The authors also showed that cholesterol affects lipid fluidity in a similar way for DPPC and DSPC and more strongly between cholesterol and SM. The driving force for lipid raft formation in the plasma membrane might be due to hydrogen bonding between SM and cholesterol, which was not observed with PC-cholesterol. The phase diagrams of ternary (and even quaternary) mixtures are now extensively studied on GUVs.^{187–189}

Hybrid polymer-only vesicles are rarer.^{190–192} In the last decade, the group of Battaglia has specialized in studying the surface topology of LUVs using mixed polymers. For example, diverse ratios of PMPC₂₅-*b*-PDPA₇₀ mixed with PEO₂₃-*b*-PDPA₁₅ were analyzed regarding their domain formation in polymersomes (200nm) and their internalization in cells.¹⁹¹ The authors showed that by mixing 25 mol% PEO-*b*-PDPA and 75 mol% PMPC-*b*-PDPA, PEO-rich domains become visible. As a result, the PMPC block shrinks which affects the overall cell uptake kinetics of the polymersomes. Fewer hybrid polymersomes are internalized than the pure PMPC polymersomes suggesting that PMPC chains have a higher affinity for cell membranes than the PEO chains. When using the opposite ratio (75 mol% PEO-*b*-PDPA and 25 mol% PMPC-*b*-PDPA), PMPC domains are formed within a PEO matrix and surprisingly its uptake kinetics is very similar to that of the pure PMPC polymersomes and faster uptake kinetics for larger polymersomes. Thus, the presence of domains and their size affect the cellular uptake of these polymersomes. Discher and coworkers reported that the PAA block of PAA-*b*-PBO can be cross-linked by calcium or copper chelators in hybrid polymersome GUVs ($10\text{--}20 \mu\text{m}$) (Fig. 19).¹⁹⁰ When PAA-*b*-PBO was mixed in different ratios with fluorescently labelled PB-*b*-PEO in the presence of Ca^{2+} , demixing from the neutral amphiphiles was observed in accordance with the PAA-*b*-PBO and PB-*b*-PEO ratios. The presence of the calcium chelator even allowed the thermodynamic stability of the domains.

In terms of polymer-lipid hybrid vesicles, domain formation is frequent as the chemical incompatibilities between lipids and polymer are greater than those for their respective identical amphiphiles (compare Section IIe). Most hybrid vesicles are made from diblock, triblock, or grafted polymers with PDMS, PB, or PIB (polyisobutylene) as the hydrophobic block and PEO or PMOXA as the hydrophilic blocks. In terms of lipids, PE or PC is almost exclusively used.⁸ Whether the membrane is homogeneous, heterogeneous or completely separated is difficult to determine pre-emptively and is highly sensitive to conditions. Chemin *et al.* described PIB-*b*-PEO and DPPC to form a heterogeneous membrane at a narrow ratio of 20–28% of block copolymer when using PIB₈₇-*b*-PEO₁₇ and in contrast only homogeneous vesicles when using PIB₃₇-*b*-PEO₄₈ regardless of the lipid/polymer molar composition.¹²⁷ Nam *et al.* also showed that when using POPC (> 30 mol%) and PB₄₆-*b*-PEO₃₀ (> 70 mol%) HLPs are obtained; POPC (35–65 mol%) and PB₄₆-*b*-PEO₃₀



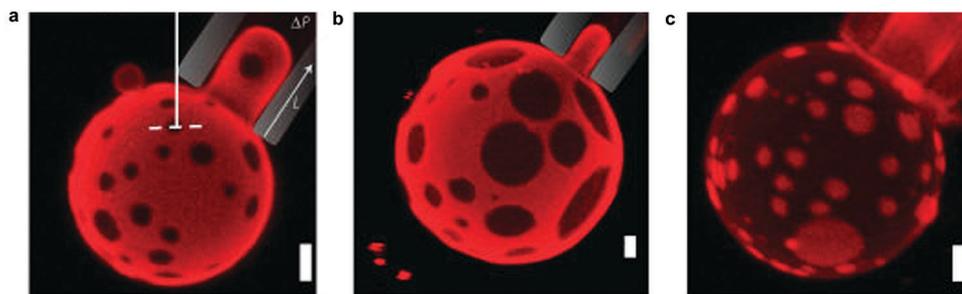


Fig. 19 Lateral phase separation of PAA-*b*-PBO and PB-*b*-PEO with 25% (a), 50% (b) and 75% (c) of PAA-*b*-PBO. Scale bar 2 μm . Reprinted with permission from Springer Customer Service Centre GmbH: Springer Nature, D. A. Christian, A. Tian, W. G. Ellenbroek, I. Levental, K. Rajagopal, P. A. Janmey, A. J. Liu, T. Baumgart and D. E. Discher, Spotted vesicles, striped micelles and Janus assemblies induced by ligand binding, *Nat. Mater.*, Copyright (2009).¹⁹⁰

(35–65 mol%) give no vesicles; and POPC (70–100 mol%) and PB₄₆-*b*-PEO₃₀ (0–30 mol%) give a mixture of liposomes and HLPs.⁷¹ Similar to lipid-only and polymer-only hybrid vesicles, domain formation in polymer-hybrid vesicles can also be controlled by external stimuli or membrane composition. Beales and coworkers could control domain formation and properties of PB-*b*-PEO and DLPC (1,2-dilauroyl-*sn*-glycero-3-phosphocholine), DCPC (1,2-dicetyl-*sn*-glycero-3-phosphocholine) or DPPC GUVs (20 μm) *via* thermally driven phase separation, inclusion of cholesterol or macromolecular additives (surfactants, enzymes, cyclodextrins).¹⁷⁴ When cooled slowly, only a few large phospholipid-rich domains are formed, while fast cooling generates numerous small domains.

Domain instability in heterogeneous membranes may result in equilibration by fission into separate vesicles.^{72,127} However, this challenge is only observed for polymer-lipid hybrid vesicles presumably due to the much greater chemical differences between lipids and polymers than lipid-lipid or the few polymer-polymer vesicles that have been tested. The formation of stable domains in GUVs is however essential for protocells. Using PDMS-*g*-(PEO)₂ and DPPC, Dao *et al.* described the formation of stable nanodomains in hybrid polymer/lipid SUVs (100 nm).¹⁷⁷ Lipid rich or polymer rich domains co-existed in the same vesicles. Domain formation in lipid/polymer hybrid membranes was also obtained in GUVs. Unfortunately, in the case of GUVs fission of the vesicles was observed after a few hours resulting in pure liposomes and polymersomes as the most entropically stable phase of the amphiphile self-assembly.¹²⁷ The different outcome between SUVs and GUVs suggests that the curvature of the membrane greatly affects domain stabilization and stable micro-domain formation in polymer-lipid GUVs remains a challenge.

Membrane transport. Plasma membranes use many proteins and specialized structures in order to constantly regulate membrane transport. A cell could not survive without any signaling to and from its environment. Thus for cell mimicking, complete hermetic compartmentalization is undesirable. The diffusion of small molecules through the membranes is generally difficult to achieve with polymersomes due to their low permeability. In contrast, liposome membranes are highly fluid vesicles but to such an extent that retaining molecules can be challenging (non-specific and high molecular cut off) (see Section IIb for more details). Thus, the ideal cell membrane mimics lie in the silver

lining between permeable and hermetic compartments. One solution to this problem has been found in stimuli-responsive vesicles (Section III). However, such compartments only change permeability upon a trigger and frequently become highly porous or even disassemble. Such vesicles are not good models for protocells, which require constant controlled membrane transport. Integration of membrane channel proteins has been found to be a favorable solution for controlled active transport of small molecules particularly in hermetic polymersomes. Initial models use channel proteins, which lack specificity, but new examples are coming to light, which can tune the permeability of the synthetic membrane with great specificity mimicking cellular membrane transport.^{193–198} Unfortunately, many examples of membrane modifications are only shown on SUVs despite the parallel drawn to living cells.

Bassereau and her group worked extensively on the incorporation of transmembrane proteins such as Ca²⁺-ATPsynthase, the photoactivable proton pump bacteriorhodopsin (BR) or the voltage-dependent K⁺ channel, KvAP, into liposome GUVs.^{199–201} These advanced liposome GUVs (10–100 μm) were generated by the initial detergent-mediated reconstitution of solubilized membrane proteins into proteoliposome SUVs (100–200 nm). The SUVs were then fused by electroformation or gel-assisted swelling. The proteins could be oriented into the membrane by a transfer mechanism from the mixed micelles to the lipid membrane (more details of this method are given below). Thus the proteins retained their biological activity for example when using BR and valinomycin, another K⁺ selective carrier, large transmembrane pH gradients can be generated upon light stimulus (Fig. 20a and b).²⁰⁰ Internal pH variations could be measured based on the fluorescence of pyranine, a photosensitive dye. In the presence of BR alone, a positive transmembrane electrical potential is generated upon light stimulus, retro-inhibiting the proton pumping. The GUVs' permeability to protons was low (Fig. 20c). Valinomycin compensates for the electrical potential by releasing K⁺ externally, allowing for the acidification of the vesicles.

A major challenge of reconstitution of most transmembrane proteins into vesicles is their orientation.^{202,203} In 2017, Mavelli and coworkers reconstituted a photosynthetic reaction center (RC), an integral membrane protein, into liposome GUVs (20 μm) by droplet emulsion transfer.²⁰² RC was initially extracted



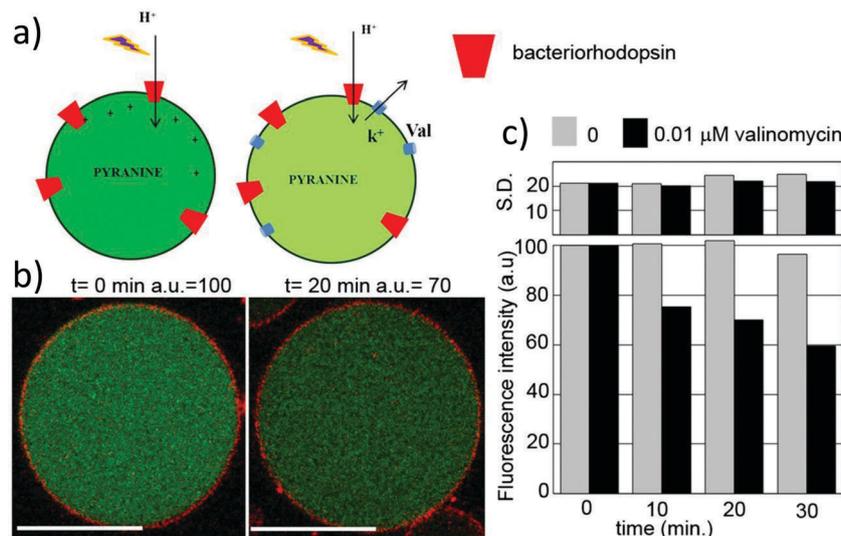


Fig. 20 Incorporation of transmembrane proteins into liposome GUVs. (A and B) Light-induced proton pumping induces an internal acidification of GUVs. (C) Normalized pyranine fluorescence intensity in the presence or absence of valinomycin. Reprinted with permission from M. Dezi, A. Di Cicco, P. Bassereau and D. Lévy, Detergent-mediated incorporation of transmembrane proteins in giant unilamellar vesicles with controlled physiological contents. *Proc. Natl. Acad. Sci. U. S. A.*, 2013, **110**(18), 7276–7281.²⁰⁰

from purple bacteria using lauryldimethylamine *N*-oxide (LDAO) in aqueous solution generating micellar-stabilized RC. The RC-micelles were then emulsified in a lipid-rich oil phase yielding w/o droplets, which were subsequently transferred into an oil/water interphase forming the GUVs by centrifugation. The RC-micelles are thought to deliver their cargo during the w/o droplet phase driven by hydrophobic interactions. The major strength of this study comes from the asymmetry of the RC-micelles in terms of hydrophilicity/hydrophobicity, which guides the orientation of the membrane yielding 90% physiological orientation of the transmembrane proteins into the GUVs. The RC@lipid-GUVs were then used to generate a light-induced pH gradient across the membrane.

In terms of polymersomes, in 2016, Messager *et al.* described the use of membrane-spanning DNA nanopores NP-3C in PMPC₂₅-*b*-PDPA₇₂ polymersomes (100–200 nm).¹⁹⁷ The NP-3C were fabricated through the self-assembly of oligonucleotides and the pore diameters could be customized. The polymersomes exhibited size-dependent permeability: substrates can be freely transported in and out of the polymersomes while larger enzymes were retained. As a proof of principle, the hydrolytic enzyme trypsin was encapsulated and the substrate peptide B-NAR-AMC (Boc-Gln-Ala-Arg-7-amido-4-methylcoumarin) was added outside the vesicle. Trypsin cleaved the substrate B-NAR-AMC to release the fluorescent product AMC that could then be quantified. In 2017, Castiglione and coworkers described the use of mutant Outer Membrane Protein F (OmpF G119D) channels in a PMOXA-*b*-PDMS-*b*-PMOXA nanoreactor (110 nm) used for an incompatible multi-enzyme cascade (Fig. 21).²⁹ One enzyme (*N*-acyl-D-glucosamine 2-epimerase (AGE)) and its allosteric activator ATP were encapsulated inside the polymersome while the other enzymes of the cascade (*N*-acetylneuraminase lyase (NAL) and CMP-sialic acid synthetase (CSS)) were immobilized on the outer surface of the vesicle as well as their respective cofactors

(pyruvate and cytidine triphosphate (CTP)). The cascade is normally incompatible as AGE is strongly inhibited by CTP. The OmpF G119D had the advantage over the wildtype OmpF that the exchange of the glycine at position 119 to aspartate introduces a negative charge in the channel, increasing cation selectivity. As a result, selective massive transport of the neutrally charged substrates (*N*-acetylglucosamine (GlcNAc) and *N*-acetylmannosamine (ManNAc)) remained possible while the problematic negatively charged allosteric cofactors (CTP and ATP) were segregated outside and inside the polymersome respectively proving the value of compartmentalization. In 2018, from the same perspective Palivan and coworkers used extensively modified OmpF notably by adding cysteine residues to make the membrane channel redox-responsive.²⁰⁴ Impressively the author used these modified OmpF polymersome SUVs (100 nm) to generate artificial organelles that could be integrated *in vitro* in HeLa cells and *in vivo* in zebrafish embryos, exemplifying the potential of cell mimicking in medicine and materials science.

In 2018, van Hest and coworkers described breathing polymersomes (100 nm) that could also be used to switch the membrane permeability on and off dependent on the pH.²⁰⁵ Upon intravesicular addition of acidic fuels (HCl and urea) and a substrate, the polymersomes expanded and horseradish peroxidase (HRP) could react with its substrate. Simultaneously, modification of urea into ammonia by urease effectively lowered the system's pH-value and thus switched off the permeability of the nanoreactor once a critical pH-value was reached. The systems lack transport selectivity compared to transmembrane proteins, but clearly demonstrated the feedback induced temporal control of the HRP's activity based on controllable permeability of the polymersome membranes.

In cells, small molecules are not exclusively transported *via* channels spanning the membrane. Many ions, molecules and



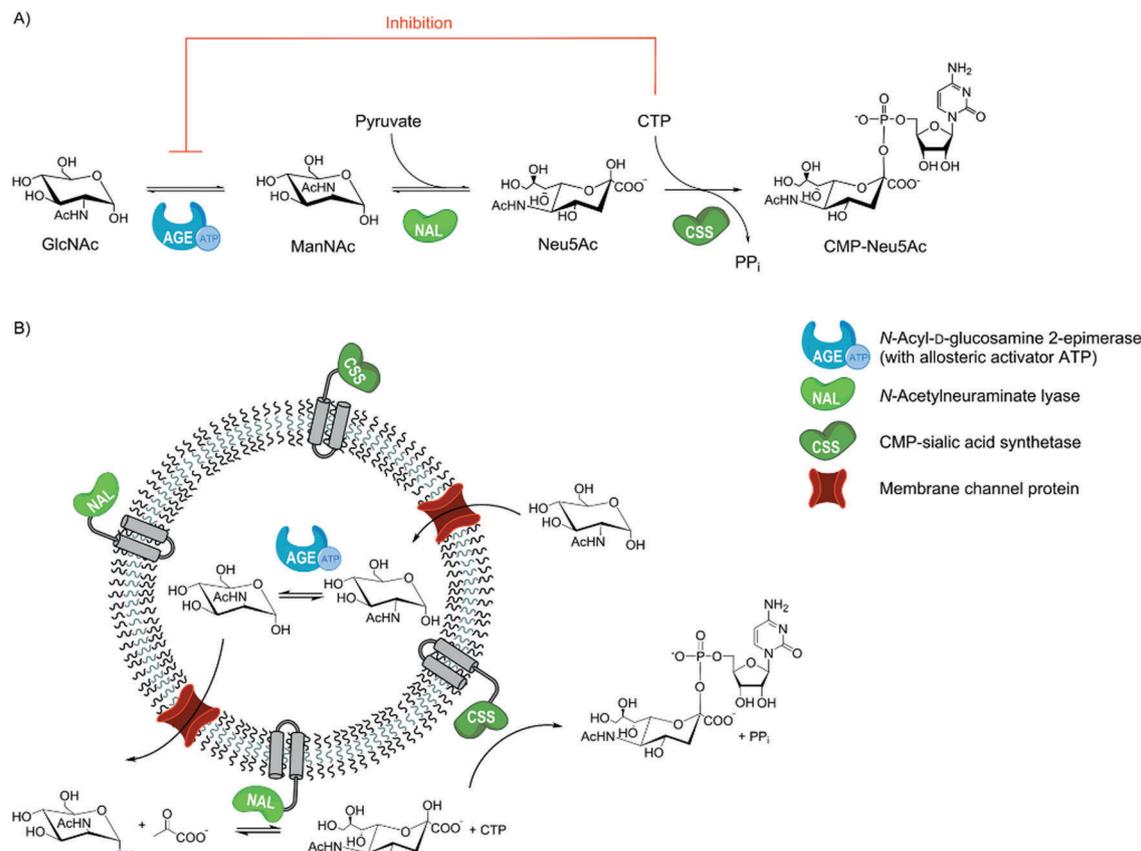


Fig. 21 Multi-enzyme cascade carried out in a polymersome nanoreactor. Reprinted with permission from L. Klermund, S. T. Poschenrieder and K. Castiglione, *ACS Catal.*, 2017, 7(6), 3900–3904.²⁹ Copyright (2017) American Chemical Society. <https://pubs.acs.org/doi/abs/10.1021/acscatal.7b00776>.

even viruses or bacteria are selectively transported through the membrane by endocytosis. Endocytosis is an important mechanism of cells but due to its complexity it is not well understood. Landfester and coworkers described the uptake of PS (16 nm) or SiO₂ nanoparticles (NPs) (14 to 57 nm) into PDMS-*b*-PMOXA polymersomes (~100 nm) (Fig. 22).^{206,207} These systems allow

us to understand transmembrane transport in general as a physical process in the absence of any protein or supplementary energy and also how the uptake of engineered nanoparticles can be optimized for biomedical applications. The NPs are encapsulated in vesicles in four steps (Fig. 22d): (1) recognition of the NPs at the vesicle surface, (2) engulfing initiation, (3) entire coverage of

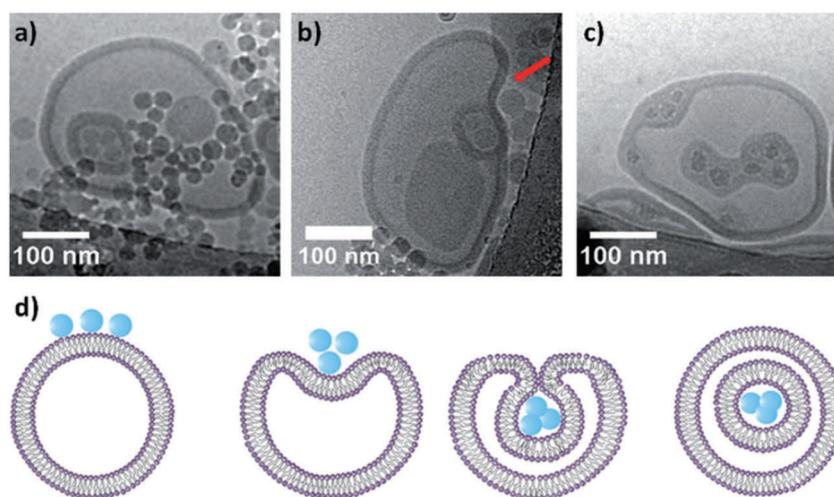


Fig. 22 Cryo-TEM micrographs of the encapsulation of PS and SiO₂ nanoparticles into polymersomes. Reprinted with permission from Jaskiewicz, A. Larsen, I. Lieberwirth, K. Koyanov, W. Meier, G. Fytas, A. Kroeger and K. Landfester, *Angew. Chem., Int. Ed.*, 2012, 51, 4613–4617. Copyright (2012) Wiley.²⁰⁶



the NPs by the membrane, and (4) complete internalization. The internalization of these particles was proven to be faster for SiO₂ particles (8 min) compared to PS nanoparticles (15 min), and to be strongly concentration and size dependent. Similarly, Meinel *et al.* studied the induced phagocytic uptake of PS particles (1 μm) in egg-PC liposome GUVs (20 μm).²⁰⁸ In order to engulf particles, the GUV needs to be significantly deformed and this was induced by optical tweezers. The optical tweezers has the advantage of applying controlled 3D force (femto-Newton) without mechanical contact.

Membrane molecular recognition. Molecular recognition in the extracellular environment by cell receptors is crucial for cell signaling and is involved in processes as broad as cell–cell communication, immune response, hormonal cellular response, receptor mediated endocytosis, virus' infection *etc.* For example, the major histocompatibility complex (MHC) expression on the extracellular-side of antigen-presenting cells is essential for the molecular recognition of pathogens.^{209–211} Therefore it is not surprising that decorating the outer membrane of vesicles has been vastly described for targeted drug delivery and release.^{212–218} For example, PEO-*b*-PCL polymersomes (100 nm) conjugated with mouse-anti-rat monoclonal antibody OX26 (OX26-PO) were used to deliver drugs to the brain as the antibody can bind to a specific receptor which initiates transcytosis through the blood–brain barrier.²¹⁹ However from the same perspective, surface selective recognition of small molecules by receptor–ligand binding or docking is thus an important aspect of cell mimicry. Surface receptor mimicry has nevertheless been poorly studied on GUVs from an artificial cell perspective at the expense of medical applications and to date only few examples exist of molecular recognition from a cellular mimic perspective^{220–223} and even less on GUVs.^{97,106,224–226}

Terminally-modified liposomes for recognition or docking are rare as difficult to modify. Thus liposomes can be decorated by incorporating surface proteins, like MHC, in a similar fashion by which transmembrane transporters are added to the membrane or by PEGylation as discussed in Section IV but has always been done from a medical perspective. For block copolymers, the addition of small moieties at their terminal end has a small impact on their self-assembly properties compared to liposomes. Commonly, compatible click moieties (azide/alkyne or maleimide/thiol), or Michael addition partners (carboxyl and amines) are often implemented on the terminal position of the block copolymers to add commercially available entities pre or post self-assembly to the polymersomes. Biotin is a commonly used ligand for biological functionalization due to its high binding affinity for the proteins streptavidin and avidin. In 2018, Landfester and coworkers used acrylate functionalized PB-*b*-PEO GUVs (20 μm) to conjugate amine-functionalized biotin post-assembly.¹⁰⁶ These biotinylated GUVs were used to mimic the adhesive properties of cells by characterizing the interaction of polymersomes with neutravidin-coated glass surfaces. Similarly, Broz *et al.* showed that biotin-functionalized PMOXA-*b*-PDMS-*b*-PMOXA polymersomes (100–250 nm) would bind to streptavidin and oligonucleotide polyguanylic acid (polyG), which specifically targets the macrophage A1 scavenger receptor and thus could be

used for diagnostic or therapeutic use.²²⁷ Kubilis *et al.* described glycopolymer (p(NβGluEAM₅-*b*-BA₅₀)) protocells (20 μm) capable of interacting with lectin-functionalized particles.⁹⁷ Carbohydrate–lectin interaction is a common means of recognition between extracellular ligands and cells and notably initiates processes such as inflammatory cascade, fertilization, cancer cell metastasis and virus docking. The glycol-GUVs were shown to bind to glucose-specific lectin Concanavalin A (Con A)-functionalized polystyrene beads and not carboxylate-modified PS beads. The authors made the parallel of this specific membrane recognition to the binding of virus to mammalian cells for subsequent uptake and infection.

(b) Compartmentalized processes

We have just discussed how the membranes of vesicles themselves have been modified to mimic the cell plasma membrane. The simplest membrane can also be used to encapsulate processes with increasing complexity just like living cells. Compartmentalization of modules allows protection from the extra-vesicular environment, especially when dealing with incompatible processes, concentration of diverse functionality in a single entity rather than in bulk or diluted in solution, and is the basis of all cellular processes (metabolism, division, growth, communication, motility *etc.*).^{90,107,154}

Multi-compartmentalization. Multi-compartmentalization is the quintessence of cells: small compartments (organelles) with a defined function and design encapsulated into a protective membrane. Thus building multi-layered systems has been extensively studied and it allows a great variety of combinations.^{105,146,228–235}

In 2014, Peters *et al.* developed a fully synthetic polymersomes-in-polymersome multi-compartmentalized system mimicking cell hierarchical construct (organelles in cells).²²⁹ The authors chose a cascade reaction involving the initial oxidation of non-fluorescent substrate **1** by the phenylacetone monooxygenase (PAMO) and its NADPH cofactor followed by ester hydrolysis by *Candida antarctica* lipase B (CalB) or alcalase yielding an alcohol (Fig. 23). The alcohol is then oxidized to an aldehyde using alcohol dehydrogenase (ADH) and NAD⁺ and undergoes spontaneous β-elimination to yield fluorescent resorufin **5**. Alcalase is sometimes used as a substitute for CalB because this protease is incompatible with other enzymes and a successful reaction cascade proves the importance of compartmentalization. In order to use this cascade to mimic a cell, CalB/alcalase and ADH were encapsulated independently in PS₄₀-*b*-PIAT₅₀ (PIAT: poly(3-(isocyanato-L-alanyl-aminoethyl)thiophene)) polymersomes (187 nm and 318 nm respectively). The CalB/alcalase nanoreactors and ADH nanoreactors were then encapsulated in a large PB-*b*-PEO polymersome (20–60 μm) along with substrate **1**, PAMO and NAD⁺. The cascade conversion was observed by confocal fluorescence microscopy by following the quantitative increase of fluorescence over time. The fluorescence was confined to inner compartments as ADH interacts electrostatically with **5** preventing its diffusion out of the nanoreactors. Similarly, Voit and co-workers also studied enzymatic cascade reactions in pH-switchable adamantyl-functionalized polymersomes as artificial organelles into large polymersomes (1 μm) possessing temperature- and pH-responsiveness and PEG surface functionalization.²³¹



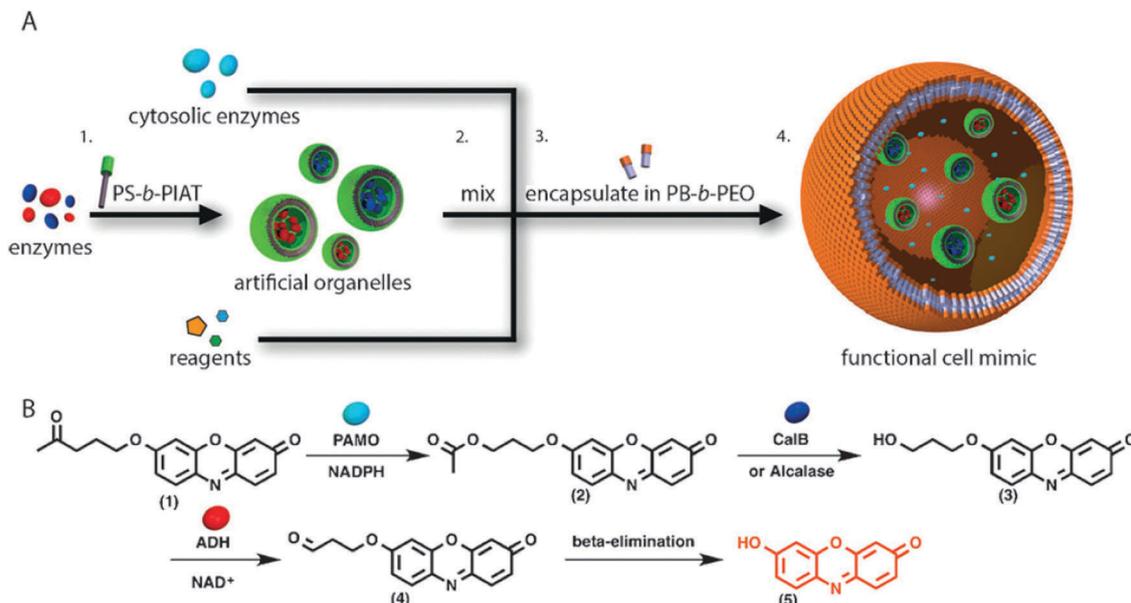


Fig. 23 Reaction cascade performed in a complex multi-hierarchical protocell. Reprinted with permission from R. J. Peters, M. Marguet, S. Marais, M. W. Fraaije, J. C. van Hest and S. Lecommandoux, *Angew. Chem., Int. Ed.*, 2014, **53**, 146–150. Copyright (2013) Wiley.²²⁹

Another example of multi-compartmentalization was demonstrated in 2017 by generating hybrid liposomes-in-polymerosomes.¹⁴⁶ POPC, DMPC, diC15-PC (1,2-dipentadecanoyl-*sn*-glycero-3-phosphocholine), or DPPC SUVs (<400 nm) were first prepared by film hydration and subsequently used as the inner emulsion droplet for the emulsion-centrifugation method to form PB-*b*-PEO GUVs (20 μ m). The loading and co-loading of the different liposomes in polymerosome GUVs was established. Moreover because lipids have a different T_m , hydrophilic cargo can selectively be released from SUVs when the liposomes are subjected to temperature above their T_m . The temperature-controlled release of first methylene blue from diC15-PC SUVs at 37 °C and then fluorescein from DPPC SUVs at 45 °C into the polymerosome GUV lumen was demonstrated, exemplifying the significant progress made in recent years to build ever-more complex systems with different 'organelles' within a single GUV.

As a matter of fact, artificial organelles have even been implemented in living cells, going beyond the generation of protocells for medical purposes.²³³ In this study for example, the authors generate liposomes/fluorescent gold nanoclusters in a polymer shell (capsosomes, 2–3 μ m) which are readily taken up by macrophages. The capsosomes are composed of both glucose oxidase (GOx) liposomes and HRP liposomes effectively converting glucose into the fluorescent resorufin, retaining its activity within the macrophages. In contrast, Elani *et al.* attempted to bridge the gap between artificial protocells and biological cells by building living-cells-in-liposomes.²³⁵ By microfluidics, a variety of human cells as well as *E. coli* could be encapsulated into POPC GUVs (70 μ m). A three step cascade converting lactose into fluorescent resorufin was subsequently demonstrated by coupling the cellular pathways with non-cellular free enzymes. Furthermore, the liposomes were also demonstrated to provide an effective protective barrier for the living cells against toxins such as Cu²⁺

thus demonstrating the advancement artificial systems can bring to living cells.

ATP regeneration. ATP is arguably one of the most essential features of a cell as it is the energy source for all biological processes.²³⁶ In the cell, ATP is generated by F₀F₁-ATP synthase in the cytosol, mitochondria, or chloroplast. This transmembrane enzyme catalyzes the otherwise unfavorable generation of ATP from ADP (adenosine diphosphate) and inorganic phosphate using chemical energy, a proton gradient across the membrane. It is thus not surprising that many studies focus on the reconstitution of ATP-synthase in lipid, polymer or hybrid vesicles.^{13,236–241}

In 2005, Choi and Montemagno designed one of the first biomimetic polymerosomes: PETOz-*b*-PDMS-*b*-PETOz (PETOz: poly(2-ethyl-2-oxazo-line)) SUVs (<200 nm) capable of generating ATP by coupling the activity of photoactive BR and F₀F₁-ATP synthase (Fig. 24a).²³⁸ BR is a light-driven proton pump, which creates a proton gradient across the polymerosome. This gradient is then used by the ATP-synthase to synthesize ATP. Montemagno's model proved that polymerosomes can be used for biosynthesis through the coupled effect of two transmembrane proteins in a single vesicle and thus that a fundamental cellular module can be recreated in a complete synthetic approach towards the generation of functional synthetic organelles.

In 2017, from the same perspective, Sundmacher and coworkers detailed the generation of artificial mitochondria by coupling ATP-synthase with cytochrome bo3 quinol oxidase into PDMS-*g*-PEO and hybrid PDMS-*g*-PEO/POPC or soy PC SUVs (100 nm) and GUVs (50 μ m) (Fig. 24b).¹³ Like BR, bo3 acts as a proton pump by oxidizing ubiquinol and generating water from O₂, driving the production of ATP into the outer media. PDMS-*g*-PEO was described as particularly advantageous over other polymers as generating only 5 nm wide membranes and also having a fluidity comparable to liposomes, contrary to other



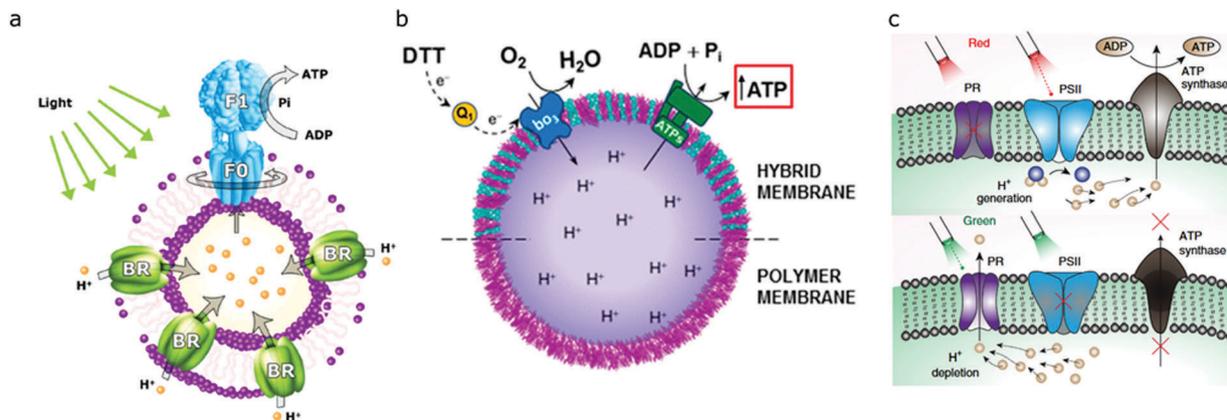


Fig. 24 ATP-synthase based proteovesicles as cellular mimics for the generation of ATP using an electrochemical proton gradient. (a) Bacteriorhodopsin (BR) and ATP-synthase reconstituted in PETOz-*b*-PDMS-*b*-PETOz polymersomes. Adapted with permission from H. J. Choi and C. D. Montemagno, *Nano Lett.*, 2005, **5**, 2538–2542. Copyright (2005) American Chemical Society.²³⁸ (b) Cytochrome bo₃ quinol oxidase (Bo₃) and ATP synthase reconstituted in PDMS-*g*-PEO polymersomes and PDMS-*g*-PEO/lipid hybrid vesicles. Reprinted (adapted) with permission from L. Otrin, N. Marusic, C. Bednarz, T. Vidakovic-Koch, I. Lieberwirt, K. Landfester and K. Sundmacher, *Nano Lett.*, 2017, **17**, 6816–6821. Copyright (2017) American Chemical Society.¹³ (c) Complementary photoconverters (PSII and PR) and an ATP-synthase reconstituted in lipid SUVs inside lipid GUVs. Reprinted by permission from Springer Customer Service Centre GmbH: Springer Nature, K. Y. Lee, S. J. Park, K. A. Lee, S. H. Kim, H. Kim, Y. Meroz, L. Mahadevan, K. H. Jung, T. K. Ahn, K. K. Parker and K. Shin, Photosynthetic artificial organelles sustain and control ATP-dependent reactions in a protocellular system, *Nat. Biotechnol.*, Copyright (2018).²³⁶

commonly used polymers, accounting for the retained bo₃'s activity in the membrane.

In 2018, Lee *et al.* designed a switchable photosynthetic liposome SUVs (100 nm) in liposome GUVs (10–100 μm) as a superb advancement towards artificial cells (multi-compartmentalization, ATP-regeneration and cytoskeleton formation).²³⁶ The authors used two photoconverters, a plant-derived photosystem II (PSII) and a bacteria-derived proteorhodopsin (PR), to control the trans-membrane proton gradients in the SUVs and thus ATP generation from ADP by ATP-synthase as artificial mitochondria (Fig. 24c). Red light activates PSII's electron-transport chains in the SUVs facilitating ATP synthesis in the GUV lumen while green light activates direct proton pumping by PR, impeding ATP synthesis.

Other processes. In addition to the commonly studied multi-compartmentalization and energy production, artificial vesicles have also been used to mimic a plethora of other cellular processes such as protein expression, metabolism, growth, motility *etc.* The array of processes applied to liposomes and polymersomes is large^{242–252} and we will only describe a few here.

The cytoskeleton has many functions, amongst which the most important ones are cell shape and support, endocytosis, mobility and division, and thus has been gathering extensive focus in cell mimicking.^{113,236,253–261} In the study by Lee *et al.* described above, the photo-switchable generation of ATP was coupled to ATP-dependent actin polymerization leading to morphological changes in the lipid GUV membrane.²³⁶ This process was achieved by embedding magnesium ionophores in the GUV membrane and G-actin in the lumen. Upon ATP synthesis, G-actin nucleation was triggered in the presence of Mg²⁺ and formed actin filaments. The growth of these filaments led to the formation of F-actin spheres which deformed the membrane eventually rupturing the vesicles. The membrane deformation by the polymerization of actin could also be controlled by forming liposomes with liquid-disordered (Ld) domains

composed of polyunsaturated phospholipids which established favorable interactions with actin and liquid-ordered (Lo) domains composed of sphingomyelin, cholesterol and polysaturated-PEG2000-PE weakly interacting with actin. The Ld domain's strong attraction between Ld and actin filaments locally deformed the membrane into teardrops (Fig. 25). Weiss *et al.* used lipid/polymer droplet-stabilized GUVs (dsGUVs) as a protocellular platform.¹¹³ The reconstitution of actin filaments and microtubules was achieved by sequential pico-injection of G-actin or tubulin. The authors showed the importance of pico-injection over pre-mixing in the vesicles as the encapsulation of microtubules inhibited the formation of the dsGUVs.

As a very elegant mimic of a cell, Martino *et al.* reproduced the cell-free expression of bacterial proteins within polymersomes.²⁴³ The GUVs (126 μm) were synthesized from PEO-*b*-PLA (PLA: poly(lactic acid)) and a PLA homopolymer which was used to strengthen the membrane using a microfluidic capillary device (Fig. 26a). The polymersome encapsulated an *E. coli* ribosomal expression kit with the DNA plasmid of a MreB fusion protein with red fluorescent protein (MreB-RFP). Assessing protein expression was achieved by monitoring the resulting increasing fluorescence. The production of protein is a pillar in cell mimicry but cells also deliver their expressed protein to the extracellular environment. Thus, there is also interest in triggered-release. Using negative osmotic shock, pores were formed in the membrane and thus the polymersomes were rendered semi-permeable allowing the triggered release of the newly expressed proteins while retaining the polymersomes by subsequent self-sealing (Fig. 26b). The light-induced osmotic shock release of SUV liposomes and polymersomes (100 nm) from PB-*b*-PEO GUVs (10 μm) has also been described.²⁵¹

Most artificial cell studies focus on the basic processes common to all cells. However, because cells are specialized (blood cells, neurons, skin cells *etc.*), certain functionalities are specific to a



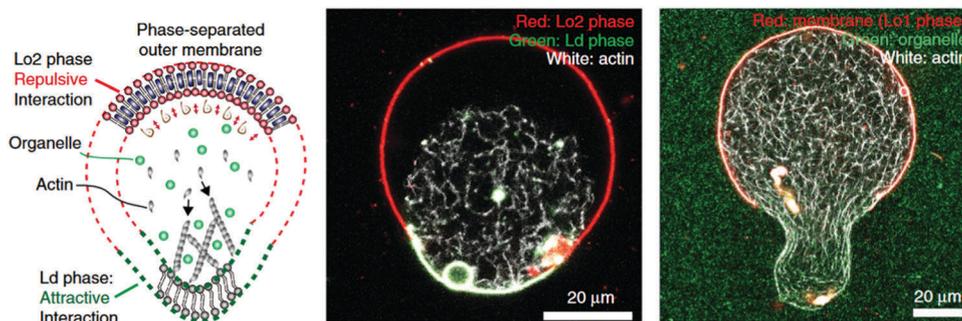


Fig. 25 Control of actin polymerization and membrane shape of the protocellular systems. Liquid-disordered domain (Ld); liquid-ordered domains (Lo1 or Lo2). Reprinted with permission from Springer Customer Service Centre GmbH: Springer Nature, K. Y. Lee, S. J. Park, K. A. Lee, S. H. Kim, H. Kim, Y. Meroz, L. Mahadevan, K. H. Jung, T. K. Ahn, K. K. Parker and K. Shin, Photosynthetic artificial organelles sustain and control ATP-dependent reactions in a protocellular system, *Nat. Biotechnol.*, Copyright (2018).²³⁶

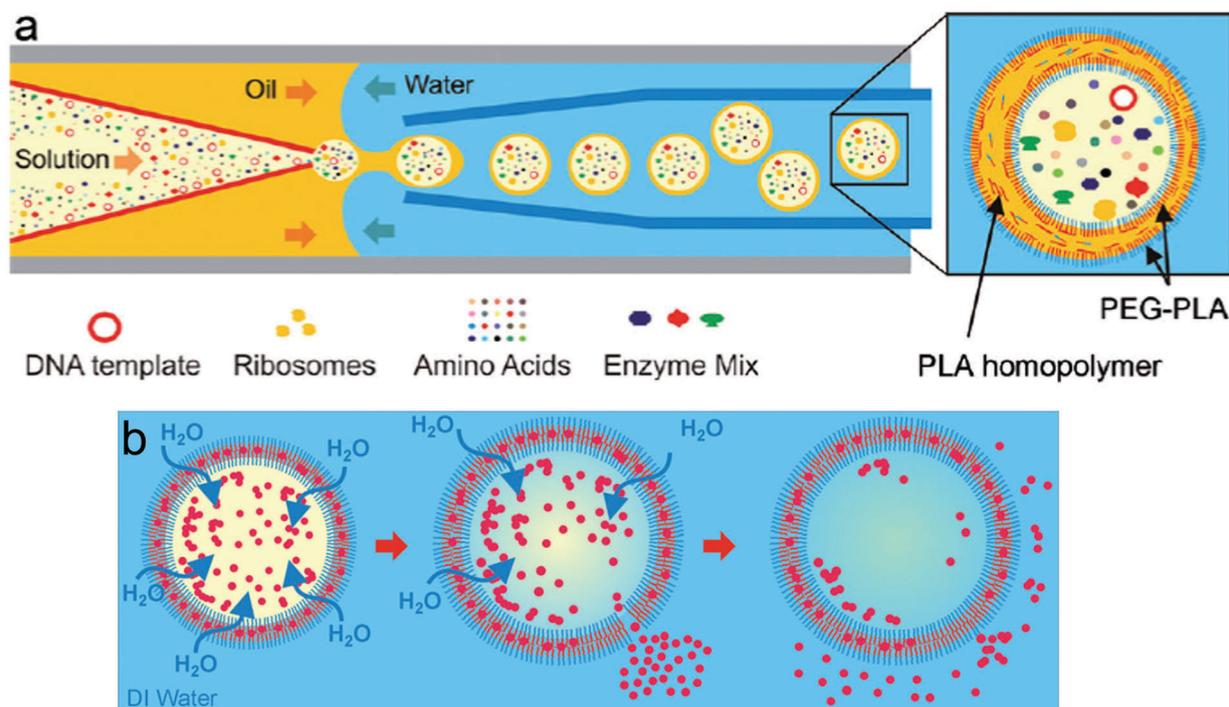


Fig. 26 Protein expressing polymersome GUVs generated by microfluidics. Schematic representation of (a) the methodology used to generate these artificial cells and (b) protein release under osmotic shock and subsequent self-sealing. PEG: poly(ethylene glycol). PLA: poly(lactic acid). Reprinted with permission from C. Martino, S. H. Kim, L. Horsfall, A. Abbaspourrad, S. J. Rosser, J. Cooper and D. A. Weitz, *Angew. Chem., Int. Ed.*, 2012, **51**, 6416–6420. Copyright (2012) Wiley.²⁴³

type of cell.²⁶² Red blood cells are of particular interest because they can also be used in medicine as specialized oxygen carriers. Multiple studies using vesicles mimicking red blood cells by encapsulating hemoglobin (Hb) have been described.^{263–266} Encapsulating hemoglobin does not affect its oxygen binding properties and O₂ off-loading can be regulated by membrane thickness of appropriately selected block copolymers to obtain a moderate O₂ release and avoid side effects. Arifin and Palmer demonstrated that PB-*b*-PEO (250 nm) encapsulating bovine Hb exhibited oxygen affinities comparable to human erythrocytes consistent with values required for efficient oxygen delivery in the systemic circulation and unlike liposomes composed of

natural lipids, these polymersomes did not induce Hb oxidation.²⁶³ However, this example and many others exclusively involved SUVs. Najer *et al.* recently developed red blood cell (RBC) membrane mimics as a potential cure for malaria.²⁶⁶ During infection, malaria parasites enter RBCs by binding to heparan sulfate in order to reproduce. By using polymersomes expressing RBC receptor-like molecules on their outer layer, the parasites would bind to the polymersome and hence their reproduction cycle would be inhibited. PMOXA₅-*b*-PDMS₅₈-*b*-PMOXA₅/PDMS₆₅-*b*-heparin₁₂ hybrid polymersomes (7 μm) were shown to specifically bind to the parasite ligand and even interact with the viable parasite.



VI. Conclusions

Vesicles are interesting carriers as offering a great variety in terms of size, permeability, fluidity, methods of preparation *etc.* and thus have been used in an array of applications ranging from nanoreactors to cell mimicry *via* FDA-approved medicines. Liposomes and polymersomes present many different properties. Liposomes are closer mimics to eukaryotic cell membranes but are difficult to handle and use whereas polymersomes are tougher, malleable, stable vesicles. Nonetheless stability does not always rhyme with cell mimicry as fluidity is a key parameter of cell membranes. Thus the silver-lining to design accomplished artificial cells lies in controllability. However, despite the arsenal of polymer chemistry, polymersome studies, especially polymeric GUVs, are so far mostly restricted to commercially available amphiphilic polymers in a similar way that liposomes are only self-assembled from natural phospholipids. Regardless of their properties, liposomes and polymersomes have both been extensively used to build artificial cells. Recent years have seen a wide range of cell processes and functionalities implemented in these synthetic membranes, which are just starting to be combined into state-of-the-art vesicles slowly building up to generating protocells. Lipid/polymer hybrid vesicles have gained interest as a good compromise between liposomes and polymersomes for a greater control and adaptability of physicochemical properties to any desired functionality and applications, optimal for cell mimicry. As polymersomes have shown great potential, we predict that the era of polymersomes, in parallel with hybrid vesicles, has yet to come when more interdisciplinary works are established to allow the full versatility of polymers' elaborate chemistries to be used in the field of cell mimicry.

Abbreviations

ADH	Alcohol dehydrogenase	DMPC	1,2-Dimyristoyl- <i>sn</i> -glycero-3-phosphocholine (lipid)
ADP	Adenosine diphosphate	DMPE	1,2-Dimyristoyl- <i>sn</i> -glycero-3-phosphoethanolamine (lipid)
AGE	<i>N</i> -Acyl-D-glucosamine 2-epimerase	DMPG	1,2-Dimyristoyl- <i>sn</i> -glycero-3-phospho-(1'- <i>rac</i> -glycerol) (lipid)
AMC	7-Amido-4-methylcoumarin	DMPS	1,2-Dimyristoyl- <i>sn</i> -glycero-3-phospho-L-serine (lipid)
anti-EGFR	Anti-epidermal growth factor receptor	DOPC	1,2-Dioleoyl- <i>sn</i> -glycero-3-phosphocholine (lipid)
ATP	Adenosine triphosphate	DOPE	1,2-Dioleoyl- <i>sn</i> -glycero-3-phosphoethanolamine (lipid)
<i>B</i>	Block	DOPG	1,2-Di-(9 <i>Z</i> -octadecenoyl)- <i>sn</i> -glycero-3-phospho-(1'- <i>rac</i> -glycerol) (lipid)
B-NAR-AMC	Boc-Gln-Ala-Arg-7-amido-4-methylcoumarin	DOPS	1,2-Dioleoyl- <i>sn</i> -glycero-3-phosphoserine (lipid)
bo3	Cytochrome bo3 quinol oxidase	DP	Degree of polymerization
BR	Bacteriorhodopsin	DPPC	1,2-Dipalmitoyl- <i>sn</i> -glycero-3-phosphocholine (lipid)
CalB	<i>Candida antarctica</i> lipase B	DPPE	1,2-Dipalmitoyl- <i>sn</i> -glycero-3-phosphoethanolamine (lipid)
Con A	Concanavalin A	DPPG	1,2-Dipalmitoyl- <i>sn</i> -glycero-3-phospho-(1'- <i>rac</i> -glycerol) (lipid)
CSS	CMP-sialic acid synthetase	DPPS	1,2-Dipalmitoyl- <i>sn</i> -glycero-3-phospho-L-serine (lipid)
CTAs	Chain-transfer agents	dsDNA	Double-stranded DNA
CTP	Cytidine triphosphate	dsGUVs	Droplet-stabilized GUVs
DC-FCCS	Dual-color fluorescence cross-correlation spectroscopy	DSPC	1,2-Distearoyl- <i>sn</i> -glycero-3-phosphocholine (lipid)
DCPC	1,2-Dicetyl- <i>sn</i> -glycero-3-phosphocholine (lipid)	DSPG	1,2-Distearoyl- <i>sn</i> -glycero-3-phospho-(1'- <i>rac</i> -glycerol) (lipid)
DEX	Dextran (hydrophilic block)	eGFP	Enhanced green fluorescent protein
diC15-PC	1,2-Dipentadecanoyl- <i>sn</i> -glycero-3-phosphocholine (lipid)	Egg PC	1- α -Phosphatidylcholine (egg, chicken) (lipid)
DLPC	1,2-Dilauroyl- <i>sn</i> -glycero-3-phosphocholine (lipid)	Egg SM	Sphingomyelin (egg, chicken) (lipid)
		<i>F</i>	Volume fractions
		F-Actin	Filamentous actin
		FDA	Food and drug administration
		FRET	Förster resonance energy transfer
		f_w	Hydrophilic weight fraction
		<i>G</i>	Grafted
		G-actin	Globular actin
		GlcNAc	<i>N</i> -Acetylglucosamine
		GOx	Glucose oxidase
		GUV	Giant unilamellar vesicle
		Hb	Hemoglobin
		HLP	Hybrid liposome-polymersome
		HPMA	2-Hydroxypropyl methacrylate
		HRP	Horseradish peroxidase
		HUV	Unilamellar hybrid vesicles
		Hydro Soy PC	1- α -Phosphatidylcholine, hydrogenated (Soy) (lipid)
		ITO	Indium titanium oxide
		KvAP	Voltage-dependent K ⁺ channel from <i>Aeropyrum Pernix</i>
		LCST	Lower critical solution temperature
		Ld	Liquid-disordered
		LDAO	Lauryldimethylamine <i>N</i> -oxide
		Liss Rhod	1,2-Dioleoyl- <i>sn</i> -glycero-3-phosphoethanolamine- <i>N</i> -Lissamine rhodamine B sulfonyl (lipid)
		Lo	Liquid-ordered
		LUV	Large unilamellar vesicle
		lyso-PC	Lysophosphatidylcholines (lipid)
		ManNAc	<i>N</i> -Acetylmannosamine



MHC	Major histocompatibility complex	PNIPAM	Poly(<i>N</i> -isopropylacrylamide) (responsive block)
MLV	Multilamellar vesicles	polyG	Polyguanylic acid (hydrophilic block)
M_n	Number average molecular weight	POPC	1-Palmitoyl-2-oleoyl- <i>sn</i> -glycero-3-phosphocholine (lipid)
MSPC	(1-Myristoyl-2-stearoyl- <i>sn</i> -glycero-3-phosphocholine) (lipid)	POPS	1-Palmitoyl-2-oleoyl- <i>sn</i> -glycero-3-phospho-L-serine (lipid)
MVV	Multivesicular vesicles	PPE	Poly(phosphonate) (responsive block)
MW	Molecular weight	PPS	Polyphenylene sulfide (responsive block)
N	Degree of polymerization	PR	Proteorhodopsin
NAD ⁺	Nicotinamide adenine dinucleotide	PS	Polystyrene (hydrophobic block)
NADPH	Reduced nicotinamide adenine dinucleotide phosphate	PSe	Phosphatidylserine (lipid)
NAL	<i>N</i> -Acetylneuraminase lyase	PSII	Plant-derived photosystem II
NP-3C	DNA nanopore	PVA	Poly(vinyl alcohol)
NPs	Nanoparticles	PVP	Poly(<i>N</i> -vinylpyrrolidone)
o/w	Oil-in-water	RAFT	Reversible addition fragmentation transfer
OmpF	Outer membrane protein F	RBC	Red blood cell
P2VP	Poly(2-vinylpyridine) (hydrophilic block)	RC	Reaction center
P4VP	Poly(4-vinyl pyridine) (hydrophilic block)	ROS	Reactive oxygen species
PAA	Polyacrylic acid (hydrophilic block)	SIM	Structured illumination microscopy
PALM	Photo-activated localization microscopy	SM	Sphingomyelin/Sphingosylphosphorylcholine (lipid)
Palmitoyl-SM	<i>N</i> -Palmitoyl-D-erythro-sphingosylphosphorylcholine (lipid)	SNOM	Scanning near-field optical microscopy
PAMO	Phenylacetone monooxygenase	SOPC	1-Stearoyl-2-oleoyl- <i>sn</i> -glycero-3-phosphocholine (lipid)
PAzoM	Poly{6-[4-(4-methylphenylazo)-phenoxy]hexyl acrylate} (responsive block)	Soy PC	1- α -Phosphatidylcholine (Soy) (lipid)
PBA	Poly(butyl acrylate) (hydrophobic block)	STED	Stimulated emission depletion microscopy
PBD or PB	Polybutadiene (hydrophobic block)	SUV	Small unilamellar vesicle
PC	Phosphatidylcholine (lipid)	TEM	Transmission electron microscopy
P_c	Critical packing parameter	TEMPO	(2,2,6,6-Tetramethylpiperidin-1-yl)oxyl
PCL	Polycaprolactone (hydrophobic block)	T_g	Glass-transition temperature
PDA	Polydiacetylenes (amphiphilic block)	T_m	Gel-liquid crystal transition temperature/ Melting temperature
PDEA	Poly(2-diethylaminoethyl methacrylate) (responsive block)	UCST	Upper critical solution temperature
PDLLA	Poly(D,L-lactide) (hydrophobic block)	UV	Ultraviolet
PDMS	Polydimethylsiloxane (hydrophobic block)	w/o	Water-in-oil
PDPA	Poly(2-(diisopropylamino)ethyl methacrylate) (responsive block)	w/o/w	Water-in-oil-in-water
PE	Phosphatidylethanolamine (lipid)	X	Flory-Huggins parameter
PEE	Poly(ethyl ethylene) (hydrophobic block)		
PEO or PEG	Poly(ethylene oxide) or Poly(ethylene glycol) (hydrophilic block)		
PEtOz	Poly(2-ethyl-2-oxazo-line) (hydrophilic block)		
PG	Phosphatidylglycerol (lipid)		
PGA	Poly(glycolic acid) (hydrophobic block)		
PI	Phosphatidylinositol (lipid)		
PIAT	Poly(3-(isocyano-L-alanyl-aminoethyl)thiophene) (hydrophilic block)		
PIB	Polyisobutylene (hydrophobic block)		
PISA	Polymerization-induced self-assembly		
PLA	Poly(lactic acid) (hydrophobic block)		
PLys	Poly-L-lysine (responsive block)		
PMA _{SH}	Thiol-functionalized poly(methacrylic acid)		
PMMA	Poly(methyl methacrylate) (hydrophobic block)		
PMOXA	Poly-2-methyl-2-oxazoline (hydrophilic block)		
PMPC	Poly(2-(methacryloyloxy)ethyl phosphorylcholine) (hydrophilic block)		

Conflicts of interest

There are no conflicts to declare.

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