Antibiofouling polymer interfaces: poly(ethylene glycol) and other promising candidates

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Antibiofouling polymer interfaces: poly(ethylene glycol) and other promising candidates

Antibiofouling polymer interfaces: design criteria for biomedicine and nano-therapeutics

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TOC one-line summary:

This review highlights antibiofouling polymer interfaces with emphasis on the latest developments using poly(ethylene glycol) and the design new polymeric structures.

Abstract

Nonspecific protein adsorption and/or microbial adsorption on biomedical materials adversely affects the efficacy of a range of biomedical systems, from implants and biosensors to nanoparticles. To address this problem, antibiofouling polymers can be coated on

biomedical devices or built into nanoparticles to confer protein and/or microbial repellent properties. The current review provides an overview of the range of synthetic polymers currently used to this end and explores their biomedical potential. The most widely-used antifouling polymer, poly(ethylene glycol) (PEG) is reviewed alongside several promising alternatives, including zwitterionic polymers, poly(hydroxyfunctional acrylates), poly(2-oxazoline)s, poly(vinylpyrrolidone), poly(glycerol), peptides and peptoids. For each material, notable applications for both nanomedicine and macroscopic surface coatings are highlighted.

Introduction

Biofouling from nonspecific ~~protein~~[biomolecule](#) or microbial adsorption is a ubiquitous and persistent challenge for any interface that is exposed to biological fluids or tissue. On *in vivo* devices, such as implants, catheters, biosensors or tissue engineered scaffolds, protein adsorption can lead to adverse immunological responses and may promote bacterial colonization and infection.^{1,2} Furthermore, on *in vitro* devices which come into contact with protein-rich biological material, such as microarrays or filtration membranes, nonspecific protein adsorption can either partially or entirely compromise functionality.² ~~Protein adsorption~~[Biomolecule adsorption](#) is also a major problem for nanoparticles, such as micelles, liposomes, and nanocapsules, used for drug delivery and bioimaging. ~~These systems often require long circulation times to sufficiently accumulate within target tissue or tumors.~~ Particles without antifouling moieties, ~~however,~~ are usually quickly coated in [biomolecules such as lipids, proteins, and sugars when they enter the bloodstream, creating a biomolecular corona.](#)³ ~~The major constituent of the corona is a small subset of blood proteins,~~³ ~~and this nonspecific adsorption of blood proteins is termed opsonisation.~~⁴ ~~Opsoblood proteins in a process called opsonization, forming a protein corona that reduces functionality.~~³ ~~Opse~~onization can adversely affect the chemistry of conjugated drugs, causing aggregation or charge neutralization, and, moreover, typically leads to a rapid removal of nanoparticles from

the blood by phagocytic cells of the immune system.⁵⁻⁷ Given the given that nonspecific protein adsorption is a key problem facing biomedical surfaces, this review will examine the key dominant major polymer-based strategies to reduce it.

A common strategy to reduce protein fouling is to attach hydrophilic or zwitterionic synthetic polymers to surfaces. These polymers' electrical neutrality can help reduce electrostatic interactions with charged protein domains.⁸ Their hydrophilic nature helps reduce nonpolar interactions between proteins and hydrophobic surfaces.⁹ Moreover, these polymers may prevent protein adsorption by steric hindrance,¹⁰⁻¹² resulting in increased circulation times. Antifouling polymers have found applications as coatings for both macroscopic surfaces (such as implants and biosensors) and in nanoparticulate systems, endowing them with "stealth properties" (i.e., the ability to evade the body's normal clearance mechanisms).

The field of antifouling polymers is a fast growing area with a number of strategies and new polymer structures designed for this purpose. Poly(ethylene glycol) (PEG) has been the dominant polymer used for antifouling surfaces. However, PEG has limitations which have led to the development of a range of alternative polymeric materials, several of which are shown in **Figure 1**. While several comprehensive reviews examine specific classes of antifouling polymers such as poly(ethylene glycol) PEG,¹³⁻¹⁹ zwitterionic materials^{19, 20} and poly(2-oxazolines),²¹⁻²⁵ the full range of antifouling polymers is rarely considered in one place. When it is, discussion typically focuses exclusively on nanoparticulate drug delivery systems²⁶⁻²⁹ or macroscopic surface coatings^{9, 12, 30, 31} rather than the whole range of nonfouling, biomedical applications.

This review will therefore provide a high level overview of the antifouling polymeric suite and its applications for biomaterials. It discusses both PEG and the most common synthetic PEG alternatives, with examples given for each material on both macroscopic surfaces and in

nanoparticulate systems. Emphasis is placed on the most recent advances and key foundational research. Focus will be placed on general antifouling polymer studies, rather than studies which focus on specialty applications like membranes and bioassays. Further, discussion will be limited to solution phase synthesis, rather than solid phase synthesis approaches such as chemical vapour deposition.³² Nevertheless, the review will examine a broad range of some of the most salient and clinically applied antifouling polymers.

Figure 1. (A) Chemical structure of PEG and PEG alternatives designed as antifouling coatings. (B) A macroscopic surface without an antifouling polymer coating is quickly coated in proteins when it comes into contact with blood. (C) Coating the surface with PEG or PEG alternatives reduces protein adsorption to the surface. (D) As with macroscopic surfaces, nanoparticles (NPs) without antifouling polymers suffer from nonspecific protein adsorption in the body. Protein adsorption leads to phagocytosis by macrophages and removal of NPs from the bloodstream. (E) Attaching antifouling polymers to NP surfaces allows the particles to resist protein adsorption and phagocytosis and increases their blood circulation half-life.

1. Poly(ethylene glycol)

PEG has been the most widely used polymer for antifouling applications, such that it has often been termed the “gold standard” of antifouling polymers.^{22, 28} In terms of surface applications, Prime and Whitesides first reported the potential for PEG-derivatives, showing that oligo(ethylene glycol) (OEG) self-assembled monolayers (SAMs) were effective protein-repellent coatings.^{33, 34} However, OEG SAMs are prone to surface defects and have limited application to metal surfaces.³⁵ Given the limitations of these SAMs, alternative means of attaching PEG to surfaces have been explored.

The effectiveness of PEG, and antifouling polymers in general, depends upon the surface grafting technique ~~and achievable grafting densities, and the polymer architecture~~.^{36, 37} ~~For this reason~~ Thus, a variety of grafting approaches have been developed to improve the effectiveness of PEG and other antifouling coatings. For example, ~~firstly, for example, linear PEG can be~~ can be covalently attached to surfaces, ~~for example, using “grafting to” approaches to form linear polymer brushes~~³⁸⁻⁴⁰. ~~Secondly, PEG methacrylates can also be~~ can be polymerized from ~~initiator pre-treated surfaces using a “grafting from” techniques~~ approach (^{34, 35} ~~For this reason, a variety of grafting approaches have been developed to improve the effectiveness of PEG and other antifouling coatings. These include polymerization techniques such as~~ se.g. Surface-Initiated Atom Transfer Radical Polymerization (SI-ATRP)).^{31, 41-44} ~~Using this approach~~ technique, ~~effective antifouling brushes are~~ can be formed consisting of linear polymers with PEG side chains. ~~but also novel~~ Novel and biomimetic ~~Several novel approaches~~ grafting strategies developed specifically for antifouling applications ~~have also been developed~~. For example, a biologically inspired approach using components of muscle adhesive protein (i.e., L-3,4-

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dihydroxyphenylalanine, L-DOPA) has been effectively used to attach PEG to surfaces via catechol groups (see **Figure 2** **Figure 2 (B)**).⁴⁵⁻⁵¹ In a separate approach, the amino acid lysine has been used as a polymer brush backbone with PEG.^{47, 52-54} ^{42, 47-49} -PEG side chains are covalently linked through the primary amine on the lysine residues to form poly(lysine)-*graft*-PEG (PLL-*g*-PEG) polymers (see **Figure 2** **Figure 2 (A)**). Some lysine residues on the backbone are left in their native, cationic state, and these residues can adsorb to an anionic metal surface such as titanium oxide. ~~The Further, the~~ lysine side chain can ~~also~~ be functionalized with the integrin ligand, Arg-Asp-Gly (RGD), to endow the surface with the ability to specifically bind to host cells.⁵³

The poly(lysine) strategy can be combined with a catechol strategy on the same polymer to create a multifunctional polymer anchor. Saxer et al.⁴⁷ developed ~~such a polymer~~ (**Figure 2** **Figure 2**), showing that the catechol groups led to increased coating resistance to high ionic salt solutions compared to PLL-*g*-PEG alone.⁴⁷

Figure 2. Poly(L-lysine)-*graft*-(3,4-dihydroxyphenylacetic acid, poly(ethylene glycol)). When bound to a surface, the polymer's PEG groups (a) conferred resistance to plasma fibrinogen adsorption, while the catechol (b) and lysine-containing (c) monomers anchored the polymer to the surface. ~~Reproduced from ref~~ [Reprinted with permission from S. Saxer, C. Portmann, S.](#)

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[Tosatti, K. Gademann, S. Zürcher and M. Textor, *Macromolecules*, 2010, **43**, 1050-1060. Copyright 2010 publisher.](#)⁴⁴

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Despite significant research into PEG's applications on surfaces, its mechanism of action are still not fully understood.^{9-11, 55} PEG's antifouling properties likely stem from steric repulsion associated with a loss of entropy when proteins attempt to adsorb to surfaces.^{10, 11}

Furthermore, it is likely PEG's hydrophilic nature leads to the formation of a hydration layer which inhibits protein encroachment.⁹ Several variables which determine PEG's effectiveness include the grafting density (with higher densities being associated with greater protein repellence), chain length, and type of PEG branching architecture.^{37, 56}

In addition to being useful for coating macroscopic surfaces, PEG has been effectively used in a variety of nanoparticulate systems.¹³⁻¹⁹ Attaching PEG to nanoparticles, known as PEGylation, gained popularity when liposomes were first PEGylated in the early 1990s.⁵⁷⁻⁵⁹ PEGylated liposomes encapsulating the anti-cancer drug doxorubicin became the first commercially available polymeric particulate drug delivery system (Doxil) in 1995.⁶⁰ In addition to liposomes, PEG has been used across the complete range of nanoscopic biomaterials, such as: drug conjugates,^{61, 62} inorganic nanoparticles,⁶³ polymeric nanoparticles,⁶⁴⁻⁷¹ micelles,⁷²⁻⁷⁴ star and hyperbranched polymers,^{65, 75-79} and knedel-like polymers.⁸⁰

PEGylation of nanoparticles confers a variety of benefits, including improved water solubility as well as reduced opsonization and associated improvements in blood circulation time.⁵ Presumably due to this reduced opsonization and increased stability, PEG can increase the plasma half-life of conjugated drugs from several minutes to several hours.⁵⁷

However, PEG has several notable weaknesses in biomedical contexts.^{15, 28, 81} PEG can suffer oxidative damage in biological media and fluids, which limits its use for long term applications.^{82, 83} Moreover, because PEG is not biodegradable, there are concerns about bioaccumulation in the lysosomes of healthy cells.^{28, 84} Additionally, although PEG is typically ~~regarded as~~considered to be immunologically inert~~stealth polymer~~~~non-immunogenic~~, it has nevertheless ~~being been~~ shown to ~~be immunogenic~~provoke an immune response in some circumstances. For instance, when PEG has been used with drug delivery particles, antibodies to the PEG polymer have been shown to be generated.⁸⁵ Anti-PEG antibodies may result in faster clearance *in vivo*, reducing the effectiveness of PEGylated drugs. Moreover, it could lead to carrier-induced epitope specific suppression,⁸⁶ an effect where pre-immunization with a carrier can lead to a reduced immune response to antigens later presented on that same carrier. This phenomena would reduce PEG's applicability as an antigen carrier.

In some cases, PEGylated liposomes or other nanoparticles have greatly reduced blood circulation time after the first dose in animal models. This response appears to be IgM-mediated^{87, 88} and is termed the accelerated blood clearance (ABC) phenomenon (see **Figure 3****Figure 3 (A)**).⁸⁹⁻⁹¹ Interestingly, the effect is not observed for PEGylated liposomes encapsulating anti-cancer drugs; this is potentially because these drugs are cytotoxic to antibody secreting cells.^{90, 91}

Figure 3. (A) Gold nanoparticles coated in PEG (PEG-GNP) stimulate the production of anti-PEG IgM over five days after the first injection [in rats](#). Production of IgM leads to the production of IgG. (B) When GNPs are coated with ultralow fouling poly(carboxybetaine) (PCB), they resist the adsorption of IgM and thus the production of IgG. (C) The ABC phenomenon was tested by injecting rats with PCB-GNPs or PEG-GNPs at two time points (data represent average values across 6 rats). The PEG-GNP's half-life markedly decreased from 8.7 h in the first dose to 5.2 h in the second dose, while the half-life for PCB-GNPs remained relatively constant between the first and second dose (55.8 h and 55.6 h, respectively). Adapted from ref. ⁸⁷. Reprinted with permission from W. Yang, S. Liu, T. Bai, A. J. Keefe, L. Zhang, J.-R. Ella-Menye, Y. Li and S. Jiang, *Nano Today*, 2014, 9, 10-16. Copyright 2014 publisher Elsevier.

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2. Zwitterionic polymers

Zwitterionic polymers have recently emerged as promising alternatives to PEG.^{8, 20, 92}

Zwitterionic polymers are electrically neutral materials which contain both a positively charged species and a negatively charged species on the same monomer or, in the case of polyampholytic materials,^{8, 93-95} on different monomers. The charged species appear

associated with an even stronger hydration effect than that created by PEG, which may in turn enhance the zwitterionic material's antifouling properties.^{20, 96}

Several studies have indicated that zwitterionic surface coatings can have highly effective protein repellence. Surfaces coated in zwitterionic poly(2-methacryloyloxyethyl phosphorylcholine) brushes have shown protein resistance which surpasses comparable PEG coatings.⁹⁷ Similarly, poly(sulfobetaine methacrylate) (pSBMA)⁹⁸ and poly(carboxybetaine methacrylate) (pCBMA)^{99, 100} have shown ultralow fouling properties. Use of an acrylamide^{101, 102} or a norbornene¹⁰³ polymer backbone with zwitterionic betaine side chains can lead to undetectable levels of protein adsorption from undiluted human blood serum or plasma (see [Figure 4](#) ~~Figure 4~~ for tests using carboxybetaine).

Zwitterionic surface coatings have recently been shown to have promise in clinically relevant implant models.^{104, 105} For example, Liu et al.¹⁰⁵ reported that grafting pSBMA on to titanium alloy orthopaedic/dental implants led to increased osteointegration by promoting mineralization of the implant surface.

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Figure 4. (A) Protein adsorption on different gold surfaces coated with various antifouling materials, including three self assembled monolayers (SAMs) based on ω -oligo(ethylene glycol) alkanethiols (EG4/COEG6, EG2, EG6), and polymer brushes produced using surface-initiated atom-transfer radical polymerization (ATRP) of oligo(ethylene glycol) methacrylate (HOEGMA), 2-hydroxyethyl methacrylate (HEMA), carboxybetaine acrylamide (CBAA) and *N*-(2-hydroxypropyl) methacrylamide (HPMA). The surfaces were exposed to undiluted blood plasma (B) or cerebrospinal fluid (C) for 15 minutes and the nonspecific protein adsorption was measured using surface plasmon resonance (SPR). Protein adsorption below the SPR detection limit ($0.03 \text{ ng} \cdot \text{cm}^{-2}$) was reported as zero. [Reprinted with permission from C. Rodriguez-Emmenegger, M. Houska, A. B. Alles and E. Brynda, *Macromol. Biosci.*, 2012, **12**, 1413-1422.](#) Copyright 2012 publisher Wiley-VCH. Adapted from ref. ⁹⁸.

In addition to surface-grafted polymer architectures, several new hydrogels with zwitterionic polymers appear to be particularly promising.^{104, 106, 107 95-98, 99} Zhang et al.,¹⁰⁴ for example, developed a pCBMA hydrogel with long-term resistance to the foreign body reaction, ~~the body's immunological response to implanted foreign objects.~~ [In the foreign body reaction,](#)

nonspecific protein adsorption is thought to signal the presence of the implant to the immune system and set off a cascade of events, setting off a cascade of events leading to the which leads to the triggering the formation of a dense collagen layer around the implant to sequester the implant from surrounding tissue. This collagen layer can prevent effective mass transport and electrical communication between an implant and the body. Remarkably, the pCBMA hydrogels implanted in mice resisted the foreign body reaction for three months. Zhang et al. The investigators showed that pCBMA hydrogels resisted the foreign body reaction for three months when implanted in mice. These zwitterionic polymers could reduce the formation of dense collagenous material at the implant site. This was presumably because the surface strongly resisted nonspecific protein adsorption. This effect was attributed, an effect attributed to the surface's strong protein repellent properties. The authors hypothesized that by resisting preventing protein adsorption, allowed the pCBMA hydrogels were able to prevent the chain of events leading to the formation of the collagen layer, as protein adsorption is thought to be the first step in the immune response which lead to the formation of the dense collagen layer. From an applications viewpoint, this would be preventing the foreign body reaction and the formation of a collagen layer, these pCBMA materials could potentially improve mass transport and electrical communication between an implant and the body.

Zwitterionic polymers' chemical structure can allow for several unique functionalities. Carboxybetaines, for example, contain multiple functional groups amenable to conjugation. Further, due to the presence of oppositely charged species on each monomer, zwitterionic materials can be used as switchable surfaces, capable of shifting between cationic, zwitterionic, and anionic states based on the environmental pH.¹⁰⁸

Zwitterionic polymers have also been successful at conferring protein resistant properties to nanoparticles.^{20, 109} To this end, zwitterionic polymers have been used in a variety of nanoparticulate systems, including surface coatings for gold nanoparticles¹¹⁰⁻¹¹² and nanorods,¹¹³ silica nanoparticles,¹¹⁴ iron oxide nanoparticles,^{115, 116} block copolymer micelles,¹¹⁷ knedel-like polymers,¹¹⁸ dendrimers,¹¹⁹ polymeric immunobeads,^{120, 121} [and quantum dots](#)^{122, 123}^{111, 112} and [in](#) protein conjugates.¹²⁴ In these applications, zwitterionic particles [can](#) strongly resist protein fouling.^{110, 113-118} A direct comparison between PEG-coated and pCBMA-coated knedel-like nanoparticles found the two had similar size stability characteristics and biodistribution profiles in mice.¹¹⁸

Jiang and co-workers in fact found that [in rats](#), pCBMA-coated gold nanoparticles ([pCB-GNPs](#)) had a longer half-life in the blood stream [in rats](#) than PEG-coated [gold nanoparticles \(PEG-GNPs\)](#)¹¹²

[A biodistribution study showed that after 5 days after the first injection, 30% of the original dose of pCBMA particles were remained found in the blood stream, while almost all of the PEGylated particles PEG-GNPs had accumulated in the been removed to the liver and other organs.](#)

[Moreover, unlike the PEGylated particles PEG-GNPs, the pCBMA particles were able to avoid the ABC phenomenon.](#)

[As shown in \(Figure 3Figure 3Figure 3Figure 3Figure 3Figure 3Figure 3, for PEG-GNPs the rate of clearance PEG particles was greater after the second dose of PEG-GNPs than the first dose. The clearance rate of pCBMA, on the other hand, did not appear to acceleratechange between the first and second dosesafter the first dose between doses.\)](#)

Consistent with this the blood residence time data, blood samples taken after injection showed that the pCBMA-PEG nanoparticles were associated with lower associated with elevated blood levels of IgM and IgG, two antibodies thought to underlie the ABC phenomenon. These two antibodies associated with opsonization and removal of nanoparticles from the bloodstream which trigger the immune removal of foreign particles, and presumably cause the accelerated clearance of the PEGylated particles. The pCB-GNP-injected rats had similar levels of IgM and IgG as control rats who were not injected with nanoparticles.

Moreover, unlike the PEGylated particles, the pCBMA particles were able to avoid the ABC phenomenon (Figure 3). Thus, zwitterionic particles were able to avoid the ABC phenomenon (Figure 3). Given such data, zwitterionic species appear to be a promising class of antifouling polymers which may be readily modified at their reactive functional groups for particular biomedical applications.

3. Poly(hydroxyfunctional acrylates)

Poly(hydroxyfunctional acrylates) include polymers such as poly(2-hydroxyethyl methacrylate) (pHEMA), poly(hydroxypropyl methacrylate) (pHPMA), and poly(N-hydroxyethylacrylamide) (pHEAA). Poly(hydroxyfunctional acrylates) are similar to other PEG alternatives in that they are electrically neutral and hydrophilic.

These polymers have a long history as biomaterials. pHEMA hydrogels, for example, have been conventionally used in a range of biomedical applications, including implants, tissue engineering scaffolds, and contact lenses, for over 25 years.^{125, 126, 144, 145}

Recent work has explored both polymer brushes^{101, 102, 127-131} and hydrogels^{101, 132, 133} containing hydroxyfunctional acrylates for surface antifouling applications. *In vitro* protein adsorbency tests with pHEMA demonstrate that it has protein repellence comparable to PEG.¹²⁷ Moreover, use of the acrylamide backbone (i.e., pHEAA) can produce surfaces with undetectable protein adsorption from undiluted human blood serum.¹⁰¹ Similarly, pHPMA brushes can produce ultralow fouling properties similar to zwitterionic poly(carboxybetaine acrylamide)¹⁰² (see [Figure 4](#)).

While polymers like pHEMA^{129, 134, 135} and pHPMA¹³⁶⁻¹³⁸ have been incorporated into nanoparticles to confer antifouling properties, pHPMA drug conjugates have been the most prominent application of hydroxyfunctional acrylates for nanomedicine.^{139, 140} pHPMA has been conjugated to cancer drugs (e.g., doxorubicin) in order to reduce drug toxicity or increase circulation time. In addition, by increasing a drug's effective molecular weight, pHPMA conjugation can be used to take advantage of the enhanced permeability and retention (ERP) effect associated with tumors.^{139, 141-145} pHPMA is notable in that several commercial clinical trials involving pHPMA drug conjugates have commenced.²⁸

4. Poly(vinylpyrrolidone)

Like the poly(hydroxyfunctional acrylates), poly(vinylpyrrolidone) (PVP) has a long history as a biocompatible polymer, being used as a plasma substitute in the 1930s and later as a food additive. PVP is a highly hydrated species,^{146, 147} which likely accounts for its protein-repellence.¹⁴⁸

Although macroscopic surfaces coated in PVP have received relatively less research attention, PVP does hold promise for surface applications.¹⁴⁹⁻¹⁵¹ Notably, Serrano et al.¹⁵¹ recently tested PVP against human serum and found protein adsorption similar to PEG and

poly(2-ethyl-2-oxazoline), supporting PVP's utility for biomedical and similar antifouling applications.

PVP has been extensively examined in the context of nanoparticulate systems. A major application for PVP is in micellar systems,¹⁵²⁻¹⁵⁷ although PVP has also been used in liposomes,¹⁵⁸⁻¹⁶⁰ nanostructured capsules,¹⁶¹⁻¹⁶⁴ and drug conjugates.^{165, 166}

PVP appears to have favourable qualities for drug delivery, including resistance to opsonization and phagocytosis. For example, PVP-conjugated tumor necrosis factor- α (TNF- α) showed a longer blood circulation time and twofold increase in antitumor effect compared to PEGylated TNF- α .¹⁶⁷ However, while most studies in mouse models of cancer find a positive effect of PVP conjugation or coating,^{165, 168-171} some report that PVP underperforms

compared to PEG controls in terms of circulation time or immunogenicity. This has been observed for with conjugation of PVP to the enzyme uricase-¹⁶⁵ and for with coating of poly(D,L-lactide) (PLA)-based nanoparticles with PVP-(Gaucher et al., 2009).¹⁷¹ Although the reason for PVP underperformance is unclear, it may be a function of the capacity of the polymer to sterically hinder protein adsorption in the - the steric hindrance provided by a polymer in any particular system - under consideration. For example, as a brush-like coating for -with the PLA nanoparticles, the relatively bulky side chain of PVP may would may confer rigidity which prevents it from freely rotating in space and sterically hindering protein adsorption conformational rigidity.¹⁷¹ (Gaucher et al., 2009) This may prevent PVP from forming a dense, conformational cloud which provides steric hindrance to protein adsorption. -and prevent the polymer from - PEG, on the other hand, lacking this side chain, would be is more flexible and - Being able to freely rotate in space, PEG may be better therefore able to provide this steric hindrance avoid this able to form forming a dense conformational cloud providing steric

~~hindrance necessary to sterically hinder protein adsorption. That said, (Gaucher et al., 2009).~~
~~That said~~~~However~~~~That~~~~That said, that~~The ability of PVP to form such a confirmation cloud is
a function of its molecular weight. Therefore, op-~~and~~ optimisation of this parameter may
yield more effective PVP coatings.~~gs.~~

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^{151, 158} Thus, PEG as with most antifouling polymers, further characterization of the immunological qualities of PVP is justified.

5. Poly(2-oxazoline)s

Poly(2-oxazoline)s (POxs) such as poly(2-methyl-2-oxazoline) and poly(2-ethyl-2-oxazoline) have recently become prominent PEG alternatives.²¹⁻²⁵ POxs have generated considerable interest in part because they are able to maintain their antifouling character for longer periods than PEG, suffering less oxidative damage in biological and oxidative media.¹⁷²

POxs have successfully been used as surface coatings, and POxs have similar protein repellence as PEG when the grafting density is optimised.²² As with PEG, a variety of surface attachment techniques are available,²¹ including “grafting-to,”^{173, 174} “grafting-from,”¹⁷⁵ and the use of poly(lysine)-g-POx brushes.¹⁷⁶

Figure 5. The synthesis and *in vivo* testing of micellar drug carriers constructed from poly(2-oxazoline)s (POxs). Four diblock or ABA-type triblock polymers were first produced entirely from POxs (A). To tune the hydrophilic/hydrophobic nature of the POx monomers, the length of the 2-alkyl side chains was varied. A methyl or ethyl side chain was used to produce a hydrophilic monomer and a butyl side chain was used to produce a hydrophobic monomer. When combined in an aqueous solution with hydrophobic drugs, these polymers self-assembled into micelles (B). Using this strategy, one of the polymers (polymer P2) was used to encapsulate the anticancer drug Paclitaxel (PTX) (P2-PTX micelles). These micelles were tested against empty P2 micelles, a saline control, and a commercially available PTX carrier, Cremophor EL/ethanol (CrEL) in a mouse model of lung cancer (C). At all time points, the P2-PTX formulation performs similarly or better than the CrEL control. [Reprinted with permission from R. Luxenhofer, A. Schulz, C. Roques, S. Li, T. K. Bronich, E. V. Batrakova, R. Jordan and A. V. Kabanov, *Biomaterials*, 2010, **31**, 4972-4979. Copyright 2010 Elsevier. Adapted from ref. 170.](#)

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POxs have been used in place of PEG in several nanoparticulate systems. Notably, POxs appear promising as the hydrophilic component of block copolymer micelles designed to deliver hydrophobic drugs (see [Figure 5](#) for an example).¹⁷⁷⁻¹⁸³ POxs have also been coated on the surfaces of liposomes^{184, 185} and nanocapsules¹⁸⁶ to successfully impart stealth properties.

POxs appear to have similar *in vivo* characteristics as PEG in terms of circulation time and uptake by the reticuloendothelial system.^{184, 187} Several recent studies have found that POxs have limited cytotoxicity or immunogenicity *in vitro*.^{183, 188-193} For instance, Pulkkinen et al.¹⁹³ have shown *in vivo* that poly- ϵ -caprolactone cross-linked with 2, 2-bis(2-oxazoline) is biocompatible, non-toxic, and is slowly enzymatically degraded over a 12 week period when implanted in rat heart, lung, liver, kidney, spleen and brain tissue. These studies help to allay previous uncertainties about POxs' biocompatibility²⁸ and encourage further exploration of POxs for nanomedicine and controlled drug release applications.

6. Poly(glycerol)

Poly(glycerol) (PG) is a common, biocompatible polyol widely used in the food and pharmaceutical industries. Its polyether structure is similar to PEG but also contains hydroxyl groups amenable to functionalization. PG has received attention as an antifouling surface coating over the last 10 years. While surfaces with linear PG have been synthesized,¹⁹⁴⁻¹⁹⁶ the hyperbranched form appears to be more resistant to protein adsorption¹⁹⁶ and is more extensively studied. PG has been affixed to surfaces both as SAMs¹⁹⁴⁻¹⁹⁹ and as polymer grafts.²⁰⁰⁻²⁰⁵ While PG coatings have strong resistance to model proteins such as fibrinogen and albumin,^{201, 202, 204, 205} they have not yet been extensively tested in complex biological media such as human blood plasma.

PG has been used to impart stealth properties to a number of nanoparticulate systems, including inorganic nanoparticles,^{206, 207} liposomes,²⁰⁸⁻²¹⁰ nanogels,²¹¹ and micelles.²¹² *In vitro*^{213, 214} and *in vivo*, mice studies^{215, 216} show that PG is biocompatible and [immunologically inert](#) [evades the immune system](#). For instance, Kainthan and Brooks report a plasma half-life for hyperbranched glycerol of approximately 57 hours with no notable toxicity.²¹⁶ However a significant build-up and retention of hyperbranched glycerol in organs of the reticuloendothelial system (i.e., the liver and spleen) was observed, an effect which increased with molecular weight. This was presumably due to a lack of biodegradable groups in the molecule, a limitation shared with PEG.

7. Peptides and peptoids

Synthetic peptides and peptoids may serve as effective antifouling polymers and may overcome the biodegradability issues of PEG and other PEG alternatives. Indeed, peptide-based SAMs,^{217, 218} peptoid-based SAMs²¹⁹ and peptoid-based polymer brushes^{36, 220-225} have recently been found to be highly effective as antifouling macroscopic surface coatings.

Peptoids, or poly(*N*-substituted glycine) (see [Figure 1](#)[Figure 4](#)), are synthetically produced mimics of natural peptides with the side chain attached to the nitrogen atom, rather than the α -carbon. The shift of the side chain appears to endow peptoids with increased resistance to proteolytic degradation.^{226, 227}

Statz and co-workers developed peptoid-coated surfaces that were able to resist protein adsorption from repeated washings with fresh serum for up to five months (**Figure 5**).²²⁰ The antifouling character arises from the peptoid's methoxyethyl side chain, whose structure resembles PEG. The impressive long-term resilience of the surface may be attributed to the anchoring group, a peptide designed to mimic mussel adhesive protein.

Peptides have a distinct appeal as building blocks for nanoparticles. Because of their biodegradability, peptide-based nanoparticles can potentially avoid the bioaccumulation problems associated with PEG. In addition, peptides can be designed such that they are degraded by proteases at target sites, a trait that can be used for drug targeting.²³⁴

By combining poly(amino acids) and PEG, advantages of both can be obtained in a single molecule.²³⁵ For example, ring-opening polymerization of amino acid N-carboxyanhydrides

has been effectively used to produce peptide star polymers with a PEG corona and a core containing cationic^{236, 237} or anionic²³⁸ poly(amino acids). These star peptides have high biocompatibility and can be functionalized to target cancer cells²³⁶ and deliver and release drug cargo in a pH-dependent manner.²³⁸

Peptides and proteins are commonly used as signalling molecules within the body, a feature which can be utilized to create stealth particles. Rodriguez et al.²³⁹ designed peptides based on human CD47 protein, a molecule which signals to the immune system that its carrier is part of the body and should not be removed by phagocytes. Conjugating these “self-peptides” to nanobeads led to improved stealth-like properties, with approximately four times as many self-peptide presenting nanobeads remaining in circulation than PEGylated controls at 40 minutes post-injection into mice.

Peptoid-based nanoparticles may also have a nonfouling character. The non-ionic, hydrophilic poly(sarcosine) has been used in block copolymers with poly(γ -methyl L-glutamate)²⁴⁰ and poly(L-lactic acid)²⁴¹ to make opsonization-resistant vesicles for bioimaging.^{240, 241} However, despite the apparent promise of peptoids for biomedical applications, limited research has specifically examined their biocompatibility²²⁵ and future work will be needed in this area.

Conclusion

Several polymeric materials, including PEG and a range of PEG alternatives, have been developed to combat protein-based fouling. Interestingly, antifouling properties observed on relatively flat, macroscopic surfaces are maintained at the nanoscale and on surfaces with a variety of geometries, including spherical interfaces.

Although the outlook for antifouling polymers is positive, much of the work is at an early-stage of application. For instance, many of the macroscopic surface coatings reviewed (e.g.,

polyglycerols) have been tested primarily *in vitro* with a limited number of model proteins, rather than whole blood fluids. Additional research will be required to test these surfaces against complex biological media and *in vivo*.

Considerable work has examined protein repellent nanoparticles in clinical settings; noteworthy examples include PEG and HPMA, which have in fact entered the commercial pharmaceutical product development pipeline. PEG has been the most successful polymer in this regard, and its application has served as a model for nanoparticulate drug delivery systems. Given its success, PEG has become the standard antifouling polymer. A challenge for nanomedicine is to bring many of the promising PEG alternatives, such as zwitterionic materials and POxs, to the clinic.

There is still significant scope to develop more effective materials based on novel structures. We believe numerous research opportunities exist to develop new zwitterionic monomers or monomers based on peptoids or unnatural amino acids. In addition to the development of new monomers, scope also exists for developing improved grafting/surface anchoring strategies. Continued application of existing anchoring strategies (e.g., using biomimetic catechol groups) as well as the development of new approaches will likely continue to be a basis for new antifouling surfaces. Indeed, combining these surface attachment strategies with new or existing monomers will pave the way for the next generation of antifouling polymer architectures.

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