Poly (glycerol adipate) (PGA) Backbone Modifications with a Library of Functional Diols: Chemical and Physical Effects

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Abstract

Enzymatically synthesised poly(glycerol adipate) (PGA) has shown a palette of key desirable properties required for a biomaterial to be considered a ‘versatile polymeric tool’ in the field of drug delivery. PGA and its variations can self-assemble into nanoparticles (NPs) and interact at different levels with small active molecules. PGA derivatives are usually obtained by functionalising the glyceryl side hydroxyl group present along the main polymer scaffold. However, if the synthetic pathways are not finely tuned, the self-assembling ability of these new polymeric modifications might be hampered by the poor amphiphilic balance. For this reason, we have designed a straightforward one-pot synthetic modification, using a small library of diols in combination with glycerol, aimed at altering the backbone of the polymer without affecting the hydrophilic glyceryl portion. The diols introduce additional functionality into the backbone of PGA alongside the secondary hydroxyl group already present. We have investigated how extra functionalities along the polymer backbone alter the final polymer reactivity as well the chemical and biological properties of the nanoparticles. In addition, with the intent to further improve the green credentials of the enzymatic synthesis, a solvent derived from renewable resources, (2-methyl tetrahydrofuran, 2-MeTHF) was employed for the synthesis of all the PGA-variants as a replacement for the more traditionally used and fossil-based tetrahydrofuran (THF). In vitro assays carried out to evaluate the potential of these novel materials for drug delivery applications demonstrated very low cytotoxicity characteristic against NIH 3T3 model cell line.

Introduction

Glycerol is an untapped bio-renewable feedstock that is mainly produced as a by-product of biodiesel and bioethanol production and also the soap industry and crude oil. The global production of this polyol has been predicted to exceed 4000 million litres by 2026. Glycerol is applied in a huge variety
of industries such as food, cosmetics, tobacco and pharmaceuticals.\textsuperscript{2} Despite this, the growing production of glycerol, mainly from the production of biodiesel is not matched by usage in other industrial sectors and, there is a need to develop new strategies for utilisation of this valuable resource through synthesis of glycerol-based polymers.

Glycerol can act as a monomer and will polymerise to yield both linear and highly branched polymers predominantly via step-growth mechanisms in combination with diacid monomers to synthesise poly(glycerol-diacid).\textsuperscript{5} The range of diacids, or derivatives, that have been polymerised with glycerol range from C\textsubscript{4} to C\textsubscript{8} aliphatic acids (e.g. sebacic acid, succinic acid, glutaric acid, suberic acid, azelaic acid and adipic acid)\textsuperscript{6-8} in addition to aromatic diacids and aliphatic-aromatic diacids.\textsuperscript{9} The polycondensation of glycerol with this palette of diacids can afford linear, branched and cross-linked materials according to the reaction conditions adopted (e.g. nature of solvent, comonomer and catalyst, reaction temperature).\textsuperscript{10-12} In this regard, the chemo- and regioselectivity of lipases have enabled elegant, and adaptable production of linear glycerol-based polyesters under mild reaction conditions (different reaction media, low temperature and ambient pressure) preserving the secondary glyceryl hydroxyl moiety along the backbone of the resulting macromolecules.\textsuperscript{13} In this way, the available hydroxyl moiety gives rise to a polymer that can undergo a variety of further functionalisations via simple chemistry avoiding tedious and lengthy protection/deprotection intermediate steps.\textsuperscript{14,15}

In particular, the enzymatically catalysed reaction of glycerol and divinyl adipate (DVA) can produce poly(glycerol adipate) (PGA), a functionalisable, biocompatible (\textit{in vitro} and \textit{in vivo}) and biodegradable polymer.\textsuperscript{16-18} Furthermore, thanks to the chemical nature of its repetitive unit, PGA shows an amphiphilic balance capable of self-assembly into NPs in aqueous media, without the use of additional stabilizers.\textsuperscript{19} Based on this evidence, PGA is a promising functionalisable, (bio)degradable and (bio)compatible polymeric carrier ideal for drug delivery applications with enhanced chemical, physical and biological properties.\textsuperscript{16,20,21} To date, post-polymerization-functionalisation is the most common strategy to tailor the final properties of PGA and broaden its chemical diversity for specific applications. In fact, the free hydroxyl group of PGA has been coupled to a variety of molecules, with a wide range of chemical and biological properties (e.g. fatty acids, amino acids, drugs, etc.)\textsuperscript{22-27} in order to tune its ability to self-assemble into NPs for delivery of lipophilic and hydrophilic drugs as well as to produce polymeric active pro-drugs.\textsuperscript{28-34} However, if the degree of coupling is not controlled, according to the nature of the added functionalities, the unbalancing of the amphiphilic ratio and the loss of free hydroxyl moieties may negatively affect the self-assembling ability.

Others have performed one-pot enzymatic polycondensation/ring-opening polymerization (ROP) by adding pentadecalactone (PDL) to the initial DVA/glycerol mixture in order to modify the PGA backbone. The resultant PGA-co-PDL polymer showed different thermal properties and amphiphilicity when compared to the original PGA.\textsuperscript{35,36} PGA-co-PDL (and its PEGylated variant) is one of the few, if not the only example of PGA backbone modification adopted for the encapsulation of small and large active molecules.\textsuperscript{37,38} With the intent to both preserve the glyceryl portion and broaden the range of chemistries added along the PGA backbone, in this paper we set out to design a straightforward one-pot synthetic modification, using a small library of diols in combination with glycerol and divinyl adipate. In particular, the glycerol moiety was partially replaced by the diols according to the functional groups involved, maintaining a 1:1 stoichiometry molar ratio (considering that only the primary alcohols of glycerol would react in the polycondensation step). The selected diols (ethylene glycol, PEG400, 1,4-butynediol, 1,4-butanediol, 1,3-benzendimethanol, 2-hydroxyethyl disulfide and 1,6-n-hexanediol) may introduce either additional functionalities able to tune the amphiphilic balance of the final polymeric materials or functionalisable chemical handles directly into the backbone of PGA.
We aim to perform a series of backbone modifications with well-known chemistries and investigate some of the effects on polymer reactivity as well as the physical and biological properties of the nanoparticles that can be formed. Additionally, in order to further improve the green credentials of the enzymatic synthesis, a solvent derived from renewable resources, (2-methyl tetrahydrofuran, 2-MeTHF) will be employed, the first time for PGA synthesis, to replace the more traditionally used and fossil-based tetrahydrofuran (THF). 39–42

Experimental Section

Materials

Novozym 435 lipase ([9001-62-1], derived from C. antarctica (>5000 U/g) and immobilized on an acrylic macroporous resin, was kindly donated by Novozymes A/S, Denmark. Glycerol and all the novel diols (ethylene glycol, PEG400, 1,4-butynediol, 1,4-butenediol, 1,3-benzendimethanol, 1,6-n-hexanediol and 2-hydroxyethyl disulfide) were purchased from Sigma–Aldrich UK. Divinyl adipate [4074-90-2] was purchased from TCI America and SPI supplies. All chemicals were used as received. Solvents were purchased from Fischer Scientific UK and used without further purification unless otherwise stated. Water was deionised before use. Solvent evaporation was performed using a rotary evaporator under reduced pressure.

General Methods and Instrumentation

Nuclear Magnetic Resonance Spectroscopy (NMR): Polymer formation and repetitive unit chemical structure assignment were determined using 1H-NMR spectroscopy. Approximately 4 mg of sample were dissolved in 2 mL of acetone-d₆ or CDCl₃ and analysed using a Bruker DPX 400 MHz spectrometer operating at 400 MHz (1H), assigning chemical shifts in parts per million (ppm). MestReNova 6.0.2 copyright 2009 (Mestrelab Research S. L.) was used for analysing the spectra.

Gel Permeation Chromatography (GPC): was performed in THF (HPLC grade, Fisher Scientific) as the eluent at 40 °C using two Agilent PL-gel mixed-D columns in series, an injection loop of 50 μL, with a flow rate of 1 mL min⁻¹. A differential refractometer (DRI) was used for the detection of samples (solution containing approximately 4 mg dissolved in 2 mL of THF, filtered in 0.22 μm Teflon filter). The system was calibrated using poly(methyl methacrylate) standards with average molecular weight in the range from 540 to 1.02·10⁶ g mol⁻¹ and dispersity (Ð) close to 1.0

Differential Scanning Calorimetry (DSC): was used to determine the thermal transition of the polymers produced. The analysis was performed on a TA-Q2000 (TA instruments), which was calibrated with indium and sapphire standards under N₂ flow (50 mL min⁻¹). The sample (5-10 mg) was weighed into a T-zero sample pan (TA instruments) with a reference T-zero pan remaining empty. Both lid pans were pin-holed and all samples were heated at a rate of 10 °C min⁻¹, from -90 °C to 200 °C. To remove any thermal history of the individual samples two heating cycles were recorded and the second heating cycle was used to determine the glass transition temperature (Tₛ).
Dynamic Light Scattering (DLS): Particle size analyses were performed by DLS utilizing a Zetasizer Nano spectrometer (Malvern Instruments Ltd) equipped with a 633 nm laser at a fixed angle of 173°. Nanoparticles were prepared at a concentration of 1 mg/mL adopting a simple solvent displacement methodology (acetone/PBS ratio 1:5). Samples were equilibrated at 25 °C for 30 seconds prior to measurements. All experiments were performed in duplicate averaging 10 scans per run of the same sample.

Water Contact Angle (θ): Water contact angle (WCA) values were measured at 25 °C using a Kruss DSA 100 equipped with dedicated software. Samples were prepared by coating microscopic glass slides with polymer thin films by solvent evaporation of 3 mg/mL polymer solutions in acetone. Six measurements were recorded for each polymer.

“Traditional” and “Greener” PGA Synthesis

PGA was synthesized (Figure 1) by enzymatic polymerization of divinyl adipate (DVA) and glycerol following a protocol adapted from Taresco et al. Glycerol (12.5 mmol) and DVA (12.5 mmol) were poured into a 20 mL glass vial and dissolved into THF (10 mL), or in the greener alternative 2-MeTHF (10 mL). To this mixture, Novozym 435 (0.11 g) was added. The resulting mixture was stirred at 200 rpm with a magnetic stirrer and submerged in an oil bath at 50 °C for 1.5 h, 3 h, 5 h or 24 h. A needle was inserted through the rubber septum in order to facilitate the release of acetaldehyde. The removal of this side product favoured the polymerisation process due to the step-growth nature of the process. The reaction was stopped by simply removing the immobilised enzyme by filtration, followed by evaporation of the solvent under reduced pressure. The residual material was kept under vacuum at around 25 °C for three days to remove the residual solvent. The resultant highly viscous yellow liquid was stored at -20 °C in order to minimise possible hydrolysis side reactions.

One-Pot PGA Backbone Diol-Alteration Synthesis in 2-MeTHF

PGA-diol was synthesized (Scheme 1) by enzymatic polymerization of divinyl adipate (DVA), glycerol and one of the diols from a protocol adapted from Taresco et al. Glycerol (6.25 mmol) and the selected functionalised diols (6.25 mmol) were added to DVA (12.5 mmol) in a 20 mL glass vial and dissolved into 2-MeTHF (10 mL). In particular, the glycerol moiety was partially replaced by the diols according to the functional groups involved, maintaining a 1:1 stoichiometry molar ratio (considering that only the primary alcohols of glycerol would react in the polycondensation step). To this mixture, Novozym 435 (0.11 g) was added. The resulting mixture was stirred at 200 rpm with a magnetic stirrer and submerged in an oil bath at 50 °C for 3 h. A needle was inserted through the rubber septum in order to facilitate the release of acetaldehyde. The removal of this side product favoured the polymerisation process due to the step-growth nature of the process. The reaction was stopped by simply removing the immobilised enzyme by filtration, followed by evaporation of the solvent under reduced pressure. The residual material was kept under vacuum at around 25 °C for three days to remove the residual solvent. The resultant polymer was stored at -20 °C in order to minimise possible hydrolysis side reactions.
Thiol–ene chemistry of PGA-alkene and PGA-alkyne

The reactivity of the ene- and yne- functionalities introduced in PGA-alkene and PGA-alkyne was probed by a radical thiol–ene reaction with benzyl mercaptan as the model thiol. To this aim, PGA-alkene or PGA-alkyne (15 mg) were dissolved in 100 μL of DMSO-\textsubscript{d6}. 2,2-Dimethyl-2-phenyl acetophenone (DMPA) (25 μL aliquot of a 80 mg mL\textsuperscript{-1} solution in DMSO-\textsubscript{d6}) and benzyl mercaptan (10 equiv.) were added and the reactions were then irradiated by UV at 365 nm for 120 minutes. A final \textsuperscript{1}H NMR spectrum was recorded.

Nanoparticle formation

Nanoparticles were prepared by a nanoprecipitation method.\textsuperscript{20} Polymers (10 mg) were dissolved in acetone (1 mL). The polymeric solution was then added dropwise to deionised water (10 mL) under constant stirring at 550 rpm. Nanoparticle dispersions were formed through solvent exchange between water and acetone. The final dispersion was then left stirring overnight at room temperature in order to reach complete acetone evaporation, final NPs concentration of 1 mg mL\textsuperscript{-1}.

Coumarin-6 (Cou6) encapsulation study

The encapsulation procedure of Cou-6 was adopted from previous reported protocols.\textsuperscript{43,44} Polymer (10 mg) was dissolved in acetone (1 mL). An aliquot (0.5 mL) of a stock solution of Cou6 (3.21 mM in acetone) was added to the polymer solution. The final acetone solution was added dropwise into deionised water (10 mL) under constant stirring at 550 rpm. The final dispersion was then left stirring overnight at room temperature in order to reach complete acetone evaporation. Nanoparticles/dye dispersions were finally filtered with a 0.22 μm filter. Cou6 control was formulated in the same way but without the addition of any polymer.

ΔA% determination

The absorbance of NPs/dye suspensions was measured by using an UV–vis multi-well plate reader at λ=470 nm. The apparent-solubility (ΔA%) value of each formulation was calculated according to the previously developed equation:\textsuperscript{45}

\[
\Delta A\% = \frac{\Delta A}{A_0} \times 100 = \left(\frac{A - A_0}{A_0}\right) \times 100
\]

where \(A\) is the absorbance of the NPs dispersion and \(A_0\) corresponds to the absorbance of the drug formulation in water.
Cytotoxicity of polymer formulations

Presto Blue™ cell viability assay was performed to assess the effect of the formulations on primary mouse embryonic fibroblast cells, NIH 3T3 cell line. The cells were grown in Dulbecco’s modified eagle medium (DMEM) supplemented with 10% (v/v) foetal calf serum and 1% antibiotics (100 units mL⁻¹ penicillin, 100 mg mL⁻¹ streptomycin, and 0.25 mg mL⁻¹ amphotericin B; Life Technologies). Cells were cultured until they reached 80% confluency and subsequently detached from the culture surface using trypsin/EDTA (0.25%/0.02% w/v), centrifuged at 200 x g for 5 min and resuspended in the culture medium. Cells were seeded in a 48 well plate at a density of 10,000 cells per well and allowed to attach for 24 h before the viability experiments. Formulations were dissolved in a complete medium at a concentration of 100 µg mL⁻¹ and 500 µL of each formulation were transferred to the wells containing the cells (n=4). Cells were exposed to the formulations for 24 and 48 h. Cells cultured in a complete culture medium only were used as a negative control. After 24 or 48 h, media was removed and cells were then washed twice with warm PBS and 100 µL 10% (v/v) PrestoBlue reagent diluted in phenol red-free medium applied per well for 30 minutes. The resulting fluorescence was measured at 560/600 nm (λex/λem). Relative metabolic activity was calculated by setting values from the negative control as 100%.

Results and Discussions

Analysis of PGA synthesised in THF and 2-MeTHF and effect of reaction time

PGA was successfully synthesised in both the commonly used THF and 2-MeTHF. This was confirmed by ¹H NMR and GPC analyses. The disappearance of the divinyl peaks at 7.30, 4.87 and 4.59 ppm and change in shape followed by a variation in chemical shift of the glycerol peaks at 4.00 and 4.26 ppm confirmed that the polymerisation occurred (Figure 1 and Figure S1).
The presence of 1,3-disubstituted (target reaction) 1,2-disubstituted and 1,2,3-trisubstituted glyceride groups (side events) in PGA has been described before, with the latter leading to polymer branching as well as to the decrease of the number of polymer hydroxyl groups. However, the 1,2-disubstituted unit, as for the 1,3-disubstituted, provides a final structure bearing a free hydroxyl group, therefore, it was not considered in the branching calculation. On the other hand, the methine proton from the 1,2,3-trisubstituted glycerol unit at 5.30 ppm (Figure 1, inset) was used to calculate the degree of branching of the PGA polymers, following a previously reported strategy to calculate the amount of trisubstituted glycerol in PGA. The integrals of the methine proton at 5.30 ppm and the CH$_2$ peaks of the adipic repeating units at 1.66 and 2.39 ppm were compared to estimate the degree of branching. In order to evaluate the effects of the solvent on the final polymer architecture, we calculated the degree of branching for each polymer produced in the two solvents. This was found to be circa 7 and 12% (Table 1) for PGAs synthesised at 50 °C for 24 h in THF and 2-MeTHF, respectively. In particular, a batch-to-batch variation in 1,2,3-trisubstituted glycerin was observed with a slight increase related to the reaction time (variation from 1 to 3% when increasing the reaction time up to 24h). In addition, the small variation might be due to different degradation events in the two solvents during the screened reaction period, presumably promoted by lipase during long contact times.

Figure 1. (TOP) PGA synthetic scheme and reagent conditions. Where R takes into account possible branching. (BOTTOM) Stacked $^1$H NMR of PGA produced in 2MeTHF (red spectrum) and THF (black spectrum). Inset: effect of the reaction medium on PGA degree of branching.
Indeed, for polymerisations carried out for 5 h, at the same reaction temperature, in each solvent, the degree of branching was found to be similar, 4% (in THF) and 6% (in 2-MeTHF) (Table 1). These preliminary observations confirmed that the adoption of 2-MeTHF as a reaction medium did not significantly affect the molecular weight distribution or the degree of branching, at least for short reaction times. In addition, with the intent of determining the optimum reaction time, further kinetic studies were carried out in this “greener” alternative solvent for 1.5 h and 3 h (at 50 °C). After 1.5 h the degree of branching was only 3%, and the number-average molecular weight of the polymer was lower than the 5 h and 24 h comparisons. In addition to this, the residual vinylic protons from the unreacted DVA or, due to the lower molecular weight, from the end group could be seen at 7.30, 4.87 and 4.59 ppm in the 1H NMR spectrum (Figure S2). On the other hand, after 3 h, the degree of branching was found to be around 6% and the resulting PGA molecular weight was around 3200 g mol⁻¹ (D = 2.70). It can be noticed that both the degree of branching and the molecular weight of the polymer after 3 h polymerisation were comparable with those obtained after 5 h polymerisation and for this reason we adopted 3 h as the best compromise between reaction time and conversion in final polymer for all the PGA backbone alteration polymerisations. It is important to mention that the reaction temperature and timing adopted in the present work are within the working window of Novozym 435 lipase previously reported for 2-MeTHF. In fact, it has been demonstrated that Novozym 435, in 2-MeTHF, can successfully catalyse both ring opening polymerisation and polycondensations with temperatures ranging from 30 to 85 °C and for reaction periods over 6h.39,42

**Table 1.** Number-average molecular weight (Mₐ), dispersity (D) and degree of branching of PGAs prepared in THF and 2-MeTHF at different reaction times at 50°C.

<table>
<thead>
<tr>
<th>Reaction Media</th>
<th>Reaction time (h)</th>
<th>Mₐ (g mol⁻¹)</th>
<th>D</th>
<th>Degree of branching (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>THF</td>
<td>24</td>
<td>3000</td>
<td>2.70</td>
<td>~7±2</td>
</tr>
<tr>
<td>THF</td>
<td>5</td>
<td>3600</td>
<td>2.80</td>
<td>~4±1</td>
</tr>
<tr>
<td>2-MeTHF</td>
<td>24</td>
<td>3000</td>
<td>3.40</td>
<td>~12±3</td>
</tr>
<tr>
<td>2-MeTHF</td>
<td>5</td>
<td>3500</td>
<td>2.90</td>
<td>~6±1</td>
</tr>
<tr>
<td>2-MeTHF</td>
<td>3</td>
<td>3200</td>
<td>2.70</td>
<td>~6±1</td>
</tr>
<tr>
<td>2-MeTHF</td>
<td>1.5</td>
<td>2600</td>
<td>2.10</td>
<td>~3±1</td>
</tr>
</tbody>
</table>

* Molecular mass determined by GPC at 40 °C, using THF as the mobile phase. **Due to the lack of purification, oligomers peaks can be noticed. These have been taken into account in the total integration. b) Polymer branching, referred to the 1,2,3-trisubstituted unit, calculated by ¹H NMR analysis (comparison between the integrals of the methine CH peak of trisubstituted glycerol at 5.30 ppm, to that at 1.66 or 2.39 ppm of CH₂CH₂C(O) of adipic acid repeating units). Branching measurements taken out of two different reactions performed in the same conditions.

**Synthesis of PGA with different diols in the backbone**

The stoichiometric addition of functionalised diols, to replace part of the glycerol, was adopted to investigate the possible range of chemical and physical properties of the final PGA derivatives. Only commercially available diols were selected (Scheme 1) and used without further purifications.
H NMR and GPC analyses confirmed that the new functionalities have been integrated into the polymer backbone. In particular, the O-CH₂ diol peak is shifted upon esterification with the adipic moiety (Figure S3) and a general peak broadening was observed (Figure S3, PGA-1,4-butynediol example). Moreover, the addition of a second diol noticeably decreases the final degree of branching, from ca. 6% to 1-4% (Table 2). This is likely due to the decrease in the number of secondary hydroxyl groups that can take part in the branching. Similarly, GPC traces showed a broad (2.00 ≤ D ≤ 2.90) polymeric peak for each derivative, ranging from about Mn 2200 up to 6400 g/mol and traces of oligomers as previously observed for the bare PGA backbone (Figure S3 shows successful modification, further examples are also presented in the SI). The modified PGAs were found to be amorphous materials apart from the 1,6-n-hexanediol variation which showed both a Tg at around -50 °C and a weak melting transition at around 13 °C (DSC analysis of the polymers section in supporting info). This phenomenon might be due to the symmetry between the hexanediol and adipic chains, in fact, when the disulfide group was added no melting transition could be observed. The set of polymers displayed a single glass transition temperature (Tg) in the range of -29 to -50 °C (Table 2); close to the Tg of -28 °C for the original PGA which is usually a viscous waxy liquid at room temperature. In general, the use of aliphatic alkyl diol (ethylene glycol and hexanediol) improved polymer flexibility with a consequent decrease in Tg with respect to the pure PGA. As expected, the length of the alkyl chain was found to affect polymer Tg. Specifically, PGA-hexanediol showed a lower Tg than PGA-ethylene glycol. Controversially, when PEG400 was used in place of hexanediol we did not observe a further decrease in Tg. Presumably, in this case, the presence of ether groups able to establish H-bonds with the PGA hydroxyl groups, in some ways, reduced the freedom of movement of polymer chains.

The propensity of a drop of water to wet the surface of a material is defined as contact angle of the specific material (θ). The introduction of the functionalised diol substituents will alter the hydrophobic/hydrophilic behaviour. In practice high contact angles corresponding with hydrophobic surfaces, and low angles more hydrophilic surfaces. PGA variants possess experimental θ ranging from circa 19 to 70°. In particular, high hydrophilic diols such as PEG400 conferred high hydrophilicity increasing the wettability of the material surface when compared to the pristine PGA (58°) and the other variants, confirming the rearrangement of the hydrophilic PEGylated chain towards the water droplet. On the other hand, more hydrophobic moieties like 1,3-benzendimethanol and 1,6-n-hexanediol endowed the final polymers with a hydrophobic property translated in higher contact angles (62.5 to 66.9).
Table 2. Number-average molecular weight (Mn), dispersity (D), degree of branching, Tg and water contact angle of PGA-derivatives.

<table>
<thead>
<tr>
<th>Entry</th>
<th>PGA-modification</th>
<th>Acronyms</th>
<th>M&lt;sub&gt;n&lt;/sub&gt; (g mol&lt;sup&gt;-1&lt;/sup&gt;)&lt;sup&gt;a&lt;/sup&gt;</th>
<th>D&lt;sup&gt;a&lt;/sup&gt;</th>
<th>Degree of branching (%)&lt;sup&gt;b&lt;/sup&gt;</th>
<th>T&lt;sub&gt;g&lt;/sub&gt; (°C)&lt;sup&gt;c&lt;/sup&gt;</th>
<th>Water contact angle (°)&lt;sup&gt;d&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>PGA</td>
<td>PGA</td>
<td>3200</td>
<td>2.90</td>
<td>~6</td>
<td>-28</td>
<td>58.2 ± 0.4</td>
</tr>
<tr>
<td>2</td>
<td>PGA-ethylene glycol</td>
<td>PGA-EG</td>
<td>2200</td>
<td>2.00</td>
<td>~2</td>
<td>-41</td>
<td>59.3 ± 0.3</td>
</tr>
<tr>
<td>3</td>
<td>PGA-PEG400</td>
<td>PGA-PEG400</td>
<td>3500</td>
<td>1.70</td>
<td>~2</td>
<td>-50</td>
<td>19.2 ± 0.6</td>
</tr>
<tr>
<td>4</td>
<td>PGA-1,4-butenediol</td>
<td>PGA-1,4BY</td>
<td>2700</td>
<td>2.10</td>
<td>~2</td>
<td>-30</td>
<td>57.9 ±1.4</td>
</tr>
<tr>
<td>5</td>
<td>PGA-1,4- butenediol</td>
<td>PGA-1,4BE</td>
<td>2600</td>
<td>2.60</td>
<td>~3</td>
<td>-47</td>
<td>48.6 ± 0.4</td>
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<tr>
<td>6</td>
<td>PGA-1,3-benzendimethanol</td>
<td>PGA-1,3Ph</td>
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<td>PGA-1,6-n-hexanediol</td>
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<td>~3</td>
<td>-50</td>
<td>66.9 ± 1.6</td>
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<td>8</td>
<td>PGA-2-hydroxyethyl disulfide</td>
<td>PGA-SS</td>
<td>4400</td>
<td>1.65</td>
<td>~2</td>
<td>-38</td>
<td>58.0 ± 1.4</td>
</tr>
</tbody>
</table>

<sup>a</sup> Molecular mass determined by GPC at 40 °C, using THF as the mobile phase. <sup>b</sup> Due to the lack of purification, oligomers peaks can be noticed. These have been taken into account in the total integration. <sup>c</sup> Polymer branching calculated by <sup>1</sup>H-NMR (comparison between the integrals of the methine CH peak of trisubstituted glycerol at 5.30 ppm, to that at 1.66 or 2.39 ppm of CH<sub>2</sub>CH_{2}C(O) of adipic acid repeating units). <sup>d</sup> Measured by DSC. <sup>e</sup> Measured by Kruss DSA 100.

PGA reactive and responsive variations

A particular example of modification is the incorporation of alkene and alkyne moieties into the polymer backbone. This can then lead to a highly versatile approach to a variety of additional functionalities by exploiting a series of well-established reaction pathways, including azide-alkyne cycloadditions and thiol-ene modifications.<sup>48,49</sup> To prove the reactivity of these ene- and yne-functionalities and to highlight the possibility to endow the PGA backbone with additional properties, we probed the reactivity of the double bond by a radical thiol–ene reaction with benzyl mercaptan as the model thiol (Scheme 2). The macromolecules were reacted with 10 equivalents of the thiol with DMPA as a photo-initiator, followed by irradiation at 365 nm for 120 minutes.<sup>50,51</sup> The intensity of the resonance related to the alkenyl protons, at 5.9 ppm, reduced by half when compared to the CH<sub>2</sub> peaks of the adipic repeating units at 2.39 ppm (Figure S4).

![Scheme 2. PGA-1,4BE thiol-ene reaction scheme.](image)

In addition to the abovementioned modifications, we also investigated the insertion of an additional moiety into the backbone to bring additional functionality. 2-Hydroxyethyl disulfide is a readily available and well-known redox-responsive linear diol.<sup>52,53</sup> As a proof of concept and with the aim of producing a multiresponsive (redox responsive and biodegradable) polymer in a straightforward synthetic pathway, we substituted 1,6-n-hexanediol with 2-hydroxyethyl disulfide. The incorporation of this diol was successful as confirmed by <sup>1</sup>H NMR and GPC (Figure 2). In addition, the polymer can
self-assemble in water producing NPs with sizes around 115 nm and PDI of 0.08. This modification may provide interesting opportunities in the future as a smart multiresponsive polymeric platform for drug delivery applications.

Figure 2. (LEFT) Stacked $^1$H-NMR spectra of PGA vs PGA-disulfide and (RIGHT) GPC chromatographs of PGA and PGA-disulfide (presence of oligomers can be noticed in both chromatograms). Due to solubility reasons, the PGA-disulfide variation has been synthesised in THF.
Polymer Cytotoxicity

These novel materials show potential for pharmaceutical and more specifically for drug delivery applications, so an initial assessment of their in vitro toxicity on NIH 3T3 model cell line was conducted. Cytocompatibility tests of the modified PGAs resuspended in cell culture aqueous medium at a fixed concentration of 100 μg mL⁻¹ showed no significant in vitro cytotoxicity; no decrease in cellular metabolic activity over 24-48 h on the cell type studied (Figure 3a and 3b). One should bear in mind that in accordance with ISO 10993-5:2009 if the cell viability is greater than or equal to 70% in relation to control group (100% viability) the polymer material can be considered non-cytotoxic.

Figure 3. Cytocompatibility of NPs on NIH 3T3 cells after a) 24h and b) 48h. No significant drop in vitality below 80% was observed. Culture medium DMEM was used as positive control. Data are presented as mean ± S.D. (n = 4).
Self-assembling ability of hydrophilic/hydrophobic PGA variants

Amphiphilic balancing to tune polymer self-assembly and interaction with small molecules was achieved in the synthesized PGA variants by incorporating a range of hydrophilic and hydrophobic moieties into the polymer backbone. In the nanoprecipitation step the different modified PGAs self-assembled in water into nanoaggregates, with diameters ranging from ca. 88 nm up to 120 nm and very narrow particle size distributions exhibiting PDI equal or below 0.1 (Figure 4A, Table 3). PGA-ethylene glycol showed similar size and PDI values of the bare PGA, hinting at a minimal effect on the final self-assembling behaviour of the short diol chain. While both the 1,6-n-hexanediol and PEG400 PGA variations gave small NPs (around 88 and 92 nm, respectively) the two modifications provided opposite hydrophobic/hydrophilic behaviours from their contact angle values. This highlights the importance of the amphiphilic modification of the repeating unit. On the other hand, the incorporation of a bulky aromatic ring, in the case of PGA-1,3-benzenedimethanol, led to an increase of around 15 nm of the hydrodynamic size of the final NPs compared to the bare PGA (Table 3).

Table 3. Size and PDI of empty and Cou6 loaded nanoparticles. a

<table>
<thead>
<tr>
<th>PGA-modification</th>
<th>Empty nanoparticles</th>
<th>Cou6 loaded nanoparticles</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Average size (nm)</td>
<td>PDI</td>
</tr>
<tr>
<td>PGA</td>
<td>102.8 ± 4</td>
<td>0.104 ± 0.01</td>
</tr>
<tr>
<td>PGA-EG</td>
<td>108.6 ± 2</td>
<td>0.077 ± 0.01</td>
</tr>
<tr>
<td>PGA-PEG400</td>
<td>92.1 ± 1</td>
<td>0.102 ± 0.01</td>
</tr>
<tr>
<td>PGA-1,3Ph</td>
<td>117.6 ± 1</td>
<td>0.079 ± 0.01</td>
</tr>
<tr>
<td>PGA-1,6H</td>
<td>87.5 ± 2</td>
<td>0.052 ± 0.03</td>
</tr>
</tbody>
</table>

a All data were produced by DLS measurements.
Coumarin6 enhancing solubility in an aqueous environment.

To prove the encapsulation ability of the novel PGA-variants, the highly hydrophobic Coumarin-6 (Cou6) fluorescent dye was used as a model. Cou6 is a water-insoluble small molecule that can be used as a model to mimic the behaviour of lipophilic drugs for initial evaluation of drug delivery- and drug release experiments. Cou6 was co-nanoprecipitated in water with and without (control) PGA modifications. In order to quickly rank the polymers in terms of Cou6 apparent-solubility enhancement, ΔA% values were calculated (Figure 4b).

PEG400, showed the lowest ΔA% likely due to a shorter hydrophobic counterpart in the repetitive unit compared to the rest of the library. PGA-EG and PGA-1,3Ph, instead, showed higher ΔA% (2292 and 2566, respectively) and therefore a higher ability to solubilise/encapsulate Cou6 when compared to PGA (1633) and PGA-PEG400 (1075). This is likely because of the increased hydrophobicity portion in the repetitive units able to interact with the small model dye. However, the aromatic modification, despite a higher hydrophobicity contribution compared to the other modifications, did not show a remarkable dye interaction, possibly due to the bulkiness of the ring. The aromatic ring might reduce the space available inside the hydrophobic core of the nanoaggregates for the Cou6. On the other hand, PGA-1,6-H ΔA% was around 4000, almost 2.5 times higher than that of the bare PGA. The addition of the 1,6-n-hexanediol moiety seemed to be a perfect balance of amphiphilicity, chain length

Figure 4. a) DLS intensity of nanoparticles formed from PGA and its variants b) ΔA% of Cou6 vs the panel of PGA-variants.
and chain flexibility which allowed for self-assembling of well-defined NPs as well as providing an improvement of the apparent solubility in water of Cou6.

Conclusions

We have demonstrated that it is possible to alter the chemical composition of the PGA backbone by utilising a small library of functionalised/functionalisable diols with a straightforward one-pot synthetic modification of the common enzymatic pathway. The selected diols introduce additional chemical handles, such as double/triple bonds, PEGylated chains and aromatic rings, directly into the backbone of PGA. In addition, tuning of the PGA amphiphilic polymer backbone was demonstrated by adding a series of diols with different hydrophilic/hydrophobic behaviours. The reactivity of the ene-PGA variants was probed by a simple UV activated thiol-ene reaction. Self-assembly and the encapsulation of a water-insoluble model molecule were demonstrated to be directly related to the nature of the added functionalised diol. Furthermore, in vitro cell-viability assays demonstrated that the produced PGA variants are nontoxic and may be considered appropriate vehicles for future investigation as drug delivery carriers. In addition, in order to enhance the green credentials of the enzymatic synthesis, a solvent derived from renewable resources, (2-methyl tetrahydrofuran, 2-MeTHF) was employed for the synthesis of all the PGA-variants as replacement of the more traditionally used and fossil-based tetrahydrofuran (THF).

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