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New Aldehyde-Functional Methacrylic Water-Soluble Polymers

Emma E. Brotherton, Craig P. Jesson, Nicholas J. Warren, Mark J. Smallridge, and Steven P. Armes*

Abstract: Aldehyde groups enable facile conjugation to proteins, enzymes, oligonucleotides or fluorescent dyes, yet there are no literature examples of water-soluble aldehyde-functional vinyl monomers. We report the synthesis of a new hydrophilic cis-diol-based methacrylic monomer (GEO5MA) by transesterification of isopropylidenglycerol penta(ethylene glycol) using methyl methacrylate followed by acetone protection via acid hydrolysis. The corresponding water-soluble aldehyde monomer, AGE5OMA, is prepared by aqueous periodate oxidation of GEO5MA at 22°C. RAFT polymerization of GEO5MA yields the water-soluble homopolymer, PGE5OMA. Aqueous periodate oxidation of the terminal cis-diol units on PGE5OMA at 22°C affords a water-soluble aldehyde-functional homopolymer (PAGE5OMA). Moreover, a library of hydrophilic statistical copolymers bearing cis-diol and aldehyde groups was prepared using sub-stoichiometric periodate/cis-diol molar ratios. The aldehyde groups on PAGE5OMA homopolymer were reacted in turn with three amino acids to demonstrate synthetic utility.

Introduction

Aldehydes are extremely useful functional groups in synthetic organic chemistry: they can be oxidized to give carboxylic acids, reduced to afford alcohols, undergo Schiff base chemistry and also form (hemiacetals). In the field of synthetic polymer chemistry, aldehyde-based initiators have been utilized to prepare various types of aldehyde-functional polymers. Alternatively, Bilgic and Klok derivatized poly(2-hydroxyethyl methacrylate) brushes under oxidative conditions in order to introduce aldehyde groups for subsequent oligonucleotide conjugation. However, surprisingly few aldehyde-functional monomers have been reported in the literature. Most of these examples are hydrophobic (e.g. 4-vinylbenzaldehyde) and hence produce water-insoluble polymers. This is unfortunate, because aldehyde groups enable facile conjugation to peptides/proteins and water-soluble dyes in aqueous solution under mild conditions. In principle, this problem can be circumvented by statistical copolymerization of the hydrophobic aldehyde-functional monomer with a sufficiently hydrophilic comonomer. Alternatively, the incorporation of a terminal protected aldehyde moiety onto a poly(ethylene glycol) (PEG) chain has been utilized to confer aldehyde functionality under aqueous conditions. Nevertheless, despite the remarkable progress made in synthetic polymer chemistry over the past few decades, there seem to be few, if any, literature examples of hydrophilic aldehyde-functional vinyl monomers and their corresponding water-soluble homopolymers.

One well-known route to aldehyde-terminated water-soluble polymers is the selective oxidation of the minor fraction of cis-diol units within poly(vinyl alcohol). This water-soluble polymer can be obtained via hydrolysis of poly(vinyl acetate), which contains such cis-diols as defect sites resulting from a small amount of head-to-head coupling during the free radical homopolymerization of vinyl acetate. Oxidation is readily achieved in aqueous solution under mild conditions using sodium periodate to afford aldehyde-capped poly(vinyl alcohol) chains. Inspired by this well-established chemistry, we recently decided to investigate the periodate oxidation of poly(glycerol monomethacrylate) (PGMA) to produce an aldehyde-functional methacrylic polymer (Scheme 1; Supporting Information). However, periodate oxidation of a 10% w/w aqueous solution of PGMA at 22°C merely produced a macroscopic precipitate. This suggests that the target aldehyde-functional methacrylic homopolymer (PAGMA) is actually hydrophobic. In principle, such precipitation could be the result of reaction between

[NaIO₄][cis-diol] = 1.0
10% w/w, H₂O, 22 °C
- formaldehyde

Scheme 1. Selective oxidation of a water-soluble PGMA₃₉ homopolymer precursor using a stoichiometric amount of sodium periodate in aqueous solution at 22°C affords PAGMA₃₉ as a water-insoluble precipitate.
the cis-diol and aldehyde units at intermediate conversion. However, reaction exotherms (data not shown) and visual inspection of the reaction mixtures suggest that the timescale required for the cis-diol oxidation is much shorter than that for precipitation. Thus, it seems more likely that intermolecular crosslinking occurs between geminal diols and aldehydes (Supporting Information, Scheme S1).

In view of these problems, we designed a new cis-diol-based methacrylic monomer (GEO5MA; Scheme 2a). We envisaged that the pendant oligo(ethylene glycol) moiety in GEO5MA should confer sufficient hydrophilic character to ensure water solubility after converting its terminal cis-diol group into an aldehyde via periodate oxidation to form either AGE5MA monomer (Scheme 2b) or the corresponding PAGE5MA homopolymer.

**Results and Discussion**

The two-step synthesis of GEO5MA monomer was conducted on a 1.2 kg scale via 1) transesterification of isopropylidene glycerol penta(ethylene glycol) using methyl methacrylate to afford IPGEO5MA (Scheme 2a) and 2) acid hydrolysis to remove the acetone protecting group (Supporting Information). The chemical structure of this new methacrylic monomer was confirmed by \(^1\)H and \(^{13}\)C NMR spectroscopy (Figure 1a; Supporting Information, Figure S1a), mass spectrometry, elemental microanalysis and FT-IR spectroscopy (Supporting Information). The integrated signals in the \(^1\)H NMR spectrum are consistent with the proposed monomer structure. Its \(^{13}\)C NMR spectrum contained ten distinct signals. A characteristic signal at 166 ppm was assigned to the ester carbonyl carbon; its relatively low intensity is attributed to the slow relaxation time for such quaternary carbon atoms.\(^{[45]}\) The presence of a methacrylate group is confirmed by signals at 135 and 127 ppm. Several signals between 62.6 and 71.3 ppm are assigned to the pendant oligo(ethylene glycol) chain and include characteristic signals for the carbons attached to hydroxyl groups. According to mass spectrometry, the number of ethylene glycol units per oligo(ethylene glycol) group ranged from 2 to 7, with a mean value of 5.

Oxidation of a 10% w/w aqueous solution of GEO5MA using a NaIO\(_4\)/cis-diol molar ratio of unity (Scheme 2b) led to essentially complete oxidation of the terminal cis-diol units within 5 min at 22°C, as confirmed by \(^1\)H NMR spectroscopy (Figure 1). The structure of this new AGE5MA monomer was confirmed by mass spectrometry, elemental microanalysis, \(^1\)H and \(^{13}\)C NMR (Figure 1b; Figure S1b) and FT-IR spectroscopy (Figure S3a). Two new signals appear at 9.52 and 5.09 ppm in the \(^1\)H NMR spectrum for AGE5MA, corresponding to an aldehyde group and a geminal diol, respectively. The aldehyde/geminal diol molar ratio was 0.034, which indicates that AGE5MA exists primarily in its hydrated geminal diol form in D\(_2\)O (Figure 1b). Similar observations have been reported for other hydrophilic aldehydes in aqueous solution, such as acetaldehyde (Figure S2).\(^{[46–49]}\) During the periodate oxidation of GEO5MA to form AGE5MA, the starting material can in principle react with the product to generate dimethacrylate species via (hemi)acetal chemistry.\(^{[1]}\) In practice, the final product contains less than 1% dimethacrylate impurity as estimated by \(^1\)H NMR spectroscopy. The \(^{13}\)C NMR spectrum also shows the appearance of two new signals at 169.5 and 88.0 ppm, which correspond to the aldehyde carbon and the geminal diol carbon, respectively. After purification by extraction with CH\(_2\)Cl\(_2\), the RAFT aqueous solution polymerization of AGE5MA was conducted using a dicarboxylic acid-functionalized water-soluble RAFT agent (CECPA) to
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![Diagram of polymerization](image)

### Table 1: Extent of oxidation, DMF GPC molecular weight and dispersity data for the selective oxidation of PGEO5MA<sub>n</sub> in aqueous solution at 22°C using (sub-)stoichiometric NaIO<sub>4</sub>/cis-diol molar ratios ranging between 0 and 1.0.

| NaIO<sub>4</sub>/cis-diol molar ratio | Extent of oxidation [%] | M<sub>n</sub> [kgmol<sup>-1</sup>] | D |
|-------------------------------------|------------------------|----------------------------------|---|
| 1.00                                | > 99                   | 16.5                             | 1.22 |
| 0.75                                | 78                     | 15.9                             | 1.24 |
| 0.50                                | 49                     | 16.8                             | 1.21 |
| 0.10                                | 11                     | 17.4                             | 1.22 |
| 0.00                                | 0                      | 17.2                             | 1.18 |

However, only a minor fraction of monomer repeat units may need to be converted into aldehyde groups for certain applications. Thus, partial oxidation of a PGEO5MA<sub>n</sub> precursor using sub-stoichiometric quantities of NaIO<sub>4</sub> oxidant relative to its cis-diol groups was also investigated (schematic in Figure 3a).

Accordingly, utilizing NaIO<sub>4</sub>/cis-diol molar ratios of 0.10, 0.50 or 0.75 produced a series of water-soluble P(GEO5MA<sub>stat</sub>-AGEO5MA<sub>n</sub>) statistical copolymers with approximate degrees of aldehyde functionality of 0.11, 0.49 and 0.78 respectively, as estimated from <sup>1</sup>H NMR spectroscopy studies (Table 1, Figure 3). Thus, the target degree of aldehyde functionality is always achieved (within experimental error). DMF GPC analyses confirmed that neither partial nor full oxidation of the PGEO5MA<sub>n</sub> homopolymer had a significant effect on its molecular weight distribution (Table 1; Figure S4). Moreover, using a slight excess of NaIO<sub>4</sub> relative to the pendent cis-diol groups also resulted in partial loss of the diithioctoate end-groups. Similarly, a PGEO5MA homopolymer (M<sub>n</sub> = 124.1 kgmol<sup>-1</sup>; D = 4.55) was synthesized via free-radical polymerization in aqueous solution at 70°C for 18 h. Selective oxidation of the cis-diol groups on this homopolymer also had minimal effect on its (broad) molecular weight distribution (Figures S5 and S6).

To investigate the scope of such new water-soluble aldehyde-functional polymers for conjugation with biologically-relevant compounds, PAGEO5MA<sub>n</sub>, homopolymer was reacted in turn with three amino acids (glucose, lysine or cysteine; amino acid/aldehyde molar ratio = 1.0) to form the corresponding Schiff base, followed by in situ reductive amination using excess NaCNBH<sub>3</sub> (Scheme 3). These aqueous reaction mixtures were stirred at 35°C for 48 h, with <sup>1</sup>H NMR spectroscopy studies indicating very high extents of reaction (> 99%) in each case (Figure S7). Aqueous GPC analysis of the resulting water-soluble polymers indicated that molecular weight distributions remained relatively narrow after this two-step one-pot derivatization (Figure S8).

This protocol was then extended to water-soluble diblock copolymers. A series of neutral, zwitterionic, cationic or anionic double-hydrophilic diblock copolymers was prepared in which one of the blocks was PGEO5MA (Scheme 4). For the neutral diblock copolymer, a trithiocarbonate-capped PEO<sub>113</sub> precursor was simply chain-extended via RAFT aqueous solution polymerization of GEO5MA at 50°C. For the synthesis of the ionic diblock copolymers, a PGE05MA<sub>37</sub> precursor was chain-extended via RAFT aqueous solution polymerization of 2-(methacryloyloxy)ethyl phosphoryl...
choline (MPC), [2-(methacryloyloxy)ethyl] trimethylammonium chloride (METAC) or ammonium 2-sulfatoethyl methacrylate (SEM) at 70°C. Each polymerization was allowed to proceed overnight to ensure high monomer conversion (>98% in all cases, as confirmed by 1H NMR spectroscopy; Table 2).

DMF GPC analysis indicated a high blocking efficiency for the RAFT solution polymerization of GEO5MA using the PEG113 macro-CTA and the resulting PEG113-PGEO5MA50 diblock copolymer had a relatively low dispersity (D = 1.20; Table 2; Figure S9a). However, aqueous GPC analysis was required to assess the molecular weight distributions of the ionic diblock copolymers. (Table 2; Figures S9b–d). Oxidation of the pendant cis-diol groups on the PGEO5MA, chains was investigated using a NaIO4/cis-diol molar ratio of unity at a diblock copolymer concentration of 20% w/w. According to 1H NMR analysis, the extent of derivatization was at least 99% in all cases (Table 2). DMF GPC analysis confirmed that periodate oxidation had minimal effect on the molecular weight distribution (D = 1.22; Figure S9a) in the case of the PEG113-PGEO5MA50 diblock copolymer. Similar results were obtained for the zwitterionic, cationic and anionic diblock copolymers when using aqueous GPC (Table 2; Figures S9b–d).

Figure 3. a) Reaction scheme for the (partial) oxidation of a near-monodisperse PGEO5MA37 precursor in aqueous solution using NaIO4 at 22°C. Adjusting the NaIO4/cis-diol molar ratio (x) between 0.1 and 1.0 generates a library of aldehyde-functional water-soluble statistical copolymers.
b) Offset 1H NMR spectra (D2O) recorded for PGEO5MA37, P(GEO5MA37-stat-AGEO5MA37) (where m = 0.11, 0.49 and 0.78), and PAGEO5MA37.

Scheme 3. Schiff base reaction of PAGEO5MA37 with an amino acid (e.g., glycine, lysine, or cysteine) followed by reductive amination using excess aqueous NaCNBH3 at 35°C to afford a series of new zwitterionic homopolymers via a two-step one-pot wholly aqueous protocol.
Conclusion

In summary, we have reported the atom-efficient synthesis of a new \textit{cis}-diol-based methacrylic monomer (GEO5MA) that is readily converted into a hydrophilic aldehyde-functional monomer (AGEO5MA) via selective oxidation using NaIO$_4$ in aqueous solution. Unlike almost all other literature examples of aldehyde-based vinyl monomers, this latter monomer is water-soluble and can be polymerized with good control via RAFT aqueous solution polymerization. Alternatively, homopolymerization of the GEO5MA precursor under similar conditions affords a well-defined water-soluble PGEO5MA precursor that can be converted into PAGEO5MA under mild conditions using a stoichiometric amount of NaIO$_4$ oxidant. On the other hand, using sub-stoichiometric quantities of NaIO$_4$ relative to the pendent \textit{cis}-diol units produces a range of water-soluble aldehyde-functional statistical copolymers. New PAGEO5MA-based double-hydrophilic diblock copolymers can be prepared and model Schiff base reactions have been conducted in aqueous solution under mild conditions using various amino acids to introduce zwitterionic groups. We anticipate that this new hydrophilic aldehydic vinyl monomer and its corresponding copolymers will offer a range of potential applications in the fields of cell biology and biomaterials.

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Conflict of interest

The authors declare no conflict of interest.

Keywords: aldehyde-functional methacrylic monomers · block copolymers · periodate oxidation · RAFT polymerization

Table 2: Summary of monomer conversions, extents of \textit{cis}-diol oxidation and GPC molecular weight data for a series of neutral, zwitterionic, cationic and anionic diblock copolymers (with reference homopolymers included for comparison).

| GPC eluent | Polymer composition | Monomer conversion [%] | Extent of \textit{cis}-diol oxidation [%] | $M_n$ [kg mol$^{-1}$]$^c$ | $D$ |
|------------|---------------------|------------------------|--------------------------------------|-----------------|---|
| DMF        | PEG$_{113}$         | –                      | –                                    | 5.0             | 1.13 |
| DMF        | PEG$_{113}$-PGEO5MA$_{17}$ | > 99                | –                                    | 27.7            | 1.20 |
| DMF        | PEG$_{113}$-PAGEO5MA$_{17}$ | –                 | > 99                                 | 26.2            | 1.22 |
| Aqueous$^a$| PEGO5MA$_{17}$      | –                      | –                                    | 5.8             | 1.29 |
| Aqueous$^a$| PEGO5MA$_{17}$-PMPC$_{30}$ | > 99               | –                                    | 13.1            | 1.34 |
| Aqueous$^a$| PEGO5MA$_{17}$-PMPC$_{30}$ | –                  | 99                                   | 13.4            | 1.38 |
| Aqueous$^a$| PEGO5MA$_{17}$-PMPC$_{30}$ | –                  | –                                    | –               | –   |
| Aqueous$^a$| PEGO5MA$_{17}$-PMPC$_{30}$ | 98                   | –                                    | 24.3            | 1.12 |
| Aqueous$^a$| PEGO5MA$_{17}$-PMPC$_{30}$ | –                   | > 99                                 | 23.3            | 1.11 |
| Aqueous$^a$| PEGO5MA$_{17}$-PMPC$_{30}$ | –                   | –                                    | 5.7             | 1.34 |
| Aqueous$^a$| PEGO5MA$_{17}$-PMPC$_{30}$ | > 99                 | –                                    | 11.0            | 1.30 |
| Aqueous$^a$| PEGO5MA$_{17}$-PSEM$_{30}$ | –                   | > 99                                 | 12.7            | 1.36 |

[a] 0.2 M NaIO$_4$, 0.05 M TRISMA buffer, pH 7. [b] 0.5 M acetic acid, 0.3 M NaH$_2$PO$_4$, pH 2. [c] Relative to PEG/PEO standards.

Scheme 4. a) Reaction scheme for the synthesis of PEG$_{113}$-PGEO5MA$_{17}$ via RAFT aqueous solution polymerization of GEO5MA at 40% w/w solids using a PEG$_{113}$/VA-044 molar ratio of 5.0 at 50°C. b) Reaction scheme for the synthesis of PEGO5MA$_{17}$-PX$_{50}$ diblock copolymers (where X = MPC, METAC or SEM) at 20% w/w solids using a PEGO5MA$_{17}$/ACVA molar ratio of 5.0.
A water-soluble aldehyde monomer (AGEOSMA) was prepared by oxidation of GEOSMA at 22 °C. RAFT aqueous polymerization yielded PAGEOSMA homopolymer, which was oxidized using sub-stoichiometric periodate/cis-diol molar ratios to yield hydrophilic statistical copolymers with cis-diol and aldehyde groups. Three PGEOSMA-based diblock copolymers were converted into aldehyde-functional copolymers, which were derivatized with amino acids.