Deep Eutectic Solvents in The Synthesis of Polymers

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Abstract

Deep Eutectic Solvents (DESs), a subclass of ionic liquids, have emerged as sustainable and green solvents in chemical reactions. In this mini-review article, deep eutectic solvents utilization in the polymer synthesis will be discussed briefly.

Keywords: Deep eutectic solvents; Polymer synthesis; Green chemistry

Abbreviations: DES: Deep Eutectic Solvents; HBA: Hydrogen Bond Acceptor; HBD: Hydrogen Bond Donor; PDCs: Poly Diol-Co-Citrate; PAAc: Poly (Acrylic Acid); PMAAc: Poly Methacrylic Acid; APS: Ammonium Persulfate; EM: Eutectic Mixture; FP: Frontal polymerization

Introduction

Solvent utilization creates the largest volume of auxiliary waste in polymer synthesis. However, increasing awareness of the environmental problems has led to the use of alternative reaction media to reduce or eliminate organic solvent use. Deep eutectic solvents (DESs) are systems formed from a eutectic mixture of Lewis or Bronsted acids and bases which can contain a variety of anionic and/or cationic species [1]. They incorporate a hydrogen bond acceptor (HBA) and a hydrogen bond donor (HBD), which are able to give a eutectic with a melting point much lower than either of the individual components (Figure 1) [2]. One of the most significant deep eutectic phenomenon’s was observed for a mixture of choline chloride and urea in a 1:2 molar ratio. The resulting mixture has a melting point of 12°C, which makes it liquid at room temperature. DESs are obtained by complexion of quaternary ammonium salts with hydrogen bond donors. The charge delocalization occurring through hydrogen bonding between the halide anion and the hydrogen donor moiety is responsible for the decrease in the freezing point of the mixture relative to the melting points of the individual components (Figure 2) [3].

DESs share many characteristics of conventional ILs (e.g. non-volatile, nonreactive with water, and biodegradable) while it offers certain advantages. The preparation of eutectic mixtures in a pure state can be accomplished more easily than that of ILs with no need of post-synthesis purification due to the purity of the resulting DES will simply depend on the purity of its individual components. Moreover, the low cost of those eutectic mixtures based on readily available components (for instance, urea and choline chloride are a well-known example) makes...
Green Polycondensations

The polycondensation of citric acid with 1,8-octanediol for the synthesis of Poly (Diol-Co-Citrates) (PDCs), were first reported by Yang et al. [5,6] in 2004 and, similar to other biocompatible polyesters, have indicated tremendous versatility as polymeric networks for regenerative medicine. The conventional synthetic process first consists of citric acid melting by thermal treatment at 160-165°C for a few minutes (Figure 1). Then, 1,8-octanediol is dissolved in this molten phase and polycondensation starts. The temperature is maintained at 140°C over 60 min, and then reduced to 80°C to let the reaction to proceed over several days until completion. Deep eutectic solvent-assisted syntheses offer an interesting opportunity to fits well with green chemistry. For example, the mixture of 1,8-octanediol and lidocaine, which is a local anesthetic, in stoichiometric molar ratios forms a eutectic with a melting point of about 40°C. Citric acid can easily be dissolved in this eutectic mixture and polycondense with 1,8-octanediol at temperatures far below those described above for regular polycondensations (Figure 2). Lidocaine was finally entrapped with a high loading due to the stoichiometric in which it formed part of the eutectic mixture into the resulting polyesters. Thus, we were able to obtain drug-eluting materials, the controlled release of which into an aqueous medium was simply based on the biodegradable character of the polyester network. It is worth noting that lidocaine would decompose (at least, partially at 155°C) following regular polycondensation [7,8].

Polymer inhibition of bacterial growth against E. coli was measured as the diameter of the zone of inhibition around polymer discs normalized by the grams of polymer. A clear zone of inhibition was observed around the discs, with values varying from 1.76 (POC-C) to 68.90 (POC-M) mm g⁻¹ of polymer as the minimum and maximum inhibition zones, respectively.

DESs have been used in polycondensations, where one of the components was also a reactant, thus called all-in-one systems [10]. Electrochemical polymerization of conductive monomers has also been explored taking advantage of the ionic conductivity of DESs [11-15], including the FeCl³⁻-catalyzed oxidative polymerization of 3-octylthiophene [14]. Recently, DESs were reported to play the role of curing agents in epoxy resins [16], monomer [17,18] and initiator [19] in ring-opening polymerization.
Free-radical polymerization of DESs: deep eutectic monomers

In free-radical polymerization of DESs, monomers able to polymerize undergo free-radical polymerization while taking part in a DES, as hydrogen bond donor or ammonium salt, so called DES monomers [20]. The introduction of DESs in free-radical polymerizations was reported by the synthesis of Poly (Acrylic Acid) (PAAc) and Poly (Methacrylic Acid) (PMAAc) monoliths by frontal polymerization [21]. Frontal polymerization (FP) is normally performed with neat monomers but can be accomplished in solution [22-25]. Bednarz et al. [26], described the free radical copolymerization of itaconic acid-forming DES with ChCl−and AAm by in situ polymerization-crosslinking of the DES (Figure 5). Itaconic acid is a bio-based monomer containing an unsaturated moiety that slowly polymerizes by free radicals; it also has two carboxylic groups available as HBDs [27]. As result of Itaconic Acid (IA) transformation into DES, copolymerization initiated by persulfate proceeded faster and at lower temperature than in water and produced higher crosslinked hydrogels, pointing to a catalytic effect of ChCl. Based on the results of itaconic acid free-radical polymerization initiated by Ammonium Persulfate (APS) in water, it was concluded that DES acted as the solvent of itaconic acid and also as catalyst in free-radical polymerization processes [26]. According to this study, the copolymerization-crosslinking of itaconic acid with N,N'-methylenebisacrylamide in the DES was a useful method for the preparation of hydrogels. Comparative studies indicate that itaconic acid undergoes faster copolymerization in DES than in water, and probably for that reason the obtained product has higher cross-link density [26].

![Figure 5: Preparation of poly (itaconic acid) crosslinked by BAA by: a) polymerization in DES (P-DES) and b) polymerization in water (P-WATER), and tentative structures of resulting composites (C-DES and C-WATER) and hydrogels (H-DES and H-WATER). Reproduced with permission from Reference [26].](image-url)

DES assisted atom transfers radical polymerization

A green solvent system, composed of a eutectic mixture (EM) and ethanol, was used in the supplemental activator and reducing agent atom transfer radical polymerization (SARA ATRP) of methyl acrylate (MA), yielding well-defined polymethyl acrylate (PMA) chains (Figure 6) [27]. In addition, this eutectic-based system allows a straightforward separation of the entire catalytic system, with possibility of reutilization, which is a much advantageous feature for a possible industrial implementation of the SARA ATRP process [27]. In this study, ethanol/reline (choline chloride: urea) mixtures showed to be excellent solvent for the polymerization, providing well-defined PMA (dispersity, D(Mw/Mn) <1.2) as well as affording the recycling of the entire catalytic system Cu(0)/CuBr₂/Me₆TREN, which could be reused at least two times in new SARA ATRP experiments.

Conclusion

Deep eutectic solvents provided a reaction medium, so that polymerizations were ultimately performed in a solventless manner. DESs also provided new precursors that favored more efficient polymerizations by decreasing the energy input required for the reaction to proceed. The use of mild reaction conditions in combination with the compositional versatility of DESs, which allows the selection of low toxic components, is also of interest from the viewpoint of green chemistry due to it opens up the way to the design of eco-friendly synthetic methods. DES-assisted polymer syntheses met more green principles than those of conventional ones, so, polymers obtained from greener processes have more opportunities to be feasible for green applications than polymers obtained from less green ones. These encouraging results suggest that DES-assisted polymerizations can help to provide synthetic designs and products with a greener character.
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References
