ABSTRACT: Producing polymers from renewable resources via more sustainable approaches has become increasingly important. Herein we present the polymerization of monomers obtained from biobased renewable resources, employing an environmentally friendly photoinduced iron-catalyzed atom transfer radical polymerization (ATRP) in low-toxicity solvents. We demonstrate that renewable monomers can be successfully polymerized into sustainable polymers with controlled molecular weights and narrow molar mass distributions ($D$ as low as 1.17). This is in contrast to reversible addition−fragmentation chain-transfer (RAFT) polymerization, arguably the most commonly employed method to polymerize biobased monomers, which led to poorer molecular weight control and higher dispersities for these specific monomers ($D_s$ ∼ 1.4). The versatility of our approach was further highlighted by the temporal control demonstrated through intermittent “on/off” cycles, controlled polymerizations of a variety of monomers and chain lengths, oxygen-tolerance, and high end-group fidelity exemplified by the synthesis of block copolymers. This work highlights photoinduced iron-catalyzed ATRP as a powerful tool for the synthesis of renewable polymers.

Directed polymerization methodologies toward more sustainable pathways is of paramount and ever-increasing importance. However, a plethora of factors should be considered to improve the feasibility of a sustainable polymerization methodology. To begin with, the most important factor is the origin of monomers to be polymerized. Currently, the vast majority of monomers used for the synthesis of polymers are based on fossil fuel feedstock. Recently, in search for more sustainable alternatives, the synthesis of polymers from renewable resources has attracted significant attention, showing great promise in counter-balancing the use of fossil fuel feedstock. Indeed, biomass-derived materials have been employed as an alternative and renewable resource for the synthesis of monomers. For example, lignocellulose has gained great popularity as an inexpensive renewable waste product which can be produced in high abundance. In particular, lignin can provide multiple phenol derivative building blocks, the secondary alcohol of which can be easily modified under mild conditions to offer polymerizable building blocks. Another promising family of renewable resources that can be used for monomer production are terpenes, which can be extracted from plants, providing interesting biological properties.

The synthesis of polymers from renewable resources is not only from a sustainability viewpoint, but also because it leads to the production of novel polymeric materials with unique properties. To maximize access over the range of polymeric materials that can be attained, reversible deactivation radical polymerization (RDRP) has recently been employed to polymerize a range of renewable monomers. For the majority of cases, reversible addition−fragmentation chain-transfer (RAFT) polymerization has been employed for the polymerization of biomass-based monomers, as it is one of the most versatile RDRP techniques. However, RAFT polymerization of some renewable methacrylate monomers often leads to relatively broad molar mass distributions ($D_s$ ∼ 1.3−1.7). In parallel, copper-mediated atom transfer radical polymerization (ATRP) has also been employed in polymerization of renewable monomers, but to a relatively lesser extent. It is noted that both polymerization methods typically employ toxic components/solvents that prevent the development of a more sustainable polymerization procedure. Arguably, one of the most environmentally friendly RDRP methodologies is iron (Fe) ATRP. Fe is one of the most abundant metals on earth and is inexpensive, nontoxic, and biocompatible. The possibility to utilize light (rather than heat) as an external stimulus to mediate Fe ATRP is also advantageous from a sustainability point of view. In a similar fashion to conventional RAFT and copper ATRP, highly toxic solvents such as acetonitrile, anisole, or trifluoroethanol are...
often required to conduct a successful Fe ATRP.\textsuperscript{38,39} In this work, we aim to develop a greener and efficient approach, which will satisfy as many of the 12 principles of green chemistry as possible,\textsuperscript{3} to polymerize renewable monomers by employing the environmentally friendly photoinduced iron-catalyzed ATRP in low-toxicity solvents while maintaining narrow molar mass distributions for all the synthesized sustainable polymers.

We first synthesized six different methacrylic monomers through the esterification of lignin derivatives (phenol, p-cresol, guaiacol, vanillin, syringol) and a thyme-derived terpene compound (thymol) using methacrylic anhydride, which is less toxic than methacryloyl chloride, the more frequently employed compound in this type of reaction (Scheme 1, S1, Spectra S1–S6)). 2-Methyltetrahydrofuran was used as the solvent for the methacrylation, a “greener” alternative when compared with more toxic and commonly employed organic solvents, such as dichloromethane.\textsuperscript{40} The monomers were purified thoroughly in order to avoid any contamination with the initial alcohols, which in some cases may be hazardous. After the successful synthesis of the renewable monomers, we sought to perform polymerizations via photoinduced iron-catalyzed ATRP. Phenyl methacrylate (PheMA) was used as the model monomer, tetraethylene glycol dimethyl ether (TEGDME) as the model low-toxicity solvent, FeBr\textsubscript{3} as the metal source, tetrabutylammonium bromide (TBABr) as the ligand, and methyl \( \alpha \)-bromophenylacetate (MBPA) as the initiator (Scheme S2). Initial experiments were conducted with a ratio of [MBPA]/[FeBr\textsubscript{3}]/[TBABr] = 1:0.1:0.1 and a targeted degree of polymerization (DP) of 50. All experiments were performed under blue light LED irradiation (48 W, \( \lambda \text{ = } 465 \text{ nm (±5 nm)} \)) in a homemade box (Figure S1). Under the aforementioned conditions, well-defined PPheMA could be obtained within 90 min with good control over the molecular weight and low dispersity as determined by size-exclusion chromatography (SEC; \( M_n = 5100 \), \( D = 1.17 \), Figure 1a, Table S1, entry 1). It can therefore be concluded that TEGDME does not decrease the catalyst’s activity (by ligation to Fe)\textsuperscript{41,42} and as such is an excellent solvent choice for the photoinduced iron-catalyzed ATRP of renewable monomers. The possibility to utilize alternative low-toxicity and green solvents was also investigated using 2-methyltetrahydrofuran and cyclopentyl methyl ether.\textsuperscript{40} Under otherwise identical conditions, both solvents fully solubilized the catalyst, resulting in PPheMA with very similar control over the molecular weight and dispersity as in the case of TEGDME (Figure 1a, Table S1, entries 2 and 3). Although the remaining experiments were conducted in TEGDME, the other two solvents were proven equally efficient to mediate a successful photoinduced iron-catalyzed ATRP, thus, suggesting no competing solvent—catalyst complexation.

Next, we were interested in whether an “on/off” temporal control is possible during the polymerization of these renewable monomers. To assess this possibility, we monitored the growth of PPheMA chains during alternating periods of light and dark, using \(^1\)H NMR to calculate the monomer conversion. Under the previously established conditions ([PheMA]/[MBPA]/[FeBr\textsubscript{3}]/[TBABr] = 50:1:0.1:0.1), negligible polymerization was observed during the dark periods (<3\%, Table S2), whereas a clear increase in monomer conversion was observed when the reaction was exposed to visible light irradiation (Figure 1b). The small percentage of polymerization noticeable during the dark periods was attributed to the relatively high catalyst loading and is in agreement with previous reports.\textsuperscript{38,43} To fully eliminate the conversion during the “off” cycles, a further decrease of the catalyst concentration is recommended, albeit at the expense of higher dispersity polymers.\textsuperscript{44}

To explore the potential of our technique to control the polymerization of higher molecular weight polymers, a range of DPs were targeted. Good control and low dispersities were observed regardless of the initial chain length targeted (Figure 1c, Table S3). Considering the recent interest in oxygen-tolerant polymerizations,\textsuperscript{35–37} we also attempted our polymerizations in the absence of any external deoxygenation by simply minimizing the reaction vessel’s headspace to reduce the amount of the initially present oxygen. Pleasingly, a well-controlled polymerization took place, thus further simplifying the polymerization procedure (Table S4). Next, we wanted to investigate whether our method allows the low-volume synthesis of polymers as this may be of high interest to applications that require the use of low reaction scales (i.e., biological studies, bioconjugations, etc.). For this purpose, we conducted the polymerization of PheMA in reaction volumes of 50 and 100 \( \mu \)L (25 and 50 mg of monomer, respectively) and observed good control over the polymerization (Figure S2), thus highlighting the versatility of our method. Finally, we wanted to investigate the end-group fidelity achieved through our methodology. To assess this, a PPheMA macroinitiator (\( M_n = 6500 \), \( D = 1.17 \)) was chain-extended, furnishing higher molecular weight polymers while maintaining low dispersity (\( M_n = 12300 \), \( D = 1.24 \), Figure 1d, Table S5).

Upon establishing optimized conditions, we then successfully polymerized a range of renewable monomers, as indicated in Scheme 1. It is noted that for the liquid monomers, including cresol methacrylate (CreMA), guaiacol methacrylate (GuMA), and thymol methacrylate (ThyMA), a ratio of monomer to solvent 1:1 was employed. Instead, the polymerization of solid monomers (i.e., vanillin methacrylate (VaMA) and syringol methacrylate (SyrMA)) required an increased solvent loading to fully dissolve the initial monomer (1.5 equiv with respect to monomer). With these modifications, all monomers were efficiently polymerized, yielding controlled molecular weights and low dispersities (Figure 2 and Table S6). An additional block copolymer was also targeted, consisting of PPGuMA (\( M_n = 6500 \), \( D = 1.17 \), Figure 2c, and Table S7, entry 1) as the first block. In the presence of 10\% of catalyst, PPGuMA was chain-extended with PheMA resulting in a diblock with a higher dispersity (\( D = 1.35 \)) (Figure S3a and Table S7, entry 2). However, further lowering

Scheme 1. Synthesis of Sustainable Biomass-Derived Monomers and Their Corresponding Polymers

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of the diblock’s dispersity was possible by doubling the catalyst concentration (\(D = 1.24\); Figure S3b and Table S7, entry 3).

Intrigued by the excellent control over the dispersities attained through our photoinduced iron-catalyzed ATRP, we were interested in a direct comparison with conventional thermal RAFT polymerization, which is the most commonly employed method to polymerize such renewable monomers.

By replicating the experiments in the presence of 2-cyano-2-propyl benzodithioate and 10\% of AIBN, PheMA was effectively polymerized by RAFT polymerization albeit the final dispersity was as high as 1.43 (Table S8, entry 4). Similar results were also obtained for the polymerization of other monomers such as SyrMA, ThyMA, and VaMA (Figure S4). Detailed kinetic analysis was conducted to compare photoinduced iron-catalyzed ATRP with thermal RAFT polymerization under otherwise identical conditions (same monomer/targeted DP/solvent). Although both systems displayed features of a controlled polymerization such as a linear increase of \(\ln[M_0]/[M]\) over time (Figure 3b,e) and comparable reaction rates, some important differences were also observed. The first observation was a clear discrepancy between theoretical and experimental molecular weights. For instance, in the case of photoinduced iron-catalyzed ATRP an \(M_n\) of 5200 was obtained by SEC at 62\% of conversion which is in close agreement with the theoretical \(M_n\) (5200). Instead, when RAFT polymerization was employed, a higher experimental \(M_n\) (\(M_{SEC} = 6500\)) was observed at a similar conversion (i.e., 60\%). We hypothesized that this discrepancy could be explained by the incomplete consumption of the RAFT agent. Indeed, the UV-SEC detector confirmed that the RAFT agent was not fully consumed, even at higher monomer conversions (>90\%, Figure S5), which verified our original hypothesis. This discovery may also be associated with our second observation in that polymers synthesized by RAFT polymerization showed significantly higher dispersity values (1.37 and 1.43 at \(\sim 60\) and 90\% conversion, respectively, as opposed to 1.18 and 1.20 for Fe-ATRP; Figure 3a,c,e,f and Tables S8 and S9). The higher dispersities observed by RAFT polymerization are attributed to the slow consumption of the RAFT agent as a result of less-efficient fragmentation and the
Instead, the lower $D$s obtained via photoinduced iron-catalyzed ATRP indicate a faster and more complete initiator consumption. To further understand our data, we conducted two additional control experiments. First, we polymerized PheMA using photoinduced electron-energy transfer (PET) RAFT instead of thermal RAFT. Under otherwise identical conditions, PET RAFT gave rise to a similarly high dispersity ($D = 1.34$, Figure S6), thus further supporting insufficient fragmentation with the selected RAFT agent. Second, to examine whether the polymerization temperature can affect the polymerization control, we conducted in parallel a polymerization of PheMA with thermal RAFT polymerization at 70 °C, and also iron-catalyzed photoinduced ATRP at the same temperature. The results show the superiority of iron-catalyzed photoinduced ATRP over RAFT in the polymerization of PheMA ($D = 1.2$ vs $D = 1.34$, Figure S6).

Figure 2. SEC traces of renewable polymers synthesized via photoinduced iron-catalyzed ATRP: (a) PPheMA, (b) PCreMA, (c) PGuMA, (d) PThyMA, (e) PVaMA, and (f) PSyrMA.

Figure 3. Polymerization kinetics of PheMA utilizing (a−c) photoinduced iron-catalyzed ATRP and (d−f) conventional thermal RAFT polymerization.

potential hybrid behavior RAFT may have. Instead, the lower $D$s obtained via photoinduced iron-catalyzed ATRP indicate a faster and more complete initiator consumption. To further understand our data, we conducted two additional control experiments. First, we polymerized PheMA using photoinduced electron-energy transfer (PET) RAFT instead of thermal RAFT. Under otherwise identical conditions, PET RAFT gave rise to a similarly high dispersity ($D = 1.34$, Figure S6), thus further supporting insufficient fragmentation with the selected RAFT agent. Second, to examine whether the polymerization temperature can affect the polymerization control, we conducted in parallel a polymerization of PheMA with thermal RAFT polymerization at 70 °C, and also iron-catalyzed photoinduced ATRP at the same temperature. The results show the superiority of iron-catalyzed photoinduced ATRP over RAFT in the polymerization of PheMA ($D = 1.2$ vs $D = 1.34$, Figure S6).
renewable methacrylate monomers than conventional RAFT reactions can be performed without any deoxygenation, through light/dark iterations. Importantly, the polymerization temporal control over the polymerization could be achieved high end-group fidelity of the polymers was demonstrated via chain extensions and block copolymers. Additionally, good ATRP to polymerize a variety of renewable monomers in low-ology, whereby we leveraged photoinduced iron-catalyzed European Research Council (ERC) under the European 2020.0324). This project has received funding from the Swiss Government Excellence Scholarship (ESKAS No. FZQ051-1/2020-2021. H.S.W. acknowledges the award of K.P. thanks the Onassis Foundation, as this scientific paper was A.A. gratefully acknowledges ETH Zurich for financial support. K.P. thanks the Onassis Foundation, as this scientific paper was supported by the Onassis Foundation-Scholarship ID: FZQ051-1/2020-2021. H.S.W. acknowledges the award of the Swiss Government Excellence Scholarship (ESKAS No. 2020.0324). This project has received funding from the European Research Council (ERC) under the European Union’s Horizon 2020 research and innovation program (DEPO: Grant No. 949219).

Notes
The authors declare no competing financial interest.

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