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Biobased, thermoreversibly crosslinked polyesters

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Chapter 2

Bio-based aromatic polyesters reversibly crosslinked via Diels- Alder reaction

2.1 Abstract

Diphenolic acid is functionalized with furfuryl amine and subsequently incorporated in a (partly) bio-based polyester through interfacial polycondensation with terephthalic chloride. The furan groups present in the resulting polyester are able to form a thermoreversible covalent network with different bismaleimide moieties via the Diels-Alder (DA) reaction. Analysis of the polymer network by $^1\text{H-NMR}$ clearly shows formation of both stereoisomers (endo and exo) possible from Diels-Alder coordination of furan and maleimide. Furthermore, it was found that these isomers can be reversibly interchanged at temperatures below the reported retro Diels-Alder reaction temperature, a phenomenon often claimed, yet up to date never directly observed, for thermally reversible polymeric systems. Finally, a proof-of-principle for reversibility and recyclability is shown.

Keywords: bio-based, short-chain, polyester, furan, bismaleimide, thermoreversibility, Diels-Alder, endo-exo regeoisomers, recyclability.

2.2 Introduction

The uncertain prices of crude oil, the increasing awareness of carbon emissions as well as the resulting desire to become gradually independent of fossil fuels constitute strong driving forces behind the search for sustainable alternatives to fossil-fuel based chemicals¹. An ever-increasing number of processes are “going green”. The first, and most obvious approach towards this end is the direct substitution of petrol-based chemicals by their bio-based counterparts. There are various examples of commercially available chemicals, obtained from renewable resources, like polyethylene² and succinic acid³. These *drop-in* products require no special adaptation before using them as they are essentially the same chemical, merely isolated from a different feedstock. The transition towards the use of green chemicals is not always smooth since obtaining these chemicals is hindered by a few complicating factors. Most predominantly, it is often cumbersome to obtain them in high purities as bio based chemicals are often obtained from a mixture of usually very similar and hard to separate chemicals⁴. This problem is particularly pressing for green monomers as these require very high purities in order to be able to undergo successful polymerization reactions⁵. Fortunately,

there are cases where a mixture of products has successfully been used as green component. Indeed, up to 75% of phenol used in wood adhesives can be replaced by a mixture of various phenolic compounds obtained from lignin⁶ and polyols from vegetable oils have successfully been employed as crosslinker in polyurethanes⁷. However, these are rare exceptions rather than common practice. Nevertheless, despite these complications, several successful implementations of green chemicals as monomers have been reported⁸⁻¹⁰, reducing both the carbon footprint of the associated materials and their fossil-fuel dependency. Unfortunately, despite these positive aspects, simple substitution of oil-based chemicals with biomass-derived ones does not provide an exhaustive answer to the sustainability issue as waste-streams (i.e. the waste produced at the end of the product life) remain virtually unaffected. Green thermoplastic materials might be recyclable in a “cradle-to-cradle” fashion as this is generally true for polymeric materials with physical (e.g. van der Waals) interactions between the chains¹¹. However, thermoset materials are notoriously and factually impossible to recycle according to a “cradle-to-cradle” approach¹². This is obviously true independently of their origin, i.e. whether oil- or bio-based. In the past decade many efforts have been made in order to increase the recyclability of these materials, and a promising recent advancement is the use of thermoreversible crosslinking by means of the Diels-alder reaction⁸⁻¹⁰. The reversible nature of the Diels Alder reaction has been successfully applied in self-healing polymers¹³⁻²¹. Most popular is the reaction between a furan and maleimide^{12,22} mainly due to fast kinetics and wide availability of the reacting groups¹², although systems based on cyanofumarate and vulvene²³ or anthracene and maleimide²⁴ have also been reported. The application of the Diels-Alder reaction in crosslinking has many advantages: as the formed bonds are covalent, the superior properties attributed to thermosets will be retained (e.g. due to their crosslinked structure, these usually possess superior barrier and mechanical properties as well as higher chemical resistance than most thermoplastic ones). Furthermore as the reaction is an equilibrium one and its extent can be easily influenced by changing temperature, it constitutes an ideal candidate for use as crosslinking reaction. Finally, the temperatures for bond formation, and the reverse reaction, which yields the de-crosslinked product, are sufficiently far apart (namely 50-80°C and 110-170°C respectively for furan and maleimide) to ensure that the obtained materials have a large application window.

The conceptual combination of thermal reversibility with the “green” character of available monomers constitutes a possible solution to the problems outlined above, thus providing in principle a lower carbon footprint as well as recyclability (*cradle-to-cradle*) of the end product. In this context, the presence of functional groups (e.g. –OH ones), often considered a major drawback of green chemicals, when compared to fossil-fuel based ones, might be conveniently exploited to provide the chemicals with an added functionality. A paradigmatic example is constituted by diphenolic acid (DPA), a structural analog to bisphenol A²⁵ obtained via a condensation reaction of levulinic acid²⁶ with phenol. The only difference with bisphenol A is the presence of an extra carboxylic acid. Bisphenol A is widely used as monomer in polycarbonate and epoxy resin synthesis, but also employed as rigidifier in polyester resins. Apart from the fact that it is currently obtained from non-renewable resources, it is also a toxic chemical with shown oestrogenic properties. Polycarbonates and polyesters have been successfully synthesized using unmodified DPA^{27,28}, or after protection of the acid group²⁹. However, the acidic group might also be employed to provide this monomer with the desired (*vide supra*) Diels-Alder active moiety.

Aim of this work is to ideally combine the two strategies towards sustainability: i.e. the bio-based character and the possibility for thermally reversible crosslinking and thus possibility for recycling. To this end, DPA was modified with furfuryl amine to obtain a furan-functionalised diol. The incorporation of this monomer into a fully aromatic novel polyester is described based on known polymerization concepts. Finally the obtained polymer was crosslinked by employing the Diels-Alder reaction with a bismaleimide in order to provide a preliminary proof of principle for the possibility to recycle the end product.

2.3 Experimental section

2.3.1 Chemicals:

4,4'-bis(hydroxyphenyl)valeric acid (Diphenolic acid, DPA) 95%, 2-methyltetrahydrofuran (MeTHF) anhydrous, N,N'-Carbonyldiimidazole (CDI) 97%, Terephthaloylchloride 99+%, 1,1(methylenedi-4,1-phenylene)bismaleimide 95% were purchased from Sigma-Aldrich and used as received. Furfurylamine 99+%

was purchased from Sigma-Aldrich and distilled prior to use. Dodecyl bismaleimide was synthesized as described in literature³⁰.

2.3.2 Equipment:

¹H-NMR spectra were recorded on a Varian Mercury Plus 400 MHz using DMSO-*d*₆ as a solvent. ¹H-NMR spectra of samples at elevated temperature were recorded on a Varian Mercury Plus 500 MHz using DMSO-*d*₆ as a solvent. PPM values are given relative to tetramethylsilane (TMS). Thermogravimetric analysis was performed on a Mettler Toledo TGA. The samples were weighed (ca. 10 mg) and placed in the analyser. Subsequently the temperature was raised from 25 °C to 900 °C at a heating rate of 10 °C per minute under N₂ atmosphere. Differential scanning calorimetry (DSC) was performed on a Perkin Elmer differential scanning calorimeter Pyris 1 under N₂ atmosphere. Before DSC's the sample was weighed (ca. 12 mg) and subsequently heated from 25 °C to 180 °C. Multiple cycles were performed at a heating rate of 10 °C/min throughout the measurements. DMTA measurements were performed using a Rheometrics scientific solid analyzer (RSA II) under air using dual cantilever mode at an oscillation frequency of 1 Hz and a heating rate of 5 °C/min. GPC measurements were performed on a HP1100 equipped with three 300x7.5 mm PLgel 3 μm MIXED-E columns in series using a GBC LC 1240 RI detector. Average molecular weight calculations were performed with the PSS WinGPC Unity software from Polymer Standards Service. The following conditions were used: THF as eluent at a flow rate of 1 ml min⁻¹; 140 bar, a column temperature of 42 °C, 20 μl injection volume and a 10 mg ml⁻¹ sample concentration. Toluene was used as a flow marker and polystyrene samples with different molecular weights were used as calibration standard. Pressing of samples was performed in a Taunus-Ton press type VS up 150A.

2.3.3 GPC

GPC measurements were performed to determine the average chain length and while absolute values cannot be given due to the absence of a suitable reference calibration, the spectra obtained across various samples confirm a consistent distribution of chain lengths between batches.

2.3.4 Synthesis of DPA-fur

Diphenolic acid (5 g, 17.5 mmol), 2-methylhydrofuran (30 mL) and *N,N'*-carbonyldiimidazole (1.2 equivalent, 3,398 g, 21 mmol) were charged to a dry three-neck round bottomed flask connected to a cooler under N₂. The reaction was stirred under reflux conditions (90°C) for two h. Furfurylamine (1.2 equivalent, 1.85 mL, 21 mmol) was added and the reaction was stirred at 90°C for an additional 5 hours. The reaction mixture was then allowed to cool to room temperature, diluted with additional 2-methylhydrofuran (45 ml). The solution was washed with 1M HCl (2 x 37 mL), 0.1 M NaOH (2 x 47 mL) and brine (1 x 15 mL) successively. Evaporation of the organic layer yields a yellow oil. Final purification by recrystallization from ethanol gave 3.26 g (51,1%) of white solid.

¹H NMR (300 MHz, DMSO-d₆) δ 9.16 (s, 2H, OH) 8.19 (t, 1H, NH) 7.53 (s, 1H, fur-*p*) 6.93 (d, 4H, benzene-*o*) 6.62 (d, 4H, benzene-*m*) 6.35 (t, 1H, fur-*m*) 6.18 (d, 1H, fur-*o*) 4.19 (d, 2H, N-CH₂) 2.17 (t, 2H, -CH₂-) 1.86 (t, 2H, -CH₂-) 1.45 (s, 3H, -CH₃).

2.3.5 Polymerization (DPA-fur/tereph)

DPA-fur (1.755 g, 4.8 mmol), tert-butyl ammonium bromide (0.152 g, 8.7 wt.% on DPA-fur), sodium hydroxide (0.384 g, 9.6 mmol) and water (45 mL) were charged to a one-neck round bottomed flask (250mL). The mixture was stirred vigorously. Terephthaloyl chloride (0.975 g, 4.8 mmol) dissolved in chloroform (45 mL) was added to the reaction mixture. The reaction was stirred vigorously for 2 h. Precipitation in methanol (1800 mL) yielded 2.20 g (95.6%) of white solid.

¹H NMR (300 MHz, DMSO-d₆) δ 8.42-7.95 (m, 4H, aromatic not next to ester) 7.53 (s, 1H, fur-*p*) 7.42-7.12 (m, 8H, aromatic next to ester) 6.34 (s, 1H, fur-*m*) 6.19(s, 1H, fur-*o*) 4.21 (d, 2H, N-CH₂) 2,38(broad s, 2H, -CH₂-) 1.95 (broad s, 2H, -CH₂-) 1.63 (s, 3H, -CH₃).

2.3.6 DMTA sample preparation via compression molding

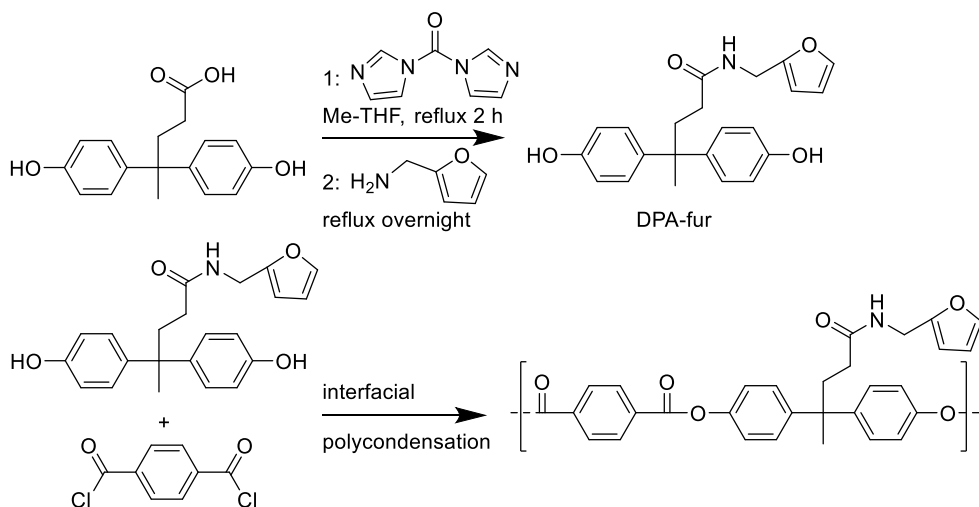
First, a mixture of polymer and bismaleimide was prepared by dissolving the desired amount of bismaleimide in a minimal amount of chloroform (e.g. 0.25 g polymer, 0.09 g of bismaleimide in 1 ml of CHCl₃). Mixing and successive solvent removal by rotary evaporation yielded a cross-linked film of polymer. This film was dried overnight in an oven at 60°C then frozen in liquid nitrogen and ground to particles using an IKA industrial hand grinder. This powder was pressed into

bars (6 mm wide, 54 mm long and 1 mm thick). Pressing was performed at 4 MPa, at a temperature of 150°C for 15 minutes. The material was expected to and did behave as a flowing thermoplastic polymer as this temperature is above the RDA decoupling temperature as well as above the T_g of the un-cross-linked polymer. The samples are cooled slowly inside the press (~30 min) to allow formation of a rigid network through DA coupling. Afterwards the samples were placed in an oven at 50 °C for 24 hours to ensure complete crosslink formation.

2.4 Results and discussion

The overall strategy for the synthesis of the desired polymers relies on the possibility of modifying DPA and functionalizing it (Scheme 1, top) with a Diels Alder reactive group (in this case furan). The employed reaction conditions, based on similar ones for amidation reactions³¹, entail the use of an intermediate (the imidazole peptide), subsequently reacted with fufurylamine to yield the desired product. This synthetic strategy allows for very controlled reaction conditions and high purity and acceptable yield of the end product. Indeed, after crystallization the DPA-fur is obtained in 51.1% yield as an off-white powder characterized by a ¹H NMR spectrum with no relevant traces of impurities (Figure 1). DPA-fur is then subsequently polymerized with terephthaloyl chloride in a two-phase system (Scheme 1, bottom) as described in the literature³².

Scheme 1: top: modification of DPA to DPA-fur, bottom: polymerization of DPA-fur with adipoyl chloride



Clean polymer formation is evident from the complete disappearance of the signals at δ 9.4 ppm in ^1H NMR (corresponding to the OH group on the DPA-fur monomer). Furthermore, aromatic protons from both terephthaloyl chloride and the DPA-fur shift significantly. Two distinct groups of peaks can be identified, one at δ 7.42-7.12 ppm belonging to the aromatic protons next to an ester bond containing 8 protons (4 *o*-from the DPA-fur, and 4 from the terephthaloyl monomer), the other group at δ 8.42-7.95 ppm contains 4 protons (the *o*-protons from DPA-fur away from the ester bond).

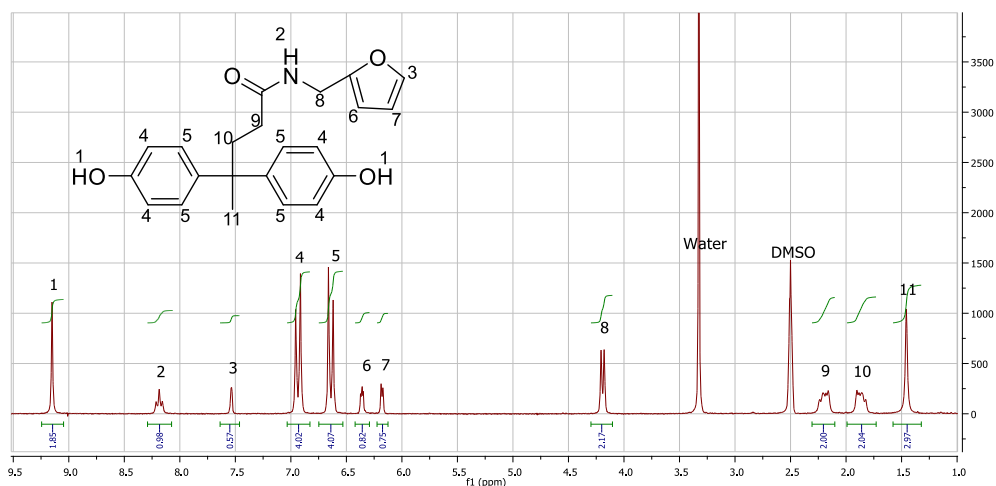
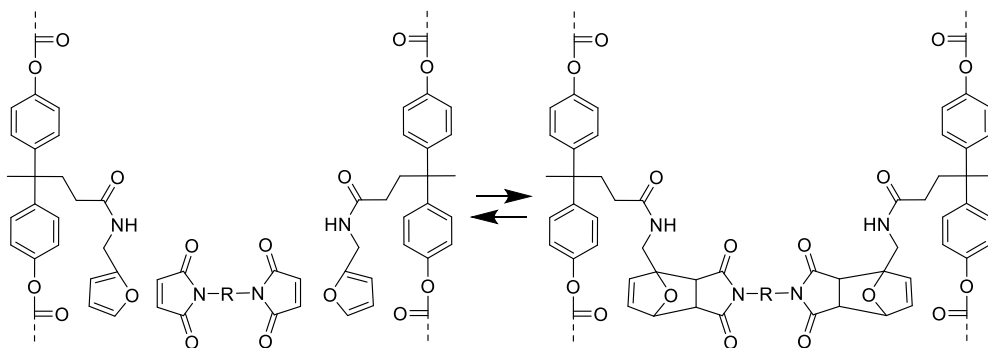


Figure 1: NMR spectrum of DPA-fur

Crosslinking of the prepared polymer via Diels Alder has been achieved by mixing it with an equimolar amount of bismaleimide in DMSO. The system is expected to crosslink by reaction of the pendant furan moieties with the maleimide groups (Scheme 2).

Scheme 2: Diels-alder crosslinking of polymer chains. $R=(CH_2)_{12}, (C_6H_4)-CH_2-(C_6H_4)$ (see Table 1)



The reaction was initially carried out in DMSO in order to visually observe gel formation (indicative of the formation of a network structure) as well as preliminarily investigate its reversibility as a function of temperature. Two types of bismaleimide were added in various ratios and the gelation time (defined as the time it takes the stirrer to stop spinning), a rough indication for the gel formation kinetics, was determined (Table 1).

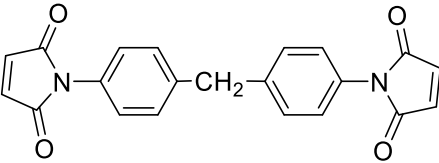
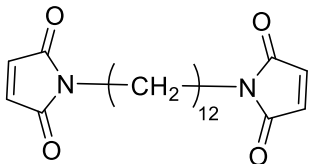
Ratio maleimides to furanes (mol:mol)	1:1	2:1	3:1	4:1
	Gelation time (min)			
 1,1'-(methylenedi-1,4-phenylene)bismaleimide	45	32	25	15
 Dodecyl bismaleimide	368	226	190	143

Table 1: Gelation times for various ratios of two different bismaleimides

In first instance it is worth noticing how both bismaleimides are able to induce gel formation, thus indicating network formation. It is also evident that the aromatic bismaleimide displays faster kinetics than the aliphatic one. The significant difference in gelation times might be related to different reaction kinetics. On the other hand this might also be due to the possibility for the aliphatic crosslinker (because of its flexible nature) of backbiting coordination where both maleimide groups attach to the same polymer chain. The aromatic bismaleimide is expected to be far too rigid to be able to coordinate in this fashion³³. Furthermore increasing the maleimide to furan ratio leads to a decrease in gelation time. This has been observed for other polymeric systems crosslinked in the same way¹² and is most probably related to the second-order kinetics (first order in the bismaleimide) of the crosslinking reaction.

In an attempt to elucidate the mechanism of crosslinking at molecular level, various NMR experiments were performed. First a sample containing uncrosslinked polymer and bismaleimide in a 1:1 ratio of maleimide to furan groups in DMSO-d₆ was prepared. Immediately after mixing an NMR spectrum was recorded (spectrum I in Figure 2). Next the sample was heated to 50°C for 3 hours; this should ensure complete formation of the crosslinks. Afterwards another NMR spectrum was recorded and the signals attributed to the DA-adduct

of furan and maleimide could clearly be detected (spectrum II in Figure 2). There have been various reports describing^{12,34} the assignments of these signals based on model compounds. Interestingly there are two clearly distinct peaks for the *endo* and *exo* adducts visible. To the best of our knowledge this is the first direct confirmation for the formation of both species since similar works on different systems describe an NMR spectrum of a crosslinked polymer network³⁴, but only see a single peak which is attributed to both isomers. When the sample is subsequently heated to 150°C for 5 minutes and a new spectrum is recorded the adduct signals disappear indicating the RDA reaction has completely occurred (spectrum III in Figure 2).

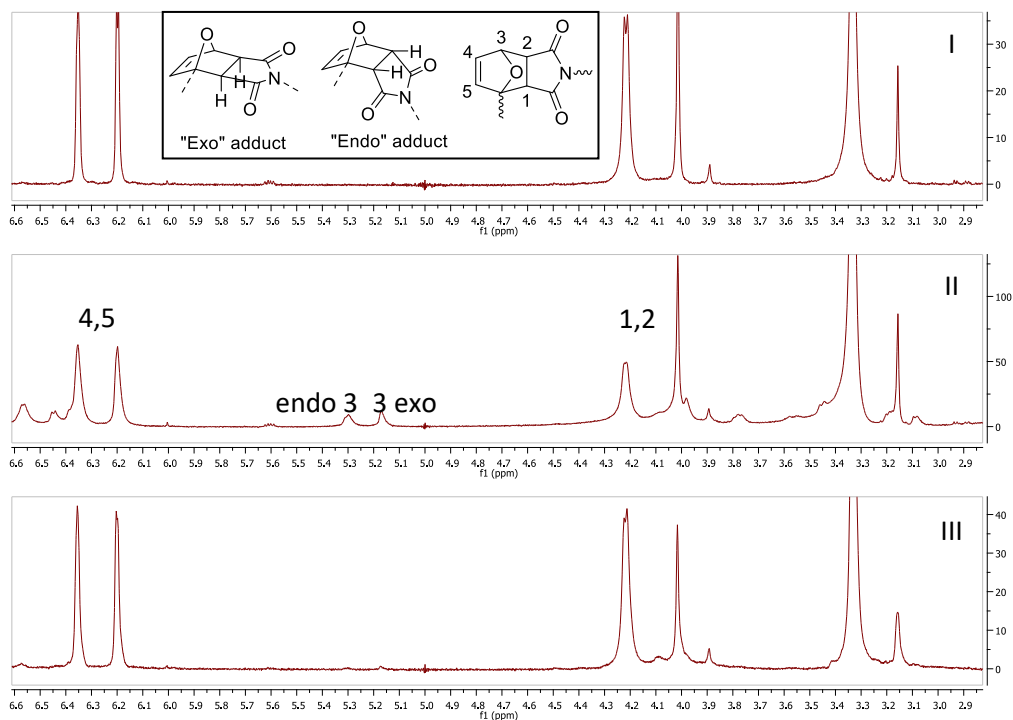


Figure 2: NMR spectra of uncrosslinked polymer and bismaleimide; I: no reaction, II: 3 h at 50°C, III: 5 min at 150°C. Inset: both regioisomers and assignments.

This is further proof, besides the gelation process and its reverse (*vide supra*), of the reversibility of this crosslinking system in solution.

Being able to observe the two DA adducts in the NMR, an attempt was made to quantify their relative ratio as function of temperature. This is quite interesting as such a shift in the ratio of these two adducts has been frequently invoked in literature in order to clarify the thermal behavior of thermal reversible networks^{12,35} and yet no clear proof has ever been reported for a crosslinked polymeric system. For the present system, in another experiment, the same solution containing uncrosslinked polymer and bismaleimide in a 1:1 maleimide:furan group ratio was created. A spectrum immediately after mixing was recorded (spectrum I in Figure 3). Next the sample was heated to 65°C for 2 h, and a new spectrum recorded (spectrum II in Figure 3). Once again the signals belonging to the DA adduct are found and the ratio of *endo:exo* adducts is 58:42. It is well known that the DA reaction is an equilibrium reaction, that the *exo* isomer is thermodynamically favored and the *endo* adduct is kinetically favored. When the sample is kept as 65°C for 48 h and another spectrum is recorded (spectrum III in Figure 3), the ratio of *endo:exo* isomers has shifted significantly to 11:88. This constitutes, to the best of our knowledge, the first direct confirmation of the two adducts formation in a as well as of their nature (thermodynamically and kinetically favored) for furan maleimide crosslinking reactions on a polymeric system. Indeed, other studies until now showed a similar behavior, but only model compounds or singular adducts have been used³⁶.

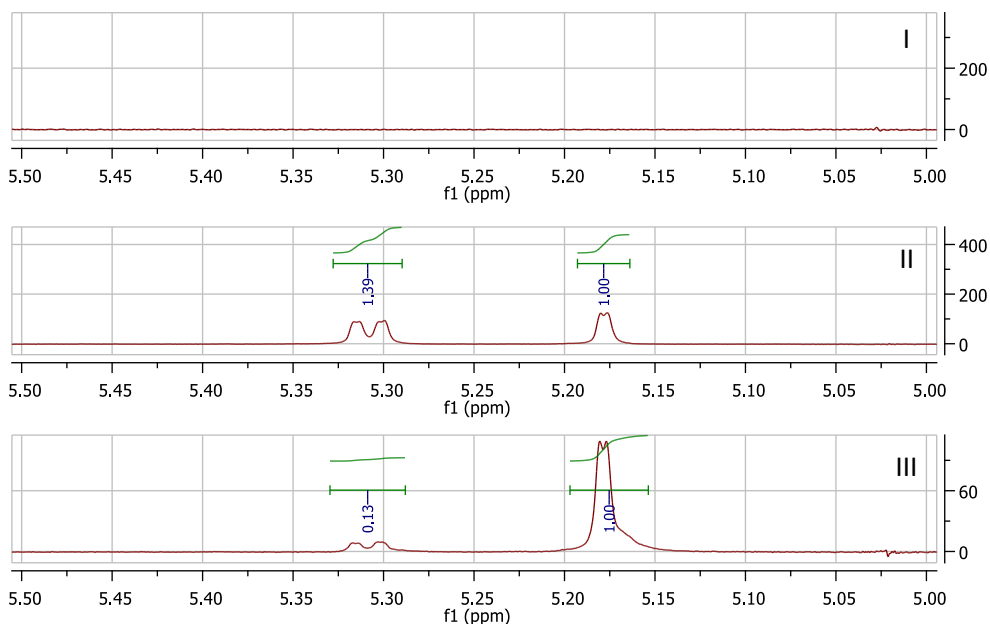


Figure 3: ^1H -NMR spectra of unreacted polymer and crosslinker (top) after 2 hours (middle) and 48 hours (bottom) at 65°C (only the endo/exo protons are shown)

The formation of the two DA adducts and their relative ratio might be of pivotal importance for thermal reversibility of the system, as suggested for related ones³⁷.

Thermal reversibility was tested by cyclical DSC measurements, according to which the material is heated to 180°C and subsequently cooled to room temperature for 8 consecutive times (Figure 4 for DPA-fur/tereph). The DA coupling is exothermic, while the RDA decoupling is an endothermic process. The integral of the corresponding peaks is an indication of the extent of the reaction.

The results suggest that the system shows thermoreversible coupling and decoupling: the heat capacity of the material does not change significantly over the course of all cycles, which is an indication of full reversibility^{12,37}. The peak at 150°C is related to the RDA decoupling reaction, while the DA coupling is not immediately apparent in this graph as it occurs gradually while cooling the sample. Over the course of all cycles, the maximum of the RDA peak at 150°C

shifts to higher values. A similar shift has been reported for related systems^{12,37}, for which this was firstly and hypothetically attributed to the RDA reaction being too slow with respect to the DSC timescale. However, this was later disproven and likely the difference lies in a change in the ratio of endo and exo DA-adducts. It has previously been established³⁸ that the exo isomer is thermodynamically favored, which makes it likely that the kinetically favored endo adduct is initially formed in excess. Nevertheless, during the course of heating and cooling this equilibrium shifts towards the more favored exo product, which in turn should have a higher temperature of RDA decoupling. This is in striking agreement with the NMR experiments (*vide supra*) and thus provides direct confirmation of the formation and importance of the adducts on the thermal behavior.

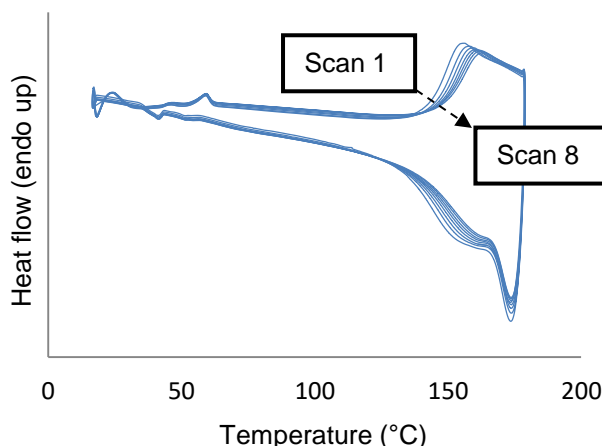


Figure 4: DSC measurements of reversible network formation for DPA-fur/Tereph

In order to assess the mechanical properties of the material, DMTA measurements were performed. The formation of solid, homogeneous bars is an indication of the reversibility of the compound, as without reversible crosslinking taking place only a compressed powder would be obtained. In order to elucidate the effect of the crosslinking degree on the softening temperature ($\tan(\delta)$ peak), various samples containing different maleimide:furan ratios were prepared. Figure 5 shows G' and G'' as well as the $\tan(\delta)$ for samples with 1:1 1:2 and 1:10 mol:mol ratios of maleimide:furan groups.

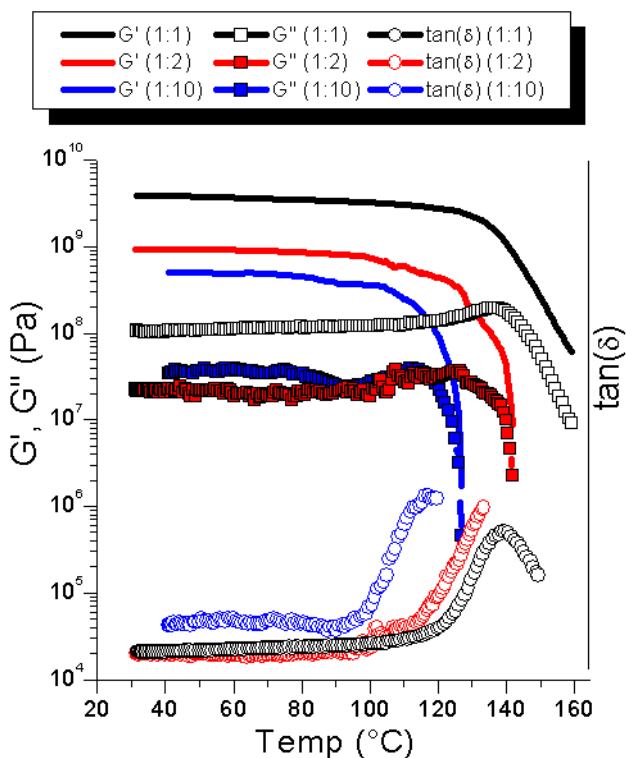


Figure 5: DMTA measurements of polymers containing various maleimide to furan ratios.

The softening temperature is 137°C, 135°C and 120 °C for the 1:1, 1:2 and 1:10 ratios of maleimide:furan respectively. The effect of the MAL:FUR ratio on the softening point is less expressed than that reported for related systems (Figure 6). A highly substituted (60%) polyketone-based system (PK50) displays a drop in softening temperature from 102°C to 86°C upon changing the MAL:FUR ratio from 1:1 to 1:2 (halving the amount of bismaleimide). An additional halving of the amount of bismaleimide leads to an even larger decrease of the softening point (22°C lower). A similar trend is observed when looking at the system obtained by the copolymerization of bis(hydroxymethyl)furane and succinic acid (PFS)³⁹: when halving the amount of bismaleimide the softening point drops from 38°C to 9°C. By noticing that this value is below the T_g of the uncrosslinked material (15°C)⁴⁰, the authors attribute this to a disturbance of the chain packing by the presence of crosslinker. For DPA-fur the effect of halving the amount of bismaleimide on the softening temperature is practically insignificant; only a drop of 2°C is observed.

Probably the large difference in softening point for the other systems stems from the low T_g of uncrosslinked polymer. When the amount of crosslinker is reduced, the material behaves more and more like a branched polymer structure than a three-dimensional network. The T_g of the material will become closer to that of the uncrosslinked polymer as fewer crosslinker is added. Interestingly, when reducing the amount of crosslinker even further to the ratio of 1:10 MAL:FUR the softening point of DPA-fur also drops below the T_g of the uncrosslinked polymer. The aromatic groups present in DPA-fur are likely involved in π - π stacking interactions which are disturbed by the presence of small amounts of crosslinker. Apparently the π - π -interactions have a larger influence on the softening point than the small amount of crosslinker present. Finally, the softening point observed for a 1:1 MAL:FUR DPA-fur network is much closer to the theoretical RDA temperature for furane and maleimide groups (e.g. 150⁴¹), which could be an indication of a more stable crosslink. Furthermore, as the material remains crosslinked for a larger range of temperatures (e.g. decrosslinking occurs at higher temperatures) this should provide a larger window of application for the resulting material than the PFS and PK50 based materials.

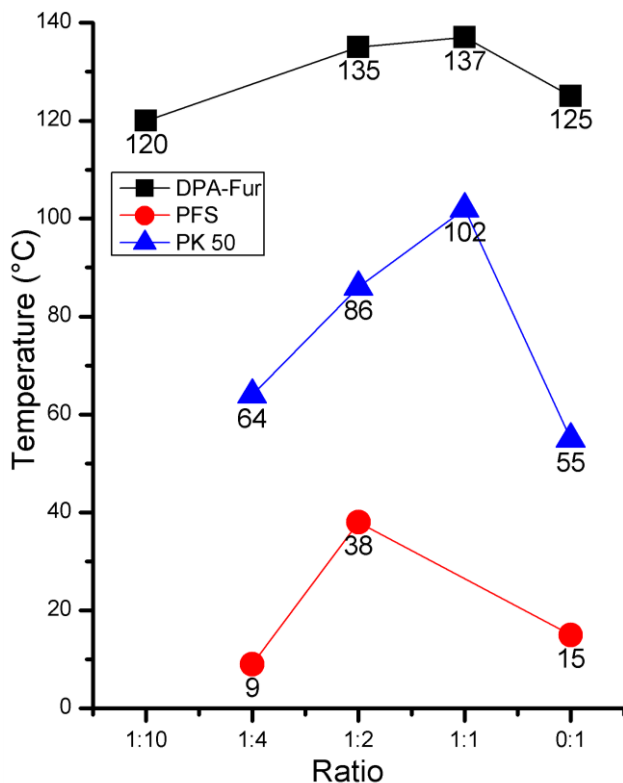


Figure 6: Softening temperature of various DA-systems for varying MAL:FUR ratios (0:1 refers to the uncrosslinked polymer)

Another way of influencing the thermal behavior is the use of a different crosslinker³⁹. When the aromatic bismaleimide was substituted by an aliphatic one containing a C12 chain between the maleimide groups, a drop of the softening point by 10°C was observed (Figure 7). These results are comparable to findings reported by Zeng *et al* where a drop of approximately 10°C was reported as well for the 1:2 ratio of MAL:FUR when substituting the aromatic bismaleimide for an aliphatic one containing a C₂OC₂OC₂ ether bridge⁴². This behavior likely comes from the increased flexibility of the aliphatic bismaleimide compared to its aromatic counterpart. This increased flexibility should result in a less rigid network. Furthermore, as suggested by the gelation experiments, the aliphatic crosslinker should be able to participate in backbiting coordination, limiting its availability for the formation of crosslinks. Independently of the exact mechanism, the observed results indicate that also in the case of the present polymer, the

amount and structure of the crosslinker might be conveniently used to manipulate the mechanical behavior of the end product.

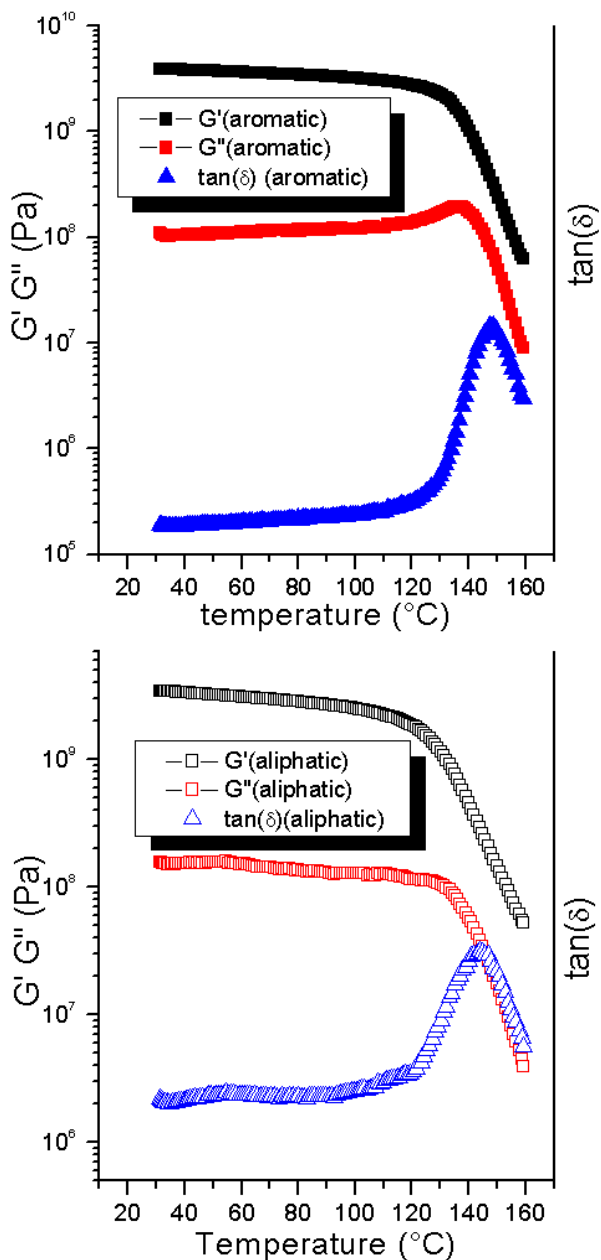


Figure 7 DMTA measurements of aromatic (top) and aliphatic (bottom) crosslinkers.

In order to assess the mechanical reworkability and thermal recovery of the material, cyclical DMTA measurements were performed. Samples were heated to 160°C and successively cooled back to room temperature (20°C) for 4 cycles (Figure 8). During cooling the material is expected to re-form the DA crosslinks which have been broken at high temperatures by the RDA reaction (which is dominant at these elevated temperatures). If the DA reaction is performed consistently, the material should display the same mechanical properties in all subsequent cycles. A relatively large change in properties is observed between the first and second cycle, this is consistent with the shift observed in DSC, and the explanation given of a kinetically favored orientation (endo vs exo) still holds true. However, after each consecutive cycle the softening point of the material increases. If the only equilibrium involved would be the changing from endo to exo isomer the difference should not be as large as observed. Most likely a more permanent change occurs in the material under these circumstances. A well-known side reaction of the Diels-alder adduct of furan and maleimide is the formation of an aromatic ring via the elimination of water⁴³. The mechanical properties of the material are not expected to change significantly in the low temperature region as in both cases (reversible crosslinked and aromatized crosslinked) a covalently crosslinked network is obtained. However, as the aromatization leads to an irreversible network under these conditions less crosslinks will be broken on each consecutive cycle explaining the ever increasing softening point. This indicates that the observed system is only able to display full reversibility for a few thermal cycles.

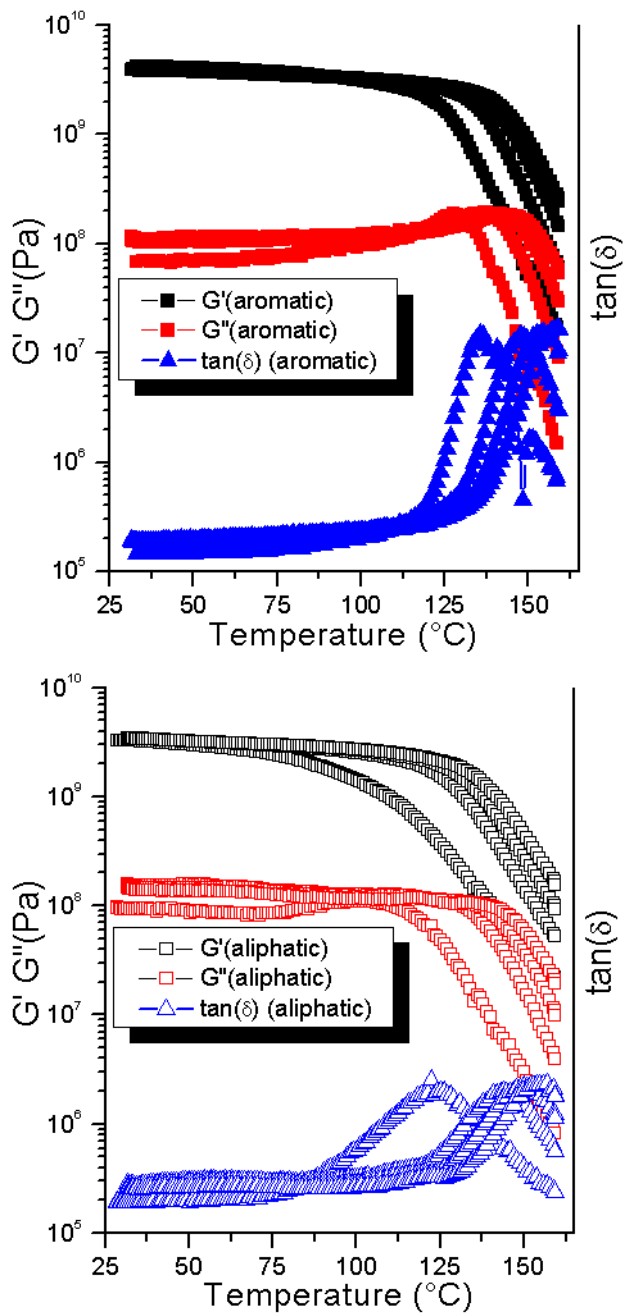


Figure 8: Cyclical DMTA measurements using the aromatic crosslinker (top) and the aliphatic crosslinker (bottom). Only the heating steps are shown.

2.5 Conclusions

In this work, diphenolic acid has successfully been functionalized with furfurylamine, yielding a furan-bearing diol. This has been incorporated into an aromatic polyester by reacting it with terephthaloyl chloride in an interfacial polycondensation reaction. The polymer obtained contains pendant furan groups which are able to participate in reversible crosslinking, forming a thermally reversible network.

Reversibility is shown in solution by gelation experiments where a gel is formed between 15 and 45 minutes when employing the aromatic crosslinker, depending on the ratio of maleimide to furan groups. Gelation with an aliphatic crosslinker takes significantly longer: 143 to 368 minutes, depending on the maleimide to furan ratio. Decoupling (evident by the transition of gel to solution) occurs for all samples in approximately 3 minutes at 150°C for all experiments.

Reversibility in the solid state is shown by cyclical DSC measurements where the peak for RDA decoupling has a maximum at 154°C for the first cycle. The maximum shifts slightly towards 160°C for the 8th cycle but all other thermal properties of the material remain constant. The small shift in the maximum is attributed to differences in the ratio of endo and exo adducts formed. NMR measurements showed a clear shift from a kinetic equilibrium to a thermodynamical one.

The material displays recovery of mechanical properties, as evident from cyclical DMTA measurements, after an initial change the polymer displays a high degree of recovery. However the increase in softening point after each cycle suggests that irreversible aromatization occurs of the furanes and maleimides.

2.6 References

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