

# Polymer Chemistry

rsc.li/polymers



ISSN 1759-9962

**PAPER**

Dina Maniar, Talita Martins Lacerda *et al.*  
Synthesis of semi-rigid-biobased polyesters from renewable  
furanic cyclobutane diacid



Cite this: *Polym. Chem.*, 2025, **16**, 3808

## Synthesis of semi-rigid-biobased polyesters from renewable furanic cyclobutane diacid

Luan Moreira Grilo,<sup>a,b</sup> Sara Faoro,<sup>b</sup> Beatriz Agostinho,<sup>d</sup> Andreia F. Sousa,<sup>d</sup> Nathanael Guigo,<sup>e</sup> Katja Loos,<sup>a</sup> Dina Maniar\*<sup>a</sup> and Talita Martins Lacerda\*<sup>b</sup>

The pursuit of novel sustainable materials is driving advancements in polymer science, with the consolidation of furfural and hydroxymethylfurfural derivatives as key renewable building blocks. 3,4-Di(furan-2-yl)cyclobutane-1,2-dicarboxylic acid (CBDA) is a highly promising rigid-structure biobased monomer that is readily synthesized from furfural. In this study, we expanded the scope of CBDA-based polymers by synthesizing this platform molecule and investigating its polymerization with a series of aliphatic diols of varying chain lengths. CBDA was successfully synthesized from furfural-derived 3-(2-furyl)acrylic acid through a green and efficient UV-mediated solid-state dimerization reaction. Subsequent polymerization was carried out in bulk via a two-step method. The success of the polymerization was confirmed through ATR-FTIR, <sup>1</sup>H NMR, and <sup>13</sup>C CP/MAS NMR spectroscopy. The resulting polymers presented average molecular weights ( $\overline{M}_n$ ) of up to 11 200 g mol<sup>-1</sup>. Thermogravimetric analysis (TGA) revealed good thermal stability, with a  $T_{d10\%}$  ranging from 263 to 284 °C and 50% weight retention observed up to 388 °C. Furthermore, DSC analysis indicated that the glass transition temperature of the polymers could be tailored, varying from 6 to 52 °C depending on the chain length of the utilized diol. These results underscore the potential of CBDA as a renewable rigid monomer for the development of sustainable biobased materials.

Received 15th May 2025,  
Accepted 31st July 2025

DOI: 10.1039/d5py00485c

rsc.li/polymers

## Introduction

Humans' dependence on fossil resources as feedstocks for fuels, chemicals, and materials has led to numerous environmental concerns, such as the direct relationship between the exploitation of such resources and greenhouse gas emissions, which severely affect the global climate.<sup>1,2</sup> Furthermore, the sources of these raw materials, namely petroleum and gas reserves, are unequally distributed worldwide, and their supply and price are volatile and subjected to political speculation.<sup>3</sup> Thus, lignocellulosic biomass has become an increasingly attractive feedstock, owing to its widespread availability from agricultural, forestry, and certain industrial waste streams, with an estimated annual global production exceeding 200

billion metric tons.<sup>4,5</sup> In addition to being inexpensive and abundant, its renewable nature plays a crucial role in promoting a more circular and sustainable economy.<sup>6</sup> As a result, research on biomass valorization has flourished in recent decades, driven by the growing importance of developing strategies to convert this complex feedstock into value-added chemicals.<sup>7</sup>

In this context, the search for biobased platform chemicals to develop innovative renewable polymers is progressing rapidly. Notably, furfural (F) and hydroxymethylfurfural (HMF), derived from lignocellulosic feedstocks, stand out as versatile platform molecules with broad applications in the synthesis of high-value biobased compounds.<sup>8–13</sup> With the advancement of a new generation of biobased polymers to potentially replace their fossil-based counterparts, certain F and HMF derivatives have gained popularity as renewable building blocks. These include molecules such as furfuryl alcohol, employed in resins and composites;<sup>14–17</sup> furfurylamine, used in polyamides and polyureas;<sup>18–21</sup> furfuryl methacrylate, utilized in polymers and composites;<sup>22–24</sup> 2,5-bis(hydroxymethyl)furan (BHMF), applied in polyesters;<sup>25–28</sup> and 2,5-furandicarboxylic acid (FDCA), used in polyamides and polyesters, most notably poly(ethylene furanoate), a renewable alternative to poly(ethylene terephthalate) (PET).<sup>29–35</sup> These relevant furan building blocks are depicted in Fig. 1.

<sup>a</sup>Macromolecular Chemistry and New Polymeric Materials, Zernike Institute for Advanced Materials, Faculty of Science and Engineering, University of Groningen, The Netherlands. E-mail: d.maniar@rug.nl

<sup>b</sup>Biotechnology Department, Lorena School of Engineering, University of São Paulo, Brazil. E-mail: talitalacerda@usp.br

<sup>c</sup>Center for Education and Research on Macromolecules, CESAM Research Unit, University of Liege, Belgium

<sup>d</sup>CICECO-Aveiro Institute of Materials, Department of Chemistry, University of Aveiro, Portugal

<sup>e</sup>Mines Paris, PSL University, CNRS, CEMEF (UMR 7635), 06904 Sophia Antipolis Cedex, France



Fig. 1 Common furan building blocks for biobased materials.

3-(2-Furyl)acrylic (FAA) acid is another furfural derivative with interesting potential applications. As a photoactive molecule, it can easily undergo a [2 + 2] photocycloaddition reaction under UV light to produce its dimer 3,4-di(furan-2-yl)cyclobutane-1,2-dicarboxylic acid (CBDA).<sup>36</sup> This dimerization phenomenon is also observed in other molecules, such as cinnamic acid and its analogs, including 3-(2-thienyl)acrylic acid and 3-(3-pyridyl)acrylic acid, as well as in heteroarylene-vinyls and related compounds.<sup>37–40</sup> These reactions consistently yield substituted cyclobutane derivatives, which are of significant interest because of their versatile applications. The cyclobutane structure, in particular, has found unique utility across a wide range of research fields, including materials science, fuel technology, and pharmaceutical synthesis.<sup>41–45</sup> CBDA has relevant potential to serve as a biobased rigid diacid building block, with previous studies demonstrating its suitability as a polycondensing monomer.<sup>36,46</sup> Moreover, when used as hardener with a flexible epoxidized vegetable oil, it formed a crosslinked network that exhibited high glass transition temperature, high strength, and relatively good toughness.<sup>47,48</sup> Additionally, in many of these cyclobutane molecules, the ring-forming bonds are reversible and can be cleaved under specific photochemical, mechanical, and/or thermal stimuli.<sup>49–53</sup> These characteristics make them promising candidates as platform molecules for the design of degradable or recyclable polymers, which are in growing demand as part of strategies to achieve more sustainable end-of-life management of polymeric waste.<sup>54–57</sup>

Therefore, this work further explores CBDA as a platform molecule for polymer synthesis, expanding the library of linear CBDA-based polyesters by pairing it with an assortment of aliphatic diols and obtaining polymers with tunable thermal properties. CBDA was synthesized *via* the solid-sated suspension photocycloaddition of FAA, using commercially available UVA LEDs and recycling the reaction medium of the process. Then,

CBDA was polymerized in bulk with different diols to avoid the excess use of organic solvents, using titanium(IV) isopropoxide (TTIP) as catalyst. Enzymatic polymerization of the ethyl ester of CBDA was also attempted, employing immobilized *Candida antarctica* lipase B (CALB) as catalyst. ATR-FTIR, <sup>1</sup>H NMR, <sup>13</sup>C CP/MAS NMR, TGA, DSC, and WAXD were used to characterize the synthesized polymers.

## Experimental section

### Materials

3-(2-Furyl) acrylic acid (FAA, 99%), 1,4-butanediol (99%), 1,6-hexanediol (99%), 1,8-octanediol (98%), 1,10-decanediol (98%), titanium(IV) isopropoxide (TTIP, 99.999%), trifluoroacetic acid (TFA, 99%), triethylamine (TEA, 99.5%), oxalyl chloride (99%), Novozyme 435 (N435, *Candida antarctica* lipase B immobilized in acrylic resin, recombinant, expressed in *Aspergillus niger*, ≥5000 U g<sup>-1</sup>), molecular sieves (4 Å), deuterated dimethyl sulfoxide (DMSO-d<sub>6</sub>, 99.9%), and deuterated chloroform (CDCl<sub>3</sub>, 99.8%), were purchased from Sigma-Aldrich. *N*-Hexane (99%), chloroform (CHCl<sub>3</sub>, 99.5%), methanol (anhydrous, 99.8%), ethanol (anhydrous, 99.8%), tetrahydrofuran (THF, 99.5%), toluene (anhydrous, 99.8%), *N,N*-dimethylformamide (DMF, 99.8%), and ethyl acetate (99%) were purchased from Avantor. N435 was pre-dried in the presence of phosphorus pentoxide in a desiccator at room temperature under a high vacuum for 12 h. The molecular sieves (4 Å) were pre-activated at 250 °C under vacuum. The other chemicals were used as received without further purification.

### Analytical methods

Proton nuclear magnetic resonance (<sup>1</sup>H NMR) spectra were acquired at 25 °C using a Bruker Avance NMR spectrometer operating at 600 MHz. Chemical shifts were recorded in parts

per million (ppm) relative to the solvent peak. Spectral analysis was performed using MestreNova 14 software.

Solid-state carbon-13 cross-polarization/magic angle spinning nuclear magnetic resonance ( $^{13}\text{C}$  CP/MAS NMR) spectra were acquired at 25 °C *via* a Bruker Avance NMR spectrometer operating at 400 MHz. Spectral analysis was performed *via* MestreNova 14 software.

Attenuated total reflection-Fourier transform infrared (ATR-FTIR) spectra were recorded *via* a Bruker VERTEX 70 spectrometer equipped with a Platinum-ATR diamond single-reflection accessory. Measurements were conducted with a resolution of 4  $\text{cm}^{-1}$ , and spectra were collected over a 4000–400  $\text{cm}^{-1}$  range, with 16 scans per sample. Atmospheric compensation and baseline correction were applied to the spectra using OPUS spectroscopy software (v7.0, Bruker Optics).

Thermogravimetric analysis (TGA) was conducted *via* a TA Instruments D2500 TGA instrument. The samples were heated from 25 °C to 700 °C at a rate of 10 °C  $\text{min}^{-1}$  under a nitrogen atmosphere (at a  $\text{N}_2$  flow rate of 50  $\text{mL min}^{-1}$ ). Prior to the TGA measurement, each sample was preheated to 100 °C and held at that temperature for 20 min to remove residual moisture and solvents.

Differential scanning calorimetry (DSC) was employed to assess the thermal transitions of the synthesized polyesters. Measurements were carried out using a TA Instruments Q1000 DSC. Samples weighing 10–15 mg were placed in nonhermetically sealed aluminum pans. Heating-cooling-heating cycles were conducted from –50 °C to 200 °C at a rate of 10 °C  $\text{min}^{-1}$ , under a mild nitrogen flow rate of 25  $\text{mL min}^{-1}$ .

Wide-angle X-ray diffraction (WAXD) patterns of the synthesized polyesters were recorded using a Bruker D8 Advance diffractometer with  $\text{CuK}\alpha$  radiation ( $\lambda = 0.1542 \text{ nm}$ ). Data were collected at room temperature in the angular range of 5–50° ( $2\theta$ ).

### Synthesis of 3,4-di(furan-2-yl)cyclobutane-1,2-dicarboxylic acid (CBDA)

CBDA was synthesized as previously reported.<sup>36,48</sup> 3-(2-furyl) acrylic acid powder (FAA, 12 g, 86.9 mmol) was dispersed in 250 mL of hexane and transferred to a quartz Erlenmeyer flask. The suspension was stirred and irradiated with three consumer-grade LED UV lamps (365 nm, 135 W) for 48 h. Occasionally, the walls of the flask were scraped off to remove some attached solid material, and extra hexane was added to keep the volume approximately constant, considering its evaporation. To confirm that the FAA conversion exceeded 95% and determine when to stop the reaction, 1 mL samples were collected after 24 h and 48 h for  $^1\text{H}$  NMR analysis. The product was isolated *via* filtration and dried under vacuum. CBDA was obtained as a pale-beige solid, and the remaining hexane was recovered for reuse.

### Melt polymerization of CBDA

CBDA was bulk polymerized with different diols—1,4-butanediol, 1,6-hexanediol, 1,8-octanediol, or 1,10-decanediol—in a

two-step process, producing the polymers PBCB, PHCB, POCB, and PDCB, respectively. Firstly, 2 g (7.24 mmol) of CBDA, 7.24 mmol of diol, and a magnetic stirrer were added to a two-neck round bottom flask attached to a short path vacuum distillation setup, and the whole system was flushed with  $\text{N}_2$ . The temperature was gradually increased to 160 °C at a rate of about 4 °C  $\text{min}^{-1}$ , after which 0.09 mmol of TTIP was added. The mixture was kept under  $\text{N}_2$  atmosphere at 160 °C for 4 h with constant stirring. Then, the temperature was increased to 180 °C at approximately 2 °C  $\text{min}^{-1}$ , followed by vacuum application for 2 h. After the second reaction step ended and the mixture had cooled down to room temperature, approximately 25 mL of chloroform and approximately 5 drops of TFA were added to the reaction flask. The mixture was stirred vigorously and left to dissolve overnight. However, in all the cases, the polymers did not fully dissolve in the solvent mixture, resulting primarily in a suspension of small polymer particles in the chloroform/TFA solution. This suspension was subsequently precipitated in cold methanol. Then, the polymers were separated from the supernatant by centrifugation and were left to dry overnight in a vacuum oven at 45 °C.

### Enzymatic polymerization of CBDA-based diester

First, CBDA was converted to its diethyl ester (CBDE), as previously reported.<sup>36</sup> In a round-bottom flask, 2.5 g of CBDA (9.1 mmol) was dissolved in 90 mL of dry THF and purged with nitrogen. Then, 3.11 mL of oxalyl chloride (36.4 mmol) and two drops of DMF were added to the flask, and the solution was stirred for 1 h under an inert atmosphere. After this period, the THF and excess oxalyl chloride were removed under vacuum to obtain CBDA acyl chloride. TEA (2.86 mL, 20.5 mmol) was dissolved in 46 mL of anhydrous ethanol (0.78 mol) and slowly added, *via* a dropping funnel, into the flask containing CBDA acyl chloride over an ice bath. After this addition, the reaction mixture was stirred for 2 h at room temperature. The mixture was then filtered to remove any solid impurities, and the solvent in the filtrate was evaporated under vacuum, yielding CBDE as a viscous brownish-yellow oil. Flash column chromatography (ethyl acetate : hexane = 2 : 98) was used to purify the diester monomer further.

Following the CBDE synthesis, its enzymatic polymerization with aliphatic diols was carried out based on the one-step procedure reported by Jiang *et al.* (2016).<sup>33</sup> First, in a 25 mL round-bottom flask, 0.5 g of CBDE (1.5 mmol) and 1.5 mmol of a diol (1,4-butanediol, 1,6-hexanediol, 1,8-octanediol, or 1,10-decanediol) were dissolved in anhydrous toluene (500 wt% relative to all monomers). Then, pre-dried N435 (20 wt%) and pre-activated 4 Å molecular sieves (200 wt%) were added, and the flask was purged with nitrogen. The flask was sealed and placed in an oil bath at 90 °C for 72 h under magnetic stirring. After this period, the toluene was evaporated under a stream of air at room temperature, and the resulting mixture was dissolved in 15 mL of chloroform. Next, N435 and molecular sieves were removed by filtration, and the filter paper and its contents were washed three times, with 10 mL of chloroform each. The excess chloroform in the filtrate was

removed under vacuum, and the concentrated solution was added dropwise into an excess of cold methanol to precipitate the synthesized polymers. However, no precipitation occurred, so the crude reaction mixture was kept for  $^1\text{H}$  NMR analysis.

**3,4-Di(furan-2-yl)cyclobutane-1,2-dicarboxylic acid (CBDA).**  $^1\text{H}$  NMR (600 MHz, DMSO,  $\delta$ , ppm): 12.55 (2H, br), 7.40 (2H, d), 6.25 (2H, m), 6.09 (2H, d), 4.05 (2H, m), 3.67 (2H, m); ATR-FTIR ( $\text{cm}^{-1}$ ): 3050 (O–H stretching vibrations), 1695 (C=O stretching vibrations), 1414 (O–H bending vibrations).

**Poly(butylene cyclobutane-1,2-dicarboxylate) (PBCB).**  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ ,  $\delta$ , ppm): 7.21 (2H, m), 5.93–6.19 (4H, m), 4.21 (2H, m), 4.09–4.13 (4H, m), 3.83 (2H, m), 1.67–1.74 (4H, m);  $^{13}\text{C}$  CP/MAS NMR (400 MHz,  $\delta$ , ppm): 171.4, 152.8, 142.1, 110.3, 107.4, 64.9, 42.4, 39.0, 25.4.

**Poly(hexamethylene cyclobutane-1,2-dicarboxylate) (PHCB).**  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ ,  $\delta$ , ppm): 7.22 (2H, m), 5.93–6.19 (4H, m), 4.22 (2H, m), 4.06–4.17 (4H, m), 3.83 (2H, m), 1.34–1.61 (8H, m);  $^{13}\text{C}$  CP/MAS NMR (400 MHz,  $\delta$ , ppm): 171.4, 152.8, 142.1, 110.5, 107.5, 65.1, 43.1, 38.8, 28.6, 25.7.

**Poly(octamethylene cyclobutane-1,2-dicarboxylate) (POCB).**  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ ,  $\delta$ , ppm): 7.22 (2H, m), 5.93–6.19 (4H, m), 4.22 (2H, m), 4.07–4.17 (4H, m), 3.83 (2H, m), 1.30–1.61 (12H, m);  $^{13}\text{C}$  CP/MAS NMR (400 MHz,  $\delta$ , ppm): 171.3, 153.0, 142.0, 110.4, 107.4, 65.2, 43.3, 38.8, 29.0, 26.3.

**Poly(decamethylene cyclobutane-1,2-dicarboxylate) (PDCB).**  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ ,  $\delta$ , ppm): 7.22 (2H, m), 5.93–6.19 (4H, m), 4.22 (2H, m), 4.07–4.17 (4H, m), 3.84 (2H, m), 1.26–1.61 (16H, m);  $^{13}\text{C}$  CP/MAS NMR (400 MHz,  $\delta$ , ppm): 171.4, 153.0, 142.0, 110.1, 107.5, 65.2, 43.0, 38.9, 29.3, 26.4.

**CBDA-based polyesters ATR-FTIR ( $\text{cm}^{-1}$ ).** 2926–2957, 2852–2897 (asymmetric and symmetric C–H stretching vibrations); 1724–1726 (C=O stretching vibrations); 1163–1161 (C–O stretching vibrations); 1012 (=C–O–C= ring vibrations); 796–800, 733–727 (=C–H out-of-plane deformation vibrations).



Scheme 1 Synthesis of CBDA.



Scheme 2 Two-step bulk polycondensations of CBDA and different diols.

## Results and discussion

CBDA was successfully synthesized *via* photocycloaddition of FAA (Scheme 1) based on a previously reported procedure.<sup>36,48</sup>

The reaction conversion was determined to be approximately 95% (calculated by  $^1\text{H}$  NMR – Fig. S1). It is important to highlight the green aspects of this process, including the recyclability of the reaction medium and the use of commercially available UVA LEDs. These LEDs can achieve an electrical efficiency of 40–50%, surpassing that of traditional mercury lamps.<sup>58</sup> Furthermore, LEDs have a significantly lower environmental impact than mercury lamps do,<sup>59</sup> mainly due to the impacts of the production of the latter.<sup>60</sup>

Polyesters were synthesized from CBDA and different diols in a two-step bulk polycondensation process based on literature reports (Scheme 2).<sup>29,61–63</sup> In the first step, consisting on a prepolymerization, a catalyst is introduced to promote the esterification reaction, forming low-molecular-weight products (oligomers). This step is critical as it enables the monomers to react under milder conditions, producing short chains with higher boiling points and improved thermal stability compared to the starting monomers. Subsequently, elevated temperatures and reduced pressure are typically applied to achieve higher molecular weight polymers. Since esterification is an equilibrium reaction, removing the byproduct—water, in this case—is crucial to shift the equilibrium forward and drive the formation of polymers. The duration of the second step was determined by visually monitoring the viscosity of the reaction mixture in the PBCB sample, stopping the reaction when the mixture solidified. This timeframe was then applied consistently across the other samples. At the end of each polymerization reaction, the recovered polymers were weighed, and the yields were calculated (Table 1). Notably, the obtained polymers exhibited partial solubility in the standard  $\text{CHCl}_3$ /TFA mixture used for recovery and were insoluble in other common organic solvents, including DMSO, DMF, and THF.

As previously mentioned, the enzymatic polymerization of the diethyl ester of CBDA (CBDE) and aliphatic diols was also investigated. The enzyme chosen for this experiment was *Candida antarctica* lipase B (CALB), specifically the commercially available Novozyme 435, which is immobilized in acrylic resin. This widely used biocatalyst has been reported to effectively catalyze the synthesis of FDCA- and BHMF-based polymers.<sup>26–28,33,34</sup> First, the esterification of CBDA was per-

**Table 1** CBDA-based polyesters and their corresponding isolation yields, number-average degree of polymerization, and number-average molecular weight

| Polyester   | Abbreviation | Average yield <sup>a</sup> (%) | $\overline{DP}_n^b$ | $\overline{M}_n^b$ (g mol <sup>-1</sup> ) |
|---|--------------|--------------------------------|---------------------|---|
| Poly(butylene cyclobutane-1,2-dicarboxylate)      | PBCB         | 64.1                           | 7                   | 2700                                      |
| Poly(hexamethylene cyclobutane-1,2-dicarboxylate) | PHCB         | 43.1                           | 10                  | 4000                                      |
| Poly(octamethylene cyclobutane-1,2-dicarboxylate) | POCB         | 66.3                           | 28                  | 11 200                                    |
| Poly(decamethylene cyclobutane-1,2-dicarboxylate) | PDCB         | 61.6                           | 12                  | 5400                                      |

<sup>a</sup> These values are the average of two experimental results. <sup>b</sup> These values were estimated through <sup>1</sup>H NMR analysis *via* the following equations:

$\overline{DP}_n = \frac{I_C \times m_I \times n_I}{I_I \times m_C}$  and  $\overline{M}_n = n_R \times M_R + M_E$ , where  $I_C$  is the integral intensity of the peak assigned to the furanic protons of the repeating unit ( $C$ , at 5.93 ppm),  $I_I$  is the integral intensity of the peak assigned to the methylene protons of the hydroxymethyl end groups ( $I$ , approximately 3.45 ppm),  $m_C$  is the number of protons  $C$  present in the repeating unit,  $m_I$  is the number of protons  $I$ ,  $n_I$  is the number of hydroxymethyl end groups,  $n_R$  indicates the number of repeating units ( $\overline{DP}_n$ ),  $M_R$  is the molecular weight of one repeating unit,  $M_E$  is the combined molecular weight of the end groups. Here, we assumed that all polyesters were terminated with one hydroxymethyl group and one carboxylic acid group.<sup>68</sup>

formed as described by Wang *et al.* (2018).<sup>36</sup> CBDE was chosen over CBDA for this reaction because of its superior solubility in the reaction medium (toluene).<sup>34,64</sup> The use of CBDA was also avoided because certain acidic substrates have been shown to deactivate the enzyme, thereby hindering the esterification process.<sup>65</sup> The one-step solution polymerization of CBDE with aliphatic diols was carried out following the method outlined by Jiang *et al.* (2016).<sup>33</sup> However, no polymeric product was recovered at the end of the reactions. <sup>1</sup>H NMR analysis of the reaction mixture confirmed this, revealing only the signals corresponding to the starting reagents, as illustrated in Fig. S2. The primary hypothesis for the failure of the enzymatic polymerization centers on the specificity of the enzyme. The active-site cavity of CALB is reportedly shaped like a tunnel, which restricts the steric positioning of bulky or branched substrates.<sup>66,67</sup> Given the cyclobutane structure of CBDE, with its pendant furan groups, the substrate is likely too sterically demanding for CALB's active site, preventing effective catalysis of the transesterification reaction with the aliphatic diols.

The synthesized polyesters were first analyzed *via* ATR-FTIR, and their spectra, along with those acquired from CBDA, are presented in Fig. 2. The shift of the carbonyl C=O stretching band from 1695 cm<sup>-1</sup> to 1725 cm<sup>-1</sup> and the C–O stretching peak at 1162 cm<sup>-1</sup> present in all the spectra of the polymers demonstrate the formation of ester bonds between CBDA and the diols. The disappearance of the O–H stretching band at 3050 cm<sup>-1</sup> and the O–H bending band at 1414 cm<sup>-1</sup> are also consistent with the formation of ester bonds. Additionally, the intensities of the asymmetric and symmetric C–H stretching bands at approximately 2942 cm<sup>-1</sup> and 2874 cm<sup>-1</sup>, respectively, increased progressively in the polymer samples as the aliphatic chain of the employed diol increased, as expected. Furthermore, furan ring bands can be observed in the fingerprint region of the polymers and CBDA spectra, specifically the =C–O–C= ring vibrations at 1012 cm<sup>-1</sup> and the =C–H out-of-plane vibrations at approximately 798 and 730 cm<sup>-1</sup>.

To further assess the molecular structures of the synthesized polymers, the polyesters were analyzed through <sup>1</sup>H NMR in CDCl<sub>3</sub>. Owing to the material's low solubility, it was allowed to dissolve overnight in CDCl<sub>3</sub> with a few drops of TFA. Before analysis, the insoluble fraction was filtered off. The resulting spectra are depicted in Fig. 3. The signals labeled A, B, and C – at 7.22, 6.19, and 5.93 ppm, respectively – are attributed to the protons from the furan ring (–CH=). The protons of the cyclobutane core generate signals D and E (4.21 and 3.83 ppm). The protons from the methylene units of the aliphatic diols (–CH<sub>2</sub>–) give rise to signals at 4.11 ppm (F), 1.61–1.67 ppm (G), and 1.26–1.40 ppm (H). The presence of peak F, in particular, indicates ester bond formation, providing additional evidence for the formation of polymers/oligomers. The peaks at 0.83 and 1.25 ppm, which are mainly detectable in the spectrum of PBCB, arise from impurities, most probably due to vacuum grease contamination of the samples. The low-intensity signals at 6.22–7.10 ppm and 7.27–7.49 ppm may come from minor impurities that are also present in the CBDA <sup>1</sup>H NMR spectrum (see the SI).

The small peak I at 3.45 ppm can be attributed to the methylene protons of the hydroxymethyl end-groups (HO–CH<sub>2</sub>–). In addition to peak C—the most well-defined signal

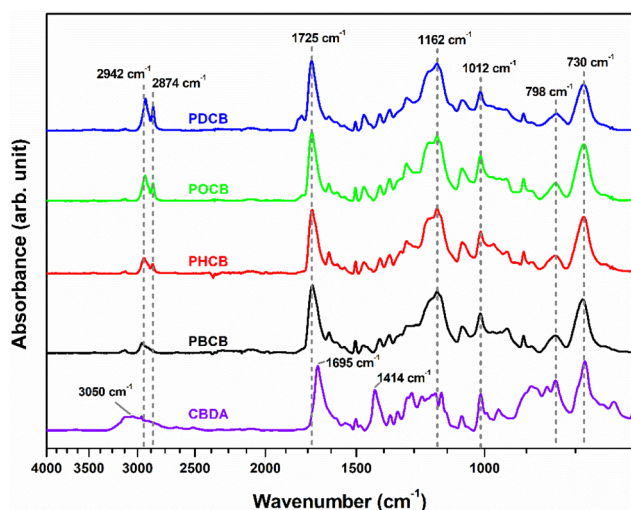
**Fig. 2** ATR-FTIR spectra of CBDA and its corresponding polyesters.



Fig. 3  $^1\text{H}$  NMR spectra of the polyesters soluble fractions.



Scheme 3 Possible (a) mechanism of C5 electrophilic substitution, and (b) resulting crosslinked polymer structure for PBCB.

originating from the repeating unit structure—this peak was used to estimate the number-average degree of polymerization ( $\overline{DP}_n$ ) and the number-average molecular weight ( $\overline{M}_n$ ) of the soluble portion of the polymers (Table 1).<sup>68</sup> Naturally, an increasing trend in the  $\overline{M}_n$  of the samples is expected, which is simply due to the differences in the molecular weights of the diols used. Moreover,  $\overline{DP}_n$  slightly increases with increasing diol chain length, as observed in the progression from PBCB to PHCB to PDCB, with average repeating units per chain of 7, 10, and 12, respectively. However, POCB exhibits a significantly higher  $\overline{DP}_n$  of 28. This deviation could stem from a limitation in the estimation method or may suggest that a diol chain length of 8 carbons is more favorable for the reaction than a length of 4, 6, or 10 carbons. This preference could be due to factors such as catalyst selectivity or improved miscibility in the reaction mixture.

Considering that the highest obtained  $\overline{DP}_n$  was 28, it is evident that all the samples consisted primarily of oligomers. Wang *et al.* reported the synthesis and further polymerization of a similar cyclobutane monomer with ethylene glycol, obtaining both oligomers and polymers with  $\overline{M}_n$  values of 1968 and 22039, respectively.<sup>61</sup> Furthermore, their polymer exhibited low solubility in common organic solvents. On this basis, it can be inferred that the insoluble fraction of the materials synthesized in this study likely consists of molecules with longer chain lengths than those detected by <sup>1</sup>H NMR analysis. This increased chain length likely correlates with the observed insolubility.

Another possible explanation for the low solubility of these polyesters in common organic solvents is the occurrence of crosslinking during the polymerization reaction. The C5 carbon in the furan ring is fairly reactive and capable of undergoing electrophilic substitution, and the possibility of it reacting with the diols and producing interchain bonds is not null

(Scheme 3).<sup>69,70</sup> Therefore, solid-state <sup>13</sup>C CP/MAS NMR analysis was employed to obtain more detailed structural data and evaluate the degree of such crosslinking in the material.

The <sup>13</sup>C CP/MAS NMR spectra of the synthesized polymers are depicted in Fig. 4. For all samples, the carbons in the furan ring—labeled A, B, C, and D—give rise to signals at approximately 142.0, 110.4, 107.5, and 152.0 ppm, respectively, with signals for carbons B and C partially overlapping. The carbons in the cyclobutane structure (E and F) produced partially overlapping peaks between 43.0 and 38.8 ppm, and the carbonyl carbon (G) appeared as a signal at 171.4 ppm. The carbon at the ester linkage (H) generates a peak at 65.2 ppm, whereas the remaining aliphatic carbons (I) yield broad signals in the range of 29.3–25.4 ppm. Thus, no new signals indicating covalent crosslinking were detected, suggesting that low solubility might be an innate feature of these polymers. Additionally, as previously mentioned, an analogous study reported the low solubility of similar structured polyesters in common organic solvents without the occurrence of crosslinks.<sup>61</sup> Nonetheless, the analysis results should be considered carefully, given the technique's resolution limitations.

The thermal stability of the polymers was assessed *via* TGA, and the resulting thermograms are presented in Fig. 5. The very small weight loss (up to 0.96%) at approximately 100 °C, which is more evident in Fig. 5b, can be attributed to the elimination of residual solvent/moisture. The polymers showed overall good thermal stability, with  $T_{d5\%}$  values between 250 and 267 °C and  $T_{d10\%}$  values between 263 and 284 °C, with 50% weight retention observed up to 388 °C. All the polyester samples exhibited two main degradation steps, with the derivative of weight loss peaking between 277–310 °C and 390–405 °C. The first degradation step corresponded to a



Fig. 4 Polyesters' <sup>13</sup>C solid-state NMR spectra.



Fig. 5 (a) Weight (%) vs. temperature (°C); (b) derivative weight vs. temperature of the synthesized polymers.

weight loss of 38–49%, whereas the second step resulted in a weight loss of 29–43%. These observations suggest that two distinct thermal decomposition phenomena occur. Additionally, the char yield of the samples ranged between 18–22%.

The thermal properties of the synthesized polyesters were analyzed *via* DSC. The first heating and cooling cycle was employed to eliminate the samples' thermal history, and the second heating curves are depicted in Fig. 6. The glass transition temperature ( $T_g$ ) of the polymers were determined to be 52, 18, 10, and 6 °C for PBCB, PHCB, POCB, and PDCB, respectively, indicating a decreasing trend as the length of the linear monomer increased. This highlights that, as expected, the diol chain length can be strategically selected to tailor the  $T_g$  of the resulting materials. Additionally, the influence of the  $T_g$  could be observed during polymer handling, as PBCB was brittle at room temperature, whereas the other samples were increasingly rubbery at room temperature. Furthermore, none of the DSC curves showed a melting endotherm, which is an evidence that all the polymers were amorphous. This was further substantiated by WAXD analysis, which did not reveal the presence of crystalline domains (Fig. S3).



Fig. 6 Polymers' DSC second heating curve.

## Conclusion

Novel polyesters from furfural-derived CBDA and aliphatic diols were successfully synthesized, achieving yields of up to 66%. The reactions were confirmed *via* multiple analytical techniques, including ATR-FTIR,  $^1\text{H}$  NMR, and  $^{13}\text{C}$  CP/MAS NMR. The resulting polymers exhibited limited solubility in conventional organic solvents, which prevented a thorough assessment of their molecular weights. However,  $^1\text{H}$  NMR analysis allowed for an estimation of the  $\overline{M}_n$  of the soluble polymer fraction, with values reaching up to  $11\,200\text{ g mol}^{-1}$ . One hypothesis for the limited solubility was the potential crosslinking of the materials. However, detailed analyses, particularly  $^{13}\text{C}$  CP/MAS NMR, did not provide any evidence to support this. This finding indicates that low solubility in conventional organic solvents may be an inherent property of the synthesized polymers.

Thermogravimetric analysis of the synthesized polymers demonstrated good thermal stability, with  $T_{d10\%}$  ranging from 261 to 282 °C and 50% weight retention observed up to 386 °C. Moreover, DSC revealed tunable glass transition temperatures varying from 6 to 52 °C depending on the length of the diol used. Additionally, there were no observable melting or crystallization peaks, and wide-angle X-ray diffraction (WAXD) analysis confirmed that the polyesters were 100% amorphous.

Further studies are necessary to obtain a deeper understanding of the physical and mechanical properties of the materials. Such investigations should provide a comprehensive characterization of these biobased polymers and their potential applications. Nevertheless, this work offers valuable insights into the potential of CBDA as a renewable rigid monomer, contributing to the advancement of knowledge in the field of sustainable biobased materials.

## Conflicts of interest

The authors declare no competing financial interest.

## Data availability

The data supporting this article have been included as part of the SI.

Supplementary information containing the  $^1\text{H}$  NMR spectra of CBDA in DMSO- $d_6$ , the  $^1\text{H}$  NMR spectra of the CBDE and 1,4-butanediol enzymatic polycondensation reaction mixture in  $\text{CDCl}_3$ , and the WAXD patterns of PBCB, PHCB, and POCB is available. See DOI: <https://doi.org/10.1039/d5py00485c>.

## Acknowledgements

LMG acknowledges the financial support from the RUG and USP double-degree PhD program and from FAPESP (Process No. 2021/02903-4).

This work was supported by COST Action FUR4Sustain, CA18220, supported by COST (European Cooperation in Science and Technology). This work was developed within the scope of the project CICECO-Aveiro Institute of Materials, UIDB/50011/2020 (DOI: 10.54499/UIDB/50011/2020), UIDP/50011/2020 (DOI: 10.54499/UIDP/50011/2020) & LA/P/0006/2020 (DOI: 10.54499/LA/P/0006/2020), financed by national funds through the FCT/MCTES (PIDDAC). The FCT is acknowledged for the research contract to AFS (CEECINSTLA/00002/2022) and a doctorate grant to BA 2020.04495.BD (DOI: 10.54499/2020.04495.BD). LMG and SF thank COST for the short-term scientific mission (STSM) grant under the framework of COST Action FUR4Sustain (CA18220).

The authors thank Jur van Dijken for the TGA analysis.

## References

- 1 P. T. Benavides, J. B. Dunn, J. Han, M. Bidy and J. Markham, Exploring Comparative Energy and Environmental Benefits of Virgin, Recycled, and Bio-Derived PET Bottles, *ACS Sustainable Chem. Eng.*, 2018, **6**, 9725–9733.
- 2 S. R. Nicholson, N. A. Rorrer, A. C. Carpenter and G. T. Beckham, Manufacturing energy and greenhouse gas emissions associated with plastics consumption, *Joule*, 2021, **5**, 673–686.
- 3 Y. Yang, Z. Liu, H. B. Saydaliev and S. Iqbal, Economic impact of crude oil supply disruption on social welfare losses and strategic petroleum reserves, *Resour. Policy*, 2022, **77**, 102689.
- 4 A. Pandey and Y. C. Sharma, Advancements in biomass valorization in integrated biorefinery systems, *Bioprod. Biorefin.*, 2024, **18**, 2078–2090.
- 5 W. Deng, Y. Feng, J. Fu, H. Guo, Y. Guo, B. Han, Z. Jiang, L. Kong, C. Li, H. Liu, P. T. T. Nguyen, P. Ren, F. Wang, S. Wang, Y. Wang, Y. Wang, S. S. Wong, K. Yan, N. Yan, X. Yang, Y. Zhang, Z. Zhang, X. Zeng and H. Zhou, Catalytic conversion of lignocellulosic biomass into chemicals and fuels, *Green Energy Environ.*, 2023, **8**, 10–114.
- 6 E. J. Cho, L. T. P. Trinh, Y. Song, Y. G. Lee and H.-J. Bae, Bioconversion of biomass waste into high value chemicals, *Bioresour. Technol.*, 2020, **298**, 122386.
- 7 Y. Liao, S.-F. Koelewijn, G. Van den Bossche, J. Van Aelst, S. Van den Bosch, T. Renders, K. Navare, T. Nicolaï, K. Van Aelst, M. Maesen, H. Matsushima, J. M. Thevelein, K. Van Acker, B. Lagrain, D. Verboekend and B. F. Sels, A sustainable wood biorefinery for low-carbon footprint chemicals production, *Science*, 2020, **367**, 1385–1390.
- 8 J. Zhu and G. Yin, Catalytic Transformation of the Furfural Platform into Bifunctionalized Monomers for Polymer Synthesis, *ACS Catal.*, 2021, **11**, 10058–10083.
- 9 C. Xu, E. Paone, D. Rodríguez-Padrón, R. Luque and F. Mauriello, Recent catalytic routes for the preparation and the upgrading of biomass derived furfural and 5-hydroxymethylfurfural, *Chem. Soc. Rev.*, 2020, **49**, 4273–4306.
- 10 V. P. Kashparova, D. V. Chernysheva, V. A. Klushin, V. E. Andreeva, O. A. Kravchenko and N. V. Smirnova, Furan monomers and polymers from renewable plant biomass, *Russ. Chem. Rev.*, 2021, **90**, 750–784.
- 11 A. Jaswal, P. P. Singh and T. Mondal, Furfural – a versatile, biomass-derived platform chemical for the production of renewable chemicals, *Green Chem.*, 2022, **24**, 510–551.
- 12 A. Gandini and T. M. Lacerda, in *Furan Polymers and their Reactions*, Wiley, 2023.
- 13 M. Annatelli, J. E. Sánchez-Velandia, G. Mazzi, S. V. Pandeirada, D. Giannakoudakis, S. Rautiainen, A. Esposito, S. Thiyagarajan, A. Richel, K. S. Triantafyllidis, T. Robert, N. Guigo, A. F. Sousa, E. García-Verdugo and F. Aricò, Beyond 2,5-furandicarboxylic acid: status quo, environmental assessment, and blind spots of furanic monomers for bio-based polymers, *Green Chem.*, 2024, **26**, 8894–8941.
- 14 J. Ye, S. Ma, B. Wang, Q. Chen, K. Huang, X. Xu, Q. Li, S. Wang, N. Lu and J. Zhu, High-performance bio-based epoxies from ferulic acid and furfuryl alcohol: synthesis and properties, *Green Chem.*, 2021, **23**, 1772–1781.
- 15 D. C. Odiy, T. Sharif, R. S. Choudhry, S. Mallik and S. Z. H. Shah, A review of advancements in synthesis, manufacturing and properties of environment friendly bio-based Polyfurfuryl Alcohol Resin and its Composites, *Composites, Part B*, 2023, **267**, 111034.
- 16 A. O. C. Iroegbu and S. S. Ray, On the chemistry of furfuryl alcohol polymerization: A review, *J. Polym. Sci.*, 2024, **62**, 1044–1060.
- 17 H. A. S. Valentino, P. de Tarso Laia dos Reis e Silva Pupi, A. Gandini and T. M. Lacerda, Furfuryl alcohol/tung oil matrix-based composites reinforced with bacterial cellulose fibres, *Cellulose*, 2021, **28**, 7109–7121.
- 18 Y. Jiang, S. Yang and X. Pan, Biomass-Based Polyureas Derived from Rigid Furfurylamine and Isomannide, *ACS Appl. Polym. Mater.*, 2022, **4**, 2197–2204.
- 19 M.-L. Yang, Y.-X. Wu, Y. Liu, J.-J. Qiu and C.-M. Liu, A novel bio-based AB<sub>2</sub> monomer for preparing hyperbranched polyamides derived from levulinic acid and furfurylamine, *Polym. Chem.*, 2019, **10**, 6217–6226.

- 20 H.-H. Shu, Y. Liu, S.-L. Han, X.-Q. Fang, C. Wang and C.-M. Liu, A novel amine-first strategy suitable for preparing both functional and engineering bio-polyamides: furfurylamine as the sole furan source for bisfuranic diamine/diacid monomers, *Polym. Chem.*, 2024, **15**, 4433–4446.
- 21 E. Trovatti, I. G. Gonçalves, A. J. Carvalho and A. Gandini, The contribution of bisfurfurylamine to the development and properties of polyureas, *Polym. Int.*, 2020, **69**, 688–692.
- 22 Z. Cao and X. Zuo, Bio-Based Self-Healing Polymeric Materials Derived from Furfuryl Alcohol Based on the Diels-Alder Reversible Reaction, *Polym. Sci., Ser. B*, 2023, **65**, 450–456.
- 23 S. Raza, J. Zhang, M. Raza, X. Li, H. Wen and C. Liu, Biomass furfural-derived green polymer microspheres: Synthesis and applications for the removal of environmental pollutants from wastewater, *Microporous Mesoporous Mater.*, 2021, **318**, 110966.
- 24 R. Techie-Menson, C. K. Rono, A. Etale, G. Mehlana, J. Darkwa and B. C. E. Makhubela, New bio-based sustainable polymers and polymer composites based on methacrylate derivatives of furfural, solketal and lactic acid, *Mater. Today Commun.*, 2021, **28**, 102721.
- 25 C. Post, D. Maniar, V. S. D. Voet, R. Folkersma and K. Loos, Biobased 2,5-Bis(hydroxymethyl)furan as a Versatile Building Block for Sustainable Polymeric Materials, *ACS Omega*, 2023, **8**, 8991–9003.
- 26 D. Maniar, Y. Jiang, A. J. J. Woortman, J. van Dijken and K. Loos, Furan-Based Copolyesters from Renewable Resources: Enzymatic Synthesis and Properties., *ChemSusChem*, 2019, **12**, 990–999.
- 27 Y. Jiang, A. J. J. Woortman, G. O. R. Alberda van Ekenstein, D. M. Petrović and K. Loos, Enzymatic Synthesis of Biobased Polyesters Using 2,5-Bis(hydroxymethyl)furan as the Building Block, *Biomacromolecules*, 2014, **15**, 2482–2493.
- 28 C. Post, D. Maniar, J. A. Jongstra, D. Parisi, V. S. D. Voet, R. Folkersma and K. Loos, Enzymatic bulk synthesis, characterization, rheology, and biodegradability of bio-based 2,5-bis(hydroxymethyl)furan polyesters, *Green Chem.*, 2024, **26**, 8744–8757.
- 29 S. Thiyagarajan, W. Vogelzang, R. J. I. Knoop, A. E. Frissen, J. van Havenen and D. S. van Es, Biobased furandicarboxylic acids (FDCA): effects of isomeric substitution on polyester synthesis and properties, *Green Chem.*, 2014, **16**, 1957–1966.
- 30 S. Singhal, S. Agarwal, M. P. Mudoi, N. Singhal and R. Singh, Chemical conversion of furan dicarboxylic acid to environmentally benign polyesters: an overview, *Biomass Convers. Biorefin.*, 2023, **13**, 15619–15636.
- 31 M. R. Miah, Y. Dong, J. Wang and J. Zhu, Recent Progress on Sustainable 2,5-Furandicarboxylate-Based Polyesters: Properties and Applications, American Chemical Society, 2024, preprint, DOI: [10.1021/acssuschemeng.3c06878](https://doi.org/10.1021/acssuschemeng.3c06878).
- 32 M. Kamran, M. G. Davidson, S. de Vos, V. Tsanaktis and B. Yeniad, Synthesis and characterisation of polyamides based on 2,5-furandicarboxylic acid as a sustainable building block for engineering plastics, *Polym. Chem.*, 2022, **13**, 3433–3443.
- 33 Y. Jiang, D. Maniar, A. J. J. Woortman and K. Loos, Enzymatic synthesis of 2,5-furandicarboxylic acid-based semi-aromatic polyamides: enzymatic polymerization kinetics, effect of diamine chain length and thermal properties, *RSC Adv.*, 2016, **6**, 67941–67953.
- 34 Y. Jiang, A. J. J. Woortman, G. O. R. Alberda van Ekenstein and K. Loos, A biocatalytic approach towards sustainable furanic-aliphatic polyesters, *Polym. Chem.*, 2015, **6**, 5198–5211.
- 35 K. Loos, R. Zhang, I. Pereira, B. Agostinho, H. Hu, D. Maniar, N. Sbirrazzuoli, A. J. D. Silvestre, N. Guigo and A. F. Sousa, A Perspective on PEF Synthesis, Properties, and End-Life, *Front. Chem.*, 2020, **8**, 585.
- 36 Z. D. Wang, Q. Elliott, Z. Wang, R. A. Setien, J. Puttkammer, A. Ugrinov, J. Lee, D. C. Webster and Q. R. Chu, Furfural-Derived Diacid Prepared by Photoreaction for Sustainable Materials Synthesis, *ACS Sustainable Chem. Eng.*, 2018, **6**, 8136–8141.
- 37 B. B. Yagci, B. Munir, Y. Zorlu and Y. E. Türkmen, Access to Symmetrical and Unsymmetrical Cyclobutanes via Template-Directed [2 + 2]-Photodimerization Reactions of Cinnamic Acids, *Synthesis*, 2023, 3777–3792.
- 38 C. Body, G. Wery, L. Gubbels, K. Robeyns and T. Leyssens, Control of Regioselectivity in the Dimerization of trans-Cinnamic Acid and Its Derivatives Using Cocystal Engineering, *Cryst. Growth Des.*, 2024, **24**, 2117–2125.
- 39 M. Lahav and G. M. J. Schmidt, Topochemistry. Part XVIII. The solid-state photochemistry of some heterocyclic analogues of trans-cinnamic acid, *J. Chem. Soc. B*, 1967, 239–243.
- 40 V. Baret, A. Gandini and E. Rousset, Photodimerization of heteroarylene-vinylenes, *J. Photochem. Photobiol., A*, 1997, **103**, 169–175.
- 41 M. R. van der Kolk, M. A. C. H. Janssen, F. P. J. T. Rutjes and D. Blanco-Ania, Cyclobutanes in Small-Molecule Drug Candidates, John Wiley and Sons Ltd, 2022, preprint, DOI: [10.1002/cmhc.202200020](https://doi.org/10.1002/cmhc.202200020).
- 42 G. K. Kole and M. H. Mir, Isolation of elusive cyclobutane ligands via a template-assisted photochemical [2 + 2] cycloaddition reaction and their utility in engineering crystalline solids, Royal Society of Chemistry, 2022, preprint, DOI: [10.1039/d2ce00277a](https://doi.org/10.1039/d2ce00277a).
- 43 J. Zhang, E. Xiu-Tian-Feng, B. Yang, X. Ma, X. Wang, N. Chang, Q. Zhang, J. J. Zou and J. Xie, Synthesis of Highly Strained Cyclobutane Fuels via Room-Temperature-Integrated Oxidation/[2 + 2] Cycloaddition of Lignocellulose-Based Cyclic Ketones and Cyclic Alcohols, *ACS Sustainable Chem. Eng.*, 2024, **12**(49), 17647–17655.
- 44 E. Lee-Ruff and G. Mladenova, Enantiomerically pure cyclobutane derivatives and their use in organic synthesis, *Chem. Rev.*, 2003, **103**, 1449–1483.
- 45 A. Sokolova, A. Pavlova, N. Komarova, O. Ardashov, A. Shernyukov, Y. Gatilov, O. Yarovaya, T. Tolstikova and N. Salakhutdinov, Synthesis and analgesic activity of new  $\alpha$ -truxillic acid derivatives with monoterpenoid fragments, *Med. Chem. Res.*, 2016, **25**, 1608–1615.
- 46 Z. Wang, M. Scheuring, M. Mabin, R. Shahni, Z. D. Wang, A. Ugrinov, J. Butz and Q. R. Chu, Renewable Cyclobutane-

- 1,3-dicarboxylic Acid (CBDA) Building Block Synthesized from Furfural via Photocyclization, *ACS Sustainable Chem. Eng.*, 2020, **8**, 8909–8917.
- 47 J. Tellers, N. Sbirrazzuoli and N. Guigo, A rigid plant oil-based thermoset with a furfural-derived cyclobutane cross-linker, *Green Chem.*, 2021, **23**, 8053–8060.
- 48 M. Jamali Moghadam Siahkali, N. Guigo and N. Sbirrazzuoli, Cross-Linking Mechanisms of a Rigid Plant Oil-Based Thermoset from Furfural-Derived Cyclobutane, *Macromolecules*, 2023, **56**, 404–415.
- 49 H. Amjaour, Z. Wang, M. Mabin, J. Puttkammer, S. Busch and Q. R. Chu, Scalable preparation and property investigation of a cis-cyclobutane-1,2-dicarboxylic acid from  $\beta$ -trans-cinnamic acid, *Chem. Commun.*, 2019, **55**, 214–217.
- 50 B. Jin, H. Song, R. Jiang, J. Song, Q. Zhao and T. Xie, Programming a crystalline shape memory polymer network with thermo- and photo-reversible bonds toward a single-component soft robot, *Sci. Adv.*, 2018, **4**(1), 3865.
- 51 D. Tunc, C. Le Coz, M. Alexandre, P. Desbois, P. Lecomte and S. Carloti, Reversible Cross-Linking of Aliphatic Polyamides Bearing Thermo- and Photoresponsive Cinnamoyl Moieties, *Macromolecules*, 2014, **47**, 8247–8254.
- 52 Y. Lin, T. B. Kouznetsova and S. L. Craig, Mechanically Gated Degradable Polymers, *J. Am. Chem. Soc.*, 2020, **142**, 2105–2109.
- 53 G. Kaur, P. Johnston and K. Saito, Photo-reversible dimerisation reactions and their applications in polymeric systems, *Polym. Chem.*, 2014, **5**, 2171–2186.
- 54 A. Rahimi and J. M. García, Chemical recycling of waste plastics for new materials production, *Nat. Rev. Chem.*, 2017, **1**, 0046.
- 55 R. Hatti-Kaul, L. J. Nilsson, B. Zhang, N. Rehnberg and S. Lundmark, *Designing Biobased Recyclable Polymers for Plastics*, Elsevier Ltd, 2020, **38**, (1), pp. 50–67, DOI: [10.1016/j.tibtech.2019.04.011](https://doi.org/10.1016/j.tibtech.2019.04.011).
- 56 N. Singh and T. R. Walker, Plastic recycling: A panacea or environmental pollution problem, *npj Mater. Sustain.*, 2024, **2**(1), 17.
- 57 S. Kumar, S. Kumar, S. Kumar, T. Yadav, P. S. Dhapola and P. K. Singh, Reducing Environmental Plastic Pollution by Designing Polymer Materials for Managed End-of-Life, *Macromol. Symp.*, 2024, **413**(1), 2300146.
- 58 M. Martín-Sómer, C. Pablos, C. Adán, R. van Grieken and J. Marugán, A review on LED technology in water photodisinfection, *Sci. Total Environ.*, 2023, **885**, 163963.
- 59 R. Pizzichetti, M. Martín-Gamboa, C. Pablos, K. Reynolds, S. Stanley, J. Dufour and J. Marugán, Environmental life cycle assessment of UV-C LEDs vs. mercury lamps and oxidant selection for diclofenac degradation, *Sustainable Mater. Technol.*, 2024, **41**, e01002.
- 60 J. J. Rytuba, Mercury from mineral deposits and potential environmental impact, *Environ. Geol.*, 2003, **43**, 326–338.
- 61 Z. Wang, M. Scheuring, M. Mabin, R. Shahni, Z. D. Wang, A. Ugrinov, J. Butz and Q. R. Chu, Renewable Cyclobutane-1,3-dicarboxylic Acid (CBDA) Building Block Synthesized from Furfural via Photocyclization, *ACS Sustainable Chem. Eng.*, 2020, **8**, 8909–8917.
- 62 R. K. Shahni, M. Mabin, Z. Wang, M. Shaik, A. Ugrinov and Q. R. Chu, Synthesis and characterization of BPA-free polyesters by incorporating a semi-rigid cyclobutanediol monomer, *Polym. Chem.*, 2020, **11**, 6081–6090.
- 63 S. Zaidi, S. Thiyagarajan, A. Bougarech, F. Sebti, S. Abid, A. Majdi, A. J. D. Silvestre and A. F. Sousa, Highly transparent films of new copolyesters derived from terephthalic and 2,4-furandicarboxylic acids, *Polym. Chem.*, 2019, **10**, 5324–5332.
- 64 H. Azim, A. Dekhterman, Z. Jiang and R. A. Gross, Candida antarctica Lipase B-Catalyzed Synthesis of Poly(butylene succinate): Shorter Chain Building Blocks Also Work, *Biomacromolecules*, 2006, **7**, 3093–3097.
- 65 F. Hollmann, P. Grzebyk, V. Heinrichs, K. Doderer and O. Thum, On the inactivity of Candida antarctica lipase B towards strong acids, *J. Mol. Catal. B: Enzym.*, 2009, **57**, 257–261.
- 66 Y. L. de los Santos, Y. L. Chew-Fajardo, G. Brault and N. Doucet, Dissecting the evolvability landscape of the CalB active site toward aromatic substrates, *Sci. Rep.*, 2019, **9**(1), 15588.
- 67 P. B. Juhl, K. Doderer, F. Hollmann, O. Thum and J. Pleiss, Engineering of Candida antarctica lipase B for hydrolysis of bulky carboxylic acid esters, *J. Biotechnol.*, 2010, **150**, 474–480.
- 68 J. U. Izunobi and C. L. Higginbotham, Polymer molecular weight analysis by  $^1\text{H}$  NMR spectroscopy, *J. Chem. Educ.*, 2011, **88**, 1098–1104.
- 69 A. Gandini and M. N. Belgacem, in *Monomers, Polymers and Composites from Renewable Resources*, Elsevier, 2008, pp. 115–152.
- 70 L. I. Belen'kii, T. G. Kim, I. A. Suslov and N. D. Chuvylkin, Substrate and positional selectivity in electrophilic substitution reactions in pyrrole, furan, thiophene, and selenophene derivatives and related benzoannelated systems, *Russ. Chem. Bull.*, 2005, **54**, 853–863.