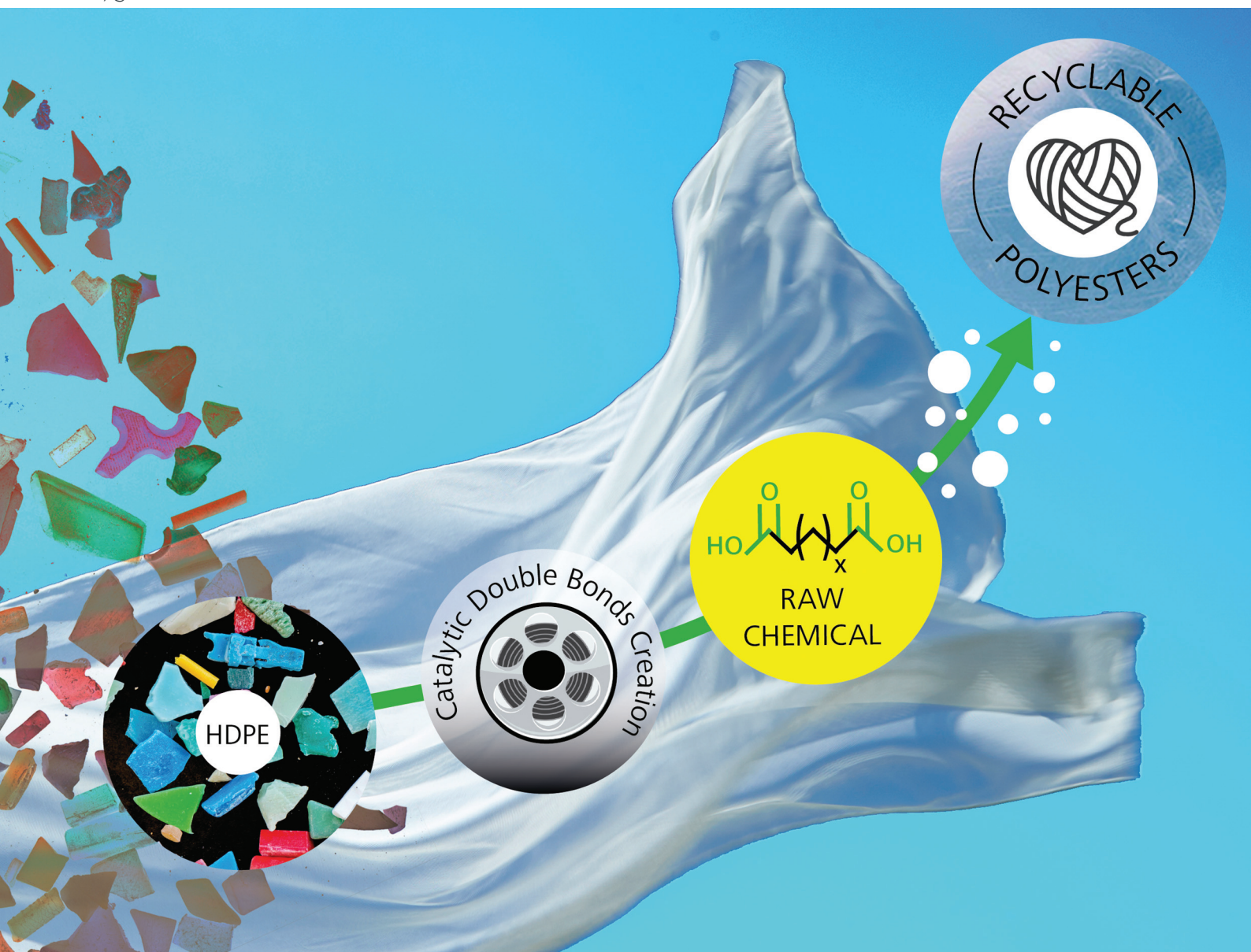


Green Chemistry

Cutting-edge research for a greener sustainable future

rsc.li/greenchem



ISSN 1463-9262

PAPER

Wenyu Wu Klingler *et al.*

A facile methodology for side- and upcycling of HDPE waste
via partial creation of unsaturated double bonds



Cite this: *Green Chem.*, 2024, **26**, 10422

A facile methodology for side- and upcycling of HDPE waste *via* partial creation of unsaturated double bonds†

Wenyu Wu Klingler,^a Lucie Perret,^a Patrick Rupper,^a Sandro Lehner,^a Xiaoyu Zhou,^b Henrik Eliasson,^c Rico Muff,^d Manfred Heuberger^{a,e} and Sabyasachi Gaan^a

Upcycling is emerging as a crucial strategy for enhancing the value of polymers, driving the transition toward a circular material economy. In this study, we present a facile chemical method for converting high-density polyethylene (HDPE) waste into valuable long-chain dicarboxylic acids (LDCAs) *via* a key enabling unsaturated double-bond formation step. That is, we propose introducing unsaturation points in HDPE using commercially available heterogeneous catalysts (e.g. Pt/Al₂O₃). The unsaturation level through C=C double bond formation is around 10%, as confirmed by FTIR, Raman and X-ray photoelectron spectroscopy. A process of microwave-assisted oxidation is demonstrated to break down the dehydrogenated HDPE into a mixture of aliphatic diacids. This approach enhances the recyclability and value of HDPE by the transformation of polymer waste into bifunctional monomers for potentially novel polyester synthesis. This process offers a sustainable and value-added alternative to conventional recycling methods. While further optimizations are needed, initial estimates of the *E*-factor and sEF provide promising indicators of the potential environmental benefits of this upcycling approach.

Received 26th June 2024,
Accepted 22nd August 2024

DOI: 10.1039/d4gc03108c

rsc.li/greenchem

Introduction

In the past century, petroleum-based plastics have undergone remarkable economic growth. However, their beneficial stability turns into a disadvantage once disposed as waste in the environment with alarming consequences.^{1,2} The growing visibility of climate change emphasizes the urgent necessity for interventions, particularly in waste management and circular material design, two crucial areas with substantial potential to mitigate these impacts.³ For converting polymer waste into valuable new materials on a large scale, complex and energy-intensive procedures are envisioned; along these lines efficient

catalytic approaches have been established.⁴ By 2020 only 14% of common polymers (8.3 billion metric tons have been produced since the 1950s), like polyethylene and polyesters, were mechanically recycled, mainly resulting in lower-value products that end up as depreciated waste after one more life cycle.^{5,6} Polyolefins in general and, in particular high-density polyethylene (HDPE, 50 million tons per year), are major constituents of solid waste where only about 30% of HDPE is mechanically recycled (Fig. 1a). This general downcycling is limited by the significant degradation of quality after a few cycles. Developing new, high-quality side-/upcycling methods for HDPE is thus a crucial strategy in a sustainable and value-added circular economy.

Catalysts play a significantly important role in side-/upcycling and even in pyrolysis methods (Fig. 1a).^{7,8} For side-cycling, the creation of certain “functional” points along the inert PE chain is a key step for repurposing the material (Fig. 1b).^{9,10} Upcycling, which transforms HDPE into lower-molecular-weight products, also offers a promising alternative to mechanical recycling.^{8,9,11,12} One such widely studied approach is ‘pyrolysis’, in which plastics are decomposed to mixed oil or wax at high temperatures (>400 °C), often assisted by the presence of a catalyst.¹³ However, efficient and economical chemical processes for HDPE-to-monomer depolymerization are still in their early stages.¹⁴ Great efforts have been

^aLaboratory for Advanced Fibers, Empa, Swiss Federal Laboratories for Materials Science and Technology, Lerchenfeldstrasse 5, 9014 St Gallen, Switzerland.
E-mail: wenyu.wu@empa.ch

^bDepartment of Chemistry and Applied Biosciences, ETH Zürich, Vladimir Prelog Weg 1–5, 8093 Zürich, Switzerland

^cElectron Microscopy Center, Empa, Swiss Federal Laboratories for Materials Science and Technology, Überlandstrasse 129, 8600 Dübendorf, Switzerland

^dTransport at Nanoscale Interfaces Laboratory, Empa, Swiss Federal Laboratories for Materials Science and Technology, Überlandstrasse 129, 8600 Dübendorf, Switzerland

^eDepartment of Materials, ETH Zurich, 8092 Zurich, Switzerland

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4gc03108c>



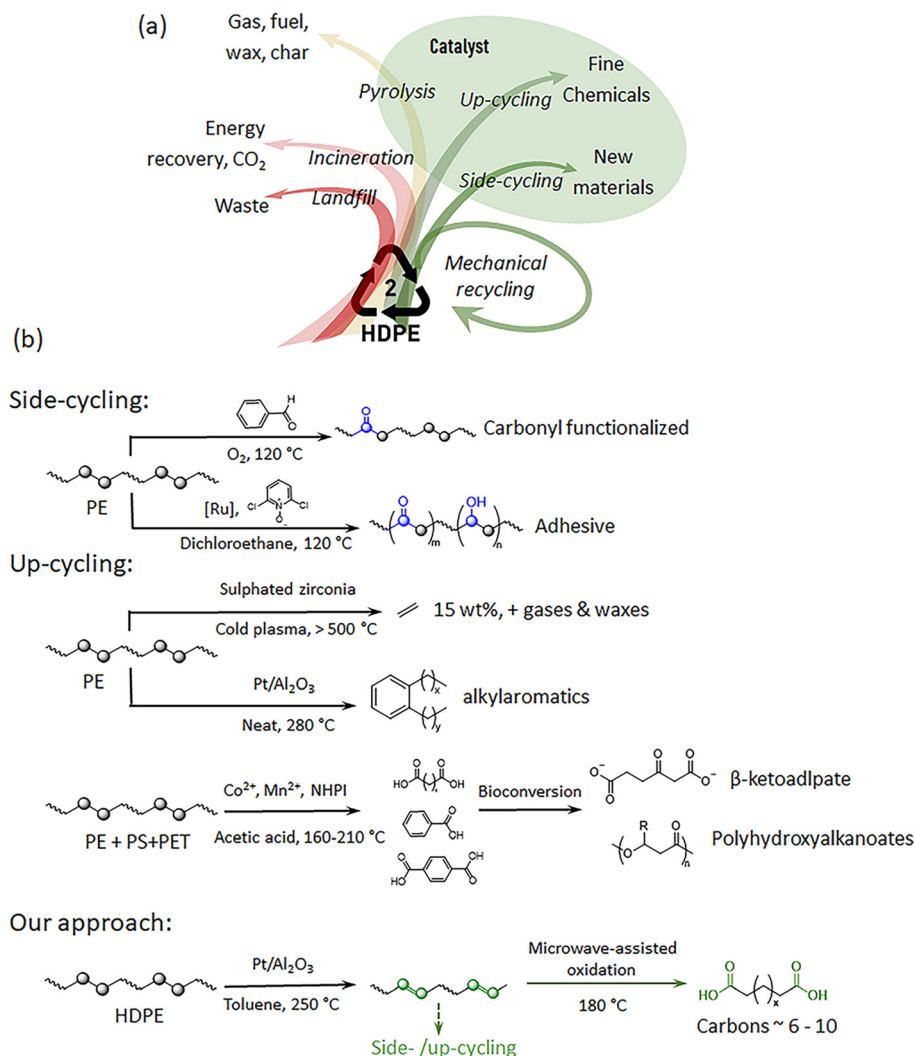


Fig. 1 (a) The general post-consumption flow of HDPE, and (b) selected existing side-/upcycling methods for PE treatment, including oxidation,⁹ catalytic functionalization,¹⁰ plasma-assisted cracking,⁴⁰ tandem hydrogenolysis/aromatization,²² chemical–biological upcycling of mixed polymers,²⁹ fragmentation–oxidation upcycling process,²⁵ and lastly our dehydrogenation and microwave-assisted oxidation approach, for obtaining aliphatic bifunctional monomers.

devoted to energy-efficient catalytic depolymerization methods to recycle HDPE to value-added products (*inter alia* surfactants, functional molecules).^{7,15–23} For example, the market price of surfactants (\$2.3–\$10 per kg) is significantly higher than that of virgin PE (\$1.2 per kg).²³ The technical capability to upconvert polymeric materials could act as a long-term strategy towards reaching sustainable cycling, where chemical feedstocks are repeatedly polymerized and after their life phase depolymerized, refined and re-polymerized to useful new products. Notably, the possibility to increase the economic value *via* re-polymerization into new functional materials is increasingly recognized as a central asset allowing evolution in a recycling scenario.^{24,25}

Innovative approaches by introducing and adjusting the HDPE unsaturation points as cleavable linkages showed great potential to introduce functional upgrades into the polymer chemical structure.^{8,26,27} Further breakdown of the C=C bonds

by oxidation, metathesis, and other processes could result in telechelic polymers.^{8,26,28} Recently, a dedicated co-catalytic system, based on tandem dehydrogenation–metathesis reactions at low temperature (~150 °C), was applied to depolymerize polyethylene (PE).¹⁸ The depolymerization relies on the selectivity of one catalyst to partially dehydrogenate PE, forming internal double bonds, while an olefin metathesis catalyst cleaves the polymer into smaller segments at the double bonds. Co(II)/Mn(II)-catalyzed oxidation supported by an *N*-hydroxyphthalimide initiator, for the production of oxygenated small molecules that are advantaged substrates for biological conversion, has also gained increased prominence more recently.²⁹ A similar method *via* catalytic oxidation and bioconversion has demonstrated the valorization of HDPE waste first to dicarboxylic acids (DCAs) and then to pharmacologically active compounds.³⁰

Long-chain aliphatic diacids, diols, and diamines are essential for creating polymers that fill the niche between poly-



olefins and traditional polycondensates.^{28,31–33} Specifically, long-chain dicarboxylic acids (LCDAs) with chain lengths of 10 or more, such as sebacic and dodecanedioic acids, are crucial for synthesizing nylons and other polycondensates, providing benefits like high modulus and heat resistance due to their ability to crystallize.³⁴ These materials, resembling polyethylene, offer potential for degradability in various applications.^{28,35–37} Currently, LCDAs are produced through chemical synthesis and biotechnological methods, which are often unsustainable or costly.³¹ There is a significant need for more efficient, economical, and sustainable production methods, potentially through side- and upcycling of polymer waste, to access these monomers more effectively.^{8,35,38}

In accordance with these concepts, we developed a two-step process that enables production of a valuable new class of long-chain aliphatic monomers *via* catalytic depolymerization of HDPE waste products (Fig. 1b, our approach). By partial creation of unsaturated double bonds, we can enable a range of new possibilities for side- and upcycling. The proposed procedure requires a sequence of dehydrogenation and oxidation steps. Dehydrogenation of HDPE creates reactive points (unsaturated double bonds, 5–10%) along the otherwise inert polymer backbone using industrially available catalysts. A platinum catalyst was employed, inspired by its established use in the dehydrogenation of alkanes, due to its high catalytic activity and stability, *e.g.* in the Oleflex technologies of UOP.³⁹ In a second step, microwave-assisted oxidation of the unsaturated bonds can directly lead to carboxylic acid-terminated long-chain monomers, which could be re-polymerized to long chain polycondensates. This work has resulted in the development of a new chemical methodology for (partial) depolymerization of HDPE into long-chain aliphatic monomers, which not only has potential to minimize polymer waste but also adds further value supporting the economics and technical evolution of a circular economy. It contributes to providing diverse solutions for chemical recycling; however, some processes within the technology still require further optimization.

Results and discussion

Characterization of catalysts

In order to obtain dehydrogenated HDPE rather than cracking products, selective C–H activation is needed, yet this remains a very challenging process. The cleavage of C–C bonds is more easily activated, since the average bond energy of the C–C bond (347 kJ mol^{−1}) is weaker than that of the C–H bond (414 kJ mol^{−1}).⁴¹ Pt is the well-utilized noble metal for the direct dehydrogenation of light alkanes, owing to its facile C–H bond activation over C–C bond scission.⁴¹ Al₂O₃ is a stable carrier of Pt catalysts, in which the coordinatively unsaturated pentacoordinate Al³⁺ sites on the γ -Al₂O₃ (100) surface can anchor Pt to make it resistant to migration, such as in the Oleflex process developed by Honeywell UOP.⁴² Therefore, a commercially available Pt/Al₂O₃ catalyst was selected deliberately to demonstrate the feasibility and potential scalability of

this work (Fig. 2a). The catalyst was firstly characterized using various methods before its use. The surface area of the Pt/Al₂O₃ catalyst was determined to be 84 m² g^{−1} based on the nitrogen adsorption–desorption isotherm at 77 K using the Brunauer–Emmett–Teller (BET) theory (Fig. S2a and S2b†).⁴³ This is smaller than that of commonly reported alumina matrix, which could be due to the wet impregnation methods used.⁴⁴ Fig. S2c† gives the measured amounts of total and strongly chemisorbed hydrogen (H₂) at 298 K. Strongly chemisorbed H₂ was determined from the amount that did not desorb during evacuation at the analysis temperature. It was measured and found to be 0.529 mL g^{−1}, which indicates dispersion of roughly 93% for the supported Pt nanoparticles. The total H/Pt ratios of Pt/Al₂O₃ at room temperature was close to one, which is often observed for small metal clusters on a supporting matrix.^{45–47}

The XRD profiles of the alumina-loaded catalysts showed the characteristic diffraction peaks of the Al₂O₃ support reported at $2\theta = 38.5^\circ$, 45.8° and 67.0° (Fig. S3†). Although the introduction of platinum usually induces the appearance of a small peak of metallic platinum, it is not tangible in the present case, indicating the well dispersed Pt on the matrix. It is also a sign that Pt particles should be very small. Consistent with BET and XRD analyses, the TEM and HAADF-STEM images of untreated catalyst show less dense and mesoporous Al₂O₃ with Pt nanoparticles around 1.0–1.5 nm (Fig. 2b and Fig. S4a–c†). The hydrogen activation of the catalyst introduced certain amounts of aggregation, yet sizes of Pt nanoparticles still remain at around 2 nm (Fig. S4d–f†). The porous nature of the matrix and the low amount of catalyst loading can also be confirmed by SEM and EDX (Fig. S5†). Furthermore, HDPE is a very viscous liquid when molten, which is a critical property that affects its processability for end-of-life treatment. To reduce the viscosity of the reaction mixture and to facilitate the dehydrogenation process by loosening the polymer chains, toluene was used as solvent for improving the homogeneity of the reaction and eventually achieving better conversion. After the reaction, the catalyst was separated and the solvent was recovered using rotavapor equipment for reuse.

Characterization of the dehydrogenated HDPE

Following the toluene-assisted dehydrogenation process, HDPE was treated using the hydrogen-activated Pt/Al₂O₃ catalyst. This reaction was conducted at 250 °C for varying durations of 36, 48, and 72 hours. Upon completion of the respective reaction times, the treated HDPEs, designated as **HDPE-36**, **HDPE-48**, and **HDPE-72**, were collected and dried for subsequent characterization. The structural simplicity of HDPE, with its repetition of CH₂ units, does not reflect the complexity and properties of the real material. Ideally, the dehydrogenation treatment of HDPE should introduce unsaturation while maintaining the polymer's molecular weight; chain scission is expected to be negligible at this point. This assertion is supported by the analysis of number-averaged molecular weight (M_n), weight-averaged molecular weight (M_w), and the polydispersity ($PD = M_w/M_n$) values, which were



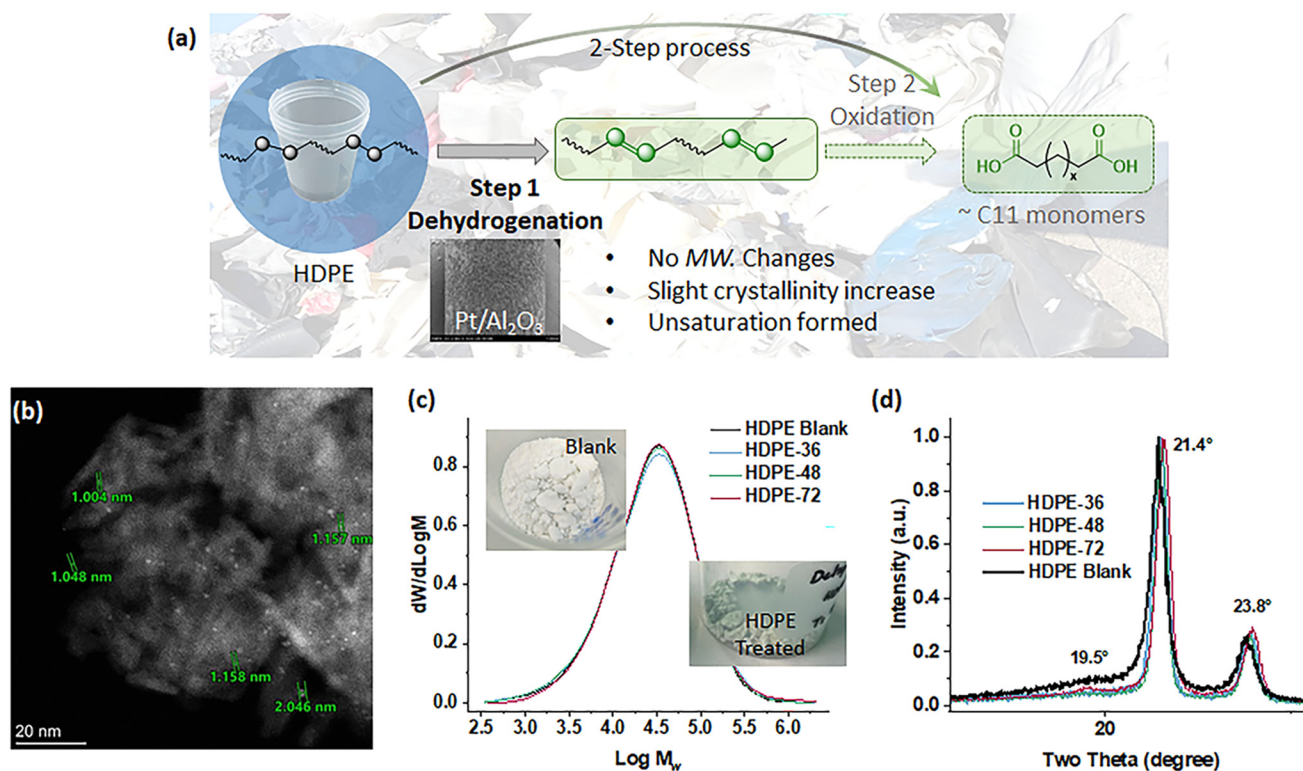


Fig. 2 Upcycling of HDPE via a two-step process: (a) schematic flow with the first dehydrogenation step using 1 wt% Pt/Al₂O₃ in dry toluene and treated for 36–72 hours at 250 °C under N₂; (b) STEM micrograph of the catalyst (Pt/Al₂O₃) used; (c) GPC traces of the HDPEs with images before and after treatment; and (d) XRD patterns of HDPEs with enlarged area of the spectra between 16° and 26°.

quantified using high-temperature gel permeation chromatography (GPC), with results as depicted in Fig. 2c. Consistency in signal profiles across treated and untreated samples was meticulously noted, indicating no significant molecular weight variation or degradation; M_n values ranged approximately from 13k to 15k, M_w from 50k to 55k, and PD from 3.6 to 4.0, as catalogued in Table S1.† The chromatographic findings suggest that the integrity of the polymer chains remained unaffected by the catalytic treatment, as demonstrated by the alignment of retention times, peak shapes, and intensities with those of the control sample.

HDPE exists generally as a mixture of crystalline and amorphous phases. One of the important questions that came up was whether the HDPE morphology changed after relatively harsh dehydrogenation reaction conditions, especially the branching and *trans-cis* properties.^{48,49} In principle, melting points of alkenes are similar to those of alkanes. However, since *trans*-configured alkenes are more symmetrical and hence less polar, they have normally higher melting points than those of *cis*-alkenes. Additionally, alkenes display weak dipole-dipole interactions due to the electron-attracting sp² carbon, which might influence their melting points. Differential scanning calorimetry (DSC) analyses (Fig. S6†) further elucidate the polymer's thermal response (*i.e.* melting and crystallization) to dehydrogenation treatment, using the cooling and second heating curves. HDPE, characterized by

minimal to no chain branching, exhibits a highly crystalline structure, reflected in the narrow range of endothermic peaks between 115 and 142 °C during melting. The similarity in melting temperature (T_m) and crystallization temperature (T_c) values, and the slightly increased crystallinity (X_c) calculated by melting heat of fusion (ΔH_m) values before and after treatment, as documented in Table S1,† imply negligible alterations in the thermal characteristics following the treatment. This preservation of thermal behavior, which is likely attributable to the maintenance of lamellar thickness within the polymer crystals, is indicative of the treatment's efficacy in retaining the polymer's structural integrity.⁵⁰ Thermogravimetric analysis (TGA, Fig. S7†) corroborates the thermal stability of HDPE, further reinforcing the conclusion that the dehydrogenation process compromises neither the polymer's molecular weight nor its thermal properties, thereby maintaining its intrinsic material complexity.

In addition, XRD is used to confirm the crystallinity of the untreated HDPE blank and its change after dehydrogenation (Fig. S8†). Two sharp crystal diffraction peaks can be observed at ~21.4° and 23.8°, which correspond to diffraction from the (110) and (200) planes of the orthorhombic crystal structure in the HDPEs, respectively.⁴⁹ Also, the broad scattering peak lying below the two sharp peaks represents the amorphous portion. The crystalline phase generally increases with increasing molecular weight and decreases with increasing chain branching.



There is a subtle decrease in diffraction intensities of the amorphous portion of treated HDPEs (Fig. 2d), which corresponds to the mild change in the crystallinity of HDPE before and after the dehydrogenation treatment, as observed in the thermal analysis.

Importantly, the formation of unsaturated double bonds is observed both in FTIR and Raman spectra (Fig. 3). The FTIR pattern and absorptions are almost identical to those of the blank HDPE (Fig. 3a), except for the appearance of =CH-stretching at around 3000 cm^{-1} , which is obvious (Fig. 3c). The =CH-bending located at around 890 cm^{-1} is also observable (Fig. 3d), while the C=C stretching peak (*i.e.* that between 1680 and 1630 cm^{-1}) is too subtle to be observed. This may result from the C=C *trans*-configuration. Since it is symmetrically substituted, the C=C stretching is a symmetric stretching of an almost symmetric molecule, similar to the symmetric stretching of CO_2 . In this case the peak intensity is zero. Furthermore, no observable vibrational peaks near 1100 cm^{-1} and 1650 cm^{-1} of the C-O and C=O bonds appear, showing no indication of chain oxidation. Meanwhile, HDPE treated at lower temperature (200°C for 36 and 48 hours) showed no evidence of =CH-stretching or =CH-bending signals (Fig. S9†). The crystallinity can be estimated from the FTIR spectra by using Zerbi's equation,⁵¹ comparing absorbance peaks at 730 cm^{-1} and 720 cm^{-1} with that of the internal reference 2019 cm^{-1} peak. According to Zerbi *et al.*,⁵¹ the bending

vibrations at 1474 and 730 cm^{-1} (crystalline phase) and 1464 and 720 cm^{-1} (amorphous phase) correspond to the frequency behavior of CH_2 symmetric stretching, which is also a function of *trans-cis* conformation (Fig. 3b). The amorphous phase indicated by the peak at 720 cm^{-1} slightly decreases.

Raman spectra for the untreated HDPE blank and dehydrogenated HDPE in the region between 200 and 3500 cm^{-1} showed some divergences. For the blank HDPE, the relevant vibrational bands due to skeletal C-C stretching (1050 – 1150 cm^{-1}), =CH-bending (scissoring between 1245 and 1277 cm^{-1}), CH_2 and CH_3 bending (1400 – 1500 cm^{-1}), C=C stretching (1600 – 1700 cm^{-1}), and C-H stretching vibrations (2800 – 3000 cm^{-1}) are shown in Fig. 3e. For the dehydrogenated HDPE, the Raman spectra possess differences in fingerprint (C=C stretching) regions;⁵² therefore, it confirms the existence of double bonds on the HDPE backbones (Fig. 3f). Visually inspecting the spectra offers a fast initial understanding of unsaturation formation. The total unsaturation can be estimated semi-quantitatively by measuring the ratio of the C=C stretching ($\nu\text{ C=C}$) centered at 1601 cm^{-1} to the CH_2 scissoring deformation ($\delta\text{ CH}_2$) at 1444 cm^{-1} , *via* Gaussian fitting and integration of the identified band areas at 1540 – 1670 cm^{-1} and 1390 – 1510 cm^{-1} (Fig. S10†).⁵³ Calculating band intensity ratios is a direct approach to quantifying total unsaturation, and here we obtained roughly 10% unsaturation yield. Additionally, Raman spectroscopy can offer

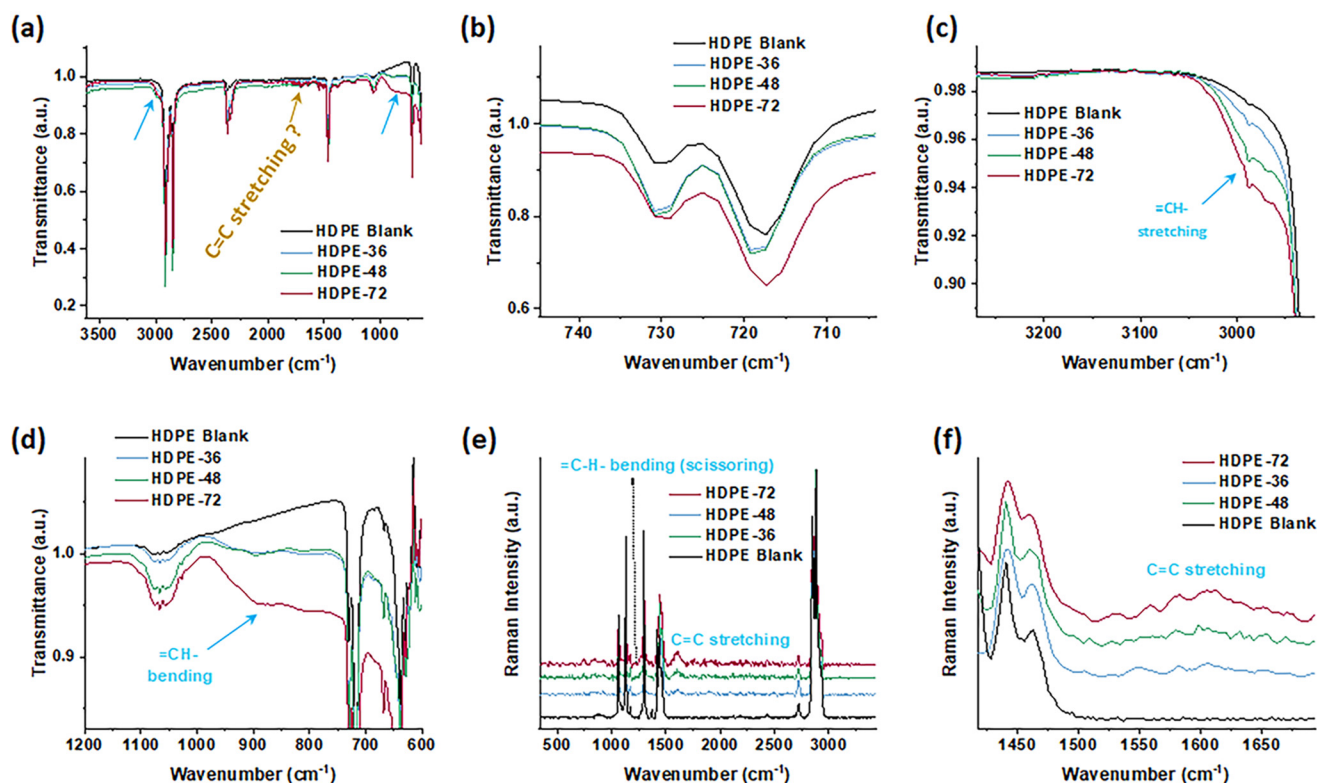


Fig. 3 (a) FTIR spectra of the HDPE before (blank) and after dehydrogenation (HDPE-36, 48, 72), with (b) enlarged area of the CH_2 stretching range, (c) the =CH- stretching range, (d) the =CH- bending geminal disubstituted alkene range, (e) Raman spectra of the HDPEs before and after dehydrogenation treatment, and (f) the enlarged area at the C=C stretching range.



information on the structural characterization of PE. The modified Strobl and Hagedorn method is generally accepted and applied in the investigation of PE morphological structure.⁵⁴ The normalized transmittance of the 1416 cm^{-1} band could be used to estimate the crystalline fraction in HDPEs (Fig. S11†), which indicates a slight increase of the crystallinity after treatment (Table S2†).

Fig. 4a depicts the XPS survey scans for HDPE in its native form as well as that for HDPE-72 with the most double bonds as characterized by Raman spectroscopy. As expected, both contain only carbon (C) in significant amounts. In addition, XPS can further confirm double bond formation by distinguishing carbon atoms at different hybridization states.⁵⁵ Both the C 1s photoelectron peak as well as the X-ray excited C KLL Auger peak can be used to characterize the hybridization fractions of carbon electrons (sp^2 and sp^3).⁵⁶ Due to only a small difference of 0.3–1.0 eV in binding energy from C 1s sp^2 to sp^3 , the C 1s peak only gives a semi-quantitative determination of the sp^2/sp^3 ratio.^{55,57–59} In the C 1s spectrum fitting, we have fixed the two components of C sp^2 and C sp^3 to the ranges of literature values of 284.3–284.6 eV and 284.9–285.2 eV, respectively,^{55,56,58} and restricted the fitted FWHM to be the same in both components.

After treatment, HDPE exhibited a slight binding energy shift towards lower energies (see Fig. 4b). This shift results in an increased part of the sp^2 component, which is localized at lower binding energies, as is observable from Fig. 4c. The C 1s component towards higher binding energy (at around 285.9

eV) represents vibrational fine structure as described for a reference HDPE spectrum.⁵⁸ The sp^2/sp^3 ratio can be more quantitatively evaluated using the shape of the Auger C KLL spectrum (excited by X-ray photons), which represents a fingerprint of the carbon states.⁶⁰ The method consists of determining the distance (the so-called D parameter) between the most positive maximum and the most negative minimum of the first derivative of the C KLL spectrum (see the ESI†).⁶¹ The D parameter is independent of absolute binding energy calibration and hence, the determined sp^2 content from it represents a more quantitative value than the one resulting from the fitting of the C 1s peak with components lying close together in binding energy. Fig. 4d depicts the C KLL Auger spectrum for the two investigated samples and Fig. 4e shows the corresponding determination of the D parameter *via* the first derivative of the C KLL spectrum. The width of the C KLL Auger peak is increased for the HDPE-72 sample. The value of 14.0 eV for the D parameter results in 17.3% of sp^2 for HDPE-72, which is consistent with the value obtained from the C 1s peak (see Table S3†). Therefore, successful partial dehydrogenation of the HDPE and formation of desired double bonds are confirmed using various techniques.

Analysis and application of the oxidized products

After achieving the desired partial unsaturation, microwave-assisted oxidation was conducted to facilitate depolymerization towards the target functional monomers. The efficacy of

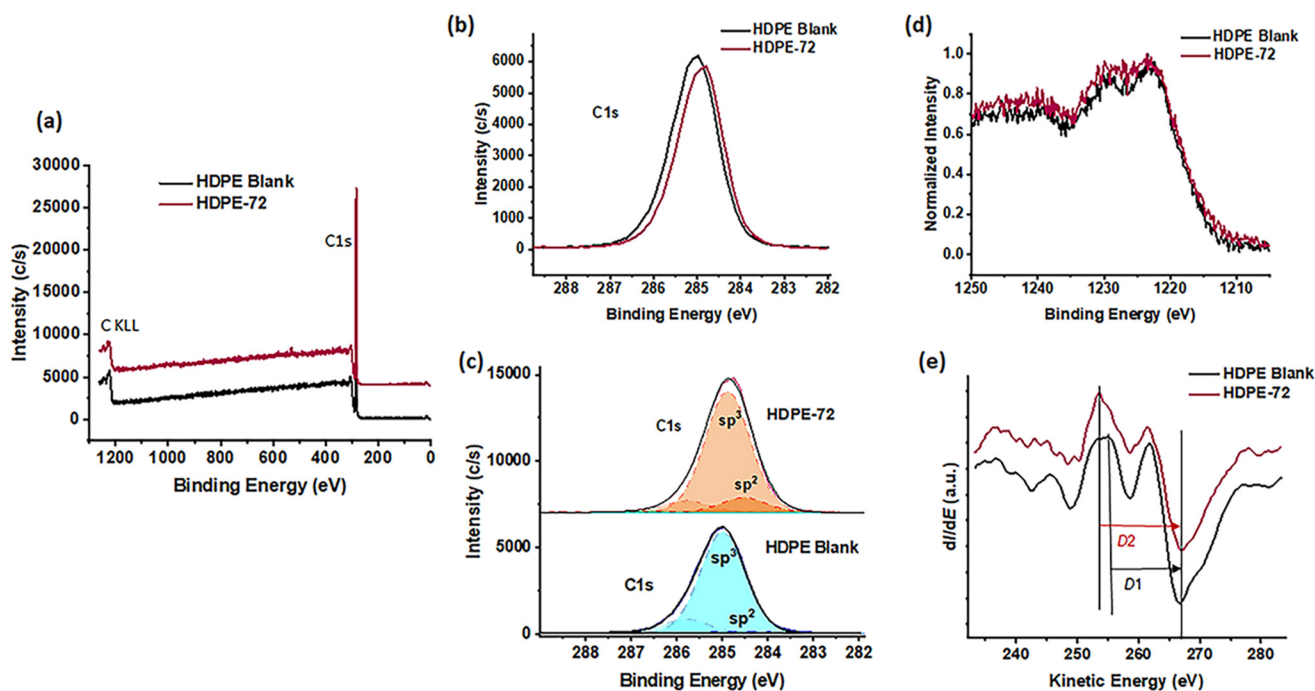


Fig. 4 Comparison of the XPS spectra of the blank HDPE and HDPE-72. In (a), (c) and (e), the two spectra are offset in the y-direction for better visualization. (a) Survey scans; (b) high-resolution scans for C 1s and (c) curve fitting of the envelope into composite peaks of sp^2 , sp^3 and vibrational fine structure (dashed lines); (d) X-ray excited carbon KLL Auger spectra; (e) first derivative of the carbon KLL Auger spectra obtained by numerical differentiation.



strong oxidation catalysts, specifically diluted nitric acid solution (0.10 g mL^{-1} in water), was assessed in promoting the oxidative degradation of the dehydrogenated HDPE (1 g) into a mixture of dicarboxylic acids. The reaction was carried out in a Milestone SyntWAVE microwave reactor for 1 hour at 180°C , and the pressure was kept constant at 60 bar under an N_2 atmosphere. The depolymerized solutions of untreated and dehydrogenated HDPEs were directly analyzed using UPLC-MS, which revealed distinct differences between the products derived from dehydrogenated HDPE and those from the blank HDPE. The blank HDPE primarily underwent surface carbonization with a low concentration of detectable organic fragments (Fig. S12†). HDPE-48 was depolymerized into fragments that are completely water-soluble (Fig. 5). This was further confirmed by UPLC-MS analysis, in which the extracted products *via* total ion chromatography (TIC) and UV spectroscopy channels indicated significant variances in their chemical signatures (Fig. 5b and Fig. S13†). Mass spectrometry (MS) analysis suggested that the primary degradation products were long-chain aliphatic diacids, predominantly those with carbon chain lengths shorter than C13 (Fig. S14†), exhibiting a notable distribution of chain lengths (C9–C13). The distribution of oxidation products is directly influenced by the con-

centration of double bonds in the treated HDPEs. The oxidation solution of HDPE-36 exhibited a broader range of mass and UV signals compared to HDPE-48 (Fig. S15†), likely due to a lower concentration of double bonds within the HDPE chains. Conversely, the oxidation solution of HDPE-72 demonstrated a narrower product distribution though with relatively lower concentration. Given its relatively narrow product distribution and sufficient concentration, HDPE-48 was selected for further analysis. The resulting diacids were collected, further characterized, and repolymerized.

After oxidation of HDPE-48, the crude solution was neutralized using NaOH (1 M) solution for further separation. Chloroform was employed to separate the longer chain DAs from the mixture. Post-separation and drying, 0.84 g of DAs was derived from 1 g of HDPE, resulting in a carbon conversion yield of approximately 58%. In future work, we plan to separate more fragments from the water solution for better recovery of the materials. The confirmation of the isolated LCDAs was further achieved through attenuated total reflection FTIR spectroscopy (Fig. 5c). This analysis highlighted the presence of carboxylic acid groups, as evidenced by characteristic absorption regions indicative of carbonyl ($\text{C}=\text{O}$) and hydroxyl ($-\text{OH}$) groups. The oxidized products, both pre- and post-isolation,

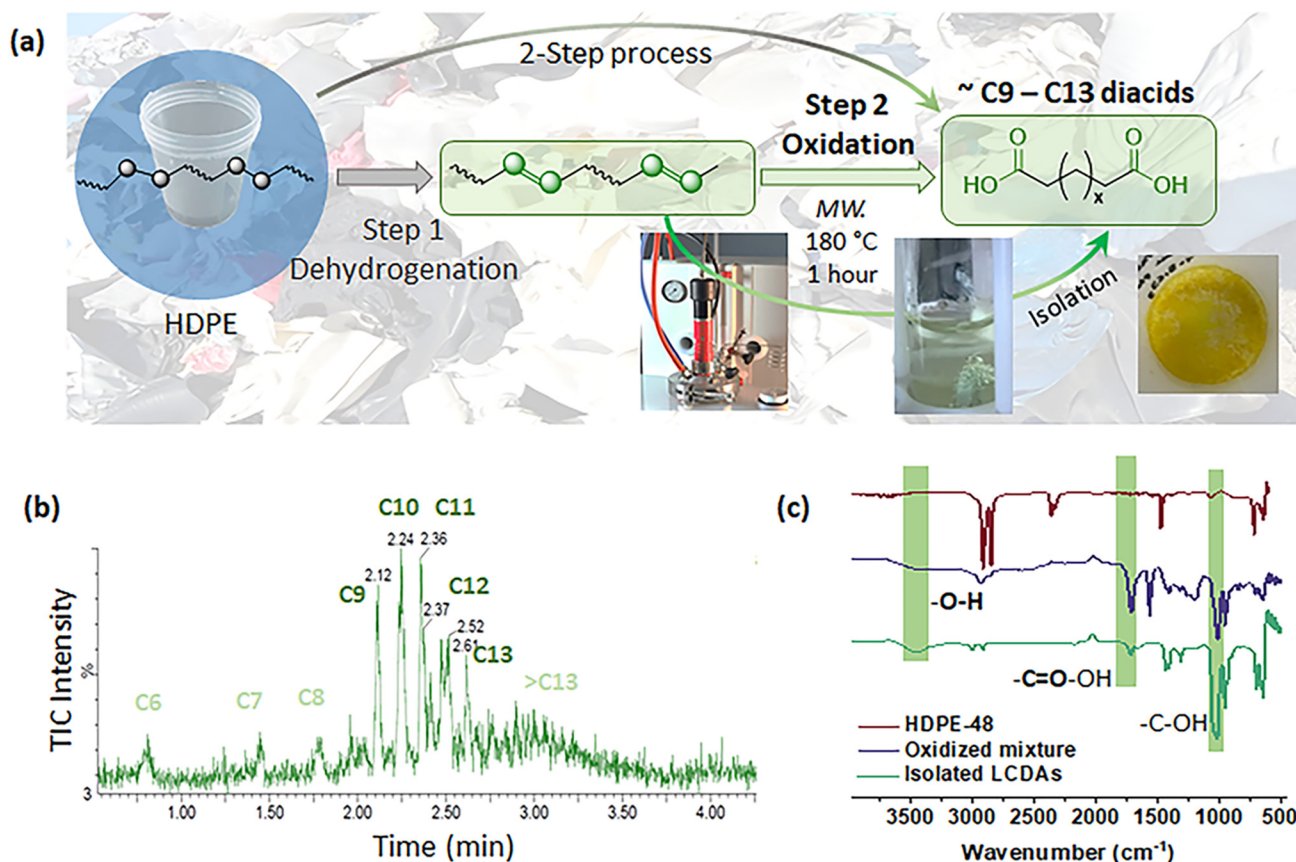


Fig. 5 (a) The microwave-assisted oxidation step of dehydrogenated HDPE to LCDAs using nitric acid as the oxidation reagent in a microwave reactor (Shelton, CT) for 1 hour at 180°C under an N_2 atmosphere; (b) UPLC-MS chromatograms of the oxidized fragments detected in the TIC channel, under the ESI-method; and (c) the FTIR spectra comparison of the dehydrogenated HDPE and the oxidative products.



displayed a prominent peak in the range of $3430\text{--}3300\text{ cm}^{-1}$, attributed to O–H stretching vibrations, and an asymmetric stretching vibration of the C=O bond at around 1710 cm^{-1} . Additionally, the C–O stretching vibration was identified at approximately 1015 cm^{-1} . Post-extraction spectra indicated the absence of nitric acid-related signals, further confirming the purity of the isolated products. The aliphatic nature of these compounds was corroborated by absorption peaks associated with CH_2 stretching modes ($2950\text{--}2850\text{ cm}^{-1}$) and scissoring vibrations ($1450\text{--}1400\text{ cm}^{-1}$), as well as a C–H bending vibration at 700 cm^{-1} . Nuclear magnetic resonance (NMR) spectroscopy provided further characterization of the diacids, revealing a predominantly linear structure with negligible branching, as inferred from the minimal presence of methyl groups in the ^1H and ^{13}C NMR spectra, as shown in Fig. S16.† In summary, the successful formation of LCDAs *via* this oxidative process has been validated through multiple analytical techniques, with the observed chain-length distribution primarily ranging between C9 and C13. The potential applications, *i.e.* raw material for polycondensation, and further processing of these diacids are under exploration.

After obtaining functional monomers, and in order to demonstrate an initial application of the isolated LCDAs, they were employed in a molten catalytic condensation process

using titanium(IV) butoxide catalyst alongside 1,5-pentanediol, utilizing a two-step polycondensation technique (Fig. 6a).⁶² The oligomers produced in the first step and the polyester created in the second step appear yellowish and opaque (Fig. 6b). In comparison with HDPE-48, both the oligomer and polyester exhibited notably smaller molecular sizes with results as depicted in Fig. 6c. However, following the two-step polycondensation, the M_n values of polyester increased from 1k to 3k, and M_w from 6k to 10k (Table S4†), quantified using high-temperature GPC. The polyester's thermal properties also presented complex behavior, with noticeable improvements over that of the oligomers (Fig. 6d and e). Despite the polycondensation processes, the FTIR spectra displayed consistent patterns (Fig. 6f). This trial confirms the potential of LCDAs for polyester synthesis, setting the stage for further exploration.

Future studies could systematically explore the effects of diol chain length and the impact of even-odd chain lengths on the chemical, thermal, crystalline, morphological, and mechanical properties of the resultant polyesters. While the recycling approaches for polyesters are more advanced compared to those for polyolefins,^{19,25} significant hurdles persist, particularly in the separation and decontamination processes. Considerable efforts are essential in order to transition from a linear polymer economy to a more sustainable, circular model.

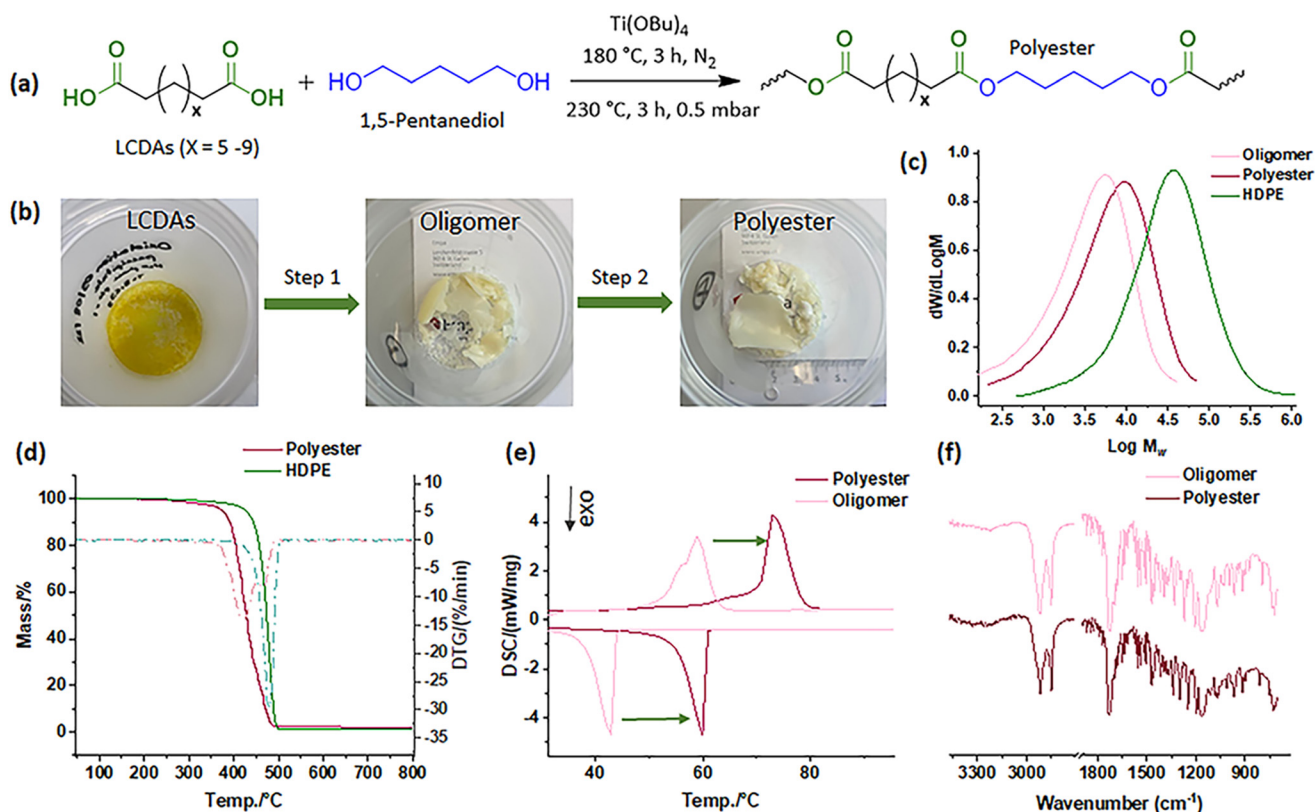


Fig. 6 (a) Demonstration of polyester synthesis using a two-step molten polycondensation process of the LCDAs with pentanediol and catalytic amount of $\text{Ti}(\text{OBu})_4$ at $180\text{ }^\circ\text{C}$ for 3 hours, then with reduced pressure at $230\text{ }^\circ\text{C}$ for another 3 hours; (b) the appearances of the oligomer and polyester; (c) comparison between the GPC traces of HDPE-48, oligomer and polyester; (d) TGA and DTG curves of the HDPE-48 and polyester; and (e) the melting and crystallization behaviors and (f) the FTIR spectra comparison of the oligomer and the polyester.



Environmental consideration

For the developed two-step upcycling process, we have considered specific environmental indicators. Since the *E*-factor (environmental factor) concept has been instrumental in addressing the issue of waste generation,⁶³ it has driven the development of cleaner processes for sustainable chemical manufacturing. It is defined as the mass ratio of waste to desired product, including reagents, solvent losses, and all process aids, except water (eqn (1)). We reviewed the two-step process and calculated the waste generation accordingly (Fig. S17†) with an estimated 10% solvent loss per step. Depending on the stage of development of the process, the use of simple *E*-factor (sEF) values for early route scouting activities, without taking solvents and water into account, is also appropriate (eqn (2)).⁶⁴

$$E\text{-factor} = (m_{\text{waste}} + m_{\text{solvent loss}}) / m_{\text{product}} \quad (1)$$

$$\text{Simple } E\text{-factors} = (\sum m_{\text{raw materials}} + \sum m_{\text{reagents}} - m_{\text{product}}) / m_{\text{product}} \quad (2)$$

The *E*-factor was calculated and was roughly 9.2, which is in the range of the fine chemical sectors. If solvents are excluded, the sEF was found to be 3.17. Compared with a few selected examples of functional molecule synthesis (Table S4†), this waste-based valorization process has demonstrated similar

environmental factors. However, further optimization is essential for upscaling the process in future work. For instance, in the dehydrogenation step, a mechanical process can be envisioned to replace the solvent-assisted solution to address the viscosity issue of polymer melts. Additionally, in the oxidation step, alternative catalytic oxidizing reagents will be screened for effective depolymerization.

Reclaiming of catalyst

In addition to the abovementioned points, the ability to reuse the catalyst holds significant economic relevance. Thus after the dehydrogenation step, the recovered catalyst was collected and underwent regeneration through calcination and hydrogen activation for subsequent utilization, as depicted in Fig. S1.† The TGA and DTG analyses indicated that the degradation of the polymer residue on the catalyst surface requires up to 400 °C in air (Fig. S18†); therefore a relatively harsh regeneration temperature (500 °C) for two hours with a temperature ramp of 3 °C min^{−1} was applied to reactivate the catalyst. However, TEM and EDX analyses, as presented in Fig. 7, confirmed not only further aggregation of the nanoparticles but also the accrual of a modest amount of carbon residues. The Pt nanoparticles displayed heterogeneity and formed aggregates with dimensions surpassing 10 nm, a significant increase from the approximately 2 nm observed in the freshly

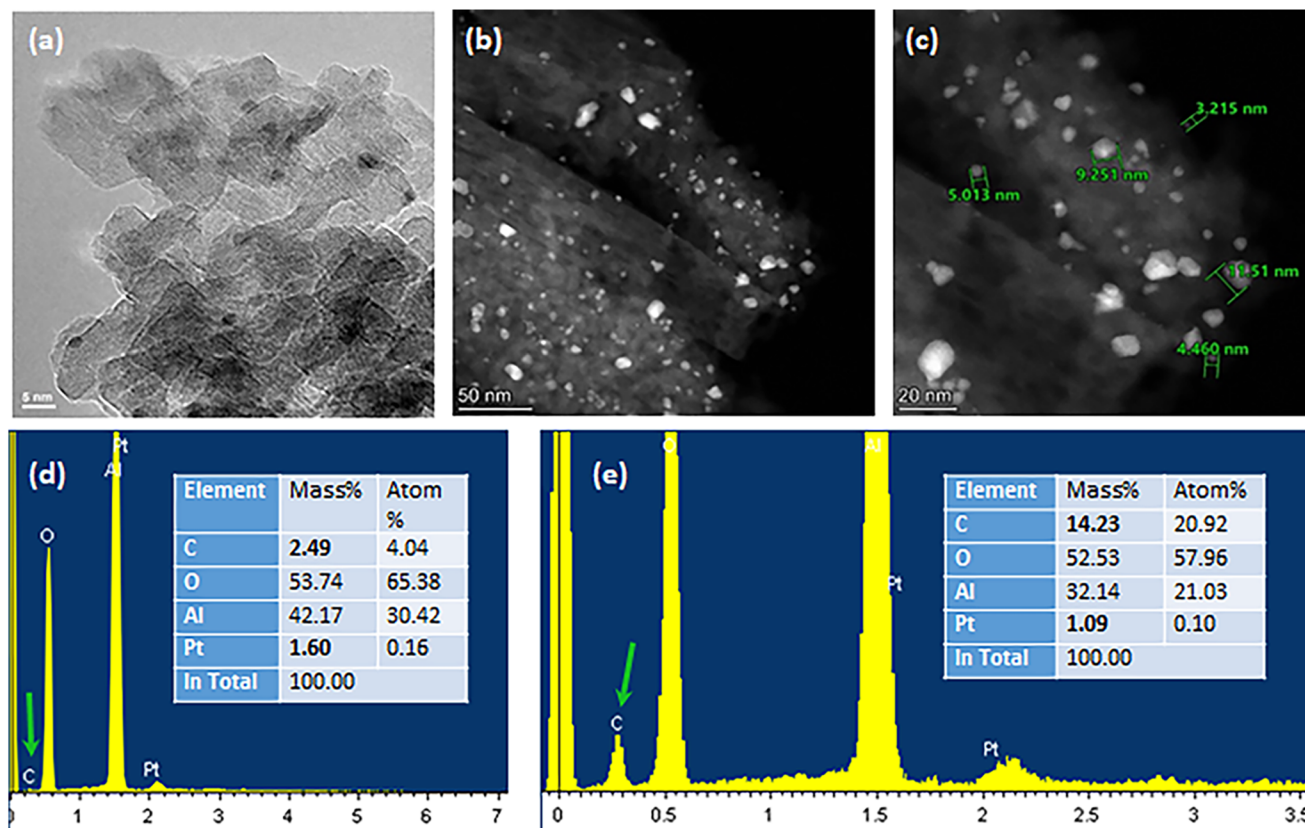


Fig. 7 (a) TEM image and (b and c) HAADF-STEM images of the recycled catalyst after one dehydrogenation use; (d) and (e) SEM-EDX spectra with a semi-quantitative estimate of the catalyst composition before and after use.



activated catalyst prior to use. EDX spectroscopy indicated a notable accumulation of carbon residues after a single catalytic cycle, as illustrated in Fig. 7 and Fig. S19.†

To address nanoparticle aggregation⁶⁵ and carbon buildup, future investigations should focus on fine-tuning the regeneration conditions, reducing Pt loading on the matrix to less than 0.5%, and incorporating a secondary metal component, such as tin (Sn), gallium (Ga), or indium (In),^{66,67} into the catalyst composition. This can be achieved using wet impregnation or plasma methods.⁶⁸ This additional metal is anticipated to form an alloy with Pt under operational conditions, enhancing the catalytic conversion, impeding metal sintering, and crucially, minimizing coke formation, thereby extending the catalyst's operational lifespan. The enhancements observed with platinum alloys are attributed to both geometric effects, which modify the crystalline structure of the catalyst, and electronic effects, influencing the reactivity of platinum's valence electrons and reducing the interaction between olefins and the catalyst.⁶⁹ Given the superior dehydrogenation efficacy of platinum alloys compared to monometallic platinum, their suitability in the HDPE dehydrogenation process warrants further exploration.

Conclusion

This study has demonstrated a chemical strategy for side- and upcycling of HDPE, addressing the critical need for treatment of plastic waste. By employing a two-step catalytic depolymerization process, we have successfully converted HDPE into value-added functional monomers, specifically long-chain aliphatic diacids, through the introduction of unsaturation imperfections *via* C=C double bond formation within the polymer chains, using a commercially available heterogeneous Pt/Al₂O₃ catalyst. The use of FTIR, Raman spectroscopy, and XPS for the characterization of the chemically modified HDPE has provided crucial insights, confirming the formation of C=C double bonds and enabling a semi-quantitative assessment of unsaturation levels. The successful production of long-chain aliphatic diacids from HDPE-48, with a diverse range of chain lengths (C9–C13), not only underscores the viability of this approach but also highlights the potential of using polymer waste as a chemical feedstock. This represents a step towards the realization of a circular economy by transforming plastic waste into critical resources, thereby mitigating environmental impact and contributing to sustainability. The utilization of these double bonds will be further investigated for HDPE functionalization or repurposed applications.

Meanwhile, this study acknowledges several challenges that require further attention, including the investigation of realistic recycling scenarios with impurities, optimization of the catalyst's lifetime, the purification of diacids, development of cleaner processes, and utilization of the resultant chemicals. The simplified environmental effect of the two-step process is estimated using the *E*-factor (9.2), which indicates potential to further improve process efficiency and environmental sustain-

ability. Additionally, employing diacids in polycondensation for application development remains valuable for future research. The goals of this research are to refine and expand the applicability of chemical recycling methods for HDPE. By continuing to explore innovative approaches to (partially) depolymerize HDPE more directly, efficiently, and cleanly, we can further contribute to the development of sustainable materials management practices and advance a circular economy.

Data availability

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

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

This work was supported by the Swiss Federal Office for the Environment (FOEN), with grant number UTF 666.14.21 and UTF 719.07.23. X. Z. thanks the National Centre of Competence in Research (NCCR) Catalysis (grant 180544) for financial support. We thank Dr. Edith Perret from Empa, St Gallen, Switzerland for assistance during the Raman measurements, and the help of Mr. Robin Pauer from Empa, Dübendorf, Switzerland for the STEM measurements.

References

- 1 P. Stegmann, V. Daioglou, M. Londo, D. P. van Vuuren and M. Junginger, *Nature*, 2022, **612**, 272–276.
- 2 E. MacArthur, *Science*, 2017, **358**, 843.
- 3 H. Li, H. A. Aguirre-Villegas, R. D. Allen, X. Bai, C. H. Benson, G. T. Beckham, S. L. Bradshaw, J. L. Brown, R. C. Brown, V. S. Cecon, J. B. Curley, G. W. Curtzwiler, S. Dong, S. Gaddameedi, J. E. García, I. Hermans, M. S. Kim, J. Ma, L. O. Mark, M. Mavrikakis, O. O. Olafasakin, T. A. Osswald, K. G. Papanikolaou, H. Radhakrishnan, M. A. Sanchez Castillo, K. L. Sánchez-Rivera, K. N. Tumu, R. C. Van Lehn, K. L. Vorst, M. M. Wright, J. Wu, V. M. Zavala, P. Zhou and G. W. Huber, *Green Chem.*, 2022, **24**, 8899–9002.
- 4 R. Wu, W. Wu Klingler, L. Stieglitz, S. Gaan, B. Rieger and M. Heuberger, *Coord. Chem. Rev.*, 2023, **474**, 214844.
- 5 R. Geyer, J. R. Jambeck and K. L. Law, *Sci. Adv.*, 2017, **3**, e1700782.
- 6 Y. Weng, C.-B. Hong, Y. Zhang and H. Liu, *Green Chem.*, 2024, **26**(2), 571–592.
- 7 K. Faust, P. Denifl and M. Hapke, *ChemCatChem*, 2023, **15**, e202300310.



- 8 A. Arroyave, S. Cui, J. C. Lopez, A. L. Kocen, A. M. LaPointe, M. Delferro and G. W. Coates, *J. Am. Chem. Soc.*, 2022, **144**, 23280–23285.
- 9 J. Y. Q. Teo, C. W. S. Yeung, T. T. Y. Tan, W. W. Loh, X. J. Loh and J. Y. C. Lim, *Green Chem.*, 2022, **24**, 6287–6294.
- 10 L. Chen, K. G. Malollari, A. Uliana, D. Sanchez, P. B. Messersmith and J. F. Hartwig, *Chem*, 2021, **7**, 137–145.
- 11 M. Chanda, *Adv. Ind. Eng. Polym. Res.*, 2021, **4**, 133–150.
- 12 G. Celik, R. M. Kennedy, R. A. Hackler, M. Ferrandon, A. Tennakoon, S. Patnaik, A. M. LaPointe, S. C. Ammal, A. Heyden, F. A. Perras, M. Pruski, S. L. Scott, K. R. Poeppelmeier, A. D. Sadow and M. Delferro, *ACS Cent. Sci.*, 2019, **5**, 1795–1803.
- 13 I. Barbarias, M. Artetxe, G. Lopez, A. Arregi, L. Santamaria, J. Bilbao and M. Olazar, *Catalysts*, 2019, **9**, 414.
- 14 M. Chu, W. Tu, S. Yang, C. Zhang, Q. Li, Q. Zhang and J. Chen, *SusMat*, 2022, **2**, 161–185.
- 15 J. M. Garcia and M. L. Robertson, *Science*, 2017, **358**, 870–872.
- 16 X. Zhao and F. You, *AIChE J.*, 2021, **67**, e17127.
- 17 S. C. Kosloski-Oh, Z. A. Wood, Y. Manjarrez, J. P. de los Rios and M. E. Fieser, *Mater. Horiz.*, 2021, **8**, 1084–1129.
- 18 X. Jia, C. Qin, T. Friedberger, Z. Guan and Z. Huang, *Sci. Adv.*, 2016, **2**, e1501591.
- 19 M. Baur, N. K. Mast, J. P. Brahm, R. Habé, T. O. Morgen and S. Mecking, *Angew. Chem., Int. Ed.*, 2023, **62**, e202310990.
- 20 C. Jehanno, M. M. Pérez-Madriral, J. Demarteau, H. Sardon and A. P. Dove, *Polym. Chem.*, 2019, **10**, 172–186.
- 21 B. Whajah, N. da Silva Moura, J. Blanchard, S. Wicker, K. Gandar, J. A. Dorman and K. M. Dooley, *Ind. Eng. Chem. Res.*, 2021, **60**, 15141–15150.
- 22 F. Zhang, M. Zeng, R. D. Yappert, J. Sun, Y. H. Lee, A. M. LaPointe, B. Peters, M. M. Abu-Omar and S. L. Scott, *Science*, 2020, **370**, 437–441.
- 23 Z. Xu, N. E. Munyaneza, Q. Zhang, M. Sun, C. Posada, P. Venturo, N. A. Rorrer, J. Miscall, B. G. Sumpter and G. Liu, *Science*, 2023, **381**, 666–671.
- 24 C. Jehanno, J. W. Alty, M. Roosen, S. De Meester, A. P. Dove, E. Y. X. Chen, F. A. Leibfarth and H. Sardon, *Nature*, 2022, **603**, 803–814.
- 25 D. Parida, A. Aerts, K. Vanbroekhoven, M. Van Dael, H. Mitta, L. Li, W. Eevers, K. M. Van Geem, E. Feghali and K. Elst, *Prog. Polym. Sci.*, 2023, 101783, DOI: [10.1016/j.progpolymsci.2023.101783](https://doi.org/10.1016/j.progpolymsci.2023.101783).
- 26 R. J. Conk, S. Hanna, J. X. Shi, J. Yang, N. R. Ciccio, L. Qi, B. J. Bloomer, S. Heuvel, T. Wills, J. Su, A. T. Bell and J. F. Hartwig, *Science*, 2022, **377**, 1561–1566.
- 27 L. D. Ellis, N. A. Rorrer, K. P. Sullivan, M. Otto, J. E. McGeehan, Y. Román-Leshkov, N. Wierckx and G. T. Beckham, *Nat. Catal.*, 2021, **4**, 539–556.
- 28 M. Häußler, M. Eck, D. Rothauer and S. Mecking, *Nature*, 2021, **590**, 423–427.
- 29 K. P. Sullivan, A. Z. Werner, K. J. Ramirez, L. D. Ellis, J. R. Bussard, B. A. Black, D. G. Brandner, F. Bratti, B. L. Buss, X. Dong, S. J. Haugen, M. A. Ingraham, M. O. Konev, W. E. Michener, J. Miscall, I. Pardo, S. P. Woodworth, A. M. Guss, Y. Román-Leshkov, S. S. Stahl and G. T. Beckham, *Science*, 2022, **378**, 207–211.
- 30 C. Rabot, Y. Chen, S. Bijlani, Y.-M. Chiang, C. E. Oakley, B. R. Oakley, T. J. Williams and C. C. C. Wang, *Angew. Chem., Int. Ed.*, 2023, **62**, e202214609.
- 31 F. Stempfle, P. Ortmann and S. Mecking, *Chem. Rev.*, 2016, **116**, 4597–4641.
- 32 M. Piccini, J. Lightfoot, B. C. Dominguez and A. Buchard, *ACS Appl. Polym. Mater.*, 2021, **3**, 5870–5881.
- 33 S. Pandey, B. S. Rajput and S. H. Chikkali, *Green Chem.*, 2021, **23**, 4255–4295.
- 34 Z. Wang, M. S. Ganewatta and C. Tang, *Prog. Polym. Sci.*, 2020, **101**, 101197.
- 35 Z. Petrovic, *Polym. Rev.*, 2008, **48**, 109–155.
- 36 M. Oshinowo, J. R. Runge, M. Piccini, F. Marken and A. Buchard, *J. Mater. Chem. A*, 2022, **10**, 6796–6808.
- 37 T. F. Nelson, D. Rothauer, M. Sander and S. Mecking, *Angew. Chem., Int. Ed.*, 2023, **62**, e202310729.
- 38 E. Bäckström, K. Odelius and M. Hakkarainen, *ACS Sustainable Chem. Eng.*, 2019, **7**, 11004–11013.
- 39 J. J. Sattler, J. Ruiz-Martinez, E. Santillan-Jimenez and B. M. Weckhuysen, *Chem. Rev.*, 2014, **114**, 10613–10653.
- 40 L. S. Diaz-Silvarrey, K. Zhang and A. N. Phan, *Green Chem.*, 2018, **20**, 1813–1823.
- 41 C. Li and G. Wang, *Chem. Soc. Rev.*, 2021, **50**, 4359–4381.
- 42 J. H. Kwak, J. Hu, D. Mei, C.-W. Yi, D. H. Kim, C. H. F. Peden, L. F. Allard and J. Szanyi, *Science*, 2009, **325**, 1670–1673.
- 43 S. Brunauer, P. H. Emmett and E. Teller, *J. Am. Chem. Soc.*, 1938, **60**, 309–319.
- 44 H. Jeong, H. B. Bathula, T. W. Kim, G. B. Han, J. H. Jang, B. Jeong and Y.-W. Suh, *ACS Sustainable Chem. Eng.*, 2021, **9**, 1193–1202.
- 45 E. Bus and J. A. van Bokhoven, *Phys. Chem. Chem. Phys.*, 2007, **9**, 2894–2902.
- 46 C. Copéret, A. Comas-Vives, M. P. Conley, D. P. Estes, A. Fedorov, V. Mougel, H. Nagae, F. Núñez-Zarur and P. A. Zhizhko, *Chem. Rev.*, 2016, **116**, 323–421.
- 47 G. Bergeret and P. Gallezot, in *Handbook of Heterogeneous Catalysis*, 2008, pp. 738–765. DOI: [10.1002/9783527610044.hetc0038](https://doi.org/10.1002/9783527610044.hetc0038).
- 48 H. W. Starkweather Jr. and R. H. Boyd, *J. Phys. Chem.*, 1960, **64**, 410–414.
- 49 T. Furukawa, H. Sato, Y. Kita, K. Matsukawa, H. Yamaguchi, S. Ochiai, H. W. Siesler and Y. Ozaki, *Polym. J.*, 2006, **38**, 1127–1136.
- 50 J. Hoffman, G. Davis and J. Lauritzen, in *Treatise on Solid State Chemistry*, ed. N.B. Hannay, Plenum, 1976, vol. 3, p. 497.
- 51 G. Zerbi, G. Gallino, N. Del Fanti and L. Baini, *Polymer*, 1989, **30**, 2324–2327.
- 52 J. R. Beattie, S. E. J. Bell and B. W. Moss, *Lipids*, 2004, **39**, 407–419.



- 53 H. Wu, J. V. Volponi, A. E. Oliver, A. N. Parikh, B. A. Simmons and S. Singh, *Proc. Natl. Acad. Sci. U. S. A.*, 2011, **108**, 3809–3814.
- 54 G. R. Strobl and W. Hagedorn, *J. Polym. Sci., Polym. Phys. Ed.*, 1978, **16**, 1181–1193.
- 55 S. Turgeon and R. W. Paynter, *Thin Solid Films*, 2001, **394**, 43–47.
- 56 B. Lesiak, J. Zemek, J. Houdkova, P. Jiricek and A. Jóźwik, *Polym. Degrad. Stab.*, 2009, **94**, 1714–1721.
- 57 S. Kaciulis, *Surf. Interface Anal.*, 2012, **44**, 1155–1161.
- 58 G. Beamson and D. Briggs, *High Resolution XPS of Organic Polymers: The Scienta ESCA300 Database*, Wiley, 1992.
- 59 B. Lesiak, J. Zemek and J. Houdkova, *Polymer*, 2008, **49**, 4127–4132.
- 60 S. Kaciulis, A. Mezzi, P. Calvani and D. M. Trucchi, *Surf. Interface Anal.*, 2014, **46**, 966–969.
- 61 Y. Mizokawa, T. Miyasato, S. Nakamura, K. M. Geib and C. W. Wilmsen, *Surf. Sci.*, 1987, **182**, 431–438.
- 62 L. Genovese, M. Gigli, N. Lotti, M. Gazzano, V. Siracusa, A. Munari and M. Dalla Rosa, *Ind. Eng. Chem. Res.*, 2014, **53**, 10965–10973.
- 63 R. A. Sheldon, *Green Chem.*, 2007, **9**, 1273–1283.
- 64 R. A. Sheldon, *Green Chem.*, 2017, **19**, 18–43.
- 65 W. Zhou, S. R. Docherty, C. Ehinger, X. Zhou and C. Copéret, *Chem. Sci.*, 2023, **14**, 5379–5385.
- 66 M. Abdelgaid, J. Dean and G. Mpourmpakis, *Catal. Sci. Technol.*, 2020, **10**, 7194–7202.
- 67 H. Yan, K. He, I. A. Samek, D. Jing, M. G. Nanda, P. C. Stair and J. M. Notestein, *Science*, 2021, **371**, 1257–1260.
- 68 D. Hegemann, M. Amberg, A. Ritter and M. Heuberger, *Mater. Tech.*, 2009, **24**, 41–45.
- 69 S. Chen, X. Chang, G. Sun, T. Zhang, Y. Xu, Y. Wang, C. Pei and J. Gong, *Chem. Soc. Rev.*, 2021, **50**, 3315–3354.

