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# Flow synthesis of hypercrosslinked polymers with additional microporosity that enhances CO<sub>2</sub>/N<sub>2</sub> separation†

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Hypercrosslinked polymers (HCPs) are typically synthesised over 24 hour batch reactions, limiting productivity rates during scale-up production. Continuous flow synthesis can potentially overcome this limitation. However, the formation of insoluble HCP products, compounded by HCP expansion due to solvent adsorption during synthesis can clog flow reactors. Here, we overcome clogging issues through reactor design and optimisation of synthesis parameters. Using this reactor, we synthesised HCPs via internal, post-, and external crosslinking strategies underpinned by Friedel–Crafts alkylation over various synthesis parameters – residence time, substrate concentration, reagent ratio, and temperature. The space-time-yield (STY) values, a key parameter for productivity rates, of flow synthesis were 32–117 fold higher than those in batch reactions. HCPs produced via internal crosslinking in flow synthesis contained additional microporosity that enhanced CO<sub>2</sub>/N<sub>2</sub> selectivity at 298 K by 850% when compared to HCPs produced in batch reactions. Outcomes from this work could enable high productivity scale-up production of HCPs for post-carbon capture.

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## Introduction

Hypercrosslinked polymers (HCPs) are excessively crosslinked, amorphous polymers typically synthesised via Friedel–Crafts alkylation (Fig. 1). In this reaction, aromatic protons are replaced with methylene bridges that are linked to an adjacent aromatic molecule. This is facilitated by an electrophilic attack on the aromatic ring by a carbocation. This can be achieved by post-, internal, and external crosslinking.<sup>1</sup> Post-crosslinking uses halogenated crosslinkers to link up the chains of polymers such as polystyrene or polystyrene-*co*-divinylbenzene. Internal crosslinking is akin to a condensation reaction in which halogenated compounds are crosslinked. External crosslinking exploits formaldehyde dimethyl acetal (FDA) as the crosslinker to link up simple aromatic compounds.<sup>2</sup> During the crosslinking process, solvent molecules pre-occupy the free spaces between the newly linked up polymer chains. Removing

solvent molecules from the product by drying leads to the formation of porous architectures that underpin Brunauer–Emmett–Teller (BET) surface areas as high as 2000 m<sup>2</sup> g<sup>-1</sup>.<sup>3</sup> As such, HCPs are widely deployed in the industry as ion-exchange resins<sup>4</sup> such as Puro-lite's Macronet™,<sup>5</sup> DowDupont Inc.'s Amberlite™, and Lanxess AG's Lewatit® series, and potentially for gas storage and separations.<sup>6,7</sup> The global market value of HCPs in 2020 was USD\$1.4 billion and is expected to reach USD\$2 billion by 2028.<sup>8</sup>

Most HCPs are produced via batch reactions over 18–24 hours<sup>9</sup> to ensure extensive crosslinking<sup>10</sup> and this may be inadequate to produce enough HCPs to meet the growing global demand. In the first hour of batch reactions, more than 80% of the reagents are used up in cross-linking. The pores of the initial polymer will adsorb solvent molecules, creating a gel-like structure. More solvent is then added to disperse the initially-

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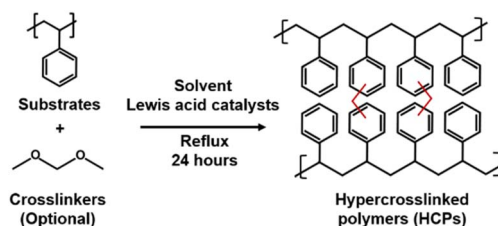


Fig. 1 Generic Friedel–Crafts alkylation for HCP synthesis.



formed polymers, breaking up the gel-like material to ensure complete crosslinking.

Most studies in this field are focused on developing new HCP materials with unprecedented adsorption capacity,<sup>11,12</sup> identifying alternative reagents (solvents, monomers) for the sustainable synthesis of HCPs,<sup>13</sup> and improving batch productivity rates.<sup>14</sup> For example, low HCP productivity rates production can be improved by reducing synthesis time from 24 hours to 5–35 minutes *via* solvent-free mechanochemical ball milling and liquid-assisted grinding.<sup>15</sup> This significant reduction in synthesis duration does not impact HCP quality, with specific surface areas reaching 625–1720 m<sup>2</sup> g<sup>-1</sup>. Another technique suitable for improving the productivity of HCP production is flow synthesis.

The key benefit of replacing batch reactions with flow synthesis is improving productivity rates. To date, flow chemistry had been used for synthesising microporous materials, such as polymers of intrinsic microporosity (PIM),<sup>16</sup> metal-organic frameworks (MOFs),<sup>17</sup> covalent organic frameworks (COFs),<sup>18</sup> and HCPs.<sup>10</sup> Flow syntheses of MOFs and COFs enhanced the space-time yield (STY) – a key indicator of productivity rates, comprising product yield, operating time, and reactor volume,<sup>17</sup> by 30 fold,<sup>19</sup> and 94 fold,<sup>18</sup> respectively. These significant STY improvements were attributed to a reduced production time. Meanwhile, Fritsch deployed continuous flow reactions to reduce the PIM synthesis time by 90%.<sup>16</sup> The first reported example of flow synthesis of HCPs was reported in 2000 when Vincent and co-workers reported using a tubular reactor to control the particle size of HCPs formed *via* post-crosslinking during suspension polymerisation.<sup>20</sup> However, this work mainly focused on narrowing the particle size distribution. Our group previously deployed flow synthesis to produce HCPs *via* the external crosslinking method.<sup>10</sup> A key limitation of this proof-of-concept was reactor clogging by the newly formed insoluble HCP products that expanded upon solvent adsorption within the reactor.

In this work, we overcome reactor clogging by optimising reactor design and synthesis parameters – residence time, temperature, reagent concentration, and ratio for flow synthesis of HCPs *via* internal crosslinking of  $\alpha,\alpha'$ -dichloro-*p*-xylene (DCX), post- and external crosslinking of waste Styrofoam in a bespoke coil reactor. To demonstrate the effects of flow synthesis on HCP quality and productivity rates, we compared the specific surface areas, micropore volumes, and STY value of HCPs produced from the flow synthesis with batch reactions. Results from this work support the theory that flow synthesis could be used to scale the production of good-quality HCPs. This is different from existing approaches that rely on HCP functionalisation,<sup>21–23</sup> utilisation of bespoke<sup>24</sup> or novel<sup>25,26</sup> monomers in HCP synthesis, and modified synthesis protocols.<sup>15</sup> Our approach establishes a new method to tailor and enhance HCP CO<sub>2</sub>/N<sub>2</sub> selectivity *via* reactor design and flow synthesis.

## Experimental

### Materials and equipment

Styrofoam or waste-expanded polystyrene (PS) was obtained from plastic waste collected at The University of Edinburgh. 1,2-

Dichloroethane (DCE) and iron(III) chloride (FeCl<sub>3</sub>) were purchased from Alfa Aesar. Formaldehyde dimethyl acetal (FDA) and  $\alpha,\alpha'$ -dichloro-*p*-xylene (DCX) were purchased from Sigma-Aldrich. Methanol, chloroform, and acetone were purchased from Thermo Fisher Scientific. All chemicals were used as is.

Polytetrafluoroethylene (PTFE) tubes with dimensions of 1/8 in × 0.063 in × 50 ft were purchased from Supelco Inc., while PTFE tube fittings (Y-junction connectors, female union connectors, ferrules, and nuts) were purchased from Nanjing Runze Fluid Control Equipment Co., Ltd. Polypropylene syringes were purchased from MB Fibreglass, while 21G needles were supplied from Becton, Dickinson, and Company. A tube holder was fabricated from polylactic acid (PLA) filaments using a Prusa i3 MK3S 3D printer. The design was based on two different shapes with 7 individual parts, a central ring, and holder wings, which can be hinged on to each other. A peristaltic dosing pump (SEKO Kronos 50) was used to pump the monomer solution as well as cleaning solvents after the operation. A syringe pump (Cole-Parmer® SP210iwz) was deployed to inject the catalyst solution into the reactor.

### HCP synthesis in batch reactors

HCPs synthesised in batch reactors were used as control materials here. All batch syntheses were performed in a 250 mL three-neck round-bottom glass flask connected to a reflux condenser. HCPs were synthesised *via*:

(i) Internal crosslinking – 2.884 g of DCX and 2.672 g of FeCl<sub>3</sub> were dissolved in 100 mL DCE. The mixture was heated to 373 K (ref. 24) and continuously stirred with a magnetic stirrer for 24 hours.

(ii) Post-crosslinking – 0.385 g of PS, 0.385 g of FeCl<sub>3</sub>, and 0.385 g of DCX were first dissolved in 100 mL of DCE. The mixture was heated to 353 K (ref. 27) *via* an oil bath and continuously stirred with a magnetic stirrer for 24 hours.

(iii) External crosslinking – 0.762 g of PS, 3.810 g of FeCl<sub>3</sub>, and 3.810 g of FDA were dissolved in 100 mL of DCE. The mixture was heated to 353 K (ref. 28) *via* an oil bath and continuously stirred with a magnetic stirrer for 24 hours.

### HCP synthesis in flow reactors

Fig. 2 and S1† show the experimental set-up of the flow reactor used in this work for HCP synthesis *via* post-, internal, and external crosslinking. The details of the synthesis *via* the flow system are described as follows:

(i) Internal crosslinking – the substrate solution was prepared by dissolving 1.154 g of DCX in 20 mL of DCE. A

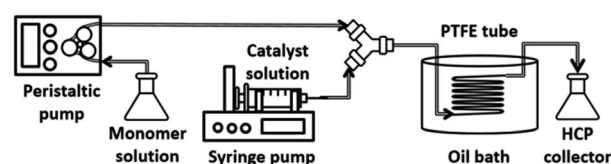


Fig. 2 Schematic showing the equipment used in our flow reactor.



peristaltic pump was used to feed the substrate solution into the system. The catalyst solution was prepared by mixing 1.069 g of FeCl<sub>3</sub> in 20 mL of DCE and then loaded into a syringe and pumped into the system by a syringe pump. All reactants were mixed in a Y-junction connector prior to the entry into the reactor at the total volumetric flow rate of 5.40 mL min<sup>-1</sup>. The reactor was submerged in a heated oil bath setting at 343 K. The slurry product was collected at the end of the reactor.

(ii) Post-crosslinking – the substrate solution was prepared by dissolving 0.154 g of PS in 20 mL of DCE. The catalyst solution was prepared by mixing 1.524 g of FeCl<sub>3</sub> and 1.524 g of DCX in 20 mL of DCE and then loaded into a syringe. The substrate solution was fed into the system *via* a peristaltic pump, while the catalyst solution was fed *via* a syringe pump. Both solutions were fed into the reactor at the total volumetric flow rate of 2.94 mL min<sup>-1</sup> and mixed at a Y-junction. The reactor was submerged in a pre-heated oil bath at 343 K. The slurry product was collected at the end of the reactor.

(iii) External crosslinking – the substrate solution was prepared by dissolving 0.305 g of PS in 20 mL of DCE. The catalyst solution was prepared by mixing 1.524 g of FeCl<sub>3</sub> and 1.524 g of FDA in 20 mL of DCE and loaded into a syringe. A peristaltic pump was used for feeding substrate solution into the reactor. A syringe pump was used for feeding the catalyst solution into the system. The total volumetric flow rate was 1.94 mL min<sup>-1</sup>. Both solutions were mixed at the Y-junction before entering the reactor, which was pre-heated inside the oil bath at 343 K. The slurry product was collected at the end of the reactor.

The reactor volume was approximately 30 mL (1.58 mm inner diameter, 15.21 m length). Our reactor comprised a 50 mm pitch helical coil with multiple curvature radii (34, 39, 44, 49, and 54 mm) designed to minimise the spatial footprint.

Since the reactor volume (*V*) was fixed, a change in the residence time (*τ*) depended on the total volumetric feed flow rate (*F*), as shown in the following equation:

$$F = \frac{V}{\tau}$$

The operating time of flow synthesis (*t*<sub>op</sub>) was calculated using the following equation where *V*<sub>feed</sub> was denoted as the volume of the feed solution:

$$t_{\text{op}} = \frac{V + V_{\text{feed}}}{F}$$

Upon completion of each synthesis process, either in batch or flow, products were washed with chloroform and methanol until the washings became clear. Then, the samples were soaked overnight in methanol and rewashed in the same way with methanol and acetone. Finally, samples were dried in an oven at 333 K for 8 hours and collected for further characterisation.

### Space-time-yield (STY) of HCP synthesis

HCP yield was calculated by dividing the weight of the dried powder produced by the weight of monomers or substrates. The

amount of external crosslinkers was not included in this calculation as the crosslinking could be formed from other halogenated compounds used in synthesis, *i.e.*, the chloro group in DCX (substrate) or DCE (solvent).<sup>27</sup> This could lead to a yield value of more than 100%.<sup>29</sup> Hence, HCP yield was calculated using the following equation:<sup>30</sup>

$$\text{Product yield} = \frac{\text{weight of dried particles}}{\text{weight of monomers or substrates}}$$

STY is the term used in flow chemistry to determine how much product, *i.e.*, yield is produced in a reactor volume within a time period. As batch reactions here took place over 24 hours, the operating period for the calculation of the flow system was then fixed to per day unit. The STY calculations for batch and flow systems<sup>17,31</sup> were shown in the following equations:

$$\text{STY}_{\text{batch}} = \frac{\text{substrate weight per batch} \times \text{product yield}}{\text{volume of solvent per batch}}$$

$$\text{STY}_{\text{flow}} = \frac{\text{inlet substrate weight per day} \times \text{product yield}}{\text{reactor volume}}$$

### Characterisation

The pore size distributions of HCPs characterize in this work were characterized with N<sub>2</sub> gas adsorption at 77 K using various gas analysers at The University of Edinburgh, Strathclyde University, and CSIRO, Australia. These isotherms were used to determine the specific surface area and for porosity analysis using the BET theory and density functional theory (DFT) method, respectively, using the AsiQwin software. Rouquerol's four criteria were applied for BET surface area correction for microporous materials (Fig. S4†).<sup>32,33</sup> Samples were degassed in a vacuum at 373 K for 12 hours before characterisation.

Fourier transform infrared spectroscopy (FTIR) was used to investigate the chemical structure of the reagents and HCPs synthesised in this work. This was achieved with an ATR-FTIR (Nicolet iS10 with Smart iTX Diamond accessory, Thermo Fisher Scientific, Waltham, MA) with 64 scans per sample and a resolution of 0.48 cm<sup>-1</sup>. The chemical structures of the synthesised HCPs were validated with solid-state <sup>13</sup>C cross polarisation (CP) magic angle spinning (MAS) nuclear magnetic resonance (NMR) spectroscopy using a Bruker Avance III spectrometer with a 9.4 T superconducting magnet.

N<sub>2</sub> and CO<sub>2</sub> adsorption studies at 298 K were performed and the adsorption isotherms were used to calculate CO<sub>2</sub>/N<sub>2</sub> selectivity (*S*<sub>CO<sub>2</sub>/N<sub>2</sub></sub>) based on the ideal adsorption solution theory (IAST). The standard conditions<sup>34</sup> of 298 K and 15 : 85 molar ratio of the inlet gas mixture (*y*<sub>CO<sub>2</sub></sub> = 0.15, *y*<sub>N<sub>2</sub></sub> = 0.85) were chosen for this calculation using following equations:

$$S_{\text{CO}_2/\text{N}_2} = (x_{\text{CO}_2}/y_{\text{CO}_2}) / (x_{\text{N}_2}/y_{\text{N}_2}) = \frac{P_{\text{N}_2}^0}{P_{\text{CO}_2}^0}$$

$$P_i y_i = P_1^0 x_i$$



$$x_{\text{CO}_2} + x_{\text{N}_2} = 1$$

$$y_{\text{CO}_2} + y_{\text{N}_2} = 1$$

$$\int_0^{P_{\text{CO}_2}^0} \frac{Q_{\text{CO}_2}}{P} dP = \int_0^{P_{\text{N}_2}^0} \frac{Q_{\text{N}_2}}{P} dP$$

$x_i$  and  $y_i$  denote the mole fraction of gas  $i$  (where  $i = \text{CO}_2, \text{N}_2$ ) in the adsorbed phase and gas phase, respectively.  $P_i^0$  is the equilibrium pressure for pure  $i$  assuming that spreading pressure ( $\pi$ ) was constant during isothermal adsorption. The single-site Langmuir–Freundlich model (SSLF) was selected to associate the adsorbed amount of gas ( $Q_i$ ) with the function of pressure ( $P$ ), as shown in the following equation:

$$Q = \frac{qkP^n}{1 + kP^n}$$

The parameters from the SSLF model equation,  $q$ ,  $k$ , and  $n$ , represented the maximum-adsorbed amount of pure gas at the adsorption site, the affinity parameter, and the solid heterogeneity respectively. These parameters were obtained from the regression analysis of the experimental adsorption isotherm data.

Transmission electron spectroscopy (TEM) was deployed to characterise HCP morphology. The samples were dispersed in DCE and deposited onto carbon-coated copper grids (Agar Scientific). Excess solvent was blotted away and the grid was air-dried. Samples were imaged in an FEI F20 microscope operating at 200 kV. The images were taken under low-dose conditions using a Gatan Rio CCD camera. The micrographs (Fig. S8) are shown in ESI.†

## Results and discussions

### The productivity of the flow processes

Here, we show that HCP synthesis duration could be reduced, from 24 hours to 5–15 minutes in a flow reactor *via* internal-, post-, and external crosslinking strategies (Table 1). This was attributed to the enhanced heat and mass transfer rates associated with the higher specific area (the ratio of the surface area to volume)<sup>35</sup> of the flow reactor deployed in this work. The specific area of the flow reactor deployed in this work was 5087  $\text{m}^2 \text{m}^{-3}$ , 68 fold higher than the 75  $\text{m}^2 \text{m}^{-3}$  of the batch reactor used here (Table S4†). Reactors with a high surface area to volume ratio<sup>35</sup> typically possess the advantages of rapid heat and mass transfers that enhance the reaction kinetics.<sup>36</sup>

In our work, heat transfer was mainly impacted by this significant difference in reactor-specific surface area. The larger specific surface area of the flow reactor enhanced heat transfer, *i.e.*, the energy flow from the oil bath to the reactor walls that were in close contact with the reactants inside the reactor. At the same time, the small volume of the flow reactor offered short diffusion pathways (better mass transfer) to enable faster mixing.<sup>37</sup> The combination of these factors was favourable for polymerisation/crosslinking kinetics. The effects of the thermal conductivity of the reactor material and heat source temperature were negligible here. The difference in thermal conductivities between the glass batch reactor ( $1 \text{ W m}^{-1} \text{ K}^{-1}$ )<sup>38</sup> and polytetrafluoroethylene (PTFE) flow reactor ( $0.25 \text{ W m}^{-1} \text{ K}^{-1}$ )<sup>39</sup> was 4 times, while the difference in heat source temperatures used in both batch and flow syntheses was only 10 K.

In terms of mass transfer, a higher degree of mixing is favourable for the reaction as reagent particles have more access to the reactor surface where the highest heat energy is provided.<sup>17</sup> The mixing efficiency could be correlated to the

Table 1 Operating parameters of HCP synthesis in batch and flow systems and their corresponding yield

Crosslinking approach	Substrate–crosslinker type	Substrate concentration [% w/v]	Substrate : FeCl <sub>3</sub> : crosslinker	Temperature [K]	Volumetric flow rate [mL min <sup>-1</sup> ]	Reaction time <sup>a</sup> [min]	HCP yield [%]	Sample <sup>b</sup>
Internal crosslinking	DCX–N.A.	2.884	1 : 1 : 0 (mol)	373	N.A.	1440	58.50	I-B-HCP
		2.884	1 : 1 : 0 (mol)	343	5.40	5	7.24	I-F-HCP
Post-crosslinking	PS–DCX	0.385	1 : 1 : 1 (wt)	353	N.A.	1440	155.38	P-B-HCP
		0.385	1 : 1 : 1 (wt)	343	2.94	10	126.72	P-F-HCP
External crosslinking	PS–FDA	0.762	1 : 5 : 5 (wt)	353	N.A.	1440	156.63	E-B-HCP
		0.762	1 : 5 : 5 (wt)	343	1.94	15	114.30	E-F-HCP

<sup>a</sup> Residence time is an equivalent term for reaction time in the flow system. <sup>b</sup> The samples were named X-Y-HCP in which X denotes hypercrosslinking approach (I for internal crosslinking, P for post crosslinking, E for external crosslinking) and Y denotes synthesis system (B for batch reactions, F for flow synthesis).



dimensionless Reynolds number ( $Re$ ) which describes fluid flow patterns – laminar, transient, and turbulence. Turbulent flow is typically associated with good mixing as an external driving force is applied to induce particle diffusion, rather than relying solely on molecular diffusion.<sup>17</sup> In a stirred tank reactor, the turbulent region is where  $Re > 10\,000$ ,<sup>40</sup> while turbulent fluid flow in a helical coil reactor is where  $Re > 40$ .<sup>41</sup>

Using an equation for a propeller turbine mixer tank,<sup>42</sup> we estimated that the  $Re$  number in our batch reactor reached a value of 6524 (see ESI†). This indicated transient fluid flow (the transition region between laminar and turbulent flow pattern,  $100 < Re < 10\,000$ ) in our batch reactor, *i.e.*, moderate mixing efficiency. Meanwhile, the  $Re$  number in our coil flow reactor (Table S5†) during the synthesis of E-F-HCP (15 minutes residence time) was estimated to be 65. This indicated turbulent mixing in our flow reactor. Clearly, the coiled reactor used in this work provided better mixing rates, enhancing polymerisation/crosslinking kinetics.<sup>20</sup>

The advantage of short reaction times in flow synthesis resulted in an increase in STY (Fig. 3). With flow synthesis, the STY value improved by 32 fold for internal crosslinking, 117 fold for post-crosslinking, and 68 fold for external crosslinking. This addressed the issue of low HCP productivity rates due to time constraints required in batch reactions that we have now overcome with flow synthesis.

### Chemical structures of flow-produced HCPs

The chemical structures of these HCPs were characterised and validated with FTIR and <sup>13</sup>C-NMR analyses. HCPs synthesised in batch reactors were used as control materials. The nomenclature for HCPs synthesised in this work is summarised in Table 1, alongside the type of reagents and operating conditions.

Comparing the FTIR spectra of HCPs synthesised by internal crosslinking in batch (I-B-HCP) and continuous flow (I-F-HCP) reactions, we observed that in both samples, the infrared peak intensities correlated to C–Cl vibrations<sup>27,43,44</sup> at 668 and 1260  $\text{cm}^{-1}$  and C–H bending<sup>44</sup> at 854  $\text{cm}^{-1}$  were reduced when compared to the unreacted DCX precursor (Fig. 4a). This was ascribed to chloromethyl substitution at the *para* position, suggesting that the chloro group was eliminated during Friedel–Crafts alkylation. The intensities of peaks correlated to the skeletal benzene rings, such as aromatic C–H stretching<sup>43</sup> at 2971 and 2866  $\text{cm}^{-1}$  and aromatic C=C stretching<sup>43,45</sup> at 1421, 1445, and 1512  $\text{cm}^{-1}$  were also reduced. This suggested the

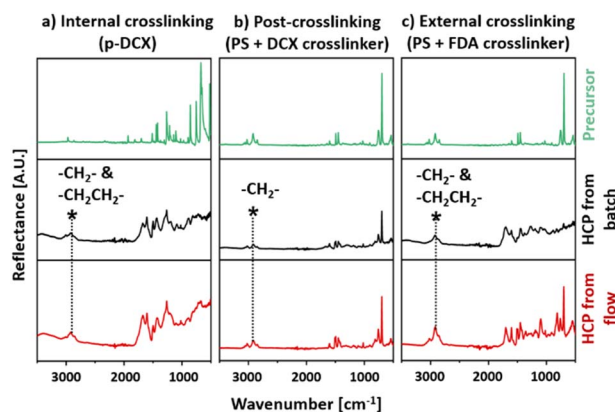


Fig. 4 FTIR spectra of the HCPs and their precursors: (a) internal crosslinking (p-DCX), (b) post-crosslinking (PS + DCX crosslinker), (c) external crosslinking (PS + FDA crosslinker).

success of DCX crosslinking.<sup>45</sup> At the same time, the peak corresponding to C–H stretching<sup>45</sup> at 2923  $\text{cm}^{-1}$  from HCP broadened when compared to that of DCX. This was due to the overlapping original C–H stretching peak with the newly formed methylene bridge ( $-\text{CH}_2\text{CH}_2-$ ) crosslinks.<sup>45</sup> This type of cross-linked bridge was formed from DCE molecules that functioned as both a solvent and a crosslinker reagent.<sup>46</sup>

The FTIR spectra of HCPs synthesised *via* post-crosslinking, P-B-HCP, and P-F-HCP (Fig. 4b), resembled that of PS. We observed peaks that corresponded to aromatic C–H stretching at 3087, 3061, 3027  $\text{cm}^{-1}$ , C–H bending at 756, 696  $\text{cm}^{-1}$ , aromatic C=C stretching at 1603, 1493, 1453  $\text{cm}^{-1}$ , C–H bending at 854  $\text{cm}^{-1}$  and backbone C–H stretching at 2921 and 2849  $\text{cm}^{-1}$ .<sup>28</sup> However, these peaks were less intense than those of PS. These indicated that the aromatic benzene rings of PS were modified during post-crosslinking. These trends were also presented in the FTIR spectra of HCPs synthesised *via* external crosslinking, E-B-HCP, and E-F-HCP (Fig. 4c). The broadened peak corresponding to C–H stretching<sup>45</sup> centred at 2921  $\text{cm}^{-1}$  similar to the spectra of internally crosslinked HCPs was observed in these spectra. This indicated that  $-\text{CH}_2\text{CH}_2-$  crosslinks were also formed in the externally crosslinked HCP. The presence of newly formed crosslinking methylene bridges in these HCPs was also validated with <sup>13</sup>C-NMR.

The <sup>13</sup>C-NMR spectra of I-B-HCP and I-F-HCP (Fig. 5a) contained major peaks centred at 138, 35, and 16 ppm. These peaks were correlated to the substituted aromatic carbon,<sup>47</sup> the crosslinked bridges from *via* FDA<sup>28,47</sup> and DCE,<sup>27</sup> respectively. There were also unreacted DCX<sup>24,48</sup> in these samples as we also observed peaks centred at 43 ppm (carbon in  $-\text{CH}_2\text{Cl}$ ) and 128 ppm (non-substituted aromatic ring). The spectra of P-B-HCP and P-F-HCP (Fig. 5b) also contained a peak centred at 146 ppm correlated to the quaternary non-substituted aromatic carbon<sup>48,49</sup> from the parent PS molecule. These spectra also contained peaks at 136 and 128 ppm, which corresponded to substituted aromatic carbon and non-substituted aromatic rings from both PS and DCX,

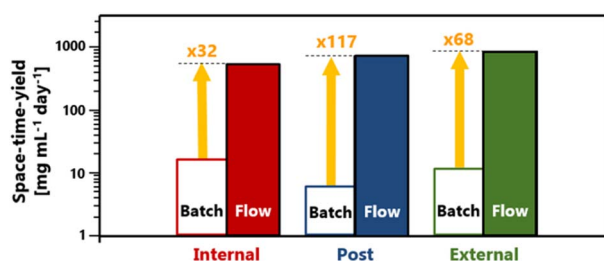


Fig. 3 STY of HCP from batch and flow production.



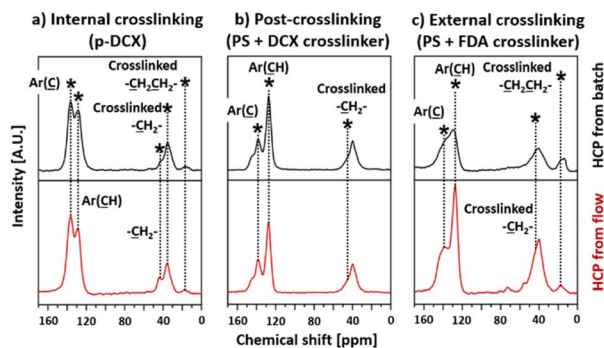


Fig. 5  $^{13}\text{C}$ -NMR spectra of the HCPs: (a) internal crosslinking (p-DCX), (b) post-crosslinking (PS + DCX crosslinker), (c) external crosslinking (PS + FDA crosslinker).

respectively. The broad peak centred at 40 ppm in these spectra was a mixture of signals attributed to aliphatic methylene carbon, methine backbone carbon, and methylene crosslink bridges.<sup>48</sup> We did not observe any peaks corresponding to chloromethyl carbon in these samples. The  $^{13}\text{C}$ -NMR spectra of E-B-HCP and E-F-HCP (Fig. 5c) were like those of HCPs produced from post-crosslinking.

### HCP microporosity

Porosity in HCPs is formed as a function of the crosslinking degree.<sup>50</sup> As the degree of crosslinking increases, the BET surface area increases, resulting from the creation of micropores (<2 nm).<sup>46,50</sup> However, as the micropores are excessively developed within the HCPs, this then results in the decline of BET surface area when a certain degree of crosslinking is achieved.<sup>1</sup> Here we observed that the BET surface areas of the flow-produced HCPs were in the range between 800–1160  $\text{m}^2 \text{g}^{-1}$  similar to reported batch-produced HCP counterparts from literature.<sup>24,27,28</sup> Regardless of crosslinking strategies, the BET surface areas were lower than those of HCPs synthesised in batch reactors by 1.5–9.6%, while the total pore volume of flow-produced HCPs was on average 21% lower (Table 2). However, the enhancement of micropore volume in the flow-

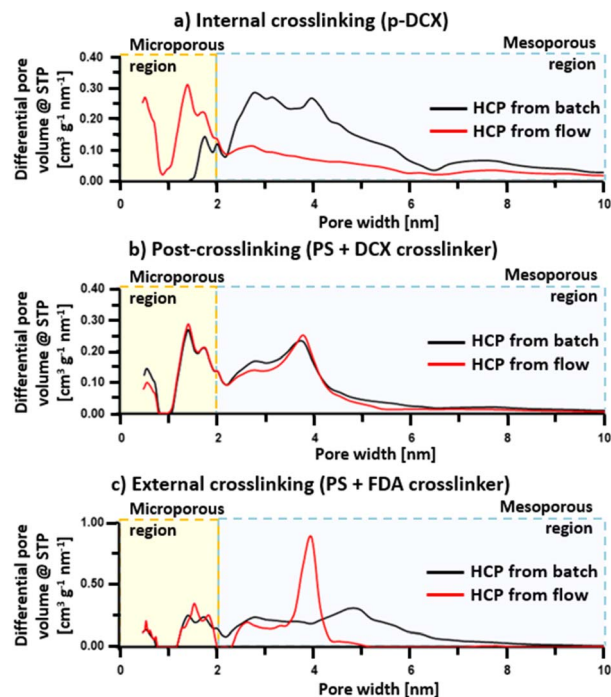


Fig. 6 Pore size distribution of flow-produced HCPs and their batch equivalents.

produced HCPs synthesised *via* internal and external crosslinking leads to the formation of narrower pores between 0–10 nm (Fig. 6). These results indicate that flow synthesis created more microporosity in such HCPs. This difference in micropore volume between HCP synthesised from batch and flow reactions could be attributed to different heat-mass transfer rates of reactor types, leading to different crosslinking rates and intensities. For batch synthesis, we used a 250 mL three-neck round-bottom glass flask as a reactor. Meanwhile, for flow synthesis, we used a 30 mL PTFE helical coil tube as a reactor. The smaller volume of the coiled tube induced faster mixing of reactants,<sup>36</sup> and enhanced mass transfer<sup>35</sup> when compared to the glass flask. The helical coil also increased the vortex flow pattern inside the tube.<sup>41</sup> Moreover, the contact area for transferring heat energy from the oil bath to the reaction mixture is larger in the flow reactor. The increment in microporosity in flow-produced HCP could potentially benefit applications, such as gas separation and storage.

The existence of micropores was confirmed through the steep slope at lower relative pressure, where  $P/P_0 < 0.05$  (Fig. S5†). Based on IUPAC classification,<sup>51</sup> the isotherms of internally crosslinked HCPs were a mixture of type I and IV. Moreover, the presence of macropores (>50 nm) in the internally crosslinked HCPs could be observed in the isotherm at  $P/P_0 > 0.9$ . It should be noted that this was the only HCP type in this study that formed macropores. The isotherms of HCPs synthesised by post-crosslinking and external crosslinking were classified as type IV, where ink bottle-like pore shapes were more prevalent. The hysteresis loops of these HCPs indicated

Table 2 Textural properties of each HCP sample<sup>a</sup>

Samples	BET surface area [ $\text{m}^2 \text{g}^{-1}$ ]	DFT total pore volume [ $\text{cm}^3 \text{g}^{-1}$ ]	DFT micropore volume [ $\text{cm}^3 \text{g}^{-1}$ ]
I-B-HCP	964	1.263	0.042
I-F-HCP	950 (−1.5%)	1.014 (−19.7%)	0.281 (+569%)
P-B-HCP	903	0.788	0.203
P-F-HCP	816 (−9.6%)	0.650 (−17.3%)	0.195 (−3.9%)
E-B-HCP	1162	1.219	0.184
E-F-HCP	1092 (−6.0%)	0.917 (−24.8%)	0.293 (+59.2%)

<sup>a</sup> DFT micropore volume data were obtained from the cumulative pore volume of the pore size up to 2 nm.



that there were significant amounts of small mesopores (2–10 nm) within these HCPs, which correlated to the broad peaks in the pore size distributions (Fig. 6).

The microporosity of HCPs studied here could be tailored by varying synthesis parameters: residence time, feed solution composition and concentration, and temperature. Here we observed that at a fixed substrate concentration, reagent ratios, and temperatures, an increase in residence time from 10 to 15 minutes typically enhanced the micropore volume of externally crosslinked HCPs by 595%. However, a further increase of residence time to 20 minutes reduced micropore volume by 65.5% (Table S3†). Similar trends were observed in batch reactions over 12 and 24 hour periods.<sup>1</sup> Prolonged reaction times could lead to excessive crosslinking, forming pores that are smaller than the kinetic diameter of N<sub>2</sub>, the gas used to probe porosity content during gas adsorption experiments. Another reason for an optimised micropore volume in HCPs produced *via* 15 minutes of residence time in flow reactors could be the mixing intensity. Our calculation (Table S5†) showed that with a residence time of 15 minutes, the Reynolds number of our reactor reached a value of 65. This was closest to the optimal Reynolds number of 40 in helical reactors.<sup>41</sup> Here it is important to highlight that the optimisation of HCP porosity by tailoring residence time did not drastically affect STY, *i.e.*, productivity rate. The highest STY value of 965 mg mL<sup>-1</sup> day<sup>-1</sup> was achieved with a residence time of 10 minutes. The increase of residence time to 15 minutes reduced this STY value by 28.3%, reaching 692 mg mL<sup>-1</sup> day<sup>-1</sup>.

The enhancements in both micropore volume and STY were observed (Table S3†) when the weight ratio of PS by FeCl<sub>3</sub> by FDA increased from 1 : 1 : 1 to 1 : 5 : 5 while fixing the other three synthesis parameters. No solid was formed at the weight ratio of 1 : 1 : 1. This indicates that crosslinking did not occur. Although a similar test in the batch system showed that the crosslinking using a 1 : 1 : 1 ratio of PS to FeCl<sub>3</sub> to FDA is achievable,<sup>28</sup> shorter residence time and lower temperature in flow reactors might be insufficient to induce crosslinking when there is a low ratio of catalyst and crosslinker to the substrate. HCP products were obtained when the weight ratio was increased to 1 : 3 : 3, with a STY of 981 mg mL<sup>-1</sup> day<sup>-1</sup>. However, the micropore volume of 0.006543 cm<sup>3</sup> g<sup>-1</sup> was the lowest among all the externally crosslinked HCPs. The increase in micropore volume and the STY when increasing the ratio from 1 : 3 : 3 to 1 : 5 : 5 was 1493% and 47.5%, respectively. Similar trends showing the improvement of the specific surface area, as a result of pore generation, and product yield were also observed in batch HCP synthesis.<sup>24,28</sup> The higher amount of the catalyst and crosslinker in the system indicated a higher possibility of the substrate getting involved in the reaction. Hence, the suitable weight ratio of substrate to the catalyst to crosslinker for flow synthesis in our system was 1 : 5 : 5.

Increasing the reaction temperature while fixing the residence time, substrate concentration, and chemical ratio did not enhance STY (average of 830 mg mL<sup>-1</sup> day<sup>-1</sup>), however, micropore volumes were enhanced. The increase in the temperature from 333 to 343 K increased the micropore volume by 84.8% from 0.059 to 0.109 cm<sup>3</sup> g<sup>-1</sup>. This indicated that

elevating temperature by a 10 K difference in our system affected the crosslinking degree in HCPs.

The impact of PS concentration on micropore volume was subtle compared with the other three synthetic parameters. The amount of micropore volume was increased by just 37.5% from 0.4% to 0.6% w/v of PS. Increasing the substrate concentration from 0.6 to 0.8% w/v and then to 1.0% w/v slightly increased and decreased micropore volume by 5.7% and 9.5%, respectively. This trend was also observed in batch synthesis,<sup>24,46</sup> which suggested that the fabrication of microporous polymers, *i.e.*, HCP is not sensitive to the monomer concentration as much as macroporous polymers.<sup>24</sup> On the other hand, the substrate concentration caused the largest influence on STY among all four synthetic parameters, as the correlation between PS concentration and STY was directly proportional. Increasing the PS concentration from 0.4% to 1.0% w/v enhanced STY by 3 fold from 457 to 1447 mg mL<sup>-1</sup> day<sup>-1</sup>.

With these optimal synthesis parameters, we were able to enhance the STY of HCP synthesis using flow reactions, as shown earlier in Fig. 3. This inferred that we were able to not only enhance the microporosity of the internally crosslinked HCPs but also improve the STY of such HCPs.

### CO<sub>2</sub>/N<sub>2</sub> separation

Due to higher total pore volumes in batch-produced HCPs (Table 2), the CO<sub>2</sub> and N<sub>2</sub> uptakes (Fig. 7) of such HCPs, regardless of crosslinking strategies, were higher than those synthesised under a continuous flow. Amongst all the HCPs studied here, the total pore volume of HCPs synthesised *via* internal crosslinking in batch reactions were the highest, reaching 1.26 cm<sup>3</sup> g<sup>-1</sup>. As such, the CO<sub>2</sub> and N<sub>2</sub> uptakes of HCPs synthesised *via* internal crosslinking reached 0.99 and

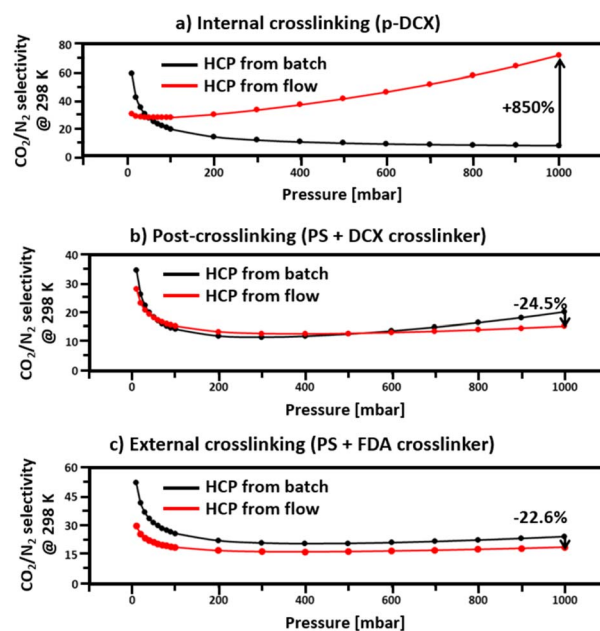


Fig. 7 CO<sub>2</sub>/N<sub>2</sub> selectivity at 298 K, 1 atm, 15 : 85 of CO<sub>2</sub> : N<sub>2</sub> inlet gas mixture.



0.27 mmol g<sup>-1</sup>, respectively. With a total pore volume of 1.22 cm<sup>3</sup> g<sup>-1</sup>, the CO<sub>2</sub> and N<sub>2</sub> uptakes of HCPs synthesised in batch reactions *via* external crosslinking reached 0.8 and 0.1 mmol g<sup>-1</sup>, respectively. These were 19% and 63% lower than those of I-B-HCP. With the lowest total pore volume of 0.79 cm<sup>3</sup> g<sup>-1</sup>, HCPs synthesised through post-crosslinking in batch reactions only uptake 0.55 and 0.05 mmol g<sup>-1</sup> of CO<sub>2</sub> and N<sub>2</sub>, respectively. Although continuous flow synthesis reduced the total pore volume of HCPs by 17–25% (Table 2), leading to lower CO<sub>2</sub> and N<sub>2</sub> uptakes, this approach enhanced the micropore volumes in internally and externally crosslinked HCPs by 569% and 59%, respectively. The 569% increase in the micropore volume in internally crosslinked HCPs underpinned a CO<sub>2</sub>/N<sub>2</sub> selectivity (at 298 K and 1000 mbar) of 71.5, 850% higher than that of batch-produced internally crosslinked HCPs, which was the lowest amongst all samples studied here, at 7.5 (Fig. 7). This significant difference in CO<sub>2</sub>/N<sub>2</sub> selectivity between the flow and the batch produced internally crosslinked HCPs could be attributed to the amount and type of pores present. With 25% more total pore volume, more CO<sub>2</sub> and N<sub>2</sub> could be adsorbed in I-B-HCPs. However, with only 3.2% of micropores, the ability to separate CO<sub>2</sub> from N<sub>2</sub> in I-B-HCPs is diminished. Due to the similar kinetic diameters of N<sub>2</sub> (3.64 Å) and CO<sub>2</sub> (3.30 Å),<sup>52</sup> materials with small pore sizes and narrow pore size distributions are required for effective CO<sub>2</sub>/N<sub>2</sub> separations, especially for post-carbon capture<sup>53</sup> and ideally, the pore sizes should be between 5 to 7.8 Å.<sup>53</sup> In this work, such pores were only observed in HCPs synthesised in flow reactions *via* internal crosslinking. Hence, the CO<sub>2</sub>/N<sub>2</sub> selectivity (calculated using IAST theory with 0.15 bar CO<sub>2</sub> partial pressure, 298 K and 1 atm) of these HCPs were the highest amongst all HCPs studied here. This outcome demonstrated that HCP CO<sub>2</sub>/N<sub>2</sub> selectivity could be improved through optimisation of the reaction parameters *via* reactor design, *i.e.*, from batch to flow reactor type. This is unique from existing strategies such as HCP functionalisation,<sup>21–23</sup> the use of bespoke monomers<sup>25,26</sup> and tailoring synthesis conditions (heat, duration, *etc.*).

## Conclusions

This study highlights two key benefits of the implementation of continuous flow synthesis in HCP production. First, we observed higher microporosity in HCPs synthesised in flow reactors. Regardless of their lower total pore volumes and BET surface areas, high volumes of micropores within flow-produced HCPs enhanced CO<sub>2</sub>/N<sub>2</sub>, reaching a value of 71.5. We also verified that the porosity can be tailored through the alteration of flow synthetic parameters. When comparing our optimised internally crosslinked HCPs with their batch counterpart, the HCP from flow synthesis showed an increase in micropore volume by 569% resulting in an 850% increase in CO<sub>2</sub>/N<sub>2</sub> separation. Second, the optimal residence time was reduced to 5 minutes with flow synthesis, instead of 24 hours as in a batch operation. This enhanced the productivity of STY by 32 fold, which resulted in a HCP production rate of 563 mg cm<sup>-3</sup> day<sup>-1</sup>. We also demonstrated the versatility of flow reactions for HCP synthesis *via* post-crosslinking and external

crosslinking approaches. The findings from this work show the potential to scale up HCP synthesis with flow reactors to meet the global demand for such materials, and the potential for carbon capture. The pilot scale of this process should be the next step, focusing on the optimisation of operating parameters.

## Author contributions

The study was proposed by C. H. L. The conceptualised idea was discussed between N. C. and C. H. L. The synthesis process was conducted by N. C. The contribution of N<sub>2</sub> gas analysis characterisation was shared between L. D. and A. J. F. FTIR characterisation was conducted by S. L. and S. S. <sup>13</sup>C-NMR characterisation was done by D. M. D. CO<sub>2</sub> and N<sub>2</sub> adsorption tests for CO<sub>2</sub>/N<sub>2</sub> separation were performed by K. K., and M. R. H. All the results were analysed by N. C. The paper was prepared by N. C. with supervision from C. H. L.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- 1 H. Liu, S. Li, H. Yang, S. Liu, L. Chen, Z. Tang, R. Fu and D. Wu, *Adv. Mater.*, 2017, **29**, 1700723.
- 2 Y. Luo, B. Li, W. Wang, K. Wu and B. Tan, *Adv. Mater.*, 2012, **24**, 5703–5707.
- 3 J. Huang and S. R. Turner, *Polym. Rev.*, 2018, **58**, 1–41.
- 4 V. A. Davankov and M. P. Tsyurupa, *React. Polym.*, 1990, **13**, 27–42.
- 5 M. P. Tsyurupa and V. A. Davankov, *React. Funct. Polym.*, 2006, **66**, 768–779.
- 6 C. H. Lau, X. Mulet, K. Konstas, C. M. Doherty, M.-A. Sani, F. Separovic, M. R. Hill and C. D. Wood, *Angew. Chem., Int. Ed.*, 2016, **55**, 1998–2001.
- 7 H. Ramezanipour Penchah, H. Ghanadzadeh Gilani and A. Ghaemi, *J. Chem. Eng. Data*, 2020, **65**, 4905–4913.
- 8 Materials and Chemicals - Ion Exchange Resin Market, *Ion Exchange Resin Market Study, By Type (Cationic Resins, Anionic Resins, Other), By End Use Industry (Power, Chemical and Petrochemical, Water and Wastewater Treatment, Food and Beverages, Pharmaceuticals, Electrical and Electronics, Metal and Mining), Forecasts To 2028*, 2021, available at: <https://www.reportsanddata.com/report-detail/ion-exchange-resin-market>, accessed: 9th July 2021.
- 9 L. Tan and B. Tan, *Chem. Soc. Rev.*, 2017, **46**, 3322–3356.
- 10 C. H. Lau, T.-d. Lu, S.-P. Sun, X. Chen, M. Carta and D. M. Dawson, *Chem. Commun.*, 2019, **55**, 8571–8574.



- 11 Y. Liu, X. Chen, X. Jia, X. Fan, B. Zhang, A. Zhang and Q. Zhang, *Ind. Eng. Chem. Res.*, 2018, **57**, 17259–17265.
- 12 T. Ratvijitvech, *J. Polym. Environ.*, 2020, **28**, 2211–2218.
- 13 F. Björnerbäck and N. Hedin, *ChemSusChem*, 2019, **12**, 839–847.
- 14 J.-S. M. Lee, T. Kurihara and S. Horike, *Chem. Mater.*, 2020, **32**, 7694–7702.
- 15 S. Grätz, S. Zink, H. Krafczyk, M. Rose and L. Borchardt, *Beilstein J. Org. Chem.*, 2019, **15**, 1154–1161.
- 16 N. B. McKeown, *Sci. China: Chem.*, 2017, **60**, 1023–1032.
- 17 M. Rubio-Martinez, T. D. Hadley, M. P. Batten, K. Constanti-Carey, T. Barton, D. Marley, A. Mönch, K.-S. Lim and M. R. Hill, *ChemSusChem*, 2016, **9**, 938–941.
- 18 Y. Peng, W. K. Wong, Z. Hu, Y. Cheng, D. Yuan, S. A. Khan and D. Zhao, *Chem. Mater.*, 2016, **28**, 5095–5101.
- 19 M. Rubio-Martinez, M. P. Batten, A. Polyzos, K.-C. Carey, J. I. Mardel, K.-S. Lim and M. R. Hill, *Sci. Rep.*, 2014, **4**, 5443.
- 20 P. J. Dowding, J. W. Goodwin and B. Vincent, *Colloid Polym. Sci.*, 2000, **278**, 346–351.
- 21 S. Krishnan and C. V. Suneesh, *J. Solid State Chem.*, 2021, **299**, 122152.
- 22 Y. Liu, X. Jia, J. Liu, X. Fan, B. Zhang, A. Zhang and Q. Zhang, *Appl. Organomet. Chem.*, 2019, **33**, e5025.
- 23 Y. Qiao, Z. Zhan, Y. Yang, M. Liu, Q. Huang, B. Tan, X. Ke and C. Wu, *Mater. Today Commun.*, 2021, **27**, 102338.
- 24 C. D. Wood, B. Tan, A. Trewin, H. Niu, D. Bradshaw, M. J. Rosseinsky, Y. Z. Khimyak, N. L. Campbell, R. Kirk, E. Stöckel and A. I. Cooper, *Chem. Mater.*, 2007, **19**, 2034–2048.
- 25 Q. B. Meng and J. Weber, *ChemSusChem*, 2014, **7**, 3312–3318.
- 26 M. Saleh, H. M. Lee, K. C. Kemp and K. S. Kim, *ACS Appl. Mater. Interfaces*, 2014, **6**, 7325–7333.
- 27 Z. Fu, I. M. A. Mohamed, J. Li and C. Liu, *J. Taiwan Inst. Chem. Eng.*, 2019, **97**, 381–388.
- 28 X. Dong, A. Akram, B. Comesaña-Gándara, X. Dong, Q. Ge, K. Wang, S.-P. Sun, B. Jin and C. H. Lau, *ACS Appl. Polym. Mater.*, 2020, **2**, 2586–2593.
- 29 L. Tan, B. Li, X. Yang, W. Wang and B. Tan, *Polymer*, 2015, **70**, 336–342.
- 30 V. Rajangam, A. Abidov, M. Peng, C. Babu, M. Palanichamy, W. Cha and H. Jang, *Fibers Polym.*, 2015, **16**, 1458–1467.
- 31 J. Wegner, S. Ceylan and A. Kirschning, *Chem. Commun.*, 2011, **47**, 4583–4592.
- 32 J. Rouquerol, P. Llewellyn and F. Rouquerol, *Stud. Surf. Sci. Catal.*, 2007, **160**, 49–56.
- 33 K. Shi, E. E. Santiso and K. E. Gubbins, in *Porous Materials: Theory and its Application for Environmental Remediation*, ed. J. C. Moreno-Piraján, L. Giraldo-Gutierrez and F. Gómez-Granados, Springer International Publishing, Cham, 2021, pp. 315–340, DOI: [10.1007/978-3-030-65991-2\\_12](https://doi.org/10.1007/978-3-030-65991-2_12).
- 34 A. Hassanpouryouzband, J. Yang, B. Tohidi, E. Chuvilin, V. Istomin, B. Bukhanov and A. Cheremisin, *Environ. Sci. Technol.*, 2018, **52**, 4324–4330.
- 35 R. L. Hartman, J. P. McMullen and K. F. Jensen, *Angew. Chem., Int. Ed. Engl.*, 2011, **50**, 7502–7519.
- 36 F. E. Valera, M. Quaranta, A. Moran, J. Blacker, A. Armstrong, J. T. Cabral and D. G. Blackmond, *Angew. Chem., Int. Ed.*, 2010, **49**, 2478–2485.
- 37 D. Wilms, J. Klos and H. Frey, *Macromol. Chem. Phys.*, 2008, **209**, 343–356.
- 38 Y. Cengel, *Heat and Mass Transfer: Fundamentals and Applications*, McGraw-Hill Higher Education, 2014.
- 39 D. M. Price and M. Jarratt, *Thermochim. Acta*, 2002, **392**, 231–236.
- 40 G. Tchobanoglous, F. L. Burton, H. D. Stensel and F. Burton, *Wastewater Engineering: Treatment and Reuse*, McGraw-Hill Education, 2003.
- 41 M. Mansour, P. Khot, D. Thévenin, K. D. P. Nigam and K. Zähringer, *Chem. Eng. Sci.*, 2020, **214**, 114522.
- 42 J. J. Derksen, M. S. Doelman and H. E. A. Van den Akker, *Exp. Fluids*, 1999, **27**, 522–532.
- 43 Z. Wei, Q. Chen and H. Liu, *New J. Chem.*, 2021, **45**, 11607–11617.
- 44 Y. Liu, X. Fan, X. Jia, B. Zhang, H. Zhang, A. Zhang and Q. Zhang, *J. Mater. Sci.*, 2016, **51**, 8579–8592.
- 45 L. Sun, K. Chai, L. Zhou, D. Liao and H. Ji, *Int. J. Biol. Macromol.*, 2021, **175**, 396–405.
- 46 Z. Fu, J. Jia, J. Li and C. Liu, *Chem. Eng. J.*, 2017, **323**, 557–564.
- 47 Z.-A. Qiao, S.-H. Chai, K. Nelson, Z. Bi, J. Chen, S. M. Mahurin, X. Zhu and S. Dai, *Nat. Commun.*, 2014, **5**, 3705.
- 48 R. Joseph, W. T. Ford, S. Zhang, M. P. Tsyurupa, A. V. Pastukhov and V. A. Davankov, *J. Polym. Sci., Part A: Polym. Chem.*, 1997, **35**, 695–701.
- 49 C. Dalla Valle, M. Zecca, F. Rastrelli, C. Tubaro and P. Centomo, *Polymers*, 2020, **12**, 600.
- 50 Q.-Q. Liu, L. Wang, A.-G. Xiao, H.-J. Yu and Q.-H. Tan, *Eur. Polym. J.*, 2008, **44**, 2516–2522.
- 51 M. Thommes, K. Kaneko, A. V. Neimark, J. P. Olivier, F. Rodriguez-Reinoso, J. Rouquerol and K. S. W. Sing, *Pure Appl. Chem.*, 2015, **87**, 1051–1069.
- 52 E. S. Kentish, A. C. Scholes and W. G. Stevens, *Recent Pat. Chem. Eng.*, 2008, **1**, 52–66.
- 53 L. Zou, Y. Sun, S. Che, X. Yang, X. Wang, M. Bosch, Q. Wang, H. Li, M. Smith, S. Yuan, Z. Perry and H.-C. Zhou, *Adv. Mater.*, 2017, **29**, 1700229.

