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# Diels–Alder cycloaddition polymerization for porous poly-phenylenes with exceptional gas uptake properties†

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**We report the synthesis of two-dimensional and three-dimensional porous polyphenylenes (2D/3D-pPPs) via the Diels–Alder cycloaddition polymerization reaction. The resulting 2D and 3D-pPPs showed surface areas up to 1553 m<sup>2</sup> g<sup>-1</sup>, pore volumes of 1.45 cm<sup>3</sup> g<sup>-1</sup> and very high H<sub>2</sub> uptake capacities of 7.4 and 7.1 wt% at 77 K, respectively, along with a competitive high-pressure CO<sub>2</sub> and CH<sub>4</sub> uptake performance.**

Diamond, renowned as the hardest known material, boasts remarkable physiochemical stability due to the fact that its carbon atoms are interconnected by robust covalent bonds.<sup>1</sup> In the pursuit of replicating its chemical structure, researchers have uncovered a class of materials known as porous aromatic frameworks (PAFs) as a subclass of porous organic polymers (POPs), which not only emulate the molecular arrangement of diamond but also exhibit exceptional porosity. PAFs represent a distinct category of porous materials, featuring covalently linked phenyl groups that endow them with ultrahigh thermal stabilities often exceeding 400 °C,<sup>2</sup> chemical robustness, and high surface areas.<sup>3</sup> For instance, PAF-1 exhibits a surface area of 5600 m<sup>2</sup> g<sup>-1</sup> along with a high thermal stability up to 450 °C and chemical robustness.<sup>2</sup> Despite their amorphous structure, PAFs exhibit significantly superior physiochemical stability. These features have led to their utilization in various applications, including gas storage and capture,<sup>2–5</sup> pollutant removal,<sup>4,6,7</sup> catalysis,<sup>4,8–10</sup> and sensing.<sup>4,8</sup> Furthermore, two-dimensional PAFs hold great promise in separation applications owing to their graphene-like structures.<sup>11,12</sup> Despite their promising features, PAFs have been predominantly synthesized through cross-coupling reactions, such as the Yamamoto-type Ullman,<sup>2</sup> Sonogashira–Hagihara,<sup>13</sup> and Suzuki–Miyaura<sup>14</sup> reactions, which require the use of either

costly precious-metals or other metals in stoichiometric amounts as catalysts.<sup>15–17</sup>

For example, the PAF-1 synthesis requires the use of 5.2 equivalents of the bis(1,5-cyclooctadiene)nickel(0) catalyst ([Ni(cod)<sub>2</sub>].<sup>2</sup> Moreover, the earlier iterations of PAFs synthesized by employing 3–6 equivalents of the same Ni catalyst exhibited varying surface areas.<sup>5,18,19</sup> In the cases of Sonogashira–Hagihara and Suzuki–Miyaura cross-coupling reactions, the most prevalent choice of catalysts are Pd(PPh<sub>3</sub>)<sub>4</sub>, and analogous Pd-based catalysts.<sup>7,13,20–22</sup> Efforts to synthesize PAFs using cost-effective catalysts, such as AlCl<sub>3</sub>, and FeCl<sub>3</sub>, *via* Friedel–Crafts<sup>23</sup> and Scholl reactions,<sup>24</sup> have been relatively limited. These catalysts, are also used in a significant excess, thus increasing the environmental impact of the process. Moreover, these alternative catalysts have yielded PAFs characterized by notably reduced thermal stability and complex chemical structures, arising from uncontrolled side reactions.<sup>23–25</sup> More recently, methanesulfonic acid has been employed both as a catalyst and a solvent in the synthesis of PAFs *via* an aldol triple condensation reaction<sup>26</sup> starting from acetylbenzenes. However, porous polymer networks (PPNs) prepared using this approach exhibited substantial oxygen contents. These results highlight the fact that PAF synthesis often relies on the use of either expensive precious-metal catalysts or leads to the formation of polymer networks with lower stability owing to incomplete polymerization and/or side reactions. In this direction, while the catalyst-free synthesis of linear polyphenylenes through the Diels–Alder cycloaddition polymerization reaction presents an interesting strategy,<sup>27</sup> the synthesis of porous polyphenylenes using this approach has yet to be realized. Here, we introduced a new strategy for the synthesis PAFs *via* Diels–Alder cycloaddition polymerization reaction by employing 5,5'-(1,4-phenylene)bis(2H-pyran-2-one) (Bispyrone) as a diene, and 1,3,5-triethynylbenzene (TEB) and tetrakis(4-ethynylphenyl)methane (TETPM) as dienophiles (Scheme 1). The resulting 2D and 3D porous polyphenylenes (pPPs) exhibited high surface areas of up to 1553 m<sup>2</sup> g<sup>-1</sup> with a pore volume of 1.45 cm<sup>3</sup> g<sup>-1</sup>, and a remarkable high-pressure H<sub>2</sub> uptake capacities of 7.4 and 7.1 wt% at 77 K, respectively.

Pyrones are recognized for their ability to undergo *in situ* diene formation *via* thermal CO<sub>2</sub> elimination, subsequently

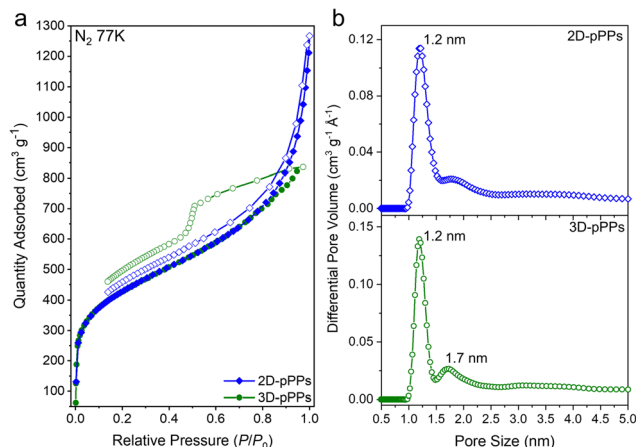
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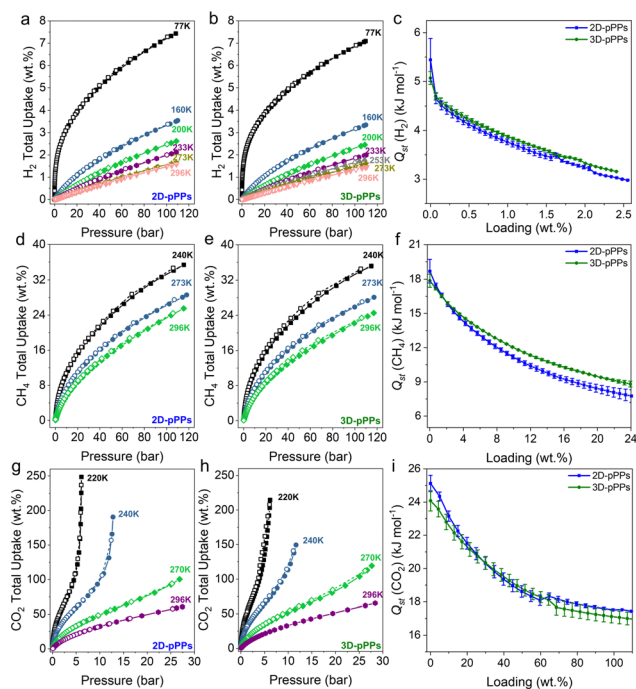


**Fig. 2** Porosity analysis of polyphenylenes, pPPs. (a)  $N_2$  uptake isotherms of 2D and 3D-pPPs measured at 77 K. Filled and empty symbols represent adsorption and desorption branches, respectively. (b) NLDFT pore size distribution of 2D-pPPs (top) and 3D-pPPs (bottom).

3D-pPPs exhibited distinct microparticles (Fig. S6, ESI†). The porosity of pPPs was examined by measuring  $N_2$  adsorption-desorption isotherms at 77 K (Fig. 2a). Both 2D- and 3D-pPPs exhibited a type II isotherm, a characteristic typically observed in carbon materials featuring a combination of micro and mesoporosity. The emergence of mesopores was expected due to the length of the employed organic linkers. However, as the linker size increases, the interpenetration of the polymer networks has been commonly observed, which results in the formation of smaller pores and the co-existence of both micro- and mesopores in the pPPs. While 2D-pPPs displayed a small hysteresis in the desorption branch, the 3D-pPPs exhibited a distinctive H2 type hysteresis loop, indicative of the presence of ink-bottle-shaped pores (see Fig. 2a). In order to assess the surface area, the Brunauer-Emmett-Teller (BET) theory was employed within the specified pressure range, determined from the Rouquerol plots (Fig. S7 and Table S4, ESI†). The 2D- and 3D-pPPs exhibited similar BET surface areas of 1553 and 1536  $m^2 g^{-1}$ , respectively. Notably, 2D-pPP displayed a higher pore volume of 1.45  $cm^3 g^{-1}$  in contrast to the 1.27  $cm^3 g^{-1}$  observed in 3D-pPPs (Table S4, ESI†). This difference can be attributed to the reduced accessibility of pores within 3D-pPPs arising from the interpenetration of the polymer network, as is commonly observed for the 3D-POPs and COFs.<sup>29</sup> The pore size distribution (PSD) of the polymers was calculated by using non-local density functional theory (NLDFT). PSD plots of both porous polyphenylenes exhibited micropores with an average pore size of 1.2 nm, which agrees well with the significant  $N_2$  uptake in the low-pressure range (Fig. 2b). The evaluation of mesopores was performed by employing the Barrett-Joyner-Halenda (BJH) model, which indicated the existence of mesopores in both polymers (Fig. S8, ESI†).

The high surface areas and pore volumes of 2D/3D-pPPs prompted us to investigate their high-pressure small gas uptake performance. Both 2D- and 3D-pPPs showed very high  $H_2$  uptake capacities of 7.4% and 7.1%, respectively, at 77 K, 110

bar (Fig. 3a and b). These results position 2D and 3D-pPPs as highly promising materials when compared to other porous polymers such as PAFs and PPNs, especially considering their higher surface areas (Fig. S9 and Table S5, ESI†). This high  $H_2$  uptake capacity is attributed to the combined effect of the large abundance of micropores<sup>30</sup> and high BET surface area. Whereas all the previously reported PAFs showed a linear correlation between their surface areas and  $H_2$  uptake capacities, both 2D/3D-pPPs showed a significant deviation from this and outperformed their counterparts with similar surface areas (Fig. S9, ESI†). 3D-pPPs exhibited slightly higher  $H_2$  excess uptake of 2.9 wt% compared to 2D-pPPs' 2.7 wt% at the same temperature (Fig. S10, ESI†). This difference in total uptake can be attributed to the varying pore volumes in the two structures, while the higher excess uptakes in 3D-pPPs could be due to the swelling of the polymer network.<sup>31</sup> The difference in gas uptake between the two structures becomes less significant at high temperatures (Fig. 3a and b and Table S7, ESI†). Additionally, both 2D and 3D-pPPs exhibited very similar  $H_2$  heats of adsorption ( $Q_{st}$ ) values. The 2D-pPPs showed a  $H_2$   $Q_{st}$  value of 6.6  $kJ mol^{-1}$  at zero coverage, while 3D-pPPs had a slightly lower  $Q_{st}$  of 6.3  $kJ mol^{-1}$  (Fig. 3c). We also investigated the high-pressure  $CH_4$  uptake performance of both polymers. 2D and 3D-pPPs exhibited good high-pressure  $CH_4$  uptake capacities of 25.6 and 24.6 wt%, respectively, at 296 K (Fig. 3d and e). The presence of an aromatic backbone along with the high pore volumes of both polymers are likely to contribute to the high affinity of towards  $CH_4$ .<sup>32</sup> The heats of adsorption for  $CH_4$  were



**Fig. 3** High pressure gas uptake performance of polyphenylenes.  $H_2$  uptake isotherms obtained at different temperatures for (a) 2D-pPPs and (b) 3D-pPPs.  $CH_4$  uptake isotherms obtained at different temperatures for (d) 2D-pPPs and (e) 3D-pPPs.  $CO_2$  uptake isotherms obtained at different temperatures for (g) 2D-pPPs and (h) 3D-pPPs. The heat of adsorption ( $Q_{st}$ ) for 2D and 3D-pPPs for (c)  $H_2$ , (f)  $CH_4$  and (i)  $CO_2$ .



18.7 and 17.9 kJ mol<sup>-1</sup> for 2D- and 3D-pPPs, respectively, indicative of a physisorption mechanism (Fig. 3f). 2D-pPPs consistently demonstrated higher total uptake, while 3D-pPPs displayed greater excess uptake (Fig. S10 and Table S7, ESI†).

Finally, considering the substantial micropore content, we also explored the CO<sub>2</sub> uptake performance of 2D- and 3D-pPPs. 2D-pPPs exhibited an impressive CO<sub>2</sub> uptake capacity of 248.6 wt% at 220 K, 25 bar, while the 3D-pPPs showed 214.5 wt% (Fig. 3g and h). Notably, 3D-pPPs displayed a higher uptake of 119.3 wt% at 270 K and 65.2 wt% at 296 K, in contrast to 2D-pPPs, which showed uptake capacities of 100.4 wt% and 60.7 wt%, respectively (Fig. 3g and h). These values align favorably with those reported for other PAFs and POPs, following the trend of CO<sub>2</sub> and CH<sub>4</sub> uptake in relation to surface area (Fig. S9, S10 and Table S6, ESI†). Furthermore, the heats of adsorption for CO<sub>2</sub> for both 2D and 3D-pPPs were found to be 25.1 and 24.1 kJ mol<sup>-1</sup>, respectively. These values position polyphenylenes as potential candidates for CO<sub>2</sub> storage (Fig. 3i).<sup>33</sup>

In conclusion, we introduced a new approach for the synthesis of porous polyphenylenes through Diels–Alder cycloaddition polymerization reaction, effectively eliminating the need for metal catalysts in the preparation of carbon-based frameworks. This approach yielded porous polyphenylenes that exhibit exceptional H<sub>2</sub> uptake capacities along with good CH<sub>4</sub> and CO<sub>2</sub> uptake performance. In a broader context, these results will stimulate the development of environmentally friendly approaches for the synthesis of porous organic polymers.

T. A. synthesized the precursors and polymers and conducted all characterizations. P. W. F. assisted in synthesizing the precursors and characterizing the polymers. T. Y. performed high-pressure gas uptake measurements. A. C. secured funding, supervised the project, and co-wrote the paper with all the authors.

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## Conflicts of interest

There are no conflicts to declare.

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