A “click-based” porous organic polymer from tetrahedral building blocks†

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A thermally and chemically stable “click-based” porous organic polymer with high surface area was synthesized from two tetrahedral building blocks.

Highly crosslinked, microporous organic polymers1–7 have recently received significant attention as potential materials for a variety of applications ranging from gas storage8–10 and gas separation,11,12 to chemical catalysis,13 chemical sensing,14 and capture of improvised chemical threats such as ammonia.15 Depending on the design, synthetic strategy, and properties, these materials have been classified into several subsets including amorphous hyper-crosslinked polymers (HCPs),14,17 polymers of intrinsic microporosity (PIMs),12,18 conjugated microporous polymers (CMPs),19 porous organic polymers (POPs),11,20 porous organic frameworks (POFs),21 and (crystalline) covalent organic frameworks (COFs).1 Regardless of the degree of crystallinity, the pore properties of these materials can often be modulated based on the choice of the building blocks. Notably, organic materials with high surface areas can be made from all-planar building blocks22 as well as from combinations of non-planar “struts”.1,11 Herein, we report the synthesis of a highly stable, microporous “click-based” porous organic polymer (POP) via copper-catalyzed alkyne–azide coupling (CuAAC) between two rigid tetrahedral building blocks (Td + Td).22

Alkyne–azide “click” chemistry has been widely employed in the synthesis of both linear and crosslinked polymers with diverse applications ranging from drug delivery23–28 to electronics.26,27 Our groups have applied this strategy in the modification of porous materials such as MOFs29,30 and nanobins.30 Given the aromatic, rigid nature of the 1,2,3-triazole group that results from alkyne–azide coupling, we reasoned that highly crosslinked organic polymers synthesized using this strategy would possess permanent microporosity and be attractive materials for gas storage. In addition, the high stability of the 1,2,3-triazole linkage could give “click-based” POPs thermal- and chemical-stability advantages over other polymers obtained via alternative bond-construction strategies.

The CuAAC reaction between tetrakis(4-ethynylphenyl)methane (C1) and tetrakis(4-azidophenyl)methane (D1) (Scheme 1) occurs readily in dimethylformamide (DMF) at both room temperature and 100 °C to give several versions of POP C1–D1 as brown powders. Since the surface area of the POP synthesized at 100 °C (1260 m² g⁻¹) was much higher than that for the version synthesized at 25 °C (440 m² g⁻¹), further optimization of the synthesis was carried out at the higher temperature (Table S1, entries 3 and 6 and Fig. S2 in the ESI†). The yield for POP C1–D1 (Table 1, entry 1a) was almost quantitative, suggesting that crosslinking was nearly complete with no soluble oligomers. However, care must be taken in isolating the product as a fraction in fine particulates can be lost if the filtration is carried out over filter paper (Table 1, entries 1b–5, see also Table S1 entries 1b–6 in the ESI†). Loss of this component during the filtration process can result in a slight decrease of the specific surface area of the overall product (1360 m² g⁻¹ vs. 1440 m² g⁻¹, Table 1 cf. entries 1a and 1b) but does not significantly affect the micropore and total pore volumes.

Because the progress of CuAAC depends on the availability of CuI intermediates, which in turn depends on the amount of sodium ascorbate reducing agent, we optimized the porosity of the product POP C1–D1 by adjusting the amount of sodium ascorbate between 10 and 70 mol%, while keeping the amount of copper catalyst constant (10 mol%). Surprisingly, the surface area of POPs C1–D1 systematically decreased with increasing amount of sodium ascorbate in the reaction mixture (Table 1 and Fig. S1 in the ESI†). Treatment of these POPs via Soxhlet extraction (to remove any sodium ascorbate possibly present in the pores) did not result in improvement in the original surface areas (Table S2 in the ESI†). Thus, we attribute the aforementioned ascorbate-induced lowering of surface area to an increase in catalyst availability: at higher loadings of ascorbate more CuI is available, allowing for more crosslinking to occur within the porous network and thus lowering surface areas. This hypothesis is supported by a progressive decrease in micropore volume and total pore volume as the amount of ascorbate is increased.

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The FTIR spectrum of POP C1–D1 (Table 1, entry 1b) compares well with that of a model compound (MC, Scheme 1). A broad N==N stretch can be observed at 1610 cm⁻¹ (Fig. 1), together with a weak C==CH band at 3029 cm⁻¹, consistent with the formation of the expected triazole ring. However, IR stretches from unreacted azide could still be observed at 2114 cm⁻¹, suggesting that the azide–alkyne coupling is incomplete. Interestingly, a strong and broad stretch at ~3300 cm⁻¹ is also observed, suggesting the presence of significant amounts of physisorbed water. The solid state ¹³C CPMAS NMR spectrum of POP C1–D1 (Table 1, entry 1b) matches well with that of MC, further supporting the formation of cyclic triazole linkages (Fig. 2).

Thermogravimetric analysis (TGA) of dried POP C1–D1 shows that this microporous material is thermally stable, losing only 20 wt% of its mass below 500 °C (Fig. S8 in the ESI†). The thermal stability of this uncapped material approaches those of the most stable metal-organic framework analogues containing carborane or Zr building blocks. The powder X-ray diffraction spectrum of POP C1–D1 is featureless, implying (as expected) that it is amorphous.

Consistent with the anticipated chemical stability of the 1,2,3-triazole linkage, POP C1–D1 is remarkably stable. Immersing C1–D1 in 6 N NaOH (aq) or 6 N HCl (aq) at 60 °C for 8 h leads to essentially no change in internal surface area (Table S3 in the ESI†). Increasing the immersion temperature to 80 °C and time to 24 h leads to only slight decreases (10–12%) in surface area (Table S3 in the ESI†). This acid- and base-stability rivals that of zeolitic imidazolate frameworks and greatly exceeds that of imine-linked POPs having comparable initial surface areas; the surface areas of the latter rapidly decrease following exposure to water. Its chemical stability is also superior to those of diimide-linked POPs, which are stable in relatively dilute aqueous acids (0.1 N) but not in bases.

High uptake of nitrogen at low pressure, as observed in the N₂ adsorption-desorption isotherms for the various preparations of POP C1–D1 (Fig. 3, see also Fig. S1 in the ESI†), suggests that these materials are mainly microporous. However, the isotherms of POPs prepared with smaller amounts of sodium ascorbate (10–30 mol%) tend to exhibit more hysteresis than those prepared with larger amounts (50–70 mol%). The slopes of the isotherm adsorption branches for the latter materials are also less steep than those for the former. Particularly interesting is the presence of a pronounced

<table>
<thead>
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<th>Entry</th>
<th>Sodium ascorbate (mol%)</th>
<th>Specific surface area (m² g⁻¹)</th>
<th>Micropore volume (cm³ g⁻¹)</th>
<th>Total pore volume (cm³ g⁻¹)</th>
<th>Yield (%)</th>
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<tr>
<td>1a</td>
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<td>0.56</td>
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<td>&gt;99</td>
</tr>
<tr>
<td>1b</td>
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<td>0.53</td>
<td>0.74</td>
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<td>1320</td>
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<td>0.73</td>
<td>62</td>
</tr>
<tr>
<td>3</td>
<td>30</td>
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<td>0.49</td>
<td>0.67</td>
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<tr>
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<tr>
<td>5</td>
<td>70</td>
<td>1090</td>
<td>0.42</td>
<td>0.52</td>
<td>77</td>
</tr>
</tbody>
</table>

a All reactions were carried out in DMF at 100 °C using 10 mol% CuSO₄ $\cdot$ 5H₂O and at 0.04 M solution of azide functional group (1 : 1 alkyne-to-azide functional group ratio). b Obtained using workup procedure II. c Obtained using workup procedure I.

![Fig. 1](link) FTIR spectra of the starting materials (C1 and D1), POP C1–D1 (Table 1, entry 1b), and model compound (MC).

![Fig. 2](link) ¹³C CPMAS NMR spectra of the model compound (MC) and POP C1–D1 (Table 1, entry 1b).

![Fig. 3](link) N₂ adsorption-desorption isotherms at 77 K for two versions of POP C1–D1 (Table 1, entries 1b and 5). A complete set of isotherms for all samples can be found in Fig. S1 in the ESI†.
hysteretic step in the N$_2$ isotherm desorption branch for the POPs prepared with low amounts (10–30 mol%), Table 1, entries 1a–3) of sodium ascorbate (Fig. 3, see also Fig. S1 in the ESI†). In contrast, there is no hysteretic step in the N$_2$ isotherm desorption branch for the POPs prepared by using higher amounts (50–70 mol%), Table 1, entries 4 and 5) of sodium ascorbate (Fig. 3, see also Fig. S1 in the ESI†).

For the samples of POP C1–D1 whose isotherms contain no hysteretic desorption step (Fig. 3 and Table 1, entries 4 and 5), the parallel, almost-horizontal nature of the two branches of the isotherm implies the presence of only narrow slit-like pores.\(^{37,38}\) In contrast, the samples whose isotherms contain a hysteretic desorption step (Fig. 3 and Table 1, entries 1a–3) likely possess more complex pore geometry.\(^{37,38}\) The near-horizontal desorption behavior in the 0.5 < $P/P_\theta$ < 1 region and the abrupt switch into a slightly steeper desorption behavior in the $P/P_\theta$ < 0.5 region (for N$_2$ adsorption at 77 K, this switch often occurs near the $P/P_\theta = 0.42$ point\(^{\text{35}}\)) indicate that the pores are not all slit-like. The upward-sloping trend in the adsorption branch of these isotherms, which becomes steeper at higher pressures, has been attributed by others to the presence of pores of different shapes and sizes (mainly microporous with some higher micropores and lower mesopores).\(^{37}\)

Despite the aforementioned indications of complex pore geometry, non-local density functional theory (NLDFT) pore size distribution analysis of POP C1–D1 prepared using 10 mol% sodium ascorbate (Table 1, entry 1b) indicates a mainly microporous material with a narrow pore-size distribution centered around a primary pore width of 9.2 Å (Fig. 4). This width is much smaller than expected (~21 Å) for an idealized diamond-like single network formed by simply “clicking” together monomers C1 and D1. This disparity constitutes strong evidence for interpenetration in the hyper-crosslinked networks constituting POP C1–D1.

Given the high surface area and narrow pore-size distribution of POP C1–D1, we explored its potential to adsorb the gases CO$_2$ and H$_2$. The NLDFT surface area obtained for POP C1–D1 (Table 1, entry 1a and Fig. S7 in the ESI†) based upon adsorption of CO$_2$ at 273 K is 1080 m$^2$ g$^{-1}$. This POP has a reasonably good H$_2$ adsorption capacity at 77 K and 1 bar (1.6 wt%, Fig. 5a), comparable to a range of MOFs,\(^{39}\) carbon materials,\(^{40,41}\) and microporous polymers\(^{42–46}\) featuring similar or higher surface areas. The isosteric heat of hydrogen adsorption for this material, which is an approximate measure of the strength of its interaction with H$_2$, was found to be 7.9 kJ mol$^{-1}$ at low surface coverage, decreasing to 4.0 kJ mol$^{-1}$ at 1 bar (Fig. 5b). These values are typical for H$_2$ adsorption in micropores of materials that can offer only London dispersion-type interactions.

In summary, we have shown that a permanently microporous POP can be assembled readily from a complementary pair of tetrahedral building blocks (T$_A$ + T$_D$) using CuAAC chemistry. The resulting material is thermally stable, as well as chemically stable, easily surviving exposure to highly acidic and basic solutions. The excellent stability of POP C1–D1 suggests more generally that cyclic-triazole-linked POP materials can serve as excellent engineering materials under a wide range of conditions. At 1 bar, the micropores of this material have reasonably good ambient-temperature adsorption of CO$_2$ and cryogenic adsorption of H$_2$. We are actively exploring structure-property relationships for these types of materials and efforts to turn them into high-surface-area and morphologically well-defined nano- and microparticle structures, similar to infinite coordination polymer particle analogues.\(^{42–46}\) Such capabilities will allow researchers to tune the properties of this novel class of materials for optimal performance in many practical applications, including gas storage, catalysis, and biological probe design.

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**Notes and references**


![Fig. 4](image-url) Pore size distribution plot for POP C1–D1 (Table 1, entry 1b) according to NLDFT analysis.

![Fig. 5](image-url) (a) H$_2$ adsorption–desorption isotherm for POP C1–D1 (Table 1, entry 1a) at 77 K, up to ca. 1 bar. (b) Isosteric heat of H$_2$ adsorption of POP C1–D1 (Table 1, entry 1a). Measurements were carried out at 77 K and 87 K.

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As the manuscript by Holst et al. appeared on the use of tetrahedral monomers to create conjugated microporous polymers (CMPs) focused primarily on CMPs derived from metal-catalyzed coupling, which was "click" chemistry. ...