A hybrid porous material from a pillar[5]arene and a poly(ionic liquid): selective adsorption of n-alkylene diols†

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† Electronic supplementary information (ESI) available: Synthetic procedures, characterizations, crystal data. See DOI: 10.1039/c3cc49108k

Poly(ionic liquid)s (PILs) are polyelectrolytes composed of ionic liquid (IL) repeating units.1 Their physical properties can be easily and broadly adjusted simply by counterion exchange. This unique merit has enabled a multitude of applications in many fields.2 Recently, we created a simple solution method to prepare porous polymer materials via in situ ionic complexation of a PIL, poly-(3-cyanomethyl-1-vinylimidazolium bis(trifluoromethanesulfonyl)imidie) (2 in Fig. 1, PCMVImTf2N) and multivalent carboxylic acids under alkaline conditions.3 These porous PIL complex networks with satisfactory specific surface area, large pore volume and stable pore structures have already led to applications in heterogeneous catalysis3a and adsorption.3b Here we successfully prepared a porous material 3 with a specific surface area of up to 350 m² g⁻¹ based on 2 and a fully carboxylic acid-substituted pillar[5]arene 1. This porous material preserved the solution-state host–guest properties of the pillar[5]arene in the solid state and exhibited unique selectivity in adsorption of n-alkylene diols.

To prepare the pillar[5]arene-containing porous material 3, 10 wt% of 2 and 1 (equimolar carboxylic acid and imidazolium units) were mixed in DMSO first, forming a homogeneous solution (Fig. 1). They did not form cross-linked networks by ionic bonding because carboxylic acid groups were electroneutral. When the solution was added dropwise into isopropanol containing 0.5 wt% of ammonia...
under stirring and sonication, the COOH groups were converted into carboxylate anions via deprotonation and initiated the in situ ionic complexation between negatively charged 1 and positively charged 2. It immediately produced the insoluble pillar[5]arene-containing poly(ionic liquid) complex material 3. The precipitate was collected by centrifugation, washed three times, and dried at 50 °C under high vacuum for 12 h. We used a cross-linked porous poly(ionic liquid) complex material 4 based on 1,2,4,5-benzenetetracarboxylic acid and 2 as a control material (Scheme S3, ESI†).

Fourier transform infrared (FTIR) spectra of 1 and 3 (Fig. S3a, ESI†) revealed that the adsorption band at 1710 cm⁻¹ attributed to the C═O bonds of the carboxylic acid groups on 1 shifted to 1590 cm⁻¹ upon formation of carboxylate anions in 3. X-ray diffraction measurements of 1 and 3 (Fig. S3b, ESI†) were performed to characterize the solid structures. The diffraction curve of 3 has lost the sharp characteristic crystalline peaks of 1 and is typical for a slightly ordered, amorphous material. These findings indicated that most of the carboxylic acid groups were converted into carboxylate anions in 3, a noncrystalline solid that is typical for polymeric complexes formed by electrostatic complexation.¹⁴ Scanning electron microscopy (SEM) was employed to analyze the morphology of 3 (Fig. 1 and Fig. S4, ESI†). Nano-sized particles (20–50 nm) as secondary tectonic units are observed. The randomly packed nanoparticles possibly provided the interstitial pore system as large transport channels which increase the mass-transfer efficiency and the specific surface area for heterogeneous binding events.

To further investigate the porous characteristics of 1, 2 and 3, nitrogen sorption measurements were performed (Fig. 2). The isotherms show that the Brunauer–Emmett–Teller (BET) specific surface area and the pore volume of 3 are 350 m² g⁻¹ and 1.22 cm³ g⁻¹, respectively. This indicated that the material is highly porous. The shape of the adsorption and desorption branches showed that the porosity of 3 is predominantly surface porosity with large pores in the macropore regime. The latter pores are the lower tail of the interstitial pores. The larger transport pores and the pores inside pillar[5]arene are not accessible by nitrogen sorption. (Fig. S9, ESI†). We found that neither powder 2 nor the fully carboxylic acid-substituted pillar[5]arene 1 shows any detectable porosity themselves (Fig. 2). It is an interesting question if nitrogen sorption really can detect the ca. 0.5 nm sized inner pores of the pillararenes. The sorption isotherm shows no or only very weak indications for those pores, i.e. they are presumably still filled with solvent, which can be tightly bound to those cavities.

Due to the organic character of the sorption material, vigorous high temperature drying (>150 °C) was not applied.

Based on the reported studies,²⁺²⁺₂⁺ the electron rich cavity of the pillar[5]arene can complex neutral guests with linear alkyll groups in solution driven by multiple C-H···π interactions. For example, dibutylpillar[5]arene includes n-hexane into its cavity to form a pseudorotaxane-type threaded structure in the solid state (Fig. S6a, ESI†). Here, in addition to the C-H···π interactions, two hydrogen bonding donors at the two ends of the guest can be added to strengthen the pillararene–guest interactions (Fig. S6b, ESI†). Therefore we studied the host–guest binding of a fully carboxylate anion–functionalized pillar[5]arene 5 and 1,6-hexanediol in D₂O (Fig. S7, ESI†). It is found that all signals from 1,6-hexanediol disappeared, caused by the strong complex-induced increase in relaxation times. Peaks assigned to the aromatic protons of 5 shifted significantly downfield.

Fig. 2 Nitrogen sorption isotherms of 1, 2 and 3. All samples were degassed at 80 °C for 20 h before measurements.

The affinity of the porous material 3 towards different guests was further tested the adsorption capability for 1,6-hexanediol by using equivalent amounts of 1, 2, 3 and 4 (Fig. S10, ESI†). We found that neither powder 2 nor the porous reference material 4 without pillar[5]arene cavities adsorbed 1,6-hexanediol in the CDCl₃ solution. Both 1 and 3 containing the pillar[5]arene were active. The adsorption capacity of 3 is in fact about 4.5 times higher than that of equivalent 1, which we attribute to the improved access to the pores in the amorphous superstructure as compared to the ground crystalline powder. This means that it was indeed successful in heterogenizing the ability for molecular recognition of pillar[5]arene from the solution to the solid state.

Using the same method, the adsorption kinetics and maximum adsorption capacities of 3 with 1,6-hexanediol were analyzed (Fig. S11, ESI†). The affinity of the porous material 3 towards different guests was analyzed from their CDCl₃ solutions using the above-described method (Fig. 3). The results showed that 3 has the highest affinity to 1,4-butanediol. This agrees well with the work of Li et al. who have revealed that a symmetric guest which contains a linear alkyl chain with four methylenes usually shows the highest binding constant.
with pillar[5]arenes relative to similar compounds with longer or shorter allyl chains.\(^{2,9,13}\) Interestingly, the maximum adsorption capacity of 1,4-cyclohexanediol is only slightly smaller than that of 1,6-hexanediol. The relative binding of 2,3-butanediol is half that of 1,4-butanediol, which is a significant selectivity, but also still a reasonable binding at the same time. More relevant, the porous material 3 showed only a very weak affinity for 1-butanol and 1-hexanol. Finally, 3 cannot adsorb cyclohexanol and tert-butanol at all, because they stretch the host–guest rules too much.

In conclusion, a macro-/mesoporous polymeric material with homogeneous pillar[5]arene cavities was prepared by \textit{in situ} ionic condensation between a negatively charged pillar[5]arene and a positively charged poly(ionic) liquid in organic solvents. SEM and XRD results showed that this noncrystalline porous material is composed of secondary nanoparticles, about 20–50 nm in size, which form a sponge-like superstructure with interstitial meso- and macro- pores improving liquid transport. Nitrogen sorption measurements revealed a specific surface area of 350 m\(^2\) g\(^{-1}\), and a pore volume of 1.22 cc g\(^{-1}\). On top of this secondary porosity, the pillar[5]arene cavities provide 3 with designated affinities with certain substrates. Binding experiments revealed that at least 80% of the primary pores are accessible from the outside and are not blocked. The porous hybrid material can for instance adsorb 1,6-hexanediol from CDCl\(_3\), where a reference complex without supramolecular pores cannot. Variation of the substrate reveals good binding to the family of aliphatic diols, with the highest affinity to 1,4-butanediol. Considering the broad family of pillararenes, the present study not only provides a useful method to introduce host–guest property of macrocycles into solid-state materials but also offers in a more general frame base for the construction of functional materials with controllable affinity to designated substrates.

This work was supported by National Basic Research Program (2013CB834502), the National Natural Science Foundation of China (21125417), the Fundamental Research Funds for the Central Universities and Alexander von Humboldt Fellowship.

Notes and references


