

RSC Advances



This is an *Accepted Manuscript*, which has been through the Royal Society of Chemistry peer review process and has been accepted for publication.

Accepted Manuscripts are published online shortly after acceptance, before technical editing, formatting and proof reading. Using this free service, authors can make their results available to the community, in citable form, before we publish the edited article. This *Accepted Manuscript* will be replaced by the edited, formatted and paginated article as soon as this is available.

You can find more information about *Accepted Manuscripts* in the [Information for Authors](#).

Please note that technical editing may introduce minor changes to the text and/or graphics, which may alter content. The journal's standard [Terms & Conditions](#) and the [Ethical guidelines](#) still apply. In no event shall the Royal Society of Chemistry be held responsible for any errors or omissions in this *Accepted Manuscript* or any consequences arising from the use of any information it contains.

ARTICLE

Novel Porous materials based on Oligospiroketal (OSK)

Pablo Wessig,^{*a} Maik Gerngroß,^a Simon Pape,^a Philipp Bruhns^a and Jens Weber^b

Received (in XXX, XXX) Xth XXXXXXXXXX 20XX, Accepted Xth XXXXXXXXXX 20XX

DOI: 10.1039/b000000x

Abstract

New porous materials based on covalently connected monomers are presented. The key step of the synthesis is an acetalisation reaction. In the last years we used acetalisation reactions extensively to build up various molecular rods. Based on this approach, investigations towards porous polymeric materials were conducted by us. Here we wish to present the results of these studies in the synthesis of 1D polyacetals and porous 3D polyacetals. By scrambling experiments with 1D acetals we could prove that exchange reactions occur between different building blocks. (evidenced by MALDI-TOF mass spectrometry). Based on these results we synthesized porous 3D polyacetals under the same mild conditions.

Introduction

Porous materials attracted the attention of scientists of all times due their manifold use in various technologies such as separation or catalysis. A high specific surface area is often beneficial and is realized especially in microporous materials, *i.e.* materials having pore sizes below 2 nm. Traditional examples are zeolites and activated carbons.¹ Metal-organic frameworks (MOF) or porous coordination polymers (PCP) have been introduced by various research groups during the last years as an addition to the classic materials.²⁻⁴ Most recently, interest in purely organic, but well-defined microporous materials arose and microporous organic polymers (MOP) have been developed following various concepts such as hypercrosslinking, intrinsic microporosity.⁵⁻¹² Beside these amorphous materials, highly porous crystalline covalent organic frameworks (COF) and low-molecular weight cages have been reported.^{13,14}

A common feature of all microporous polymers is the necessity of highly rigid strands and chains in order to prevent pore collapse. To ensure high microporosity in the final materials they need to feature either regular sites of contortion (such as a spirocenter or a tetraphenyl methane building block) to prevent dense chain packing or they need to be synthesized under well-controlled phase separation conditions (porogenic solvent) in order to fix the loose structure instead of forming dense polymer phases.

In the geometric sense, many of the above mentioned organic frameworks could be imagined to be composed of more or less long stiff molecular rods. Such molecules are characterized by a

high aspect ratio and conformational rigidity and are the object of intense research efforts. Numerous applications both in life- and material-sciences were indeed already developed, which were summarized in some reviews.¹⁵ Basically, molecular rods may be subdivided in conjugated and non-conjugated rods. Whereas the former are important for organic electronics, the latter are predestined for biological and biochemical applications. Some years ago we developed a new class of non-conjugated molecular rods whose key elements are ketal structures. We therefore called these rods oligospiroketal (OSK) rods.¹⁶⁻²² The general structure of OSK rods (A) is depicted in Figure 1. The backbone of the rods consists of spirocyclically joined 1,3-dioxane and cyclohexane rings, which are formed by ketalization of cyclohexan-1,4-dione E (or synthetic equivalents) with pentaerythritol B. Terminal functionalities are introduced by piperidine-4-one building blocks D. Furthermore, solubility enhancing building blocks (so-called "sleeves", C) are often used due to the scarce solubility of longer OSK rods (Fig. 1).^{17,20} Meanwhile, we developed a series of mainly biochemical applications of OSK rods.^{18,19,21,22}

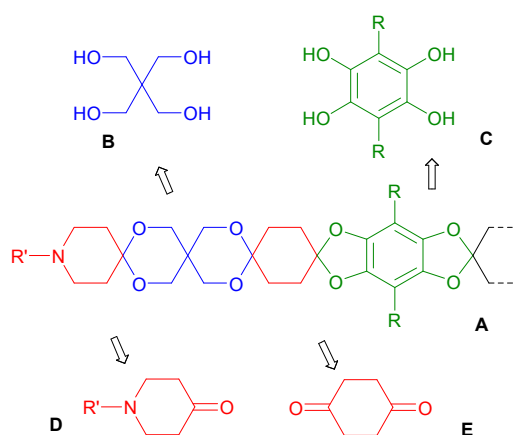


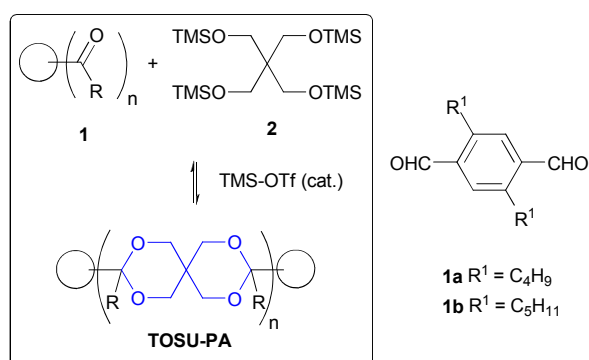
Fig. 1 General structure of OSK rods (A) and building blocks (B-E).

Building on the experience with OSK rods and their structural similarity to the basic requirements of microporous polymers, we report here on some novel porous materials with polyacetals (PA) structure whose common structural element is the 2,4,8,10-

tetraoxaspiro[5.5]undecane (TOSU) skeleton. Therefore we call these structures TOSU-PA. The synthesis of the rods is analyzed in detail before the synthesis of microporous structures and their characterization by means of gas adsorption is discussed.

Results and discussion

The synthesis of TOSU-polyacetals relies on the acetalization method developed by Noyori.²³ The bis-trimethylsilyl ethers of 1,2- and 1,3-diols react with aldehydes or ketones, catalyzed by trimethylsilyl trifluoromethanesulfonate (TMS-OTf), under very mild conditions to yield the corresponding acetals. A special feature of this method is that the reactants can be used in equimolar amounts in contrast to the “classic” method, which involves normally an excess of one of the reactants. The key reagent for the preparation of TOSU-polyacetals is the tetrakis-trimethylsilyl ether of pentaerythritol **2**.²⁴ This compound was treated with five different carbonyl compounds **1a-e** with varying number of carbonyl groups. The synthetic route to TOSU-polyacetals TOSU-PA-a-e and the structure of building blocks **1a-e** is summarized in Scheme 1. Depending on the number of carbonyl groups we expect linear 1D (**1a, b**) and cross-linked (**1c, d, e**) polyacetals.

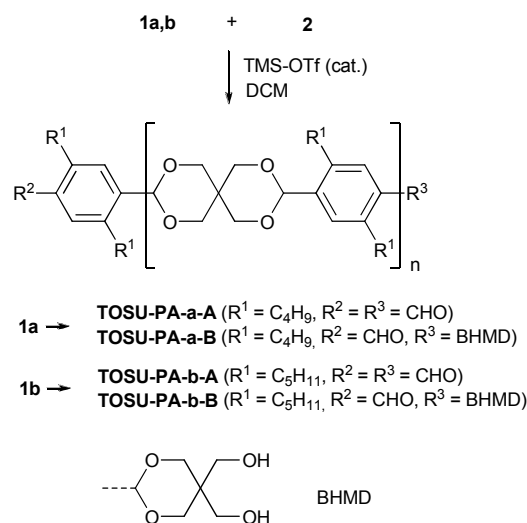


Scheme 1. Synthesis of TOSU-polyacetals and structure of building blocks **1a-e** (the TOSU skeleton is marked blue in the formula of TOSU-PA).

1D Polyacetals

A linear 1D model system was analyzed first to investigate the optimum conditions and the mechanism of the polyacetalization, especially the issue of reversibility. Reaction between terephthalaldehydes **1a, b**, bearing solubility enhancing alkyl

groups in 2- and 5-position,²⁰ and tetrasilylether **2** under Noyori conditions gave the polymers TOSU-PA-a and -b respectively (Scheme 2). Compounds TOSU-PA-a,b are white solids, which are soluble in dichloromethane. Investigation with MALDI-TOF mass spectrometry (for details see the experimental section) revealed the molar mass dispersion of the products, which have up to 10 monomer repeat units. The maximum of the distribution was centred around 4-6 monomer repeat units. Exemplarily, the MALDI-TOF MS spectrum of TOSU-PA-a ($R^1 = C_4H_9$) is depicted in Figure 2. The most intensive peaks in the mass spectra (marked with ♦) originate from polymer TOSU-PA-a-A, bearing free formyl groups in the terminal positions (as Na^+ adducts).



Scheme 2. Synthesis of 1D polyketals TOSU-PA-a and TOSU-PA-b.

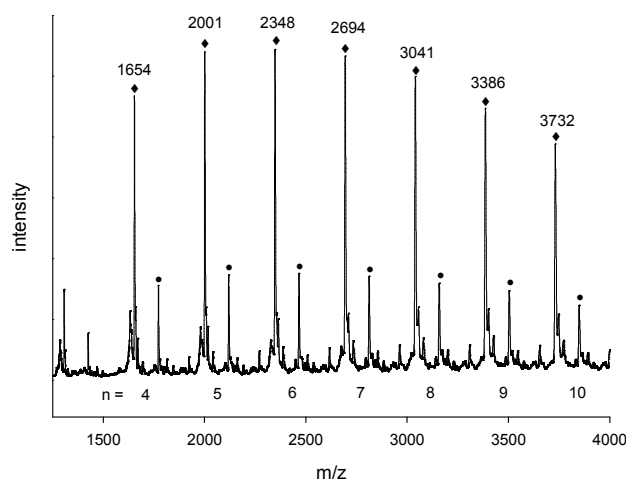


Fig. 2 MALDI-TOF MS spectrum of polyacetal TOSU-PA-a (♦ = a-A, • = a-B)

The next lower peaks (marked with •) have a difference of 118 Dalton relative to the main peaks and can be explained by

replacement of one of the formyl groups in TOSU-PA-a-A by a 5,5-bis-hydroxymethyl-1,3-dioxan-2-yl (BHMD) group (TOSU-PA-a-B). An analogous composition is observed in the case of pentyl-substituted polymer TOSU-PA-b. The dependency of the degree of polymerization on the concentration of the monomers and/or the catalyst was analyzed subsequently. All synthesized polymers TOSU-PA-a, b, sc (sc = scrambling) were analyzed by MALDI-TOF-MS. Minimum influence of the monomer concentration on the degree of polymerization was found (figure 3). Monomer concentrations ranging from 6.67 mmol/L up to 40 mmol/L of monomer **1a** yielded rather similar MALDI-TOF-MS spectra. Furthermore, we varied the concentration of the catalyst TMS-OTf and observed once more no significant influence on the degree of polymerization (see ESI). We also investigated the monomer molecular ratio. But contrary to expectations that polycondensations gave the best turnover to high molecular weights at a monomer molar ratio of 1:1 we determined for our soluble systems that a monomer molar ratio of **1a/b** : **2** = 1:1.5 gave the highest molecular weights, i.e. largest number of repeating units according to MALDI-TOF analysis.

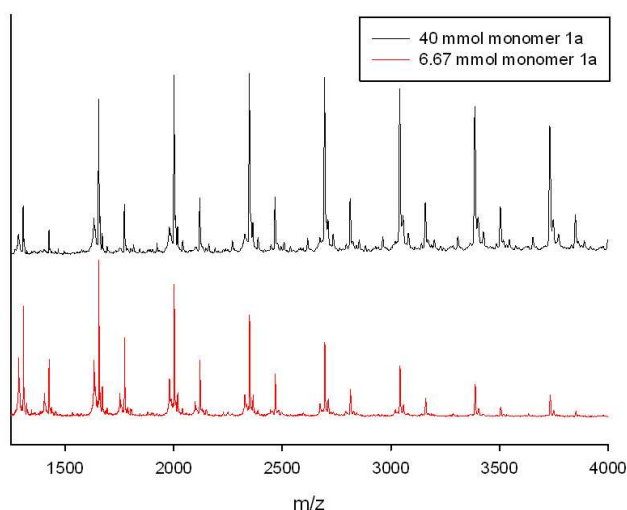
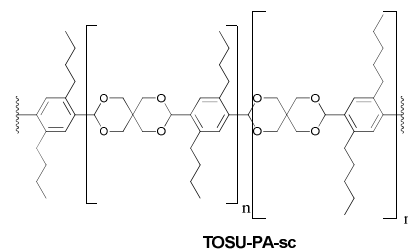


Fig. 3. MALDI-TOF MS spectra demonstrating the influence on the monomer concentration of polyacetal TOSU-PA-a.

The acetalization reaction is basically a reversible reaction. Evidence came from a scrambling experiment, in which preformed OSKs having different alkyl side chain length (TOSU-PA-a and TOSU-PA-b) were reacted with **2** and the corresponding monomer **1b** or **1a** under Noyori conditions. If the reaction is indeed reversible, the final product should have both butyl and pentyl substituted units as repeating units in a single chain. This was indeed found and evidenced by MALDI-TOF-MS (see Figure 4).



Scheme 3. Structure of TOSU-PA-sc from the scrambling experiment.

Figure 4 shows a part of the resulting MALDI-TOF-MS spectrum around repeating units with $n=4$ and $n=5$. The significant new peaks around the main peak of TOSU-PA-a with a mass difference of 28 Dalton are the evidence of the exchange of the given butyl side chains in TOSU-PA-a with pentyl side chains of monomer **1b**.

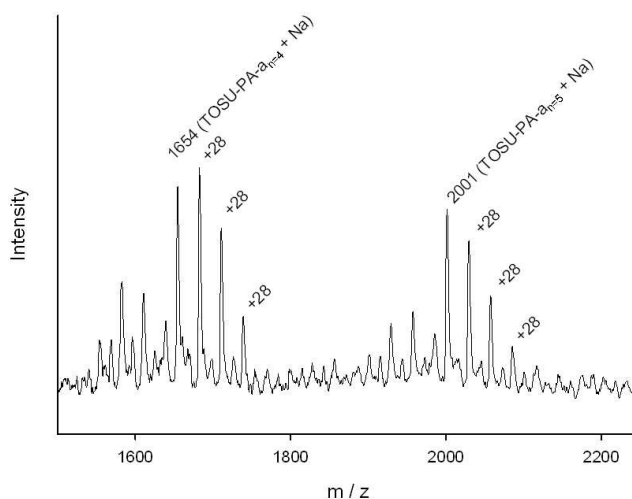
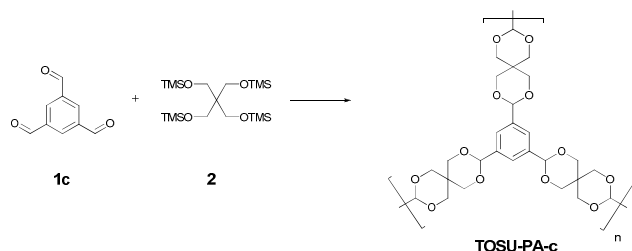


Fig. 4. MALDI-TOF MS spectrum of the scrambling with monomer **1b** and linear polymer TOSU-PA-a.

Microporous Polyacetal Networks

Having established the synthetic route towards OSK, we aimed the synthesis of microporous OSK networks (TOSU-PA-c to TOSU-PA-e) by reaction of multifunctional aldehydes or ketones (**1c-1e**) with **2** under Noyori conditions. First of all we tried to use the reaction conditions, which have been shown suitable for the synthesis of soluble TOSU-PAs. However, it was impossible to obtain micro- or mesoporous solids using those conditions, but soluble products. Instead, we used ultrasonic irradiation based on experiences that showed the possibility of conducting polymerization reactions using ultrasonic irradiation.²⁵ It turned out that the products obtained by using ultrasonic irradiation yielded indeed cross-linked micro- or mesoporous solid products. The choice of the solvent and the used catalyst did also seriously impact the final porosity of the polymer networks (see below). Depending on the conditions, which ultimately lead to phase separation, we observed also meso- and/or macroporosity in some of the materials. The finer effects of the phase separation and the impact of the reversibility of the reaction are however hard to discriminate in their effect on the meso/macroporosity and we will not discuss them in detail within this study. Initial experiments were conducted using 1,3,5-triformyl benzene (**1c**) as building block (see scheme 4).



Scheme 4. Synthesis of TOSU-PA-c.

Figure 4 shows the N₂ (77.4 K) and CO₂ (273 K) adsorption/desorption isotherms of TOSU-PA-c and TOSU-PA-d materials synthesized under varying conditions.

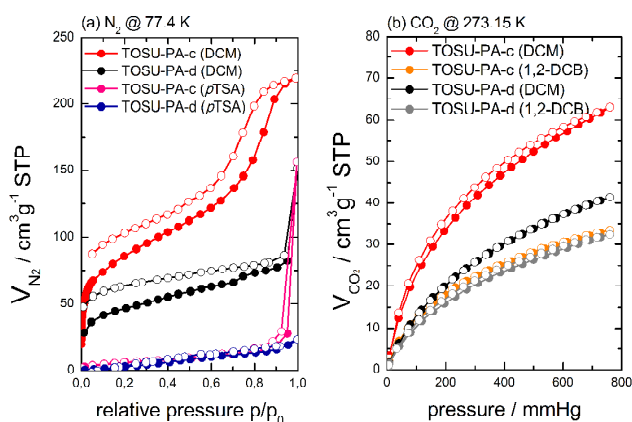


Figure 4. N₂ (77.4 K) (a) and CO₂ (273 K) (b) adsorption/desorption isotherms (adsorption with filled spheres, desorption with hollow spheres) of various TOSU-PA-c / TOSU-PA-d materials synthesized under different conditions.

A first question relates to the used acetalization conditions. Figure 4a shows clearly, that only materials synthesized under Noyori conditions (using DCM as solvent in first instance) yield porous materials. Classic conditions (*i.e.* refluxing benzene under continuous water removal and catalysis by 4-toluenesulfonic acid, pTSA) did not result in micro/mesoporous materials. This is in line with a report of Han *et al.*, who could show that rather harsh conditions were needed to obtain microporous polyketals based on pTSA catalysis (180°C, 3d, vacuum, closed flask).²⁶

A closer look on the N₂ adsorption/desorption isotherms shows that TOSU-PA-c shows some significant mesoporosity (indicated by the very steep volume increase at $p/p_0 \sim 0.8$ and the associated hysteresis) next to pronounced microporosity, which is evident from the large volume uptake at low relative pressures but also from more detailed isotherm analysis of this particular material (see ESI). TOSU-PA-d does not show signs of mesoporosity. At this stage, we will not discuss the mesoporosity in detail as it seems to originate from the finer details of the phase separation, which might vary from batch to batch. Instead, we wish to discuss the microporosity in more detail, which is expected to originate from the rigid molecular structure of the used monomers and the linking chemistry (although phase separation might influence the final microporosity to some extent as well).

Associated with the microporosity of the materials, a non-closing hysteresis loop (*i.e.* adsorption and desorption branch do not unify even at very low relative pressure) is observed, which is a characteristic feature of many microporous polymers.²⁷ The hysteresis was discussed in terms of the micropore connectivity and the softness of the materials and it was shown that CO₂ adsorption at 273 K is another (more) useful tool for the analysis of microporous polymers. The materials were hence analysed by CO₂ adsorption/desorption (see Figure 4b for examples) as well and the impact of changed synthesis conditions was analysed as well.

In a first experiment, the CO₂ adsorption capacity of TOSU-PA-c and TOSU-PA-d synthesized under Noyori conditions using either DCM or 1,2-dichlorobenzene (1,2-DCB) as solvent, was investigated (Figure 4b). The results for DCM based systems are in line with the trend obtained by N₂ physisorption at 77.4 K, *i.e.* TOSU-PA-c (DCM) and TOSU-PA-d (DCM) show high microporosity with the trimethyl benzene based system having a higher porosity (CO₂ uptake of 2.85 mmol g⁻¹ vs. 1.85 mmol g⁻¹ at 273 K and 1 bar). The use of 1,2-dichlorobenzene (1,2-DCB) resulted finally in materials with lowered porosity. Interestingly, the porosity was quite comparable for both materials (TOSU-PA-c and TOSU-PA-d) in this case. This could relate to the fact that upon synthesis in 1,2-DCB no solid product precipitated from the reaction mixture (in contrast to the use of DCM), but just upon addition of ethanol. The materials obtained by this method could however not be dissolved after drying under high vacuum (see ESI for details). The fact that no precipitate was formed initially indicates however a rather incomplete reaction, which in turn provides an explanation for the lowered porosity.²⁸ It should be mentioned that 1,4-dichlorobenzene (1,4-DCB) can basically also be used as solvent that allows the synthesis of microporous TOSU-PA-c. Its use under Noyori conditions is however not practically useful. This is due to its solid nature, which results in rather heterogenous conditions. 1,4-DCB was hence not investigated any further.

Finally, monomer **1e** was used to produce TOSU-PA-e. **1e** is basically a somewhat less reactive, but more readily available keto-based monomer. Reaction in 1,2-DCB yielded a moderately microporous polymer, while reaction in DCM led surprisingly to a lowered porosity (see ESI for isotherm data) This might be related to the different solubility of the keto monomer **1e** and oligomers build from it, compared to the formyl based monomer **1d**. Table 1 summarizes the CO₂ uptake for all materials as well as the respective surface areas and pore volume data obtained by analysis of the CO₂ data by GCMC methodology,²⁹ or by analysis of the N₂ data by QSDFT analysis.³⁰

Table 1. Summary of Porosity Parameters

Material	$S_{\text{BET}}^{\text{a)}}$ / m^2g^{-1}	$V_{\text{pore}}^{\text{a)}}$ / cm^3g^{-1}	CO_2 capacity ^{b)} / mmol g^{-1}
TOSU-PA-1c (DCM)	310	0.340	2.81
TOSU-PA-1c (1,2-DCB)	n.d.	n.d.	1.52
TOSU-PA-1c (pTSA)	22	n.d.	n.d.
TOSU-PA-1d (DCM)	233	0.128	1.85
TOSU-PA-1d (1,2-DCB)	n.d.	n.d.	1.47
TOSU-PA-1d (pTSA)	25	n.d.	0.67
TOSU-PA-1e (DCM)	14	n.d.	0.71
TOSU-PA-1e (1,2-DCB)	38	n.d.	1.38

a) from N_2 data at 77.4 K, pore volume determined from uptake at $p/p_0 = 0.95$; b) at 273.15 K and 1 bar

5 The structures of prepared TOSU-PAs were confirmed by FT-IR spectroscopy. In all spectra of the materials, bands that can be attributed to the typical bands of the monomers *e.g.* carbonyl stretching of aldehyde and acetyl monomers at 1690 cm^{-1} and the C-H stretching at 2950 cm^{-1} (Figures S6–S10, ESI) are observed.

10 The new C-O-C-O-C stretching vibration at 1160 cm^{-1} indicates a successful acetalization.

Finally, thermal analysis of selected materials indicated a reasonable stability (see ESI). The onset of decomposition under air was found at temperatures larger than 200°C . Decomposition 15 took place in a number of distinct steps, which can be related to the chemical diversity of aliphatic and aromatic units, connected by heteroatoms. Full decomposition was usually observed under air atmosphere at temperatures of $\sim 650^\circ\text{C}$, indicating that no inorganic residues are contained within the materials. The 20 thermal stability under nitrogen atmosphere was tested for a single material (TOSU-PA-e) as well. The onset of decomposition was again found at $T > 250^\circ\text{C}$, which is not very much different compared to decomposition under air, indicating that the initial steps are thermolytic cleavage of covalent bonds.

25 Again, multiple further decomposition steps were found, however the material cannot be decomposed completely under N_2 atmosphere and 32 % residue (most probably carbon materials) are found at $T \sim 1000^\circ\text{C}$.

It should be noted that the obtained TOSU-PAs are totally 30 amorphous, concluded from Wide-Angle X-ray Scattering (WAXS) measurements (see ESI).

Conclusions

In summary, we have shown that the acetalization of various 35 carbonyl compounds, each with 2-4 carbonyl groups, with tetrasilylether **2** provides polyacetals (PA) under very mild conditions (short reaction time, room temperature). The central structural element of these molecules is the conformationally rigid 2,4,8,10-tetraoxaspiro[5.5]undecane (TOSU) skeleton. In 40 the case of tri- and tetravalent carbonyl compounds (**1c-e**) we obtained TOSU-PAs with partly high porosity. They show however also a sensitivity of the observed porosity to the exact reaction conditions and it can be speculated that this complexity arises as a consequence of the different solubility of the growing 45 branched/cross-linked polymer in the various solvents, which impairs the phase-separation process as well as the principal reversibility of the reaction. Given the high CO_2 capacity of some of the materials, we believe that the presented platform of smooth and simple acetalization reactions is indeed of interest for further 50 development of porous materials.

Experimental Section

Materials and Methods

All chemicals were purchased from Aldrich or Merck and were 55 used as received. 2,5-dibutylbenzene-1,4-dicarbaldehyde (**1a**) and 2,5-dipentylbenzene-1,4-dicarbaldehyde (**1b**) were synthesized according to the reported procedure, respectively.²⁰ Benzene-1,3,5-tricarbaldehyd **1c** also was synthesized to a reported procedure.³¹ And tetrakis(4-acetylphenyl)methane (**1e**) was 60 synthesized to the reported procedure of Müller et al.³² The detailed synthesis procedure of 4,4',4'',4'''-methane-tetrayltetrabenzaldehyde (**1d**) is included in the supporting information.

Preparation of TOSU-PA-a.

To a solution of 123 mg of **1a** (0.5 μmol , 1.0 eq) and 318 mg of **2** (0.75 μmol , 1.5 eq) in 15 mL dry dichloromethane were added 70 100 μL TMSOTf. The resulting mixture was stirred for 20 h at 20°C . The solution was neutralized with a saturated solution of sodium bicarbonate, dried with magnesium sulphate and evaporated. The crude product was recrystallized from ethanol and was obtained after filtration as white solid.

75 The same procedure is followed for the preparation of TOSU-PA-b.

Preparation of TOSU-PA-sc.

80 To a solution of 24 mg of TOSU-PA-a, 6.5 mg of **1b** (8.9 μmol) and 3.8 mg of **2** (8.9 μmol) in 5 mL dry dichloromethane were added 20 μL TMSOTf. The resulting mixture was stirred for 20 h at 20°C . The solution was neutralized with a saturated solution of sodium bicarbonate, dried with magnesium sulphate and the 85 solvent was evaporated off. The crude product was recrystallized from ethanol and was obtained after filtration as white solid.

Preparation of TOSU-PA-c.

To a solution of 100 mg of **1c** (617 μmol , 1 eq) and 393 mg of **2** 90 (925 μmol , 1.5 eq) in 50 mL dry dichloromethane were added 100 μL TMSOTf. After the reaction in an ultrasonic bath for 20 h the obtained white-gray precipitate was filtered and washed with sodium bicarbonate, water, ethyl acetate, acetone, dichloromethane and dried in high vacuum. 95 The same reaction conditions in 1,2-DCB as solvent did not lead to a solid, but with the addition of ethanol after reaction there could be obtained a gray precipitate which was filtered and washed with sodium bicarbonate, water, ethyl acetate, acetone, dichloromethane and dried in high vacuum.

100 The same procedure is followed for the preparation of TOSU-PA-d-e.

Instrumentation

105 Nitrogen (N_2) adsorption/desorption experiments were performed with a Quantachrome Quadrasorb machine at liquid nitrogen temperature (77.4 K), CO_2 adsorption was measured at 273.15 K using a Quantachrome Autosorb-MP1 machine. High-purity gases were used in all cases. All samples were degassed at 70°C 110 for 20 hours before gas adsorption measurements. Initial data analysis (calculation of surface area using the Brunauer–Emmett–Teller (BET) equation, pore volume calculation and analysis using GCMC or QSDFT methodologies) was performed using the

Quadratorb 5.05 software package (Quantachrome Instruments). MALDI-TOF spectra were measured with the MALDI-TOF mass spectrometer Axima Assurance from Shimadzu Biotech and were plotted with the software Launchpad 2.

Fourier-transform Infrared (FTIR) spectroscopy was conducted using a Perkin Elmer UATR Two machine with single reflection diamond. Wide-Angle X-ray Scattering (WAXS) was measured using a FR590 diffractometer from ENRAF NONIUS (Cu-K-alpha).

Thermogravimetric analysis (TGA) was performed under synthetic air or nitrogen flow using a NETZSCH TG209-F1. A heating rate of 10 K/min was used.

Notes and references

^a Institut für Chemie, Universität Potsdam, Karl-Liebknecht-Str. 24–25, 14476 Potsdam, Germany Fax: +49 331977-5065; Tel: +49 331977-5401; E-mail: wessig@uni-potsdam.de

^b Present Address: Hochschule Zittau/Görlitz, Fachgruppe Chemie, Theodor-Körner-Allee 16, D-02763 Zittau, Germany

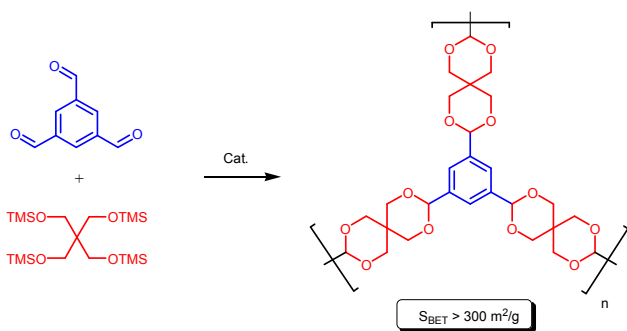
Fax: +49 3583 61-1740; Tel: +3583 61-1705, E-mail: j.weber@hszg.de

Previous address: Department of Colloid Chemistry, Max Planck Institute of Colloids and Interfaces, Research Campus Golm, D-14476, Germany;

† Electronic Supplementary Information (ESI) available: (Additional synthetic procedures and analytical data (IR, TGA, gas adsorption analysis). See DOI: 10.1039/b000000x/

- Davis, B. H. and Sing, K. S. W. (2002) Historical Aspects, in Handbook of Porous Solids (eds F. Schüth, K. S. W. Sing and J. Weitkamp), Wiley-VCH Verlag GmbH, Weinheim, Germany.
- S. Kitagawa, R. Kitaura and S. Noro, *Angew. Chem. Int. Ed.*, 2004, **43**, 2334-2375.
- O. M. Yaghi, M. O'Keeffe, N. W. Ockwig, H. K. Chae, M. Eddaoudi and J. Kim, *Nature*, 2003, **423**, 705-714.
- K. Sumida, D. L. Rogow, J. A. Mason, T. M. McDonald, E. D. Bloch, Z. R. Herm, T.-H. Bae and J. R. Long, *Chem. Rev.*, 2012, **112**, 724-781.
- V. Davankov and M. P. Tsvurupa, *Hypercrosslinked Polymeric Networks and Adsorbing Materials: Synthesis, Properties, Structure, and Applications*, Elsevier, 2010.
- A. Thomas, *Angew. Chem. Int. Ed.*, 2010, **49**, 8328-8344.
- A. Thomas, P. Kuhn, J. Weber, M.-M. Titirici and M. Antonietti, *Macromol. Rapid Commun.*, 2009, **30**, 221-236.
- N. B. McKeown and P. M. Budd, *Macromolecules*, 2010, **43**, 5163-5176.
- Y. Xu, M. D. Smith, M. F. Geer, P. J. Pellechia, J. C. Brown, A. C. Wibowo and L. S. Shimizu, *J. Am. Chem. Soc.*, 2010, **132**, 5334-5335.
- Te-Jung Hsu, F. W. Fowler and J. W. Lauher, *J. Am. Chem. Soc.*, 2012, **134**, 142-145.
- R. Dawson, A. I. Cooper and D. J. Adams, *Prog. Polym. Sci.*, 2012, **37**, 530-563.
- S. Xu, Y. Luo and B. Tan, *Macromol. Rapid Comm.*, 2013, **34**, 471-484.
- T. Tozawa, J. T. A. Jones, S. I. Swamy, S. Jiang, D. J. Adams, Shakespaere, R. Clowes, D. Bradshaw, T. Hasell, S. Y. Chon, C. Tang, S. Thompson, J. Parker, A. Trewin, J. Basca, A. M. Z. Slawin, A. Steiner and A. I. Cooper, *Nature Mater.*, 2009, **8**, 973 - 978.
- M. Mastalerz and I. M. Oppel, *Angew. Chem. Int. Ed.* 2012, **51**, 5252-5255.
- Reviews on molecular rods: a) J. M. Tour, *Chem. Rev.*, 1996, **96**, 537-553. b) P. F. H. Schwab, M. D. Levin and J. Michl, *Chem. Rev.*, 1999, **99**, 1863-1933. c) M. D. Levin, P. Kaszynski and L. Michl, *Chem. Rev.*, 2000, **100**, 169-234. d) J. M. Tour, *Acc. Chem. Res.*, 2000, **33**, 791-804. e) P. F. H. Schwab, J. R. Smith and J. Michl, *Chem Rev.*, 2005, **105**, 1197-1279. f) N. Sakai, J. Mareda, S. Matile, *Acc. Chem. Res.*, 2005, **38**, 79-87.
- P. Wessig, K. Möllnitz and C. Eiserbeck, *Chem. Eur. J.*, 2007, **13**, 4859-4872.
- P. Wessig and K. Möllnitz, *J. Org. Chem.*, 2008, **73**, 4452-4457.
- P. Müller, J. Nikolaus, S. Schiller, A. Herrmann, K. Möllnitz, S. Czaplá, P. Wessig, *Angew. Chem. Int. Ed.* 2009, **48**, 4433-4435.
- J. Nikolaus, S. Czaplá, K. Möllnitz, C. T. Höfer, A. Herrmann, P. Wessig and P. Müller, *Biochim. Biophys. Acta (BBA) - Biomembranes*, 2011, 2781-2788.
- P. Wessig and K. Möllnitz, *J. Org. Chem.*, 2012, **77**, 3907-3920.
- A. Techen, S. Czaplá, K. Möllnitz, D. Budach, P. Wessig, M. Kumke, *Helv. Chim. Acta* 2013, **96**, 2046-2067.
- C. Grimm, T. Meyer, S. Czaplá, J. Nikolaus, H. A. Scheidt, A. Vogel, A. Herrmann, P. Wessig, D. Huster and P. Müller, *Chemistry Eur. J.*, 2013, **19**, 2703-2710.
- T. Tsunoda, M. Suzuki and R. Noyori, *Tetrahedron Lett.*, 1980, **21**, 1357-1358.
- S. Aldridge, R. J. Calder, A. Rossin, A. A. Dickinson, D. J. Willock, C. Jones, D. J. Evans, J. W. Steed, M. E. Light, S. J. Coles and M. B. Hursthouse, *J. Chem. Soc. Dalton Trans.*, 2002, 2020-2026.
- P. Kmus and T. J. Patraboy, *J. Phys. Chem.*, 1985, **89**, 3379-3384.
- Y. Han, L.-M. Zhang, Y.-C. Zhao, T. Wang and B.-H. Han, *ACS Appl. Mater. Interfaces*, 2013, **5**, 4166-4172.
- J. Jeromenok, J. Weber, *Langmuir* 2013, **29**, 12982-12989
- A high degree of conversion is usually helpful for high surface area materials, see e.g. J. Weber, Q. B. Meng, *Encyclopedia of Polymer Science and Technology* 2014, **in press**, DOI: 10.1002/0471440264.pst622
- A. Vishnyakov, P. Ravikovitch, A. Neimark, *Langmuir* 1999, **15**, 8736-8742)
- J. Landers, G. Y. Gor, A. Neimark, *Colloids Surf. A* 2013, **437**, 3-32
- M. Fourmigué, I. Johannsen, K. Boubekeur, C. Nelson and P. Batail, *J. Am. Chem. Soc.*, 1993, **115**, 3752-3759.
- T. J. Zimmermann, O. Freundel, R. Gompper and T. J. J. Müller, *Eur. J. Org. Chem.*, 2000, **19**, 3305-3312.

Table of contents entry



New materials with partly high porosity are accessible by polyacetalization under very mild conditions.