

# A non-aqueous procedure to synthesize amino group bearing nanostructured organic–inorganic hybrid materials†

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**Amino-functionalized organic–inorganic hybrid materials with a narrow distributed nanostructure of 2–4 nm in size were obtained by means of a template-free and non-aqueous procedure. Simultaneous twin polymerization of novel amino group containing twin monomers with 2,2'-spirobi[4H-1,3,2-benzodioxasiline] has been applied for this purpose. The amino groups of the organic–inorganic hybrid material are useful for post derivatization.**

Organic–inorganic hybrid materials, which contain amino groups, are of great importance for a variety of applications such as supports for catalysts, adsorption of metal ions, or as key precursors for post derivatization.<sup>1,2</sup> It is an important task to obtain nanostructured materials with accessible amino groups due to their great potential for post-functionalization reactions with electrophilic reagents.<sup>3–5</sup> Amino functionalized silica materials have been successfully used for CO<sub>2</sub>,<sup>6</sup> water<sup>7,8</sup> or dye absorption<sup>7</sup> and catalysis of Knoevenagel condensation reactions<sup>9,10</sup> and the synthesis of nitrostyrenes.<sup>11</sup> Furthermore, primary and secondary amino groups can be modified in many ways to introduce carboxyl groups from cyclic anhydrides or imine groups from aldehydes.

These materials can be synthesized by different strategies which use amino-functionalized reagents.<sup>1,2,8,12,13</sup> Suitable reagents are 3-aminopropylalkoxysilanes or amino group bearing water soluble polymers such as polyvinylamine.<sup>13</sup> The synthetic procedure employing these reagents is based on water as a co-reagent or a solvent. Therefore, co-reagents which contain hydrolytically sensitive groups such as imines or isocyanates cannot be applied when water based procedures are used. To avoid these peculiarities, a synthetic method is needed which does not require water as a solvent or a reagent.

Twin-polymerization can be carried out either in organic solvents or even in the melt without the use of any solvent.<sup>14,15</sup> This advantage overcomes all problems resulting from water chemistry. Furthermore, amino groups undergo acid–base reactions with water, and ammonium ions are formed which can have a negative effect on post-reactions of amino groups.

In this communication a specific application of twin-polymerization is presented which uses a non-aqueous procedure involving various amino group bearing twin monomers to produce nanostructured organic–inorganic hybrid materials related to amino group containing sol–gel materials.

During the last six years the so called twin polymerization has been developed to fabricate organic–inorganic hybrid materials by solely one procedure.<sup>14–18</sup> The principle of twin-polymerization is based on specific twin monomers (TMs), which contain two covalently bonded polymerizable monomeric fragments, *i.e.* A and B for each polymer. The formation of the two polymers from the twin monomers during polymerization is mechanistically coupled.<sup>17</sup> Thus, polymer  $-(A)_n-$  can only be formed when polymer  $-(B)_n-$  is also formed. This is a crucial difference in the polymerization behavior from that of hetero-bifunctional monomers whose two different polymer strands are mechanistically independent of each other during the polymerization procedure.<sup>4,5,13</sup> Simultaneous twin polymerization (STP) of two different twin monomers in one process can yield up to four different polymers (see also Fig. S1 ESI†).<sup>18</sup>

A benefit of this type of STP is that the formation of polymer  $-(A)_n-$  mediates the covalent connection of fragments B and C during polymerization within the organic–inorganic hybrid material.

The synthetic procedure for the amino group containing TM (2–5) starts from salicylic alcohol and 3-amino-*n*-propyldimethoxymethylsilanes. The transesterification reaction was catalyzed by tetra-*n*-butylammonium fluoride (TBAF) (see Fig. 2). Details of the synthetic procedures are given in the ESI.†

Objective of this basic study is the synthetic feasibility of new types of TMs and to study their polymerization by STP to produce hybrid materials bearing accessible amino groups. The STP of 2,2'-spirobi[4H-1,3,2-benzodioxasiline] (1) with amino group functionalized twin monomers has been studied.

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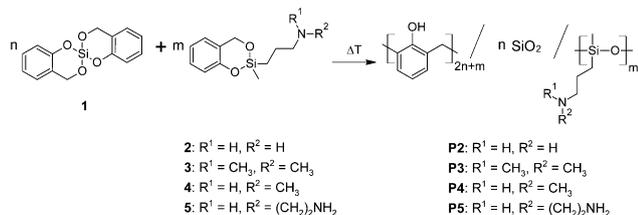


Fig. 1 Procedure for thermally induced STP of twin monomer **1** with **2**, **3**, **4** and **5**, respectively, and appellation of the resulting hybrid materials.

Pure **1** undergoes twin polymerization by either an acid- or a base-catalyzed reaction in melt or in solution.<sup>19,20</sup> However, it is also possible to polymerize **1** without the use of any catalyst at 230 °C.<sup>19</sup>

Advantageously, the amino functionalized monomers (**2–5**) used for the STP can serve as both a component and a basic catalyst. This is an elegant way of synthesis, because impurities resulting from the use of additional catalysts can be avoided. Thus, monomer **1** was simultaneously polymerized with **2**, **3**, **4** or **5** at a much lower temperature (120 °C) using a stoichiometric ratio of 1 : 1.

According to the conceptual idea of STP, an organic–inorganic hybrid material is composed of phenolic resin, silica and oligo(3-amino-*n*-propyl)methylsiloxane (OAMS) (Fig. 1).

It is apparent from silicon chemistry that silica and OAMS moieties can undergo Si–O–Si bond formation to form a class II hybrid structure among the phenolic resin/SiO<sub>2</sub>/OAMS hybrid compound.<sup>21–23</sup> This feature can be readily evidenced by solid state <sup>29</sup>Si- and <sup>13</sup>C-NMR spectroscopy shown in Fig. 2 and the ESI†

Altogether, the <sup>29</sup>Si-CP-MAS-NMR spectra of hybrid materials (P2–P5) show the expected Q- and D-signals originating from monomer **1** and the accordant amino functionalized monomers (**2–5**). In each case Q<sub>4</sub> (–110 ppm) is the most intense signal in the silica region of the spectrum, although the cross polarization technique used overrates Q<sub>2</sub> (–90 ppm) and Q<sub>3</sub> (–100 ppm) signals due to polarization transfer from <sup>1</sup>H to <sup>29</sup>Si. This result indicates that the silica network is highly condensed. The same applies to the OAMS part, as no signals from the monomer are detectable. Instead, a rather broad signal at –17.6 ppm is found,

which can be assigned as D(Q).<sup>24</sup> Parts of OAMS are covalently bonded to silica and form a copolymer, related to Co-STP as shown in Fig. 1, ESI†. The broad line widths of the D(Q)-signals are typical for a high dispersion of chemical shifts and a reduced flexibility of the OAMS moiety. The formation of phenolic resin and of OAMS can be clearly evidenced by means of <sup>13</sup>C-CP-MAS-NMR spectroscopy. The signal-structure assignments are depicted in Fig. 2. The formation of *o,o'*- and *o,p'*-linked phenolic resin structures could be evidenced by the signal pattern in the aromatic carbon region (signals 3 and 4, Fig. 2) and a broad signal of the bridging methylene groups (signal 5). An intense signal of methyl groups of OAMS arises at 0 ppm. The other carbon atoms of the alkyl chains of OAMS are found in the expected regions (10–50 ppm). The weak signal at 62 ppm originates from unreacted Si–O–CH<sub>2</sub> groups of the monomers (**1**: 66.3 ppm; **2**, **3**, **4** and **5**: ~63 ppm) or benzyl alcohol (–CH<sub>2</sub>–OH: 64.7 ppm). This is in agreement with the results of the <sup>29</sup>Si-NMR (Q<sub>3</sub>), and post-reactions above 120 °C were detected by differential scanning calorimetry (DSC) measurements (Fig. S5 ESI†). Soxhlet extraction with dichloromethane (DCM) showed extractable fractions which are attributed to low molecular weight products (see ESI†). A high degree of polymerization in the case of step growth polymerizations is only achieved with nearly complete conversion of functional groups. Thus, composites of monomers **1** and **2** were synthesized at higher temperatures (180 °C; monomer **2** solely polymerizes at 220 °C) and in different ratios to P2\_1 (1 : 2 = 95 : 05), P2\_2 (85 : 15), P2\_3 (50 : 50), P2\_4 (15 : 85) and P2\_5 (0 : 100) (experimental details are given in the ESI†). The higher polymerization temperature reduces the extractable content of the resulting hybrid material significantly from 43 wt% (P2) to 6 wt% (P2\_3), see ESI†. DSC-measurements show no more post-reactions upon thermal treatment of the materials (see ESI†). The molecular structure monitored by means of <sup>13</sup>C- and <sup>29</sup>Si-CP-MAS-NMR spectroscopy is not altered by the increased temperature (P2 Fig. S2 vs. P2\_3 Fig. S3 ESI†).

The solid state NMR spectra again evidence the formation of the phenolic resin, silica and OAMS (Fig. S3 ESI†). The different monomer ratios are reflected in the NMR spectra and the signal intensities of phenolic resin/OAMS change accordingly. The <sup>13</sup>C-NMR spectra show no differences in the chemical shifts. An increasing content of monomer **2** also leads to more intense D-signals as shown in the <sup>29</sup>Si-NMR spectra (Fig. 3). A low ratio of monomer **2** results in D(Q) species. The corresponding signal arises

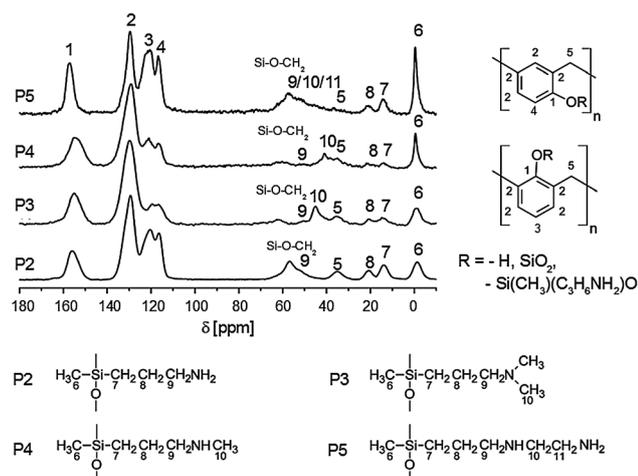


Fig. 2 Solid state <sup>13</sup>C-CP-MAS-NMR spectra of samples P2–P5.

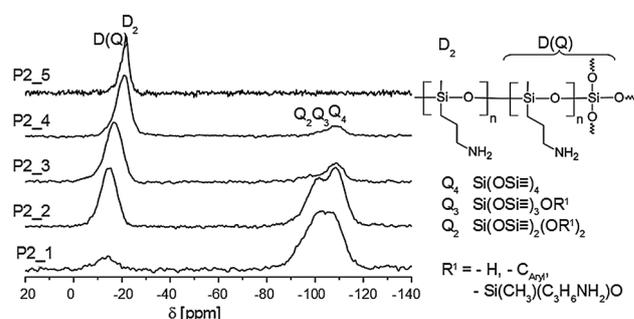


Fig. 3 Solid state <sup>29</sup>Si-CP-MAS-NMR spectra of samples P2\_1–P2\_5.



at  $-17$  ppm. Because of the formation of the silica-OAMS copolymer, the extractable content of sample **P2\_1** (0.5 wt%) is low. Longer chains (or rings) of OAMS are formed upon increasing content of monomer 2. The signal is shifted to a higher field ( $-21$  ppm) related to  $D_2$  structures.<sup>24</sup> Also a larger quantity of OAMS could be extracted (**P2\_4**: 31.2 wt%) because less covalent bonds were formed between the silica network and OAMS. Due to the absence of monomer 1 the material **P2\_5** only shows D- without any Q signals. The intensities of the Q signals of **P2\_1** to **P2\_4** are affected by the monomer ratio used. A reduction of monomer 2 leads to increased  $Q_3$  signals because the concentration of the basic catalyst (2) is decreased.

High angle annular dark field (HAADF) scanning transmission electron microscopy (STEM) images of selected hybrid materials show domain sizes of 2–4 nm (Fig. 4a). This is much lower than comparable phenolic resin/silica composites synthesized by a sol-gel process.<sup>25,26</sup> The materials are transparent and no macroscopic agglomeration could be observed in all cases (Fig. 4b and ESI†).

Thermogravimetric analysis (TGA) of the organic–inorganic hybrid materials shows a slight weight loss of 2.11 wt% (**P2\_1**) to 8.66 wt% (**P2\_5**) between 30 and 200 °C. Further decomposition of the hybrid material is comparable to that of aminopropyl-modified hybrid materials known from the literature.<sup>12</sup>

In order to check the accessibility of the amino groups in the hybrid materials after STP, the reaction with different aldehydes was studied (Fig. 5). After washing and drying procedures, the formation of the Schiff base was examined by infrared- (IR) and ultraviolet-visible (UV/Vis) spectroscopy. The treated materials show additional IR-bands at 2230, 1643 and 696  $\text{cm}^{-1}$  belonging to  $\text{C}\equiv\text{N}$  and  $\text{C}=\text{N}$  vibrations and at 1346  $\text{cm}^{-1}$  assigned to the N–O vibration of the nitro group (see ESI†). The conversion of the amino groups within the hybrid material was 14.4–66.7% determined by quantitative elemental analysis of the N-content with respect to 100% theoretical conversion (see ESI†).

The functionalized hybrid materials **SB1–3** obtained have yellow to orange color shades. UV/Vis absorption spectra of the untreated composite **P2\_3** and **SB1–3** have been recorded by means of diffuse reflectance UV/Vis spectroscopy. The UV/Vis absorption bands are non-symmetric and show several UV/Vis absorption maxima. A bathochromic shift of the UV/Vis absorption band can be observed for Schiff bases **SB1–3** in comparison to the untreated composite material **P2\_3** ( $\lambda_{\text{max}} = 294$  nm) ( $\Delta\bar{\nu}$  (**P2\_3**  $\rightarrow$  **SB3**) = 2268  $\text{cm}^{-1}$ ), which confirms the accessibility of the amino groups and the formation of Schiff bases (Fig. 6).

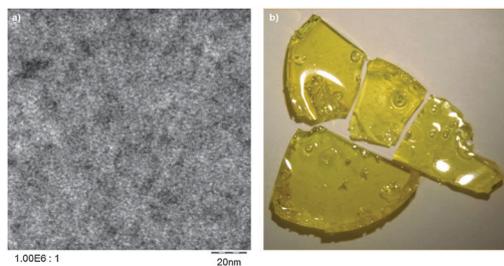


Fig. 4 (a) HAADF-STEM image of the composite material **P2** showing domains of 2–4 nm in size. (b) Macroscopic picture of the transparent, monolithic composite material.

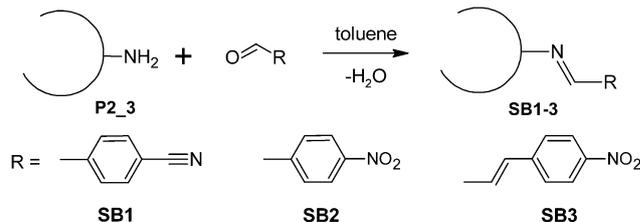


Fig. 5 Reaction of surface amino groups with *p*-cyanobenzaldehyde, *p*-nitrobenzaldehyde and *p*-nitro-cinnamaldehyde.

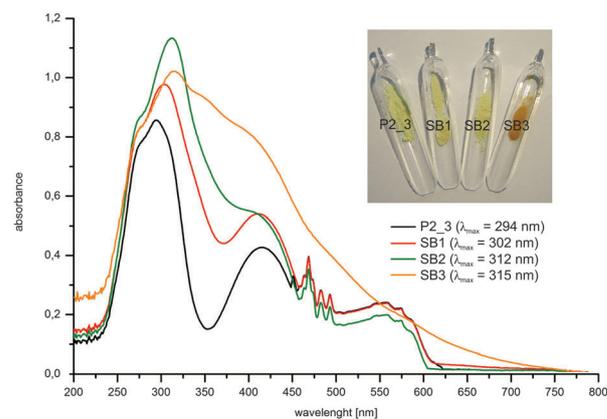


Fig. 6 UV/Vis absorption spectra of the untreated composite material **P2\_3** and **SB1–SB3**.

In addition, the UV/Vis absorption maximum of the Schiff bases shifts bathochromically from 302 nm (**SB1**) to 312 nm (**SB2**) and 315 nm (**SB3**). This is caused by an increase in the strength of the push–pull  $\pi$ -electron system with the increasing accepting capacity of the corresponding substituent ( $\text{CN} \rightarrow \text{NO}_2$ ).

We can conclude that monolithic nanostructured (phenolic resin/aminosilane/ $\text{SiO}_2$ ) organic–inorganic hybrid materials obtained from **1** (2,2'-spirobi[4*H*-1,3,2-benzodioxazoline]) and various *N*-substituted 2-(3-amino-*n*-propyl)-2-methyl-4*H*-1,3,2-benzodioxazoline derivatives (**2–5**) can be produced in the melt by self-induced simultaneous twin polymerization at 180 °C. The molecular structures of the organic–inorganic hybrid materials have been evidenced by solid state  $^{13}\text{C}$ - and  $^{29}\text{Si}$ -NMR spectroscopy. Intactness of the primary amino groups has been proved by reactions with aldehydes to achieve azomethine moieties (**SB1–3**), which can be detected readily by UV/Vis spectroscopy.

Application of this specific type of twin monomer combination in heat-induced coating, particle fabrication or other purposes will be demonstrated soon.

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