Polymers of intrinsic microporosity as high temperature templates for the formation of nanofibrous oxides†

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The highly rigid molecular structure of Polymers of Intrinsic Microporosity (PIM) – associated with a high thermolysis threshold – combined with the possibility to fill intrinsic micropores allows the direct “one-step” templated conversion of metal nitrates into nanostructured metal oxides. This is demonstrated here with PIM-EA-TB and with PIM-1 for the conversion of Pr(NO₃)₃ to Pr₆O₁₁.

Nano-templating offers rapid access to novel nano-structured materials and interfaces in particular for technologies where high surface area inorganic architectures are desirable. Template hosts such as regular opaloid structures, surfactant-based nano-structures, or novel MOF structures have been proposed. Here, polymers of intrinsic microporosity (PIM) are introduced as “high temperature templates” for “one-step” metal oxide nano-structure growth as demonstrated for the case of Pr₆O₁₁.

Polymers of intrinsic microporosity represent a novel group of polymers with a rigid backbone (see structures in Fig. 1) that prevents them from collapsing into a close-packed conformation even when heated up. Space is created within the polymer, allowing for permanent microporosity and leading to a surface area as high as 900 m² g⁻¹ for PIM-1 (ref. 7) and 1027 m² g⁻¹ for PIM-EA-TB. PIM materials are readily casted from solution into films and have been investigated for applications in gas separation membranes, catalysis, and gas storage. From thermogravimetric data (TGA, see Fig. 1) it is clear that these rigid polymer structures also show considerable high temperature stability (after some initial weight loss due to water desorption below 100 °C, decomposition onset occurs for PIM-EA-TB at 310 °C and for PIM-1 at 480 °C, both with charring). Therefore, in this study we contrast the ability of PIM-EA-TB and PIM-1 to function as template hosts for high temperature metal oxide nanostructure synthesis. A suitable model nano-structured metal oxide with promise for application in sensors and in catalysis is Pr₆O₁₁.

The synthesis of praseodymium oxides has been carried out previously by chemical vapour deposition, calcination of praseodymium hydroxide (Pr(OH)₃), electro-deposition, or by thermal transformation of a praseodymium-containing precursor compound. The products obtained through

Fig. 1 Thermogravimetric data and molecular structures for PIM-EA-TB and PIM-1.
thermal oxidation depend on both the precursor as well as oxidation conditions such as temperature and oxygen partial pressure. The oxygen deficient Pr\textsubscript{6}O\textsubscript{11} phase can be formed as dominant phase at temperatures higher than 465 °C. Bäumer and coworkers\textsuperscript{11b} investigated the formation of nanostructured praseodymium oxide via thermal decomposition of praseodymium nitrate with and without carbon-based templates. Here, we report the formation of praseodymium oxide structures not in bulk, but directly at the surface of tin-doped indium oxide (ITO) electrodes.

When the PIM host solution (1 mg mL\textsuperscript{-1} in chloroform) and Pr(NO\textsubscript{3})\textsubscript{3} solution (1 mg mL\textsuperscript{-1} in DMF) are mixed in 1 : 1 weight ratio and deposited onto ITO, calcination at 500 °C in air affords a thin film of oxide materials on ITO (see Experimental, Fig. 2).

The presence of praseodymium oxide is confirmed by EDX (see Fig. S1D\textsuperscript{†}) and by XRD (Fig. 2E, with characteristic lines\textsuperscript{15,11b}). Electron micrographs show fibrous deposits of Pr\textsubscript{6}O\textsubscript{11} on the ITO substrate (Fig. 2). With PIM-EA-TB as host template “leaf-like” nano-structures are seen. Doubling the amount of precursor deposit resulted in slightly coarser structures, which must reflect the pore geometry of the precursor at the point when solidification of the oxide precursor occurs. Changing the ratio of PIM-EA-TB to Pr(NO\textsubscript{3})\textsubscript{3} resulted in similar structures (see Fig. S1A–C\textsuperscript{†}). When investigating the PIM-1 template (see Fig. 2C and D) it became apparent that a much finer nano-structure with filaments down to 20 nm or less are formed. BET-based pore size data for PIM-EA-TB polymer (12–40 Å (ref. 18)) and for PIM-1 polymer (5 to 15 Å (ref. 19)) suggest that in both the parent polymers only comparably smaller pores are present. The feature size in the Pr\textsubscript{6}O\textsubscript{11} deposits appear considerably bigger for PIM-EA-TB but more similar to the original pore size for PIM-1. Therefore the feature size could be linked to the behaviour of the polymer template at elevated temperature. TGA data in Fig. 1 clearly show the higher thermal stability of PIM-1, which is likely to result in a finer oxide nano-structure that more closely reflects the original PIM-1 template pores.

In order to demonstrate the absence of polymer remnants, further surface analysis has been performed with XPS (Fig. 3). Apart from the underlying ITO surface elements clear evidence for Pr, C, and O is observed in the survey scan. Carbon signals are very low and assigned to adventitious surface-adsorbed molecules (or possibly remnants of the template). Oxygen signals are assigned predominantly to Pr\textsubscript{6}O\textsubscript{11}, but with some other species present at the surface. Wolffram \textit{et al.}\textsuperscript{20} have studied thin Pr\textsubscript{2}O\textsubscript{3} films made from Pr\textsubscript{6}O\textsubscript{11} targets and their work is the primary basis for fitting the O1s spectra here. Four peaks were required to curve fit the O1s spectra. The two main component at \textasciitilde 528.5 eV and \textasciitilde 531 eV most likely belong to Pr\textsubscript{2}O\textsubscript{3} and Pr\textsubscript{6}O\textsubscript{11}, respectively. Lutkehoff \textit{et al.}\textsuperscript{21} indicated that the signal at \textasciitilde 532 eV can be ascribed to Pr-based hydroxides, such as Pr(OH)\textsubscript{3}. These would be expected from the breakdown of Pr\textsubscript{6}O\textsubscript{11} in the presence of surface water (eventually leading to PrO\textsubscript{2} formation).\textsuperscript{20} The feature at \textasciitilde 529.5 eV could be either indicative of the presence of PrO\textsubscript{2} (ref. 22) or be related to surface adsorbates in the form of Pr-O-R,\textsuperscript{20} both species have O1s signal known to overlap with the Pr\textsubscript{6}O\textsubscript{11} O1s signal. As seen in Table 1, the Pr\textsubscript{6}O\textsubscript{11} and Pr-hydroxide content at surfaces seem

![Fig. 2](https://example.com/figure2.png)

**Fig. 2** Electron micrographs of (A) a 10-layer PIM-EA-TB – Pr(NO\textsubscript{3})\textsubscript{3} (1 : 1) film after calcination, (B) a 20-layer PIM-EA-TB – Pr(NO\textsubscript{3})\textsubscript{3} (1 : 1) film after calcination, (C) a 10-layer PIM-1 – Pr(NO\textsubscript{3})\textsubscript{3} (1 : 1) film after calcination and (D) a higher resolution image of the same sample. (E) XRD analysis.

![Fig. 3](https://example.com/figure3.png)

**Fig. 3** XPS survey spectra (A) of PIM-EA-TB and PIM-1 and core level data for (B) Pr\textsubscript{3}d\textsubscript{5}/\textsubscript{2} (with red and blue curve fits for Pr\textsuperscript{3+} and Pr\textsuperscript{4+} species, respectively), (C) O1s, (D) C1s.
Table 1  Oxygen composition data from XPS in as-prepared Pr$_6$O$_{11}$

<table>
<thead>
<tr>
<th>Species</th>
<th>% O1s composition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pr$_2$O$_3$</td>
<td>29.2</td>
</tr>
<tr>
<td>PrO$_2$/adsorbates</td>
<td>16.7</td>
</tr>
<tr>
<td>Pr$<em>6$O$</em>{11}$</td>
<td>45.3</td>
</tr>
<tr>
<td>Hydroxides</td>
<td>8.8</td>
</tr>
<tr>
<td>PIM-EA-TB</td>
<td>57.4</td>
</tr>
<tr>
<td>PIM-1</td>
<td>42.6</td>
</tr>
</tbody>
</table>

Complementary cyclic voltammetry data (Fig. 4B) also demonstrate the decrease in impedance as an increase in charging current. Full charging and therefore full capacitive characteristics would require more time (or a higher conductivity of the oxide). The electrochemical properties are consistent with those reported previously for Pr$_6$O$_{11}$ with potential applications in charge storage and sensing. However, the methodology for oxide nano-structure formation in PIM templates will be applicable for a much wider range of oxides and mixed oxides.

Experimental

Chemical reagents

Praseodymium nitrate hexa-hydrate, N,N-dimethylformamide (DMF) and chloroform were obtained from Sigma-Aldrich and used without further purification. Polymers with intrinsic microporosity PIM-EA-TB$^6$ and PIM-1 (ref. 23) were prepared following literature procedures. Tin-doped indium oxide glass plates (ITO) with a resistivity of 15 $\Omega$ per square were obtained from Image Optics Components Ltd (Basildon, UK). A KCl-saturated calomel (SCE) reference electrode was obtained from Radiometer.

Instrumentation

The morphology of the samples was analysed using a JEOL FESEM6301F field emission scanning electron microscopy (FE-
SEM). XPS experiments were conducted using a Thermo K Alpha (Thermo Scientific) spectrometer (operating at \( \approx 10^{-8} \)–\(10^{-9}\) Torr) with a 180° double focusing hemispherical analyser running in constant analyser energy (CAE) mode and a 128-channel detector. A mono-chromated Al K\(\alpha\) radiation source (1486.7 eV) was used. Peak fitting was conducted using XPS Peak Fit (v. 4.1) software using Shirley background subtraction. Peaks were referenced to the adventitious carbon C1s peak (284.6 eV) and peak areas were normalized to the photoelectron cross-section of the F1s photoelectron signal using atomic sensitivity factors.\(^{24}\) An Elite Thermal Systems Ltd tube furnace was used to remove the possible organic contamination on the ITO electrodes and for calcination of metal oxides. Electro-chemical testing was performed using an Ecochemie Autolab PGSTAT12 potentiostat system. TGA data were collected on a Setaram Setsys Evolution TGA instrument. The samples were heated under Ar from 20°C until 800°C at 10 K per minute.

**Procedure for nano-Pr\(_2\)O\(_{11}\) film deposition**

Tin-doped indium oxide (ITO) coated glass slides were cut into 1 cm \(\times\) 3 cm strips and cleaned by rinsing with water and ethanol, followed by calcination at 500°C for one hour. A solution of 1 mg mL\(^{-1}\) PIM in chloroform was mixed with a solution of 1 mg mL\(^{-1}\) Pr(NO\(_3\))\(_3\) \(\cdot\) 6H\(_2\)O in DMF in the desired ratio. From the resulting mixture, 25 \(\mu\)L was deposited onto a clean ITO plate covering approximately 1 cm\(^2\) and dried in an oven at 100°C for 15 minutes. This deposition process was repeated for a desired number of layers and finally followed by calcination in a tube furnace at 500°C for 1 hour.

**Conclusions**

Praseodymium oxide nano-structures have been formed in a convenient and novel “one-step” process using a high temperature template based on polymers of intrinsic microporosity. The resulting structures differ from those obtained through simple calcination and show a leaf-like or nano-fibrous structures. Finer structures are formed with the more thermally stable PIM-1 template. The results indicate that this methodology could be used beneficially for the rapid formation of a wider range of nano-structured metal oxides as well as mixed metal oxides with future applications in electronic, sensor, or solar cell components.

**Notes and references**