

## Supporting Information

# Synthesis of TiO<sub>2</sub>/SBA-15 Nanocomposites by Hydrolysis of Organometallic Ti Precursors for Photocatalytic NO Abatement

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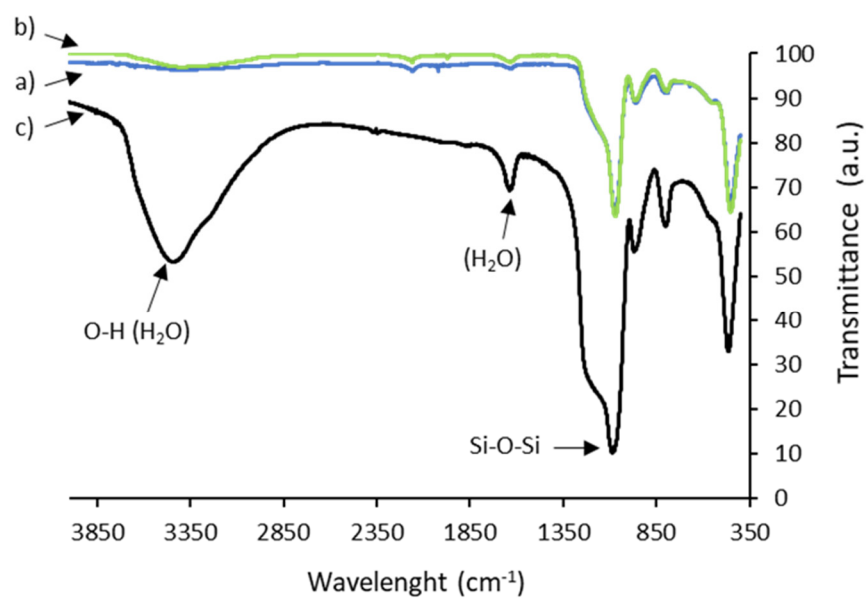
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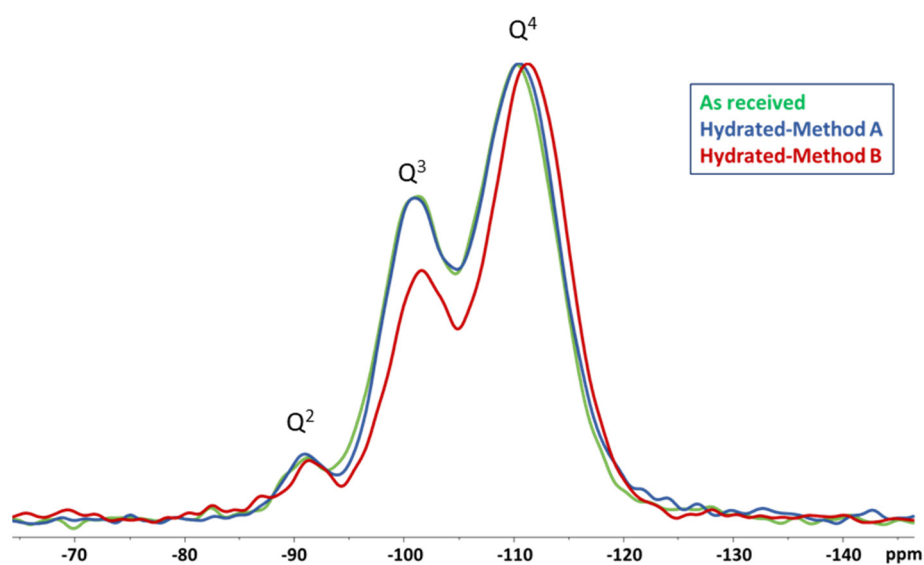
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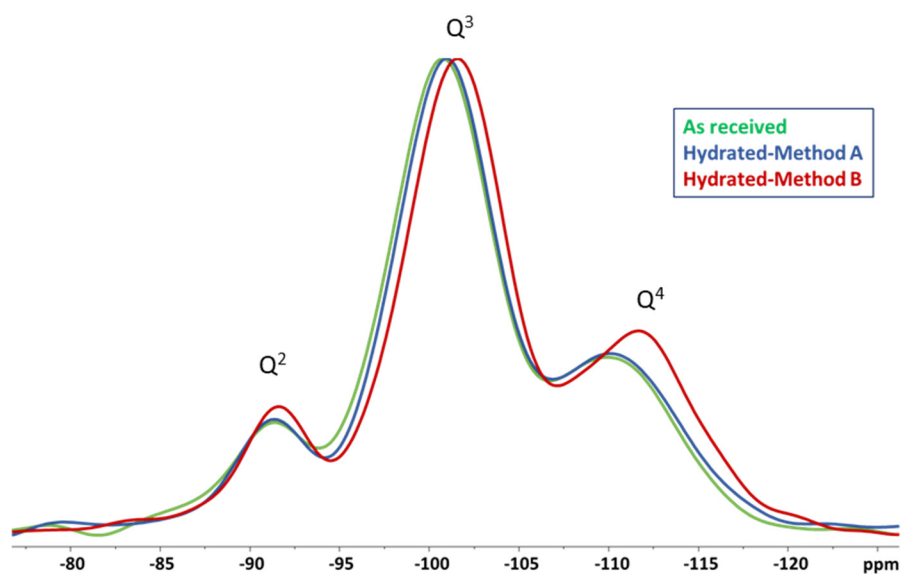
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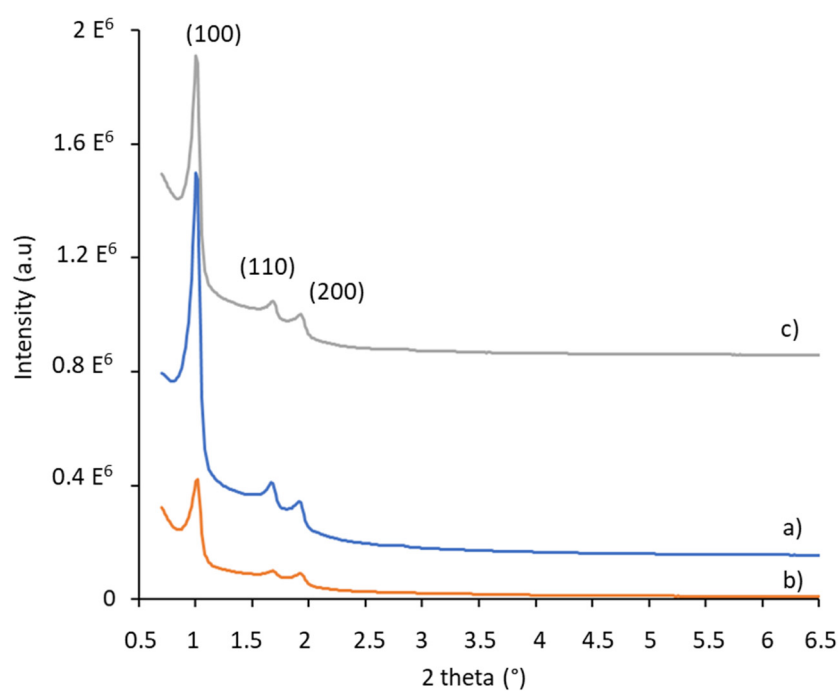
**Figure S1.** FTIR spectra for SBA-15 according to the hydration pre-treatments. a) as-received, b) hydrated by method A (atmosphere with 75 % RH), c) hydrated by method B (boiling water).



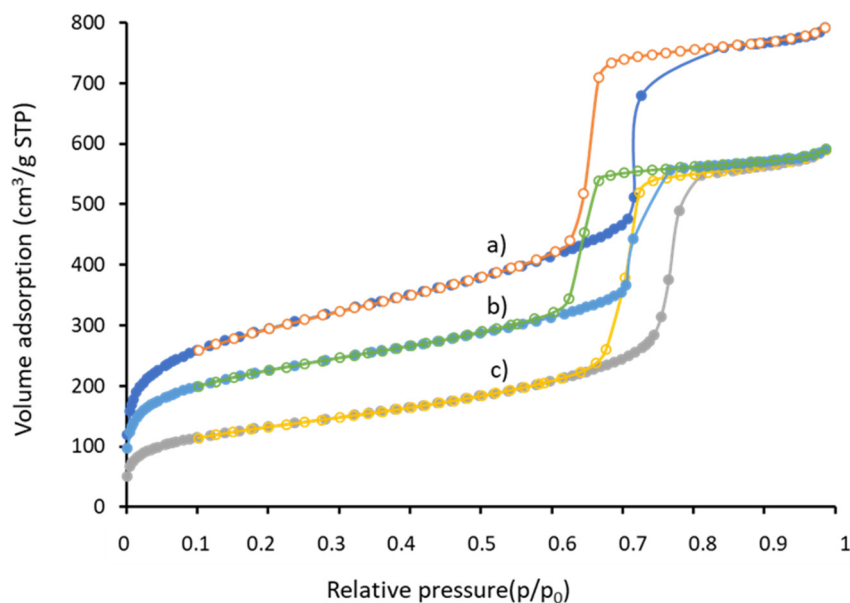
**Figure S2.**  $^{29}\text{Si}$  MAS NMR spectra of SBA-15 powders according to the hydration pre-treatments. Green curve) as-received, blue curve) hydrated by method A (atmosphere with 75 % RH), red curve) hydrated by method B (boiling water). Signal intensities were normalized to those of Q<sup>4</sup> sites.



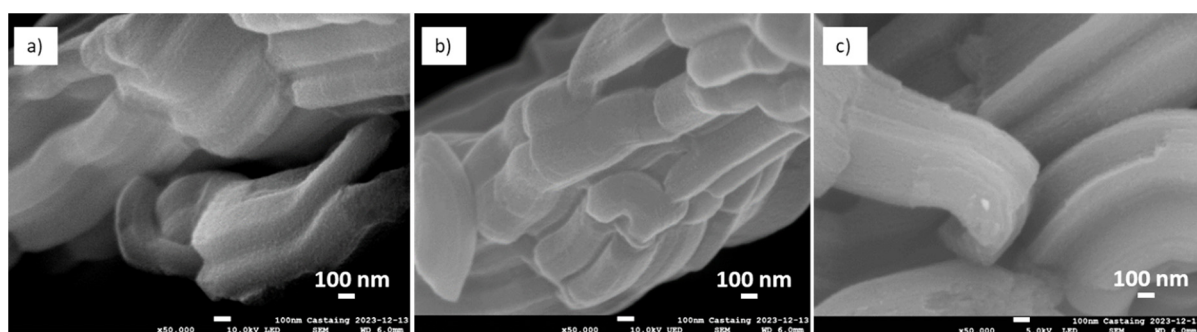
**Figure S3.**  $^{29}\text{Si}$  CP-MAS NMR spectra of SBA-15 powders according to the hydration pre-treatments Green curve: as-received, blue curve: hydrated by method A (atmosphere with 75 % RH), red curve: hydrated by method B (boiling water). Signal intensities were normalized to those of  $\text{Q}^3$  sites.



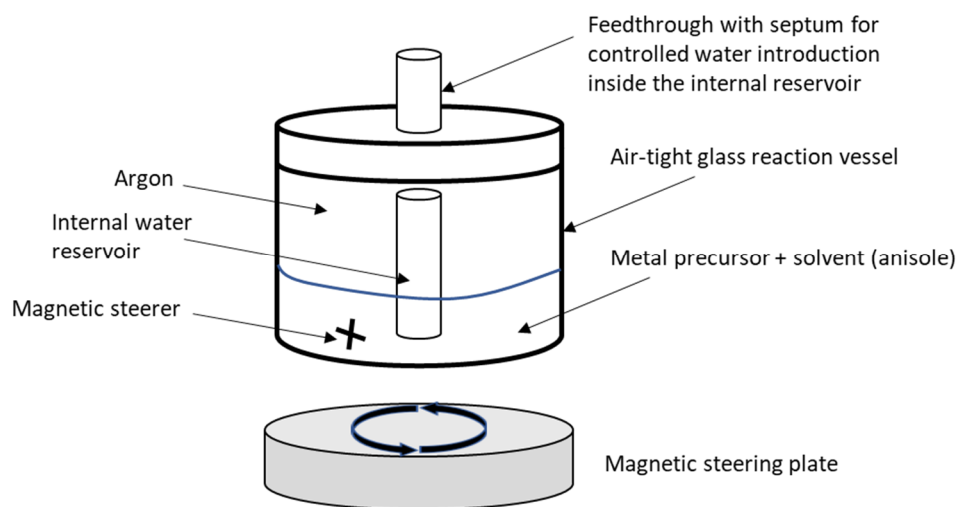
**Figure S4.** Small angle XRD diagrams of SBA-15 powders according to the hydration pre-treatments. a) as- received, b) hydrated by method A (atmosphere with 75 % RH), c) hydrated by method B (boiling water).



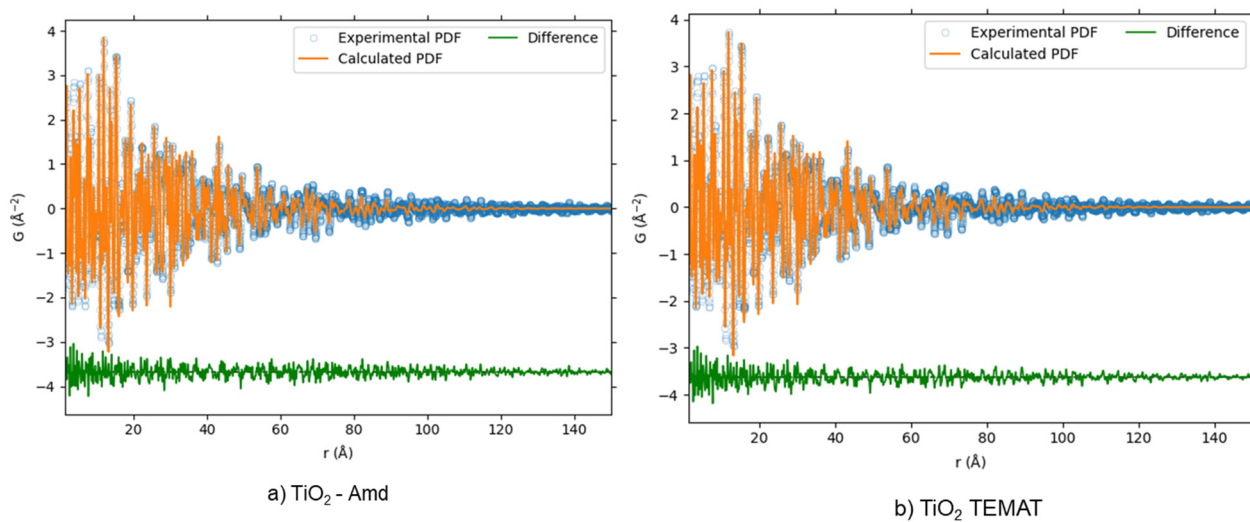
**Figure S5.** N<sub>2</sub> adsorption/desorption isotherms for SBA-15 according to the hydration method a) powder as-received, dark blue/dark orange: adsorption/desorption curve; b) hydrated by method A (atmosphere with 75 %RH), light blue/green: adsorption/desorption curve; c) hydrated by method B (boiling water), grey/light orange: adsorption/desorption curve



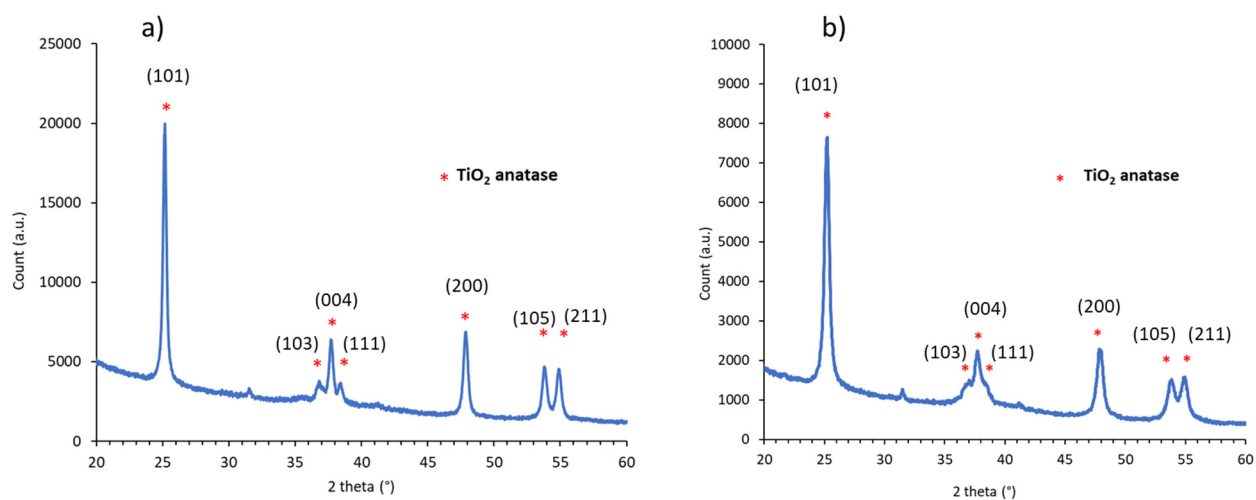
**Figure S6.** SEM images of SBA-15 powders according to the hydration pre-treatments. a) as- received, b) hydrated by method A (atmosphere with 75 % RH), c) hydrated by method B (boiling water).



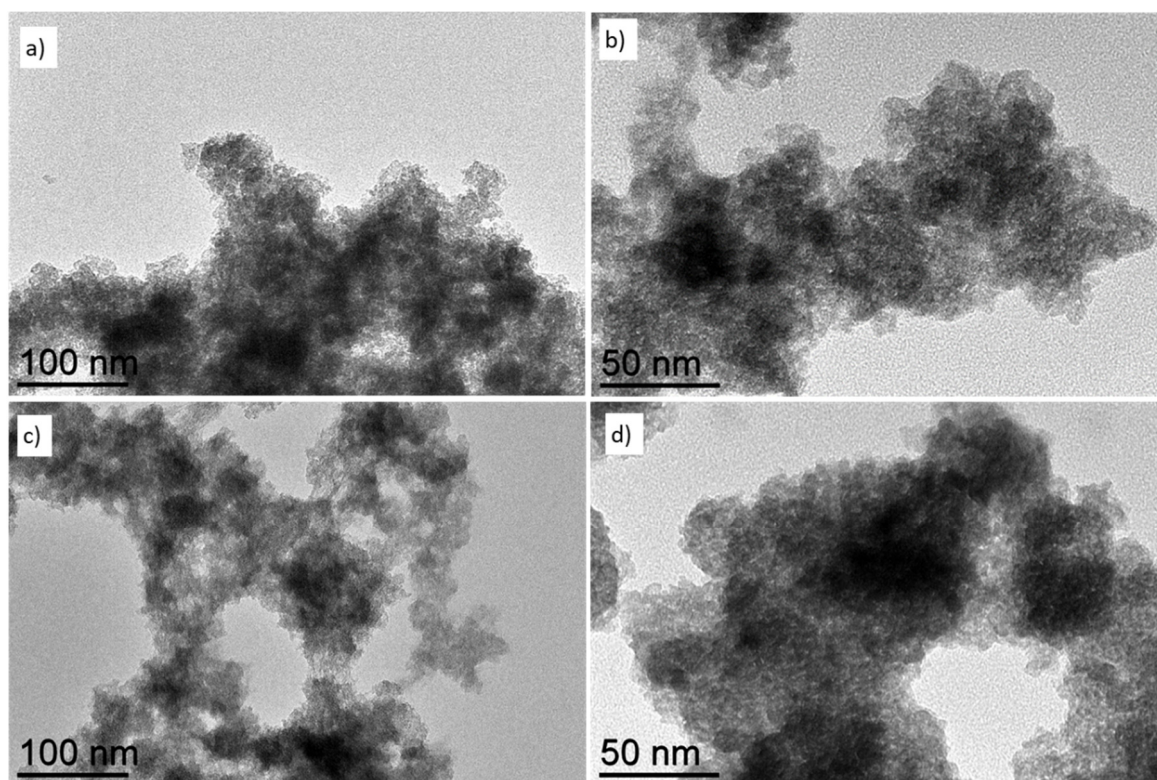
**Figure S7.** Scheme of the glass vessel used for the controlled hydrolysis of metalorganic precursor in organic solvent under argon.



**Figure S8.** Refinement of anatase structures against PDF obtained on a)  $\text{TiO}_2$ -amd and b)  $\text{TiO}_2$ -TEMAT after calcination at 500 °C.

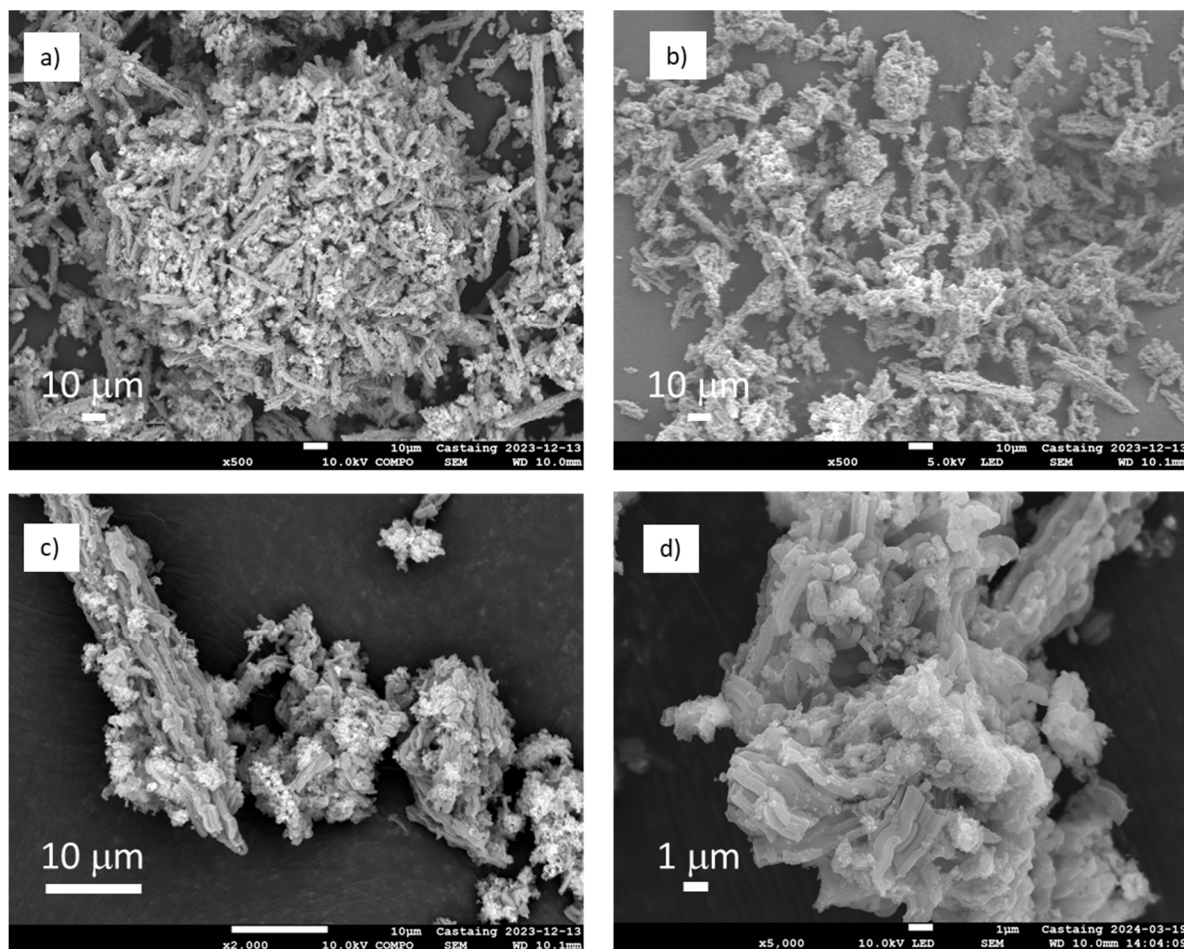


**Figure S9.** X-ray diffraction diagrams of  $\text{TiO}_2$  powders obtained after annealing at  $500^\circ\text{C}$  under air of precursor a) Ti-Amd and b) TEMAT. The anatase phase is evidenced for both samples. Small peaks at  $31.5^\circ$  and  $41.2^\circ$  are due to the mechanical support used for XRD analysis.

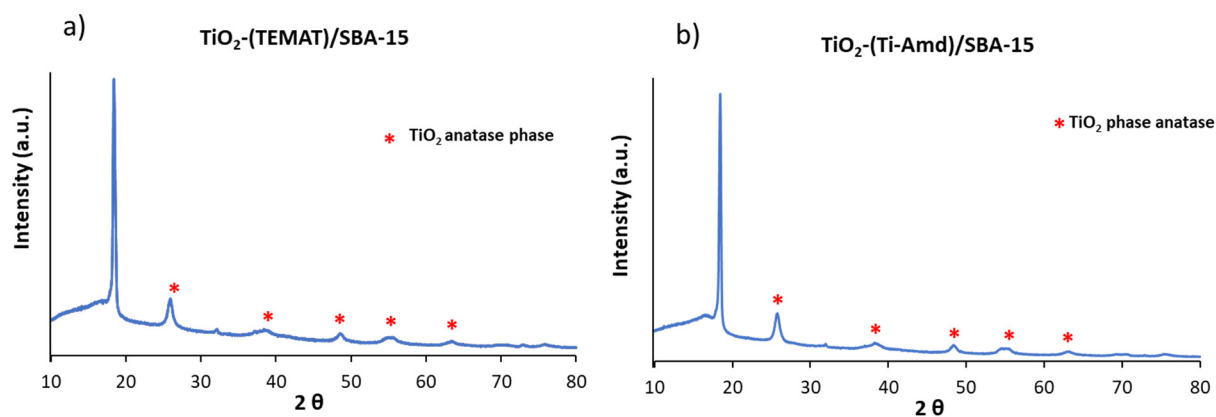


**Figure S10.** TEM images of  $\text{TiO}_x$  powders obtained from hydrolysis of TEMAT a) as-prepared, b) calcined at  $350^\circ\text{C}$  under air, and from hydrolysis of Ti-Amd c) as-prepared, d) calcined at  $350^\circ\text{C}$  under air. Magnification is  $\times 50'000$  for a) and c), and  $\times 100'000$  for b) and d).





**Figure S11.** SEM image of  $\text{TiO}_2/\text{SBA15}$  powders calcined at  $500^\circ\text{C}$  showing the homogeneous repartition of the  $\text{TiO}_2$  over the silica matrix in the sample a) and c)  $\text{TiO}_2$  obtained from TEMAT hydrolysis (magnification  $\times 500$  and  $\times 2'000$  respectively), b) and d)  $\text{TiO}_2$  obtained from Ti-Amd hydrolysis (magnification  $\times 500$  and  $\times 5'000$  respectively).



**Figure S12.** XRD analyses of nanocomposites  $\text{TiO}_2/\text{SBA-15}$  calcined at  $500^\circ\text{C}$  and a) prepared from precursor TEMAT, b) prepared from precursor Ti-Amd.