A low-cost photoactive composite quartz sand/TiO$_2$

Jonáš Tokarský$^a$, Vlastimil Matějka$^a$, Lucie Neuwirthová$^a$, Jiřina Vontorová$^b$, Kateřina Mamulová Kutláková$^a$, Jana Kukutschová$^a$, Pavla Čapková$^c$

$^a$ Nanotechnology Centre, VŠB-Technical University of Ostrava, 17. listopadu 15/2172, 708 33 Ostrava-Poruba, Czech Republic.

$^b$ Department of analytical chemistry and material testing, Faculty of Metallurgy and Materials Testing, VŠB-Technical University of Ostrava, 17. listopadu 15/2172, 708 33 Ostrava-Poruba, Czech Republic.

$^c$ Faculty of Science, J. E. Purkyně University, České mládeže 8,400 96 Ústí nad Labem, Czech Republic.

Abstract
The photoactive quartz sand/TiO$_2$ composites were prepared by thermal hydrolysis of the suspension obtained by addition of quartz sand to a titanyl sulfate solution. The required amount of TiO$_2$ in the prepared composites (i.e 9, 22 and 45 wt. %, respectively) was achieved using a variable titanyl sulfate/quartz ratio. As reference materials, pure TiO$_2$ was prepared using the thermal hydrolysis of the titanyl sulfate solution under the same conditions used during the preparation of composite quartz/TiO$_2$. The composite samples, dried at 105°C and calcined at temperatures of 500–900°C, were investigated using X-ray fluorescence spectroscopy, X-ray powder diffraction analysis, transmission electron microscopy, and Fourier transform infrared spectroscopy. Structural ordering of TiO$_2$ particles on the quartz surface was studied using atomistic simulations in a Material Studio modeling environment. Photodegradation activity of the composites was evaluated by the discoloration of Acid Orange 7 aqueous solution. The composite containing 22 wt.% of TiO$_2$ and calcined at 800°C exhibits the highest photoactivity. Higher and lower amounts of TiO$_2$ led to worse results. The quartz/TiO$_2$ composite is a promising material able to replace pure TiO$_2$ in a wide range of building materials.

Keywords: quartz sand; TiO$_2$; nanocomposite; photoactivity

*Corresponding author

E-mail address: jonas.tokarsky@vsb.cz (J. Tokarský)

Tel.: +420 597 321 519; Fax: +420 597 321 640

Abbreviations: E$_{ad}$ – adhesion energy [kcal], E$_{tot}$ – total energy of the nanocomposite [kcal], E$_{tot,SO_2}$ – total energy of the SiO$_2$ substrate [kcal], E$_{tot,TiO_2}$ – total energy of the TiO$_2$ nanoparticle [kcal], MS – Accelrys Materials Studio modeling environment, QEq – charge equilibration method, SiO$_2$(1) – pure SiO$_2$ powder dried at 105°C, SiO$_2$(6) – pure SiO$_2$ powder calcined at 600 °C, SITI – titanium oxide nanoparticles/SiO$_2$ nanocomposite, SITI11 – composite dried at 105 °C and containing 10 wt.% of TiO$_2$, SITI12 – composite dried at 105 °C and containing 20 wt.% of TiO$_2$, SITI15 – composite dried at 105 °C and containing 50 wt.% of TiO$_2$, SITI61 – composite calcined at 600 °C and containing 10 wt.% of TiO$_2$, SITI62 – composite calcined at 600 °C and containing 20 wt.% of TiO$_2$, SITI65 – composite calcined at 600 °C and containing 50 wt.% of TiO$_2$, TiO$_2$(I) – pure TiO$_2$ powder dried at 105 °C, TiO$_2$(6) – pure TiO$_2$ powder calcined at 600 °C, UFF – Universal Force Field

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1. Introduction

Titanium dioxide (TiO₂) in its nano-form is today the most frequently studied photocatalyst. The surface and structural properties of TiO₂ are summarized in detail by Diebold [1]. The TiO₂ group is composed of three modifications - anatase, brookite and rutile. Among them, the anatase is a modification with promising properties due to its high photocatalytic activity. The principle of the photodegradation mechanism of TiO₂ can be found in the article published by Fujishima and his co-workers [2].

TiO₂ nanoparticles can be prepared using a wide scale of synthesis methods [3]. The sol–gel process is the most frequently used, and the common precursors are titanium (IV) alkoxides, mainly titanium (IV) tetraisopropoxide, and titanium (IV) n-butoxide (or titanium (IV) tert-butoxide) [4,5]. The main disadvantage of these precursors is that they are too expensive to be employed for the preparation of TiO₂ nanoparticles in large amounts. Therefore, a large effort has been devoted towards the use of low cost intermediates: mainly titanium tetrachloride (TiCl₄) [6,7] and titanyl sulfate (TiOSO₄) [8,9], which are obtained during the chloride or the sulfate procedure of the TiO₂ white-pigment manufacturing.

The main application of the anatase form of TiO₂ is in the field of photocatalysis, especially degradation of environmental contaminants. This is proven by degradation of model substances presented in liquid and/or in gaseous phase. The TiO₂ can be used for the degradation of pesticides [10], phenol [11,12] and organic dyes [13,14,15] in contaminated waters (i.e. in liquid phase). In a gaseous phase the degradation of formaldehyde [16], toluene [17,18] and nitric oxides [6,19] has been studied.

The photodegradation test with organic dyes represents a fast and easy estimation of TiO₂ photodegradation activity. The rate of the degradation can be monitored by the change in the absorption of the dye solution by the UV-VIS spectrometry. In the case when TiO₂ forms a finely dispersed suspension in a dye solution, the photocatalyst has to be removed by filtration or ultracentrifugation before a measurement of UV-VIS absorption. The complete separation of TiO₂ is almost impossible if its size is in the order of nanometers. If the TiO₂ nanoparticles are fixed on the surface of a suitable particulate substrate, then the separation of the photocatalyst can be easily made by filtration or, for bigger substrate particles, by sedimentation.

Previously a large amount of work has been done in the field of toxicity of TiO₂ nanoparticles against water organisms. It was found that nanoparticles do not necessarily have to enter the cells to cause the toxicity, and that nanoparticles could cause toxic effects despite that they
formed aggregates in suspensions \[20,21\]. Zhu and his co-workers found that TiO\(_2\) nanoparticles exerted minimal toxicity to freshwater crustacean *Daphnia magna* within the traditional 48 h exposure time, but caused high toxicity when the exposure time was extended to 72 h. Therefore, it seems that TiO\(_2\) nanoparticles exposure, especially for long periods of time, may exert a negative impact on a population of aquatic organisms and on food web dynamics in aquatic systems \[22\].

An excellent method to avoid (or at least minimize) these risks is a growing and anchoring of the TiO\(_2\) nanoparticles on the surface of a suitable substrate, which prevents the release of TiO\(_2\) nanoparticles to the environment. Moreover, nanocomposites containing nanoparticles directly anchored on the carrier exhibit improved functional properties over simple mechanical mixtures \[23,24\].

Many substrates have already been studied for anchoring of the TiO\(_2\) on their surface. Preparation of a TiO\(_2\) thin layer on glass substrates by dip or spin coating is widely investigated \[25,26\]. The resulting glass exhibits anti-fogging and self-cleaning properties. Silica particles are also studied as a matrix for TiO\(_2\) growth and many authors showed that the prepared silica/TiO\(_2\) composites have improved photodegradation activity against model pollutants, such as methyl orange \[14\], methyl blue \[27\], 4-nitrophenol \[6\] and toluene \[18\]. SiO\(_2\) is very often the natural part of building materials and, therefore, represents an ideal matrix for TiO\(_2\) nanoparticles. Replacing the pure TiO\(_2\) nanoparticles by the photoactive SiO\(_2\)/TiO\(_2\) composite can help to reduce all environmental hazards and risks. But the above mentioned approach (i.e. synthesis of SiO\(_2\)), however, causes an increase of the final cost and complicates the mass production. Therefore, in the present study we made an attempt to reduce the cost of these photoactive composites by using the quartz sand instead of pure silica. We believe that these composites are useful for dealing with environmental pollution and the necessary condition for their extensive use in building industry is optimum price-efficiency ratio.

This work is focused on the preparation and characterization of the quartz sand/TiO\(_2\) composites using thermal hydrolysis of quartz sand and TiOSO\(_4\) suspension. The composites containing 9, 22 and 45 wt.% of TiO\(_2\) were prepared using a simple hydrothermal procedure. Photoactivity of the prepared composites was evaluated by means of photodegradation of Acid Orange 7 (AO7) model solution. Moreover, atomistic simulations using an empirical force field in the Materials Studio (MS) modeling environment was carried out in order to study structure and adhesion forces in the quartz/TiO\(_2\) composite.
2. Materials and methods

2.1. Sample preparation

The quartz sand / TiO$_2$ composites were prepared by thermal hydrolysis of the suspension of quartz sand in the TiOSO$_4$ solution (Precheza a.s.). The quartz sand was used as received, without any further treatment. TiOSO$_4$ contains 90 g of TiO$_2$ per 1 dm$^3$ of its suspension in concentrated H$_2$SO$_4$. The required amount of TiO$_2$ in final composite can be achieved by the setting of the appropriate quartz sand/TiOSO$_4$ ratio. The first step of composites preparation includes 1h mixing of the suspension of quartz in TiOSO$_4$ at room temperature. In the second step, the suspension is heated up to 100 °C. The third step includes 90 min long thermal hydrolysis of the suspension at 100 °C performed by addition of the appropriate volume of water (having the temperature 100°C) in the volume ratio of 1:1 with respect to the volume of TiOSO$_4$. After the thermal hydrolysis the reaction mixture is cooled down, then the supernatant liquor is separated by filtration, and the solid portion is washed several times using distilled water to achieve the conductivity of the filtrate of less than 1 mS·cm$^{-1}$. Finally, the resulting sample is dried in an oven at 105 °C. The resulting composite samples are denoted as SITI-1-X. In this assignment the number 1 means that the samples were dried at 105 °C, letter X represents TiO$_2$ content where 9, 22 and 45 means 9, 22 and 45 wt.% of TiO$_2$, respectively. Further, approximately 1 g of each of the prepared composite is calcined for 1h at selected temperatures and the samples denoted as SITI-T-X are obtained. At this assignment the letter T denotes the temperature of calcination (i.e. numbers 5, 6, 7, 8 and 9 are used for 500, 600, 700, 800 and 900 °C, respectively). Letter X has the same meaning as for the dried composites SITI-1-X. The samples of pure TiO$_2$ were assigned as TiO$_2$-T, where the letter T again denotes the temperature of heat treatment. Pure quartz sand is denoted as SiO$_2$-T in tables or figures and the letter T has the same meaning as for the TiO$_2$-T.

2.2. Characterization methods

Chemical composition of prepared samples was determined by X-ray fluorescence spectroscopy (XRFS) using SPECTRO XEPOS (SPECTRO Analytical Instruments GmbH, Germany) equipped with 50 W Pd X-ray tube. Samples were prepared in the form of pressed pellets. Wax was used as a binder.

The X-ray powder diffraction (XRPD) patterns of powder samples were recorded under CoK$_\alpha$ irradiation ($\lambda = 1.789$ nm) using Bruker D8 Advance diffractometer (Bruker AXS, Germany)
equipped with fast position sensitive detector VÅNTE 1. Powder samples were pressed in a rotational holder and measurements were carried out in the reflection mode. Lanthanum hexaboride (LaB$_6$) was used as a standard. Phase composition was evaluated using the PDF 2 Release 2004 database (International Centre for Diffraction Data). The morphology of the composite particles in the powder samples was observed on transmission electron microscope (TEM) JEOL 2010F (JEOL Ltd., Japan). LaB$_6$ crystal was used as an electron source.

Mid-infrared (IR) spectra were obtained by Fourier transform infrared spectrometer Nicolet 6700 (Thermo Fisher Scientific, USA) using the single reflection ATR technique on a diamond crystal. Measurements were performed with a resolution of 4 cm$^{-1}$ and 32 scans. Achieved data were processed using the OMNIC software.

The content of sulfur in the powder samples was determined using LECO CS244 equipped with an induction furnace HT1000 (LECO Corporation, USA). The combustion process was realized in ceramic crucibles filled with 1g of flux LECOCEL (tungsten particulates) and 1g of iron chip accelerator.

Specific surface area (SSA) was determined at liquid-nitrogen temperature using Sorptomatic 1990 (Thermo Fisher Scientific, USA), employing the BET methodology. Nitrogen gas was used as the adsorbate.

The method for testing of the photodegradation activity of the prepared composites is based on the degradation of model organic compound, Acid Orange 7 (AO7), after irradiation with UV light. In the first step two identical suspensions containing 50 mg of the composite, 65 ml of deionized water and 5 ml of the AO7 aqueous solution ($c = 6.259\times10^{-4}$ mol dm$^{-3}$) were stirred in the dark for 60 min to achieve the absorption equilibrium. After this period one suspension was filtered and the obtained filtrate was stored in the dark. Concentration of AO7 in this filtrate is denoted as $c_0$. The second suspension was subjected to 60 and 300 min long irradiation using a Pen Ray lamp (intensity of the irradiation measured at 365 nm is 2 mW cm$^{-2}$) and subsequently filtered. Concentrations of AO7 in these suspensions are denoted as $c_t$ where $t$ means the length of irradiation in minutes. The values of absorbance of filtrates at 450 nm from which the concentrations were calculated are measured using a CINTRA 303 UV-VIS spectrometer. Relative loss of AO7 (i.e. the photoactivity of the sample) is quantified using the following formula

$$\text{photoactivity} = \left(1 - \frac{c_t}{c_0}\right) \times 100$$

(1)
Both the geometry optimization and the energy calculation were carried out in *Accelrys Materials Studio (MS)* modeling environment/Forcite module. A smart algorithm was used for the geometry optimization with 50,000 iterations. Universal force field (UFF) was used for the energy calculation and the atom based summation method was used to calculate the Coulombic and the van der Waals energy.

3. Results and discussion

3.1. Chemical and phase composition of the samples

The chemical composition of quartz sand and the resulting composites SITI-1-9, SITI-1-22 and SITI-1-45 can be seen in Table 1.

<table>
<thead>
<tr>
<th></th>
<th>quartz sand</th>
<th>SITI-1-9</th>
<th>SITI-1-22</th>
<th>SITI-1-45</th>
</tr>
</thead>
<tbody>
<tr>
<td>Al₂O₃</td>
<td>1.6</td>
<td>1.4</td>
<td>1.2</td>
<td>0.835</td>
</tr>
<tr>
<td>CaO</td>
<td>0.027</td>
<td>0.052</td>
<td>0.098</td>
<td>0.195</td>
</tr>
<tr>
<td>K₂O</td>
<td>0.026</td>
<td>0.205</td>
<td>0.172</td>
<td>0.124</td>
</tr>
<tr>
<td>MgO</td>
<td>0.020</td>
<td>0.018</td>
<td>0.009</td>
<td>0.008</td>
</tr>
<tr>
<td>Na₂O</td>
<td>&lt;0.020</td>
<td>0.140</td>
<td>0.116</td>
<td>0.085</td>
</tr>
<tr>
<td>SO₃</td>
<td>0.170</td>
<td>1.5</td>
<td>4.1</td>
<td>8.1</td>
</tr>
<tr>
<td>SiO₂</td>
<td>96.9</td>
<td>86.7</td>
<td>73.0</td>
<td>47.6</td>
</tr>
<tr>
<td>TiO₂</td>
<td>0.062</td>
<td>8.0</td>
<td>18.6</td>
<td>39.0</td>
</tr>
<tr>
<td>LOD</td>
<td>0.041</td>
<td>0.890</td>
<td>2.3</td>
<td>4.5</td>
</tr>
<tr>
<td>LOI</td>
<td>0.2</td>
<td>2.5</td>
<td>6.0</td>
<td>12.1</td>
</tr>
</tbody>
</table>

The measured real content (8.0, 18.6 and 39 wt.%) of TiO₂ is lower than the expected TiO₂ content (9, 22 and 45 wt.%) showing the yield approx. 87% with respect to the theoretical content of TiO₂. It is evident that the amount of the sulfur (remaining in the samples as a remnant of the TiOSO₄ precursor) as well as LOD and LOI increases with the increase of the TiO₂ content. The increasing content of the sulfur in the composites is probably connected to the presence of free sulfuric acid, which in fact represents a dispersion medium for TiO₂ in TiOSO₄ colloid suspension. With the sulfur content growing, the amount of absorbed water (expressed by LOD) also increases. Sulfuric acid is a strongly hygroscopic compound and is responsible for the presence of the physically absorbed water, which is easily removed at relatively low temperatures. The values of LOI also increase with the TiO₂ content, and these
values comprise the presence of sulfur (in a form of sulfuric acid), and chemically bonded water in TiO$_2\cdot$nH$_2$O, which is the main product of the TiOSO$_4$ thermal hydrolysis. The dependency of the sulfur content on the amount of TiO$_2$ and calcinations temperature is shown in Table 2, whereas it is evident that the last most significant decrease in the sulfur content was observed at 800 °C.

**Table 2**
The dependence of sulfur content [wt.%] on the amount of TiO$_2$ in SITI composites and calcinations temperature.

<table>
<thead>
<tr>
<th>TiO$_2$ content in SITI [wt.%]</th>
<th>105</th>
<th>500</th>
<th>600</th>
<th>700</th>
<th>800</th>
<th>900</th>
</tr>
</thead>
<tbody>
<tr>
<td>9</td>
<td>0.599</td>
<td>0.311</td>
<td>0.086</td>
<td>0.030</td>
<td>0.025</td>
<td>0.019</td>
</tr>
<tr>
<td>22</td>
<td>1.620</td>
<td>0.922</td>
<td>0.259</td>
<td>0.055</td>
<td>0.028</td>
<td>0.018</td>
</tr>
<tr>
<td>45</td>
<td>3.240</td>
<td>2.270</td>
<td>0.449</td>
<td>0.184</td>
<td>0.044</td>
<td>0.023</td>
</tr>
</tbody>
</table>

XRPD patterns (note that all XRPD patterns were measured using Co$_{K\alpha}$ irradiation) of the quartz sand sample and the SITI composites dried at 105 °C are shown in Fig. 1.

![XRPD patterns](image)

**Fig. 1.** Comparison of X-ray powder diffractions pattern of pure quartz sand and SITI composites dried at 105 °C. Q: SiO$_2$, M: KAlSi$_3$O$_8$, A: TiO$_2$ - anatase form.

Muscovite (M; PDF number 7-0042) represents the typical mineral admixture accompanying pure quartz (Q; PDF number 85-0798). The presence of this admixture is also in agreement with the chemical composition of the original quartz sand sample (see Table 1). In the SITI
composites, the increase of TiO$_2$–anatase (A; PDF number 86-1157) content led to the
decrease of the Q and M fractions and consequently to the decrease of intensity for Q and M
peaks. This decrease is the only evidence that SITI-I-9 contains a TiO$_2$ because no distinctive
TiO$_2$–anatase peak can be found in this XRPD pattern. For SITI-I-22 and SITI-I-45 only one
broad, and not very distinctive, TiO$_2$–anatase peak (101) is evident in the region of 26–29º 20
(CoK$_\alpha$) suggesting a low crystallinity of TiO$_2$ (see Fig. 1).
The XRPD patterns of the SITI samples calcined for 1h at 600ºC are presented in Fig. 2.

![X-ray powder diffraction patterns of SITI composites calcined at 600ºC. Q: SiO$_2$, M: KAlSi$_3$O$_8$, A: TiO$_2$ - anatase form.](image)

Comparing the XRPD patterns of composites before and after the calcinations, it is clearly
observable that the quartz does not undergo any structure change while the TiO$_2$-anatase
peaks become significant. The presence of the (101) TiO$_2$-anatase diffraction peak in the
region of 26–29º 20 (CoK$_\alpha$) is clearly observable for all SITI composites calcined at 600ºC.
The presence of this peak is caused by the increase of size of the TiO$_2$ crystallites during the
calcination process. The lowering the intensity of M peak (32º 20) is caused by the decrease
of the fraction of this admixture with the increasing amount of TiO$_2$.

The XRPD patterns of the SITI samples calcined for 1h at 800ºC are presented in Fig. 3. One
can see that quartz still does not undergo any significant changes while TiO$_2$ starts to
transform from the anatase phase into the rutile phase. The comparison of the (101) TiO$_2$-
anatase diffraction peak (26–29º 20) in Figs. 2 and 3 reveals the further increase of the anatase
crystallites size (see also Table 3). The M peak (32º 20) is no longer observable because of the
presence of the much more significant (110) TiO$_2$-rutile diffraction peak.
Fig. 3. Comparison of X-ray powder diffraction patterns of SITI composites calcined at 800°C. Q: SiO$_2$, A: TiO$_2$ - anatase form, R: TiO$_2$ - rutile form.

The TiO$_2$–anatase crystallite size ($L_C$) for all SITI composites was calculated according to the (101) TiO$_2$–anatase diffraction peak using the Scherrer formula [28] and the values are shown in Table 3 (lanthanum hexaboride (LaB$_6$) was used as a standard).

Table 3
The crystallite size $L_C$ [nm] of anatase grows with the temperature of calcination and reaches the maximum for the sample SITI-9-45.

<table>
<thead>
<tr>
<th>TiO$_2$ content in SITI [wt.%]</th>
<th>temperature [°C]</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>500</td>
</tr>
<tr>
<td>9</td>
<td>18</td>
</tr>
<tr>
<td>22</td>
<td>13</td>
</tr>
<tr>
<td>45</td>
<td>13</td>
</tr>
</tbody>
</table>

It is evident that the $L_C$ grows with dependence on the calcination temperature. In the temperature range 500-800 °C the $L_C$ is not significantly affected by an increasing amount of TiO$_2$ while for SITI-9-45 the $L_C$ = 160 nm is much higher than for SITI-9-22 and SITI-9-9 ($L_C$ ~ 103 nm).

The dependence of the anatase/rutile ratio on the amount of TiO$_2$ and on the temperature of calcination is clearly shown in Fig. 4. The SITI composites with a lower amount of TiO$_2$ (i.e. containing 9 and 22 wt.%, respectively) show the appearance of the rutile phase for the first time at 800 °C, while in the case of the SITI composite containing 45 wt.% of the TiO$_2$ the rutile phase occurs even at 700 °C. For all the SITI composites 600 °C is the highest
calcination temperature at which only the anatase form is present (see Fig. 4). The amount of rutile increases with increasing content of TiO$_2$ in the SITI composite.

![Graph showing anatase and rutile content](image)

**Fig. 4.** The anatase and rutile content in all SITI samples depending on drying (105°C) and calcination (500 – 900°C) temperature.

3.2 *Infra-red spectra of the prepared composites*

Fig. 5 shows the mid-IR spectra of the samples dried at 105 °C while the mid-IR spectra of the samples calcined at 600 °C and 800 °C can be found in Fig. 6 and Fig. 7, respectively.

![Mid-IR spectra graphs](image)

**Fig. 5.** Mid-IR spectra of the samples dried at 105 °C.
Fig. 6. Mid-IR spectra of the samples calcined for 1h at 600 °C.

Fig. 7. Mid-IR spectra of the samples calcined for 1h at 800 °C.

The spectra of both dried and calcined TiO$_2$ have a broad band around 1200 cm$^{-1}$ which may be attributed to vibration of S-O bonds in sulfuric acid remaining in the TiO$_2$ sample from the used precursor TiOSO$_4$ (see Fig. 5). The spectra of the SITI samples exhibit broad bands at 1160 cm$^{-1}$ corresponding to stretching vibrations in the linear chain of Si-O-Si, and two bands around 1080 cm$^{-1}$ (vibrations of 4-fold and 6-fold SiO$_4$ rings) and 1050-1060 cm$^{-1}$ (vibrations of 4-fold SiO$_4$ rings) [29]. The spectra of the pure quartz sand and the samples with lower content of TiO$_2$ contain a doublet below 800 cm$^{-1}$ characteristic for vibrations of Si-O bonds in quartz.

The characteristic band between 910-960 cm$^{-1}$ attributed to Si–O–Ti antisymmetric stretching vibration [30] was not observed at the recorded spectra of the SITI composites, probably due
to a low abundance of these bonds in the prepared composites. Therefore, computer molecular modeling has been used to at least indirectly confirm the presence of these bonds.

3.3. Composite particles morphology and texture parameters

SEM image of SITI-6-22 in Fig. 8 reveal that the surface of quartz particles in the SITI composite is fully covered by TiO\textsubscript{2} layer. The cracks caused by the calcination are clearly visible. However, also the aggregates of pure TiO\textsubscript{2} are present. The structure of these aggregates is more obvious in TEM images (see Figs. 9a and 9b).

![SEM micrograph of SITI-6-22 composite.](image)

![TEM micrographs of (a) SITI-1-22 and (b) SITI-6-22 composites.](image)

The presence of the amorphous TiO\textsubscript{2} clusters in SITI-1-22 (Fig. 9a) and agglomerates of well-crystallized TiO\textsubscript{2} nanoparticles SITI-6-22 (Fig. 9b) is in a good agreement with the XRPD patterns in Figs. 1 and 2, respectively. Both Figs. 9a and 9b also reveal that the surface of
quartz in the SITI composite is not fully covered by TiO$_2$. SSA values for the pure quartz sand (dried and calcined at 600 °C), SITI-$I$-$X$ and SITI-$T$-$22$ composites are shown in Table 4.

Table 4
Comparison of the specific surface area values for the pure quartz sand (SiO$_2$; dried at 105°C and calcined at 600°C), SITI-$I$-$X$ and SITI-$T$-$22$ composites.

<table>
<thead>
<tr>
<th>sample</th>
<th>SSA [$\text{m}^2/\text{g}$]</th>
<th>sample</th>
<th>SSA [$\text{m}^2/\text{g}$]</th>
</tr>
</thead>
<tbody>
<tr>
<td>SiO$_2$(1)</td>
<td>&lt; 1</td>
<td>SITI-5-22</td>
<td>22.2</td>
</tr>
<tr>
<td>SiO$_2$(6)</td>
<td>&lt; 1</td>
<td>SITI-6-22</td>
<td>18.4</td>
</tr>
<tr>
<td>SITI-1-9</td>
<td>10.9</td>
<td>SITI-7-22</td>
<td>11.3</td>
</tr>
<tr>
<td>SITI-1-22</td>
<td>38.1</td>
<td>SITI-8-22</td>
<td>9.1</td>
</tr>
<tr>
<td>SITI-1-45</td>
<td>105.8</td>
<td>SITI-9-22</td>
<td>4.4</td>
</tr>
</tbody>
</table>

One can see that the SSA values increase with dependence on the TiO$_2$ amount, which is related to the origination of new TiO$_2$ surfaces. For the calcined samples the SSA values are significantly lower (for an example, comparing the SSA value for SITI-$I$-$22$ and SITI-$6$-$22$). The decrease of SSA values with increasing temperature of calcination is in a good agreement with the increasing size of the TiO$_2$ crystallites (see Tables 3 and 4).

3.4. Determination of photodegradation activity of prepared composites

For the determination of the photodegradation activity the model organic compound AO7 was used. The photodegradation activity of the prepared samples, calculated using Eq. (1), can be seen in Table 5.

Table 5
Photodegradation activities [%] for all dried and calcined SITI samples. Comparison of the photodegradation activity for the SITI-$T$-$22$ composites after 1h and 5h long irradiation with UV light can be seen in the second and third row.

<table>
<thead>
<tr>
<th>TiO$_2$ content in SITI [wt.%]</th>
<th>temperature [$^\circ\text{C}$]</th>
<th>105</th>
<th>500</th>
<th>600</th>
<th>700</th>
<th>800</th>
<th>900</th>
</tr>
</thead>
<tbody>
<tr>
<td>9</td>
<td></td>
<td>6.5</td>
<td>11.4</td>
<td>5.1</td>
<td>10.3</td>
<td>13.8</td>
<td>-</td>
</tr>
<tr>
<td>22 (1h)</td>
<td></td>
<td>7.7</td>
<td>20.2</td>
<td>23.0</td>
<td>26.5</td>
<td>27.6</td>
<td>9.3</td>
</tr>
<tr>
<td>22 (5h)</td>
<td></td>
<td>20.0</td>
<td>73.0</td>
<td>70.0</td>
<td>76.0</td>
<td>84.0</td>
<td>31.0</td>
</tr>
<tr>
<td>45</td>
<td></td>
<td>9.1</td>
<td>22.7</td>
<td>21.8</td>
<td>25.6</td>
<td>19.4</td>
<td>-</td>
</tr>
</tbody>
</table>

At first the photodegradation activity of the SITI-$T$-$X$ composites has been evaluated after the 1h long UV irradiation. Comparing the results obtained it is evident that the SITI-$T$-$9$
composites shows the lowest photodegradation activities, which ranges from 6.5 – 13.8 %. The maximum photodegradation activity after 1 h long irradiation 27.6 % was observed for the SITI-8-22 composite (the composite with 19 wt.% of TiO₂ calcined at 800 °C). Significantly higher content of TiO₂ in SITI-T-45 composite (40 wt.% of TiO₂) did not enhance the photodegradation activity as evident from Table 5, slightly higher photodegradation activity shows only the SITI-I-45 and SITI-5-45 composites. Taking into account the results of photodegradation activity obtained after 1h long irradiation, the composites with 19 wt.% of TiO₂ were selected for photodegradation tests using 5h long UV irradiation. Comparing the extent of AO7 photodegradation after 1 and 5 h long irradiation tested for composites SITI-T-22 it is evident, that the prolonged irradiation has significant effect on degradation of AO7 and that the highest extent of photodegradation of AO7 shows the composite SITI-8-22. Kamaruddin and Stephan [31] studied the photodegradation activity of quartz-TiO₂ composites (containing 10, 20, 30 and 40 wt.% of TiO₂) prepared using titanium tetrapropylate as TiO₂ precursor. They used methylene blue (MB) as the model organic compound and observed the highest photodegradation activity for the composite calcined at 650 °C and containing 20 wt.% of TiO₂ which showed approx. 85% decrease in initial MB concentration after 10h long irradiation. Although it is not possible to directly compare the photodegradation activity of the SITI composites with the composites prepared by Kamaruddin and Stephan (the setup of photodegradation testing is different) both studies shows the quartz-TiO₂ composites as photocatalytically active and similarly indicate the composites with approx. 20 wt.% of TiO₂ calcined at the temperatures higher than 600 °C as the best performing.

Analyzing the data obtained during the characterization of the prepared SITI composites it is difficult to find the explanation why the SITI-8-22 composite is the best performing one. Taking into account all of the data it can be outlined the possible synergic effect of amount of the mixed anatase-rutile structure when SITI-8-22 consists of approx. 72 % of anatase and 28 % of rutile which makes this composite the best performing one. Similar observation reported e.g. Bakardjieva et al. [32], where the authors reported the highest photodegradation activity against 4-chlorphenol for the TiO₂ sample consisting of anatase/rutile (77%/23%) mixed structure which was achieved after the calcination of the pure anatase at 825°C. The calcination of the SITI composite at the temperature 900°C promote the anatase-rutile transformation what has a detrimental effect on the photodegradation activity of the composite SITI-9-22 as is evident from Table 5.
3.5. Study of the structure and adhesion forces using molecular modeling

The molecular modeling with an empirical force field as implemented in MS was used to confirm the presence of Ti-O-Si bonds in SITI composites. Despite the fact that no new chemical bonds can form in systems treated by molecular mechanics, they are easily inferred to exist based on the distances between atoms [33]. Therefore, in addition to computing the adhesion energies, special attention has been paid to the measurement of distances between Ti atoms from the base of TiO$_2$ nanoparticles and O atoms from adjacent SiO$_2$ surface.

Three SiO$_2$ substrates (quartz alpha [34]) were built as non-periodic superstructures (see Fig. 10) by cleaving the structure along (001), (011) and (111) crystallographic planes. The formula of each substrate was Si$_{3442}$O$_{6888}$ with the total charge $x = -8$ el.

![Fig. 10](image)

Fig. 10. (a) Initial SiO$_2$(001)/TiO$_2$(001) model. Imperfect crystal structure of the SiO$_2$(001) surface can be seen under the TiO$_2$(001) nanoparticle; (b) optimized SiO$_2$(001)/TiO$_2$(001) model. Fixed and unfixed atoms are displayed in balls-and-stick and van der Waals diameter mode, respectively. Colors: O – red; Si – beige; Ti – grey.

This charge comes from the lack of atoms (4 silicon atoms and 4 oxygen atoms are missing). These atoms are removed from the SiO$_2$ surface in order to obtain the site with imperfect crystal structure (see Fig. 10a). The site with a lack of atoms may originate on the SiO$_2$ surface due to chemical attack by sulfuric acid, which is the main component of the TiOSO$_4$ colloid suspension [35].

The SiO$_2$ substrate models are denoted as SiO$_2$(001), SiO$_2$(011) and SiO$_2$(111). The average size of each SiO$_2$ substrate is 9.0 nm × 9.0 nm. The thickness of each SiO$_2$ substrate is 2 nm. In each SiO$_2$ substrate the bottom atomic layers (7737 atoms) representing bulk material are
fixed. Thicknesses of non-fixed top atomic layers are 0.5 nm (2593 atoms) for each SiO$_2$ substrate.

Three types of TiO$_2$ nanoparticles (anatase structure [36]) were prepared with various crystallographical orientations against the SiO$_2$ substrates. These orientations were (001), (100) and (110). Each TiO$_2$ nanoparticle has 113 atoms (Ti$_{39}$O$_{74}$) a and total charge of $x = +8$ el. For more details about the prepared TiO$_2$ nanoparticles, see Table 6.

**Table 6**
Characteristics of nanoparticles prepared for the initial TiO$_2$/SiO$_2$ models. Value d is the size of nanoparticles in the diagonal direction of the basal planes; h is the height of the nanoparticle; n is the number of atoms in nanoparticle.

<table>
<thead>
<tr>
<th>nanoparticles</th>
<th>n</th>
<th>d [nm]</th>
<th>h [nm]</th>
</tr>
</thead>
<tbody>
<tr>
<td>TiO$_2$(001)</td>
<td>133</td>
<td>1.7</td>
<td>1.1</td>
</tr>
<tr>
<td>TiO$_2$(100)</td>
<td>133</td>
<td>1.7</td>
<td>1.0</td>
</tr>
<tr>
<td>TiO$_2$(110)</td>
<td>133</td>
<td>2.0</td>
<td>0.6</td>
</tr>
</tbody>
</table>

The similarity of the nanoparticles with different crystallographic orientations for possible comparison was warranted by the fulfillment of the two following conditions: (a) the same number of atoms in whole nanoparticle and (b) the same number of atoms in the plane adjacent to the SiO$_2$ substrate (12 titanium and 12 oxygen atoms). Because the distribution of atoms in various crystallographic planes is not the same, the second condition constrained each nanoparticle to have a different size in the diagonal direction of the basal planes (see Table 6).

Initial models of the SITI composite were prepared by anchoring each TiO$_2$ nanoparticle onto two sites (either imperfect or perfect crystal structure) of each SiO$_2$ substrate. The surfaces of SiO$_2$ are large enough to keep the TiO$_2$ nanoparticle anchored on the site with perfect structure unaffected by the site with imperfect crystal structure and by the substrate margin (see Fig. 10). Thus it is possible to compare the adhesion forces for TiO$_2$ nanoparticles anchored on both types of sites.

QEq (charge equilibration) method in MS was used to calculate the atomic charges [37]. The interaction between nanoparticles and substrate is quantified using the adhesion energy

$$E_{ad} = (E_{tot,TiO2} + E_{tot,SiO2}) - E_{tot}$$

(2)

where $E_{tot}$ is the total energy of the composite (i.e., the TiO$_2$ nanoparticle anchored on the SiO$_2$ substrate). $E_{tot,TiO2}$ is the total energy of the TiO$_2$ nanoparticle and $E_{tot,SiO2}$ is the total energy of the SiO$_2$ substrate. These energies are expressed in the unit [kJ] and have been
calculated using the UFF [38] in the MS Forcite module. In our previous works the UFF proved to be the suitable force field for the modeling of such type (i.e. silicate/nanoparticle) of composites [33,39,40].

The models have been optimized using UFF in the MS Forcite module. A smart algorithm (i.e. the cascade of Steepest descent, Conjugate gradient and Quasi-Newton optimization algorithms) was used for the geometry optimization with 50,000 iteration cycles.

Nine initial models have been prepared, optimized and examined. This number does not include modifications via shifting and rotations of the TiO$_2$ nanoparticles along the substrate in order to obtain better optimization results. In fact, more than fifty models were studied.

The optimized models revealed that in both cases (i.e. TiO$_2$ nanoparticles anchored on perfect or imperfect SiO$_2$ sites) TiO$_2$(001) nanoparticle exhibits the strongest adhesion while TiO$_2$(110) nanoparticle exhibits weak adhesion. The highest and the lowest adhesion energies computed using Eq. (2) are in good agreement with the values of average distances between Ti atoms from the base of TiO$_2$ nanoparticles and O atoms from SiO$_2$ surfaces. The higher adhesion energies, the shorter average distance (see Table 7).

**Table 7**

Adhesion energies calculated from models containing TiO$_2$ nanoparticles anchored on either the perfect site of SiO$_2$ substrate without surface defect or the imperfect site of SiO$_2$ substrate without surface defect. Average distances between Ti atoms from the base of TiO$_2$ nanoparticle and O atoms from SiO$_2$ surface (d$_{Ti...O}$) together with the shortest measured d$_{Ti...O}$ distances are also listed.

<table>
<thead>
<tr>
<th>models</th>
<th>adhesion energy [kJ·mol$^{-1}$]</th>
<th>average $d_{Ti...O}$ (shortest $d_{Ti...O}$) [nm]</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>perfect site</td>
<td>imperfect site</td>
</tr>
<tr>
<td>TiO$_2$(001) / SiO$_2$(001)</td>
<td>36282</td>
<td>36211</td>
</tr>
<tr>
<td>TiO$_2$(100) / SiO$_2$(001)</td>
<td>34861</td>
<td>35655</td>
</tr>
<tr>
<td>TiO$_2$(100) / SiO$_2$(011)</td>
<td>29498</td>
<td>30481</td>
</tr>
<tr>
<td>TiO$_2$(110) / SiO$_2$(001)</td>
<td>23724</td>
<td>30281</td>
</tr>
</tbody>
</table>

One can see the stronger adhesion forces between TiO$_2$ nanoparticles and SiO$_2$ imperfect sites with one exception - TiO$_2$(001)/SiO$_2$(001) model - in the first row, where the adhesion energies are nearly the same in both cases of perfect and imperfect sites of SiO$_2$(001) substrate. Nevertheless, the average distance is shorter for the imperfect site. The optimized TiO$_2$(001)/SiO$_2$(001) model can be seen in Fig. 10b.

Average distances in Table 7 are presented together with values of the shortest distances measured in the models. Taking into account that the lengths of Ti-O and Si-O bonds are 0.1973 (or 0.1930) and 0.1522 nm [34,36], respectively, the formation of Ti-O-Si bonds can
be assumed in the TiO$_2$(001) / SiO$_2$(001) model and very probably also in the TiO$_2$(100) / SiO$_2$(001) model. Therefore, computer molecular modeling at least indirectly proved the presence of Ti-O-Si bonds and revealed the most probable mutual crystallographic orientations of TiO$_2$ and SiO$_2$ adjacent surfaces.

4. Conclusions

In the preparation process of the photoactive SITI composites, we used TiOSO$_4$, the widely accessible intermediate from the sulfate process of TiO$_2$ pigment manufacturing, as a cheap precursor of TiO$_2$. In order to minimize the environmental risks of TiO$_2$ nanoparticles and to make the manipulation of this material easier and safer, the TiO$_2$ nanoparticles were anchored to larger particles of pure quartz sand. It was proved that the anchoring does not affect the photoactivity of TiO$_2$ nanoparticles. Such prepared SITI composite represents a promising material, which is suitable for use in wide range of building materials containing both TiO$_2$ and sand (i.e. plasters, etc.). The future plan of our research group is to prepare the mortars based on white cement enriched with TiO$_2$.

Obtained results can be briefly summarized as follows.

1) The quartz sand/TiO$_2$ composites were successfully prepared using the low cost precursor TiOSO$_4$, whereas the yield reaches approx. 87 % with respect to the assumed content of TiO$_2$. It was observed that the content of the sulfur grows with an increase of TiO$_2$ content, and decreases with the temperature of heat treatment. Minimal temperature for anatase to rutile transformation occurs at 700°C in the composite with 39 wt.% of TiO$_2$. Anatase crystallite size increases with calcination temperature and reaches 160 nm in the case of the composite with SITI-9-45 (composite with 39 wt.% of TiO$_2$ calcined at 900°C).

2) Calcination at 800°C caused an increase in the photodegradation activity of the SITI composite containing 22 wt.% of TiO$_2$. This SITI-8-22 composite contains about 28 wt.% of rutile in the total volume of TiO$_2$, which is similar to the standard Degussa P-25 powder containing 25 wt.% of rutile. The longer irradiation time led to the significant increase (more than 100%) of the photodegradation activity.

3) The photodegradation activity of the SITI samples decreases rapidly when more than 50 wt.% of rutile is present in the total volume of TiO$_2$ in sample and also when the size of TiO$_2$ nanoparticles becomes larger than 100 nm.

4) The fact, that SITI-T-22 composites (calcined at 600°C, 700°C or 800°C) exhibits higher photodegradation activity than SITI-T-45 composites calcined at the same
temperatures is the result which is in very good agreement with our intention to find
the photoactive TiO$_2$/SiO$_2$ composite with optimum price-efficiency ratio.

5) The presence of Si-O-Ti bond was not observed at the recorded IR spectra, probably
due to a small number of these bonds in prepared SITI composites. However, the
presence of Si-O-Ti bonds has been proved indirectly by computer molecular
modeling.

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