The swelling transition of lepidocrocite-type protonated layered titanates into anatase under hydrothermal treatment

Huiyu Yuan, Rogier Besselink, Zhaoliang Liao & Johan E. ten Elshof

MESA+ Institute for Nanotechnology, University of Twente, P.O. Box 217, 7500 AE Enschede, the Netherlands.

The common facets of anatase crystals are the (001) and (101) planes. However, the phase transformation from lepidocrocite-type titanate into anatase by hydrothermal processing yields an anatase microstructure with high concentration of exposed (010) planes. The phase transformation of a lepidocrocite-type protonated layered titanate (HTO) into anatase was studied using XRD, TEM, FTIR, and measurement of pH and zeta potential. It was found that HTO is proton-deficient. The phase transformation process begins after uptake of a sufficient number of protons into the lepidocrocite-type structure. With the uptake of protons new hydroxyl groups form on the internal surfaces of the layered titanate and result in a bilayer state of HTO. The phase transformation reaction is a topotactic dehydration reaction in which anatase forms and water is expelled by syneresis.

Titanium dioxide (TiO₂) is receiving a lot of attention because of its promising photocatalytic properties that may possibly find applications in dye-sensitized solar cells, photoelectrochemical conversion cells and the photocatalytic decomposition of hazardous organic species. Anatase-type TiO₂ has been studied intensively for photocatalytic applications because of its high photoreactivity, especially on (010) planes. However, anatase bulk crystals normally expose their (001) and (101) planes rather than their (010) planes. Recently, it was found that anatase crystals prepared by hydrothermal treatment from layered lepidocrocite-type titanates H₁.₀₇Ti₁.₇₃O₄·nH₂O and H₀.₆₈Ti₁.₈₃O₄·nH₂O, or their individual 2D titanate nanosheets, preferentially expose their (010) planes at the crystal surface. Anatase prepared from layered titanates and nanosheets showed high photocatalytic activity and a high adsorption capacity of the N719 dye that is used in dye-sensitized solar cells.

The formation mechanism of anatase has been considered to involve an in situ topotactic transformation reaction in the crystal bulk of the layered titanate H₁.₀₇Ti₁.₇₃O₄·nH₂O. It has been proposed that a distorted layered structure is formed during hydrothermal treatment, which is subsequently transformed into a distorted anatase structure, and subsequently into the normal anatase structure. However, details of the topotactic phase transformation mechanism are lacking. Such information could help in preparing new materials by this approach. We studied the phase transformation from lepidocrocite-type titanates to anatase with H₁.₀₇Ti₁.₇₃O₄·nH₂O (HTO) as model compound because of its simple structure among the topotactic reactants. HTO can easily be obtained from K₀.₈[Ti₁.₇₃Li₀.₂₇O₄] (KLTO) by an ion exchange reaction. KLTO is composed of corrugated host layers of edge-sharing TiO₆ octahedra and interlayer K⁺ ions compensating for the negative charge of the TiO₆ octahedral layers, wherein Li⁺ ions occupy 13.₅% of the Ti octahedral sites in the host layers, as shown in Figure 1. Only 40% of the potassium positions shown in Figure 1 are occupied. After protonation a monolayer of water and H₃O⁺ occupies the interlayer space instead of alkali ions.

In the present paper the behavior of lepidocrocite-type protonated layered titanates derived from KLTO under hydrothermal treatment is reported. We present details of the topotactic reaction of lepidocrocite-type titanates to anatase during hydrothermal treatment. Our results show that the process first goes through a stage of protons uptake, which causes swelling of the layered material. Then, with increasing concentration of protons at a suitable temperature the topotactic phase transformation reaction occurs. It is noted that this work focuses on the phase evolution prior to and in the early stages of transformation rather than on the stage after the dehydration reaction. Details of the phase transformation after topotactic reaction can be found in the work of Wen et al.
Results

The uptake of protons and swelling of the layered structure. We performed a series of experiments to investigate the influence of hydrothermal treatment temperature on HTO pre-dried at room temperature (further denoted as HTO RT) in aqueous solutions at 80–120°C. Figure 2a shows the XRD pattern of HTO RT. It is consistent with a protonated titanate with one hydrous layer. The peak at 2θ = 9.59° indicates that the basal spacing between the layers is 0.92 nm. Figure 2b shows that after HTO RT was hydrothermally treated at 80°C, a small peak corresponding with a basal spacing of 1.15 nm was observed at 2θ = 7.72°, suggesting an expansion of the interlayer distance between the titanate planes of HTO. The increase of ~0.20 nm is approximately equal to the molecular size of water (~0.25 nm). At a hydrothermal treatment temperature of 120°C, the peak intensity increased by a factor of ~16 compared to the peak intensity of a sample prepared at 80°C (Figure 2c). This indicates that the swollen phase was the dominant phase at 120°C. The shoulder between the peaks at 2θ = 7.72° and 9.59° implies that the new phase was derived from HTO RT. Also traces of the anatase phase can be observed in Figure 2c. Figure 2d shows the XRD pattern of HTO RT after repetitive hydrothermal treatment at 120°C, which is discussed in more detail below.

The pH change after 24 h of hydrothermal treatment as a function of treatment temperature is shown in Figure 3. It is noted that no further pH changes occurred when the hydrothermal treatment process was prolonged beyond 24 h. The data revealed an increasing pH with increasing hydrothermal treatment temperature, with an anomaly around 100°C. Repeated hydrothermal experiments at 100°C resulted in final pH values of 7.2, 7.3, 7.6 and 7.7. Hence, the change of pH below 100°C was limited to around pH = 8, although the trend would predict a higher pH. The final pH increased further at temperatures above 100°C. The trend in pH indicates a decreasing proton concentration in the solution with increasing treatment temperature. The reduced proton concentration may have been caused by a competitive uptake of protons by the surface O2− groups of the titanate planes of HTO. This agrees with the XRD data, which suggest an increased spacing between the HTO layers. This swelling process is most likely caused by the effect of the uptake of H3O+ ions.

From lepidocrocite-type titanate to anatase. One sample was first given a single hydrothermal treatment at 120°C for 24 h, after which it was titrated with 2 M HNO3 acid solution to adjust the pH of the solution to 2.8. Then a second hydrothermal treatment was applied. It was observed that no more protons were absorbed after repetitive hydrothermal treatment at 120°C because the pH remained constant. The FTIR spectrum of the supernatant after hydrothermal treatment shows that no additional products were observed in the solution besides water and nitrate ions. The XRD pattern of the final state of the powder product is presented in Figure 2d. The diffractogram indicates that an anatase-type TiO2 phase has formed. The broad and low intensity peaks indicate a low degree of crystallinity.

The microscopic structure of HTO-RT after repetitive hydrothermal treatment at 120°C was further investigated by transmission electron microscopy (TEM). A typical high resolution image of the HTO. The increase of ~0.20 nm is approximately equal to the molecular size of water (~0.25 nm). At a hydrothermal treatment temperature of 120°C, the peak intensity increased by a factor of ~16 compared to the peak intensity of a sample prepared at 80°C (Figure 2c). This indicates that the swollen phase was the dominant phase at 120°C. The shoulder between the peaks at 2θ = 7.72° and 9.59° implies that the new phase was derived from HTO RT. Also traces of the anatase phase can be observed in Figure 2c. Figure 2d shows the XRD pattern of HTO RT after repetitive hydrothermal treatment at 120°C, which is discussed in more detail below.

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The microscopic structure of HTO-RT after repetitive hydrothermal treatment at 120°C was further investigated by transmission electron microscopy (TEM). A typical high resolution image of the
as-formed powder in Figure 4a indicates that the product was poly-
crystalline and the average grain size was about 10 nm. The small
grain size is in agreement with the broad peaks observed with XRD as
shown in Figure 2d. The crystal structure of each grain was analyzed
by performing a fast Fourier transform (FFT) on the high resolution
TEM (HRTEM) image. Two kinds of crystalline planes, i.e. (010) and
(001), were observed in the TEM images. Figure 4b shows an example
of a (010) oriented anatase nanocrystal. The plane distances were
determined to be \( d_{002} = 0.49 \) nm and \( d_{101} = 0.36 \) nm, indicating
an (010) oriented plane\(^{13}\). The FFT at the left top panel of Figure 4b
also revealed that the zone axis is [010] and thus the plane which is
vertical to the TEM electron beam is (010). The HRTEM image of a
(001) oriented nanocrystal is shown in Figure 4c. The FFT revealed a
lattice structure with \( d_{200} = d_{020} = 0.19 \) nm, which is the same as
the bulk value. In our TEM investigation, we found that most of the
external surface could be identified as (010) surface plane, and only a
minor fraction of the surface was found to be (001) plane. Therefore,
the nanocrystals preferentially exhibit their (010) facets.

The mechanism of phase transformation. The zeta potential of
HTO RT as function of pH is shown in Figure 5. The zeta
potential of anatase is shown for the sake of comparison. The zeta
potential is related to surface charge characteristics of the external
HTO surface. It can be seen that the point of zero charge (pzc) of
HTO RT was 3.1. In comparison, the pzc of anatase with particle sizes
of 15 nm (Figure 5) is approximately 6.0\(^{19}\). Hence, HTO is much
more acidic than anatase. At pH > 3 the external surface of the HTO
grains is proton-deficient. Remarkably, we observed substantial
uptake of protons by HTO from solutions with a starting pH of 2.8
upon hydrothermal treatment, with a resulting increase in pH up to
values of ~10. Since the external grain surfaces of HTO are proton-
deficient at pH value of 10, it must be the inner surface planes of HTO
RT that absorb protons strongly. The proton exchange behavior of
the external HTO surfaces and the internal HTO planes is therefore
completely different. Casarin et al. investigated multilayer structures
of protonic titanate nanosheets H\(_2\)Ti\(_3\)O\(_7\) with a step structure by first
principles calculations\(^{20}\). They found that two ribbon oxygen atoms
on adjacent inner surfaces of H\(_2\)Ti\(_3\)O\(_7\), which have a similar bond
configuration as the ribbon oxygen atoms in HTO, share a hydrogen
atom in the form of a hydroxyl bond and a hydrogen bond. The
results of our study are in agreement with their conclusions.

For the sake of comparison HTO dried at 80°C (further denoted as
HTO HT) was also hydrothermally treated. HTO HT has a smaller
interlayer spacing than HTO RT because it contains less interlayer
water\(^{21,22}\). Several reports concluded that dehydrated layered materials
cannot be intercalated by chemical means because of the strong
electrostatic force between the titanate layers and the proton counter
ions\(^{21–23}\). Figure 6 shows the corresponding XRD data. The peak of
HTO RT at \( 2\theta = 9.59° \) shifted to \( 2\theta = 12.42° \) in HTO HT (Figure 6a),
indicating that the basal spacing of 0.92 nm of HTO RT decreased to
0.71 nm upon dehydration (HTO HT). The decrease of 0.21 nm is
close to the ~0.25 nm thickness of a single molecular layer of water\(^{18}\).
The XRD patterns of HTO HT after hydrothermal treatment at 80°C
and 120°C are shown in Figure 6b and Figure 6c, respectively. A
similar peak appeared at \( 2\theta = 7.72° \) (\( d = 1.15 \) nm) as for HTO
RT after hydrothermal treatment at 80–120°C (Figure 2b, c). The
 correspondence indicates that a structure with the same large plane

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Figure 4 | (a) TEM image of anatase-type TiO\(_2\) derived from HTO RT with repetitive hydrothermal treatment at 120°C; (b)–(c) HRTEM images of (010) and (001) oriented nanocrystals, respectively.

Figure 5 | Variation of zeta potential of HTO RT and anatase with pH: ●, HTO RT; ○, anatase. Data for anatase are taken from ref. 19.

Figure 6 | X-ray diffraction patterns of HTO HT. (a) HTO HT in dry state, (b) HTO HT in wet state after hydrothermal treatment at 80°C, and (c) at 120°C.
Spacing formed upon hydrothermal treatment, irrespective of whether the initial sample was HTO RT or HTO HT. However, while the major part of the HTO RT material appears to transform to the state with a large lattice spacing of 1.15 nm, only a minor part of the HTO HT material phase appears to do that. The larger part of the HTO HT material appears to remain in a swollen state of the original dry HTO HT structure with a lattice spacing of 0.71 nm. As shown in Figure 6, the original peak of dry HTO HT at 20 = 12.42° shifted to slightly lower angles and broadened so that a shoulder formed at lower 20. This is a very strong indication of swelling caused by interaction with water. Apparently, the ability of the HTO HT structure to swell is smaller than that of HTO RT. This is probably due to the presence of a barrier to swelling in the HTO HT samples due to strong electrostatic forces. The initial pH of 8.3 of a dispersion of HTO HT in water increased to 8.8 and 9.5 after hydrothermal treatment at 80°C and 120°C, respectively, which indicates that HTO HT is proton-deficient and absorbs protons during hydrothermal treatment just like HTO RT. Nevertheless, the swelling of HTO HT demonstrates that the driving force for the uptake of protons is large, which implies that the process involves a chemical reaction between host layers and protons.

Based on the discussion above, we propose that the bonding configuration changes upon hydrothermal treatment: 1) Before hydrothermal treatment of HTO RT the two oxygen atoms on an adjacent titanate sheet share one hydrogen atom, which forms a hydroxyl bond with one oxygen atom and a hydrogen bond with the other oxygen; 2) During hydrothermal treatment absorbed protons form hydroxyl groups with the hydrogen-bond oxygen atoms on the inner surfaces of titanate layers. A bilayer hydroxyl bond configuration forms when all the oxygen atoms on the inner surfaces of titanate layers are bonded to their own hydrogen atom.

To validate this hypothesis, FTIR was used to characterize the bonding configuration. Figure 7 shows the FTIR results of HTO RT before and after hydrothermal treatment at 120°C. The FTIR spectra of anatase derived from HTO RT are also shown. The data indicate that two broad bands around 3400 and 1640 cm⁻¹, which are assigned to stretching and bending vibrations of H2O and/or H3O⁺, respectively, did not change intensity upon hydrothermal treatment. On the other hand, the peak at 920 cm⁻¹ shifted to 880 cm⁻¹ and increased significantly in intensity after hydrothermal treatment at 120°C. H2Ti3O7, H2Ti4O9·1.2H2O, and H2Ti5O11·3H2O with a zigzag structure have been reported to exhibit a characteristic sorption band at 950–1000 cm⁻¹, which was attributed to the bending mode of hydroxyl groups. Based on that interpretation, Sasaki et al. concluded that no hydroxyl groups are present in lepidocrocite-type protonated layered titanates since no peaks are observed in that range of 950–1000 cm⁻¹. However, it is noteworthy that the present material has a planar rather than a zigzag structure. The basicity of the oxygen atoms on the planar structure is lower than the non-shared oxygen atoms at the stepping corner in the zigzag structure. The reduced basicity results in a weaker O-H bond, thus reducing the force constant of the O-H vibration. This implies that the peak that can be assigned to the hydroxyl group is present at a lower wave number than in the zigzag structures. This agrees with our observation of a peak at 880 cm⁻¹. We recorded the FTIR spectrum of HTO-derived anatase to rule out the possibility that the peak at 880 cm⁻¹ is associated with anatase. Hence, the peak at 920 cm⁻¹ in Figure 7 may be assigned to hydroxyl groups. Upon hydrothermal treatment the intensity of the peak increased significantly, and the interpretation agrees well with the observed pH change and the XRD data. Furthermore, the peak shift from 920 cm⁻¹ to 880 cm⁻¹ indicates that the binding strength of the O-H groups weakened upon hydrothermal treatment. To our best knowledge, the stability of fully hydroxylated lepidocrocite-type titanates has not been reported till date. However, it is known that bilayer (001)-oriented anatase films can be converted through a barrierless pathway into a lepidocrocite nanosheet, which implies that the (001) surface of anatase has a similar chemistry as the (001) surface of lepidocrocite-type titanate. Arrouvel et al. found that the fully hydroxyl-covered surface state is less stable than the partially hydroxyl-covered surface state on the TiO2 (001) surface. Our results indicate that protons were absorbed and participated in the formation of hydroxyl groups, i.e. the surface of the inner HTO layers became hydroxylated. The formation of hydroxyl groups provides an explanation why HTO HT swells under hydrothermal treatment. It is noteworthy that this intermediate state is interesting for the preparation of 2-dimensional HTO nanosheets, since the charges between the interlayers are neutralized, thus facilitating the loss of coherence between HTO planes. However, since the hydroxyl groups formed on the surface of the titanate layers are not stable, they can easily participate in the reaction of hydroxyl groups on adjacent titanate layers once the surface OH concentration is saturated, following the net reaction 2·OH → -O- + H2O.

**Discussion**

The phase transformation from lepidocrocite-type titanate to anatase under hydrothermal treatment has been observed not only in layered titanates and exfoliated nanosheets, but also in titanio nanotubes. The common conditions in those preparation procedures are that: 1) the crystal structure of the host titanate is lepidocrocite-type; 2) protons are present in the solution. Wen et al. observed that the phase transformation from lepidocrocite-type nanosheets to anatase occurred preferably under low basic, neutral, or acidic conditions. Gao et al. observed that the electrical conductivity that was associated with the presence of interlayer H⁺ ions decreased at the temperature at which the phase transformation from lepidocrocite-type nanotubes to anatase occurred. These findings suggest that the concentration of surface hydroxyl groups which results from uptake of protons is a key factor for phase transformation of lepidocrocite-type titanate to anatase.

Based on our results the mechanism of phase transformation from HTO to anatase is thought to involve two steps. Firstly, protons are absorbed by the layered structure of HTO. They form hydroxyl groups of the (internal) surface of the titanate host layer of HTO. The data in figure 3 suggest that hydroxyl formation starts at low temperatures. Secondly, after the hydroxyl groups are formed, the topotactic reaction from layered HTO to anatase occurs. Adjacent hydroxyl groups located on two neighboring layers react with each other to form an oxo (-O-) bond between the layers, generating H2O as byproduct, and yielding anatase upon expulsion of water.

![Figure 7](image-url) FTIR spectrum of HTO RT before (black line) and after (green line) hydrothermal treatment at 120°C and anatase from HTO RT (red line). The vertical dotted line is placed at 920 cm⁻¹.
In summary, we found that the inner surface of dry HTO is proton deficient. Our results suggest that the protons in HTO involve a hydrogen bond configuration in which the hydrogen atom forms a hydroxyl bond with one oxygen atom and a hydrogen bond with the other. The phase transformation from lepidocrocite-type titanate to anatase is a topotactic dehydration reaction between hydroxyl groups that are located on adjacent titanate layers. The phase evolution involves 1) proton uptake into the proton-deficient layered structure of HTO and formation of hydroxyl bonds on the inner layer of HTO, and 2) reaction between adjacent hydroxyl groups located on different layers, forming anatase-type TiO₂ under expulsion of H₂O.

**Methods**

**Materials.** Titanium(IV) dioxide TiO₂ (Ridel-de Haen), molybdenum(VI) oxide MoO₃ (Sigma-Aldrich), anhydrous potassium carbonate K₂CO₃ (Fluka) and lithium carbonate Li₂CO₃ (Ridel-de Haen) had a purity of 99.0% or higher and were used as received. Deionized water was used throughout the experiments.

**Preparation of lepidocrocite-type protonated layered titanates.** The Kₓ[Liₓ,Ti₁₋ₓO₂]ₙ precursor was obtained with a flux method as described by Tanaka et al.2 The resulting KLT0 powder was washed 3 times in 250 mL water to remove KₓMoO₃. Then 5.5 g KLT0 powder was dispersed in a 2 mol/L HNO₃ solution (250 mL) at room temperature while stirring. The acidic solution was replaced daily by a fresh one by decantation. After treatment for 3 days to completely remove all alkali metal ions, the acid-exchanged crystals H₁.₀₇Ti₁.₇₃O₄ were obtained. The sample was ultrasonicated for 10 min to get a nicely dispersed solution and the pH of the samples was titrated manually with HCl aqueous solution (0.2 mol/L).

**Hydrothermal treatment of HTO.** 0.1 g of HTO RT was weighed and mixed with 30 mL water without any additives and placed in a glass bottle for hydrothermal experiments. The closed glass bottles were placed in an oven at temperatures ranging from 60 to 140 °C for 24 h. Another part of the HTO RT powder with 30 mL water was hydrothermally treated at 120 °C and then air dried, and finally slightly ground to get HTO powder (HTO RT). The pH of the HTO RT powder was dried at 80 °C to release all intercalated water in the interlayer space, so that finally a dried HTO powder was obtained (HTO HT).

**Characterization.** Powder X-ray diffraction (XRD) data were acquired on a Bruker D2 PHASER (Cu Kα radiation with a wavelength of 0.15406 nm). Measurements on wet state samples were conducted after vacuum filtration (glass microfiber filter) drying unless mentioned otherwise. Transmission electron microscopy (TEM) observation was performed on a Philips CM30ST-FEG at 300 kV, and the sample was supported on a carbon microgrid. The pH was measured by a Fisher Scientific Accumet Research AR15 pH meter. The Fourier transform infrared (FTIR) spectra were recorded by a Bruker Fourier Transform Infrared Spectrometer Model TENSOR 27 with a Pike GladiATR. The zeta potential data were measured by a Malvern Zetasizer Nano ZS. The sample was ultrasonicated for 10 min to get a nicely dispersed solution and the pH of the samples was titrated manually with HCl aqueous solution (0.2 mol/L).

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**Author contributions**

H.Y. conducted the experiments and analyzed the results. H.Y. and J.E.T.E. wrote the main manuscript text and H.Y. prepared figures 1–8. R.B. contributed to the FTIR data acquisition and useful discussion. Z.L. contributed to the analysis of TEM data and useful discussion. All authors reviewed the manuscript.

**Additional information**

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