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ARTICLE

Electrochemical synthesis of coaxial TiO₂-Ag nanowires and their application in photocatalytic water splitting

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A new method for the formation of coaxial TiO₂-Ag nanowires is presented, in which TiO₂ nanotubes were formed by the templated electrochemically induced sol-gel method, followed by thermal annealing. The as-formed TiO₂ nanotubes have been successfully filled with a Ag core using a subsequent electrodeposition step. Coaxial nanowires have a very suitable architecture for photocatalysis, solar cells or batteries due to the high contact area between the two different phases, the large outer surface area exposed to the reactant, and short electron diffusion paths. The coaxial nanowires showed a higher efficiency than empty TiO₂ nanotubes and TiO₂ nanotubes with attached Ag nanoparticles in photocatalytic water splitting. Coaxial TiO₂-Ag nanowires formed H₂ at a rate of $\sim 1.23 \cdot 10^{-3} \pm 0.3 \cdot 10^{-3}$ mol/g·h without deactivation for at least 6 h.

Introduction

One-dimensional nanostructures, such as nanowires and nanotubes, have attracted increasing attention in the last years. The unique physical and chemical properties of these nanostructures make them promising building blocks for various kinds of future applications, such as nanosensors,²⁻⁴ cell trackers,⁵⁻⁷ and self-propelling nanomotors.⁸⁻¹⁰ Because of their very high surface-to-volume ratio, some of the most interesting applications of nanowires and nanotubes are solar cells and photocatalytic water splitting.^{1, 11-13} The physical properties of most semiconductors demand the use of one-dimensional nanostructures, since the diffusion length of minority charge carriers in most semiconductors is of the order of 100 nm, while the absorption depth near the bandgap is of the order of 10 μm .¹⁶⁻¹⁹ In this respect, a maximum of solar light can be absorbed over the nanowire length, while the nanowire diameter is small enough for efficient electron and hole diffusion, thereby leading to high efficiency.²⁰

With a band gap of 3.0 eV, high chemical stability and low synthesis costs, TiO₂ in the anatase phase is a very interesting material for efficient solar hydrogen formation.²⁰⁻²² TiO₂ is therefore the most commonly used material for photocatalytic water splitting and solar cells.²³⁻²⁵ TiO₂ nanotubes are also often used for photocatalytic water splitting,^{14, 15, 26-30} although few authors present actual H₂ formation rates.^{14, 15, 29} Alternatively, TiO₂ nanowires and nanotubes can also be used as micromotors, microfireworks or micropumps upon excitation by UV light,³¹ as catalyst for methanol (electro)oxidation,^{32, 33} or as anode material in Li-ion batteries.³⁴⁻³⁶ The formation of a

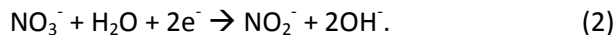
core-shell structure based on TiO₂ nanotubes may also be advantageous for such applications.

TiO₂ nanowires and nanotubes can be formed via several techniques, i.e. hydrothermal treatment,³⁷ anodization of titanium foils,³⁸⁻⁴⁰ atomic layer deposition (ALD),^{41, 42} modified sol-gel route,^{11, 43, 44} sol-gel electrophoresis,^{45, 46} electrochemical deposition,^{47, 48} and electrochemically induced sol-gel synthesis.^{49, 50} As sol-gel electrophoresis and the modified sol-gel route provide nanowires or nanotubes consisting of smaller sol-gel particles, the resulting nanostructures are very brittle with rough surfaces, which decreases the mechanical stability of the formed nanowires. Although a large area of single crystalline TiO₂ nanotubes with tuneable dimensions can be made by anodization of Ti foil, the main disadvantage of this technique is that the formed nanotubes do not contain a conductive back electrode, so that extra steps need to be applied before these nanotubes can be filled with a second material.⁵¹ In contrast, the availability of a back electrode and the synthesis of smooth films or nanostructures makes the electrochemically induced sol-gel method a very promising technique for the synthesis of core-shell nanostructures with a TiO₂ shell.

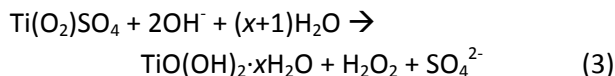
The electrochemically induced sol-gel method for the cathodic electrodeposition of titanium dioxide thin films was first introduced by Natarajan et al.⁵² and was further improved by Karuppuchamy et al.^{53, 54} This method involves dissolving titanium oxysulfate (TiOSO₄) powder in an aqueous solution of hydrogen peroxide (H₂O₂), yielding a peroxotitanate complex (Ti(O₂)SO₄) (equation (1)):



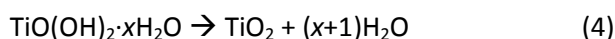
At potentials more negative than -0.9 V vs. Ag/AgCl, nitrate is reduced to form hydroxyl ions (equation (2)),⁵⁴ and consequently the pH in the vicinity of the electrode surface is increased:⁴⁹



This local increase in pH is necessary for the formation of a titanium hydroxide gel as shown in equation (3):⁵⁴



According to differential thermal analysis data of Natarajan et al., water is removed from the gel during thermal annealing around 283 °C (equation (4)), resulting in an amorphous TiO₂ phase.⁵² When the temperature is increased above 365 °C, crystallization into the anatase phase occurs.^{52, 55}



By employing the electrochemically induced sol-gel method within the pores of an anodic aluminium oxide (AAO) template, nanowires and nanotubes can be formed. Nanowires were previously made using templates with pores of 50 nm diameter and smaller.^{49, 50} In the present study, AAO templates with a diameter of 200 nm were used, which resulted in nanotube formation. The mechanism of nanotube formation was previously described by Maas et al., and occurs by removal of water from the titanium hydroxide gel.⁵⁶ The first 6 steps in Figure 1 provide a schematic representation of TiO₂ nanotube formation according to the above reactions.

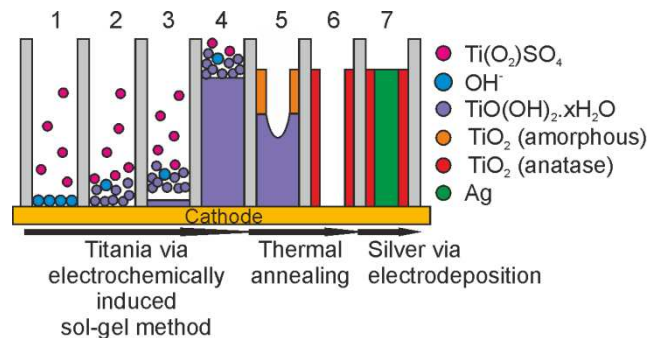


Figure 1: Schematic overview of the preparation of coaxial TiO₂-Ag nanowires using the electrochemically induced sol-gel method: formation of OH⁻ at the electrode (step 1, reaction (2)), reaction of Ti(O₂)SO₄ with OH⁻ to TiO(OH)₂·xH₂O and filling of the template (step 2-4, reaction (3)), drying and thermal annealing of the titanium hydroxide gel to TiO₂ (step 5-6, reaction (4)), and filling of the TiO₂ nanotubes with Ag (step 7).

As mentioned above, an advantage of the electrochemically induced sol-gel method for TiO₂ nanotube formation is the availability of a cathode at one end of the AAO membrane or template. This allows for the formation of coaxial nanowires by filling the TiO₂ nanotubes in a subsequent deposition step with a second phase like silver (step 7 in Figure 1). By combining an *n*-type semiconductor like TiO₂ with a noble metal, a Schottky barrier is formed when the work function of the metal is smaller than that of the *n*-type semiconductor, which diminishes electron-hole recombination.²⁰ This is schematically depicted in Figure 2. Upon illumination, electrons and holes are formed in the TiO₂ shell layer (step 1). As the electrons are transferred to

the silver core of the wires, the holes go to the surface of the TiO₂ phase (steps 2), where they are known to react with water to form H⁺ ions and O₂ (steps 3). In the Ag phase, at the outer ends of the coaxial nanowire, electrons and protons can combine to form H₂ (step 4).²⁰⁻²²

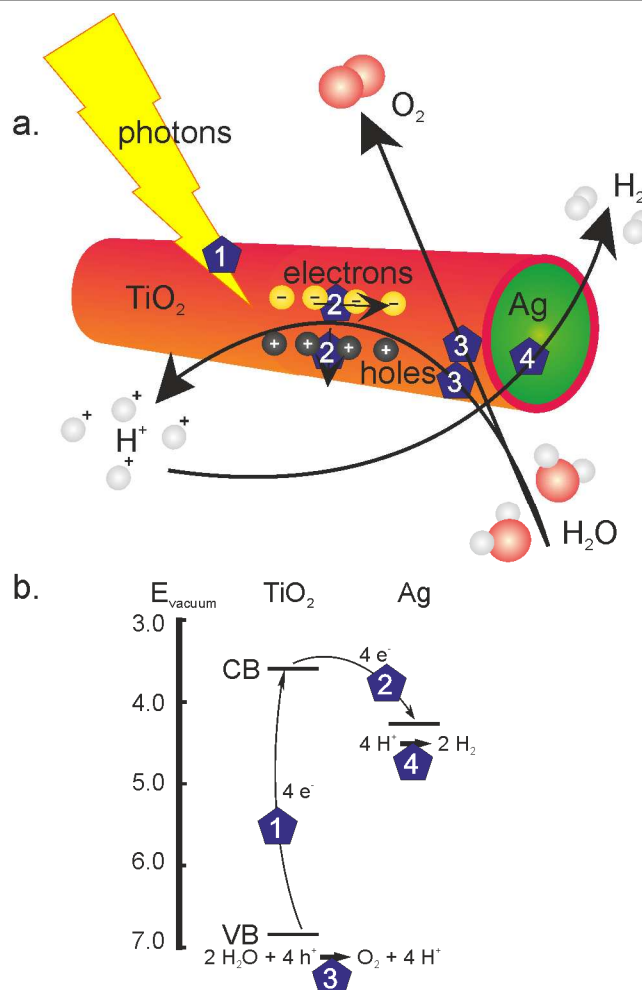


Figure 2: (a) Schematic representation and (b) band energy diagram of water splitting by coaxial TiO₂-Ag nanowires in an alkaline solution: formation of electrons and holes in the TiO₂ phase (step 1), electron-hole separation at the TiO₂-Ag interface (steps 2), water oxidation at the TiO₂ surface (steps 3) and H₂ formation at the Ag surface (step 4).

In this study, we employed the electrochemically induced sol-gel method for the formation of TiO₂ nanotubes and filled these nanotubes with a silver phase using electrodeposition. The photocatalytic activity of the coaxial TiO₂-Ag nanowires in splitting water was investigated in an alkaline solution. Furthermore, the photocatalytic activity of the coaxial TiO₂-Ag nanowires was compared with similar TiO₂ nanostructures with/without Ag that were also prepared in this study, and with values reported in other studies.

Results and discussion

Since it was found that annealing within a template resulted in crystallization at a higher temperature than in films, we investigated the crystallinity of different samples annealed at variable temperatures (Figure 3). The figure shows that TiO₂ within an AAO template crystallized into a mixture of anatase,

rutile and brookite between 525 °C and 550 °C, while unconstrained TiO₂ gel made via the same procedure crystallized below 400 °C (data not shown, see e.g. Natarajan et al).⁵² Also for TiO₂ powder made via a different sol-gel process, it is known that TiO₂ crystallizes below 400 °C.⁵⁷ A possible explanation for the large difference found in crystallization temperatures between TiO₂ powder and the nanotubes formed in this study may be clamping of the TiO₂ phase on the sidewalls of the AAO template. Clamping was also previously found to retard the crystallization of BaTiO₃ and SrRuO₃ films.⁵⁸

After annealing the titania gel inside an AAO membrane for 2 h at 650 °C, the TiO₂ gel was crystallized into a polycrystalline mixture of the anatase, rutile and brookite polymorphs of TiO₂, with no preferred orientation. It is noted that the presence of all three polymorphs of TiO₂ is known to have a positive influence on the photocatalytic activity as previously discussed in several other studies.^{59, 60} After annealing at 550 °C, the formation of an unidentified impurity phase was observed, which is denoted with an asterisk. For the TiO₂ nanotubes, crystallite sizes of approximately 42 and 45 nm were found when annealed at 550 and 650 °C, respectively, showing negligible crystallite growth. For TiO₂ films made via the same method we observed exponential crystallite growth with sizes of 19 and 28 nm when annealed at 550 and 650 °C, respectively.

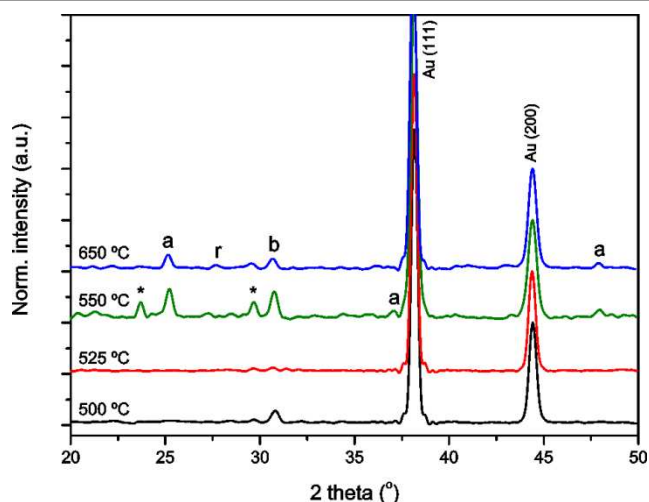


Figure 3: XRD patterns of TiO₂ nanotubes annealed for 2 h at 500 °C (black line), 525 °C (red line), 550 °C (green line), and 650 °C (blue line), with the tubes still inside an AAO membrane; Diffraction peaks were assigned to Au (PDF # 00-004-0784), anatase TiO₂ (a, PDF # 00-021-1272), rutile TiO₂ (r, PDF # 00-021-1276) and brookite TiO₂ (b, PDF # 00-029-1360); An unknown impurity phase is denoted with an asterisk (*).

The cross section of a membrane containing TiO₂ nanotubes after annealing at 650 °C shows a gradual decrease in thickness of the TiO₂ coating on the AAO pore walls from ~55 nm at the bottom of the membrane (Figure 4b), which is the cathode side, to ~10 nm at the top of the membrane (Figure 4a). The thickness gradient can be explained by the formation of a thicker gel near the electrode as OH⁻ is formed here. To put the thickness of the TiO₂ phase into perspective of the proposed application in solar water splitting, a wall thickness less than 20 nm would provide efficient charge carrier separation as the minority carrier diffusion length in TiO₂ is approximately 20 nm.⁶¹ This means that the most efficient charge carrier separation takes place only in the upper part of these nanotubes.

On the other hand, the chance that a photon is absorbed by the material is higher when the titania wall is thicker. It is expected that a lower concentration of KNO₃ in the electrolyte solution can provide TiO₂ nanotubes with thinner walls, but a trade-off has to be made between carrier diffusion length, mechanical stability of the nanotubes and degree of photon absorption. After dissolving the membrane, a droplet of nanotube solution was placed on a silicon wafer for visualization with SEM. These nanotubes appear to be smooth at first sight, but a closer look reveals that they are composed of very small grains (Figure 4c).

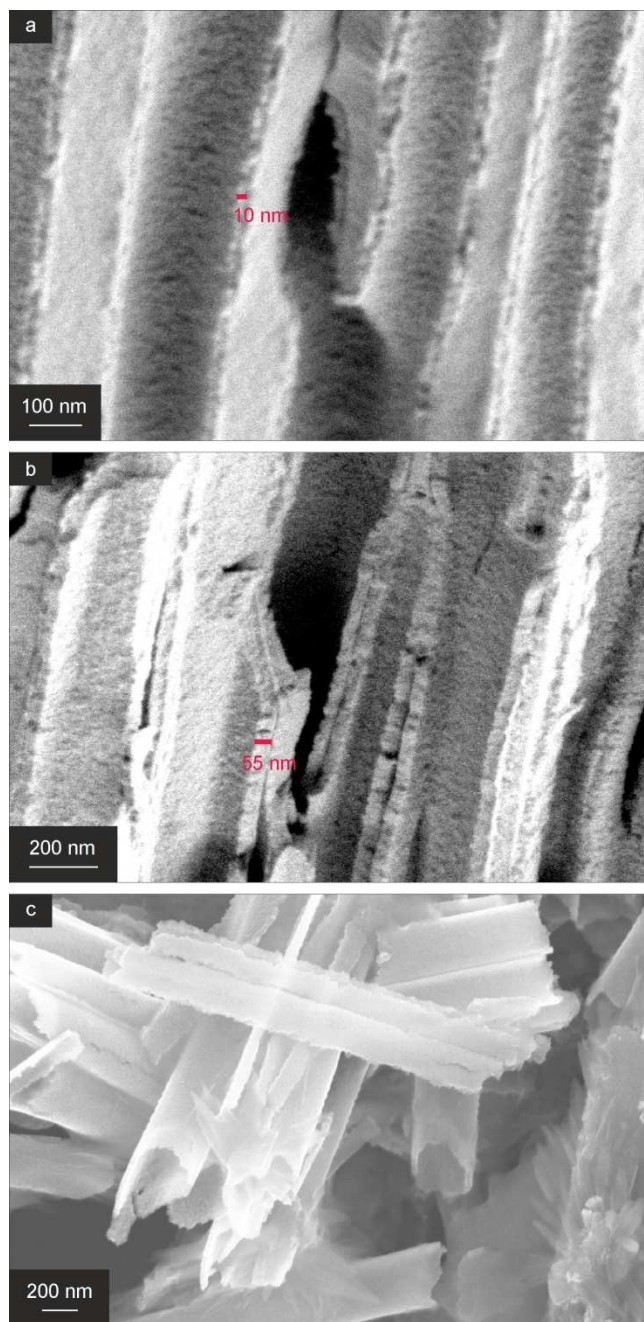


Figure 4: Cross section SEM images of TiO₂ nanotubes inside the membrane (a) at the top of the membrane, (b) at the bottom of the membrane, and (c) TiO₂ nanotubes after dissolving the membrane.

After formation of a titania gel inside the AAO membrane, a fraction of the samples was dried overnight at 100 °C to remove water, while the other fraction was annealed for 2 h at 650 °C to form crystalline nanotubes. When silver was deposited inside a membrane after drying at 100 °C, TiO₂ nanotubes containing isolated Ag nanoparticles (TiO₂/Ag nanotubes) were formed after Ag electrodeposition and subsequent annealing (Figure 5a). The reason for this is that the Ag⁺ ions could penetrate the gel, because the titania phase was not completely dense after exposure to 100 °C. As a result, Ag nuclei formed where such ions combined with an electron. During subsequent heat treatment at 650 °C, these Ag nanoparticles were transported to the pore walls along with the collapse of the titania gel.

However, when Ag was electrodeposited after high temperature annealing of the TiO₂ gel at 650 °C, solid TiO₂ nanotubes formed first, and dense silver nanowires were deposited inside these tubes. This resulted in the formation of coaxial TiO₂-Ag nanowires (Figure 5b and Figure 6).

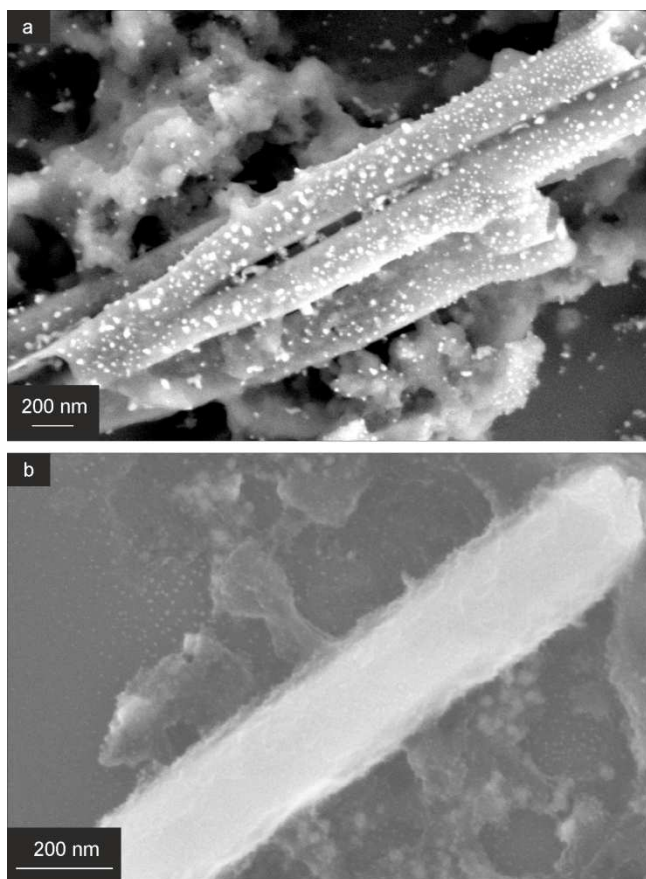


Figure 5: SEM images of (a) TiO₂/Ag nanotubes from two step electrodeposition process with intermediate TiO₂ gel drying at 100 °C, and (b) a coaxial TiO₂-Ag nanowire from two step electrodeposition process with intermediate TiO₂ tube solidification at 650 °C. See main text for details.

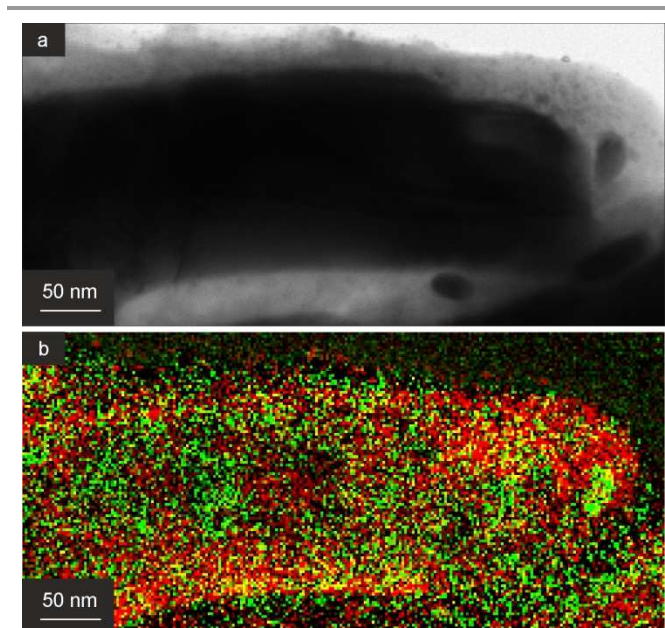


Figure 6: (a) TEM image and (b) EELS TEM map of a TiO₂ nanotube (red) filled with Ag (green)

A TEM image of a TiO₂ nanotube with Ag core is shown in Figure 6a. Especially the Electron Energy Loss Spectroscopy (EELS) TEM map in Figure 6b clearly shows the architecture of the core-shell structure, with a higher concentration of Ti (red) at the outside of the coaxial nanowire, and a higher concentration of Ag (green) at the interior of the tube.

The photocatalytic activities in photocatalytic water splitting of coaxial TiO₂-Ag nanowires, empty TiO₂ nanotubes and TiO₂/Ag nanotubes with isolated Ag nanoparticles were compared in Figure 7a after 3.5 h of irradiation. The coaxial TiO₂-Ag nanowires formed significantly more H₂ than the TiO₂ nanotubes and the TiO₂/Ag nanotubes. H₂ formation was absent in the reference reactor containing aqueous 1 M NaOH solution. Only the intensity of the H₂ signals from the coaxial TiO₂-Ag nanowires was sufficient for automatic integration, so for this system a long-term measurement was performed (Figure 7b). Continuous H₂ formation was observed for a period of 6 h at a rate of $6.2 \pm 1.5 \cdot 10^{-6}$ mol/h, after which the reaction was terminated.

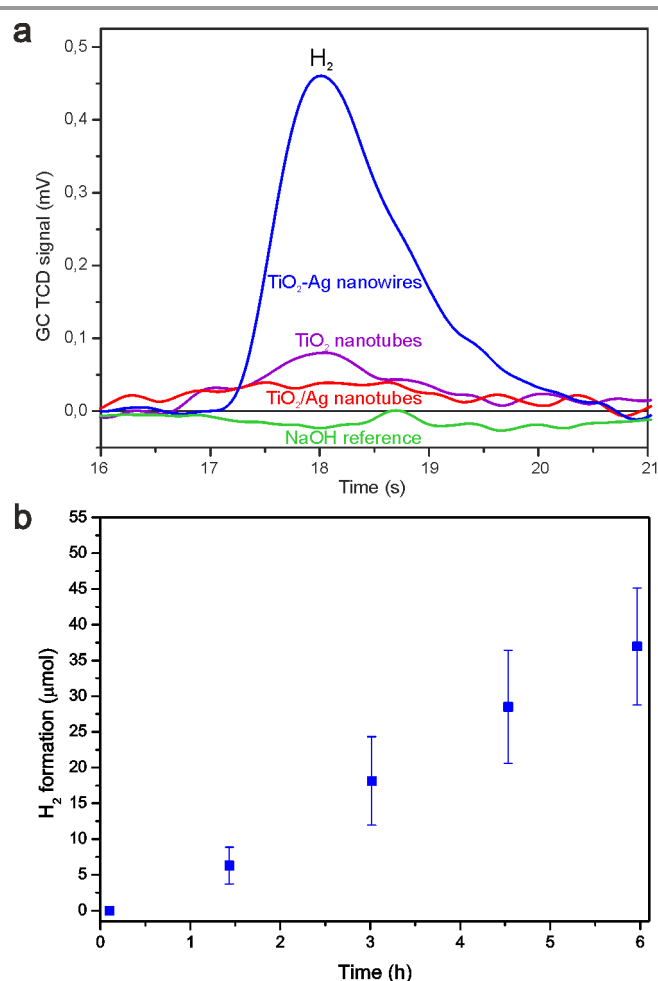


Figure 7: (a) GC TCD peaks showing H₂ formation from TiO₂ nanotubes (purple line), TiO₂/Ag nanotubes (red line), TiO₂-Ag nanowires (blue line) and NaOH baseline (green line). (b) Moles of H₂ formed from coaxial TiO₂-Ag nanowires during 6 h measurement in 1M NaOH.

Several possible explanations can be given for the higher H₂ formation rate by the coaxial TiO₂-Ag nanowires compared to the TiO₂ nanotubes: (1) The availability of an H₂ evolution catalyst such as Ag, which decreases the chance for electron-hole recombination, (2) the use of the metallic Ag core as a mirror to reflect photons back into the TiO₂ shell, which could approximately double the number of photons captured by the photocatalyst, and/or (3) the use of the Ag core for enhancement of the light absorption due to localized surface plasmon resonance of the Ag nanowire.

Comparing the TiO₂/Ag nanotubes with the coaxial nanowires, several possible explanations can be given for the higher H₂ formation rate of the latter: (1) The availability of a larger amount of Ag catalyst with an increased interface between the photocatalyst and H₂ evolution catalyst, which shortens the electron diffusion length and decreases the chance for electron-hole recombination in the coaxial nanowires, and/or (2) since most of the Ag particles are located at the outside of the TiO₂ nanotubes, these particles may prevent some of the photons from reaching the TiO₂ phase which decreases the efficiency of this system. It is however unlikely that the second possibility alone can explain the large difference.

For comparison with other data from literature, the observed H₂ formation rate was corrected for either the weight of nanowires that was put in solution or the initial surface area of the template, resulting in an H₂ formation rate of $1.2 \cdot 10^{-3} \pm 0.3 \cdot 10^{-3}$ mol/g·h and $0.18 \cdot 10^{-6} \pm 0.04 \cdot 10^{-6}$ mol/g·h, respectively. As can be seen from Table 1, this was higher than what we measured previously using segmented ZnO|Ag nanowires, where H₂ formation at a rate of $6.9 \cdot 10^{-6}$ mol/g·h was observed.¹ Especially the increased surface area between the photocatalytic oxide phase and the H₂ evolution catalyst, and the shorter diffusion length for electrons to cross the oxide/metal junction are likely explanations for the difference of ~3 orders of magnitude. Obviously, the use of different photocatalytic materials and the fact that methanol was used as hole scavenger by the ZnO|Ag nanowires also influence the efficiency.

Table 1: Comparison of H ₂ formation from different nanowire and nanotube systems.						
Nanowire/ tube system	Reference	H ₂ formation mol/g·h	H ₂ formation mol/g·h	Dimensions diameter (TiO ₂ thickness) x length	Electrolyte	Light source
Coaxial TiO ₂ -Ag nanowires	This work	$1.2 \cdot 10^{-3}$	$0.18 \cdot 10^{-6}$	200 nm (10-50 nm) x 40 μm	1.0 M NaOH	10 W Hg lamp
Coaxial TiO ₂ -Ag nanowires by ALD	This work	$0.26 \cdot 10^{-3}$		175 nm (20 nm) x 8 μm	1.0 M NaOH	10 W Hg lamp
Segmented Ag ZnO nanowires	1	$6.9 \cdot 10^{-6}$		200 nm x 6 μm	4:1 v/v MeOH/H ₂ O	60 W Hg lamp
Anodized TiO ₂ with Pt mesh counter electrode	14	$2.83 \cdot 10^{-3}$		60 nm (10 nm) x 1.5 μm	0.01 M KOH 1.0 mM KI	400 W Hg lamp
Anodized TiO ₂ with Pt counter electrode in second compartment	15		$97 \cdot 10^{-6}$	75 nm (20-70 nm) x 6 μm	1.0 M KOH	350 W Xe lamp
Anodized TiO ₂ containing evaporated Pt	29		$14 \cdot 10^{-6}$	150 nm (30 nm) x 650 nm	0.01 M KOH 1.0 mM KI	400 W metal hydride lamp

We also compared the H₂ formation rate from the coaxial TiO₂-Ag nanowires with an alternative coaxial TiO₂-Ag nanowire system in which the TiO₂ phase was prepared by Atomic Layer Deposition (ALD), and the Ag phase by electrodeposition. An SEM image of TiO₂ nanotubes made by ALD is shown in Figure 8. Compared to the SEM image in Figure 4c, the nanotubes made by both techniques have approximately the same dimensions, but the nanotubes made by ALD appear much smoother. It was confirmed by XRD that the TiO₂ nanotubes made by ALD are also polycrystalline without preferred orientation, the difference being that they are composed of large anatase crystallites with a size of 250-300 nm as shown in ref.⁶² Using the same setup for photocatalytic water splitting, we found a H₂ formation rate of 0.26·10⁻³ mol/g·h for the ALD-based system, which is ~4.7 times lower than the rate found for the electrochemically deposited TiO₂-Ag system and also lower than the TiO₂ nanotubes without Ag. Since the TiO₂ nanotubes deposited by ALD were polycrystalline anatase with a larger crystallite size, it was expected *a priori* that the photocatalytic activity of these nanowires would be higher, since the concentration of grain boundaries that can diminish electron-hole recombination is lower. On the other hand, it is known that photocatalytic water splitting is more efficient on specific facets, so when these facets are not exposed to the solution this will lead to a lower H₂ evolution rate. Of course, the different TiO₂ polymorphs that were observed in both systems, purely anatase for the ALD-based system vs. a mixture of anatase, rutile and brookite for the system prepared via the electrochemically induced sol-gel method, might also influence the photocatalytic activity. Also other effects like impurities or secondary phases can have a positive effect on the photocatalytic properties of the electrochemically synthesized system. As listed in Table 1, there are also some other minor differences between the two systems in nanowire length, diameter and TiO₂ shell thickness, but it is not expected that these differences alone can explain the large difference in photocatalytic activity.

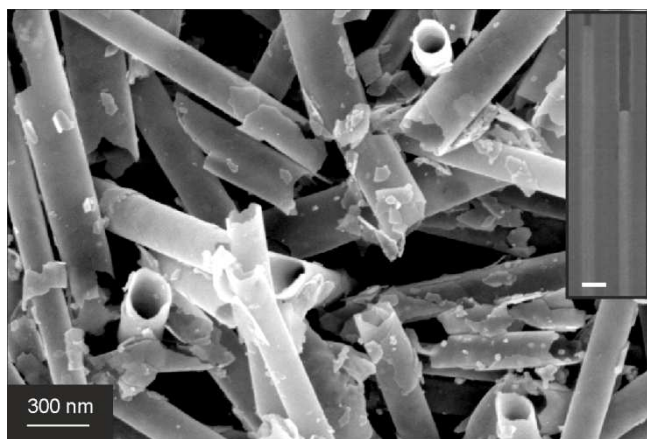


Figure 8: SEM image of TiO₂ nanotubes made by ALD before Ag filling, after release from the template. The inset shows coaxial TiO₂-Ag nanowires inside the membrane, where the TiO₂ phase is visible as a bright white phase covering the pore walls at the top of the figure, and the Ag phase is visible as a solid nanowire at the bottom of the figure.

Only a few other publications report H₂ evolution from TiO₂ nanotubes, and these are listed in Table 1. The geometries and process conditions vary widely between these studies, so that direct comparison of these data should be done with care. Besides one other publication on hydrogen evolution by

autonomous nanowires in methanol,¹ all other studies involve photoelectrochemical cells with compositionally distinct, and in some cases spatially separated, hydrogen and oxygen evolution catalysts. Only one publication reports the H₂ evolution rate per weight of used material for TiO₂ nanotubes, resulting in an evolution rate of 2.83·10⁻³ mol/g·h H₂.¹⁴ This is a factor of 2.3 higher than our current result, and it might be explained by the difference in crystallinity of TiO₂, as anodization can result in the formation of single crystalline TiO₂ nanotubes, while polycrystalline TiO₂ nanotubes were made in the current research. Two other main differences are that a light source with higher energy was used in ref.¹⁴, and that an IO₃⁻/I⁻ redox couple was used as a redox mediator to prevent recombination of H₂ and O₂ into H₂O. Both these differences also drastically influence the H₂ evolution rate.

Others calculated the H₂ formation rate per unit surface area of the anodized Ti foil, resulting in H₂ formation rates of 97·10⁻⁶ mol/g·h,¹⁵ and 14·10⁻⁶ mol/g·h,²⁹ which are both much higher than our current result using coaxial TiO₂-Ag nanowires made via the electrochemically induced sol-gel method. Part of this difference may be explained by the different methods used to measure the surface area: the surface area of the AAO template before dissolution vs. the anodized surface area of the Ti foil. Both authors used Pt as counter electrode,¹⁵ or as thin layer on top of the nanotubes for enhanced photocatalytic activity.²⁹ Also here a light source with higher energy was used in both studies and an IO₃⁻/I⁻ couple was used as redox mediator in one of the two studies to prevent recombination of H₂ and O₂ into H₂O.²⁹ Another method to prevent recombination of H₂ and O₂ is to split the electrodes in a two-compartment setup.¹⁵ On the other hand, even though efficiencies could be increased by preventing H₂ and O₂ recombination, we have shown that coaxial TiO₂-Ag nanowires are able to split water autonomously without addition of sacrificial agents, redox mediators or external power supply. This can be of advantage for self-fuelling nanodevices in aqueous environments, or to generate H₂ at a specific location.

Next to the possible explanation of different crystallinity, use of redox mediators and two-compartment setups, it is still difficult to directly compare these results, because every author uses different light sources and electrolytes that both have a large impact on the efficiency for H₂ formation. In that respect, a better way for comparison would be to keep the nanowires attached to the substrate for photocurrent measurements. Also the applied potential should be compensated for the reference electrode and pH of the solution, as illustrated by Seabold et al.⁶³ Unfortunately, with the AAO templates available for this study, it was not possible to keep the nanotubes or nanowires standing on the substrate after dissolving the template because the nanowire/nanotube connection to the bottom was mechanically too weak. Additionally, standardized simulated AM1.5 solar illumination with an energy of 100 mW/cm² should be used in all measurements, but unfortunately these solar simulators are expensive and not available in every lab.

Several strategies can be pursued to further improve the activity and H₂ formation rate. One of these strategies includes changing the alkaline electrolyte for water splitting to either an acidic or neutral electrolyte.³⁰ The second method includes the use of a sacrificial agent, e.g. methanol, or redox mediator, e.g. KI, to increase the H₂ formation rate.^{26, 64-66} As a third option, the activity for water splitting can be improved by adding an oxidation catalyst, e.g. Mn₃O₄, for more efficient O₂ formation. In the fourth method, the dimensions of the coaxial TiO₂-Ag nanowires could be changed in order to find an optimum in

total diameter, TiO₂ shell thickness and nanowire length. These methods should be systematically investigated in future research.

Experimental

All chemicals were purchased from commercial sources and used without further purification. Titanium oxysulfate (TiOSO₄, synthesis grade), hydrogen peroxide (H₂O₂, 35 wt%), boric acid (H₃BO₃, purity 99.97%), sodium hydroxide (NaOH, purity >98%) and titanium(IV)isopropoxide (99%) were purchased from Sigma-Aldrich. Silver nitrate (AgNO₃, purity >99%), nitric acid (HNO₃, 65%), potassium nitrate (KNO₃, purity >99%), chromium(VI)oxide (CrO₃, purity 99.5%) and phosphoric acid (H₃PO₄, >85%) were purchased from Acros Organics. Oxalic acid (C₂H₂O₄, p.a. grade) was purchased from Merck Millipore. Silver sulfate (Ag₂SO₄, purity 99.999%), diammoniumcitrate (purity >99%) and potassium thiocyanide (KSCN, purity 98%) were purchased from Alfa Aesar. Milli-Q water with a resistivity of 18.2 MΩ·cm was used in all experiments.

Anodized aluminium oxide (AAO) membranes and templates with a pore diameter of 200 nm were either purchased commercially (Anopore®, Whatman Inc.), or custom-made. The commercial AAO membranes contained open pores on both sides and are therefore referred to as “membrane” in the text. A back electrode of AuPd was sputtered on one side of the commercial membranes as cathode using a tabletop sputterer (SC7640, Quorum Technologies). Prior to deposition, the backside of the AuPd electrode was isolated to ensure exclusive deposition inside the pores of the membrane and avoid deposition on the external surface of the back electrode.

The custom-made AAO templates (referred to as “template”) contained self-ordered arrays of aligned cylindrical pores with narrow diameter distribution and is closed on one side. Self-ordered AAO was prepared by two-step anodization following procedures introduced by Masuda et al.⁶⁷ In the first step, an Al chip (Goodfellow, purity > 99.999 %) was anodized at 195 V for 10 h using 1 wt% H₃PO₃ cooled to 0-1 °C as electrolyte solution. The resulting porous alumina layer was selectively removed by etching with an aqueous solution of 0.18 M CrO₃ and 0.72 M H₃PO₄. As a result, patterned Al substrates with hexagonal arrays of indentations were obtained. In the second step, hierarchical pores consisting of macroporous segments branching into dendritic mesopores were obtained by adapting a method introduced by Nielsch et al.⁶⁸ Using 1 wt% H₃PO₄ at 0-1 °C as electrolyte solution, the anodization voltage $U(t)$ was directly reduced from 195 V to 80 V according to

$$U(t) = U_0 e^{-t/\tau}, \quad (5)$$

where U_0 is the start voltage of 195 V, τ is a constant, i.e. $\tau = 191.7$ min in this case, and t is the anodization time (171 min to reach a voltage of 80 V). Then, the electrolyte was replaced by an aqueous solution of 0.3 M C₂H₂O₄, and the temperature was adjusted to 10 °C. Following the profile of equation (5) with $U_0 = 80$ V and $\tau = 14.0$ min, the anodization voltage was further reduced to a target value of 0.01 V, which was reached after 125 min. Subsequently, the AAO layer was rinsed with deionized water. Prior to electrodeposition, the aluminium disks were isolated with nail varnish to ensure exclusive deposition inside the pores of these AAO templates.

The AAO membranes and templates were used as working electrode in a three electrode setup, with a Pt sheet (Metrohm Autolab) as counter electrode and Ag/AgCl in 3 M KCl (Metrohm Autolab) as reference electrode. All potentials are given with respect to the reference electrode. These electrodes were connected to an Autolab PGSTAT 128N potentiostat. TiO₂ nanotubes were formed from an electrolyte containing 0.02 M TiOSO₄, 0.03 M H₂O₂, 0.05 M HNO₃ and 0.25 M KNO₃ with a pH of 1.4. Deposition occurred at -1.00 V for 6 h. After deposition, the gel was either dried overnight at 100 °C or annealed for 2 h at 650 °C in a tube furnace. Ag was electrodeposited for 15 min at +0.10 V in an aqueous electrolyte containing 0.20 M AgNO₃ and 0.10 M H₃BO₃ with pH adjusted to 1.5 using HNO₃. In the final steps, the nanowires or nanotubes were collected by dissolving the AAO membranes or templates in an aqueous solution of 1 M NaOH or in an aqueous solution of 0.18 M CrO₃ and 0.72 M H₃PO₄, respectively, followed by detachment from the back electrode by gentle swirling. In order to get rid of the NaOH or CrO₃ in the nanowire/nanotube solution, the solution was washed at least 5 times with water using a Hermle Z36HK centrifuge at 10,000 rpm for 5 min.

The TiO₂ nanotubes made by Atomic Layer Deposition (ALD) were also prepared inside custom-made AAO membranes with the same dimensions as the templates used for TiO₂ nanotube formation by the electrochemically induced sol-gel method. Amorphous TiO₂ was deposited at 120 °C using titanium(IV)isopropoxide preheated at 70 °C as metal containing reactant and water vapor as oxygen source. The porous membranes were coated using the exposure-mode sequence due to the high aspect ratio of these nanostructures, where a full ALD cycle consisted of 2 s pulse, 40 s exposure and 60 s purging for the titanium precursor, followed by 0.5 s pulse, 40 s exposure, and 60 s purging for the water vapor. Dried Ar was used as carrier and purging gas and the process was carried out in a commercial ALD reactor SAVANNAH 100 (Cambridge NanoTech Inc., USA). After thermal annealing for 3 h at 400 °C and Au sputtering, Ag was electrodeposited at -1.2 V in a two-electrode setup using Pt as the counter electrode from an aqueous electrolyte containing 0.027 M Ag₂SO₄, 0.88 M diammoniumcitrate and 1.08 M KSCN at pH 4.5. The procedures used for dissolution of the AAO membrane and washing were the same as above.

Samples were measured with X-ray powder diffraction (XRD) to confirm the formation of crystalline TiO₂ phases in the membranes using a PANalytical X'Pert Pro (PANalytical B.V., Almelo, The Netherlands) instrument. Samples were measured from $2\theta = 20$ -50°, with step sizes of 0.026° and 600 s per step using a PIXcel 1D scanning line detector. The spectra were further analyzed using the X'Pert Highscore Plus software package (version 3.0e). Scanning Electron Microscopy (SEM) images were taken using a Zeiss HR-LEO 1550 FEG SEM or a Zeiss Merlin HRSEM. Transmission Electron Microscopy (TEM) images and Electron Energy Loss Spectroscopy (EELS) maps were taken with an Analytical TEM from FEI Instruments.

The photocatalytic activity of TiO₂ nanotubes with and without Ag core was investigated by water splitting experiments in a custom-made setup consisting of 8 top-illuminated reactors with a volume of 36.4 mL each, connected to a Compact gas chromatograph (GC) (Interscience BV) equipped with a thermal conductivity detector (TCD). 5-14 mg of nanotubes/wires (1 completely filled AAO template with a diameter of ~3.5 cm) was dispersed in 15 mL of a 1 M NaOH

stock solution (pH 13) that was bubbled with Ar gas for at least 30 min to remove a large part of the O₂ from the solution. Then the reactors were evacuated to 3 mbar and subsequently filled with Ar to remove all air from the head space in the reactors. After three evacuations, a 120 Watt high pressure Mercury lamp (Dr. Gröbel UV-Elektronik GmbH) was turned on, of which the emission was divided over 12 optical fibers. During illumination, the formed H₂ was measured subsequently in each reactor. In a standard procedure, the composition in each reactor was measured every 15 min for 13 times. In a long-term measurement, the formed H₂ gas was measured 5 times using a measurement interval of 1.5 h.

Conclusions

The electrochemically induced sol-gel method was used to prepare anatase TiO₂ nanotubes in AAO membranes and templates with a diameter of 200 nm. By varying the intermediate thermal annealing process of the titania gel phase, the location where Ag deposited during the subsequent electrodeposition step in the same template could be tuned. Intermediate heating at 100 °C resulted in the formation of Ag nanoparticles incorporated in the shell of the TiO₂ nanotubes, while thermal annealing at 650 °C led to TiO₂ nanotubes filled with a continuous Ag phase, thus forming coaxial TiO₂-Ag nanowires. These coaxial TiO₂-Ag nanowires showed improved photocatalytic activity for direct water splitting due to the favorable geometry of core/shell systems, which is characterized by a large contact area between the photoactive material (TiO₂) and the H₂ evolution catalyst (Ag), photon reflection by the Ag core to capture the maximum number of photons in the titania phase, and short diffusion paths from the TiO₂ phase to the Ag phase in order to separate the exciton efficiently. During a long-term experiment, H₂ was formed at a rate of 1.2·10⁻³ mol/g·h for at least 6 h.

This technique is not only limited to the formation of TiO₂-Ag nanowires, but basically any combination of core-shell materials can be envisaged as long as electrodeposition or a related technique results in the formation of a nanotube for the shell material and a nanowire for the core material. Just as the choice of materials is not limited to what is described in this paper, also the possible applications are not limited to photocatalysis, but they can be extended to other areas, e.g. solar cells and batteries.

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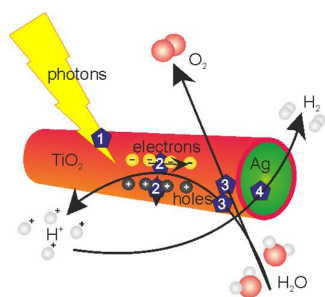
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Coaxial TiO_2 -Ag nanowires were formed that show improved water splitting due to the high contact area between the phases, the large outer surface area, and short electron diffusion paths.