Phosphorus stimulated unidirectional growth of TiO$_2$ nanostructures†

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Previously reported TiO$_2$ nanowire fabrication from Ni catalysts shows a surprising amount of phosphorous (P) contamination incorporated into the seed particle. We proposed this unintentional P-doping of Ni particles aids the mechanism for nanowire growth and occurs by an alternative pathway from the Vapor–Liquid–Solid (VLS) mechanism. To confirm this new mechanism, mixed phase NiP/Ni$_2$P (Ni$_x$P$_y$) and Ni$_2$P nanoparticles were fabricated and the central role of phosphorous in TiO$_2$ nanowire synthesis confirmed. This newly developed P-assisted fabrication method yielded crystalline rutile TiO$_2$ nanowires. In this mechanism solid, quasi-spherical catalyst particles attached to the ends of nanowires and surrounded by a Ni/P liquid shell are responsible for the nanowire growth. The growing end of the nanowire appears to form a “tangent-plane” to the solid catalyst core with the liquid shell wetting and occupying the interstice between the catalyst and the nanowire. In Ni$_x$P$_y$ assisted growth, nanowire diameters occurred as small as 12.3 nm, some of the thinnest yet reported TiO$_2$ nanowires resulting from atmospheric-pressure chemical vapor deposition (APCVD) growth.

Introduction

Recent investigations of titanium dioxide nanowires (NWs) highlight these nanostructures as attractive candidates for gas sensors, charge transport materials in photovoltaic devices, and photocatalysts.\textsuperscript{1–4} Specifically, rutile phase TiO$_2$ nanowires have been studied for excellent charge transport capabilities and high surface area and tested as anode materials to improve lithium ion batteries.\textsuperscript{5} Titanium dioxide (TiO$_2$) has achieved large scale real-world applications due to its low cost, low toxicity and because it is environmentally safe. Currently, typical nanowire syntheses methods include atmospheric-pressure chemical vapor deposition (APCVD).\textsuperscript{6,7} However, optimization of growth conditions, and hence bulk production, is difficult since their growth mechanism is still uncertain. Wagner and Ellis (1964) first recognized the Vapor–Liquid–Solid (VLS) mechanism for nanowire fabrication as being the primary pathway for growth. VLS growth utilizes catalytic metal nanoparticles (NPs) that form low-melting point eutectic alloys with one of the reactants.\textsuperscript{8} The liquid catalyst nanodroplet ultimately becomes the nucleus on which preferential and unidirectional crystal growth occurs. In addition, subsequent research provided evidence of alternative mechanisms such as Vapor–Solid–Solid (VSS)\textsuperscript{4,6} and Solid– Liquid–Solid (SLS)\textsuperscript{9} where nanowire growth occurs well below the expected melting point of the catalyst nanoparticles. Furthermore, recent reports by Kim \textit{et al.} describe another variation of the growth mechanism which they describe as “tangent plane growth”.\textsuperscript{10}

Previously, we reported that an unexpected amount of phosphorus impurity was involved in APCVD growth of TiO$_2$ nanowires. In those experiments, a 2 nm film of Ni was annealed onto a Ti/Si substrate where the Ni film was the catalyst (seed particle) for nanowire growth. Despite the fact that phosphorus was not intentionally added in those experiments, high resolution transmission electron microscopy (HRTEM) revealed post-growth catalyst particles composed of a solid Ni core with an outer shell of Ni$_x$P$_y$. Since highly crystalline rutile TiO$_2$ nanowires were grown from these phosphorus-contaminated Ni catalysts, we hypothesized that phosphorus, and possibly other dopants, could be enhancing the growth mechanism.

In this work, we report the results of using synthesized Ni/P alloys for fabricating TiO$_2$ nanowires by APCVD. Specifically, the goal is to understand the role of phosphorus in the TiO$_2$ nanowire growth mechanism by using mixed phase NiP/Ni$_2$P...
and Ni$_2$P nanoparticles under growth conditions similar to those used in previous experiments. Furthermore, we compare the resulting nanowire growth by the intentional addition of phosphorus to those obtained previously in order to obtain a better understanding of the growth mechanism. In our previous work, we showed unintended P-impurities exhibited a remarkable influence on TiO$_2$ nanowire growth. By doing so we are able to demonstrate comparatively smaller nanowire diameters and control over TiO$_2$ phase (consistently rutile phase). We show P-doping has a positive influence on the control and fabrication of TiO$_2$ nanowires and suggest that this method may be optimized and tuned for a variety photovoltaic and sensing applications.

**Experimental section**

**Synthesis of NiP/Ni$_2$P solid crystalline nanoparticles**

Initially, we synthesized solid crystalline NiP/Ni$_2$P nanoparticles (referred to as Ni$_x$P$_{2-x}$ for brevity) following a similar procedure to that reported by Chiang and Chiang.$^{18}$ This is a one pot synthesis of Ni$_x$P$_{2-x}$ nanoparticles that uses nickel acetate (tetrahydrate, 1 mmol) [Sigma-Aldrich], trioctylphosphine oxide (TOPO) (10 mmol) [Sigma Aldrich], oleylamine (20 mmol) [Sigma Aldrich 70%], trioctylphosphine (TOP) (2 mmol) [Sigma Aldrich 90%] and a growth temperature of 250 °C under inert atmosphere for 1 h. The resulting nanoparticles are precipitated whereupon they were repeatedly (3–5 times) redissolved and separated by centrifugation and further separated the NPs from the solvent. Nanoparticles were dispersed in hexane, approximately 3 drops of colloidal nanoparticle solution were spin coated onto the Si/SiO$_2$ substrate at 3000 rpm for 40 s using a WS-400-6NPP-LITE spin coater. The number of drops of solution was held constant while the NP concentration was varied to observe effects of nanoparticle surface concentrations on nanowire growth.

**CVD nanowire growth**

NW growth was carried out in a horizontal quartz tube furnace, 2.5 cm in diameter and 122 cm long. Approximately 0.2 g of Ti(II) O mesh (99.95%, Sigma-Aldrich) were placed ~5 mm upstream from prepared substrates in the center of a quartz boat. The quartz boat was situated at the center of the quartz tube furnace and heated to 950 °C in air for ~10 min and maintained at 950 °C for another 10 min. APCVD growth was carried out for 2 h under flowing high-purity Ar (99.998%) at 300 sccm. For analysis by transmission electron microscopy (TEM), substrates containing nanowires were sonicated for 60 s in 1 mL EtOH and added dropwise to a 200 mesh carbon coated copper TEM grid.

**Nanoparticle and nanowire analysis**

The resulting nanowires were observed directly on the Si/Ti substrate by scanning electron microscopy (SEM) using a FEI XL40 Sirion FEG microscope with attached Oxford Inca X-ray system for energy dispersive spectroscopy (EDS). The parameters used for obtaining high resolution SEM images were: 5 kV electron energy, working distance of 5 mm and 98 pA electron current. EDS spectra were obtained using an electron energy of 20 kV, a point dwell time of 60 s and a 62 nA electron current. EDS spectra were obtained using an electron energy of 20 kV, a point dwell time of 60 s and a 62 nA electron current. EDS spectra were obtained using an electron energy of 20 kV, a point dwell time of 60 s and a 62 nA electron current.

**Results**

NiP/Ni$_2$P (Ni$_x$P$_{2-x}$) nanoparticles were synthesized according to the procedure outlined in the Experimental section. Three varying concentrations (~1.00 wt%, 2.00 wt% and 7.50 wt%) of Ni$_x$P$_{2-x}$ in hexane solution were spin coated onto three identically prepared Si/SiO$_2$/Ti wafers. APCVD nanowire growth was attempted on all three samples with similar conditions as described in the Experimental section for all three substrates. In addition, it should be noted that CVD growth conditions for new phosphorus-assisted growth are similar to those described in earlier work.$^7$
Atomic force microscopy (AFM) (ESI S1†) and TEM imaging confirmed a solid structure for the resulting Ni$_x$P$_y$ nanoparticles and average diameters ranging from 10–12 nm (Fig. 2). TEM and EDS analysis confirmed the nanoparticles synthesized were composed of Ni and P (ESI S2†). SAED patterns collected over the nanoparticles confirmed a crystalline Ni$_2$P$_{12}$ composition mixed with NiP$_{13}$ (ESI S3†).

Ni$_x$P$_y$ assisted growth

The three concentrations of Ni$_x$P$_y$ in hexane were spin coated onto three Si/SiO$_2$/Ti substrates prepared under identical conditions. Nanowire fabrication was attempted on all three samples using the previously described APCVD method where all samples were heated to 950 °C. Fig. 3 shows HRSEM images of the resulting nanowire growth. Analysis of these images confirms that the lowest concentration of Ni$_x$P$_y$ on the surface of the substrate produces the least amount of nanowire growth and as the concentration of Ni$_x$P$_y$ is increased the number of nanowires grown on the surface increases. Because the same amount of solution was used to spin coat all three substrates for all three solutions, the lowest concentration of Ni$_x$P$_y$ in solution likely results in a lower number of seed particles on the substrate surface. Therefore, fewer seeds are available for growth and relatively fewer nanowires are observed for the lowest Ni$_x$P$_y$ solution concentration (1.00 wt%). More importantly, uniform high quality TiO$_2$ nanowire growth is facilitated, and not inhibited by, the addition of phosphorus to the catalyst particle. Typically, nanowire growth by chemical vapor deposition is thought to occur by the VLS mechanism where the diameter of the nanowires is closely controlled by the diameter of the starting catalytic particle. Since all three samples were spin coated with Ni$_x$P$_y$ nanoparticles from the same batch but with differing concentrations, one might expect to see similar diameters of nanowire growth for all three substrates. However, as shown in Fig. 3, the average diameter of the TiO$_2$ nanowires increases from 12–20 nm (Fig. 1 and 3b) for 2.00 wt% Ni$_x$P$_y$ to 30–80 nm for 7.50 wt% Ni$_x$P$_y$. Since the increasing wt% of Ni$_x$P$_y$ translates into nanoparticles resting closer together on the substrate surface, the increasing nanowire diameter may result from increased NP aggregation with increasing nanoparticle surface concentration, i.e. NPs may be more likely to aggregate to form particles with larger diameters (greater than 10–12 nm) during the initial heating step before nanowire growth occurs. Thus, subsequent nanowire growth from large catalyst particles

![Fig. 1](image1)

**Fig. 1** HRSEM imaging of TiO$_2$ nanowire grown from Ni$_x$P$_y$ catalyst confirms 12.3 ± 0.4 nm diameter sizes. The scale bar represents 100 nm.

![Fig. 2](image2)

**Fig. 2** BF TEM imaging of Ni$_x$P$_y$ nanoparticles confirming average 10–12 nm diameter sizes (<20 nm). The scale bar represents 20 nm.

![Fig. 3](image3)

**Fig. 3** HRSEM imaging shows (a) little to no TiO$_2$ nanowire growth with ~1.00 wt% Ni$_x$P$_y$ in hexane solution (b) NWs grown from ~2.00 wt% Ni$_x$P$_y$ in hexane solution occur 12–30 nm in diameter and 2–3 μm in length and (c) NWs grown from ~7.50 wt% Ni$_x$P$_y$ in hexane solution ranging from 30–80 nm in diameter and 6–10 μm in length. All Ni$_x$P$_y$ solutions were dispersed onto a Si/SiO$_2$/Ti substrate.
leads to larger nanowire diameters. Energy dispersive X-ray spectroscopy (EDS) on resulting nanowires revealed the presence of Ni and P in the catalyst particle and Ti and O in the nanowires. EDS spectra showed a semi-quantitative Ni : P atomic% ratio of nearly 3 : 1 (Fig. 4) – a similar ratio to the crystalline Ni₃P catalytic particles produced by CVD growth on the ends of TiO₂ nanowires that prompted the present investigation. Measured g vectors from TEM diffraction patterns obtained for nanowires matched those of rutile TiO₂ (Fig. 5). This result is consistent with previous phosphorus contaminated experiments where the TiO₂ nanowires produced were also found to be rutile phase.

It should be noted that while Ni₃P batches synthesized by the same procedure (i.e. similar to that of Chiang and Chiang) usually produced similar nanowire growth on Si/SiO₂/Ti substrates, other batches yielded amorphous head particles with higher occurrence of kinks and bends in the resulting nanowires. To better understand this, we performed TEM imaging on nanoparticles that led to kinked nanowire growth. Images revealed that the nanoparticles produced formed a mixture of both hollow and solid crystalline phases (ESI S4†). Since our previous results suggested that a core–shell structure facilitates the nanowire growth through the transport of materials to the surface of the substrate, we conclude here that the hollow nanoparticles were inhibiting the nanowire growth and causing the kinked structures observed. We determined that the procedure outlined in the Experimental methods section, as well as the nanoparticle synthesis outlined by Chiang and Chiang, does not consistently produce solid crystalline NPs but occasionally yields a mixture of both solid crystalline as well as hollow crystalline NPs. This result is supported by subsequent research by Wang et al. describing similar mixtures of hollow and solid nanoparticles when following the Chiang and Chiang procedure. Wang et al. addresses the problem by conducting a series of experiments that lead to more controlled methods of producing exclusively hollow or solid nanoparticles. They report the fabrication of crystalline solid Ni₁₂P₅/Ni₂P nanoparticles exclusively (with no hollow NPs) by adjusting the reactant P : Ni ratios as well as the temperature ramp sequence and solvent. Therefore, following the procedure outlined by Wang et al., solid crystalline Ni₁₂P₅/Ni₂P NPs were fabricated for use as catalysts to further investigate phosphorus-assisted TiO₂ nanowire growth. Moreover, we sought to observe the effect of changing the concentration of phosphorus in the nanoparticle (NiP/NiP₂ versus Ni₁₂P₅/Ni₃P), on nanowire growth. This was undertaken using identical substrates and growth conditions (950 °C) as in previous experiments with Ni₃P.

**Analysis of Ni₁₂P₅/Ni₂P**

Upon synthesis of Ni₂P solid crystalline nanoparticles, four dilutions of Ni₂P in hexane were prepared for spin-coating onto Si/SiO₂/Ti substrates, 14.26 wt%, 7.13 wt%, 3.56 wt% and 1.78 wt%. A dilute solution of Ni₂P dispersed in hexane was also added dropwise onto a 200 mesh carbon coated copper grid for later TEM analysis.

EDS and SAED were used to confirm the crystal structure of the synthesized seed particles following the work of Wang et al. EDS spectra revealed that nanoparticles are composed of Ni and P (ESI S5†). Cu and carbon peaks observed on EDS spectra can be attributed to background from the TEM grid. Measured g vectors from synthesized nanoparticle SAED patterns appear to fall mainly on values for Ni₃P, suggesting the Ni₁₂P₅–Ni₂P mixture is primarily Ni₁₂P (ESI S6†). This result is further supported by the Ni–P binary phase diagram which indicates a relatively narrow compositional range for pure Ni₁₂P₅ directly adjacent to the formation of Ni₃P (Fig. 9). Therefore we refer to the Ni₁₂P₅–Ni₂P mixture as Ni₃P for clarity.

**Ni₃P assisted growth**

Nanowire fabrication was attempted for all four dilutions of NPs (as described above) under identical CVD conditions as
described in the Experimental methods section. Similar to \( \text{Ni}_x \text{P}_y \) assisted growth, the greatest occurrence of nanowire growth resulted from the highest concentration of particles (14.26 wt%) and no nanowires were observed for the lowest (1.78 wt%) concentration of \( \text{Ni}_2\text{P} \). However, nanowire diameters grown from \( \text{Ni}_2\text{P} \) seeds exhibited a wider range compared to \( \text{Ni}_x \text{P}_y \) growth, with diameters from 26–161 \( \pm \) 5 on the 14.26 wt% substrate. Nevertheless, average diameters are similar to those observed on the \( \sim 7.50 \) wt% \( \text{Ni}_2\text{P} \) substrate, ranging from \( \sim 50–80 \pm 2 \) nm (Fig. 6). The diameters of nanowires and their corresponding attached catalyst particle were obtained from SEM analysis. We note in one example (ESI S7†) that two different catalyst particle diameters (197 nm and 470 nm) yielded closely similar nanowire diameters (85.4 nm and 88.9 nm respectively).

High resolution STEM imaging and EDS analysis revealed non-homogeneous compositions in some nanowires produced from \( \text{Ni}_2\text{P} \) batch solutions. Shown in Fig. 6 and 7, most nanowires were confirmed to be the \( \text{TiO}_2 \) rutile phase. Semi-quantitative data from EDS line scans suggest a \( \text{TiO}_2 \) NW composition and a \( \text{Ni}–\text{P} \) spherical catalyst (Fig. 8a). Moving from left to right across the \( \sim 20 \) nm line shown in Fig. 8b, EDS spectra reveal a relatively high abundance of \( \text{Ti} \) and \( \text{O} \) (68.48% and 31.52% respectively) while the signal from \( \text{Ni} \) and \( \text{P} \) are nearly zero. Near the center of the particle and away from the nanowire, the composition shifts along the line to mostly \( \text{Ni} \)-rich with \( \text{Ni}, \text{P}, \text{Ti} \) and \( \text{O} \) signals amounting to 83.66%, 2.28%, 0.98% and 3.45% respectively. \( \text{Si} \) was also observed (9.62% abundance near the center of the particle) in the amorphous shell of this catalyst particle. Although abundant in the shell observed in Fig. 8, \( \text{Si} \) was observed only in some particles (roughly half of those analyzed by EDS) while in others it was completely absent. The reduced signals from \( \text{Ti} \) and \( \text{O} \) at the end of the line nearest to the center of the catalyst particle are comparable to those observed in ref. 8 and suggest that an oxygen-rich material forms a shell around the nanoparticle. The spectrum collected over the interface region between nanowire and nanoparticle (from 11–20 nm) shows a steep increase in oxygen content at the interface where the ratio of oxygen to titanium approaches to nearly 2 : 1. Complementary HRTEM imaging over this oxygen-rich region indicates that the material is amorphous and extends around the entire exterior surface of the catalyst. The \( \text{Ti} \) signal also shows a similar steep increase near the same region on the line (4–20 nm). This amorphous \( \text{Ti}–\text{O}–\text{P} \) containing shell is consistent with previous line scans over phosphorus-contaminated particles.8 This core–shell structure is also consistent with recent studies on \( \text{Ni}–\text{P} \) nanoparticles which form core–shell structures depending on the relative kinetics of each migrating species (\( \text{Ni} \) and \( \text{P} \)).13

![Image](https://example.com/image1.png)

**Fig. 6** (a) HRSEM image reveals that a greater quantity of nanowire growth occurs with higher number of \( \text{Ni}_2\text{P} \) particles dispersed onto the surface is increased. (b) HRSEM image showing nanowire diameters are relatively large.

![Image](https://example.com/image2.png)

**Fig. 7** (a) STEM image and (b) corresponding EDS spectra show a solid nickel seed particle with an amorphous layer of \( \text{P} \) and \( \text{O} \) on the exterior surface of the resulting nanoparticle after \( \text{Ni}_2\text{P} \) assisted nanowire growth. EDS 1 represents analysis on the center of the nanoparticle while EDS 2 was taken over the outermost edge of the particle. The \( \text{Si} \) peak is presumed to be an artifact of TEM analysis.
Measured \( g \) vectors from SAED patterns obtained during TEM analysis of post-growth nanowires suggest that the seed particles are solid Ni (ESI S8†). Although no phosphorus was observed by SAED, EDS line scan spectra obtained on the same Ni seeds show an increasing amount of phosphorus in the outermost layer of the nanoparticle as compared with the

Fig. 8  (a) STEM image and corresponding EDS spectra showing Ti–O nanowire composition grown from Ni\(_2\)P seed particle. (b)–(e) EDS line scan across nanowire–nanoparticle interface. Spectra indicate a TiO\(_x\) nanowire and solid Ni NP with an amorphous shell of O-rich material at the NW–NP interface.

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Fig. 9  The binary phase diagram for Ni–P (adapted from Okamoto15) shows a eutectic temperature of 891 °C for a Ni–P mixture that is much lower than the Ni melting temperature at 1455 °C.
interior (ESI S9†). Thus, EDS analysis in combination with SAED results suggests that the post-nanowire growth catalysts are composed of a solid Ni core with an outer phosphorus/nickel shell, similar to that reported in ref. 8.

In some Ni2P derived nanowires, EDS indicated catalyst particles were solid crystalline Ni with no observable phosphorus content and an outer shell of amorphous Ni–O. This result is similar to the amorphous Ni–Ti–O shell seen in ref. 8. SEM and STEM imaging show spheroidal structures protruding from the surface of other Ni2P derived nanowires. EDS spectra collected on these structures indicate that these spheroids are the result of phosphorus erupting outward from the particle and solidifying on the outer surface of the resulting solid Ni catalyst (Fig. 7). In some rare cases, we also observe nanoparticles at the ends of NWs with a hexagonal structure and EDS-calculated Ni : P ratios of 2 : 1. This ratio and hexagonal shape indicates a typical hcp structure for Ni2P suggesting that some Ni2P seed particles may remain unchanged during the nanowire growth process.

The mechanism indicated by nanowire and seed particle structures appears to be the same in both NiP and Ni2P assisted growth with the resulting seed particles remaining nearly spherical for unidirectional nanowire growth. This ball-and-stick structure differs from the typical VLS structure where catalyst particles are quasi-hemispherical with a greater surface area exposed to the direction of nanowire growth. A fully spherical structure is exhibited by our P-assisted nanowire growth and is strikingly similar to our previously proposed "tangent-plane" mechanism in P-contaminated experiments. Furthermore, the amorphous shell of material observed at the interface between the nanowire and catalyst, as seen by TEM and STEM imaging (Fig. 7 and 8), suggests a solid nickel core surrounded by a shell of Ni–P. The solid Ni core structure precludes the collapse of the catalyst particle into a quasi-hemispherical shape typical of VLS, while the liquid shell provides the active (and efficient) reaction medium. As expected based on the data given in the Ni–P phase diagram, the Ni–P mixture has a lower melting point than that of pure Ni (Fig. 9).

Discussion

In light of previously discovered phosphorus impurities in our seed particles, we intentionally incorporated phosphorus into the catalyst particle in order to experimental test our previously suggested hypothesis that phosphorus may be important in facilitating TiO2 nanowire growth. New experiments include nickel–phosphide catalysts, mixed phase NiP/Ni2P (NiP-Ni2P) and NiP nanoparticles. Nanowire growth appeared similar in both NiP-Ni2P and Ni2P growth except in cases where hollow NiP-Ni2P nanoparticles occurred. In both cases, we observe the formation of rutile phase TiO2 nanowires and quasi-spherical seed particles attached to the ends of nanowires.

Seeding TiO2 with NiP nanoparticles resulted in highly crystalline rutile phase nanowires. By increasing the number of particles dispersed onto the substrate surface we saw a general trend of increasing nanowire growth both with respect to diameter and length. In the case where ~2.0 wt% NiP in hexane was spin-coated onto the substrate, TiO2 nanowires as narrow as ~12 nm across were observed. To the best of our knowledge, this is the thinnest diameter yet reported for TiO2 nanowires grown by APCVD. Where NP batches were later found to contain a mixture of hollow and solid particles, poor nanowire growth resulted with amorphously shaped seed particles and non-unidirectional nanowire growth.

In order to control the starting crystallinity of catalyst particles and observe any changes in nanowire growth with changes in the Ni–P phase (NiP versus Ni2P), we attempted similar CVD nanowire growth using NiP nanoparticles as catalysts. In Ni2P assisted growth, we again observed higher density nanowire growth in the case where the highest wt% of Ni–P NPs in hexane solution was used. As seen in NiP and Ni-film experiments, the resulting TiO2 nanowire growth appears unidirectional with quasi-spherical seed particles surrounded by an amorphous outer shell. Overall, the range of nanowire diameters appears much larger in NiP compared to Ni2P assisted growth. However, NiP assisted growth yielded more consistently uniform nanowire growth with fewer cases of amorphous shapes or kinks in wires as observed in NiP growth experiments.

In addition to Ni–P nanoparticle experiments, we also performed experiments without the intentional addition of phosphorus. Using different quartz tubes and high purity quartz boats, we performed effectively phosphorus-free TiO2 nanowire growth under the same conditions (950 °C). In this experiment we used identical substrates (Si/SiO2 and 50 nm Ti layer) and titanium source (titanium mesh) as for those described for the Ni–P nanoparticle experiments. The Ni–P nanoparticles were replaced with a 2 nm Ni layer (as used in previous non-phosphorus doped experiments). As described previously, TiO2 nanowire growth was suppressed and no well-formed head particles or nanowires were observed (data not shown). This observation, in combination with data from the phosphorus-doped experiments described here, suggests phosphorus facilitates highly crystalline TiO2 nanowire growth.

At present, Vapor–Liquid (VLS) growth is the most common method for producing large quantities of high quality nanomaterials. However, many attempts have recently been made to challenge the growth mechanism originally proposed by Wagner and Ellis in 1964.15,16 In our previous work, we presented evidence for tangent-plane growth where the seed catalyst particle remains sphericaly intact and an amorphous shell of Ti–Ni–O material surrounds the exterior of the nanoparticle as well as the interface region between nanowire and seed. We also clearly presented a case for trace element, P, appearing on the surface of a designed Ni catalyst particle. We showed that high quality TiO2 nanowires could grow on solid (phosphorus-doped) Ni at 950 °C, a temperature far below the typical melting point of 1455 °C for bulk Ni (or of the Ni nanoparticle).

In the experiments we report here, we show consistently similar rutile TiO2 nanowire growth at the same temperature (950 °C) using phosphorus-doped nickel nanoparticles. EDS line scans through the interface area between the nanoparticle and nanowire indicate an amorphous, oxygen-rich material which forms a shell on the outer surface of a quasi-spherical NP.
HRTEM images and SAED patterns collected on the catalyst particle after nanowire growth reveal a solid Ni phase with an amorphous shell on the outer surface. The binary phase diagram (Fig. 9) indicates a eutectic temperature of the Ni–P that is 891 °C, well below our working temperature of 950 °C. The amorphous shell and quasi-spherical shape observed in our Ni–P seeds suggest that the mechanism for growth includes the formation of a liquid Ni–P shell (in agreement with the Ni–P phase diagram) which surrounds a solid Ni core. These results are consistent with the tangent-plane mechanism proposed previously in experiments where unintentional addition of phosphorus was hypothesized to aid in TiO2 nanowire synthesis. In this mechanism, the liquid shell and solid core together act as the catalyst for adsorption, diffusion, and reaction of Ti and O, resulting in TiO2 nucleation and crystallization from the liquid phase near the surface of the solid nickel nanoparticle. Formation of a typical VLS-growth quasi-hemispherical shape is inhibited by the solid Ni core while the liquid shell provides the active reaction medium. Furthermore, unlike typical VLS growth where the diameter of the nanowire is limited by the diameter of the starting catalyst, we observed nanowires of similar size grown from different diameters of Ni–P nanoparticles.

**Conclusion**

This research confirms in more detail the facilitation of TiO2 nanowire growth by means of phosphorus catalysis. Furthermore, through repeated experimentation, we show a novel phosphorus doping method for fabricating rutile TiO2 nanowires. By incorporating phosphorus into the catalytic nanoparticle, APCVD nanowire growth is accomplished by an alternative process to the vapor–liquid–solid mechanism. The observed quasi-spherical nanoparticle shape appears similar to our previously proposed “tangent-plane” mechanism. When phosphorus is removed, TiO2 nanowire growth is suppressed and no quasi-spherical seed particles are observed. This novel method of P-assisted TiO2 nanowire growth consistently results in rutile phase TiO2 nanowires. As noted earlier, rutile TiO2 is regarded as promising for lithium insertion/extraction materials due to its low production cost and high capacity for lithium intercalation. In addition, we report significantly smaller nanowire diameters (down to 12 nm) produced from the P-assisted method compared to previous non-P-assisted experiments. Such small diameters would be useful particularly in gas sensing devices. For example, Moskovits et al. (2010) recently developed an “electronic nose” device comprised of decorated metal-oxide nanowires providing improved sensitivity toward carbon monoxide, hydrogen and ethylene over bulk materials due to the higher surface area of the catalyst. Compared to previously reported nanowire synthesis, the P-assisted method similarly produces rutile phase TiO2 nanowires but smaller diameters depending on Ni–P seed particle used. Except where hollow nanoparticles were present, TiO2 nanowire fabrication from Ni–P seeds was consistent for both amounts of P tested. Therefore, these results suggest trace elements which may be present in VLS nanowire growth should not be ignored and can have a positive influence on the fabrication method. By taking impurities into account, this work represents a novel example of nanoparticle assisted nanowire growth which is inspired by P impurities. We postulate that a similar P-assisted method could be used for other metal oxide nanowire fabrication in future research.

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