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A Template and Catalyst-Free Metal-Etching-Oxidation Method to Synthesize Aligned Oxide Nanowire Arrays: NiO as an Example

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Deterministic synthesis of one-dimensional (1D) crystalline nanowires (NWs) with controlled morphologies and properties is currently a major objective in nanosciences and nanotechnologies.¹ Various synthesis routes have been developed, and their applications to NW synthesis have been intensively pursued. Compared with the solution-based methods, vapor transport growth has the advantage of being more environmentally friendly and often can give NWs with high aspect ratios, which is critical for many key applications. Vapor–liquid–solid (VLS) growth is one of the most popular synthesis methods, where liquid-state catalyst nanoparticles absorb the precursor vapor and NW growth starts when supersaturation is reached in the nanoparticles.^{2–5} Recently, a variant that uses solid-state catalysts at lower temperatures, termed as the “vapor–solid–solid” (VSS) growth, has also been devised to grow NWs.⁶ In another distinct approach without the use of catalyst, NW synthesis has been accomplished through the vapor–solid (VS) route.^{7,8} Among all the techniques to produce NWs, the hot plate method is probably the most convenient and cost-effective, where heating metal foils in either atmosphere or controlled environment was found to give 1D nanomaterials.⁹ Complementary to the above vapor-based routes, template-assisted methods have been employed to produce NWs with uniform size; for example, liquid or gas phase precursors were filled into the pores of anodic aluminum ox-

ABSTRACT Although NiO is one of the canonical functional binary oxides, there has been no report so far on the effective fabrication of aligned single crystalline NiO nanowire arrays. Here we report a novel vapor-based metal-etching-oxidation method to synthesize high-quality NiO nanowire arrays with good vertical alignment and morphology control. In this method, Ni foils are used as both the substrates and the nickel source; NiCl₂ powder serves as the additional Ni source and provides Cl₂ to initiate mild etching. No template is deliberately employed; instead a nanograined NiO scale formed on the NiO foil guides the vapor infiltration and assists the self-assembled growth of NiO nanowires *via* a novel process comprising simultaneous Cl₂ etching and gentle oxidation. Furthermore, using CoO nanowires and Co-doped NiO as examples, we show that this general method can be employed to produce nanowires of other oxides as well as the doped counterparts.

KEYWORDS: nickel oxide · nanowire · metal-etching-oxidation method · catalyst · self-assembly

ide (AAO) membranes to render either NWs or nanotubes.¹⁰

However, it is well recognized that for some materials the currently available synthesis methods are not very effective to enable the anisotropic growth of 1D NWs. This is in particular true for materials with cubic crystal symmetries which lack the structural anisotropy critically needed for the growth of single crystalline NWs. Synthesis of these oxide NWs requires developing novel routes beyond the conventional ones. Nickel oxide (NiO) is one of such notable materials. NiO is an important member in the family of binary oxides with a cubic rock salt structure and a lattice parameter of 0.4195 nm. Several decades ago, NiO played a very critical role in the development of the modern solid state physics by revealing the limitation of the free electron model and the critical role of strong electron correlation.¹¹ Today, NiO is an important p-type semiconductor

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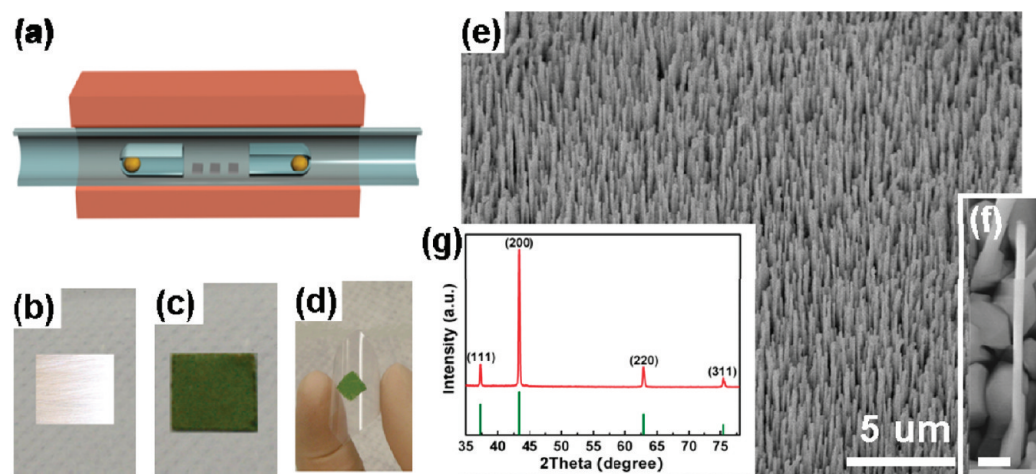


Figure 1. (a) Schematic of the experimental setup. This figure is not drawn to scale. Photographs of (b) polished Ni foil, (c) Ni foil after NiO NW growth, and (d) NiO NW arrays after being transferred from the Ni foil to a flexible plastic substrate. (e) A typical SEM image of the NiO NW arrays. (f) SEM image of an individual NiO NW. The scale bar is 200 nm. (g) XRD spectrum of the NiO NW arrays. The NiO powder diffraction data (JCPDS Card No. 78-0423) is also shown at the bottom.

material¹² and has been extensively investigated because of its myriad applications in catalysts,^{13,14} gas sensors,¹⁵ battery materials,¹⁶ electrochromic coatings,¹⁷ and so on. It is also a canonical antiferromagnetic material with a Neel temperature of 523 K,¹⁸ and often used in magnetic multilayer devices.^{19,20} Most recently, NiO showed excellent resistive switching behaviors and may find potential uses in nonvolatile memories.^{21,22} Up to now, a variety of methods have been attempted to achieve nanostructured NiO, but all of them suffer from drawbacks. For example, solution-based chemical routes²³ and sol-gel methods²⁴ were shown to produce NiO NWs in large quantity, but the NWs have no alignment. Electrochemical deposition into anodic alumina membranes^{25,26} and oxidation of NiS precursor in molten salt²⁷ were also employed to prepare NiO NWs. However, these two methods lead to only polycrystalline NWs. To the best of our knowledge, template-free synthesis of aligned single-crystal NiO NW arrays has not been reported.

In this study, we discovered a novel template- and catalyst-free method to synthesize high quality NiO NW arrays. This facile method involves controlled anisotropic etching of a nickel (Ni) foil through a NiO layer with low-pressure Cl₂ vapor, which is accompanied by continuous oxidation. It is the first time, to our knowledge, that single-crystal NiO NW arrays were synthesized with a template-free method. Furthermore, we show that this method is a general approach to produce metal oxide NW arrays and can be used to dope the NWs by adjusting the vapor source.

RESULTS AND DISCUSSION

The synthesis of the NiO NW arrays was carried out in a home-built vapor transport system comprising a horizontal tube furnace (Lindberg/Blue mini-mite) and a gas delivery and pumping system as described previously.^{28–30} The schematic of the experimental

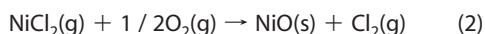
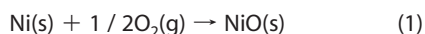
setup is shown in Figure 1a. Ni foils were cut into small pieces, cleaned, mechanically polished, and electrochemically polished. Finally, the Ni foils were annealed to decrease the internal stress and to further improve the crystalline quality (see Supporting Information Figure S1). These presynthesis treatments are critical to achieve (100) textured Ni foils with very flat surfaces. The treated Ni foils (Figure 1b) were then put in the center of the furnace between the small tubes containing the NiCl₂ powder. We need to point out here that the Ni foils serve as both the substrate and the nickel source. Nickel chloride (NiCl₂) powder was put into two small one-end-open quartz tubes. After the growth process, a layer of green products was grown on the surface of the Ni foils. Figure 1c shows the optical image of the Ni foil after growth. The green color indicates the presence of NiO on the Ni foil. We found that NiO NWs are not directly grown on the Ni foil, but imbedded under a NiO film (the growth mechanism will be discussed in details later on). Thus to reveal the NiO NW arrays, we need to bond a supporting substrate to the sample using either silver paste or double-sided tape, then the NiO NWs can be conveniently transferred to the substrate by lift-off. Since the growth and transfer processes are separated, a supplementary advantage of this method is that there is no limitation on selecting the final supporting substrate. Figure 1d shows the optical image of a sample of NiO NW arrays being transferred to a flexible plastic substrate. Figure 1e shows a typical SEM image of the NiO NW arrays. The NiO NWs are several micrometers in length and about one hundred nanometers in diameter. They are quite uniform and appear well aligned. Figure 1f is the SEM image of an individual NiO NW showing a smooth surface. The chemical composition of the NWs was characterized by XRD, and as shown in Figure 1g, only NiO related peaks were detected. The relative intensity of the peaks differs from the powder diffraction data (JCPDS Card No.

F1

145 78-0423), and the dominant (200) peak indicates a preferred (200) orientation. The absence of NiCl₂ or Ni related peaks excludes the existence of any impurity phase above the detection level.

149 The crystalline structure of the NiO NWs was further characterized by TEM. Figure 2a shows a low-resolution TEM image of a NiO NW. The diameter of this particular NW is about 130 nm. The selected area electron diffraction (SAED) pattern in Figure 2b can be indexed as the cubic structure with no detectable secondary phase. The HRTEM image (Figure 2c) was recorded along the [011] zone axis and it shows clear lattice fringes. The distance between the lattice planes along the NW growth direction is 0.208 nm, which is consistent with the lattice constant of the NiO unit cell. Both the SAED pattern and the HRTEM image suggest that the NW is single crystalline and the growth direction is along [200]. Among a dozen NiO NWs we examined using HRTEM, we also observed a few growing along [220] (Supporting Information, Figure S2). Nevertheless, we found that the majority of the NiO NWs are along the [200] direction, which is in agreement with the XRD data.

167 We now discuss the tentative formation mechanism of the NWs which grow underneath the NiO overlayer. The schematic in Figure 3a–d illustrates the growth process of the NiO NW arrays. At the growth temperature, three major reactions are involved:³¹



172 The high temperature oxidation of Ni and the formation of NiO scale have been studied for many years.³² The oxidation reaction in eq 1 accelerates as the furnace temperature increases, which is accompanied by a weight gain of the Ni foil (Supporting Information, Figure S3). The formation of NiO is also confirmed by the XRD data shown in Figure S4; the NiO [200] peak at 43.3° starts to appear and becomes stronger as the temperature increases. When the furnace temperature reaches 600 °C, NiCl₂ powder starts to evaporate into the gas form, which is evidenced by the rapid weight loss of the NiCl₂ powder above this temperature (Figure S3). The reaction 2 suggests that NiCl₂ vapor reacts with O₂ in the mixture gas to produce Cl₂ gas. The existence of Cl₂ gas can significantly affect the kinetics of the oxidation process.³¹ It is important to note that there are many closely spaced microchannels across the columnar NiO scale, and the gaseous species can penetrate the scale along the fine grain boundaries.³² Subsequently, according to reaction 3, the Cl₂ gas diffuses through the NiO layer and etches the Ni substrate underneath it. This process helps to gently sculpt the continuous Ni foil into wire-like morphology as illustrated in Figure 3c. According to the reaction 3, during

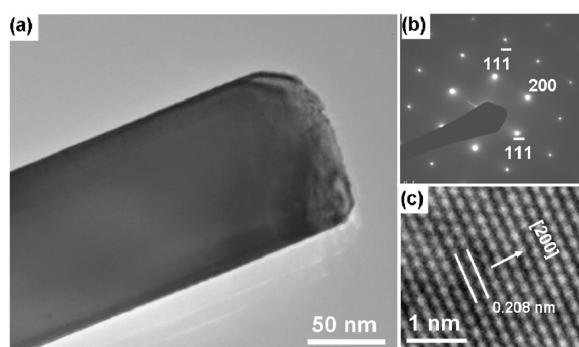


Figure 2. (a) Low resolution TEM image of a NiO NW. (b) SAED pattern and (c) HRTEM image taken on the same NW. The inter-plane distance of 0.208 nm is consistent with the lattice constant of NiO.

etching, additional NiCl₂ is produced in the gas form, which is fed back to reaction 2. Hence, NiCl₂ and Cl₂ are conserved, and the overall result is the oxidation of Ni. However, this balance is not strictly maintained in the experiments; NiCl₂ and Cl₂ in the gas form continuously escape from the samples and flow away with the carrying gas, which eventually terminates the reactions.

We should note that although no template is deliberately employed in the synthesis, the nanograined NiO scale serves as a mask guiding the infiltration of Cl₂ and O₂ gases. As illustrated in the schematics in Figure 3, concurrent to the etching process is the oxidation of Ni into the NiO NWs, which is sustained by the carrying gas. During this process, the electric field of more than 10⁶ V cm⁻¹ developed between Ni and NiO,³³ along with the giant mechanical stress at the interface,³⁴ may assist the self-organized formation of the NiO NW arrays. It has been reported that in the oxidation of highly textured Ni tapes the crystallographic orientation of the NiO grains is closely correlated with that of the Ni grains underneath.³⁵ Therefore, the annealing process prior to the NW growth helps to produce highly textured Ni foils and warrants the crystalline alignment of the produced NiO NW arrays. Furthermore, the NiO scale with fine channels limits the reaction rate and helps to protect the Ni foil from being etched excessively. If the oxidation proceeds too fast, the growth

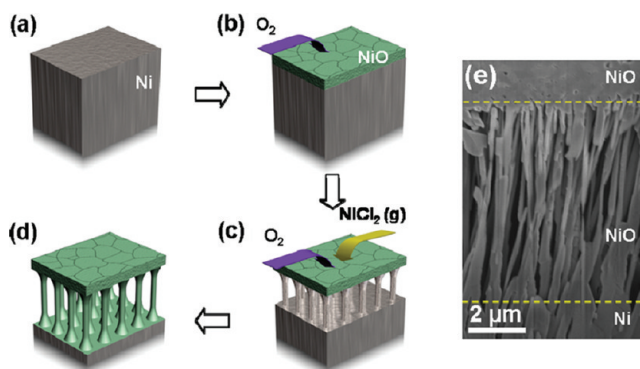


Figure 3. Schematic of the metal-etching-oxidation (MEO) mechanism proposed for the growth of NiO NW arrays. (e) Cross-sectional SEM image of the NiO NW arrays prepared by focused ion beam (FIB) milling.

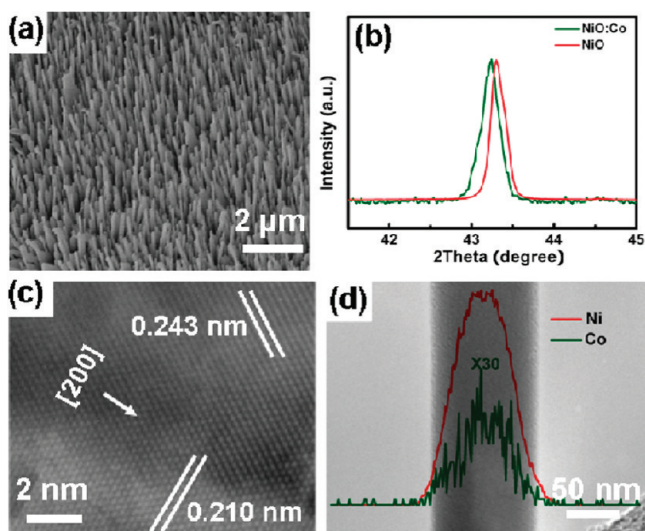


Figure 4. (a) SEM image of Co-doped NiO NW arrays obtained by using CoCl_2 as the source powder. (b) The normalized (200) peak in XRD pattern of the NiO and Co-doped NiO NW arrays. (c) HRTEM image showing the growth direction along [200]. (d) EDS mappings of Ni and Co across an individual NW.

stress may cause the oxide to separate from the metal due to the insufficient oxide plasticity.³⁶ This is analogous to the recently reported spontaneous growth of NiSi_x NWs where polygrained NiO_x overlayers act as a diffusion barrier to kinetically and spatially control the Ni diffusion during the chemical vapor growth.^{37,38} It was shown that the presence of the oxide overlayers on Ni seed layers promote the reproducible one-dimensional growth of NiSi_x single crystals. Similarly, oxide overlayers play critical roles in our experiments to control the vapor supersaturation and to give the reliable NiO NW growth.

We suggest to term this approach of producing aligned NiO NW arrays the “metal-etching-oxidation”

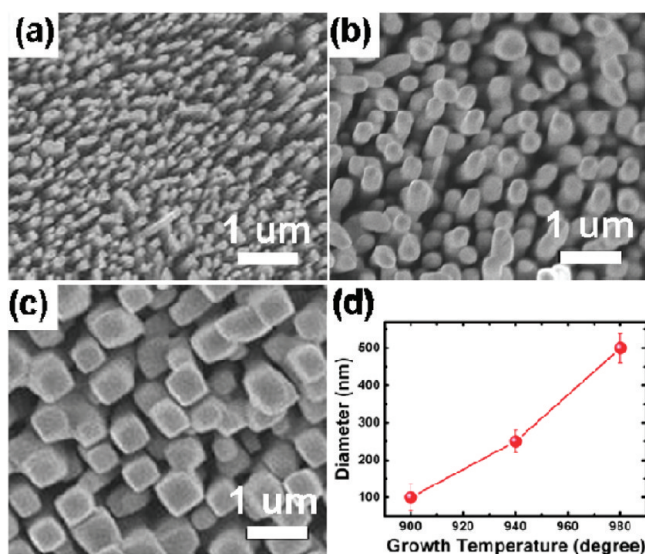
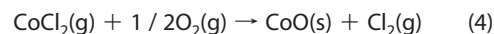


Figure 5. Effect of the growth temperature on the morphology of NiO NW arrays. Typical SEM images taken on samples grown at substrate temperatures of (a) 900, (b) 940, and (c) 980 °C. (d) Dependence of the NW diameter on the growth temperature.

(MEO) process which captures the essence of the formation mechanism. Furthermore, to achieve direct evidence to confirm the above scenario, we looked at the cross-sectional SEM image of the NW arrays exposed by using focused ion beam (FIB) milling. As shown in Figure 3e, the NiO NWs are imbedded between the NiO scale layer and the Ni foil. Even though significant breaking, melting, and fusing of the NiO NWs occurred because of the large ion milling current, the observed particular columnar structure of the NiO NW arrays supports the proposed growth mechanism. However, we should be cautious regarding the details of the exact growth mechanism owing to the lack of direct experimental evidence and real-time observation. We noted that anisotropic silicon NW arrays have been recently fabricated *via* a metal-assisted chemical etching method.^{39,40} Although both the reported method and ours rely on anisotropic etching, our approach does not involve the use of any catalyst and appears to be more suitable to prepare NWs of metal oxides.

Since etching and oxidation are the two key elements in the MEO process, chlorides other than NiCl_2 should also be capable to initiate the growth. To test this hypothesis, we chose CoCl_2 as the source powder in the synthesis. At high temperatures, Cl_2 gas is produced as the following:



Following a similar growth mechanism as in the NiCl_2 case, NW arrays were obtained. As shown in Figure 4a, the NW arrays are uniform, being 2 μm in length and about 100 nm in diameter. In addition, this change of source allows us to introduce Co as a doping element into the NiO NWs. The Co doping process occurs simultaneously with the etching and oxidation process, and the high synthesis temperature assists the efficient diffusion of Co in the NiO matrix. As shown in the XRD patterns in Figure 4b, the (200) peak of the Co-doped NiO NWs shifts toward a lower angle compared with that of the undoped counterpart, which is consistent with the fact that the radius of Co ions (0.0838 nm) is slightly larger than that of the Ni ions (0.0830 nm). The HRTEM image in Figure 4c shows that the single crystal NWs grow along the [200] direction, consistent with the results of the NiO NWs. The EDS data suggest a Co doping level of *ca.* 1%. In addition, in the EDS line scans across an individual Co-doped NiO NW (Figure 4d), the Co profile appears to follow the Ni profile very well, indicating a uniform distribution of the Co element.⁴¹

We further demonstrate that one is able to tune the morphology of the NiO NWs through adjusting the growth temperature and the oxygen content in the carrying gas. As shown in the SEM images in Figure 5a–c, the diameter of the NiO NWs increased from 100 to 500 nm as the growth temperature changed from 900 to 980 °C. This clear trend shown in Figure 5d could be

291 due to two reasons. One is that a higher growth tem-
 292 perature always results into a NiO scale with larger
 293 grains.⁴² Since the NW morphology is mainly defined
 294 by the Cl₂ gas etching through the grain boundaries,
 295 larger grains eventually lead to thicker wires. The other
 296 is that a higher temperature significantly promotes the
 297 Ni oxidation, thus circumvents the etching, which may
 298 also result in NWs with larger diameters. At 980 °C, the
 299 cross section of the NiO NWs appears to be square-like,
 300 indicating their single crystal nature.

301 We found that the growth temperature has little ef-
 302 fect on the length of the NiO NWs. Instead, the NW length
 303 can be well controlled by adjusting the gas content. As
 304 shown in the SEM images in Figure 6a–d, shorter NWs
 305 could be obtained if a carrying gas with lower oxygen
 306 content was used. The length and diameter of NWs were
 307 measured for these samples, and the statistical results are
 308 shown in Figure 6e and f. The NW length increased from
 309 1 to 10 μm with the increasing oxygen content from 0.5%
 310 to 15%, while the diameter changed only slightly. This ef-
 311 fective length control originates from the fact that a
 312 higher oxygen concentration promotes the oxidation
 313 and protects the NW core from the excessive etching.
 314 Consequently, the Cl₂ etching gas goes deeper into the
 315 Ni foil, forming longer NiO NWs. Since the diameter and
 316 the length of the NiO NWs can be independently con-
 317 trolled by the growth temperature and the carrying gas
 318 content, the synthesis route herein gives lots of flexibility
 319 in the morphology control.

320 Recently NiO thin films have been reported for their
 321 potential applications in the next generation resistive
 322 RAM (ReRAM) devices, which may serve as alternatives
 323 to the current nonvolatile flash memory technology.^{43,44}
 324 Although the switching model involving the formation
 325 and rupture of conducting filaments is widely used to ex-
 326 plain the experimental observations, the complex switch-
 327 ing mechanisms in NiO warrants further investigations.
 328 Very recently, reversible resistive switching behaviors
 329 have been demonstrated in NiO nanowires fabricated in
 330 anodized aluminum oxide membranes¹³ and MgO/NiO
 331 core/shell heterostructured NWs.⁴⁵ Our preliminary results
 332 suggest that the NiO NW arrays grown by the MEO
 333 method also show bipolar resistive switching (Figure 7).
 334 We scratched near the edge of the sample to reach the
 335 Ni foil underneath the NWs, and then the exposed Ni foil
 336 and a Pt pad deposited on the NiO NW arrays served as
 337 the electrodes for the switching experiments (the inset of
 338 Figure 7). When a voltage was applied as 0 → 4 V, the cur-
 339 rent through the Pt/NiO/Ni device increased dramatically
 340 to the μA range, indicating a “SET” process. The device
 341 can be switched back to the initial high resistance state
 342 by applying a voltage sweep as 0 → -4 V, indicating a
 343 “RESET” process. The ON/OFF ratio between the high re-
 344 sistance state and the low resistance state reached 2 or-
 345 ders of magnitude.

346 Furthermore, the novel synthesis approach pre-
 347 sented here can also be applied to other metal oxides.

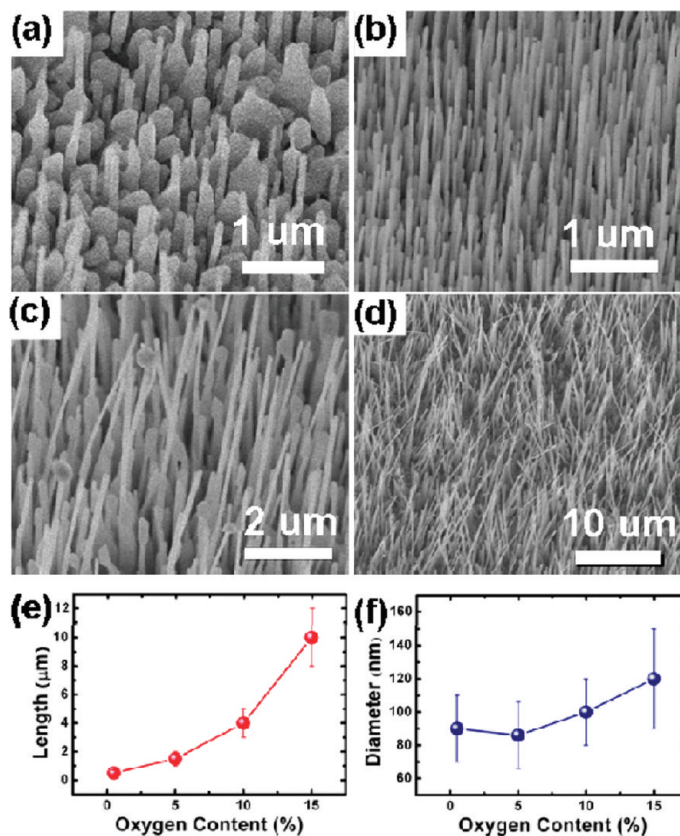


Figure 6. Morphology evolution of the NiO NW arrays with different growth atmospheres: (a) 0.5% O₂, (b) 5% O₂, (c) 10% O₂, and (d) 15% O₂ in the carrying gas. (e) Dependence of the NW length on the oxygen content. (f) Dependence of the NW diameter on the oxygen content.

348 One prerequisite of the application of this method is
 349 the formation of a thin granular oxide scale covering
 350 the metal foil which critically facilitates the etching pro-
 351 cess. Considering that the Gibbs energy of forming
 352 CoO is similar to that of forming NiO,⁴⁶ we tried our
 353 MEO method on the fabrication of CoO NW arrays. In-
 354 deed we found that CoO NW arrays can also be grown
 355 on Co foils following a similar route, as shown in Figure
 356 8a. Our preliminary results suggest that, similar to NiO,
 357 NWs of CoO can also be grown *via* the MEO method.
 358 In this case, we used Co foils as the substrate and CoCl₂
 359 as the source. The experimental growth conditions are
 360 similar to those for the growth of NiO NWs. As shown in
 361 the SEM picture, the CoO NW quality is yet to be opti-

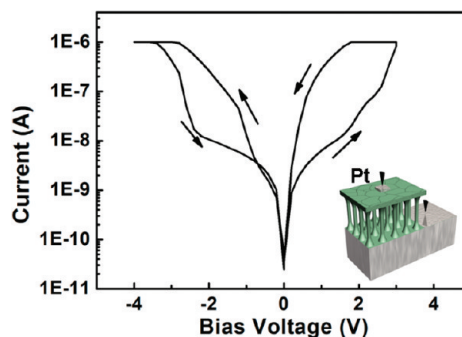


Figure 7. I–V characteristics of the NiO NW arrays. Inset shows the schematic diagram of the measurement.

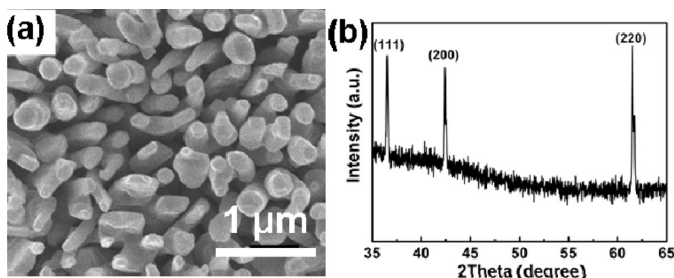


Figure 8. Top-view SEM image and XRD data of cobalt oxide (CoO) NW arrays.

mized; the NWs appear shorter than the NiO counterpart and the NWs diameters are not uniform. Nevertheless, the XRD data in Figure 8b suggests that the growth product is CoO without second phase. Thus these preliminary results encourage the application of this MEO method to the synthesis of other metal oxide nanowires.

EXPERIMENTAL SECTION

Nanowire Synthesis. The synthesis of the NiO NW arrays was carried out in a home-built vapor transport system comprising a horizontal tube furnace (Lindberg/Blue mini-mite) and a gas delivery and pumping system.^{28–30} NiCl₂ powder with a weight of 0.1 g (Aldrich, 98%) was introduced into two one-end-open quartz tubes and put opposite to each other in the middle of the furnace quartz tube. Ni foils (0.125 mm thick, Aldrich 99.9%) were cut into small pieces with a typical size of 10 mm × 10 mm and cleaned by acetone. After cleaning, the foils were mechanically polished using a Buehler MiniMet 1000 polisher, following three steps: (1) polishing with SiC paper disks (120-grit) under water cooling; (2) using rigid grinding disks (Metadi Supreme, 3-μm PC); (3) using microcloth pad with 0.05-μm alumina suspension. Subsequently, the foils were electrochemically polished in a mixture of HClO₄ and C₂H₆O (1:4 volume ratio). As a final step of foil preparation, the Ni foils were annealed at 1000 °C for 8 h in a gas mixture of 90% Ar and 10% H₂ in order to decrease the internal stress and to further improve the crystalline quality. After these polishing and annealing procedures, the Ni foils were placed at the middle of the furnace tube between the small tubes containing the NiCl₂ powder. For the growth, the quartz tube was evacuated to a background vacuum of a few mbar, and the furnace was heated up to 600 °C at a rate of 50 °C/min. Then, a mixture gas of Ar and O₂ was introduced with a constant flow rate of 100 sccm, and the pressure inside the quartz tube was maintained at 800 mbar. Finally, the furnace was heated up further to the growth temperature and maintained for 15 min. After the furnace was cooled to room temperature, a layer of green products was grown on the surface of the Ni foils.

Characterization. A JEOL JSM-6700F field emission scanning electron microscope (SEM) and a JEOL 2100 high resolution transmission electron microscope (HRTEM) were used to study the morphology and structure of the samples. The composition was determined by an attached energy-dispersive X-ray spectrometer (EDS). The X-ray diffraction (XRD) analysis was carried out using a Bruker AXS D8 Advanced powder diffractometer with Cu K α radiation. Focused ion beam (FIB) milling was performed using a XT Nova Nanolab 200. The electrical characterizations were carried out at room temperature on a probe station equipped with a Keithley 4200 semiconductor parameter analyzer.

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CONCLUSIONS

Single crystal NiO NW arrays have been synthesized for the first time through a facile vapor-based metal-etching-oxidation method. The diameter and the length of the nanowire can be well controlled by adjusting the growth temperature and the carrying gas content. Furthermore, by replacing NiCl₂ with CoCl₂ in the source powder, uniform Co doping can be introduced in the NiO NWs. We also used CoO as an example to demonstrate that this method can be applied to synthesize NWs of other oxides. In the general context, it is well-known that materials belonging to the high-symmetry cubic space groups are notoriously difficult to grow into anisotropic single crystalline nanowires. Therefore, our method presents a novel and general route to synthesize low dimensional metal oxide nanostructures, and will help to advance the research on their functionalities and applications.

Supporting Information Available: The change of Ni foil crystal quality after annealing; SEM image of the NiO NW arrays milled by focused ion beam; the TEM images of the nanowire with [220] growth direction; the weight change and the XRD patterns of the Ni foil taken at various growth stages used for further growth mechanism discussion. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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