

OPEN

Double-walled iron oxide nanotubes via selective chemical etching and Kirkendall process

João Azevedo¹, M. P. Fernández-García^{2,6}, César Magén^{3,4,5}, Adélio Mendes¹, João P. Araújo² & Célia T. Sousa²

Double-walled oxide nanotube structures are interesting for a wide range of applications, from photocatalysis to drug delivery. In this work, a progressive oxidation method to fabricate double-walled nanotube structures is reported in detail. The approach is based on the electrodeposition of metallic iron nanowires, in porous alumina templates, followed by a selective chemical etching, nanoscale Kirkendall effect, a fast oxidation and out-diffusion of the metallic core structure during thermal annealing. To validate the formation mechanism of such core-shell structure, chemical composition and atomic structure were assessed. The resulting hematite nanotubes have a high degree of uniformity, along several microns, and a nanoscopic double-walled structure.

Iron oxides have attracted much interest for their potential application in nanotechnology^{1–4}. In particular hematite (α -Fe₂O₃), being the most stable form among several Fe oxide phases⁵, is of significant scientific and technological importance for its wide application in catalysis/photocatalysis^{6,7}, gas sensors⁸ and drug delivery⁹. One of the greatest challenges with this material is its nanostructuring, which is required to increase the surface area for better reaction kinetics^{5,10}, light absorptivity^{6,11} or electronic conductivity¹². Thus, low-cost and scalable nanostructuring production methods are essential challenges in which researchers are focusing on, so that the full potential of this material is exploited.

Various α -Fe₂O₃ nanostructures with different morphologies such as nanoparticles^{8,12}, nanowires (NWs)^{13,14} and nanotubes (NTs)^{15,16} have already been successfully synthesized. Additionally, several works reported, recently, the synthesis of hollow iron oxide nanoparticles through the Kirkendall effect^{13–17}. The major part of these works refers to the fabrication of hollow spherical magnetite nanoparticles, but also nanorods and nanocubes were reported. These hollow structures are an opportunity for constructing complex nanodevices, but the reported methods were limited to spherical (or quasi-spherical) nanoparticles with many difficulties in the precise control of the final dimension and morphology of the nanostructure. Given that fundamental properties of materials can be tuned through morphology and size, precise structure control is of utmost importance^{18,19}. For example, nanostructured α -Fe₂O₃ has been studied as an anode material for lithium ion batteries for its high capacitance and it was demonstrated that higher porosity is advantageous over repetitive cycling¹⁵; this way a nanotube would provide better results than a nanorod, for example. Also, quantum confinement in very thin nanostructures, such as nanotubes with extremely thin walls, can be used to tune the band edges and band gap of a semiconductor for photoelectrochemical applications¹⁹. In this way, template-assisted methods are advantageously providing a highly flexible and controlled template that shapes the desired nanoparticle to the intended dimensions. Some reports have been published on the synthesis of α -Fe₂O₃ NTs using template-assisted methods^{20,21}, but no report of a double-walled NT structure using such methods has been made. Moreover, the reported nanostructures were relatively wide in diameter (>100 nm), decreasing the attainable surface-to-volume

¹LEPABE - Faculdade de Engenharia, Universidade do Porto, Rua Dr. Roberto Frias, 4200-465, Porto, Portugal.

²IFIMUP and Departamento de Física e Astronomia da Faculdade de Ciências da Universidade do Porto, Rua do Campo Alegre 687, 4169-007, Porto, Portugal. ³Instituto de Ciencia de Materiales de Aragón (ICMA), Universidad de Zaragoza-CSIC, 50009, Zaragoza, Spain. ⁴Laboratorio de Microscopías Avanzadas (LMA) - Instituto de Nanociencia de Aragón (INA), Universidad de Zaragoza, 50018, Zaragoza, Spain. ⁵Departamento de Física de la Materia Condensada, Universidad de Zaragoza, 50009, Zaragoza, Spain. ⁶Department of Physics, University of Oviedo, 33007, Oviedo, Spain. Correspondence and requests for materials should be addressed to C.T.S. (email: celiasousa@fc.up.pt)

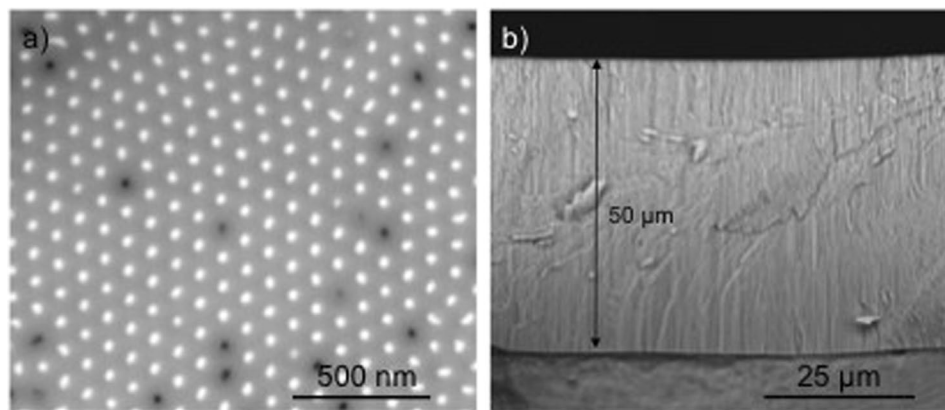


Figure 1. (a) Top view SEM images showing the high-density of Fe nanowire array. (b) Cross section reveals the thickness of the template, without any cracks.

ratio. The reason for this is the difficulty to make a controlled chemical etching to erode the nucleus and leave the walls intact.

A novel and scalable fabrication method of crystalline α -Fe₂O₃ double-walled NTs grown on porous anodic alumina (PAA) templates with controllable dimensions is here reported. For the first time, 1D Fe nanostructures grown inside PAA templates were converted into the desired α -Fe₂O₃ phase with double-walled NT morphology by combining a controlled selective chemical etching and the Kirkendall effect through an annealing route. A detailed morphological analysis was performed to reveal the growth mechanism of double-walled hollow nanostructures. The key role of the Kirkendall effect, to obtain double-walled nanostructures instead of reported single-walled, is here addressed.

Results and Discussion

Double-walled hematite nanotubes. Crystalline Fe NWs were grown inside PAA templates into pores of 35 nm diameter, typical from oxalic acid self-organized regime anodization (Fig. 1a). Oxalic acid was used due to its high reproducibility rate and because it yields nanometric pore diameter necessary for high surface areas²². Other pore sizes are currently under study and not reported in this work. The deposition was highly homogeneous up to lengths of 50 μ m, which were used to achieve large aspect ratios ($>1.4 \times 10^3$). Cross-section scanning electron microscopy (SEM) images revealed no defects along the PAA template and each wire was continuous and homogeneous (Fig. 1b).

The structure and phase composition of the samples were determined by X-ray diffraction (XRD) (Fig. S2). The positions of all Bragg reflections in Fe NWs coincide with the Bragg reflections characteristic of metallic α -Fe phase. Metallic Fe NWs show a preferential bcc-structure and only two peaks, (110) and (200), can be found in the patterns. From the most intense peak (110), a grain size of 17.8 nm was estimated.

The encapsulated NWs were removed from the PAA template, using a chemical etching solution containing H₃PO₄, known to promote selective coordination-assisted dissolution, which is discussed in detail later on in this paper. The Bragg reflections of the Fe NWs after template removal were broader than in template (Fig. S2), since the vertical alignment provided by the PAA was lost with its dissolution and the NWs became dispersed. The relative size of the peaks also changed, related to textured NWs in template versus more polycrystalline configuration in dissolved NWs.

The NWs were annealed at 600 °C in air during 6 h. The morphology of the obtained nanostructures was analyzed by scanning transmission electron microscopy (STEM) and a double-walled nanotube structure was observed – Fig. 2a. The electron energy loss spectroscopy (EELS) compositional line profiles of this double-walled NTs indicate Fe and O signals variation consistent with the double-walled structure – Fig. 2b. From the compositional line profiles, a distance between inner and outer tubes of ca. 4 nm, with corresponding diameters of 21 nm and 46 nm, was estimated. Inner and outer walls have different thicknesses (8 nm and 4 nm, respectively), which is 5 times thinner than previous reports, explained by the small features of the selected PAA template, providing a much larger surface-to-volume ratio¹³. Very thin walls are interesting for different applications due to the quantum confinement effects¹⁹. The NTs bottom ends (designated as the side originally adjacent to the Al foil before removing the PAA template) were dissolved during the alumina chemical etching, so no dendritic structure was present and most NTs were opened/broken at both ends. The inner and outer tubes do not appear to be connected at any specific point and only very thin connection bridges were observed – Fig. 2c. These double-walled nanotubes present a higher geometrical surface area than other reported nanostructures – Fig. S1.

The composition and crystallographic structure of the NTs were assessed by different techniques. Raman spectroscopy was performed on reference samples of the most common Fe oxides (maghemite, magnetite and hematite) – Fig. 3a. The NT modes positioned at 221, 285, 397, 602 and 645 nm in the spectrum (Fig. 3b) are all indexed to the characteristic bands of α -Fe₂O₃ confirming that the NTs are constituted of only one Fe oxide phase – α -Fe₂O₃. The XRD pattern of the NTs is shown in Fig. 3c. All the Bragg reflections were identified and correspond to the α -Fe₂O₃ phase. The spectrum was fitted with hematite (PDF Card No.: 2101167) and is in great

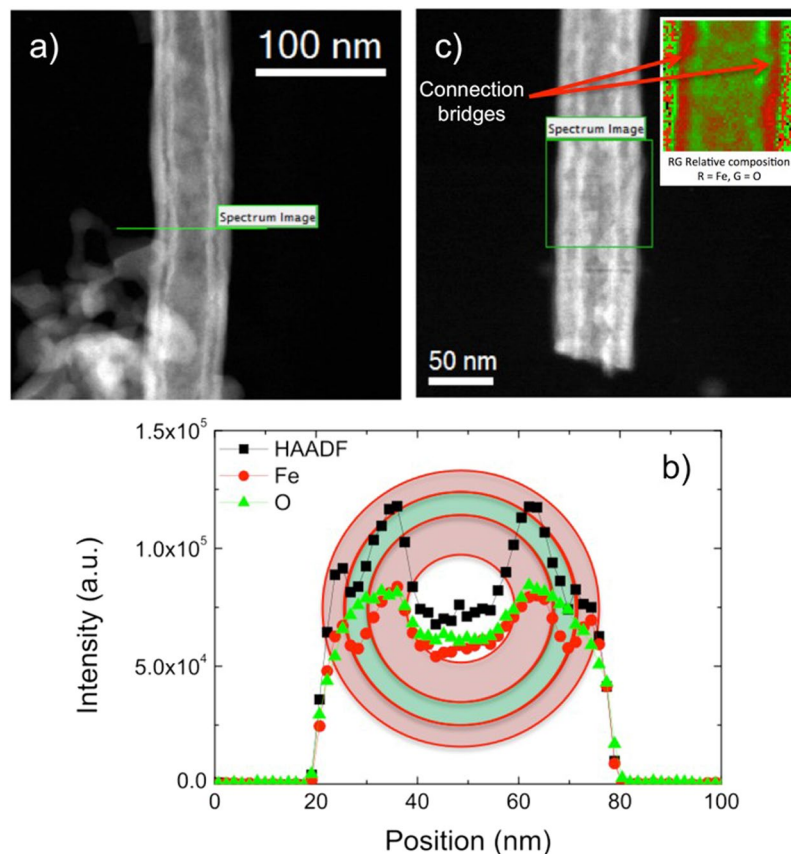


Figure 2. (a) High angle annular dark field (HAADF) STEM side-view image of a single double-walled nanotube; (b) STEM-EELS chemical profile across α - Fe_2O_3 double wall NT along the green line in (a); (c) HAADF-STEM side-view of another double-walled nanotube with composition on inset.

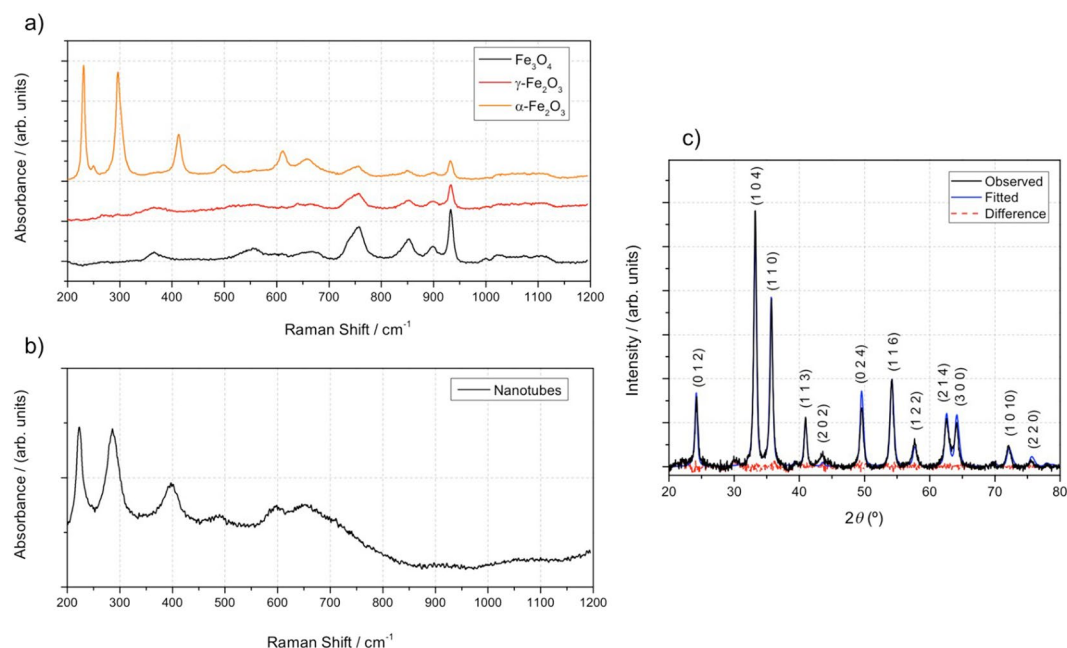


Figure 3. (a) Raman spectra of three most common iron oxide phases; (b) Raman spectra of double-walled nanotubes (peaks fit well with hematite); (c) X-ray diffraction patterns of double-walled nanotubes, with the corresponding miller indexes are marked. Peak values were compared to reference – Table S1.

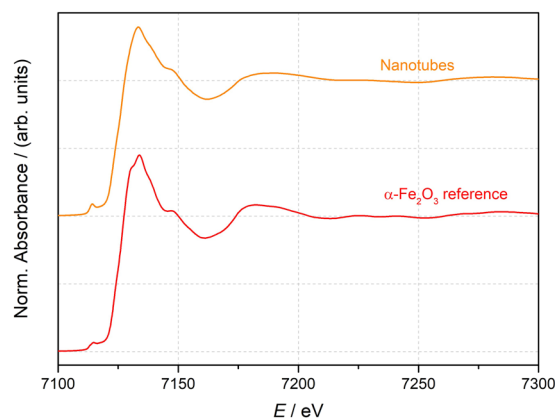


Figure 4. Experimental Fe K-edge XANES spectrum of a double-walled nanotube sample (orange line) measured at room temperature together with the hematite reference (red line).

accordance with the α -Fe₂O₃ spectrum. All XRD peaks can be indexed to α -Fe₂O₃ R-3C hexagonal cell with lattice constants of $a = b = 5.0380 \text{ \AA}$ and $c = 13.7662 \text{ \AA}$. A grain size of 11.5 nm was determined using the Scherrer equation.

To confirm the composition of the nanostructures in a precise manner, X-ray absorption spectroscopy (XAS), a technique selective to atomic species, was employed. The existence of Fe hydroxides (goethite or ferrihydrite) magnetite and maghemite was discarded by both XRD and Raman spectrometry and therefore, only hematite was considered for the data analysis of the X-ray absorption spectroscopy near edge structure (XANES) spectrum. In Fig. 4 the experimental Fe K-edge XANES spectrum of the sample is shown together with the reference of α -Fe₂O₃ measured in the same equipment. It can be observed that the experimental spectrum reproduces the shape, energy position of the different spectral features, their relative energy separation and the intensity ratio of the reference sample. This result is in accordance with the XRD measurements, validating the oxidation of the nanostructure to α -Fe₂O₃.

Growth Mechanism. From STEM it is clear that a double-walled nanostructure is attainable by a simple chemical etching and annealing route. But the formation mechanism of these structures is not straightforward. The morphology of the samples at the various formation stages was studied and is presented in Fig. 5.

Formation of heterostructures by selective chemical etching. The as-deposited α -Fe sample had a typical structure of NWs fabricated through template-assisted electrodeposition, with no visible morphology defects (Fig. 5a). After several hours of selective chemical etching in a H₃PO₄ and H₂Cr₂O₇ solution, the NWs were converted to a core-shell structure, with a very thin and smooth outer layer and a granular nucleus (Fig. 5b). In this step, two main processes occurred: the formation of an iron oxide shell and the partial dissolution. When submerged for long times in an aqueous solution, the α -Fe nanowires slowly oxidized. A compact layer of iron oxide was formed on the surface of nanowires without visible gaps (Fig. 5b). The oxide shell was passive and stable in the aqueous solution²³.

The shell diameter of ~46 nm was larger than the previous Fe NWs 35 nm diameter on the PAA template, imposed by the alumina template, which confirmed the oxidation of Fe accompanied with an expansion due to oxygen incorporation. During this intermediate stage, the nanowires break since they are no longer supported by the PAA. This yields aspect ratios ranging from ca. 22 to 109, that are the same as the final product, the double-walled nanotube. Optimization of the chemical etching step might decrease the nanowire breakage and further increase the final aspect ratio. XRD of the sample after alumina template removal (Fig. S2) indicated the presence of Fe and no trace of Fe oxide. The absence of Fe oxide phases was attributed to the amorphous nature of the outer wall that only crystallizes after annealing at high temperatures. After long exposures to etching solutions, the iron core was transformed due to the outward mass transport of iron to form the iron oxide shell combined with the dissolution of the iron shell. In this case, single-walled nanotubes can be achieved, as reported for hollow spherical nanoparticles^{14,24–26}.

To achieve double-walled nanostructures, the α -Fe nucleus was preserved with a radius between 30–35 nm by decreasing the etching time to less than 20 h. It is observed that the remaining granular nucleus has several voids and a hollow tubular structure at the tips, left from the dissolved or detached nucleus (Fig. 5b). Based on this morphology and crystallography evolution between the previous steps, it is proposed a dissolution-based mechanism for the transformation of NW to core-shell heterostructure.

It has been demonstrated that iron oxide nanoparticles, in the presence of phosphate ions, promote a selective coordination-assisted dissolution¹³. Similarly, here Fe was first oxidized²⁷ and then selectively dissolved. The presence of a smooth sheath and the voids along the structures at step 2 (Fig. 5b) supports the coordination-assisted dissolution mechanism preferentially in the grain boundaries of the iron core and in the tips of the tubes, where the oxidation of iron is a self-limiting process. Additionally, the flat outer oxide shell enables highly homogeneous and reproducible nanostructures, with many high-energy sites in the iron core that can be easily exposed to produce an internal nanotube.

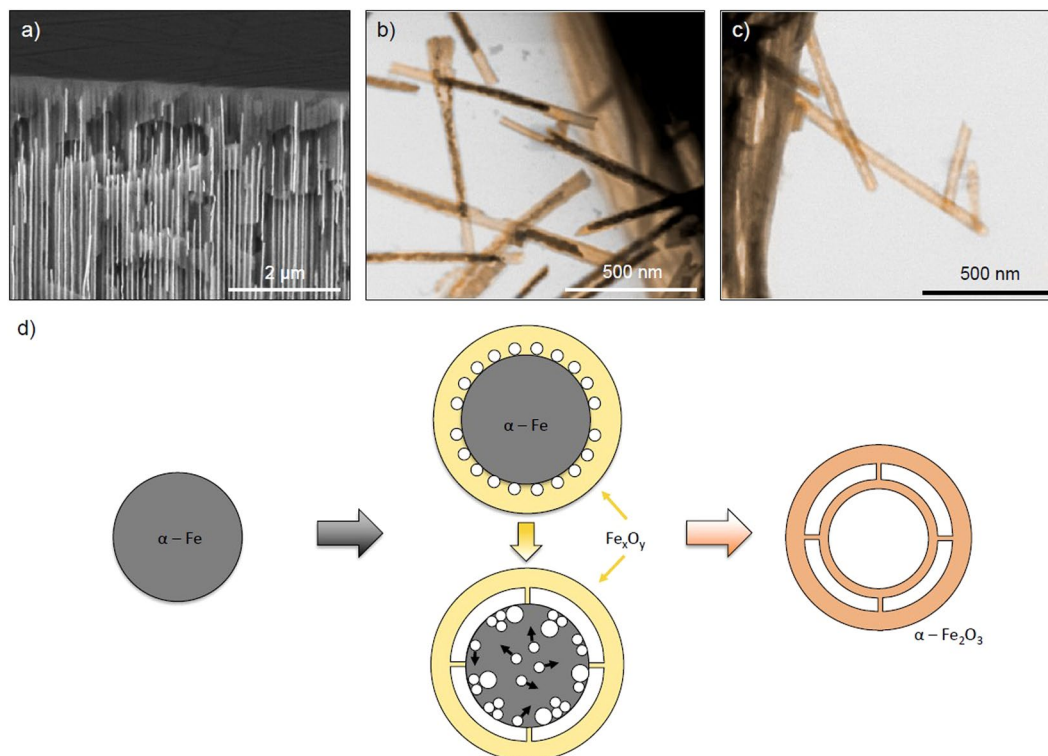


Figure 5. Morphology evolution of the nanostructures: SEM and STEM images of the products obtained (a) as prepared in template, (b) after chemical etching and (c) after annealing. STEM images were colored for better representation. (d) Schematic showing different stages of double-walled iron oxide formation: iron nanowire, iron oxide shell and selective coordination-assisted dissolution, void formation by the Kirkendall effect, complete oxidation of structure.

Oxidation in high-temperature regime. The voids created during the etching process in the core of the heterostructures increase with the annealing ramp, as reported²⁸. After 6 h of annealing at 600 °C the core structure is converted to an interior smooth oxide tube, inserted into the outer layer, created in the etching process (Fig. 5c). At this temperature the outer diameter increased once more, indicating a complete oxidation and crystallization of the structure. Annealing of Fe NWs in template was performed and no indication of tube formation was observed, as reported elsewhere²¹. On the other side, if the annealing is performed in dispersed nanoparticles or nanowires, the oxygen intake and Fe diffusion creates a hollow nanostructure^{29,30}. Only if the annealing is performed on core-shell nanostructures a double-walled can be achieved. The inner wall formed during the annealing can be explained by the Kirkendall effect, where the rapid formation and growth of vacancies along the edge of the core compared to vacancy diffusion, leads to the conversion of the iron core into a secondary hematite tube^{28,31–33}. STEM images show very tenuous connection between both tubes and it is here hypothesized that many small bridges are preserved during the vacancies formation and prevent both tubes to collapse. These connections are particular features of this method and its formation process is illustrated in Fig. 5d. With the increasing temperature, there is a continuous diffusion of the iron core and oxide shell through the interface between them, with consequent formation of excess vacancies at the center of the core and small Kirkendall voids at the iron/oxide interface³¹. Then, during the annealing time, the voids begin to collapse leading to the enlargement of the hollow core and the hollow space between the inner and the outer tubes where the skeletal bridges were preserved.

The proposed new methodology demonstrates that, if the annealing stage is performed after the NWs are immersed in a phosphate containing solution a double walled NT can be formed in a reproducible manner. Given that the template dictates the initial shape of the NW, a variety of aspect ratios should be obtained. Presently, various porous alumina templates are being tested with larger and smaller diameters. With recent advances in porous alumina template control, it should be even possible to use modulated template to a variety of length-to-diameter ratios for more complex applications^{28,34}.

Methods

Sample preparation. Double-walled iron oxide NTs were prepared from arrays of metallic Fe NWs that were fabricated as described previously^{22,35}. Briefly, a high purity thick Al foil was rinsed and electropolished to reduce surface roughness and create nanopatterns for posterior pore nucleation. The Al foils were subjected to a two-step anodization process to create a PAA template. The first one in 0.3 M oxalic acid, (COOH)₂, at 40 V and 2 °C. Secondly, to obtain high aspect ratio NWs, a second anodization was carried out during 20 h (corresponding to 50 μm template thickness). Before electrodeposition, the anodization potential was lowered to 8 V to form

dendritic channels³⁶. Pulsed electrodeposition was employed to fill the template with Fe NWs in conditions previously optimized. A mixture of 153 g·L⁻¹ Fe (II) sulfate heptahydrate, FeSO₄·7H₂O, 45 g·L⁻¹ boric acid, H₃BO₃, and 1 g·L⁻¹ ascorbic acid, C₆H₈O₆, was used as a precursor in electrodeposition. A current density of 94 mA·cm⁻² was applied. The obtained PAA templates were completely filled. To ensure that no Fe film was left on the surface of the template they were mechanically polished using alumina powder²².

Afterwards, the alumina template was dissolved in an aqueous solution of phosphoric acid, 0.4 M H₃PO₄, and dichromic acid, 0.2 M H₂Cr₂O₇, at 60 °C (removal rate: 50 nm·min⁻¹), resulting in core-shell nanostructures. These nanostructures were dispersed within an ultrasound bath and suspended in the previous solution during 1 h. After complete removal of the alumina template, the nanostructures were washed several times in ethanol. The dispersed nanostructures were then annealed in air at 600 °C for 6 h, at 2 °C/min heating and cooling ramps resulting in double-walled iron oxide NTs.

Sample characterization. The morphology and surface topography of the samples were characterized using a high-resolution SEM in a FEI Quanta 400 FEG ESEM with an in-lens detector for secondary electrons. SEM imaging was carried out with an acceleration voltage of 15 kV and a working distance of about 10 mm. Cross-sectional images were acquired from freshly cleaved samples. STEM was performed in a FEI Titan 60–300 operated at 300 kV and equipped with an aberration corrector for the probe. STEM spectrum imaging combining HAADF imaging and EELS was used for the local chemical analysis of the NTs. Chemical line profiles and maps of the NTs were obtained from EELS spectrum images by integrating the O-K and Fe L_{2,3} edges intensity after background extrapolation and subtraction.

The crystalline structure of the samples was analyzed by XRD in a Rigaku SmartLab diffractometer in the $\theta - 2\theta$ parallel beam geometry using the Cu – K α line with wavelength $\lambda = 1.540593$ Å. The spectra were acquired by moving the detector between $2\theta = [20^\circ, 80^\circ]$ at a scan rate of 0.5 deg·min⁻¹ with a step width of 0.04°. The spectra peaks were fitted using a Lorentzian function to determine the average grain size according to the Scherrer equation, as described elsewhere³⁵. Reflection patterns were matched to reference values – Table S1. To measure the crystalline structure of Fe NWs in the template, the Al substrate was removed to eliminate its contribution. For that, the samples were submerged in 0.5 M CuCl₂ and 2.8 M HCl solution.

Raman spectra were determined on a LabRAM by Jobin-Yvon micro-sampling with the excitation wavelength of 633 nm of a He-Ne laser at 100x amplification. To avoid sample damaging the laser power was 10 mW. Gratings of 1800 gr/mm were used. The measurements were from 200 to 1200 cm⁻¹ using a 100 μ m slit. Each measurement was done over 30 s at two different places in each sample to assure a correct reproducibility of the spectrum. The samples were put in a low-susceptibility plastic sample holder for measurements. X-ray absorption spectroscopy measurements at the Fe K-edge in transmission were developed at the XAFS beamline of the Elettra synchrotron with an energy range of 6812–7969 eV. An Fe metal foil spectrum was also recorded for energy calibration. For each sample, three spectra were acquired for statistics with an integration time of 5.00 s. The experimental XANES spectrum of the samples was normalized at high energy after background subtraction to eliminate thickness dependence. To quantify the composition of each sample, the spectra were fitted with reference Fe/Fe oxide absorption curves using a specifically made script.

Conclusions

Using a simple and low-cost fabrication method, the synthesis of double-walled hematite nanotubes was achieved using porous anodic alumina templates to control the overall morphology of the nanotubes. The features of these structures were assessed by STEM revealing diameters smaller than 50 nm and wall thicknesses less than 10 nm, in addition to a subtle network of connection bridges between inner and outer nanotube walls. The crystallography and composition were evaluated precisely revealing a single iron oxide phase – hematite. Finally, the growth mechanism of these double-walled nanotubes was revealed. A synergetic combination of a selective chemical etching and annealing on metallic nanowires allowed the fabrication of double-walled oxide nanotubes with controlled morphology.

References

- Laurent, S. *et al.* Magnetic iron oxide nanoparticles: synthesis, stabilization, vectorization, physicochemical characterizations, and biological applications. *Chem. Rev.* **108**, 2064–2110 (2008).
- Lu, Y., Yin, Y., Mayers, B. T. & Xia, Y. Modifying the Surface Properties of Superparamagnetic Iron Oxide Nanoparticles through A Sol–Gel Approach. *Nano Lett.* **2**, 183–186 (2002).
- Park, J. *et al.* One-Nanometer-Scale Size-Controlled Synthesis of Monodisperse Magnetic Iron Oxide Nanoparticles. *Angew. Chem.* **117**, 2932–2937 (2005).
- Wu, H. B., Chen, J. S., Hng, H. H. & Lou, X. W. Nanostructured metal oxide-based materials as advanced anodes for lithium-ion batteries. *Nanoscale* **4**, 2526–2542 (2012).
- Tadić, M. *et al.* Morphological, structural and magnetic properties of α -Fe₂O₃ nanoparticles in an amorphous alumina matrix obtained by aqueous combustion method. *JALC* **486**, 839–843 (2009).
- Steier, L. *et al.* Understanding the Role of Underlayers and Overlayers in Thin Film Hematite Photoanodes. *Adv. Funct. Mater.* **24**, 7681–7688 (2014).
- Wedegge, K., Azevedo, J., Khataee, A., Bentien, A. & Mendes, A. Direct Solar Charging of an Organic-Inorganic, Stable, and Aqueous Alkaline Redox Flow Battery with a Hematite Photoanode. *Angew. Chem. Int. Ed. Engl.* **55**, 7142–7147 (2016).
- Sun, B. *et al.* Synthesis of Mesoporous α -Fe₂O₃ Nanostructures for Highly Sensitive Gas Sensors and High Capacity Anode Materials in Lithium Ion Batteries. *The Journal of Physical Chemistry C* **114**, 18753–18761 (2010).
- Gupta, A. K. & Gupta, M. Synthesis and surface engineering of iron oxide nanoparticles for biomedical applications. *Biomaterials* **26**, 3995–4021 (2005).
- Ling, Y., Wang, G., Wheeler, D. A., Zhang, J. Z. & Li, Y. Sn-doped hematite nanostructures for photoelectrochemical water splitting. *Nano Lett.* **11**, 2119–2125 (2011).
- Sivula, K., Le Formal, F. & Gratzel, M. Solar water splitting: progress using hematite (α -Fe₂O₃) photoelectrodes. *Chem Sus Chem* **4**, 432–449 (2011).

12. Warren, S. C. *et al.* Identifying champion nanostructures for solar water-splitting. *Nat Mater* **12**, 842–849 (2013).
13. Jia, C.-J. *et al.* Iron Oxide Tube-in-Tube Nanostructures. *The Journal of Physical Chemistry C* **111**, 13022–13027 (2007).
14. An, K. *et al.* Synthesis of uniform hollow oxide nanoparticles through nanoscale acid etching. *Nano Lett.* **8**, 4252–4258 (2008).
15. Cho, J. S., Park, J. S. & Kang, Y. C. Preparation of Hollow Fe₂O₃ Nanorods and Nanospheres by Nanoscale Kirkendall Diffusion, and Their Electrochemical Properties for Use in Lithium-Ion Batteries. *Sci. Rep.* **6**, 38933 (2016).
16. Khurshid, H., Li, W., Tzitzios, V. & Hadjipanayis, G. C. Chemically synthesized hollow nanostructures in iron oxides. *Nanot* **22**, 265605 (2011).
17. Yin, Y. *et al.* Formation of hollow nanocrystals through the nanoscale Kirkendall effect. *Sci* **304**, 711–714 (2004).
18. Mohapatra, S. K., John, S. E., Banerjee, S. & Misra, M. Water Photooxidation by Smooth and Ultrathin α -Fe₂O₃ Nanotube Arrays. *Chem. Mater.* **21**, 3048–3055 (2009).
19. Vayssieres, L. *et al.* One-Dimensional Quantum-Confinement Effect in α -Fe₂O₃ Ultrafine Nanorod Arrays. *Adv. Mater.* **17**, 2320–2323 (2005).
20. Lim, J. H., Min, S. G., Malkinski, L. & Wiley, J. B. Iron oxide nanotubes synthesized via template-based electrodeposition. *Nanoscale* **6**, 5289–5295 (2014).
21. Mao, A. *et al.* Controlled synthesis of vertically aligned hematite on conducting substrate for photoelectrochemical cells: nanorods versus nanotubes. *ACS Appl Mater Interfaces* **3**, 1852–1858 (2011).
22. Azevedo, J. *et al.* Ultra-long Fe nanowires by pulsed electrodeposition with full filling of alumina templates. *Materials Research Express* **1**, 015028 (2014).
23. Sarathy, V. *et al.* Aging of Iron Nanoparticles in Aqueous Solution: Effects on Structure and Reactivity. *The Journal of Physical Chemistry C* **112**, 2286–2293 (2008).
24. Huang, J. *et al.* One-Dimensional Chainlike Arrays of Fe₃O₄ Hollow Nanospheres Synthesized by Aging Iron Nanoparticles in Aqueous Solution. *Journal of Physical Chemistry C* **113**, 12067–12071 (2009).
25. Zhang, Q., Lee, I., Ge, J., Zaera, F. & Yin, Y. Surface-Protected Etching of Mesoporous Oxide Shells for the Stabilization of Metal Nanocatalysts. *Adv. Funct. Mater.* **20**, 2201–2214 (2010).
26. Zhang, Q., Zhang, T., Ge, J. & Yin, Y. Permeable silica shell through surface-protected etching. *Nano Lett.* **8**, 2867–2871 (2008).
27. Muralidharan, V. S. & Rajagopalan, K. S. Kinetics and mechanism of corrosion of iron in phosphoric acid. *Corros. Sci.* **19**, 199–207 (1979).
28. Ren, Y. *et al.* Formation of Nickel Oxide Nanotubes with Uniform Wall Thickness by Low-Temperature Thermal Oxidation Through Understanding the Limiting Effect of Vacancy Diffusion and the Kirkendall Phenomenon. *Adv. Funct. Mater.* **20**, 3336–3342 (2010).
29. Wang, C. M. *et al.* Void formation during early stages of passivation: Initial oxidation of iron nanoparticles at room temperature. *JAP* **98** (2005).
30. Krajewski, M. *et al.* High temperature annealing of iron nanowires. *physica status solidi (a)* **212**, 862–866 (2015).
31. Tu, K. N. & Gösele, U. Hollow nanostructures based on the Kirkendall effect: Design and stability considerations. *ApPhL* **86** (2005).
32. Nakamura, R., Matsubayashi, G., Tsuchiya, H., Fujimoto, S. & Nakajima, H. Formation of oxide nanotubes via oxidation of Fe, Cu and Ni nanowires and their structural stability: Difference in formation and shrinkage behavior of interior pores. *AcMat* **57**, 5046–5052 (2009).
33. Nakamura, R., Matsubayashi, G., Tsuchiya, H., Fujimoto, S. & Nakajima, H. Transition in the nanoporous structure of iron oxides during the oxidation of iron nanoparticles and nanowires. *AcMat* **57**, 4261–4266 (2009).
34. Lee, W. & Kim, J. C. Highly ordered porous alumina with tailor-made pore structures fabricated by pulse anodization. *Nanot* **21**, 485304 (2010).
35. Azevedo, J., Sousa, C. T., Mendes, A. & Araújo, J. P. Influence of the Rest Pulse Duration in Pulsed Electrodeposition of Fe Nanowires. *Journal of Nanoscience and Nanotechnology* **12**, 9112–9117 (2012).
36. Nielsch, K., Müller, F., Li, A. P. & Gösele, U. Uniform Nickel Deposition into Ordered Alumina Pores by Pulsed Electrodeposition. *Adv. Mater.* **12**, 582–586 (2000).

Acknowledgements

J. Azevedo would like to acknowledge the Portuguese Foundation for Science and Technology (FCT) for funding (CEECIND/03937/2017). C.T. Sousa thanks FCT for financial support through the Investigador FCT program (Contract No. IF/01159/2015). M.P. Fernández-García acknowledges financial support through Spanish MINECO (research project RTI2018-094683-B-C52); Gobierno del Principado de Asturias and FICYT (through research project FC-GRUPIN-IDI/2018/000185) and University of Oviedo for R&D project 2018/00061/008 in the competitive call PAPI-18-EMERG-8. The authors are thankful to Elettra Sincrotrone for allocating beam-time and their experimental facilities. This work was partially supported by the Project POCI-01-0145-FEDER-006939 (Laboratory for Process Engineering, Environment, Biotechnology and Energy – LEPABE and “SunStorage - Harvesting and storage of solar energy”, with reference POCI-01-0145-FEDER-016387, funded by European Regional Development Fund (ERDF), through COMPETE 2020 - Operational Programme for Competitiveness and Internationalization (OPCI), by FCT - Portuguese Foundation for Science and Technology I.P. This work was financially supported by: Project PTDC/EQU-EQU/30510/2017 - POCI-01-0145-FEDER-030510 – Sunflow “Solar energy storage into redox flow batteries” funded by FEDER funds through COMPETE2020 - Programa Operacional Competitividade e Internacionalização (POCI) and by national funds (PIDDAC) through FCT/MCTES. This work was also supported by the Portuguese Fundação para a Ciência e Tecnologia (FCT) and COMPETE 2020 (FEDER) under the projects POCI-01-0141-FEDER-032527, PTDC/FIS-MAC/31302/2017, MIT-EXPL/IRA/0012/2017, PTDC/CTM-CTM/28676/2017 and PTDC/FIS-OTI/32257/2017 and the European Union’s Horizon 2020 research and innovation program under the Marie Skłodowska-Curie Grant Agreement No. 734801.

Author Contributions

J.A., M.P.F.-G. and C.T.S. performed the experiments and prepared the manuscript, while C.M. performed STEM measurements and analysis; A.M. edited the final manuscript; J.P.A. participated in data interpretation and edited the final manuscript.

Additional Information

Supplementary information accompanies this paper at <https://doi.org/10.1038/s41598-019-47704-5>.

Competing Interests: The authors declare no competing interests.

Publisher's note: Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2019