



OPEN

SUBJECT AREAS:
THEORETICAL PHYSICS
ELECTRONIC STRUCTURE
STRUCTURAL PROPERTIES
ELECTRONIC PROPERTIES AND
MATERIALS

Received 3 March 2014

Accepted 30 May 2014

Published 23 June 2014

Correspondence and requests for materials should be addressed to E.L. (elson@iq.unesp.

Structural and electronic analysis of the atomic scale nucleation of Ag on α -Ag₂WO₄ induced by electron irradiation

Juan Andrés¹, Lourdes Gracia^{1,2}, Patricio Gonzalez-Navarrete^{1,2}, Valeria M. Longo³, Waldir Avansi, Jr.⁴, Diogo P. Volanti⁵, Mateus M. Ferrer⁴, Pablo S. Lemos⁴, Felipe A. La Porta², Antonio C. Hernandes³ & Elson Longo²

¹Departament de Química Física i Analítica, UJI – Universitat Jaume I, Av. de Vicent Sos Baynat, s/n, Castelló de la Plana 12071, Spain, ²Instituto de Química, UNESP – Universidade Estadual Paulista, R. Francisco Degni, 55, Araraquara 14800-900, Brazil, ³Instituto de Física de São Carlos, USP – Universidade de São Paulo, Av. Trabalhador são-carlense, São Carlos 13560-970, Brazil, ⁴Departamento de Física, UFSCar – Universidade Federal de São Carlos, Rod. Washington Luis, km 235, São Carlos 13565-905, Brazil, ⁵Departamento de Química e Ciências Ambientais, UNESP – Universidade Estadual Paulista, R. Cristóvão Colombo, 2265, São José do Rio Preto 15054-000, Brazil.

In this work, we utilise a combination of theory, computation and experiments to understand the early events related to the nucleation of Ag filaments on α -Ag₂WO₄ crystals, which is driven by an accelerated electron beam from an electron microscope under high *vacuum*. The growth process and the chemical composition and elemental distribution in these filaments were analysed in depth at the nanoscale level using TEM, HAADF, EDS and XPS; the structural and electronic aspects were systematically studied in using first-principles electronic structure theory within QTAIM framework. The Ag nucleation and formation on α -Ag₂WO₄ is a result of the order/disorder effects generated in the crystal by the electron-beam irradiation. Both experimental and theoretical results show that this behavior is associated with structural and electronic changes of the [AgO₂] and [AgO₄] clusters and, to a minor extent, to the [WO₆] cluster; these clusters collectively represent the constituent building blocks of α -Ag₂WO₄.

ungstates of transition metals are ternary oxide semiconductors that contain a combination of covalent, ionic and metallic bonding. They have attracted considerable attention due to their unique symmetry-dependent and spontaneous polarization properties, which are technologically important in numerous applications, such as ferroelasticity, ionic conductivity and photoluminescence^{1–4}. They show, in the range of a few nanometers, a surprising tendency to evolve into ordered structures of highly sophisticated architectures. Understanding the structural and electronic properties of the materials in these systems is critical to determining their physical/chemical properties and performance.

 Ag_2WO_4 is a member of the tungstate family that crystallizes in an orthorhombic structure (α - Ag_2WO_4) with a space group of Pn2n. α - Ag_2WO_4 has been the focus of investigations by our group; we have synthesised this material using different methods (coprecipitation, sonochemistry and hydrothermal synthesis) and have studied its photoluminescence (PL) properties⁵. More recently, the formation of Ag filaments on α - Ag_2WO_4 crystal, via irradiation of electrons by an accelerated electron beam from an electron microscope under high *vacuum*, was reported⁶, and the corresponding potential applications⁷ were investigated. Our current investigation is motivated by our incomplete understanding of the underlying fundamental phenomena and mechanisms driving the nucleation and growth of Ag filaments.

While the concept of a crystalline solid as a perfect, periodic structure is at the core of our understanding of a wide range of material properties, in reality, disorder is ubiquitous and is capable of drastically influencing various properties⁸. α -Ag₂WO₄ is expected to exhibit interesting forms of structural and electronic disorder that play a central role in driving function. α -Ag₂WO₄ represents a class of inorganic compounds that exhibit unmatched structural versatility and unique intrinsic properties due to the fundamental characteristics of the interactions between its octahedral WO₆ and [AgO_x] (x = 2, 4, 6, and 7) clusters, which are its basic building blocks. These



clusters are excellent candidates to study to explore the nature of the chemical bond and the influence of the so-called finite-size effects on the physical and electronic structures of such compounds.

Our emphasis is on highlighting how the structural and electronic disorder lead to interesting and unusual physical/chemical properties in this long-established class of materials. First, because this structure exhibits different types of coordination [AgO_x] and because its synergistic coordination mode facilitates the formation of several types of clusters with low and high coordination numbers, the nature of the [WO₆] and [AgO_x] clusters and the interactions between them are inherently predisposed to the formation of flexible materials with open structures. Second, the O-Ag-O and O-W-O bonds in the clusters, which have a weak interaction between the basic units, may also be chemically modified using electron irradiation, as will be shown. Consequently, the stability of the lattice is dependent on the local bonding, and the enthalpic difference between the crystalline and amorphous forms may be very small because the local bonding arrangements are similar in both. This behavior is well established in the study of conventional glass-forming materials, and the network topology and connectivity play a critical role in stabilizing disordered states⁵. Thus, not only might these systems be somewhat predisposed to disorder, but this disorder might readily be tuned by exploiting the versatile structural chemistry of this broad family.

The goal of this work is to understand the basis of the early-nucleation and -growth of Ag filaments after the irradiation process. To investigate the $in\ situ$ growth of Ag filaments in the $\alpha\text{-}AgWO_4$ crystal, information on the structural and electronic evolution, as defined by the changes in the electron density, was obtained by density functional theory (DFT) calculations using the quantum theory of atoms in molecules (QTAIM). Experimental techniques such as transmission electron microscopy (TEM) with a high-angle annular dark field (HAADF), energy-dispersive X-ray spectroscopy (EDS), and X-ray photoelectron spectroscopy (XPS) were also employed. The results provide a valuable probe into the relationship between atomic-scale structural and electronic perturbations and their macroscopic consequences.

We focus primarily on the structural and electronic properties to answer three central questions: (i) What happens with the excess electron density, simulating the electron beam of TEM, as it approaches the surface and bulk of α-Ag₂WO₄? (ii) How are the electrons distributed in this material, and how does this distribution relate to the structural and electronic evolution? (iii) Can QTAIM properties provide insight into the strength of the bonds after electron irradiation of α-Ag₂WO₄? We studied the geometric and electronic structure of α -Ag₂WO₄ and derived a mechanism for the early events in the formation and growth of Ag filaments in the scenario of electron irradiation of $[AgO_x]$ (x = 2, 4, 6, and 7) and $[WO_6]$ clusters, which are the constituent polyhedra of α -Ag₂WO₄. We shall discuss how the analysis, provided by both the experimental and theoretical results, of the physical and electronic structure of α -Ag₂WO₄ allows us to explain the Ag-nucleation process. The discussion will address the details of image acquisition and analysis and will provide a guide to interpret the experimental results. All technical details are provided in the Supplementary Information (SI).

Results

A graphical representation of α -Ag₂WO₄ using polyhedra is presented in Fig. S1 (SI) to show [WO₆] and [AgO_x] (x = 2, 4, 6 and 7) as the building blocks of this material. The crystallographic structural characteristics of the as-prepared products investigated by XRD analysis are also presented in the Supplementary Information (Fig. S2). All of the diffraction peaks of the samples reflect an orthorhombic structure without any deleterious phases and with a space group of Pn2n and a point-group symmetry of $C_{2v}^{9,10}$; these results are in agreement with the standard values listed in the Inorganic Crystal

Structure Database (ICSD) no. 4165^{10} and with those reported in the literature 5,11 . Optimised lattice parameters and atomic positions are listed in Table S1 (SI).

The first step is the growth of metallic Ag on the α-Ag₂WO₄ crystal; this part of the process is accompanied by in situ TEM. A detailed characterisation of the chemical reactions and physical interactions involving nanomaterials requires simultaneous measurement of atomic structure and chemical composition at the nanoscale level. However, the ability to observe the formation and growth of a crystal in real time is reshaping our understanding of these molecular processes, revealing subtleties previously hidden in ensemble averages. *In situ* TEM is an elegant method for uncovering the dynamic processes in the growth of nanocrystals¹². These recent technological advancements in conjunction with high-resolution imaging provide a new opportunity to view nanoscale processes and nanocrystal growth by aggregation and coalescence has been directly observed in real time by in situ TEM studies¹²⁻²¹. The injection of electrons can also be used to modify the properties of different materials 16,22-25; for example, the localised electron probes used in TEM can be used to foster electron-induced processes on a nanometre or subnanometre scale²⁶⁻³².

TEM provides a foundation for such atomic resolution studies and is an appropriate tool for monitoring nanoparticle growth in real time^{12,18,33–36}. This *in situ* imaging allows the study of materials processes and biological, multistage reactions^{12,14,37–40}. However, the studies to date have typically been performed on samples in solution, and the environment around nanoparticles is known to change their physical and chemical properties. The formation and growth conditions of nanoparticles in solution differ markedly from those of nanoparticles in a high-vacuum environment because the surface of nanoparticles adapts to its environment. This difference has been demonstrated using *in situ* spectroscopy and microscopy techniques^{12,39,41–53}.

In a recent study⁶, we observed the growth of Ag on the α -Ag_2WO_4 structure. In the early steps of the process, the structure of α -Ag_2WO_4 was clean and smooth (Fig. 1a); a broader view of the same sample is shown in Fig. 1b. With a longer exposure (on the order of a fraction of a second) of electron irradiation, the nucleation process of several Ag nanoparticles occurred throughout the bulk structure (Fig. 1c). A greater density of Ag was observed at the end of the matrix structure, which resulted in the nucleation and growth of a thicker filament (Fig. 1d) that grew further, as shown in Figs. 1e and 1f.

A detailed EDS analysis was performed to elucidate the composition of the nanostructures. Fig. 2a presents a HAADF image of a single nanostructure on which elemental EDS analysis was performed. This structure contains nanofilaments whose growth was stimulated by the electron-beam irradiation on the α -Ag₂WO₄ surface⁶. The EDS analyses performed on regions 1 to 4 confirmed that the filaments were mainly composed of Ag (Fig. 2b). However, in the nanostructure, mainly Ag, W, and O and mainly W and O were clearly identified in regions 3 and 4, respectively. A small amount of Cu atoms was also observed, which is related to the grid.

XPS measurements were performed on the irradiated and non-irradiated $\alpha\text{-}Ag_2WO_4$ materials to compare the purity of the surface and the degree of oxidation, to gain insight into the shape evolution, and to unravel the formation mechanism of the samples. The widerange spectrum is shown in Fig. 3. The results show the presence of a large amount of carbon, and the peaks of Ag, W, and O are clearly visible, indicating that the samples were highly pure. The core-level binding energies (BEs) of O 1 s, Ag 3 d, and W 4f, which have been corrected for the surface charging effect, were determined from the respective XPS spectra and are presented in Table S2 (SI). These results are consistent with the BE values reported in the literature 54,55 .

High-resolution C 1 s, O 1 s, Ag 3 d, and W 4f XPS spectra of the as-synthesised and irradiated samples are included for comparison purposes (see Figs. S3 (a–d) (SI) and Table S3 (SI)). An analysis of the



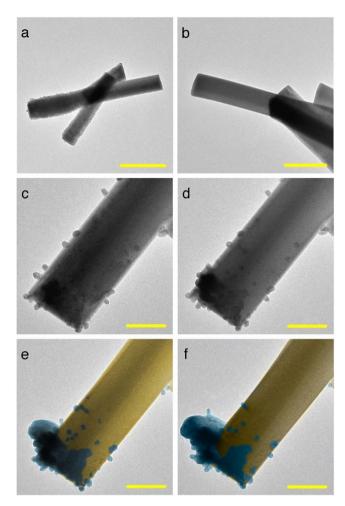
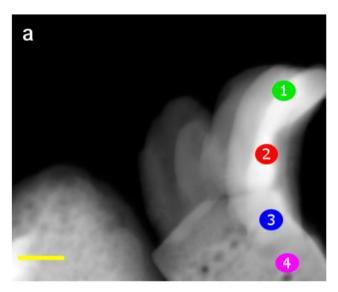


Figure 1 | TEM images of the formation of Ag filaments from the a-Ag₂WO₄ bulk. (a) and (b) TEM images obtained at different magnifications indicate a smooth and clear surface. (c)–(f) Thick Ag filaments grow at the edge of the sample, whereas other Ag nanoparticles are absorbed by the matrix. (Scale bar = 500 nm in a, 200 nm in b and, 100 nm in (c–f).

XPS results shows a strong effect on both the Ag and W atoms. The irradiated sample has a larger amount of $W^{(V)}$, which indicates greater disorder of the structure, in accordance with previous HRTEM results. The Ag $3d_{5/2}$ peak is identified at 368.10 and 367.80 eV in the spectra of the as-synthesised and irradiated samples, respectively, which suggests the presence of metallic Ag 56,57 . To confirm the presence of metallic Ag from the XPS data, the asymmetric peaks observed in the Ag $3d_{5/2}$ core-level region and the Auger $M_4N_{45}N_{45}$ peaks of $\alpha\text{-Ag}_2WO_4$ and of irradiated $\alpha\text{-Ag}_2WO_4$ were analysed (see Fig. S4 (S1)).

Our findings confirm the presence of metallic Ag, and the corresponding results agree with the results reported in the *Handbook of Monochromatic XPS Spectra* 58 . a The XPS spectra are also very similar to the spectrum of Ag $M_4N_{45}N_{45}$ of $\alpha\text{-Ag}_2WO_4$ published by Ho et al 55 . These results represent a valuable contribution to the XPS analysis of the $in\ situ$ atomic-scale nucleation of Ag filaments on $\alpha\text{-Ag}_2WO_4$.

Because the formation of metallic Ag after the electron irradiation of $\alpha\text{-}Ag_2WO_4$ is a quantum phenomenon, we performed quantum mechanical calculations to understand the structural and electronic modifications of $\alpha\text{-}Ag_2WO_4$ that were observed experimentally. In the calculations, electrons were introduced one by one up to ten in the orthorhombic unit cell of $\alpha\text{-}Ag_2WO_4$, and a redistribution of these extra electrons takes place by means of a simultaneously



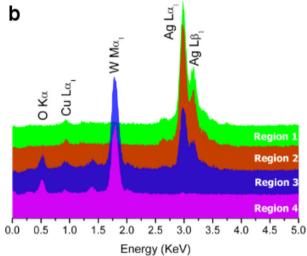


Figure 2 \mid (a) HAADF image of an isolated nanostructure. (b) EDS performed in different regions, which are illustrated in Fig. 2a. (Scale bar = 50 nm in a).

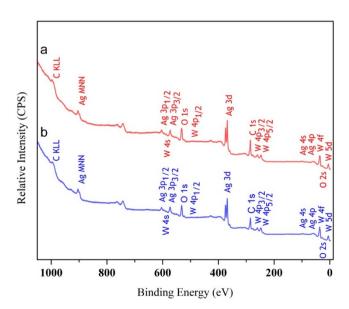


Figure 3 | XPS survey spectrum of the irradiated in (a), and non-irradiated a- Ag_2WO_4 material in (b).



geometry optimization on both the lattice parameters and the atomic positions. The change of lattice constants (a, b, and c) as a function of the number of added electrons are presented in Fig. S5 (SI). In Fig. 4a, the values of the bond distances of Ag-O and W-O in [AgO2], [AgO₄], and [WO₆] clusters are shown as a function of electrons added. An analysis and a comparison of the geometries for a neutral (N = 0) and charged (N = 10) structures show a pronounced increase in the corresponding Ag-O distances with the addition of electrons. In the [AgO₂] cluster, the Ag-O distance increases from 2.15 to 2.75 Å (in blue). In the [AgO₄] cluster, two different distances are observed: one pair (in red) exhibits behaviour similar to that of the bonds in the [AgO₂] cluster and has an O-Ag-O angle of approximately 170°; the other pair (in garnet) has an O-Ag-O angle of approximately 108° and a longer Ag-O distance, which indicates that the atoms detach as the electrons are added. In fact, when N=10, the first pair of O atoms forms an O-Ag-O angle of approximately 178°, whereas the angle of the second pair is reduced to approximately 90°, as shown in Fig. 4b. In the two types of [WO₆] clusters, we find that the W-O distances corresponding to the W2 and W3 atoms remain almost unaltered, whereas the distance of the W-O bond corresponding to W1 decreases smoothly with the addition of electrons. These results show that during electron irradiation, electronic and structural disorder was introduced into the material thus illustrating the fundamental role of cluster concepts in the formation and growth of Ag filaments.

QTAIM, which was developed by Bader and collaborators $^{59-61}$, allows the analysis of the experimental and theoretical electron density distributions, $\rho(r)$, in a molecule or solid as well as the study of the properties of $\rho(r)$; these analyses reveal the bonding interactions in a molecular or crystal system and the nature of these interactions. The electronic charge of each atom is evaluated using Bader charge analysis within the QTAIM framework, which is a way of dividing molecules or solids into atoms on the basis of electronic charge density. Finding zero flux surfaces between two atoms allows the charge of each atom to be calculated. In Fig. 5, the charge density of the Ag and W centres of the [AgO_2], [AgO_4], and [WO_6] clusters is depicted as a function of the number of electrons added. Atomic charges were calculated using integrations of the charge density

within the atomic basins, Ω , and subtracting the nuclear charge, Z, of the corresponding atom.

$$q(\Omega) = \mathbf{Z}_{\Omega} - \mathbf{N}(\Omega) \text{ with } N(\Omega) = \int_{\Omega} \rho(\Omega) dr \tag{1}$$

The average bond distances of Ag-Ag as a function of the number of electrons added are presented in Fig. 5a, and electron density contours on the (100) plane for the neutral (N = 0) structure and charged (N = 10) structures are depicted in Fig. 5b. Isodensity lines less than 0.02 a.u. are coloured white to highlight the differences between them. An analysis of the results presented in Fig. 5b reveals that the electron density distribution is enhanced between Ag4 and Ag5 at the same time that the Ag4–Ag5 contact distance is shortened when the number of added electrons is increased from N = 0 to N = 10. In addition, there is an electronic charge density enlargement in the vicinity of Ag6 atoms on going from N = 0 to N = 10.

An analysis of the results presented in Fig. 5c reveals that the Ag6 atoms of the [AgO₂] clusters are the atoms most prone to reduction. At N = 7, the Ag6 atoms are practically reduced, whereas the Ag4/ Ag5 centres require at least 10 electrons to reach the same state. This behaviour implies the existence of two different paths to obtaining metallic Ag, which are associated with the [AgO₂] and [AgO₄] clusters. In the case of W atoms that form [WO₆] clusters, W1 atoms behave differently than W2 and W3 atoms, and a minor decrease in electron density relative to that of Ag centres is observed (a decrease of 0.2 units at N = 10). Therefore, the extra electron density added to the material is transferred from one cluster to another through the lattice network, and the Ag-formation process involves both adjacent $[AgO_2]$ and $[AgO_4]$ clusters and, to a minor extent, $[WO_6]$ clusters. During electron irradiation, electronic and structural disorder is introduced into the material, indicating the fundamental role that cluster concepts play in the formation and growth of Ag filaments.

The results of the calculations of the Laplacian, $\nabla^2 \rho_{bcp}$, and the charge density, ρ_{bcp} , at the (3,-1) bond critical points (BCP) in Ag-O bonds in the $[AgO_2]$ and $[AgO_4]$ clusters are shown in Table 1. The effect of adding electrons to the material produces striking differences in the values of the Laplacian and the charge density at the (3,-1) BCP. Notably, the Laplacian and charge density values of the

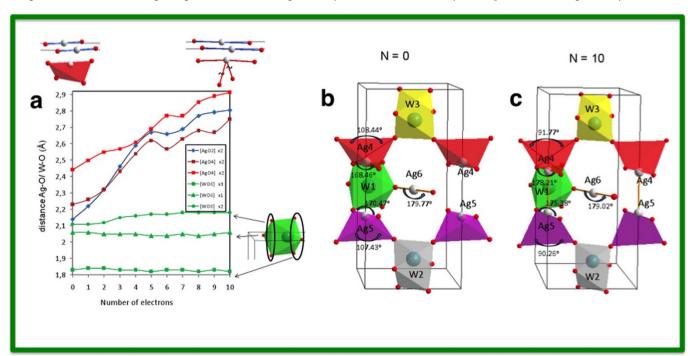


Figure 4 \mid (a) Values of Ag-O and W-O bond distances in the [AgO₂], [AgO₄], and [WO₆] clusters as a function of the number of electrons added. (b) Geometry of neutral (N = 0) and charged (N = 10) structures.



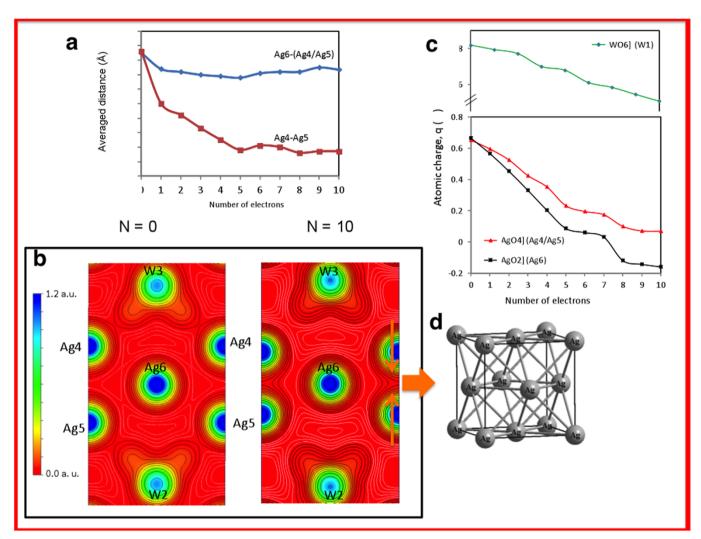


Figure 5 | (a) Average Ag-Ag distances as a function of the number of electrons added. (b) Electron density contours on the (100) plane for a neutral (N=0) structure and a charged (N=10) structure. Isodensity lines less than 0.02 a.u. are coloured white. (c) Charge density of the Ag and W centres in $[AgO_2]$, $[AgO_4]$, and $[WO_6]$ clusters as a function of the number of electrons added. $q(\Omega)$ represents the number of valence electrons minus the calculated charge density. (d) Structure of metallic Ag.

Ag-O bonds decrease considerably as electrons are added, which indicates that these bonds become weaker in favour of the formation of metallic Ag from both the $[{\rm AgO_2}]$ and $[{\rm AgO_4}]$ clusters.

Finally, we have calculated the total and projected density of states (DOS) on atoms and orbitals in order to complement the Bader charge analysis. For neutral Ag₂WO₄ (N = 0), there is essentially no visible difference in orbital character between PBE and PBE + U, however up to U = 6, the band gap values are maintained nearly constant to 2.0 eV which conveys the most satisfactory picture for both the occupied and unoccupied states. The DOS plots depicted in Fig. S6 (SI) are focused on the upper part of the valence band (VB) and the lower part of the conduction band (CB) for different Ag atoms as the number of electrons is added, for N = 0, 2, 4 and 8. For the DOS evaluation, from a total of 16 Ag atoms in the unit cell, three different types of Ag atoms have been classified according to previous results of the electron density distribution: "Ag1/2/3" (10 atoms), "Ag4/5" (4 atoms) and "Ag6" (2 atoms). An analysis of the plots shows a 50% of reduction of the band gap energy for N = 2, with a high contribution of s orbitals in the lower part of CB from Ag4/5 and Ag6. When the number of electrons is increased to 3 and up to 8 $\,$ the system becomes a conductor. For N = 4 and N = 8 the DOS images have been also depicted in Fig. S6, highlighting the same contribution around 0.0 eV for N = 4 and in the energy range -1.5-1.0 eV for N = 8. It is worth noting that the Fermi level is placed at energy of 0 eV. Therefore, a shift of the s band (and the d band as a minor contribution) of Ag4/5 and Ag6 atoms is possibly induced by the interaction of the sample with the electron beam, and

Table 1 | Laplacian and charge density at the (3,-1) BCPs in Ag-O bonds for [AgO₂] and [AgO₄] clusters as a function of the number of electrons added, N

	BCP [AgO ₂]			BCP [AgO₄]			
	Ag-O		Ag-O1		Ag-O2		
N	Рьср	$\nabla^2 \rho_{bcp}$	ρ_{bcp}	$\nabla^2 \rho_{bcp}$	ρ_{bcp}	$\nabla^2 \rho_{bcp}$	
0	0.49	6.80	0.26	3.29	0.42	4.91	
1	0.42	4.56	0.23	2.93	0.39	4.35	
2	0.34	3.88	0.21	2.67	0.35	3.86	
3	0.26	3.10	0.20	2.56	0.28	3.10	
4	0.20	2.26	0.19	2.35	0.22	2.56	
5	0.17	1.84	0.16	1.96	0.19	2.17	
6	0.17	1.76	0.15	1.83	0.19	2.08	
7	0.16	1.71	0.13	1.65	0.19	2.12	
8	0.14	1.55	0.12	1.39	0.17	1.80	
9	0.14	1.47	0.10	1.23	0.17	1.76	
10	0.13	1.40	0.10	1.14	0.16	1.51	



predicted by XPS analysis. Consequently, a structural rearrangement is caused related to the shortening of the distance Ag4–Ag5 and the Ag-Ag metallic bond formation when the number of added electrons is increased in accordance with the dynamics results measured *in situ* using TEM.

Discussion

Understanding the in situ atomic-scale nucleation induced by the electron irradiation of Ag filaments on α-Ag₂WO₄ can be achieved through a combination of experimental and theoretical analyses. We used experimental techniques (TEM, EDS, and XPS) and first-principle calculations based on the QTAIM framework to study the geometric and electronic structure of α-Ag₂WO₄. We then derived a mechanism on the basis of the electron irradiation of $[AgO_x]$ (x = 2, 4, 6, and 7) and [WO₆] clusters, which are the constituent polyhedra of α-Ag₂WO₄. The mechanism is relevant to the nucleation and growth of Ag filaments that occurs when α -Ag₂WO₄ is irradiated using an accelerated electron beam from an electron microscope under high vacuum. Our model allows us to better understand the atomic events in this process, and we are currently pursuing this relevant area of investigation. The results of the present work provide an atomic-scale picture of the formation of metallic Ag as well as an initial glimpse into how [WO₆] and [AgO_x] clusters behave in this material when it is irradiated with electrons from an external source. The strategy reported here for retrospective mapping of the growth of metallic Ag has the potential to yield valuable insights into the mechanistic aspects of the clusters, particularly when the experimental results are considered in conjunction with first-principles calculations.

Our main conclusions are as follows: During the electron irradiation process, electronic and structural disorder is introduced into the material showing the fundamental role of clusters in the nucleation and growth of Ag filaments. Extra electrons added to the material are transferred from one cluster to another through the lattice network, and Ag formation occurs via the reduction of both [AgO₂] and [AgO₄] clusters that are adjacent and, to a lesser extent, through the reduction of the [WO₆] cluster. These findings provide a valuable probe into the relationship between atomic-scale structural and electronic perturbations and their macroscopic consequences. The broader significance of this work lies in the direct and visually impressive demonstration of the conversion of electron irradiation into structural and electronic order/disorder effects to cause the formation of Ag on α-Ag₂WO₄. Therefore, the Ag nucleation mechanisms observed here can provide deep insight into the physical and electronic structure of other silver-metal-oxide-based materials under electron irradiation conditions. The present work leads to a unified understanding of the complex behaviour of these materials and opens a new direction for the rational design of materials with exciting properties.

Methods

Experimental details. The $\alpha\text{-}Ag_2WO_4$ samples were synthesised at $90^\circ C$ in 1 min by the injection of precursor ions into hot aqueous solutions according to the method reported in our previous work. TEM analysis was performed on a CM200-Philips and JEOL JEM 2100F TEM/STEM microscope operated at 200 kV. Specimens for TEM images were obtained by drying droplets of as-prepared samples from an acetone dispersion that had been sonicated for 10 min and deposited on 300-mesh Cu grids. The chemical analyses of the samples were performed by EDS using a Thermo-Noran EDS attached to the JEM 2100F and equipped with a Si detector.

XPS spectra were collected using a commercial spectrometer (UNI-SPECS-UHV). To analyse the near-surface composition and the chemical environment in the α - Ag_2WO_4 nanocrystals, the measurements were performed in a small area of each sample using the Mg K α line (hv = 1253.6 eV); the analyser pass energy was set to 10 eV. In particular, the inelastic background of the C 1 s, O 1 s, W 4f, and Ag 3 d electron core-level spectra and the Auger Ag MNN peak were subtracted using Shirley's method 62 ; the BEs were corrected using the hydrocarbon component of adventitious carbon fixed at 285.0 eV.

The X-ray diffraction measurements were performed on a D/Max-2500PC diffractometer (Rigaku, Japan) using Cu K α radiation ($\lambda=1.5406$ Å) in the 20 range

from 10° to 70° in the normal routine with a scanning velocity of 2° /min and a step of 0.02° .

Calculation details. First-principles total-energy calculations were performed within the periodic DFT framework using the VASP program^{63,64}. The Kohn-Sham equations were solved by means of the Perdew, Burke, and Ernzerhof exchangecorrelation functional⁶⁴, and the electron-ion interaction was described by the projector-augmented-wave pseudopotentials^{65,66}. Because of the well-known limitations of standard DFT in describing the electronic structure of "stronglycorrelated" compounds, a correction to the PBE wavefunctions was made (PBE + U) by including a repulsive on-site Coulomb interaction, U, according to the formula of Dudarev et al⁶⁷. PBE + U scheme contains the same PBE approximate correlation, but takes into account orbital dependence (applied to the d states of silver) of the Coulomb and exchange interactions. The value of the Hubbard parameter was tested, and a value of U = 6 eV for the Ag element was used. The plane-wave expansion was truncated at a cut-off energy of 400 eV, and the Brillouin zones were sampled through Monkhorst-Pack special k-points grids (6 \times 6 \times 6) that ensure geometrical and energetic convergence for the Ag₂WO₄ structures considered in this work. The keyword NELECT was used to increase the number of electrons in the bulk structure, and all the crystal structures were optimised simultaneously on both the volume of the unit-cell and the atomic positions. Further information, as well as details of the ab initio quantum chemical calculations, is given in the Supplementary Information.

- Heyer, O. et al. A new multiferroic material: MnWO₄. J. Phys.-Condens. Matter 18, L471–L475 (2006).
- Zhang, L., Lu, C., Wang, Y. & Cheng, Y. Hydrothermal synthesis and characterization of MnWO₄ nanoplates and their ionic conductivity. *Mater. Chem. Phys.* 103, 433–436 (2007).
- Song, X. C. et al. Hydrothermal preparation and photoluminescence of bundlelike structure of ZnWO₄ nanorods. Appl. Phys. A 94, 185–188 (2009).
- Xing, Y. et al. Microemulsion-mediated solvothermal synthesis and photoluminescent property of 3D flowerlike MnWO₄ micro/nanocomposite structure. Solid State Sci. 10, 1299–1304 (2008).
- Cavalcante, L. S. et al. Cluster Coordination and Photoluminescence Properties of alpha-Ag₂WO₄ Microcrystals. *Inorg. Chem.* 51, 10675–10687 (2012).
- Longo, E. et al. Direct in situ observation of the electron-driven synthesis of Ag filaments on alpha-Ag₂WO₄ crystals. Sci. Rep. 3, 1676 (2013).
- Longo, E. et al. Toward an Understanding of the Growth of Ag Filaments on α-Ag2WO4 and their Photoluminescent Properties: A Combined Experimental and Theoretical Study. J. Phys. Chem. C doi:10.1021/jp408167v (2014).
- Ziman, J. M. Models of disorder: the theoretical physics of homogeneously disordered systems (Cambridge University Press, Cambridge Eng.; New York, 1979).
- Cui, X. J. et al. Selective synthesis and characterization of single-crystal silver molybdate/tungstate nanowires by a hydrothermal process. Chem.-a Eur. J. 10, 218–223 (2004).
- Skarstad, P. M. & Geller, S. (W₄O₁₆)⁸⁻ Polyion in High-Temperature Modification of Silver Tungstate. *Mater. Res. Bull.* 10, 791–799 (1975).
- 11. Stone, D. *et al.* Layered atomic structures of double oxides for low shear strength at high temperatures. *Scripta Mater.* **62**, 735–738 (2010).
- Zheng, H. M. et al. Observation of Single Colloidal Platinum Nanocrystal Growth Trajectories. Science 324, 1309–1312 (2009).
- Chai, J., Liao, X., Giam, L. R. & Mirkin, C. A. Nanoreactors for Studying Single Nanoparticle Coarsening. J. Am. Chem. Soc. 134, 158–161 (2012).
- Evans, J. E., Jungjohann, K. L., Browning, N. D. & Arslan, I. Controlled Growth of Nanoparticles from Solution with In Situ Liquid Transmission Electron Microscopy. *Nano Lett.* 11, 2809–2813 (2011).
- Liao, H. G., Cui, L. K., Whitelam, S. & Zheng, H. M. Real-Time Imaging of Pt₃Fe Nanorod Growth in Solution. Science 336, 1011–1014 (2012).
- Papageorgiou, A. C., Pang, C. L., Chen, Q. & Thornton, G. Low-dimensional, reduced phases of ultrathin TiO₂. ACS Nano 1, 409–414 (2007).
- 17. Xin, H. L. L. & Zheng, H. M. In Situ Observation of Oscillatory Growth of Bismuth Nanoparticles. *Nano Lett.* **12**, 1470–1474 (2012).
- 18. Yuk, J. M. *et al.* High-Resolution EM of Colloidal Nanocrystal Growth Using Graphene Liquid Cells. *Science* **336**, 61–64 (2012).
- Kraus, T. & de Jonge, N. Dendritic Gold Nanowire Growth Observed in Liquid with Transmission Electron Microscopy. *Langmuir* 29, 8427–8432 (2013).
- Sun, M., Liao, H.-G., Niu, K. & Zheng, H. Structural and Morphological Evolution of Lead Dendrites during Electrochemical Migration. Sci. Rep. 3, 3227 (2013).
- Yuk, J. M. et al. In situ atomic imaging of coalescence of Au nanoparticles on graphene: rotation and grain boundary migration. Chem. Commun. 49, 11479–11481 (2013).
- Hamann, T., Bohler, E. & Swiderek, P. Low-Energy-Electron-Induced Hydroamination of an Alkene. Angew. Chem.-Int. Edit. 48, 4643–4645 (2009).
- Ramsier, R. D. & Yates, J. T. Electron-Stimulated Desorption Principles and Applications. Surf. Sci. Rep. 12, 243–378 (1991).
- Turchanin, A. et al. One Nanometer Thin Carbon Nanosheets with Tunable Conductivity and Stiffness. Adv. Mater. 21, 1233-+ (2009).
- Xu, W. et al. In-situ atomic-scale observation of irradiation-induced void formation. Nat. Commun. 4, 2288 (2013).



- Randolph, S. J., Fowlkes, J. D. & Rack, P. D. Focused, nanoscale electron-beaminduced deposition and etching. Crit. Rev. Solid State Mat. Sci. 31, 55–89 (2006).
- 27. Utke, I., Hoffmann, P. & Melngailis, J. Gas-assisted focused electron beam and ion beam processing and fabrication. *J. Vac. Sci. Technol. B* **26**, 1197–1276 (2008).
- van Dorp, W. F. & Hagen, C. W. A critical literature review of focused electron beam induced deposition. J. Appl. Phys. 104, 081301 (2008).
- van Dorp, W. F., Hagen, C. W., Crozier, P. A. & Kruit, P. Growth behavior near the ultimate resolution of nanometer-scale focused electron beam-induced deposition. *Nanotechnology* 19, 225305 (2008).
- van Dorp, W. F., van Someren, B., Hagen, C. W. & Kruit, P. Approaching the resolution limit of nanometer-scale electron beam-induced deposition. *Nano Lett.* 5, 1303–1307 (2005).
- van Kouwen, L., Botman, A. & Hagen, C. W. Focused Electron-Beam-Induced Deposition of 3 nm Dots in a Scanning Electron Microscope. *Nano Lett.* 9, 2149–2152 (2009).
- Walz, M. M. et al. Electrons as "Invisible Ink": Fabrication of Nanostructures by Local Electron Beam Induced Activation of SiOx. Angew. Chem.-Int. Edit. 49, 4669–4673 (2010).
- Yagi, K., Takayanagi, K., Kobayashi, K. & Honjo, G. Insitu Observations of Growth Processes of Multiply Twinned Particles. J. Cryst. Growth 28, 117–124 (1975).
- de Jonge, N. & Ross, F. M. Electron microscopy of specimens in liquid. Nat. Nanotechnol. 6, 695–704 (2011).
- Kim, J. U., Cha, S. H., Shin, K., Jho, J. Y. & Lee, J. C. Synthesis of gold nanoparticles from gold(I)-alkanethiolate complexes with supramolecular structures through electron beam irradiation in TEM. J. Am. Chem. Soc. 127, 9962–9963 (2005).
- Sohn, J. H. et al. Preparation of conducting silver paste with Ag nanoparticles prepared by e-beam irradiation. Rad. Phys. Chem. 79, 1149–1153 (2010).
- de Jonge, N., Peckys, D. B., Kremers, G. J. & Piston, D. W. Electron microscopy of whole cells in liquid with nanometer resolution. *Proc. Natl. Acad. Sci. U. S. A.* 106, 2159–2164 (2009).
- Klein, K. L., Anderson, I. M. & De Jonge, N. Transmission electron microscopy with a liquid flow cell. J. Microsc. 242, 117–123 (2011).
- 39. Tao, F. & Salmeron, M. In Situ Studies of Chemistry and Structure of Materials in Reactive Environments. *Science* **331**, 171–174 (2011).
- Williamson, M. J., Tromp, R. M., Vereecken, P. M., Hull, R. & Ross, F. M. Dynamic microscopy of nanoscale cluster growth at the solid-liquid interface. *Nat. Mater.* 2, 532–536 (2003).
- 41. Altman, M. S. CO Prefers the Aisle Seat. Science 327, 789-790 (2010).
- 42. de Smit, E. *et al.* Nanoscale chemical imaging of a working catalyst by scanning transmission X-ray microscopy. *Nature* **456**, 222–U239 (2008).
- Friebel, D., Miller, D. J., Nordlund, D., Ogasawara, H. & Nilsson, A. Degradation of Bimetallic Model Electrocatalysts: An In Situ X-Ray Absorption Spectroscopy Study. Angew. Chem.-Int. Edit. 50, 10190-10192 (2011).
- Hansen, P. L. et al. Atom-resolved imaging of dynamic shape changes in supported copper nanocrystals. Science 295, 2053–2055 (2002).
- Ketteler, G. et al. In situ spectroscopic study of the oxidation and reduction of Pd(111). J. Am. Chem. Soc. 127, 18269–18273 (2005).
- 46. Nguyen, L., Cheng, F., Zhang, S. R. & Tao, F. Visualization of Surfaces of Pt and Ni Model Catalysts in Reactive Environments Using Ambient Pressure High Temperature Scanning Tunneling Microscopy and Understanding the Restructurings of Surfaces of Model Metal Catalysts under Reaction Conditions at Near Ambient Pressure. J. Phys. Chem. C 117, 971–977 (2013).
- Park, J. et al. Direct Observation of Nanoparticle Superlattice Formation by Using Liquid Cell Transmission Electron Microscopy. ACS Nano 6, 2078–2085 (2012).
- 48. Rodriguez, J. A. *et al.* Activity of CeOx and TiOx nanoparticles grown on Au(111) in the water-gas shift reaction. *Science* **318**, 1757–1760 (2007).
- Tao, F. et al. Break-Up of Stepped Platinum Catalyst Surfaces by High CO Coverage. Science 327, 850–853 (2010).
- Tao, F., Tang, D., Salmeron, M. & Somorjai, G. A. A new scanning tunneling microscope reactor used for high-pressure and high-temperature catalysis studies. Rev. Sci. Instrum. 79, 084101 (2008).
- 51. Thostrup, P. et al. Adsorption-induced step formation. Phys. Rev. Lett. 87, 126102 (2001).
- Vestergaard, E. K. et al. Adsorbate-induced alloy phase separation: A direct view by high-pressure scanning tunneling microscopy. Phys. Rev. Lett. 95, 126101 (2005).
- Wachs, I. E. & Roberts, C. A. Monitoring surface metal oxide catalytic active sites with Raman spectroscopy. *Chem. Soc. Rev.* 39, 5002–5017 (2010).
- 54. Wagner, C. D. W. W., Riggs, W. W., Davis, L. E., Moulder, J. F. & Muilenberg, G. E. Handbook of X-ray Photoeclectron Spectroscopy (Physical Electronics Division, Perkin–Elmer Corporation, Wellesley, MA, 1979).

- 55. Ho, S. F., Contarini, S. & Rabalais, J. W. Ion-Beam-Induced Chemical-Changes in the Oxyanions (CrOYN-, MoOYN-, WOYN-, VOYN-, NbOYN-, and TaOYN-) and Oxides (CrOX, MoOX, WOX, VOX, NbOX, and TaOX). J. Phys. Chem. 91, 4779–4788 (1987).
- 56. Wang, D., Xue, G., Zhen, Y., Fu, F. & Li, D. Monodispersed Ag nanoparticles loaded on the surface of spherical Bi₂WO₆ nanoarchitectures with enhanced photocatalytic activities. *J. Mater. Chem.* 22, 4751–4758 (2012).
- Zhang, L. S. et al. AgBr-Ag-Bi₂WO₆ nanojunction system: A novel and efficient photocatalyst with double visible-light active components. Appl. Catal. A 363, 221–229 (2009).
- 58. Crist, V. Handbook of Monochromatic XPS spectra. The Elements and Native Oxides. (John Wiley & Sons Ltd., New York 2000).
- Bader, R. F. W. Atoms in molecules: a quantum theory (Clarendon Press Oxford New York, 1990).
- Matta, C. F. & Boyd, R. J. The quantum theory of atoms in molecules (Wiley-VCH, Weinheim, 2007).
- Popelier, P. L. A. Atoms in molecules: an introduction (Prentice Hall, Harlow, 2000)
- 62. Shirley, D. A. High-Resolution X-Ray Photoemission Spectrum of Valence Bands of Gold. *Phys. Rev. B* **5**, 4709–& (1972).
- Kresse, G. & Hafner, J. ab-initio Molecular-Dynamics Simulation Of The Liquid-Metal Amorphous-Semiconductor Transition In Germanium. *Phys. Rev. B* 49, 14251–14269 (1994).
- 64. Kresse, G. & Furthmuller, J. Efficiency of ab-initio total energy calculations for metals and semiconductors using a plane-wave basis set. *Comput. Mater. Sci.* 6, 15–50 (1996).
- 65. Perdew, J. P., Burke, K. & Ernzerhof, M. Generalized gradient approximation made simple. *Phys. Rev. Lett.* 77, 3865–3868 (1996).
- Kresse, G. & Joubert, D. From ultrasoft pseudopotentials to the projector augmented-wave method. *Phys. Rev. B* 59, 1758–1775 (1999).
- 67. Dudarev, S. L., Botton, G. A., Savrasov, S. Y., Humphreys, C. J. & Sutton, A. P. Electron-energy-loss spectra and the structural stability of nickel oxide: An LSDA + U study. *Phys. Rev. B* 57, 1505–1509 (1998).

Acknowledgments

The authors are grateful to FAPESP 2013/07296-2, CAPES, CNPq, Prometeo/2009/053 (Generalitat Valenciana), CTQ2012-36253-C03-02 (Ministerio de Economia y Competitividad, Spain), CTQ2012-36253-C03-02, and the Spanish Brazilian program (PHB2009-0065-PC). Special thanks are given to Dr. P. Hammer (LEFE-IQ/UNESP) for help with the XPS analyses. STEM analysis were performed in the Electron Microscopy Laboratory (LME) of the Brazilian Nanotechnology National Laboratory (LNNano).

Author contributions

L.G., P.G.-N. and V.M.L. have been carried the calculations and the theoretical part. M.F. and P.S.L. prepared the samples and performed the structural analysis; D.P.V. and W.A. performed the electron microscopy data collection. F.A.L.P. has been carried out the XPS analysis. J.A., A.C.H. and E.L. conceived the project. All of the authors participated in writting the manuscript and discussion of the results.

Additional information

Supplementary information accompanies this paper at http://www.nature.com/scientificreports

Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Andrés, J. et al. Structural and electronic analysis of the atomic scale nucleation of Ag on α -Ag₂WO₄ induced by electron irradiation. Sci. Rep. 4, 5391; DOI:10.1038/srep05391 (2014).



This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivs 4.0 International License. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder in order to reproduce the material. To view a copy of this license, visit http://creativecommons.org/licenses/by-nc-nd/4.0/