Surface Morphology and Magnetic Properties of Isolated Cylindrical Nickel Nanowires

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Abstract: In this article, the surface morphology and magnetic properties of isolated cylindrical nickel (Ni) nanowires fabricated by electrodeposition have been thoroughly investigated using scanning electron microscopy and Magneto-Optical Kerr Effect (MOKE) magnetometry, respectively. The surfaces of most nanowires were found to be homogenous, uniform, and cylindrical in shape. Some others show different diameters and surface features, including; protrusions and branches along their length. The diameter distribution of a wide range of nanowires was found to differ from their template pore diameters. These all variations are more likely due to defects exist in the internal surfaces of the pores within the template itself, or may be associated with the trapped air pockets within the pores during nanowires growth or due to the oxide formation or residual contaminants which may cover these wires. The nanowires lengths were found to differ from their actual lengths estimated during deposition growth. This was attributed to the breakage of nanowires into small sections during releasing process. The hysteresis loops obtained by applying a magnetic field at different angles with respect to the nanowires long axis showed square hysteresis loops with a sharp jump of Kerr signal during switching behaviour, as well as a high squareness ratio, indicating the dominance of shape anisotropy. These results are quite different from the measurements of high density templated nanowires reported in the literature, due to the small number of nearest neighbour nanowires, and hence no magneto-static interaction. The magnetisation reversal of such wires is well described by the non-uniform rotation of the curling model of domain reversal.

Keywords: Ferromagnetic Nanowires, Surface Morphology, Magnetic Properties, Magneto-Optical Kerr Effect (MOKE) Magnetometry, Magnetisation Reversal, Curling Model

1. Introduction

Cylindrical ferromagnetic nanowires are structures that have diameters in nanometer scale and unrestricted in length. Typical cylindrical nanowires exhibit high aspect ratios (length to diameter), and they can often be approximated as one dimensional materials [28-51]. These nanowires have many interesting properties that are not seen in a bulk or three dimensional structures. Therefore, fabrication and characterisation of such structures present interest from both fundamental and industrial points of view [14, 16, 28, 29, 37, 44, 51, 58]. As an example; they may play a significant role in future applications, in particular; ultra-high density magnetic storage devices in hard disc drives and volatile magneto resistive random access memory (MRAM) [28-29].

Different techniques have been utilized to fabricate these structures, but electrodeposition has been proved to be simple, fast and a low cost technique, as well as using this technique, it is possible to produce different materials and multilayer as: thin films, nanowires or nanotubes [5, 8, 11, 12, 14, 18, 31].

The magnetic properties of as-deposited two dimensional arrays of ferromagnetic nanowires have been investigated intensively over the last few decades using a wide range of characterising techniques [13-38]. These researches, however, have produced a significant insight into the physical behaviour of such arrays, where the magneto-static or the dipolar interactions among the wires have been found
Nevertheless, few articles have been published throughout potentially interesting, as it provides a very sensitive probe polarised laser light following its reflection from a magnetic have been widely used to study the magnetisation behaviour of packed two Ni nanowires with diameters of ~300 nm and this work discussing the magnetisation behaviour of Ni based substrates. Therefore, researchers were trying to investigate these properties by releasing such nanowires from their templates, and characterise them using MOKE magnetometry. Nevertheless, this process was accomplished by dividing the templates into small parts with an approximate area of 1×1 mm², then one or two pieces were placed in a small beaker containing two molar solutions of sodium hydroxide (NaOH), to dissolve the alumina membrane. The template was left in the solvent for a time of about 72 hours, by which the alumina template had dissolved and removed. With care, the NaOH solvent was drained out from the container and distilled water was added to wash the nanowires and remove the solvent. In order to be sure that NaOH solution was entirely removed from the beaker, this process was repeated for 4-5 times. However, care should be taken during this process that the released wires may fall out. Now, to remove the distilled water from the container, a small amount of isopropanol alcohol was added. To release the nanowires from their gold base substrate, the beaker was placed in an ultrasonic bath for a time of about 10 minutes. For further discussion on the template dissolving, nanowires releasing and deposition on silicon substrates can be found in reference [48]. Once the nanowires were released, a one drop was spilled onto silicon chips with a special micromarkers pre-fabricated by electron beam lithography. These markers were designed to be large enough to be seen in the microscope of MOKE system. After drying the solution, the wires are strongly stuck on the silicon chip surface. The position of the nanowires relative to the gold patterns was then carried out using scanning electron microscopy. The dissolution of the membrane and cleaning the wires before subsequent process is not an easy task. The objective is to achieve clean nanowires spread over the chips and sufficiently isolated from each other, to allow the magnetic measurements to be performed on an isolated nanowire. Nevertheless, this process was accomplished by dividing the templates into small parts with an approximate area of 1×1 mm², then one or two pieces were placed in a small beaker containing two molar solutions of sodium hydroxide (NaOH), to dissolve the alumina membrane. The template was left in the solvent for a time of about 72 hours, by which the alumina template had dissolved and removed. With care, the NaOH solvent was drained out from the container and distilled water was added to wash the nanowires and remove the solvent. In order to be sure that NaOH solution was entirely removed from the beaker, this process was repeated for 4-5 times. However, care should be taken during this process that the released wires may fall out. Now, to remove the distilled water from the container, a small amount of isopropanol alcohol was added. To release the nanowires from their gold base substrate, the beaker was placed in an ultrasonic bath for a time of about 10 minutes. For further discussion on the template dissolving, nanowires releasing and deposition on silicon substrates can be found in reference [48]. Once the nanowires were released, a one drop was spilled onto silicon chips with a special micromarkers pre-fabricated by electron beam lithography. These markers were designed to be large enough to be seen in the microscope of MOKE system. After drying the solution, the wires are strongly stuck on the silicon chip surface. The position of the nanowires relative to the gold patterns was then carried out using scanning electron microscopy. Imaging of nanowires and fabrication of gold markers on silicon substrates were performed using a high-resolution field emission scanning electron microscopy column on an FEI-Helios Nanolab dual beam FIB/SEM system with the electron beam energy of ~10 and ~30 KeV, respectively. More details on the scanning electron microscopy, electron to dominate over the magnetic properties of an individual nanowire.

The intrinsic magnetic state of an individual or small number of closely packed ferromagnetic nanowires is a significant role for scientific point of view and demanding issue of industrial applications. Therefore, researchers were trying to investigate these properties by releasing such nanowires from their templates, and characterise them using MOKE magnetometry. Nevertheless, this process was accomplished by dividing the templates into small parts with an approximate area of 1×1 mm², then one or two pieces were placed in a small beaker containing two molar solutions of sodium hydroxide (NaOH), to dissolve the alumina membrane. The template was left in the solvent for a time of about 72 hours, by which the alumina template had dissolved and removed. With care, the NaOH solvent was drained out from the container and distilled water was added to wash the nanowires and remove the solvent. In order to be sure that NaOH solution was entirely removed from the beaker, this process was repeated for 4-5 times. However, care should be taken during this process that the released wires may fall out. Now, to remove the distilled water from the container, a small amount of isopropanol alcohol was added. To release the nanowires from their gold base substrate, the beaker was placed in an ultrasonic bath for a time of about 10 minutes. For further discussion on the template dissolving, nanowires releasing and deposition on silicon substrates can be found in reference [48]. Once the nanowires were released, a one drop was spilled onto silicon chips with a special micromarkers pre-fabricated by electron beam lithography. These markers were designed to be large enough to be seen in the microscope of MOKE system. After drying the solution, the wires are strongly stuck on the silicon chip surface. The position of the nanowires relative to the gold patterns was then carried out using scanning electron microscopy. Imaging of nanowires and fabrication of gold markers on silicon substrates were performed using a high-resolution field emission scanning electron microscopy column on an FEI-Helios Nanolab dual beam FIB/SEM system with the electron beam energy of ~10 and ~30 KeV, respectively. More details on the scanning electron microscopy, electron 2. Materials and Methods

Two dimensional arrays of cylindrical ferromagnetic Ni nanowires were fabricated using electrodeposition technique. The fabrication process was carried out using an Autolab-30 potentiostat and a conventional three electrode cell with 20 cm² capacity. The electrolyte solution used was 0.57 M of NiSO₄ and 0.32 M of H₂BO₃. The pH of the solution was maintained to approximately 3.5-4. An alumina template of thickness ~60 µm with a nominal pore diameter of ~300 nm was used as a working electrode (cathode), after depositing a 100 nm pure gold layer (99.99%),+ on one side using thermal evaporation technique. The gold coated membranes are then used in the cell keeping the bare side of the template in front of the counter electrode. The voltage applied between the counter and reference electrodes was around–0.85 V according to linear voltametry results. Once the electrodeposition process was completed, the contact wires were removed and the template washed in distilled water for the next step. The deposition time and the current density were chosen to produce nanowires with lengths of around 8 µm. More details on the fabrication process using electrodeposition can be found elsewhere [9, 27, 30, 35, 39].

The dissolution of the membrane and cleaning the wires before subsequent process is not an easy task. The objective is to achieve clean nanowires spread over the chips and sufficiently isolated from each other, to allow the magnetic measurements to be performed on an isolated nanowire. Nevertheless, this process was accomplished by dividing the templates into small parts with an approximate area of 1×1 mm², then one or two pieces were placed in a small beaker containing two molar solutions of sodium hydroxide (NaOH), to dissolve the alumina membrane. The template was left in the solvent for a time of about 72 hours, by which the alumina template had dissolved and removed. With care, the NaOH solvent was drained out from the container and distilled water was added to wash the nanowires and remove the solvent. In order to be sure that NaOH solution was entirely removed from the beaker, this process was repeated for 4-5 times. However, care should be taken during this process that the released wires may fall out. Now, to remove the distilled water from the container, a small amount of isopropanol alcohol was added. To release the nanowires from their gold base substrate, the beaker was placed in an ultrasonic bath for a time of about 10 minutes. For further discussion on the template dissolving, nanowires releasing and deposition on silicon substrates can be found in reference [48]. Once the nanowires were released, a one drop was spilled onto silicon chips with a special micromarkers pre-fabricated by electron beam lithography. These markers were designed to be large enough to be seen in the microscope of MOKE system. After drying the solution, the wires are strongly stuck on the silicon chip surface. The position of the nanowires relative to the gold patterns was then carried out using scanning electron microscopy. Imaging of nanowires and fabrication of gold markers on silicon substrates were performed using a high-resolution field emission scanning electron microscopy column on an FEI-Helios Nanolab dual beam FIB/SEM system with the electron beam energy of ~10 and ~30 KeV, respectively. More details on the scanning electron microscopy, electron...
beam lithography and lift-off techniques are available in Refs [8, 22, 23, 56].

The magnetic analysis of isolated Ni nanowires was achieved using a highly sensitive MOKE magnetometry in the longitudinal configuration. In this system, the laser spot was focused to a diameter of approximately 5 µm. The change in the polarisation angle of the laser light after reflection from the sample was proportional to the longitudinal component of the wire magnetisation [5, 7, 32, 50]. The reflected light passed through an analysing prism and the intensity detected using a silicon photodiode connected to a high-bandwidth amplifier and a ~1 GHz-bandwidth oscilloscope. An AC electromagnet with a maximum field of ±450 Oe was used to switch the magnetisation of wires at a frequency of approximately 21 Hz. To provide information about the angular dependence of the switching fields, the experimental setup allows for a rotation of the substrate around its surface normal by an angle up to ~180 degrees, in order to apply a magnetic field in the plane of the substrate at different angles with respect to the nanowires long axis. Thus, this setup was used for hysteresis loop measurements of a collection of closely packed two Ni nanowires. More information about MOKE setup can be found in Refs [5, 7, 32, 50].

3. Results

Figure 1 show high resolution scanning electron microscopy micrograph examples of single cylindrical Ni nanowires with different diameters, lengths and surface morphologies lying on the gold premarked silicon substrates following release and deposition from the same dilute suspension. Figure 2 shows the distribution histograms of (a) diameters, and (b) lengths of such dispersed individual Ni nanowires. The insets of figure 2 are scanning electron microscopy micrograph examples of the measured nanowires.

Figure 1. High resolution scanning electron microscopy micrograph examples of cylindrical Ni nanowires with different diameters, lengths and surface morphologies lying on the gold premarked silicon substrates following releasing and deposition from same dilute suspension.
Figure 2. Distribution histograms of: (a) Diameters, and (b) Lengths of cylindrical Ni nanowires dispersed on silicon chips with nominal diameter and length of ~300 nm and ~8 µm, respectively. The figure insets are scanning electron micrograph examples of the measured nanowires.

During the course of MOKE measurements, it was difficult to get significant Kerr signals from such individual nanowires. Therefore, a collection of closely packed two cylindrical Ni nanowires with diameters of ~300 nm and ~8 µm each was chosen for the magnetic measurements. Figure 3 shows scanning electron microscopy micrograph of the measured Ni nanowires. Even, with this closely packed nanowire, it was also a big challenge to detect a reasonable amount of Kerr signal, due to their small sizes and their curved surfaces which scatter the laser light in different directions.

Figure 3. Scanning electron microscopy micrograph showing the collection of a closely packed two cylindrical Ni nanowires with ~300 nm diameter and ~8 µm lengths each lying on the pre-marked silicon substrates.

Figure 4 show examples of the corresponding hysteresis loops obtained from this closely packed nanowires when the magnetic field was applied in four different angles (0°, 15°, 30°, and 60°) with respect to the nanowires long axis. To get the coercivity distributions, the measurements were repeated in multiple locations on the same wires and for all the angles investigated here.

In all MOKE measurements, the signal to noise ratio of the hysteresis loops was found to be low. This is expected since the nanowires are very small compared to the laser spot size and the scattering effect. Importantly, the shapes of the hysteresis loops are quite different from those of two dimensional arrays of nanowires reported in the literature [6, 37, 44, 51]. There is also, a change in the shape of the loop and a reduction in the coercivity as the external magnetic field is changed from parallel to the perpendicular direction.

The effect of changing the angle of applied magnetic field with respect to the nanowires long axis on the squareness ratio (the ratio of the remanent magnetisation to the saturation magnetisation) and the coercivity is plotted in Figure 5. The bars shown in Figure 5(b) are the distribution of the coercivity obtained from repeating measurements on the same wires at a given angle. High squareness ratios (about 0.86) were found in all the hysteresis loops. A reduction in the coercivity with increasing the nanowires angle with respect to the magnetic field applied. The average coercivity was found to be ~110 Oe and ~75 Oe when the magnetic field was applied at 0° and 60° to the nanowires long axis, respectively.
Figure 4. Hysteresis loop examples obtained from a collection of closely packed two Ni nanowires when the magnetic field was applied at four different angles: (a-a) 0°, (b-b) 15°, (c-c) 30°, and (d-d) 60° with respect to the nanowires long axis.
4. Discussion

The surface morphology of a huge amount of individual cylindrical Ni nanowires spread over the silicon substrates was investigated using a high resolution scanning electron microscopy micrographs. The results demonstrate that these wires are smooth, uniform and cylindrical in shape. Some other wires showed a different diameter along their length [17, 40]. Whilst others showed surface irregularities, including; protrusions and branches, as demonstrated in the micrograph examples shown in Figure 1. This variation in the surface morphology is more likely reflects the variation or defects exist in the internal surfaces of the pores within the template itself, or in some cases, may be associated with the trapped air pockets within the pores during nanowires growth [26, 40]. Considerably, these defects may lead to changes in the actual magnetic properties of templated arrays of ferromagnetic nanowires. This is because, the surface roughness distort the spin-structure of ferromagnetic nanowires prior to the switching and it could lead to an increase or decrease in the switching field.

The high resolution scanning electron microscopy analysis, showed a wide range of diameters and lengths of nanowires dispersed on the silicon substrates. The distribution histograms were performed and are plotted in Figure 2. Clearly, a wide range of diameters (100-650 nm) and lengths (1-5.5 µm) were observed. The nanowires diameter was found to differ from their nominal template pore diameter (~300 nm), and this may be due to the oxide formation or residual templates and contaminants which may cover these wires or due to the variation in pore diameter themselves. Again, the variation in the nanowire diameters would make it difficult to entirely describe the magnetisation behaviour of two dimensional arrays of ferromagnetic nanowires.

The distribution of nanowires length was found to range from ~0.5 µm to ~5.5 µm which is far from their definite length estimated during deposition growth (~8 µm). This variation indicates that the sonication process used to separate nanowires from their base gold substrate causes the nanowires to break in different places. Hence, produce the wider distribution observed. Further analysis on the releasing process of ferromagnetic nanowires can be found in other published work [48, 50].

Despite of the difficulties arising from locating the nanowires within the MOKE illumination, the very small size of these wires with respect to laser spot size, and the rounded surfaces of these structures which leads to the scattering of reflected light into various directions, MOKE setup has been uninterruptedly applied for the measurements of hysteresis loops of a closely packed two Ni nanowires, so that it is not influenced by other nanowires at nearest neighbour. In templated nanowires, the interaction with other nanowires complicates the situation and makes it more complex to understand the exact magnetisation behaviour of a single or small number of closely packed nanowires.

The switching behaviour was found to be sharp in all angles of measurements in comparison with the hysteresis loops noticed in literature for two dimensional arrays of such Ni nanowires using other investigative techniques, including; vibrating sample magnetometer (VSM), superconducting quantum interference devices (SQUID) and alternating gradient magnetometers (AGM).

It is well known that the magnetic properties of an individual nanowire are governed by several anisotropies [15, 25], including; shape, magneto-crystalline, stress and surface anisotropies. The first anisotropy originates from the self-demagnetising field due to it is cylindrical shape. The closely packed two Ni nanowires investigated here have the aspect ratio of about 27. The demagnetisation field, along the perpendicular direction to the nanowires long axis of such an aspect ratio can be approximated to be $2\pi M_s = 4346 \text{ Oe}$ [15], where $M_s$ is the saturation magnetisation of Ni bulk. This makes the long axial direction of the nanowires as the easy axis and its perpendicular direction as a hard axis of magnetisation by this amount of anisotropy field. The Ni
nanowire is polycrystalline with a face centre cubic structure and it has a weak magneto-crystalline anisotropy (magneto-crystalline anisotropy constant \( K_1 = 10^4 J/m^3 \) \[15\]). Therefore, magneto-crystalline anisotropy of such nanowires can be neglected. Here, there is no chance to induce stress anisotropy to the nanowires, because the measurements were performed at room temperature and there is no source of stress anisotropy to introduce. Thus, stress anisotropy, can also be neglected from the analysis presented here. Lastly, the surface to volume ratio in such nanowires is low; therefore, the surface anisotropy makes a small contribution to the magnetisation behaviour of such wires due to uneven coordination number of the magnetic spins near the surface. These spins on the nanowires surface try to make the perpendicular direction as an easy axis of magnetisation. For more details on these mechanisms can be found in Refs \[15, 25, 44\].

Thus, the closely packed two Ni nanowires investigated here behaves like a single nanowire and the resultant magnetic anisotropic direction is directed by the shape anisotropy of the nanowires. As a result, squared shaped hysteresis loops were obtained when the magnetic field was applied along their long axial direction. In contrary, the hysteresis loops for two dimensional arrays of Ni nanowires seen in literature showing the perpendicular direction as an easy axis of magnetisation [6, 37, 44, 51]. This is solely due to the strong magneto-static interaction between the nearest nanowires, which makes the perpendicular direction of the templated nanowires is easy to magnetise. Here, the squareness ratio exhibits constant maximum values is expected due to the small number of nearest neighbors and hence no magneto-static interaction. The plot of coercivity against the angle of applied magnetic field shown in Figure 5 indicates that the coercivity decreases continuously with increasing nanowires angle due to a reduction in the shape anisotropy. Small distribution of the switching fields was obtained from repeating measurements at different locations on the same wires can be attributed to the variation in nanowire diameters or their misalignment with each other.

Various distinct mechanisms are able to describe the magnetisation reversal behaviour in nanoparticles, nanotubes and nanowires [2-4, 15, 46, 47]. For instance; coherent rotation of the Stoner - Wohlfarth model appears when the nanostructures are smaller than the exchange length, while curling and buckling modes occur when they are larger than the exchange length. In order to understand and explain the angular dependence of the coercivity presented here, it should first be noted that since the diameters of these nanowires (~300 nm) are much larger than the exchange length, therefore one can expect a deviations of the micromagnetic structure away from the uniform rotation behaviour that is described by the Stoner-Wohlfarth reversal model \[47\]. Since buckling behaviour is expected to occur when the nanowire diameter is comparable to the exchange length (reduced radius around unity), buckling may also be excluded from the applicability here. The shape and aspect ratio of the closely packed two Ni nanowires investigated here is far from being a chain of spheres which would reverse as in the fanning model \[15\]. Therefore, the angular dependence of the switching fields is likely to be the consequence of an incoherent rotational process following the curling model of domain reversal.

5. Conclusions

Here, the surface morphology and magnetisation behaviour of electrodeposited isolated cylindrical Ni nanowires was intensively investigated using a high resolution scanning electron microscopy and MOKE magnetometry, respectively.

The surfaces of a wide range of Ni nanowires were found to be homogenous, uniform, and cylindrical in shape. Some other nanowires showed different diameters and surface features, including; protrusions and branches along their length. These variations were attributed to defects exist in the internal surfaces of the pores within the template itself, or due to trapped air pockets within the pores during nanowires growth. The diameters distribution histograms of a wide range of nanowires were found to differ from their template pores diameters. This was attributed to the oxide formation or residual templates and contaminants which may cover these wires or due to the defects in the template itself. The distribution of lengths was found to differ from their actual lengths estimated during deposition growth was attributed to the breakage of nanowires during the releasing process.

Due to the small surface area of nanowires, the signal to noise ratio of the hysteresis loops was found to be always low. The dipolar or magneto-static interaction between the Ni nanowires was expected to be low or even not exist, due to the small number of nearest neighbour, therefore the squareness ratio was found to be high (around 0.86) at all angles of measurements investigated here. Maximum coercivity was obtained (average ~110 Oe), when the magnetic field was applied parallel to the nanowires long axis, and low coercivity (average ~75 Oe) observed when the magnetic field was applied perpendicular to the nanowires long axis and this was attributed to the reduction in the shape anisotropy. Small distribution of coercivity was obtained from repeating measurements at different locations on the same nanowires. This was attributed to the variation in nanowire diameters or their misalignment with each other. Furthermore, the angular dependence of corecivity was investigated regarding the mechanisms described in literature. The angular dependence of corecivity was likely to be the consequence of an incoherent rotational process following the curling model of domain reversal.

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