

Supporting Information for

Direct observation of dynamical magnetization reversal process governed by shape anisotropy in single NiFe₂O₄ nanowire

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Crystalline structure and chemical analysis of NiFe_2O_4 single-particle-chain nanowire

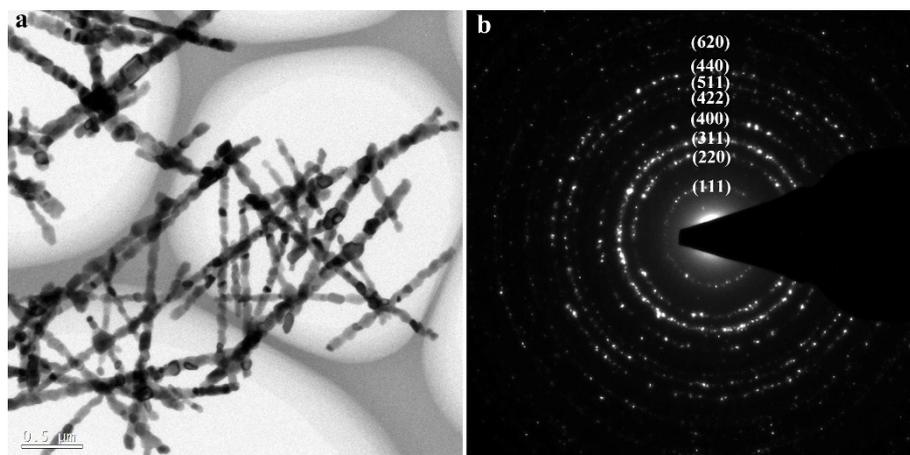


Figure S1. (a) The TEM image of NiFe_2O_4 NWs; (b) The corresponding SAED.

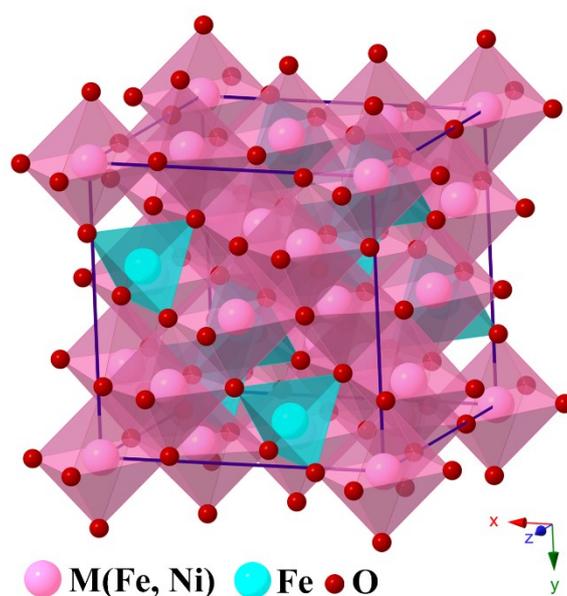


Figure S2. Polyhedral model of cubic spinel nickel ferrite, showing a face-centered cubic network of O^{2-} anions with Ni^{2+} and Fe^{3+} cations occupying two interstitial sites. Half of Fe^{3+} cations occupies tetrahedral-A sites marked by cyan. The other half of Fe^{3+} and Ni^{2+} cation randomly occupy octahedral B sites marked by pink.

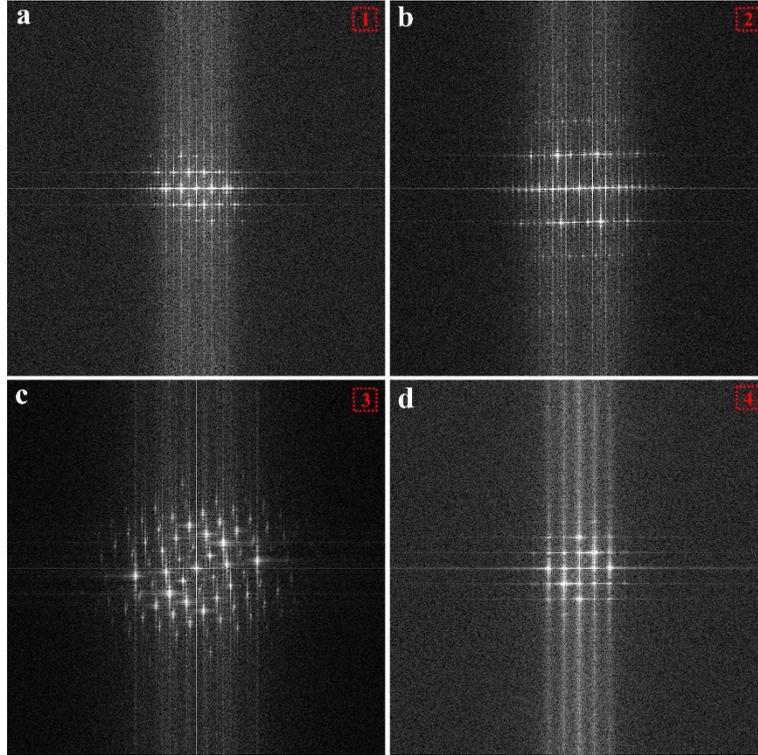


Figure S3. The corresponding FFT for experimental high angle angular dark field-scanning transmission electron microscopy (HAADF-STEM) images in Figure 1.

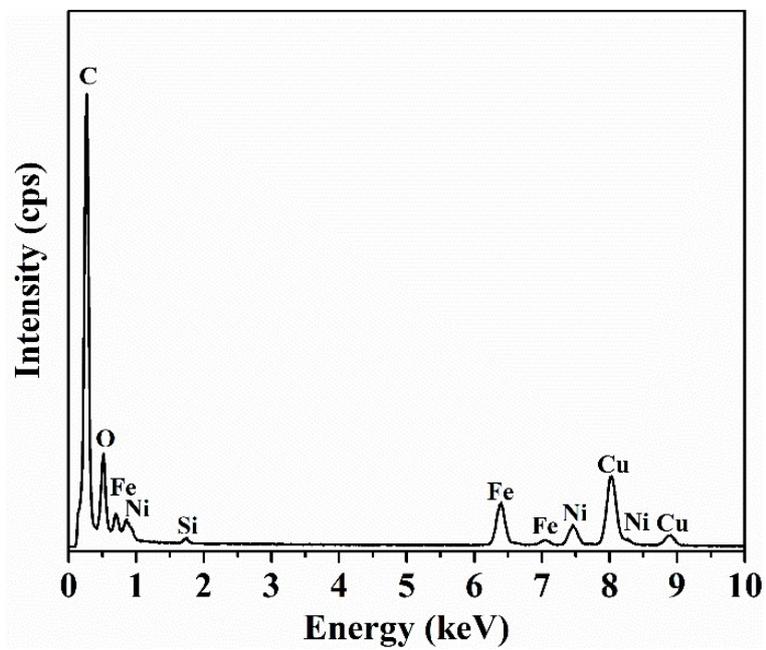


Figure S4. The EDX spectrum acquired from the area in Figure S1a.

Magnetic properties of NiFe₂O₄ nanowire.

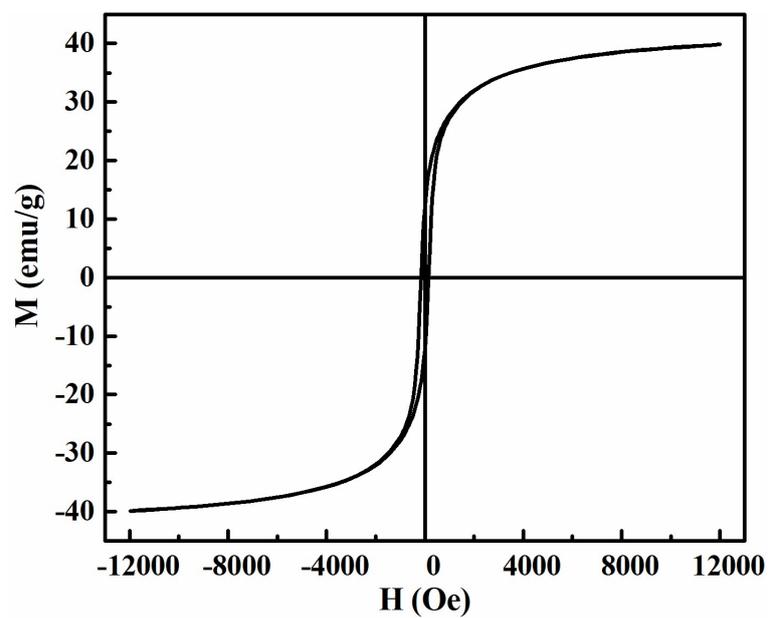


Figure S5. Hysteresis loop of the NiFe₂O₄ NWs.

Detail of the magnetization reversal mechanism

Coherent rotation diameter D ($D_{\text{coh}}=3.655l_{\text{ex}}$ for one-dimensional magnetic materials) is regarded as criteria to judge which magnetization reversal mechanism may occur in magnetic materials. When the diameter D of NW is smaller than D_{coh} , coherent rotation occurs. Conversely, curling (incoherent) rotation happens when R is larger than D_{coh} . l_{ex} is exchange length, given by:

$$l_{\text{ex}} = \sqrt{\frac{A}{\mu_0 M_s^2}} \quad (1)$$

where, A is exchange constant, 1.15×10^{-6} erg·cm⁻¹ for NiFe₂O₄, μ_0 is permeability of free space, $4\pi \times 10^{-7}$ N/A², and M_s is the saturated magnetization, 270 emu/cm³ for our samples. D_{coh} of NiFe₂O₄ NW is 81.8 nm, a little small than the diameter (91nm) of NW in Figure 3. Therefore, the magnetization reversal mechanism of this NiFe₂O₄ NW is curling rotation mode, consistent with above observed results.