

Electronic supplementary information (ESI) available

Self-Assembled Mesoporous Co and Ni-Ferrite Spherical Clusters Consisting of Spinel Nanocrystals Prepared using a Template-Free Approach

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Table S1. FWHM values of the main diffraction peaks and the crystallite size for M-CoFe₂O₄ with the respective diffraction planes.

h	k	l	2 θ (deg.)	<i>d</i> -spacing (nm)	Intensity (a.u.)	FWHM (2 θ)	Crystallite size (nm)
1	1	1	18.10	0.48	22.9	1.16	6.85
2	2	0	30.14	0.29	32.5	1.28	6.35
3	1	1	35.44	0.25	100.0	1.12	7.36
4	0	0	42.98	0.21	22.6	1.16	7.28
4	2	2	53.54	0.17	13.0	1.64	5.36
5	1	1	57.00	0.16	27.7	1.60	5.58
4	4	0	62.64	0.15	36.8	1.44	6.38

Table S2. FWHM values of the main diffraction peaks and the crystallite size for M-NiFe₂O₄ with the respective diffraction planes.

h	k	l	2 θ (deg.)	<i>d</i> -spacing (nm)	Intensity (a.u.)	FWHM (2 θ)	Crystallite size (nm)
1	1	1	17.98	0.48	32.4	1.60	4.97
2	2	0	30.06	0.29	43.3	1.48	5.50
3	1	1	35.42	0.25	100.0	1.48	5.57
4	0	0	42.92	0.21	30.7	1.04	8.12
4	2	2	53.30	0.17	15.9	1.62	5.53
5	1	1	56.74	0.16	31.7	1.00	8.93
4	4	0	62.48	0.15	42.0	1.56	5.89

Table S3. The physical properties of as-synthesized mesoporous ferrite clusters (as-M-CoFe₂O₄ and as-M-NiFe₂O₄).

Samples	Cluster size (nm) ^a	Surface area (m ² g ⁻¹)	Pore diameter (nm)	Pore volume (cm ³ g ⁻¹)
as-M-CoFe ₂ O ₄	162	103	3.42	0.14
as-M-NiFe ₂ O ₄	167	111	3.39	0.12

a: the mean ferrite cluster size was measured using DLS

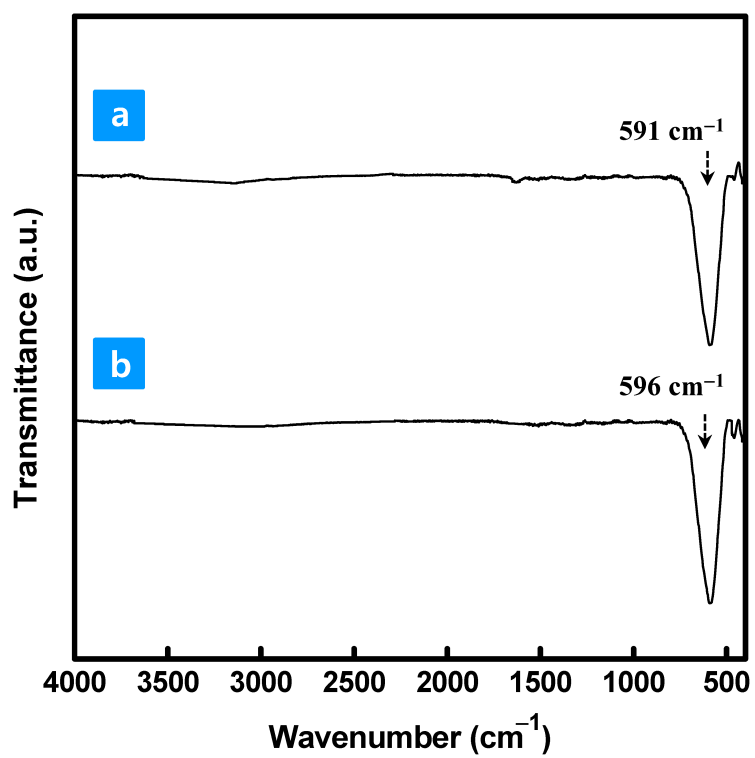


Fig. S1 FT-IR spectra of (a) M-CoFe₂O₄ and (b) M-NiFe₂O₄.

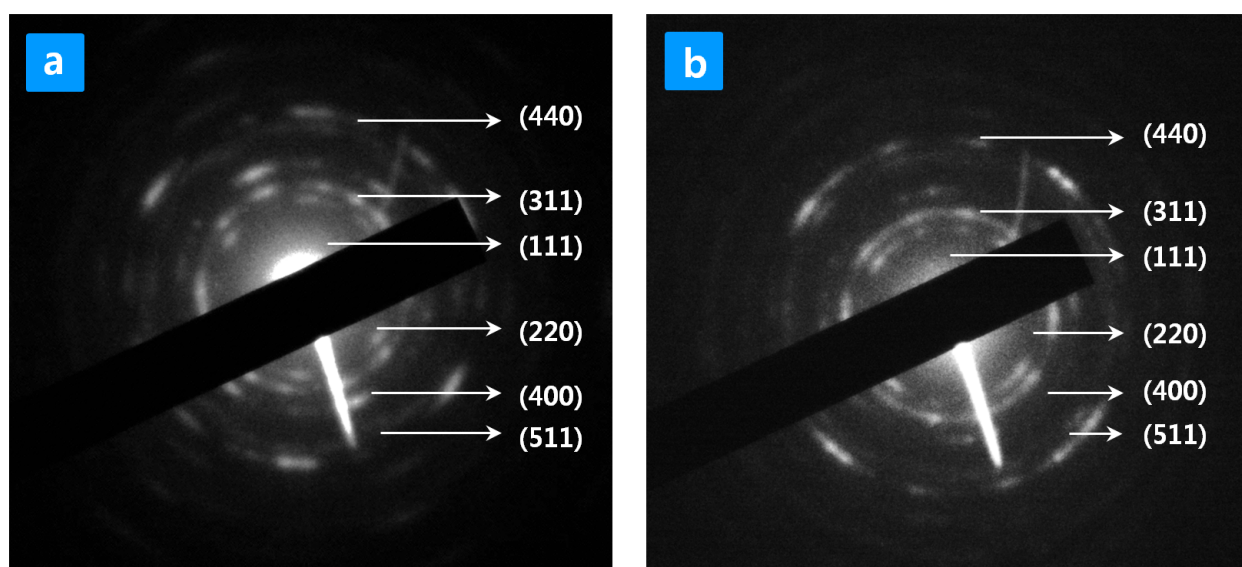


Fig. S2 Selected area electron diffraction (SAED) patterns acquired from a particular (a) M-CoFe₂O₄ and (b) M-NiFe₂O₄ cluster.

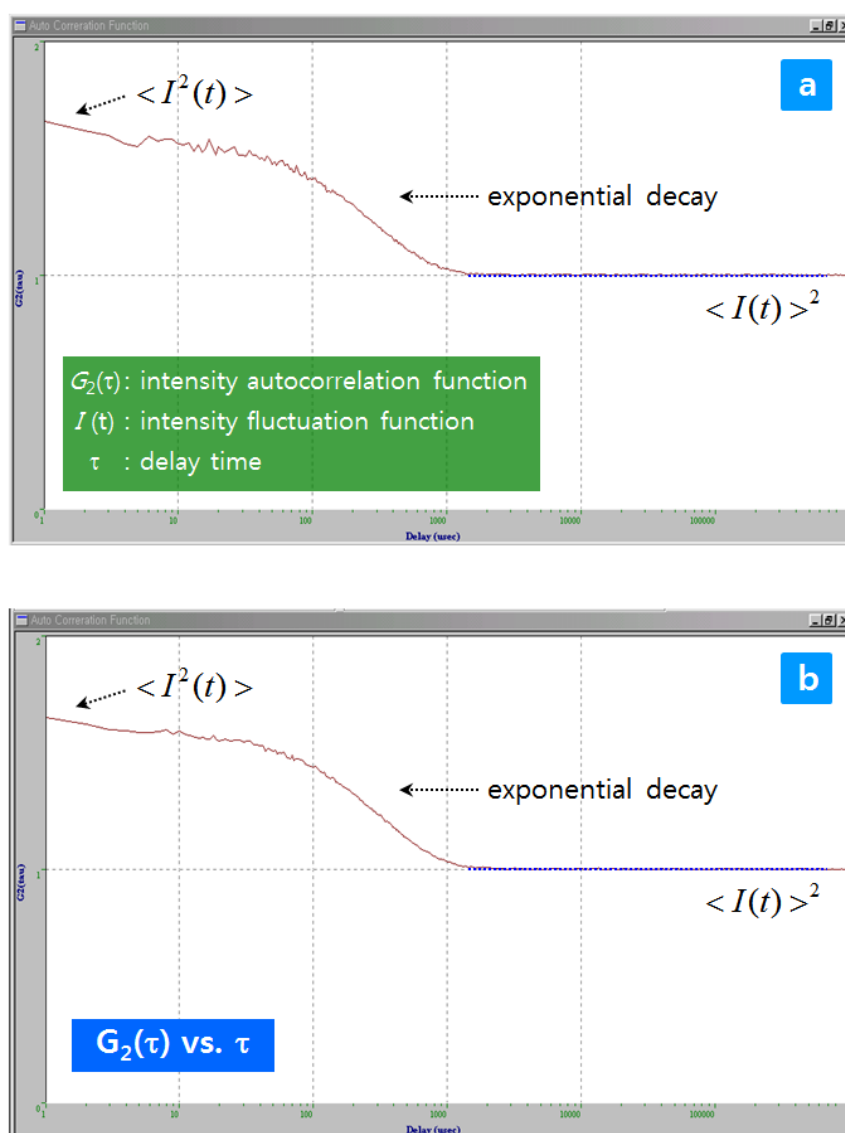


Fig. S3 The intensity auto-correlation function (ACF), $G_2(\tau)$ for (a) M-CoFe₂O₄, (b) M-NiFe₂O₄ samples in DLS.

The second-order correlation function $G_2(\tau)$ can be expressed as a function of the first-order correlation function $G_1(\tau)$ according to the Siegert relation: $G_2(\tau) = B(1 + \beta G_1(\tau)^2)$, where B is the baseline constant and β is a coherence constant. In the case of a perfect setup, both equal unity. In the case of single-exponential decay, $G_1(\tau)$ can be expressed in terms of a typical decay rate Γ and time t ; $G_1(\tau) = \exp(-\Gamma \tau)$.

The apparent translational diffusion coefficient, D , is given by equation: $\Gamma = Dq^2$, where q is the magnitude of the scattering vector $q = 4\pi n \sin(\theta/2)/\lambda$, where n is the refractive index of the solvent, θ is the scattering angle, and λ is the wavelength of the incident light. For spherical particles, the translational diffusion coefficient can be related to the hydrodynamic radius, R , according to the Stokes-Einstein equation: $D = k_B T / 6\pi\eta R$, where D is the diffusion coefficient of the Brownian motion of spherical particles, k_B is the Boltzmann constant, T is the absolute temperature, and η is the viscosity of the solvent. The hydrodynamic radius distribution of particles, $G(R)$ was estimated using the COTIN algorithm, which is conventionally used to determine the inverse Laplace transform of the measured amplitude autocorrelation function.^{1,2}

- (1) R. Finsy, *Adv. Colloid Interfac.* 1994, **52**, 79.
- (2) I. K. Voets, A. De Keizer, M. A. Cohen Stuart and P. De Waard, *Macromolecules* 2006, **39**, 5952.

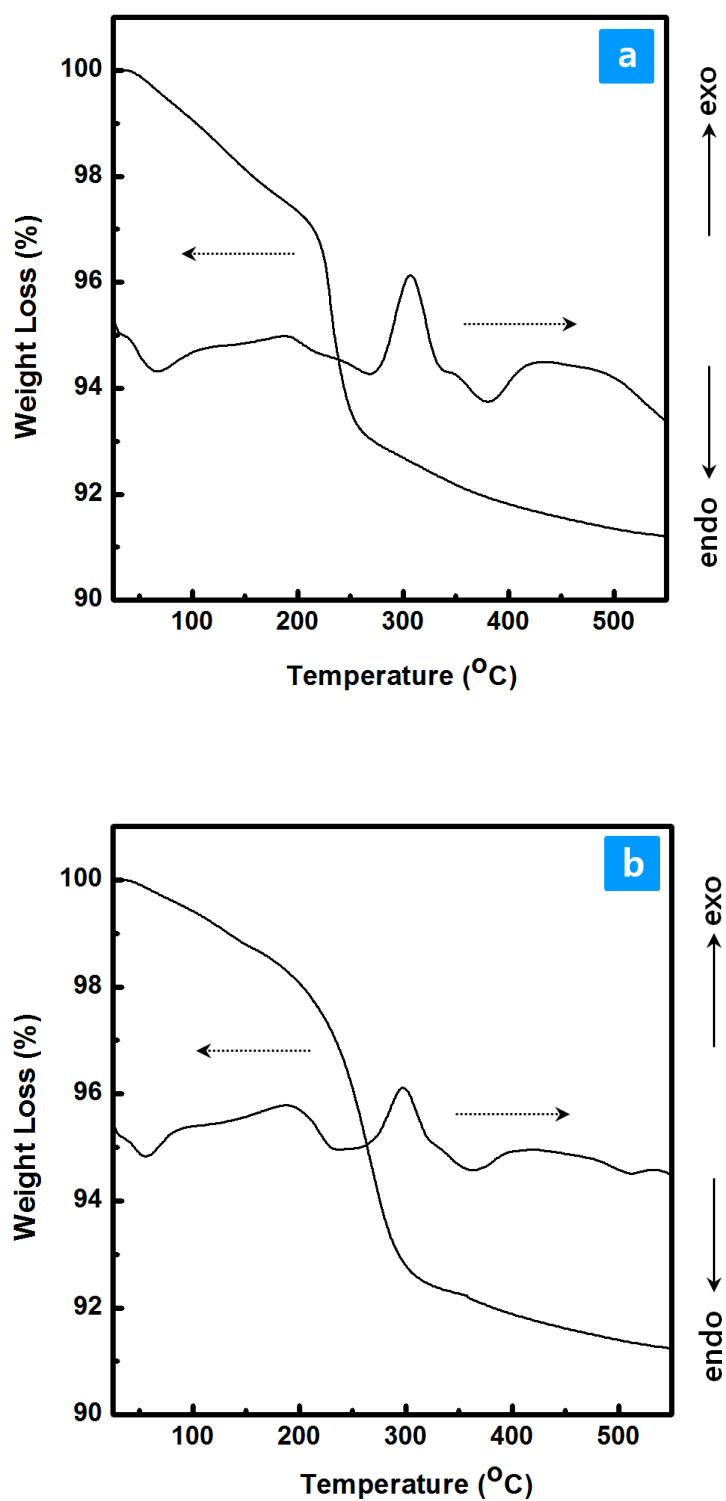


Fig. S4 TGA and DSC curves of as-prepared (a) M-CoFe₂O₄ and (b) M-NiFe₂O₄ before calcination.

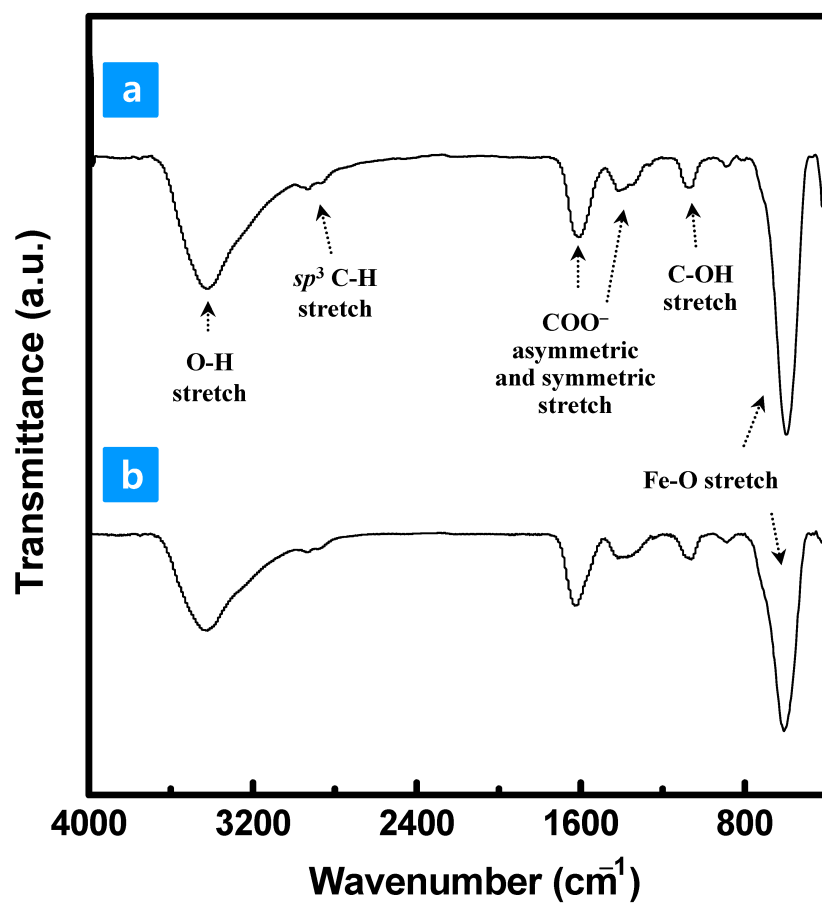


Fig. S5 FT-IR spectra of as-synthesized (a) M-CoFe₂O₄ and (b) M-NiFe₂O₄ before calcination.

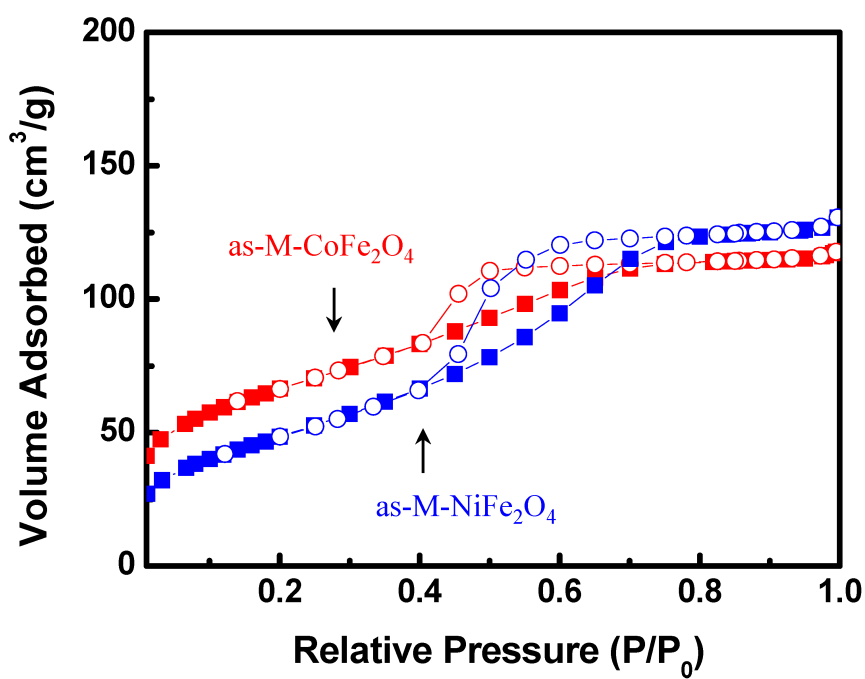


Fig. S6 N₂ adsorption(■) -desorption(○) isotherms for as-synthesized (a) M-CoFe₂O₄ and (b) M-NiFe₂O₄.

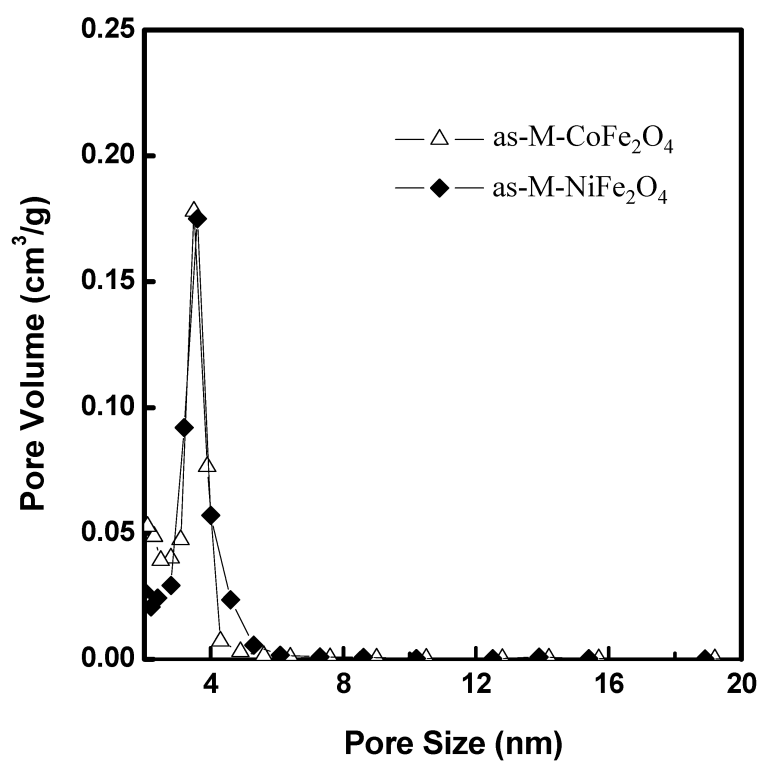


Fig. S7 The pore size distribution of as-synthesized (a) M-CoFe₂O₄ and (b) M-NiFe₂O₄.

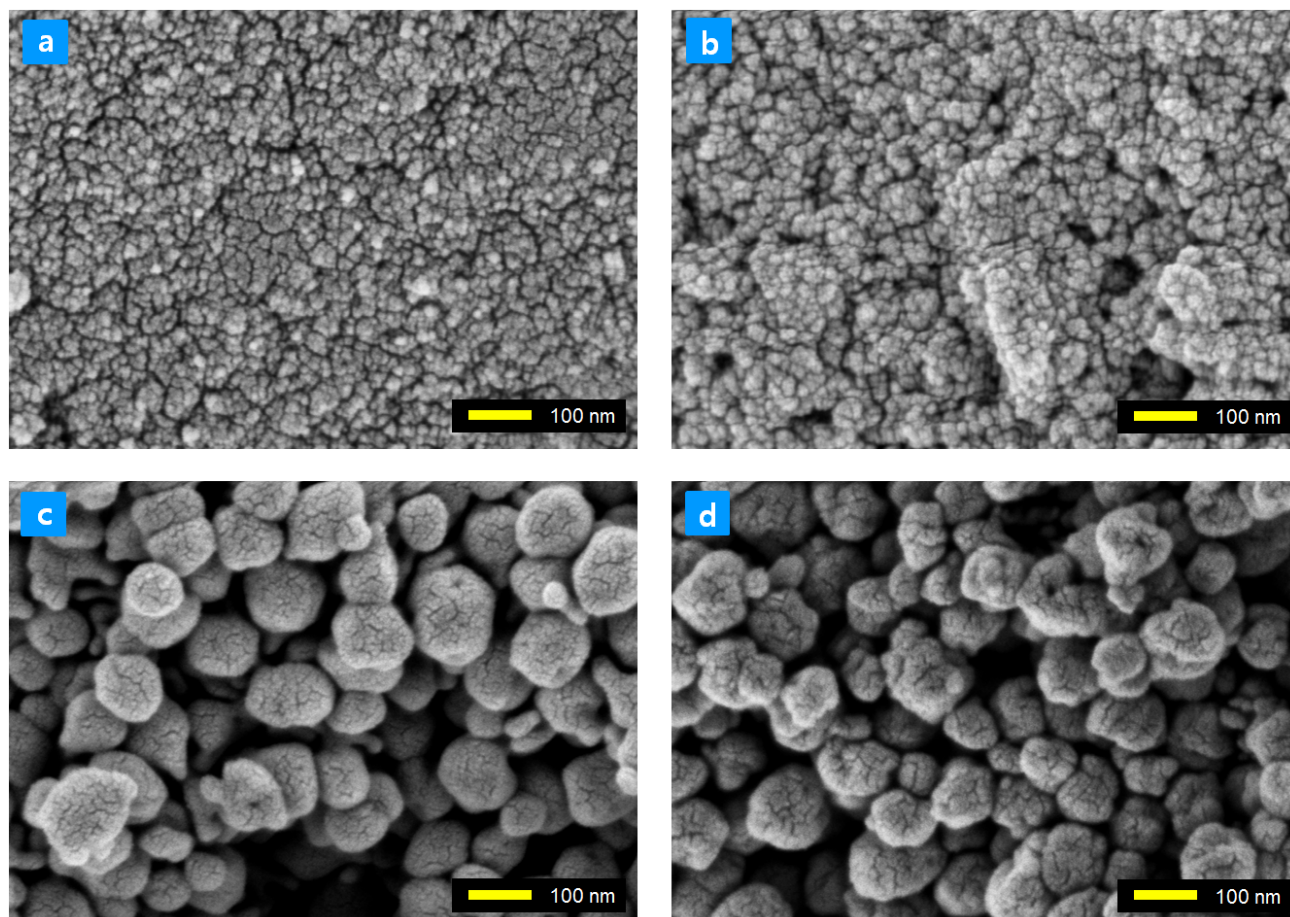


Fig. S8 FE-SEM images of (a, c) CoFe₂O₄ and (b, d) NiFe₂O₄ at different sodium acetate concentrations; (a, b) high concentration and (c, d) low concentration.

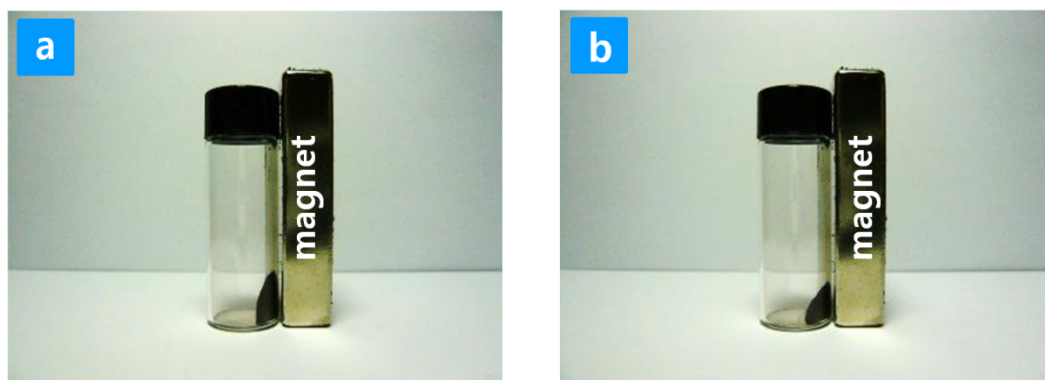


Fig. S9 Photograph of (a) M-CoFe₂O₄ and (b) M-NiFe₂O₄ particles in the presence of a magnetic field.