Resolving Anomalies in the Critical Exponents of FePt Using Finite-Size Scaling in Magnetic Fields

J. Waters,^{1,*} D. Kramer,¹ T.J. Sluckin,² and O. Hovorka¹

¹Engineering and Physical Sciences, University of Southampton, Southampton, SO17 1BJ, United Kingdom ²School of Mathematical Sciences, University of Southampton, Southampton, SO17 1BJ, United Kingdom

(Received 24 October 2018; published 11 February 2019)

FePt is the primary material being considered for the development of information storage technologies based on heat-assisted magnetic recording (HAMR). A practical realization of HAMR requires understanding the high-temperature phase transition behavior of FePt, including critical exponents and Curie temperature distributions as the fundamental HAMR media design characteristics. The studies so far found a significant degree of variability in the values of critical exponents of FePt and remain controversial. Here, we show that at the heart of this variability is the phase transition crossover phenomenon induced by two-ion anisotropy of FePt. Through Monte Carlo simulations based on a realistic FePt effective Hamiltonian, we demonstrate that in order to identify the critical exponents accurately, it is necessary to base the analysis on field-dependent magnetization data. We have developed a two-variable finite-size scaling method that accounts for the field effect. Through the use of this method, we show unambiguously that true critical exponents of FePt are fully consistent with the three-dimensional Heisenberg universality class.

DOI: 10.1103/PhysRevApplied.11.024028

I. INTRODUCTION

Heat-assisted magnetic recording (HAMR) is a rapidly developing technology, designed to address the magnetic recording "trilemma" [1]. Materials with high uniaxial anisotropy are required to increase thermal stability of magnetic grains, the most notable of these being FePt in the $L1_0$ phase. The high anisotropy of FePt is overcome during the recording process by laser heating the grains close to their Curie temperature T_c , which is dependent on the size of the FePt grains and their geometry. Understanding the dependence of T_c on the finite-size effects of magnetic grains in granular films is required for advancing HAMR technology.

The divergent behavior of thermodynamic quantities in materials near T_c is described by critical exponents and universal scaling functions, which depend on the type and dimensionality of the material. Such critical exponents and universal scaling functions are used in a variety of studies of high-temperature magnetism. Examples include the models of thermalized dynamics based on the Landau-Lifshitz-Bloch equation [2], thermodynamic characterization of materials near T_c [[3], p. 80], or the experimental identification of T_c distributions in magnetic granular media [4,5]. Knowledge of accurate critical

exponents is thus essential for a reliable high-temperature quantification and optimization of materials for HAMR.

The critical exponents of FePt have been studied by fitting the power-law behavior to the magnetization versus temperature M(T) data [6] or to the dependence of T_c on the grain size R [7,8]. Another widely used approach has to use finite-size scaling analysis, rescaling M(T) for grains of different R so that they collapse onto a single curve [9,10]. However, considerable variation of the critical exponents was found depending on the method used to identify them as illustrated in Table I. Even small variations in the critical exponents can lead to large errors in the estimation of, for instance, the T_c distributions [9]. It therefore becomes important to understand the reasons for these discrepancies and establish the values that should be used for the critical exponents.

In this work, we investigate this issue by employing large-scale Monte Carlo simulations based on the FePt effective Hamiltonian obtained earlier from *ab initio* calculations [12]. We identify that the differences in the values of the critical exponents reported previously result from the phase transition crossover effect due to the presence of the two-ion anisotropy in FePt. Although small in comparison to exchange interactions, the strength of the two-ion anisotropy is sufficient to conceal the information about the true critical exponents in the temperature region near T_c . We show that generalizing the finite-size scaling analysis to consistently include the field-dependent magnetization data allows us to circumvent the crossover

^{*}j.m.waters@soton.ac.uk

TABLE I. Values of the magnetization β and correlation length ν critical exponents and the bulk Cure temperature T_c^b collected from the literature. The values obtained in this work using the magnetic-field-dependent finite-size scaling method fully agree with the values of the 3D Heisenberg model (Table II).

	Critical paramaters			
Reference	β	ν	T_c^b (K)	
Hovorka et al. [9]	0.33 ± 0.10	0.85 ± 0.10	677 ± 11	
Lyberatos et al. [10]	Not available	1.06 ± 0.06	658 ± 4	
Rong et al. [7]	Not available	0.67 ± 0.11	775 ± 20	
Zhao et al. [8]	0.327	0.631	642.5	
This work	0.366 ± 0.001	0.72 ± 0.17	654 ± 2	
(Heisenberg)				

effect. In this way, we demonstrate unambiguously that, in the case where there is an external magnetic field, the critical exponents of FePt are fully consistent with the three-dimensional (3D) Heisenberg model.

The article is organized as follows. In Sec. II, we develop the finite-size scaling method, based on the scaling of temperature and field-dependent magnetization data, M(T, H), rather than M(T). In Sec. III, we apply this technique to study the critical parameters of FePt and establish their relationship with the Heisenberg model. In Sec. IV, we discuss the crossover effects relevant to FePt and underline the dominant role of the two-ion anisotropy. In Sec. V, we conclude with recommendations for the critical exponents to use in future work.

II. TWO-VARIABLE FINITE-SIZE SCALING

In the present study, we consider the finite-size scaling (FSS) analysis as an established and robust technique for extracting the critical exponents in finite-size simulations and small systems in general [13,14]. Considering magnetic grains of different diameters R, the FSS in its simplest form allows relating the temperature-dependent M(T) data for different R to the magnetization and correlation critical exponents, β and ν , and the universal scaling function \tilde{M} as

$$M(T;R) = R^{-\beta/\nu} \tilde{M} \left(R^{1/\nu} t \right), \qquad (1)$$

where $t = (T - T_c^b)/T_c^b$ is the reduced temperature, T_c^b the Curie temperature of a bulk system, and M represents the projection of the magnetization vector onto the *z*-axis. The critical exponents β and ν can be identified by plotting the M(T; R) data for different R as $R^{\beta/\nu}M$ versus $R^{1/\nu}t$ and systematically choosing the values of β , ν , and T_c^b until all data collapse onto a single universal curve representing \tilde{M} . A method convenient for producing such data collapses has been developed earlier [11] and for completeness is reviewed here in the Appendix.

The FSS analysis based on Eq. (1) is applicable only to M(T; R) data in zero magnetic field, H = 0. Nonzero magnetic fields introduce a rounding effect when M no longer approaches zero at the critical point but instead follows a power-law behavior $M \sim H^{1/\delta}$, where δ is the magnetic field critical exponent. To incorporate the magnetic field dependence of the data in the analysis, it is necessary to generalize Eq. (1) to include this field contribution, which results in the following two-variable FSS form:

$$M(T,H;R) = R^{-\beta/\nu} \tilde{M} \left(R^{1/\nu} t, R^{\beta\delta/\nu} H \right).$$
(2)

The exponent δ is related to β and ν through the so-called hyperscaling relation [13]:

$$\delta = d\nu/\beta - 1,\tag{3}$$

with *d* being the dimension of the spin lattice, e.g., d = 3 for three-dimensional systems such as a grain in a magnetic hard disk. The form of Eq. (2) suggests that the two-variable scaling approach is based on scaling the data seen as surfaces $R^{\beta/\nu}M(T,H;R)$ versus $R^{1/\nu}t$ and $R^{\beta\delta/\nu}H$ and then tuning the values of β , ν , δ , and T_c^b until a unique collapse onto the surface of \tilde{M} is achieved. Details of our practical implementation of the two-variable scaling method are summarized in the Appendix.

To validate the two-variable scaling approach based on Eq. (2), we consider Metropolis Monte Carlo simulations of a two-dimensional (2D) Ising model, which is well established, and rigorous analytical results for the critical exponents and T_c are available for computational benchmarking. The Hamiltonian of the field-dependent twodimensional Ising model is $\mathcal{H} = -H \sum_{i} s_{i} - \sum_{ij} J_{ij} s_{i} s_{j}$, where H is the strength of the external magnetic field being applied to the system, s_i is the *i*th spin taking values ± 1 , and $J_{ii} > 0$ is the ferromagnetic exchange interaction between s_i and s_j such that $J_{ij} = J = 1$ for neighboring spins and J = 0 otherwise. In the Monte Carlo simulations, the magnetization is sampled for three different circular cuts of square lattices with radii R = 25, 50, and 75 lattice spacings. Note that physical dimensions in this Ising-model-based test case are irrelevant. For each lattice size, the average magnetization is computed in the temperature range from $T = 1.775J/k_b$ to $2.725J/k_b$ and in the range of external field strengths of H = -0.005Jto 0.005J. This region encompasses the critical point at H = 0 and $T_c = 2.269 J/k_b$. For each field strength, lattices are annealed from high to low temperature, with 10 000 Monte Carlo sweeps (MCS) to equilibrate and then 51 200 samples taken at intervals of 50 MCS.

A. Validation of one-variable FSS

As a test case, the one-variable FSS analysis of only the H = 0 data sets gives $\beta = 0.14 \pm 0.01$, $\nu = 1.05 \pm 0.08$, and $T_c^b = 2.265 \pm 0.004 J/k_b$, which agree well with the



FIG. 1. The results of finite-size scaling of Ising model data according to Eq. (1). (a),(b) show the scaled curves when H = 0.0005 and H = 0.005, respectively. The resulting critical parameters (c) β , (d) ν , and (e) T_c^b that result from this method are plotted for a series of external field strengths. The error bars in (c)–(e) represent the uncertainties consistent with a 99% confidence level [11].

theoretical values for the 2D Ising model, $\beta = 0.125$, $\nu =$ 1, and $T_c^b = 2.269$ [13]. Next we apply the one-variable FSS to the 2D Ising model magnetization data obtained in a nonzero magnetic field $H \neq 0$. The obtained data collapses are successful for all considered field strengths, as suggested in Figs. 1(a) and 1(b) for fields differing by a factor of 10. However, despite the successful data collapses, the obtained values of the critical parameters β , ν , and T_c^b tend to increase with H as shown in Figs. 1(c)-1(e). Thus, the identified values of the critical exponents and T_c^b appear to be incorrectly field dependent despite the fact that the magnetic field is relatively weak, never greater than 0.5% of the strength of the exchange field, and therefore certainly not the dominant interaction. An analysis of the data using the zero-field ansatz in Eq. (1) thus introduces a fielddependent bias of the critical exponents, which no longer represent the true critical exponents of the 2D Ising model. We also attempt to extrapolate the field-dependent critical exponent data in Figs. 1(c)-1(e) to H = 0 and obtain the critical parameters $\nu = 0.126$, $\beta = 0.988$, and $T_c^b = 2.256$, which agree with the theoretical values for the 2D Ising model surprisingly well. Thus, the one-variable FSS combined with extrapolation to H = 0 appears to be a valid technique for extracting the accurate values of the critical exponents. Unfortunately, as demonstrated below, because of the crossover effects, this procedure is not applicable to systems with more complex Hamiltonians containing contributions from magnetic anisotropies. Since FePt suffers from crossover effects, the expropolation approach is not applicable.

B. Validation of two-variable FSS

Figure 2(a) shows the M(T, H) surfaces obtained from the Monte Carlo simulations of the 2D Ising model used above. Figs. 2(b) and 2(c) show the cuts through these data surfaces at specific values of the field and temperature. Figure 2(d) shows the data collapse of these M(T, H)data sets for all R obtained by using the scaling procedure described in the Appendix. Figures 2(e) and 2(f)demonstrate representative data cuts through the collapsed surface and suggest excellent scaling in both H and T. The scaling procedure yields $\beta = 0.136 \pm 0.018$, $\nu = 1.09 \pm$ 0.07, and $T_c^b = 2.25 \pm 0.02$, which are in very good agreement with the values of the 2D Ising model. Using Eq. (3), we find $\delta = 15.0 \pm 2.4$, consistent with the analytical $\delta = 15$. The strong agreement with analytical results demonstrates the validity of the two-variable FSS in the presence of magnetic fields, which will turn out to be essential for the analysis of the data of FePt below.



FIG. 2. (a) The magnetization surfaces in the *H*-*T* plane corresponding to the Ising model. Slices through these surfaces are shown at (b) H = 0.005 and (c) T = 2.25625. (d) The same magnetization surfaces as shown in (a) after rescaling according to Eq. (2). Slices through these surfaces are shown at (e) $R^{\beta\delta/\nu}H = 1.871$ and (f) $R^{1/\nu}(T - T_c^b)/T_c^b = -0.029$.

III. CRITICAL EXPONENTS OF FePt

To study the phase transition behavior of FePt, we consider the following realistic classical effective spin Hamiltonian:

$$\mathcal{H} = -\sum_{ij} \left(J_{ij} \mathbf{s}_i \cdot \mathbf{s}_j + d_{ij}^{(2)} s_i^z s_j^z \right) -\sum_i \left(d_i^{(0)} \left(s_i^z \right)^2 + \mu_{\rm Fe} H s_i^z \right).$$
(4)

Here, $\mu_{\text{Fe}} = 3.23\mu_B$ is the effective magnetic moment of an Fe atom, with μ_B the Bohr magneton, and *H* is the strength of the magnetic field oriented along the *z* axis. The Fe spins \mathbf{s}_i are represented as Heisenberg spins with magnitude $|\mathbf{s}_i| = 1$. The effective exchange interaction J_{ij} is not restricted to the nearest-neighbor Fe spins and, while the contribution from the short-range interactions is stronger, the long-range contributions remain significant and cannot be truncated in the phase transition studies of FePt. Overall, this amounts to up to 1358 neighbor pairs for each spin in the crystal lattice. In addition, Eq. (4) contains a single-ion energy term with strength $d_i^{(0)}$ and a two-ion term with strength $d_{ij}^{(2)}$. These anisotropy terms introduce the tendency toward the spin alignment along the *z* axis to maximize the s_i^z spin component. The magnitude of these anisotropy terms is small and they act as a perturbation to the exchange energy term. For example, as discussed in more detail below, the strength of the two-ion anisotropy is less than 1% of the strength of the exchange interaction. The effective Hamiltonian given by Eq. (4) is determined by mapping the energy of FePt in the L1₀ phase to a tetragonal lattice of Fe atoms based on *ab initio* calculations [12] and is now widely used for studying FePt for HAMR.

To simulate the magnetic field and temperaturedependent magnetization data of the model defined by Eq. (4), we use the Monte Carlo method introduced earlier. In simulations, we consider grains as spherical particles with radii R in the range from 2 to 5 nm. The magnetic field strengths normalized in the energy units as $\mu_{\text{Fe}}H$ are chosen in the range from 0 to 3 meV (0–16 T) and the temperature T range from 310 to 800 K. The



FIG. 3. The *z*-axis magnetization profiles of zero-field FePt data. Inset: The same magnetization profiles after rescaling according to Eq. (1).

temperature-dependent magnetization curve at every field value is generated by applying the annealing protocol by decreasing the temperature self-consistently in small steps starting from 800 K. An equilibration time of 10 000 MCS is applied for every field and temperature value and 6048 samples are taken for averaging at intervals of 50 MCS.

A. One-variable FSS

We first identify the critical exponents of FePt using the one-variable FSS based on Eq. (1). Figure 3 shows an example of the raw M(T) data for particles of variable R and corresponding to H = 0. Here, M represents the z component of the lattice-averaged magnetization vector. The FSS analysis of these zero-field data leads to excellent data collapse shown in the inset in Fig. 3. The values of the critical parameters are estimated as $\beta = 0.28 \pm 0.08$, $\nu = 0.94 \pm 0.15$, and $T_c^b = 678 \pm 15$ K and are reasonably consistent with our earlier work [9], as shown in Table I. Any differences could be attributed to the details of the data analysis, such as the density of data points or the choice of temperature range used in FSS. An over-reliance on temperatures close to T_c can enhance the effect of the crossover phenomena, as is discussed below. We also attempted the one-variable FSS of the field- and temperature-dependent FePt data, similarly to the 2D Ising model test case in Fig. 1. Unfortunately, we do not succeed in producing meaningful data collapses and, therefore, the one-variable FSS based on Eq. (1) does not allow us to identify the critical exponents from the FePt magnetization data for nonzero magnetic fields.

B. Two-variable FSS

Figure 4(a) shows the full set of the FePt magnetization data computed for different *T*, *H*, and *R*. Figures 4(b) and

4(c) show slices through the data surfaces at specific values of *H* and *T*. The corresponding data collapse obtained by using the two-variable FSS ansatz Eq. (2) is shown in Fig. 4(d). The representative collapsed data cuts are shown in Figs. 4(e) and 4(f) and confirm that good scaling has been achieved. The critical parameters obtained from the two-variable FSS analysis are $\beta = 0.366 \pm 0.001$, $\nu = 0.72 \pm 0.17$, and $T_c^b = 654 \pm 2$ K, and the magnetic field exponent as found from Eq. (3) is $\delta = 4.9 \pm 0.9$. These results are in very good agreement with the known exponents for the 3D Heisenberg model for which $\beta = 0.36$, $\nu = 0.71$, and $\delta = 4.8$ [13].

Table II summarizes the results of the FSS analysis of the zero- and nonzero-field two-dimensional Ising model and FePt magnetization data studied in this work. The critical exponents of the Ising model obtained by onevariable FSS of the H = 0 data and two-variable FSS of the $H \neq 0$ data are both consistent with the theoretical values. For FePt, the two-variable FSS of the M(T,H)data gives critical exponents consistent with the Heisenberg model universality class. However, the one-variable FSS of the zero-field FePt data is not consistent with the Heisenberg model. The question arises if the origin of this inconsistency, absent in the Ising model case, is inherent to the chosen data analysis or if it is of a fundamental nature. Below, we argue that it is the result of the presence of anisotropy terms in the effective Hamiltonian Eq. (4), leading to phase transition crossover effects and complicating the identification of accurate values of the critical exponents from the zero-field data.

IV. CROSSOVER EFFECT

The most common sources of the crossover behavior during phase transitions are (i) the small residual fields acting on the system, (ii) the presence of weak anisotropies and long-range interactions, (iii) the effect of disorder, (iv) and finite-size effects [14]. We focus on analyzing case (ii) only, given that simulations allow setting well-controlled external field conditions, unlike experiments, thus ruling out the residual fields in (i); atomic disorder is not considered and, therefore, its effects (iii) are irrelevant in our simulations; and the finite-system-size effects (iv) are accounted for by choosing the analysis method based on the FSS ansatz Eqs. (1) or (2), rather than naive power-law fits.

A. Effect of anisotropy

It is useful to first estimate the magnitude of contributions of the individual terms in the effective Hamiltonian Eq. (4) in the critical temperature region. The effective field acting on a spin *i* expressed in the mean-field approximation reads $\mu_{\text{Fe}}\mathbf{H}_{\text{eff},i} = -\partial \mathcal{H}_{\text{mf}}/\partial \langle \mathbf{s}_i \rangle$, where $\langle \mathbf{s}_i \rangle$ is the expectation value of the spin \mathbf{s}_i consistent with the equilibrium Boltzmann statistics. According to Eq. (4), the



FIG. 4. (a) The magnetization surfaces in the *H*-*T* plane corresponding to the field-dependent FePt data, with slices through these surfaces at (b) $\mu_{\rm Fe}H = 3$ meV and (c) T = 608.57 K. (d) The same magnetization surfaces as shown in (a) after rescaling according to Eq. (2), with slices through these surfaces at (e) $R^{\beta\delta/\nu}\tilde{\mu}_{\rm Fe}H = 18.88$ and (f) $R^{1/\nu}(T - T_c^b)/T_c^b = -0.20$.

effective field $\mu_{\text{Fe}} \mathbf{H}_{\text{eff},i}$ can be written as a sum of contributions:

$$\sum_{j} J_{ij} \langle \mathbf{s}_{j} \rangle + \sum_{j} d_{ij}^{(2)} \langle s_{j}^{z} \rangle + d_{i}^{(0)} \langle s_{i}^{z} \rangle + \mu_{\rm Fe} H, \quad (5)$$

where the first term defines the local exchange field, the second and third terms are the local two-ion and single-ion

TABLE II. The results of FSS analysis performed in this work and the expected theoretical values. For comparison, the summary of the data from previous studies is given in Table I.

	Critical parameters			
	β	ν	δ	
2D Ising model				
H = 0 [Eq. (1)]	0.14 ± 0.01	1.05 ± 0.08	Not applicable	
$H \neq 0$ [Eq. (2)]	0.136 ± 0.018	1.09 ± 0.07	15 ± 0.02	
Analytic [13]	0.125	1	15	
FePt				
H = 0 [Eq. (1)]	0.28 ± 0.08	0.94 ± 0.15	Not applicable	
$H \neq 0$ [Eq. (2)]	0.366 ± 0.001	0.72 ± 0.17	4.9 ± 0.9	
3D Heisenberg [13]	0.36	0.71	4.8	

anisotropy fields, and the last term is the external field. All fields are in units of energy. Taking $|\langle \mathbf{s}_i \rangle| = 0.05$ as a representative value of magnetization in H = 0 at 650 K near the phase transition point (Fig. 3), we estimate the exchange and two-ion and single-ion anisotropy to be approximately 11, 0.15, and 8.45×10^{-5} meV, respectively. In the following analysis, the single-ion anisotropy contribution can therefore be neglected and the two-ion anisotropy taken as a small perturbation to the exchange energy term.

The anisotropy-induced crossover effect can be understood by the following qualitative argument [14] applied to the zero-field Heisenberg Hamiltonian with the twoion anisotropy acting as a small perturbation. The scaling form of the free energy of such a system reads $f_s = |t|^{2-\alpha} \tilde{F}(\langle d_{ij}^{(2)} \rangle |t|^{-\phi})$, where \tilde{F} is the universal scaling function for the free energy, $t = (T - T_c^b)/T_c^b$ is the reduced temperature as before, $\phi > 0$, and $\langle d_{ij}^{(2)} \rangle$ is an estimate of the characteristic two-ion anisotropy strength such as an average over all $d_{ij}^{(2)}$, for example. It can be shown that asymptotically, the function $\tilde{F}(x)$ for |x| << 1 represents the effective Heisenberg exchange-dominated system, while for |x| >> 1 the behavior is Ising-like [14]. Which of the two asymptotic regimes will be attained by the system at a given temperature depends on the value of $x = \langle d_{ij}^{(2)} \rangle |t|^{-\phi}$. For temperatures sufficiently far from T_c , one expects |t| to be large and $\langle d_{ij}^{(2)} \rangle |t|^{-\phi} \rightarrow 0$, given that ϕ is positive, and thus the behavior to be Heisenberg-like. However, for measurements close to T_c , the reduced temperature $t \rightarrow 0$ and $\langle d_{ij}^{(2)} \rangle |t|^{-\phi} \rightarrow \infty$, leading to an Ising-like asymptotic regime. The results of the analysis will, therefore, strongly depend on the temperature interval selected for the analysis.

To see this, we perform one-variable FSS analysis similar to Fig. 3 but with a restricted range of temperatures from $T_c^b - \Delta T$ to $T_c^b + \Delta T$ and varying the ΔT . Figure 5 shows the exponents β and ν identified in this way. As ΔT decreases, there is a change of the trend near T = 180 K. Exponent values below about 90 K could not be obtained due to the insufficient data points available in the now narrowly restricted temperature interval, so, instead, we used naive linear extrapolation to estimate the exponents for $\Delta T = 0$. The extrapolation procedure gives $\beta \approx 0.1$ and $\nu \approx 1.08$, which are, according to Table II, indeed close to the critical exponents for the 2D Ising model. Thus, even small anisotropic effects can completely alter the estimated value of the critical exponents. Note that the values of the critical exponents in the high ΔT limit in Fig. 5 are not expected to have a physical meaning



due to the mixed contributions from both asymptotic regimes.

The anisotropy-induced crossover behavior can be overcome by performing the data analysis in the nonzero applied magnetic field, stronger than the anisotropy field but sufficiently small to avoid driving the system away from the phase transition region. For our two-variable FSS analysis, we have applied magnetic fields in the interval 0-3 meV (0-16 T), which dominate the two-ion anisotropy field (0.15 meV) and, given that the data scaling is excellent, still remain within the phase transition region. Given that the two-variable scaling analysis systematically incorporates the effects of magnetic fields in the phase transition region, this allows resolving the crossover effect and establishing correct critical exponents in the universality class of the 3D Heisenberg model. Thus the two-variable FSS based on Eq. (2) is essential in the identification of the correct exponents. Also, Fig. 4(f) shows that the data collapse is good through the entire range of magnetic fields. Therefore, in practice, smaller fields may be all that is necessary for the analysis, such as are achievable by a standard laboratory magnetometry.



FIG. 5. The values of (a) β and (b) ν found from zero-field scaling of FePt magnetization curves in zero field [according to Eq. (1)] using data in the range $[T_c^b - \Delta T, T_c^b + \Delta T]$, for increasing values of ΔT . The dashed lines shows a naive extrapolation to $\Delta T = 0$.

FIG. 6. The strength of the interactions of Fe spins based on their separation for in-plane particles (blue) and out-of-plane particles (orange). (a) Strength of the two-ion anisotropy. (b) Strength of the effective exchange.

B. Effect of long-range interactions

The long-range nature of the interaction and two-ion anisotropy energy terms as a potential contributing factor to the crossover behavior can be understood based on the following standard argument [14,15]. If the interaction strength J_{ii} between spins at locations \mathbf{r}_i and \mathbf{r}_i along a given crystallographic direction decays as a function of their separation $r_{ii} = |\mathbf{r}_i - \mathbf{r}_i|$ exponentially fast, i.e., if they can be fitted to a function $f_{\exp}(r_{ij}) = A \exp(-r_{ij}/\tau)$, then the interaction can be considered short range and not contributing to the crossover. Given that the interaction and the two-ion anisotropy profiles in the effective Hamiltonian of FePt in Eq. (4) have directional variability, we apply this argument to in-plane and out-of-plane interaction and anisotropy pairwise coupling distributions. Figure 6(a) shows the plots of in-plane and out-of-plane two-ion anisotropy couplings as a function of the spin-spin separation r_{ii} . The raw data are fitted very well by exponential function f_{exp} , confirming the short-range character of the two-ion anisotropy. Similarly, Fig. 6(b) shows the dependence of in-plane and out-of-plane interaction coupling strength $|J_{ii}|$ versus r_{ii} . The modulus is taken because of the oscillatory character of J_{ij} with increasing spin-spin seperation. Once more, the functional dependence is represented very well by the exponential decay f_{exp} and, thus, the interactions in Eq. (4) can be considered short range. Overall, this effectively rules out the long-range interactions as driving the crossover behavior and suggests that the two-ion anisotropy is the primary contributing factor.

V. CONCLUSIONS

In summary, the present study resolves the issue of the variability of critical exponents that have been reported in earlier research (Table I). The study also shows that magnetic fields can overcome the effects of crossover and that the critical exponents of FePt are fully consistent with those of the three-dimensional Heisenberg model. The discrepancies between the various groups of exponents in Table I can be attributed to the phase transition crossover behavior, which renders the identification of critical exponents highly sensitive and dependent on the details of the analysis. In particular, our study suggests that it is not possible to identify accurate values of critical exponents of FePt based on zero-field magnetization data and, instead, field-dependent data needs to be included in the analysis. The origin of the crossover behavior is in the presence of the two-ion anisotropy term in the effective Hamiltonian of FePt [Eq. (4)], which acts as perturbation to exchange energy and can become relevant in the phase transition region close to T_c . While this crossover effect dominates the zero-field magnetization data, it can be counterbalanced by applying the

external magnetic field. The two-variable FSS technique that is introduced in this paper is required to overcome this crossover effect.

Our study has broad implications for the HAMR technology design. Since significant magnetic fields are used in the writing process [16], it suggests that, in order to quantify the T_c distributions for quality control of the recording media, the Heisenberg critical exponents can be used in the design directly. Being able to use Heisenberg critical exponents directly avoid the need for further independent identification of critical exponents anytime new experimental realizations of FePt-based HAMR media become available. At the fundamental level, our study suggests that the crossover behavior of FePt during phase transitions is governed by the two-ion anisotropy energy, rather than the long-range nature of exchange interactions, and can be circumvented through application of magnetic fields accessible by using standard laboratory magnetometry.

ACKNOWLEDGMENTS

J. W. acknowledges financial support from the EPSRC Centre for Doctoral Training Grant No. EP/L006766/1. This worked used the ARCHER UK National Supercomputing Service (http://www.archer.ac.uk), accessed through the UK HEC Materials Chemistry Consortium, which is funded by EPSRC (EP/L000202). This work also made use of the IRIDIS High Performance Computing Facility and associated support services at the University of Southampton. We also thank Matthew Ellis for providing the parameterization data for the FePt Hamiltonian.

APPENDIX

The method of data collapse that is proposed in this manuscript is an adaptation of the automatic collapse proposed by Bhatacharjee and Seno [11]. A measure is proposed for the success of the collapse of various curves as a function of the critical exponents β and ν , which can be minimized in order to find the optimum values of the critical exponents. Following this method, if the function \tilde{M} is known, then when rescaling a set of points M(t) according to Eq. (1), the total residual r can be given as

$$r = \frac{1}{N} \sum \left| R^{\beta/\nu} M(t, R) - \tilde{M} \left(R^{1/\nu} t \right) \right|.$$
 (A1)

In general however, \tilde{M} is unknown. Instead, interpolated values from another rescaled curve can be used in place of \tilde{M} . If it is assumed that a perfect collapse of several curves is one where the total residual is zero, then the best collapse of an imperfect data set can be considered to be the one that

minimizes r, i.e., minimizing the quantity P_b [11], where

$$P_{b} = \frac{1}{N_{ol}} \left[\sum_{p} \sum_{j \neq p} \sum_{i,ol} \left| R_{j}^{\beta/\nu} M_{j}(t_{i}) - R_{p}^{\beta/\nu} M_{p}^{*}(t_{i}^{*}) \right|^{q} \right]^{1/q},$$
(A2)

where M_j is the set of samples of M associated with system size R_j , M_j^* is an interpolating function of the data set M_j , t_i is the *i*th member of a data set in the variable t, and t_i^* is the value of t_i , rescaled from system size R_p to R_j , i.e., $t_i^* = (R_j/R_p)^{1/\nu}t_i$. The value N_{ol} is equal to the total number of data points that overlap between all pairs of curves with different R after rescaling. Likewise, the sum over the points t_i is only performed for those points where t_i^* is within the extent of the interpolated curve M_p^* . Choosing only these points avoids errors due to extrapolation. The function P_b is minimized through variation of ν , β , and T_c^b , identifying them for that specific material. The value q, introduced in Eq. (A2), should have no effect on the results of the collapse; however, it may affect the convergence of the method in some cases. For this work, q is chosen to be equal to 1.

The errors on critical parameters that are shown in this work are the width of the minimum value of P_b to a given level. This can be calculated as

$$\Delta \beta = \eta \beta \left(2 \ln \frac{P_b(\beta \pm \eta \beta, \nu, T_c^b)}{P_b(\beta, \nu, T_c^b)} \right)^{-1/2},$$

$$\Delta \nu = \eta \nu \left(2 \ln \frac{P_b(\beta, \nu \pm \eta \nu, T_c^b)}{P_b(\beta, \nu, T_c^b)} \right)^{-1/2},$$
 (A3)

$$\Delta T_c^b = \eta T_c^b \left(2 \ln \frac{P_b(\beta, \nu, T_c^b \pm \eta T_c^b)}{P_b(\beta, \nu, T_c^b)} \right)^{-1/2},$$

where $\Delta\beta$, $\Delta\nu$, and ΔT_c^b are the widths of β , ν , and T_c^b to the level of η ; i.e., at the 1% level, $\eta = 0.01$.

A modification to Eq. (A2) is required in order to take into account the scaling with magnetic field strength; however, the minimization principle remains the same. The minimization quantity P_b is redefined so as to take into account the two-variable FSS form given in Eq. (2):

$$P_{b} = \frac{1}{N_{ol}} \left[\sum_{p} \sum_{j \neq p} \sum_{i,ol} \left| R_{j}^{\beta/\nu} M_{j}(\mathbf{x}_{i}) - R_{p}^{\beta/\nu} M_{p}^{*}(\mathbf{x}_{i}^{*}) \right|^{q} \right]^{1/q},$$
(A4)

where \mathbf{x}_i is the set of scaling variables $\{t_i, H_i\}$ for the *i*th data point and \mathbf{x}_i^* is the rescaled data set $\{(R_j/R_p)^{1/\nu}t_i, (R_j/R_p)^{\beta\delta/\nu}H_i\}$. In Eq. (A4), the function M_j and the interpolating function M_p^* are now surfaces rather than curves. The sum over \mathbf{x}_i is performed only for those points where \mathbf{x}_i^* lies within the extent of the interpolated surface M_n^* .

- [1] D. Weller, G. Parker, O. Mosendz, E. Champion, B. Stipe, X. Wang, T. Klemmer, G. Ju, and A. Ajan, A HAMR media technology roadmap to an areal density of 4 Tb/in², IEEE Trans. Magn. 50, 3100108 (2014).
- [2] Unai Atxitia, Denise Hinzke, and Ulrich Nowak, Fundamentals and applications of the Landau–Lifshitz–Bloch equation, J. Phys. D: Appl. Phys. 50, 033003 (2016).
- [3] Amikam Aharoni, *Introduction to the Theory of Ferromagnetism* (Clarendon Press, Oxford, 2000), Vol. 109.
- [4] J. Waters, A. Berger, D. Kramer, H. Fangohr, and O. Hovorka, Identification of Curie temperature distributions in magnetic particulate systems, J. Phys. D: Appl. Phys. 50, 35LT01 (2017).
- [5] H. Oezelt, A. Kovacs, J. Fischbacher, S. Bance, M. Gubbins, and T. Schrefl, Transition jitter in heat-assisted magnetic recording by micromagnetic simulation, IEEE Trans. Magn. 53, 1 (2017).
- [6] H. J. Richter, and G. J. Parker, Temperature dependence of the anisotropy field of L1₀ FePt near the Curie temperature, J. Appl. Phys. **121**, 213902 (2017).
- [7] C.-B. Rong, D. Li, V. Nandwana, N. Poudyal, Y. Ding, Z. L. Wang, H. Zeng, and J. P. Liu, Size-dependent chemical and magnetic ordering in L1₀-FePt nanoparticles, Adv. Mater. 18, 2984 (2006).
- [8] C. Zhou, T. C. Schulthess, and O. N. Mryasov, Magnetic anisotropy of FePt nanoparticles: Temperature-dependent free energy barrier for switching, IEEE Trans. Magn. 43, 2950 (2007).
- [9] O. Hovorka, S. Devos, Q. Coopman, W. J. Fan, C. J. Aas, R. F. L. Evans, Xi Chen, G. Ju, and R. W. Chantrell, The Curie temperature distribution of FePt granular magnetic recording media, Appl. Phys. Lett. 101, 052406 (2012).
- [10] A. Lyberatos, D. Weller, G. J. Parker, and B. C. Stipe, Size dependence of the Curie temperature of L1o-FePt nanoparticles, J. Appl. Phys. **112**, 113915 (2012).
- [11] S. M. Bhattacharjee, and F. Seno, A measure of data collapse for scaling, J. Phys. A: Math. General 34, 6375 (2001).
- [12] O. N. Mryasov, U. Nowak, K. Y. Guslienko, and R. W. Chantrell, Temperature-dependent magnetic properties of FePt: Effective spin Hamiltonian model, Europhys. Lett. 69, 805 (2005).
- [13] J. M. Yeomans, *Statistical Mechanics of Phase Transitions* (Clarendon Press, Oxford, 1992).
- [14] N. Goldenfeld, in Lectures on Phase Transitions and the Renormalization Group, Frontiers in Physics (Addison-Wesley, Advanced Book Program, Reading, Massachusetts, 1992).
- [15] M. E. Fisher, S.-K. Ma, and B. G. Nickel, Critical Exponents for Long-Range Interactions, Phys. Rev. Lett. 29, 917 (1972).
- [16] A. Lyberatos, D. Weller, and G. J. Parker, Switching time in laser pulse heat-assisted magnetic recording using L10-FePt nanoparticles, J. Appl. Phys. 117, 133905 (2015).