Self-organized criticality of magnetic avalanches in disordered ferrimagnetic material

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We observe multiple steplike jumps in a Dy-Fe-Ga-based ferrimagnetic alloy in its magnetic hysteresis curve at 2 K. The observed jumps are found to have a stochastic character with respect to their magnitude and the field position, and the jumps do not correlate with the duration of the field. The distribution of jump size follows a power law variation indicating the scale invariance nature of the jumps. We have invoked a simple two-dimensional random bond Ising-type spin system to model the dynamics. Our computational model can qualitatively reproduce the jumps and their scale-invariant character. It also elucidates that the flipping of antiferromagnetically coupled Dy and Fe clusters is responsible for the observed jumps in the hysteresis loop. These features are described in terms of the self-organized criticality.

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I. INTRODUCTION

In physics and materials science, many systems, when driven by a slowly varying external parameter, can show avalanches in their physical properties. They include flux penetration in type II superconductors [1], Barkhausen noise due to the domain distribution in ferromagnets [2], earthquakes [3], sand piles [4], forest fires [5], and others. One of the important characteristics of these systems is the scale invariance of the avalanche size [6]. The distribution of the jumps shows a power law behavior, $D(s) \sim s^{-\alpha}$, where D(s) is the probability of an avalanche of size *s*, and α is an exponent mostly lying between 1 and 2.

The scale-invariant avalanche dynamics of such systems are often described by the phenomenon of self-organized criticality (SOC) [7–10]. The physical systems showing SOC are generally dissipative and locally interacting. The system organizes itself into self-organized metastable states, which transform from one to another via avalanches. Despite its complexity, the SOC has basic statistical features that are described by power laws [10]. In the case of ferromagnetic systems, the magnetization shows a series of small jumps when slowly driven by a magnetic f,ield and it is interpreted on the basis of SOC. This is called the Barkhausen effect, and it is related to the sudden reversal of the ferromagnetic domains [11].

Recently, a few magnetically phase-separated materials have been reported to show multiple metamagnetic jumps under a varying magnetic field. The systems showing such jumps include various Mn-site doped manganites [12–18], Fe-site

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doped CeFe₂ [19,20], TbFeAl [21], and Gd₅Ge₄ [22,23] and its alloys. These jumps are characteristically different from the Barkhausen noise with a relatively larger variation of M. It is generally believed that these metamagnetic jumps occur due to the field-induced transition of an antiferromagnetic (AFM) cluster to a ferromagnetic (FM) one in those AFM-FM phase-separated systems. The jumps are often found to vary systematically with a sweep rate of H [24,25]. With an increasing sweep rate, the jump shifts to lower fields. This indicates *nonstationary* nature of the jumps. Even jumps are found to occur spontaneously, if one waits at a fixed temperature and magnetic field for a sufficient amount of time [26,27].

The classical theory of SOC manifests that a slowly driven system should have a stationary critical state devoid of any external fine tuning [11]. Therefore, traditionally the metamagnetic jumps in the phase-separated system have not been linked to a classical SOC phenomenon. All the above systems are characterized by strong magneto-elastic coupling, and the jumps are associated with the structural transition. Therefore, the internal strain at the interface of clusters plays an important role in the observed jumps.

The question remains, can there be a system showing large metamagnetic jumps obeying the SOC scenario? In the present work, we chose a relatively simpler system $DyFe_3$, which does not show any structural instability down to 4 K [28]. Here Dy and Fe sublattices are aligned antiparallel giving rise to a ferrimagnetic state. The Zeeman energy is supposed to be high due to the large moments at the Dy and Fe sites, which can facilitate spin flip under an applied magnetic field. We doped nonmagnetic Ga at the Fe site to introduce disorder in it. The magnetization curve of $DyFe_3$ is event-less showing small coercivity and saturation of moment above 15 kOe. On the other hand, the Ga-doped samples show clear ultrasharp metamagnetic jumps. This provides with us an

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FIG. 1. Field-cooled (FC) and zero-field-cooled (ZFC) magnetization (*M*) data as a function of temperature (*T*) measured under 100 Oe of magnetic field for $DyFe_{2.5}Ga_{0.5}$.

opportunity to study avalanches, where structural instability is unlikely to play a major role.

In the present work, we have mostly focused on the sample where 1/6 Fe is replaced by Ga (nominal composition DyFe_{2.5}Ga_{0.5}). The experimental data are supported by our classical Monte Carlo-based simulation of the magnetic hysteresis loop.

II. EXPERIMENTAL DETAILS

Polycrystalline samples of DyFe₃ and DyFe_{2.5}Ga_{0.5} were prepared by a standard argon arc melting technique and subsequent annealing. Structural characterization of the samples was performed by a room temperature powder x-ray diffraction experiment using Cu-K_{α} radiation. We found that the parent DyFe₃ compound crystallizes in a rhombohedral structure, while Ga-doped DyFe_{2.5}Ga_{0.5} has a hexagonal structure (see Fig. 10 in Appendix A). The dc magnetization (*M*) of the samples was measured using a Quantum Design SQUID magnetometer (MPMS3) as well as using the vibrating sample magnetometer of the Quantum Design Physical Properties Measurement System (PPMS). The resistivity (ρ) and the magnetoresistance were measured by a four-point technique using the same PPMS.

III. EXPERIMENTAL RESULTS

A. Magnetization

Figure 1 depicts the temperature (*T*) variation of *M* of DyFe_{2.5}Ga_{0.5} recorded in zero-field-cooled (ZFC) and field-cooled (FC) protocols under 100 Oe of the magnetic field. The ferrimagnetic nature of the undoped DyFe₃ is reported previously by Plusa *et al.* with ordering temperature and compensation point of 615 K and 525 K, respectively [29]. Ga doping in the Fe site is expected to reduce the magnetic ordering temperature. For DyFe_{2.5}Ga_{0.5}, we see large irreversibility between FC and ZFC curves, which extends up



FIG. 2. (a)–(d) Full magnetization loop recorded at different temperatures for $DyFe_{2.5}Ga_{0.5}$. The inset depicts the temperature variation of the coercive field.

to the maximum temperature of measurement (370 K) (i.e., the curves bifurcate from each other from the maximum temperature that we could reach.) This signifies that the magnetic ordering temperature is at least above 370 K. The FC-ZFC irreversibility indicates the presence of disorder in the system. In presence of quenched disorder, pinning of the magnetic clusters or domains is responsible for the bifurcation of the FC-ZFC data [30].

Figures 2(a) and 2(b) show the isothermal variation of M with the magnetic field (H) of DyFe₃ and DyFe_{2.5}Ga_{0.5} samples at 2 K, respectively. The M vs H curve for the undoped DyFe₃ at 2 K rises sharply at low fields and saturates beyond 10 kOe with a coercive field of 920 Oe. The saturation moment is found to be 4.3 μ_B . This moment arises from the antiparallel arrangements of Dy and Fe spins [28,31].

The most striking observation of the present work is found in the 2 K isotherm of DyFe_{2.5}Ga_{0.5} [Fig. 2(b)]. We find successive jumps in M as the field is swept between \pm 70 kOe. The jumps are sharp, and they are present in all five legs (see Appendix B). M eventually saturates above 60 kOe of the field with saturation moment $m_{\text{sat}} = 5.3 \mu_B$. Here m_{sat} is higher than the undoped sample because a part of Fe is replaced by nonmagnetic Ga. Since the total moment is $m_{\text{sat}} =$ $m_{\rm Dy} - m_{\rm Fe}$ (the Fe sublattice is antiparallel to Dy in the ferrimagnetic state), the reduction in Fe content will enhance m_{sat} . The coercive field of the Ga-doped sample is much higher $(\sim 7.5 \text{ kOe})$ than the parent one. Notably, the occurrence of multiple jumps vanishes as the temperature is slightly increased. The isotherm at 5 K [Fig. 2(c)] shows a smooth variation of M with H. A similar smooth isotherm is also observed at 75 K [Fig. 2(d)], albeit with the much lower value of the coercive field. The inset of Fig. 2(d) shows the temperature variation of the coercive field (H_{coer}) recorded between 2 and 75 K. H_{coer} shows a nonmonotonous variation with T with a peak around 7 K. In disordered granular systems, such a kind of variation of H_{coer} is common [32].





FIG. 3. (a) The fifth leg (0 to 70 kOe in the first quadrant, after returning from -70 kOe) of the magnetization loop recorded at 2 K for different sweep rates of the magnetic field (*H*). (b) The full loop at 2 K after the sample's being cooled in different magnetic fields. Inset of (b) depicts the fifth leg of the hysteresis loop recorded at 2 K for different cycles.

We also recorded *M*-*H* curves for DyFe_{2.5}Ga_{0.5} at different sweep rates ($\dot{H} = 100, 70, 60, 48, \text{ and } 45 \text{ Oe s}^{-1}$) in the first quadrant, after the sample being returned from -70 kOe of the field [Fig. 3(a)]. It is interesting to note that the position of the jumps does not vary systematically with the sweep rate. Even for several consecutive runs at $\dot{H} = 60 \text{ Oe s}^{-1}$, the jump fields are markedly different [see the inset of Fig. 3(b)]. Nevertheless, the observed jumps follow a certain pattern for all values of \dot{H} , and this is illustrated by the run at $\dot{H} = 70 \text{ Oe s}^{-1}$ [see the inset of Fig. 3(a)]. There is a common pattern in the jumps denoted by AA', BB', CC', DD',..., with AA' < BB', $BB' > CC', CC' \sim DD'$. Although the positions, A, B, C, D, ... are random on the H axis, they occur in the same order and arrangement. Unlike previously reported multiple metamagnetic jumps [24-27], DyFe_{2 5}Ga_{0.5} does not show a systematic sweep rate dependence.

We also studied the effect of the cooling field (H_{cool}) on the hysteresis loop. For that purpose, the *M*-*H* isotherm is recorded at 2 K, after the sample being cooled from 300 K

FIG. 4. Magnetization as a function of time for different protocols measured at 2 K after the application of a magnetic field of (a) 12 kOe and (b) 13 kOe. Before the measurement, the sample was first cooled to 2 K, and it was kept at this temperature for the rest of the measurements.

under a certain H_{cool} . The shift in the hysteresis loop on field cooling is called the exchange bias effect, and it is expressed in terms of $H_{eb} = (H_+ + H_-)/2$ where H₊ and H₋ stand for the positive and negative intercepts of the magnetization curve with the field axis, respectively. For DyFe_{2.5}Ga_{0.5}, we see an increase of the magnitude of H_- , keeping H_+ (loop spreads in the negative side of H asymmetrically). It is difficult to say whether such a shift arises from exchange bias or the shift of a jump. The jumps maintain the same pattern, but their height and magnitude vary randomly with H_{cool} .

Temporal effects are often associated with magnetization jumps. It is found that the jump can spontaneously occur if one waits for sufficient time at a point on the *M*-*H* loop at a field slightly lower than the avalanche field for a particular jump [26,27]. We studied this phenomenon by noting the variation of *M* with time (*t*) in DyFe_{2.5}Ga_{0.5}. In Fig. 4(a) we have shown the relaxation (*M* vs *t* data) after applying $H_s = 12$ kOe of field (H_s slightly lower than the field required for BB' jump). However, no spontaneous jump is seen even after waiting for 180 minutes, though *M* changes by 8% in 40 minutes. In



FIG. 5. Panels (a) and (b), respectively, show the variation of electrical resistivity (ρ) as a function of the field at 2 and 7 K. The inset of (a) represents the variation of electrical resistivity as a function of temperature. The inset of (b) shows the variation of ρ as a function of *H* at 125 K.

Fig. 4(b) measurement was performed at $H_s = 13$ kOe, where the BB' jump already took place. Interestingly, *M* drops with *t*, and a 5% relaxation is seen.

Multiple metamagnetic jumps are common among glassy magnetic systems [25,33,34]. To confirm the glassy nature in DyFe_{2.5}Ga_{0.5}, we measured the filled-cooled-field-stop memory effect (not shown here). The sample was cooled from 300 K under H = 100 Oe with intermediate zero-field stops at T_i (= 200, 150, 100, and 50 K) for 60 minutes. On reaching 2 K, the sample was heated back to 300 K in the presence of H = 100 Oe. However, no feature is observed at the stopping temperatures T_i during heating. This rules out the possibility of a glassy magnetic state in the system [35,36].

B. Resistivity

DyFe_{2.5}Ga_{0.5} shows metallic behavior in the *T*-dependent plot of resistivity (ρ) [see inset of Fig. 5(a)]. In Figs. 5(a) and 5(b) we have plotted the *H* variation of ρ measured at 2 and 7 K, respectively. In $\rho(H)$ data at 2 K, the ultrasharp jumps are also present, and they approximately correspond to the similar avalanche fields for the jump as observed in the magnetization data. Similar to the M(H) data, the jumps are absent at a higher temperature of 7 K. In both 2 and 7 K data, the full five quadrants $\rho(H)$ curves (between \pm 50 kOe) form a hysteresis loop. Such observation indicates that the electronic property of DyFe_{2.5}Ga_{0.5} is intimately correlated with the magnetic state of the system. At 125 K [inset of Fig. 5(b)], there is no loop, and $\rho(H)$ saturates above 10 kOe showing small magnetoresistance.

C. Distribution of the jumps observed in experiment

In Fig. 6(a) we have plotted the histogram of the jumps using our magnetization data depicted in Fig. 3(a). We have considered 560 jumps from 34 *M*-*H* loops. Since we do not find any correlation between the sweep rate and the jump size, we used data from all sweep rates to construct the histogram. We have used Sturge's rule ($\kappa = 1 + 3.22 \log \nu$, where ν is the total number of data points) to calculate the number of bins, κ [37]. Clearly the smaller jumps (ΔM is low) are larger in number. In Fig. 6(b) we have plotted the number of occurrences $N(\Delta M)$ as a function of jump size. This provided a power law distribution, $N \sim \Delta M^{-\alpha}$, with $\alpha = 1 \pm 0.1$.

In Fig. 6(c) we have studied the variation of jump size with the *H*. It is seen that the bigger jumps are found around 10 kOe, and the jump size is smaller at higher fields [Fig. 6(c)]. This is understandable from the isotherms recorded at 2 K. In Fig. 6(d) we have plotted the jump number as a function of jump size. We took 12 five-leg *M*-*H* loops measured at 2 K and serially counted the jumps for all the loops. A jump number is assigned to each jump. We find that the jump size span across multiple ranges, e.g., big jumps (4–5 $\mu_B/f.u.$), medium jumps (1–2 $\mu_B/f.u.$), and small jumps (below 0.25 $\mu_B/f.u.$), and the data correlate with the characteristics of Fig. 6(b). Here f.u. stands for formula unit.

IV. THEORETICAL MODEL

A vast majority of real-world materials contain impurities that introduce disorder in the system [38,39]. For such systems, the central challenge is to predict large-scale behavior from local dynamics. Developing a corresponding model of interacting spins originates from the inspection of the experimental data. The experimental observations of magnetization jumps in the Dy-Fe-Ga alloys are found to be scale invariant and stationary [11,40,41]. This prompted us to model the system for quantitative analysis of the hysteresis loops. We simulated the jumps with changing magnetic fields and quantify their distribution.

We choose the generalized random-bond Ising model. The Hamiltonian is defined as

$$\mathcal{H}(h) = -\sum_{i,j} \mathcal{J}_{ij} \sigma_i \sigma_j - h \sum_i \sigma_i - h \sum_{i \in Dy} L_i^{Dy}, \qquad (1)$$

where $\sigma_i = \pm 1$ are the local spin moment, \mathcal{J}_{ij} are the nearestneighbor interactions, and *h* is the external field. Two different types of spins corresponding to Fe and Dy are assumed on a square lattice in 3:1 ratio, respectively. The spin value σ_i is taken as unity for both Dy and Fe because their spin moments are almost equal [42]. The direction of the external magnetic field is fixed along one of the spin variables



FIG. 6. (a) Histogram of the observed jumps in the experimental magnetization vs field data. (b) Number of occurrences as a function of jump size in log-log scale. (c) Jump size as a function of the applied field. (d) Individual jumps are plotted as a function of the jump number.

(e.g., negative (-ve) or positive (+ve)). $L_i^{Dy} = 1$ (see Appendix C) is the orbital moment for Dy alone, directed along the Dy spin moment, and contributes to the Hamiltonian at specific lattice sites occupied by Dy atoms. The nearestneighbor interactions in the first term of Eq. (1) can be negative or positive (i.e., AFM or FM, respectively) depending upon the type of interacting particles. Ferromagnetic interaction between two Fe is given by FM coupling $\mathcal{J}_{ij} = 1$, while between two Dy it is $\mathcal{J}_{ij} = 0.05$ (see Fig. 7). Interaction between Fe and Dy is AFM and coupling constant $\mathcal{J}_{ij} = -2$. The \mathcal{J}_{ij} values are obtained from the relative strength of the magnetic interaction of the site disorder due to the doping of nonmagnetic Ga at the expense of Fe. For the experimentally studied samples DyFe₃ and DyFe_{2.5}Ga_{0.5}, f_{Ga} is 0 and 0.2, respectively.

V. SIMULATION DETAILS

An ensemble of Ising spins is considered on a twodimensional (2D) square lattice. Considering a layered structure of the real system (see Appendix A), a 2D modelbased calculation is a reasonable approximation [45,46]. Initially, the system is prepared with Fe and Dy randomly distributed at a 3:1 ratio on a square lattice of linear size



FIG. 7. Model schematic shows a lattice structure in two dimensions with Fe and Dy randomly distributed in a 3:1 ratio. Identical and nonidentical atoms are linked via ferromagnetic and antiferromagnetic interactions, respectively. Disorder is introduced by nonmagnetic Ga replacing Fe according to fraction f_{Ga} .

 \mathcal{L} represented by the model described in Fig. 7. The disorder is introduced by replacing randomly chosen Fe sites by Ga with fraction f_{Ga} . The system is then equilibrated using Glauber dynamics, which is a computer simulation of the Ising model (a magnetism model). Using Monte Carlo simulation without the external field, at T = 0, only the sign of the energy differences is required for the Glauber dynamics. We have simulated this system with the periodic boundary condition. The initial configuration is random, and single spin flip Glauber dynamics has been used for subsequent updating, i.e., a spin is picked up at random and flipped if the resulting configuration has lower energy, never flipped if the energy is raised, and flipped with probability 1/2 if there is no change in energy on flipping. Next, a small external magnetic field (h)is applied and the system is reequilibrated. The external field is raised slowly and the system is equilibrated for every value of h. Magnetization (m) is computed as a function of h by summing over the spins. The external field is increased until a saturation in the magnetization is observed. The process is repeated by decreasing h. The simulation yields a single hysteresis loop as a function of model parameters, $(f_{Ga}, h, \mathcal{J}_{ij})$. Note that the notations, values representing the external magnetic field and magnetization in the simulation, are to mark a quantitative difference with the experiment. We emphasize in our experimental observation that the pure sample without any random substitution by Ga atoms does not show any jumps in the magnetization data that may correspond to the self-organized criticality. The disorder is expected to play a crucial role in the observed magnetization jumps, as the jumps are present only when disordered is introduced in the stoichiometric DyFe₃ sample. Such multiple jumps are attributed to disorder in some other systems in the literature [19,26]. In the present calculation, the disorder is introduced by allowing the atoms to occupy the lattice sites randomly.

VI. NUMERICAL RESULTS

The initial system is prepared with zero external magnetic field, and then it is subjected to gradually increasing field h in step of 0.02. For each field value, the system is equilibrated up to 2×10^5 Monte Carlo steps. Figure 8(a) shows the simulated hysteresis loop for $f_{Ga} = 0.2$, which clearly demonstrates multiple steps in magnetization. The data show intermittent jumps ($\Delta m > 0$) and stationary phases ($\Delta m = 0$) while the external field is increased. A similar characteristic is observed when the external magnetic field is decreased. Figure 8(b)compares the magnetization profile for three different disorder fractions $f_{\text{Ga}} = 0$, $f_{\text{Ga}} = 0.1$, $f_{\text{Ga}} = 0.2$, and $f_{\text{Ga}} = 0.3$. Jumps observed in the magnetization are a signature of frustration in spin-spin interaction, usually characteristics of a glassy magnetic system [47–49]. It is to be noted that jumps are present even for $f_{Ga} = 0$, where there is no doping at the Fe sites. This is because the present model is inherently disordered due to the random occupancy of Fe and Ga in a 3:1 ratio. The nature of the jumps remains nearly unchanged with f_{Ga} .

Snapshots at different field strengths are shown in Figs. 8(c)-8(e) and Figs. 12(a)-12(d) in Appendix D for the $f_{Ga} = 0.2$ lattice. At h = 0, we find the finite clusters of Dy and Fe with up (+) and down(-) spin configurations. With increasing h (i.e., in the + direction), the clusters flip in the

TABLE I. List of exponents for various avalanche models.

Avalanche models	Exponent α	Reference
DyFe _{2.5} Ga _{0.5}	1.1 (experiment) 1.0 (simulation)	This study
BTW sand pile	1.0 (2D)	[8]
Superconducting vortex avalanches	1.4–2	[1]
Barkhausen noise spectra	2	[51]
Forest fire model	2	[5]
Avalanches in lung inflation	1.8	[52]
Microfracturing process	1.3	[53]

direction of the applied field, which produces jumps in the hysteresis curve.

A statistical analysis of the jump size Δm estimated from Fig. 8(a) projects the occurrence histogram of the avalanche sizes for $f_{Ga} = 0.2$ in Fig. 9(a). The data points represent an average of more than 200 samples. A comparison of this distribution with respect to the zero disorder scenario ($f_{Ga} = 0$) shows that for both cases the probability distribution of Δm for very small Δm appears to be significantly higher than large Δm [see Figs. 13(a) and 13(b) in Appendix D]. Moreover, the histogram stretches to larger Δm for the zero disorder.

The log-log plot of the distribution of magnetization $N(\Delta m)$ for disorder fraction $f_{Ga} = 0.2$ [as in Fig. 9(a)] is shown in Fig. 9(b) for four different system sizes $\mathcal{L} =$ 16, 32, 64, and 128. The graph indicates a power law distribution of $N \sim \Delta m^{-\alpha}$. Fitting the data points with the given expression yields the exponent $\alpha = 1 \pm 0.05$ (see Table I). This value of α is found to be close to the experiment [see Fig. 6(b)]. The avalanche sizes (Δm) of a single hysteresis loop are depicted in Fig. 9(c). The jump sizes obtained from the hysteresis loop are plotted sequentially showing multiple regimes corresponding to small, intermediate, and large jumps of magnetization. The data signify that the size of domain flips (avalanche) due to a marginal and systematic change in the external magnetic field is likely uncorrelated as is obvious from the power law distribution in Fig. 6(b). We have further shown Δm as a function of h. It is clear that for low fields, the magnetization jump is substantial. Due to the AFM interaction between Fe and Dy, two adjacent domains remain frustrated and merge to form a larger domain, lowering the net surface energy as the external field is increased. In contrast, jumps are small in a large field. Because the majority of the adjacent domains have already flipped along the field direction, so, with a large field, only a few small clusters or individual spins remain to flip along the field direction until saturation magnetization is reached.

VII. DISCUSSION

Our combined experimental and theoretical simulation reveal that the magnetic avalanches found in the Dy-Fe-Ga alloy are a manifestation of self-organized criticality. Experimentally, the magnetization jumps in the studied alloy are stochastic, do not show any systematic change with the change in the rate of the external driving parameter (here magnetic field), and most importantly, are scale-invariant following a power law distribution. The Monte Carlo simulation of a



FIG. 8. (a) Hysteresis loop of random-bond Ising model for disorder fraction $f_{Ga} = 0.2$ at $\mathcal{L} = 64$. Inset: Many small jumps occur at a large field in the hysteresis loops. (b) Magnetization profiles for four distinct disorder fractions $f_{Ga} = 0, 0.1, 0.2$, and 0.3 are illustrated when the external field (*h*) is reduced. (c)–(e) Snapshots of spin configurations before and after large jumps in the magnetization at field strengths of h = 1.5, 2.5, and h = 5 (as in a), respectively. Red, brown, green, yellow, and blue represent Fe(+), Fe(-), Dy(+), Dy(-), and nonmagnetic Ga, respectively. The magnetic field (*h*) is applied in the +*ve* direction. The snapshots are magnified in the middle to provide a detailed insight into cluster flipping.

model 2D analog of the real material also supports our experimental data, where the change in the magnetization appears to be discontinuous as a function of the external field and obeys a power law distribution.

Numerous pieces of evidence suggest the existence of power law distribution for natural events. The power law exponents of some avalanche models are shown in Table I. Remarkably, in some cases, the exponents of the distribution are the same for systems with very different elemental interactions [50]. SOC is attributed to the evolution of a complex system towards criticality in the presence of local interaction. For the Dy-Fe-Ga alloy, the local interaction is the magnetic correlations between Dy and Fe atoms. The exponent obtained from our analysis closely compares with the BTW sandpile model.

In $DyFe_3$, Dy and Fe spins show AFM correlation, and they are aligned antiparallel resulting in ferrimagnetism. Doping by some amount of nonmagnetic element Ga in Fe site introduces disorder in the Fe sublattice. The finite jumps in the magnetization data indicate the flipping of magnetic clusters than the individual spins. Because of the disorder, the system is characterized by the coexistence of spontaneously ordered but oppositely oriented neighboring magnetic domains of Dy and Fe, respectively. Because of the AFM interaction between Fe and Dy, two such adjacent domains remain frustrated and cannot merge to become a larger domain lowering the net surface energy. To grow a larger domain, one of the oppositely oriented domains must be flipped entirely, which can be achieved by the external magnetic field. However, due to frustration in the local interaction, a sufficiently strong external magnetic field equivalent to the surface area of the domain must be provided. Therefore, a slight change in the external magnetic field often does not alter the overall magnetization of the system. As a result, we observe a staircase-like feature in the magnetization isotherms both in the experiment (Fig. 3) and in simulation (Fig. 8).



FIG. 9. (a) Distribution of magnetization jumps showing occurrence (N) as a function of Δm for disorder fraction $f_{Ga} = 0.2$ at system size $\mathcal{L} = 32$. The data points correspond to an average of over 200 samples. (b) The power law distribution of $N(\Delta m)$ vs Δm decays with exponent $\alpha = 1 \pm 0.05$. (c) Magnetization jump sizes (Δm) of a single hysteresis loop demonstrate jump sizes ranging from small to large. (d) Jump size is plotted as a function of the external field *h*.

In Figs. 8(c) and 8(d) snapshots of the spin clusters are shown before and after the jumps respectively. At vanishing external fields, Dy and Fe clusters assume both +ve and -ve orientations and remain frustrated due to antiferromagnetic interaction [see Fig. 12(a) of Appendix A]. In this scenario, one recognizes that Dy(+) spins are surrounded by Fe(-) and vice versa. At the other extreme, when the external field is increased substantially in the + direction, several Fe(-) clusters flip toward the direction of the applied field [Fe(-) \rightarrow Fe(+)] producing a large jump in the magnetization. This is evident from the increase of red domains of Fe(+) spins at h = 5. In this case, the Zeeman energy overcomes the frustration due to the AFM coupling present between Dy and Fe clusters. Interestingly, a contrasting spin arrangement



FIG. 10. Panels (a) and (b) show powder x-ray diffraction (PXRD) patterns of $DyFe_3$ and $DyFe_{2.5}Ga_{0.5}$ samples, respectively. The insets show the perspective view of the crystal structures.



FIG. 11. Isothermal field variation of magnetization at 5 K for the compound $DyFe_{2.5}Ga_{0.5}$. All five legs are highlighted.

occurs at an intermediate field. The enlarged section of the snapshots in Figs. 8(c) and 8(d) indicates some Dy(-) clusters flip to Dy(+) as the h is increased from 1.5 to 2.5. This is accompanied by the flipping of Fe(+) to Fe(-) adjacent to the Dy. Such an arrangement is energetically favorable due to the large moment of Dy compared to Fe. The outcome is manifested by smaller jumps in the magnetization. The flipping events connect nearby domains of like spins. A cascade of spin-flip and domain rearrangement leads to the formation of bigger clusters when the Zeeman energy overcomes the interfacial AFM interaction between Fe and Dy clusters. Flipping of larger domains then gives rise to larger jumps in the magnetization for a small increase in the external magnetic field. At a very large external field, the flipping events are dominated by the external field, and the Fe(-) spins flip back to Fe(+) [enlarged part of Fig. 8(e)] leading to the saturation of magnetization. Note that, considering a purely antiparallel arrangement of Dy and Fe moments, the expected moment should be 4.30 $\mu_B/f.u.$ in DyFe_{2.5}Ga_{0.5}. However, our experimental value of the saturation moment is $5.25 \,\mu_B/f.u.$ Such discrepancy is likely due to the parallel arrangement of some Dy and Fe clusters, which takes place through jumps.

In conclusion, the magnetization jumps in the Dy-Fe-Ga compound are found to be a spectacular manifestation of self-organized criticality. Our work indicates the flipping of the finite domains of Fe and Dy sublattices in the otherwise antiferromagnetically coupled spin system. The theoretical analysis can broadly reproduce the experimental results. It should be kept in mind that the present computational analysis is a simplified approach, and predictions are qualitative. We considered a 2D system and random distribution of the spins. The experiment, however, is carried out on a 3D design. Moreover, the arrangement of the atoms is not random in the actual system. Therefore, the computational model devoid of the crystal structure of DyFe₃ contains an intrinsic disorder, unlike the real systems. A computational model on a larger system with a 3D crystal lattice can help us better understand the phenomenon.



FIG. 12. Snapshots of the spin configurations Fe(+), Fe(-), Dy(+), Dy(-), and nonmagnetic Ga is shown with increasing external field *h* for disorder fraction $f_{Ga} = 0.2$ at $\mathcal{L} = 64$.

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APPENDIX A: CRYSTAL STRUCTURE

The x-ray diffraction patterns of the two samples are shown in Fig. 10. The undoped sample DyFe₃ crystallizes in a PuNi₃-type rhombohedral structure [inset of Fig. 10(a)] with space group $R\bar{3}m$. In the literature, such structures are often represented as a hexagonal equivalent, i.e., $a = b \neq c$, $\alpha = \beta = 90^\circ$, $\gamma = 120^\circ$ [54–56]. From our refinements, we get a = 5.125 Å and c = 24.575 Å. On the other hand, the Gadoped sample DyFe_{2.5}Ga_{0.5} assumes a CeNi₃-type hexagonal structure with space group $P6_3/mmc$, where the refined lattice parameters are found to be a = 5.165 Å and c = 16.560 Å [57]. The structure consists of hexagonal layers in the *a-b* plane of the crystal as shown in the inset of Fig. 10(b).

APPENDIX B: M-H FIVE-LEG MEASUREMENT

We have shown the significance of five legs in the *M*-*H* curve in Fig. 11. Here five legs in the *M*-*H* curve, respectively, signify the following protocol of field ramping: (1) $0 \rightarrow H_{\text{max}}$ (virgin line), (2) $H_{\text{max}} \rightarrow 0$, (3) $0 \rightarrow -H_{\text{max}}$, (4) $-H_{\text{max}} \rightarrow 0$, and (5) $\rightarrow H_{\text{max}}$, where H_{max} is the maximum value of the field in the loop.

APPENDIX C: MAGNETIC MODELING

In the case of Fe, there are six electrons in the 3*d* orbital, so the total spin is $S_{\text{Fe}} = 2$, while Dy has a total spin of $S_{\text{Dy}} = 5/2$. Since the values are close, we considered the Fe and Dy spins are equal and assigned the reduced spin value $\sigma_i = \pm 1$ for both elements. The Zeeman energy in the magnetic field *h* for Dy is given by $\mathcal{H}_Z = -(\mathbf{L} + g_0 \mathbf{S})$ [42]. Here $\mathbf{L} = \text{Dy}$ orbital moment, $\mathbf{S} = \text{Dy}$ spin moment, and $g_0 = 2$ is the Landé-*g* factor for spin. We know for Dy, L = 5 and S = 5/2, and due to g_0 , their contribution are the same in the Zeeman term. Therefore, in the Hamiltonian, we can take the reduced value of effective orbital contribution to be unity $(L_i^{Dy} = 1)$.

APPENDIX D: SIMULATION SNAPS AND PROBABILITY DISTRIBUTION OF JUMP SIZES

We show steady-state snapshots at different field strengths for the $f_{Ga} = 0.2$ in Figs. 12(a)–12(d). Figures 13(a) and 13(b) show the probability distribution of jump sizes (normalized occurrence of the jump size as shown in Fig. 9) with respect to the zero disorder scenario ($f_{Ga} = 0$) and the finite disorder scenario ($f_{Ga} = 0.2$).



FIG. 13. Probability distributions of magnetization jump size derived from the hysteresis loop at system size $\mathcal{L} = 64$ with disorder fraction (a) $f_{Ga} = 0$, and (b) 0.2, respectively. The intermittent bumps in the histogram plot are due to the lack of enough sample averages. In Fig. 9 we have calculated the standard deviation which shows a larger errorbar for intermittent bump.

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