

Dynamic phase transitions in the presence of quenched randomnessErol Vatansever¹ and Nikolaos G. Fytas²¹*Department of Physics, Dokuz Eylül University, TR-35160 Izmir, Turkey*²*Applied Mathematics Research Centre, Coventry University, Coventry CV1 5FB, United Kingdom*

(Received 8 May 2018; published 25 June 2018)

We present an extensive study of the effects of quenched disorder on the dynamic phase transitions of kinetic spin models in two dimensions. We undertake a numerical experiment performing Monte Carlo simulations of the square-lattice random-bond Ising and Blume-Capel models under a periodically oscillating magnetic field. For the case of the Blume-Capel model we analyze the universality principles of the dynamic disordered-induced continuous transition at the low-temperature regime of the phase diagram. A detailed finite-size scaling analysis indicates that both nonequilibrium phase transitions belong to the universality class of the corresponding equilibrium random Ising model.

DOI: [10.1103/PhysRevE.97.062146](https://doi.org/10.1103/PhysRevE.97.062146)**I. INTRODUCTION**

In the past 60 years our understanding of equilibrium critical phenomena has developed to a point where well-established results are available for a wide variety of systems. In particular, the origin and/or the difference between equilibrium universality classes is by now well understood. This observation also partially holds for systems under the presence of quenched disorder. However, far less is known for the physical mechanisms underlying the nonequilibrium phase transitions of many-body interacting systems that are far from equilibrium and clearly a general classification of nonequilibrium phase transitions into nonequilibrium universality classes is missing.

We know today that, when a ferromagnetic system, below its Curie temperature, is exposed to a time-dependent oscillating magnetic field, it may exhibit a fascinating dynamic magnetic behavior [1]. In a typical ferromagnetic system being subjected to an oscillating magnetic field, there occurs a competition between the time scales of the applied-field period and the metastable lifetime τ of the system. When the period of the external field is selected to be smaller than τ , the time-dependent magnetization tends to oscillate around a nonzero value, which corresponds to the dynamically ordered phase. In this region, the time-dependent magnetization is not capable of following the external field instantaneously. However, for larger values of the period of the external field, the system is given enough time to follow the external field, and in this case the time-dependent magnetization oscillates around its zero value, indicating a dynamically disordered phase. When the period of the external field becomes comparable to τ , a dynamic phase transition takes place between the dynamically ordered and the disordered phases.

Throughout the years, there have been several theoretical [2–22] and experimental studies [23–27] dealing with dynamic phase transitions as well as with the hysteresis properties of magnetic materials. The main conclusion emerging is that both the amplitude and the period of the time-dependent magnetic field play a key role in dynamical critical phenomena (in addition to the usual temperature parameter). Furthermore,

the characterization of universality classes in spin models driven by a time-dependent oscillating magnetic field has also attracted a lot of interest lately [28–37]. Some of the main results are listed below:

(1) The critical exponents of the kinetic Ising model were found to be compatible with those of the equilibrium Ising model at both two dimensions (2D) and three dimensions (3D) [28–30,32,36].

(2) Buendía and Rikvold using soft Glauber dynamics estimated the critical exponents of the 2D Ising model and provided strong evidence that the characteristics of the dynamic phase transition are universal with respect to the choice of the stochastic dynamics [31].

(3) The role of surfaces at the nonequilibrium phase transitions in Ising models has been elucidated by Park and Pleimling: The nonequilibrium surface exponents were found to be different than the equilibrium critical surface ones [33].

(4) Experimental evidence by Riego *et al.* [27] and numerical results by Buendía and Rikvold [35] verified that the equivalence of the dynamic phase transition to an equilibrium phase transition is limited to the area near the critical period and for zero bias.

(5) Numerical simulations by Vatansever and Fytas showed that the nonequilibrium phase transition of the spin-1 Blume-Capel model belongs to the universality class of the equilibrium Ising counterpart (at both 2D and 3D) [37]. General and very useful features of the dynamic phase transition of the Blume-Capel model can also be found in Refs. [8,16,17,19,38,39].

The above results in 2D and 3D kinetic Ising and Blume-Capel models establish a mapping between the universality principles of the equilibrium and dynamic phase transitions of spin-1/2 and spin-1 models. They also provide additional support in favor of an earlier investigation of a Ginzburg-Landau model with a periodically changing field [10] as well as with the symmetry-based arguments of Grinstein *et al.* in nonequilibrium critical phenomena [40].

Motivated by the current literature, in the present paper we attempt to shed some light on the effect of quenched disorder on dynamic phase transitions. With the exception

of a few mean-field and effective-field theory treatments of the problem [41–46], we attempt to show this here. However, what we have mainly learned from the previous studies on the topic is that the dynamic character of a typical magnetic system driven by a time-dependent magnetic field sensitively depends on the amount of disorder, accounting for reentrant phenomena and dynamic tricritical points [43]. In the current paper we use as test-case platforms for our numerical experiment the Ising and Blume-Capel models on the square lattice under a time-dependent magnetic field, diffusing disorder in the ferromagnetic exchange interactions. For the case of the Blume-Capel model we focus on the disordered-induced continuous dynamic transition at the low-temperature regime of the phase diagram. In a nutshell, our results indicate that the dynamic phase transitions of both the random-bond Ising and the Blume-Capel models belong to the universality class of the equilibrium random Ising model.

The outline of the remaining parts of the paper is as follows: In Sec. II we introduce the disordered versions of the Ising and Blume-Capel models and in Sec. III the thermodynamic observables necessary for the application of the finite-size scaling analysis. The details of our simulation protocol are given in Sec. IV and the numerical results and discussion in Sec. V. Finally, Sec. VI contains a summary of our conclusions.

II. MODELS

We consider the square-lattice random-bond Ising and Blume-Capel (BC) models under a time-dependent oscillating magnetic field, described by the following Hamiltonians:

$$\mathcal{H}^{(\text{Ising})} = - \sum_{\langle xy \rangle} J_{xy} \sigma_x \sigma_y - h(t) \sum_x \sigma_x, \quad (1)$$

and

$$\mathcal{H}^{(\text{BC})} = - \sum_{\langle xy \rangle} J_{xy} \sigma_x \sigma_y + \Delta \sum_x \sigma_x^2 - h(t) \sum_x \sigma_x. \quad (2)$$

In the above Eqs. (1) and (2) $\langle xy \rangle$ indicate summation over nearest neighbors, and the spin variable σ_x takes on the values of $\{\pm 1\}$ for the Ising model and $\{-1, 0, +1\}$ for the Blume-Capel model, respectively. The couplings $J_{xy} > 0$ denote the random ferromagnetic exchange interactions, drawn from a bimodal distribution of the form

$$\mathcal{P}(J_{xy}) = \frac{1}{2} [\delta(J_{xy} - J_1) + \delta(J_{xy} - J_2)]. \quad (3)$$

Following Refs. [47–49], we choose $J_1 + J_2 = 2$ and $J_1 > J_2 > 0$ so that $r = J_2/J_1$ defines the disorder strength; for $r = 1$ the pure systems are recovered. A clear advantage of using the bimodal distribution (3) is that the critical temperature T_c of the random Ising model is exactly known from duality relations as a function of the disorder strength r via [50,51]

$$\sinh(2J_1/T_c) \sinh(2rJ_1/T_c) = 1. \quad (4)$$

For the case of the Blume-Capel Hamiltonian (2) Δ denotes the crystal-field coupling that controls the density of vacancies ($\sigma_x = 0$). For $\Delta \rightarrow -\infty$ vacancies are suppressed, and the model becomes equivalent to the Ising model. Finally, the term $h(t)$ corresponds to a spatially uniform periodically oscillating magnetic field so that all lattice sites are exposed to a square-

wave magnetic field with amplitude h_0 and half-period $t_{1/2}$ [30–32].

A brief description of the Blume-Capel model's phase diagram together with some necessary pinpoints of the current literature with respect to the effect of disorder on its critical behavior may be useful here: The phase diagram of the equilibrium pure Blume-Capel model in the crystal-field—temperature plane consists of a boundary that separates the ferromagnetic from the paramagnetic phase. The ferromagnetic phase is characterized by an ordered alignment of ± 1 spins. On the other hand, the paramagnetic phase can be either a completely disordered arrangement at high temperatures or a ± 1 -spin gas in a 0-spin dominated environment for low temperatures and high crystal fields. At high temperatures and low crystal fields, the ferromagnetic-paramagnetic transition is a continuous phase transition in the Ising universality class, whereas at low temperatures and high crystal fields the transition is of first-order character [52,53]. The model is thus a classic and paradigmatic example of a system with a tricritical point $[\Delta_t, T_t]$ [54] where the two segments of the phase boundary meet. A detailed reproduction of the phase diagram of the model can be found in Ref. [55], and an accurate estimation of the location of the tricritical point has been given in Ref. [56]: $[\Delta_t, T_t] = [1.9660(1), 0.6080(1)]$. A lot of work has been also devoted in understanding the effects of quenched bond randomness on the universality aspects of the Blume-Capel model especially in two dimensions where any infinitesimal amount of disorder drives the first-order transition at the low-temperature regime to a continuous transition. Quantitative phase diagrams of the random-bond Blume-Capel model at equilibrium have been constructed in Refs. [47,48] and, more recently, a dedicated numerical study at the first-order transition regime revealed that the induced under disorder continuous transition belongs to the universality class of the random Ising model with logarithmic corrections [57].

III. OBSERVABLES

In order to determine the universality aspects of the kinetic random-bond Ising and Blume-Capel models, we will consider the half-period dependencies of various thermodynamic observables. The main quantity of interest is the period-averaged magnetization,

$$Q = \frac{1}{2t_{1/2}} \oint M(t) dt, \quad (5)$$

where the integration is performed over one cycle of the oscillating field. Given that for finite systems in the dynamically ordered phase the probability density of Q becomes bimodal, one has to measure the average norm of Q in order to capture symmetry breaking so that $\langle |Q| \rangle$ defines the dynamic order parameter of the system. In the above Eq. (5), $M(t)$ is the time-dependent magnetization per site,

$$M(t) = \frac{1}{N} \sum_{x=1}^N \sigma_x(t), \quad (6)$$

where $N = L \times L$ defines the total number of spins and L defines the linear dimension of the lattice.

To characterize and quantify the transition using finite-size scaling arguments we must also define quantities analogous to the susceptibility in equilibrium systems. The scaled variance of the dynamic order parameter,

$$\chi_L^Q = N[\langle Q^2 \rangle_L - \langle |Q| \rangle_L^2] \quad (7)$$

has been suggested as a proxy for the nonequilibrium susceptibility also theoretically justified via fluctuation-dissipation relations [18]. Similarly, one may also measure the scaled variance of the period-averaged energy,

$$\chi_L^E = N[\langle E^2 \rangle_L - \langle E \rangle_L^2], \quad (8)$$

so that χ_L^E can be considered as the corresponding heat capacity. Here E denotes the cycle-averaged energy corresponding to the cooperative part of the Hamiltonians (1) and (2). With the help of the dynamic order parameter Q we may define the fourth-order Binder cumulant [28,29],

$$U_L = 1 - \frac{\langle |Q|^4 \rangle_L}{3\langle |Q|^2 \rangle_L^2}, \quad (9)$$

a very useful observable for the characterization of universality classes [58].

IV. SIMULATION DETAILS

We performed Monte Carlo simulations on square lattices with periodic boundary conditions using the single-site update Metropolis algorithm [59–61]. This approach, together with the stochastic Glauber dynamics [62], consists the standard recipe in kinetic Monte Carlo simulations [31]. Let us briefly outline below the steps of our computer algorithm:

(1) A lattice site is selected randomly among the $L \times L$ options.

(2) The spin variable located at the selected site is flipped, keeping the other spins in the system fixed.

(3) The energy change originating from this spin flip operation is calculated using the Hamiltonians of Eqs. (1) and (2) as follows: $\Delta\mathcal{H} = \mathcal{H}_a - \mathcal{H}_o$, where \mathcal{H}_a denotes the energy of the system after the trial switch of the selected spin and \mathcal{H}_o corresponds to the total energy of the system with the old spin configuration. The probability to accept the proposed spin update is given by

$$W_M(\sigma_x \rightarrow \sigma'_x) = \begin{cases} \exp(-\Delta\mathcal{H}/k_B T), & \text{if } \mathcal{H}_a \geq \mathcal{H}_o, \\ 1, & \text{if } \mathcal{H}_a < \mathcal{H}_o. \end{cases} \quad (10)$$

(4) If the energy is lowered, the spin flip is always accepted.

(5) If the energy is increased, a random number R is generated such that $0 < R < 1$: If this number R is less than or equal to the calculated Metropolis transition probability the selected spin is flipped. Otherwise, the old spin configuration remains unchanged.

Using the above scheme we simulated system sizes within the range of $L = 32$ –256. For each system size 300 independent realizations of the disorder have been generated—see Fig. 1 for characteristic illustrations of disorder averages and their relative variance—and for each random sample the following simulation protocol has been used: The first 10^3 periods of the external field have been discarded during the thermalization process, and numerical data were collected and

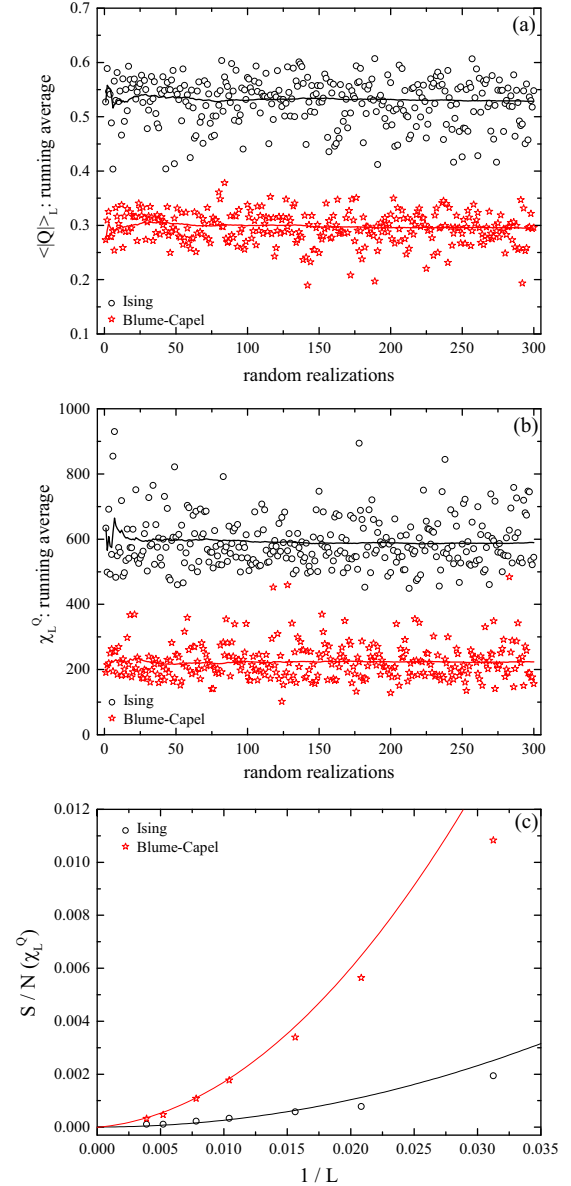


FIG. 1. Panel (a) Disorder distributions of the nonequilibrium dynamic order parameter, and panel (b) susceptibility maxima for a lattice size $L = 128$ and for both models considered in this paper. The running averages over the samples are shown by the solid lines. Panel (c) shows the signal-to-noise ratio S/N of the dynamic susceptibility, that is, the ratio of the relative variance of the distribution over the square of its mean value as a function of the inverse linear size. The solid lines are second-order polynomial fittings to $1/L$ for the larger system sizes. Clearly, $S/N(\chi_L^Q) \rightarrow 0$ as $L \rightarrow \infty$, indicating that self-averaging is restored in the thermodynamic limit for both kinetic disordered systems [63,64].

analyzed during the following 10^4 periods of the field. The time unit in our simulations was one Monte Carlo step per site (MCSS), and error bars have been estimated using the jackknife method [61]. To give a flavor of the actual CPU time of our computations we note that the simulation times needed for a single disorder realization of the kinetic Ising model on a single node of a Dual Intel Xeon E5-2690 V4 processor were 6 h for $L = 32$ and 11 days for $L = 256$. The analogous

CPU times for the kinetic Blume-Capel model were 3 h and 9 days for $L = 32$ and $L = 256$, respectively. For the Ising model we fixed the value of the disorder strength to $r = 1/7$, whereas for the Blume-Capel model we focused on the value of $\Delta = 1.975$ in the original first-order regime selecting now $r = 0.75/1.25$ following Refs. [48,49]. Appropriate choices of the magnetic-field strength $h_0 = 0.3$ and the temperatures $T^{(\text{Ising})} = 0.8 \times T_c^{(\text{Ising})}$ and $T^{(\text{BC})} = 0.6 \times T_c^{(\text{BC})}$ ensured that the system lies in the multidroplet regime [32]. Here, $T_c^{(\text{Ising})} = 1.7781$ [50,51] and $T_c^{(\text{BC})} = 0.626$ [48,49] are the equilibrium critical temperatures of the Ising and Blume-Capel models for the particular choices of r and Δ considered in this paper.

For the fitting process on the numerical data we restricted ourselves to data with $L \geq L_{\min}$. As usual, to determine an acceptable L_{\min} we employed the standard χ^2 test of goodness of fit [65]. Specifically, the p value of our χ^2 test is the probability of finding a χ^2 value which is even larger than the one actually found from our data. We considered a fit as being fair only if $10\% < p < 90\%$.

V. RESULTS AND DISCUSSION

As a starting point let us describe shortly the mechanism underlying the dynamical ordering in kinetic ferromagnets (here, under the presence of quenched randomness) as exemplified in Figs. 2–4 for the Ising model and Figs. 5–8 for the Blume-Capel model. In both cases, results for a single realization of the disorder are shown over a system size of $L = 96$.

Figure 2 presents the time evolution of the magnetization, and Fig. 3 presents the period dependencies of the dynamic order parameter Q of the kinetic random-bond Ising model. Several comments are in order: For rapidly varying fields, Fig. 2(a), the magnetization does not have enough time to switch during a single half-period and remains nearly constant for many successive field cycles as also illustrated by the black line in Fig. 3. On the other hand, for slowly varying fields, Fig. 2(c), the magnetization follows the field, switching every half-period so that $Q \approx 0$ as also shown by the blue line in Fig. 3. In other words, whereas in the dynamically disordered phase the ferromagnet is able to reverse its magnetization before the field changes again, in the dynamically ordered phase this is not possible, and therefore the time-dependent magnetization oscillates around a finite value. The competition between the magnetic field and the metastable state is captured by the half-period parameter $t_{1/2}$ (or by the normalized parameter $\Theta = t_{1/2}/\tau$ with τ being the metastable lifetime [32]). Obviously, $t_{1/2}$ plays the role of the temperature in the equilibrium system. Now, the transition between the two regimes is characterized by strong fluctuations in Q , see Fig. 2(b) and the evolution of the red line in Fig. 3. This behavior is indicative of a dynamic phase transition and occurs for values of the half-period close to the critical one of $t_{1/2}^c$ (otherwise when $\Theta \approx 1$). Of course, since the value of $t_{1/2} = 76$ MCSS used for this illustration is slightly above $t_{1/2}^c = 74.7(3)$, see also Fig. 10, the observed behavior includes as well some nonvanishing finite-size effects.

Some additional spatial aspects of the transition scenarios described above via the configurations of the local order parameter $\{Q_x\}$ are shown in Fig. 4. Below $t_{1/2}^c$, see panel (a),

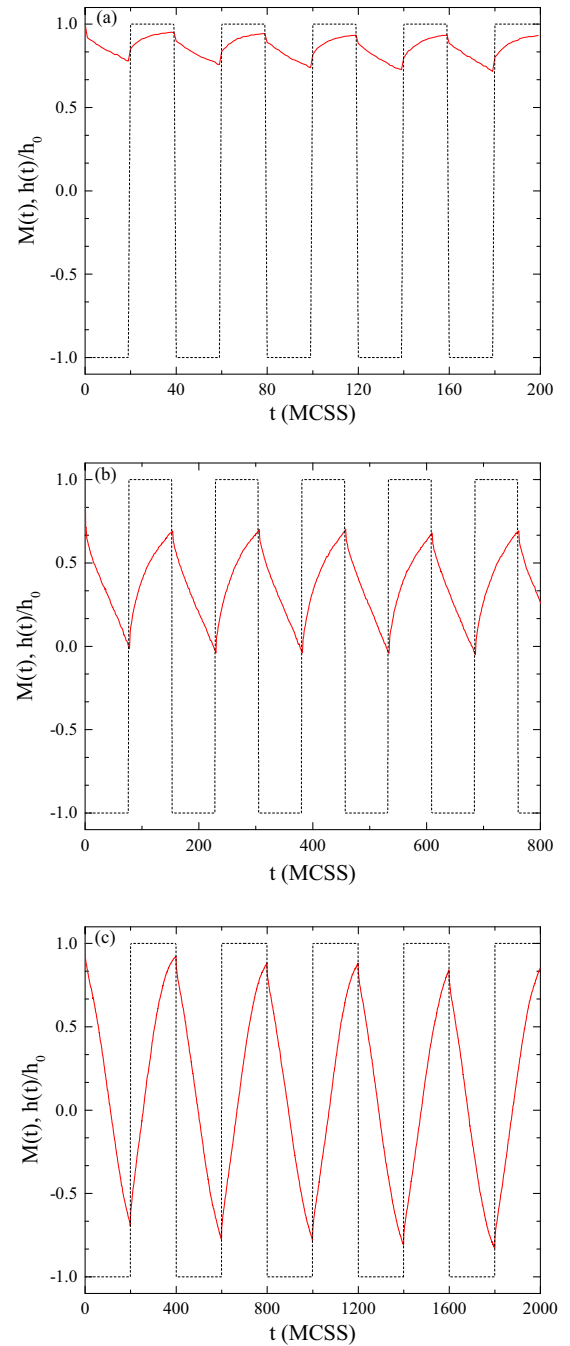


FIG. 2. Time series of the magnetization (red solid curves) of the kinetic random-bond Ising model under the presence of a square-wave magnetic field (black dashed lines) for $L = 96$ and three values of the half-period of the external field: (a) $t_{1/2} = 20$ MCSS, corresponding to a dynamically ordered phase, (b) $t_{1/2} = 76$ MCSS, close to the dynamic phase transition, and (c) $t_{1/2} = 200$ MCSS, corresponding to a dynamically disordered phase. Note that for the sake of clarity the ratio $h(t)/h_0$ is displayed.

the majority of spins spend most of their time in the $+1$ state, i.e., in the metastable phase during the first half-period and in the stable equilibrium phase during the second half-period except for equilibrium fluctuations. Thus most of $Q_x \approx +1$ and the system is now in the dynamically ordered phase. On the other hand, when the period of the external field is selected

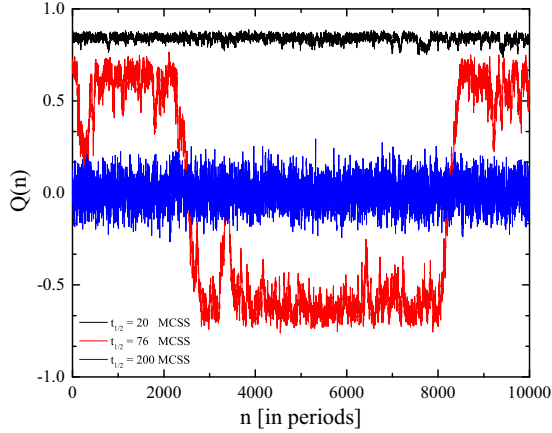


FIG. 3. Period dependencies of the dynamic order parameter of the kinetic random-bond Ising model for $L = 96$. Results are shown for the three characteristic cases of the half-period of the external field, following Fig. 2.

to be bigger than the relaxation time of the system, above $t_{1/2}^c$, see panel (c), the system follows the field in every half-period with some phase lag and $Q_x \approx 0$ at all sites x . The system lies in the dynamically disordered phase. Near $t_{1/2}^c$ and the expected dynamic phase transition, there are large clusters of both $Q_x \approx +1$ and -1 values within a sea of $Q_x \approx 0$ as shown in Fig. 4(b).

Although the discussion above concentrated on the Ising case, an analogous description and relevant conclusions may be drawn also for the dynamical ordering of the disordered-induced continuous transition of the Blume-Capel model as depicted in Figs. 5–7. Note that in this case the critical half-period of the system has been estimated to be $t_{1/2}^c = 83.6(4)$ (see also Fig. 11 below). However, we should underline here that for the case of the Blume-Capel model the value of the local order parameter $\{Q_x\}$ does not distinguish between random distributions of $\sigma_x = \pm 1$ and regions of $\sigma_x = 0$. To bring out this distinction, we present in Fig. 8 similar snapshots of the dynamic quadrupole moment over a full cycle of the external field $O = \frac{1}{2t_{1/2}} \oint q(t) dt$, where $q(t) = \frac{1}{N} \sum_{x=1}^N \sigma_x^2$. In the spin-1 Blume-Capel model the density of the vacancies is controlled by the crystal-field coupling Δ and, thus, the value of the dynamic quadrupole moment changes depending on Δ [37]. We point out that, in Fig. 8, except for the red $+1$ areas, the regions enclosed by finite values demonstrate the role played by the crystal-field coupling in the Blume-Capel model.

To further explore the nature of the dynamic phase transitions encountered in the above disordered kinetic models we performed a finite-size scaling analysis using the observables outlined in Sec. III. Previous studies in the field indicated that, although scaling laws and finite-size scaling are tools that have been designed for the study of equilibrium phase transitions, they can be successfully applied as well to far from equilibrium systems [28–32].

As an illustrative example for the case of the kinetic random-bond Ising model and for a system size of $L = 64$ we present in the main panel of Fig. 9 the finite-size behavior of the dynamic order parameter and in the lower inset the emerging dynamic susceptibility [see Eq. (7)]. The dynamic order parameter goes

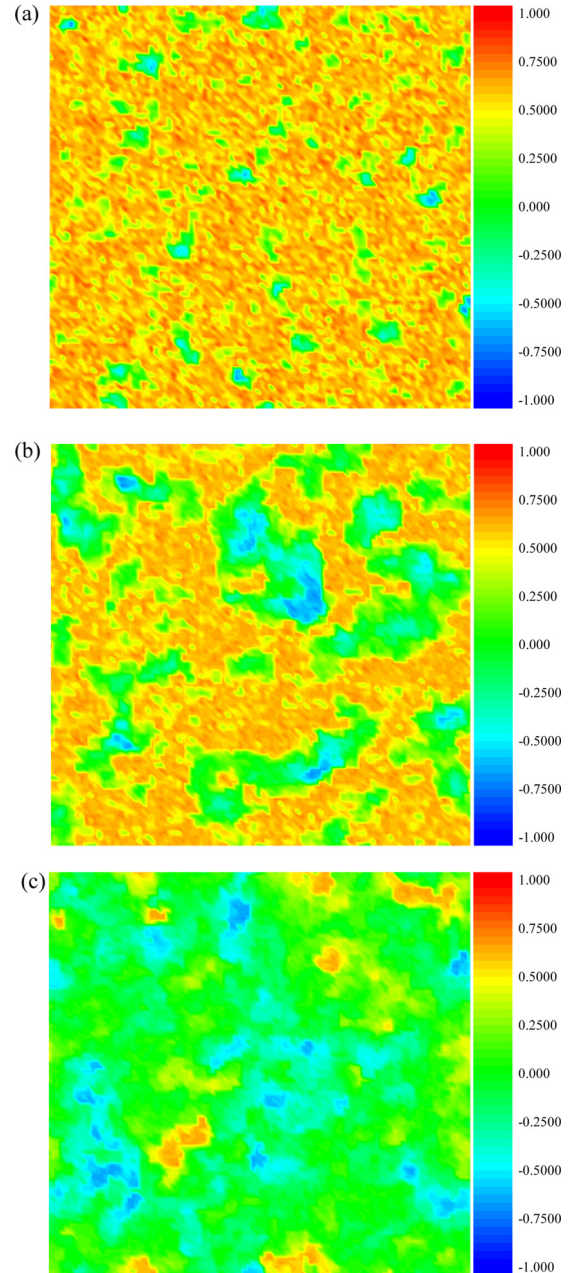


FIG. 4. Configurations of the local dynamic order parameter $\{Q_x\}$ of the random-bond kinetic Ising model for $L = 96$. The “snapshots” of $\{Q_x\}$ for each regime are the set of local period-averaged spins during some representative period. Three panels are shown: (a) $t_{1/2} = 20$ MCSS $< t_{1/2}^c$ —dynamically ordered phase, (b) $t_{1/2} = 76$ MCSS $\approx t_{1/2}^c$ —near the dynamic phase transition, and (c) $t_{1/2} = 200$ MCSS $> t_{1/2}^c$ —dynamically disordered phase.

from a finite value to zero values as the half-period increases showing a sharp change around the value of the half-period that can be mapped to the respective peak in the plot of the dynamic susceptibility. The location of the maxima in χ_L^Q may be used to define suitable pseudocritical half-periods, denoted hereafter as $t_{1/2}^*$. The corresponding maxima may be analogously denoted as $(\chi_L^Q)^*$. We also measured the energy and its scaled variance, the heat capacity χ_L^E [see Eq. (8)]. The

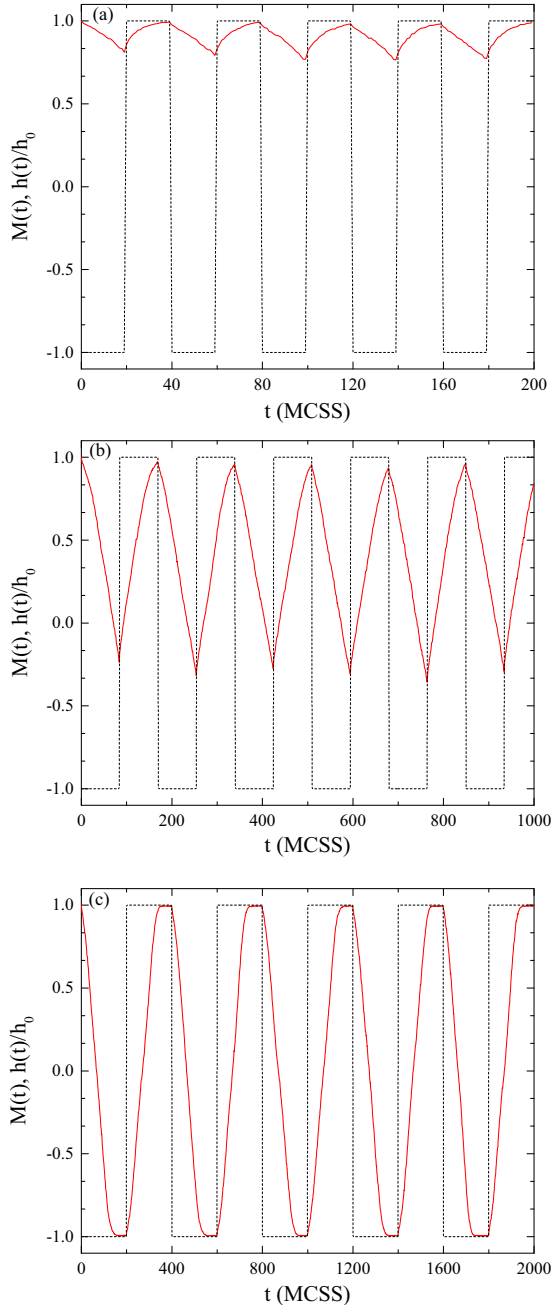


FIG. 5. Time series of the magnetization (red solid curves) of the kinetic random-bond $\Delta = 1.975$ Blume-Capel model under the presence of a square-wave magnetic field (black dashed lines) for $L = 96$ and three values of the half-period of the external field: (a) $t_{1/2} = 20$ MCSS, corresponding to a dynamically ordered phase, (b) $t_{1/2} = 85$ MCSS, close to the dynamic phase transition, and (c) $t_{1/2} = 200$ MCSS, corresponding to a dynamically disordered phase. Note that for the sake of clarity the ratio $h(t)/h_0$ is displayed.

upper inset of Fig. 9 shows the half-period dependency of the energy of the same system and the relevant heat capacity. In this case the maxima may be denoted as $(\chi_L^E)^*$.

We start the presentation of our finite-size scaling analysis with the Ising case. In the main panel of Fig. 10(a) we present the size evolution of the peaks of the dynamic susceptibility

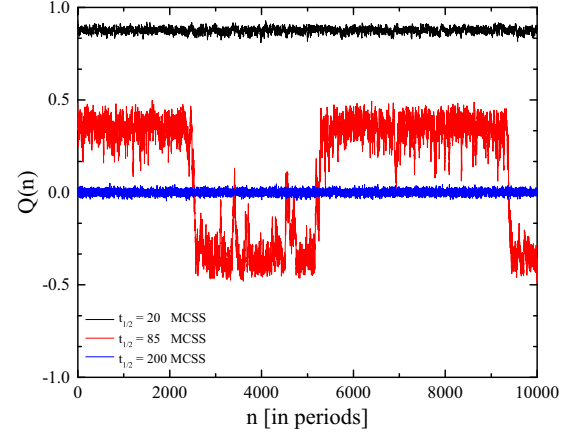


FIG. 6. Period dependencies of the dynamic order parameter of the kinetic random-bond $\Delta = 1.975$ Blume-Capel model for $L = 96$. Results are shown for the three characteristic cases of the half-period of the external field, following Fig. 5.

on a log-log scale. The solid line is a fit of the form [66]

$$(\chi_L^Q)^* \sim L^{\gamma/\nu}, \quad (11)$$

providing an estimate of 1.75(1) for the magnetic exponent ratio γ/ν , in excellent agreement to the Ising value of $7/4$. The shift behavior of the corresponding peak locations $t_{1/2}^*$ is plotted in the inset of Fig. 10(a) as a function of $1/L$. The solid line shows a fit of the usual shift form [67–69]

$$t_{1/2}^* = t_{1/2}^c + bL^{-1/\nu}, \quad (12)$$

where $t_{1/2}^c$ defines the critical half-period of the system and ν is the critical exponent of the correlation length. The obtained values of $t_{1/2}^c = 74.7(3)$ and $\nu = 1.03(4)$ are listed also in the panel and, in particular, the value of the critical exponent ν appears to be in very good agreement with the value of $\nu = 1$ of the 2D equilibrium Ising model [70]. This finding strongly supports the claim that the kinetic Ising model under the presence of quenched bond randomness shares the universality class of its corresponding equilibrium counterpart. Ideally, we would also like to observe the double logarithmic scaling behavior of the maxima of the heat capacity $(\chi_L^E)^*$. Indeed, as is shown in the main panel of Fig. 10(b), the data for the maxima of the heat capacity are adequately described by a fit of the form

$$(\chi_L^E)^* \sim \ln[\ln(L)], \quad (13)$$

as predicted by Ref. [71] for the random Ising universality class. As a comparison, we plot the same data with respect to the simple logarithm of the system size in the corresponding inset. It is obvious that a fit $(\chi_L^E)^* \sim \ln(L)$, as shown by the solid line, does not capture the full scaling behavior.

So, where do we stand at this point: We have shown that the universality class of the dynamic phase transition encountered in an Ising model under the presence of quenched bond disorder is equivalent to that of its equilibrium counterpart with the inclusion of logarithmic corrections in the scaling of the heat capacity. We turn now our discussion to the dynamic phase transition of the $\Delta = 1.975$ Blume-Capel model with bond disorder. As mentioned previously in Sec. II, only very recently

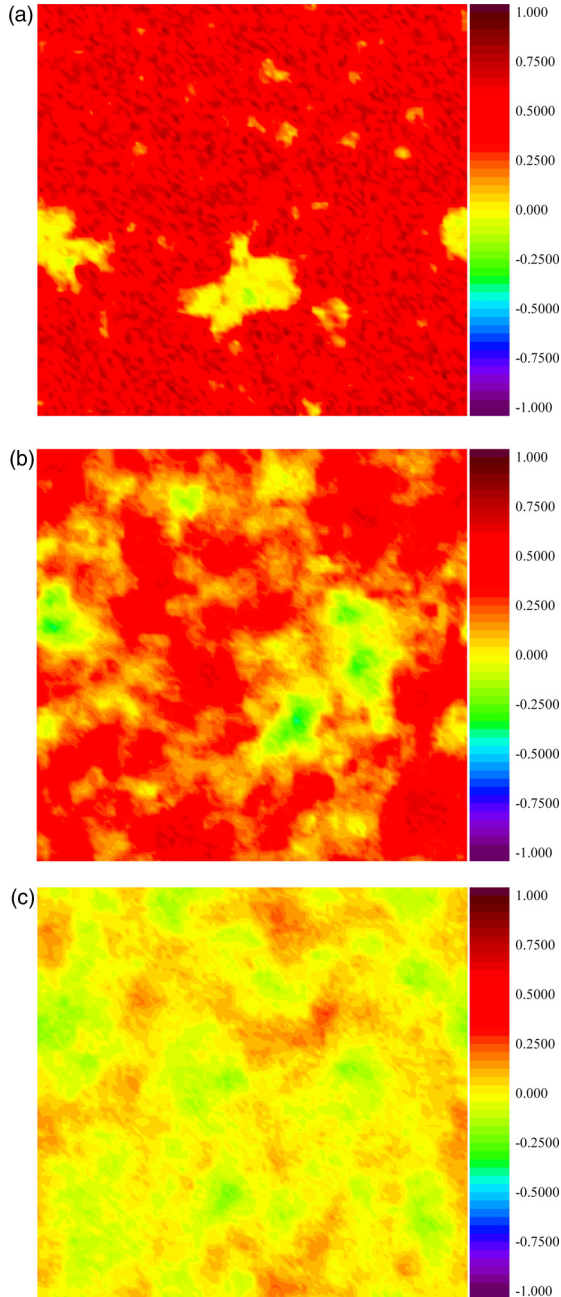


FIG. 7. Configurations of the local dynamic order parameter $\{Q_x\}$ of the random-bond kinetic $\Delta = 1.975$ Blume-Capel for $L = 96$. The snapshots of $\{Q_x\}$ for each regime are the set of local period-averaged spins during some representative period. Three panels are shown: (a) $t_{1/2} = 20$ MCSS $< t_{1/2}^c$ —dynamically ordered phase, (b) $t_{1/2} = 85$ MCSS $\approx t_{1/2}^c$ —near the dynamic phase transition, and (c) $t_{1/2} = 200$ MCSS $> t_{1/2}^c$ —dynamically disordered phase.

the claims of universality violation in the equilibrium random-bond Blume-Capel model have been dispelled, and it was shown that the induced under disorder continuous transition belongs to the universality class of the random Ising model [57]. We therefore expect, or at least hope, that the results presented in the current paper will also be relevant to this reignited problem yet from a nonequilibrium perspective.

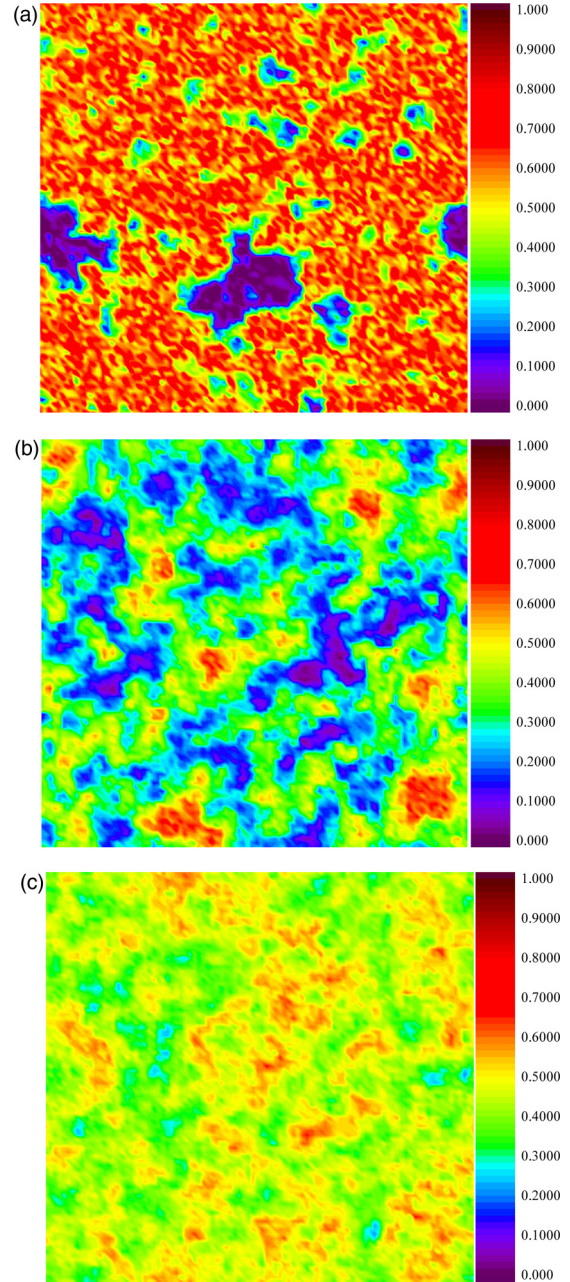


FIG. 8. In full analogy to Fig. 7 we show snapshots of the period-averaged dynamic quadrupolar moment conjugate to the crystal-field coupling Δ . The simulation parameters are exactly the same to those used in Fig. 7 for all three panels (a)–(c).

The scaling aspects of the dynamic phase transition of the kinetic random-bond Blume-Capel model at $\Delta = 1.975$ are shown in Fig. 11, following fully the presentation and analysis style of Fig. 10 and excluding the data for $L = 32$ that suffer from strong finite-size effects. In this case an estimate of $1.74(2)$ is obtained for the magnetic exponent ratio γ/ν , again compatible within errors to the Ising value of $7/4$. From the shift behavior of the corresponding pseudocritical half-periods $t_{1/2}^*$ [inset of Fig. 11(a)] the critical half-period and the correlation-length exponent are estimated to be $t_{1/2}^c = 83.6(4)$ and $\nu = 1.05(7)$, respectively. Again the estimate of ν supports

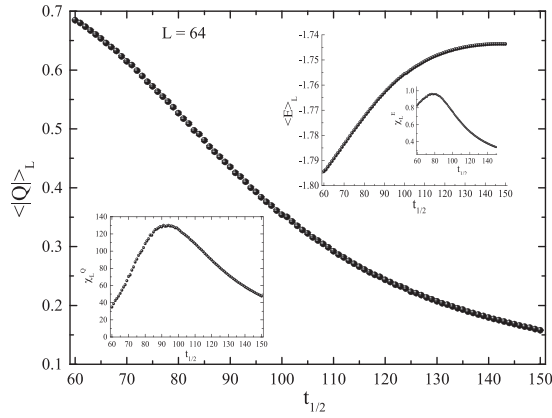


FIG. 9. Half-period dependency of the dynamic order parameter of the kinetic random-bond Ising model. The lower inset illustrates the half-period dependency of the corresponding dynamic susceptibility χ_L^Q . The upper inset shows the half-period dependency of the energy and the corresponding heat capacity χ_L^E . All results shown refer to a system size of $L = 64$ at the critical $t_{1/2}$ region.

the scenario presented above in Fig. 10 for the criticality in the dynamic phase transition of the random-bond kinetic Ising model. Last but not least, in Fig. 11(b) the maxima of the heat capacity $(\chi_L^E)^*$ are plotted versus $\ln[\ln(L)]$ (main panel) and

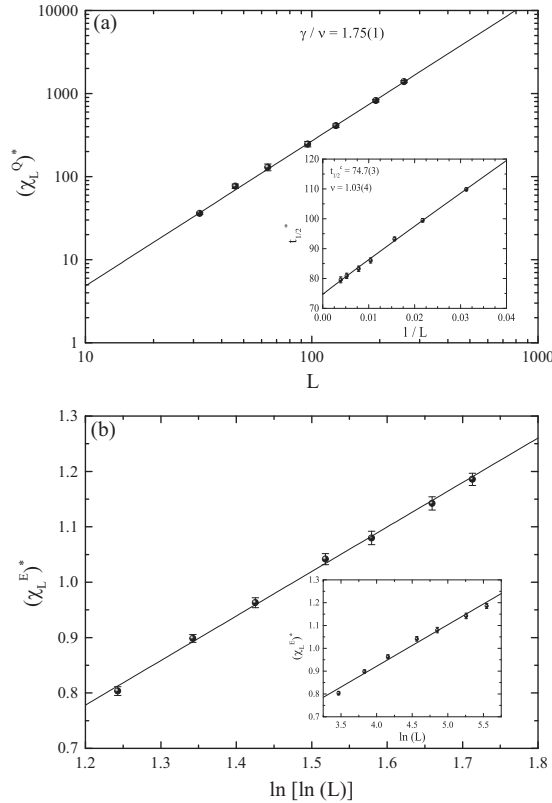


FIG. 10. Criticality in the kinetic random-bond Ising model ($r = 1/7$): (a) Finite-size scaling behavior of the maxima $(\chi_L^Q)^*$ on a log-log scale (main panel) and shift behavior of the corresponding pseudocritical half-periods $t_{1/2}^*$ (inset). (b) Double (main panel) and simple (inset) logarithmic scaling behavior of the heat-capacity maxima $(\chi_L^E)^*$. In all cases lines are linear fittings.

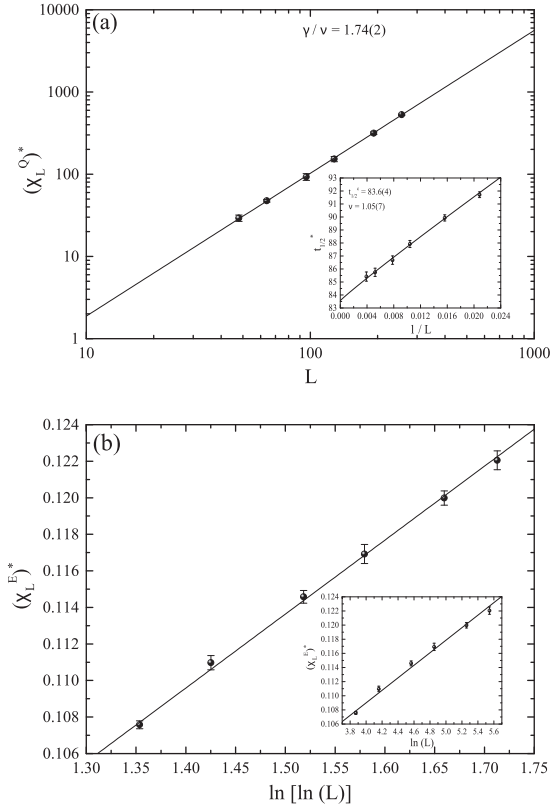


FIG. 11. Criticality in the kinetic random-bond Blume-Capel model ($r = 0.75/1.25$; $\Delta = 1.975$). The description is analogous to that of Fig. 10.

$\ln(L)$ (inset) and as in the Ising case are much better described by the double logarithmic fit (13).

An alternative test of universality comes from the study of the fourth-order Binder cumulant U_L defined in Eq. (9) for the case of the dynamic order parameter. In Fig. 12 we present our numerical data of U_L for the kinetic random-bond Ising (main

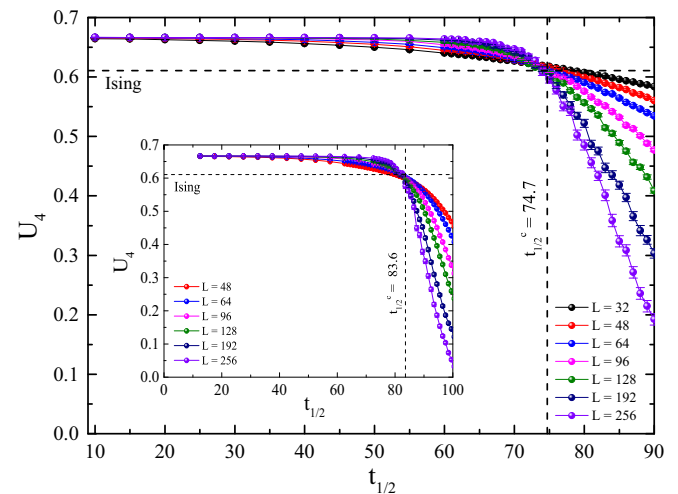


FIG. 12. Half-period dependency of the fourth-order Binder cumulant U_L of the kinetic random-bond Ising (main panel) and Blume-Capel (inset) models for a wide range of system sizes studied.

panel) and the Blume-Capel (inset) models. In both panels the vertical dashed lines mark the critical half-period value of the system $t_{1/2}^c$, and the horizontal dotted lines mark the universal value of $U^* = 0.6106924(16)$ of the 2D equilibrium Ising model [72]. Certainly, the crossing point is expected to depend on the lattice size L (as also shown in the figure), and the term universal is valid for given lattice shapes, boundary conditions, and isotropic interactions [73,74]. However, the data shown in Fig. 12 support, at least qualitatively, another instance of equilibrium Ising universality since in both panels the crossing point is consistent with the value of 0.6106924. We should note here that Hasenbusch *et al.* presented very strong evidence that the critical Binder cumulant of the equilibrium 2D randomly site-diluted Ising model maintains its pure-system value [75]. In this respect, a dedicated study along the lines of Ref. [75] for an accurate estimation of U^* in the kinetic random-bond Ising and Blume-Capel models would be welcome but certainly goes beyond the scope of the current paper.

VI. CONCLUSIONS

In the present paper we investigated the effect of quenched disorder on the dynamic phase transition of kinetic spin models in two dimensions. In particular, we considered the square-lattice Ising and Blume-Capel models under a periodically

oscillating magnetic field, the latter at its low-temperature regime where the pure equilibrium system exhibits a first-order phase transition. Using Monte Carlo simulations and finite-size scaling techniques we have been able to probe with good accuracy the values of the critical exponent ν and the magnetic exponent ratio γ/ν , both of which were found to be compatible with those of the equilibrium Ising ferromagnet. An additional study of the scaling behavior of the heat capacity revealed the double logarithmic divergence expected for the universality class of the random Ising model. To conclude, although universality is a cornerstone in the theory of critical phenomena, it stands on a less solid foundation for the case of nonequilibrium systems and for systems subject to quenched disorder. In the current paper we have studied two systems where both of the above complications merge, yet arriving at the simplest scenario. We hope that our paper will stimulate further research in the field of nonequilibrium critical phenomena in both numerical and analytical directions.

ACKNOWLEDGMENTS

The authors would like to thank P. A. Rikvold and W. Selke for many useful comments about the paper. The numerical calculations reported in this paper were performed at TÜBİTAK ULAKBİM (Turkish agency), High Performance and Grid Computing Center (TRUBA Resources).

-
- [1] T. Tomé and M. J. de Oliveira, *Phys. Rev. A* **41**, 4251 (1990).
 - [2] W. S. Lo and R. A. Pelcovits, *Phys. Rev. A* **42**, 7471 (1990).
 - [3] M. F. Zimmer, *Phys. Rev. E* **47**, 3950 (1993).
 - [4] M. Acharyya and B. K. Chakrabarti, *Phys. Rev. B* **52**, 6550 (1995).
 - [5] B. K. Chakrabarti and M. Acharyya, *Rev. Mod. Phys.* **71**, 847 (1999).
 - [6] M. Acharyya, *Phys. Rev. E* **56**, 1234 (1997).
 - [7] M. Acharyya, *Phys. Rev. E* **69**, 027105 (2004).
 - [8] G. M. Buendía and E. Machado, *Phys. Rev. E* **58**, 1260 (1998).
 - [9] G. M. Buendía and E. Machado, *Phys. Rev. B* **61**, 14686 (2000).
 - [10] H. Fujisaka, H. Tutu, and P. A. Rikvold, *Phys. Rev. E* **63**, 036109 (2001); **63**, 059903(E) (2001).
 - [11] H. Jang, M. J. Grimson, and C. K. Hall, *Phys. Rev. E* **68**, 046115 (2003).
 - [12] H. Jang, M. J. Grimson, and C. K. Hall, *Phys. Rev. B* **67**, 094411 (2003).
 - [13] X. Shi, G. Wei, and L. Li, *Phys. Lett. A* **372**, 5922 (2008).
 - [14] A. Punya, R. Yimnirun, P. Laoratanakul, and Y. Laosiritaworn, *Physica B* **405**, 3482 (2010).
 - [15] P. Riego and A. Berger, *Phys. Rev. E* **91**, 062141 (2015).
 - [16] M. Keskin, O. Canko, and U. Temizer, *Phys. Rev. E* **72**, 036125 (2005).
 - [17] M. Keskin, O. Canko, and Ü. Temizer, *J. Exp. Theor. Phys.* **104**, 936 (2007).
 - [18] D. T. Robb, P. A. Rikvold, A. Berger, and M. A. Novotny, *Phys. Rev. E* **76**, 021124 (2007).
 - [19] B. Deviren and M. Keskin, *J. Magn. Magn. Mater.* **324**, 1051 (2012).
 - [20] Y. Yüksel, E. Vatansever, and H. Polat, *J. Phys.: Condens. Matter* **24**, 436004 (2012).
 - [21] Y. Yüksel, E. Vatansever, U. Akinci, and H. Polat, *Phys. Rev. E* **85**, 051123 (2012).
 - [22] E. Vatansever, *Phys. Lett. A* **381**, 1535 (2017).
 - [23] Y.-L. He and G.-C. Wang, *Phys. Rev. Lett.* **70**, 2336 (1993).
 - [24] D. T. Robb, Y. H. Xu, O. Hellwig, J. McCord, A. Berger, M. A. Novotny, and P. A. Rikvold, *Phys. Rev. B* **78**, 134422 (2008).
 - [25] J.-S. Suen and J. L. Erskine, *Phys. Rev. Lett.* **78**, 3567 (1997).
 - [26] A. Berger, O. Idigoras, and P. Vavassori, *Phys. Rev. Lett.* **111**, 190602 (2013).
 - [27] P. Riego, P. Vavassori, and A. Berger, *Phys. Rev. Lett.* **118**, 117202 (2017).
 - [28] S. W. Sides, P. A. Rikvold, and M. A. Novotny, *Phys. Rev. Lett.* **81**, 834 (1998).
 - [29] S. W. Sides, P. A. Rikvold, and M. A. Novotny, *Phys. Rev. E* **59**, 2710 (1999).
 - [30] G. Korniss, C. J. White, P. A. Rikvold, and M. A. Novotny, *Phys. Rev. E* **63**, 016120 (2000).
 - [31] G. M. Buendía and P. A. Rikvold, *Phys. Rev. E* **78**, 051108 (2008).
 - [32] H. Park and M. Pleimling, *Phys. Rev. E* **87**, 032145 (2013).
 - [33] H. Park and M. Pleimling, *Phys. Rev. Lett.* **109**, 175703 (2012).
 - [34] K. Tauscher and M. Pleimling, *Phys. Rev. E* **89**, 022121 (2014).
 - [35] G. M. Buendía and P. A. Rikvold, *Phys. Rev. B* **96**, 134306 (2017).
 - [36] E. Vatansever, [arXiv:1706.03351](https://arxiv.org/abs/1706.03351).
 - [37] E. Vatansever and N. G. Fytas, *Phys. Rev. E* **97**, 012122 (2018).
 - [38] X. Shi and G. Wei, *Phys. Scr.* **89**, 075805 (2014).

- [39] M. Acharyya and A. Halder, *J. Magn. Magn. Mater.* **426**, 53 (2017).
- [40] G. Grinstein, C. Jayaprakash, and Y. He, *Phys. Rev. Lett.* **55**, 2527 (1985).
- [41] G. Gulpinar and E. Vatansever, *J. Stat. Phys.* **146**, 787 (2012).
- [42] G. Gulpinar, E. Vatansever, and M. Agartioglu, *Physica A* **391**, 3574 (2012).
- [43] U. Akinci, Y. Yüksel, E. Vatansever, and H. Polat, *Physica A* **391**, 5810 (2012).
- [44] E. Vatansever, U. Akinci, and H. Polat, *J. Magn. Magn. Mater.* **344**, 89 (2013).
- [45] E. Vatansever, U. Akinci, Y. Yüksel, and H. Polat, *J. Magn. Magn. Mater.* **329**, 14 (2013).
- [46] E. Vatansever and H. Polat, *Phys. Lett. A* **379**, 1568 (2015).
- [47] A. Malakis, A. N. Berker, I. A. Hadjiagapiou, and N. G. Fytas, *Phys. Rev. E* **79**, 011125 (2009).
- [48] A. Malakis, A. N. Berker, I. A. Hadjiagapiou, N. G. Fytas, and T. Papakonstantinou, *Phys. Rev. E* **81**, 041113 (2010).
- [49] A. Malakis, A. N. Berker, N. G. Fytas, and T. Papakonstantinou, *Phys. Rev. E* **85**, 061106 (2012).
- [50] R. Fisch, *J. Stat. Phys.* **18**, 111 (1978).
- [51] W. Kinzel and E. Domany, *Phys. Rev. B* **23**, 3421 (1981).
- [52] H. W. Capel, *Physica (Amsterdam)* **32**, 966 (1966).
- [53] M. Blume, *Phys. Rev.* **141**, 517 (1966).
- [54] I. D. Lawrie and S. Sarbach, in *Phase Transitions and Critical Phenomena*, edited by C. Domb and J. L. Lebowitz (Academic, London, 1984), Vol. 9.
- [55] J. Zierenberg, N. G. Fytas, M. Weigel, W. Janke, and A. Malakis, *Eur. Phys. J. Spec. Top.* **226**, 789 (2017).
- [56] W. Kwak, J. Jeong, J. Lee, and D.-H. Kim, *Phys. Rev. E* **92**, 022134 (2015).
- [57] N. G. Fytas, J. Zierenberg, P. E. Theodorakis, M. Weigel, W. Janke, and A. Malakis, *Phys. Rev. E* **97**, 040102(R) (2018).
- [58] K. Binder, *Z. Phys. B: Condens. Matter* **43**, 119 (1981); *Phys. Rev. Lett.* **47**, 693 (1981).
- [59] N. Metropolis, A. W. Rosenbluth, M. N. Rosenbluth, A. H. Teller, and E. Teller, *J. Chem. Phys.* **21**, 1087 (1953).
- [60] D. P. Landau and K. Binder, *A Guide to Monte Carlo Simulations in Statistical Physics* (Cambridge University Press, Cambridge, UK, 2000).
- [61] M. E. J. Newman and G. T. Barkema, *Monte Carlo Methods in Statistical Physics* (Oxford University Press, New York, 1999).
- [62] R. J. Glauber, *J. Math. Phys.* **4**, 294 (1963).
- [63] A. Aharony and A. B. Harris, *Phys. Rev. Lett.* **77**, 3700 (1996).
- [64] S. Wiseman and E. Domany, *Phys. Rev. Lett.* **81**, 22 (1998); *Phys. Rev. E* **58**, 2938 (1998).
- [65] W. H. Press, S. A. Teukolsky, W. T. Vetterling, and B. P. Flannery, *Numerical Recipes in C*, 2nd ed. (Cambridge University Press, Cambridge, UK, 1992).
- [66] A. M. Ferrenberg and D. P. Landau, *Phys. Rev. B* **44**, 5081 (1991).
- [67] M. E. Fisher, in *Critical Phenomena*, edited by M. S. Green (Academic, London, 1971).
- [68] V. Privman, *Finite Size Scaling and Numerical Simulation of Statistical Systems* (World Scientific, Singapore, 1990).
- [69] K. Binder, in *Computational Methods in Field Theory*, edited by C. B. Lang and H. Gausterer (Springer, Berlin, 1992).
- [70] A. E. Ferdinand and M. E. Fisher, *Phys. Rev.* **185**, 832 (1969).
- [71] Vik. S. Dotsenko and VI. S. Dotsenko, *Sov. Phys. JETP Lett.* **33**, 37 (1981).
- [72] J. Salas and A. D. Sokal, *J. Stat. Phys.* **98**, 551 (2000).
- [73] W. Selke, *J. Stat. Mech.: Theor. Exp.* (2007) P04008.
- [74] W. Selke and L. N. Shchur, *Phys. Rev. E* **80**, 042104 (2009).
- [75] M. Hasenbusch, F. Parisen Toldin, A. Pelissetto, and E. Vicari, *Phys. Rev. E* **78**, 011110 (2008).