

# Coarsening dynamics of an antiferromagnetic XY model on the kagome lattice: Breakdown of critical dynamic scaling

Sangwoong Park,<sup>1</sup> Bongsoo Kim,<sup>2</sup> and Sung Jong Lee<sup>1</sup><sup>1</sup>*Department of Physics, The University of Suwon, Kyonggi-do 445-743, Korea*<sup>2</sup>*Department of Physics, Changwon National University, Changwon 641-773, Korea*

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We find a breakdown of the critical dynamic scaling in the coarsening dynamics of an antiferromagnetic XY model on the kagome lattice when the system is quenched from disordered states into the Kosterlitz-Thouless phases at low temperatures. There exist multiple growing length scales; the length scales of the average separation between fractional vortices are found to be *not* proportional to the length scales of the quasiordered domains. They are instead related through a nontrivial power-law relation. The length scale of the quasiordered domains (as determined from the optimal collapse of the correlation functions for the order parameter  $\exp[3i\theta(r)]$ ) does not follow a simple power-law growth but exhibits an anomalous growth with time-dependent effective growth exponent. The breakdown of the critical dynamic scaling is accompanied by unusual relaxation dynamics in the decay of the fractional ( $3\theta$ ) vortices, where the decay of the vortex numbers is characterized by an exponential function of the logarithmic powers in time.

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## I. INTRODUCTION

Thermodynamic systems quenched from a high-temperature disordered phase into a low-temperature ordered phase exhibit characteristic growth of the length scale  $\ell$  of the ordered domains which in typical situations can be represented as a power law  $\ell \sim t^{1/z}$ , where the growth exponent  $1/z$  depends on the dimension of the space and of the relevant order parameter as well as on the conserved or non-conserved nature of the order parameter.<sup>1</sup> Phase ordering dynamics (or coarsening process) is usually accompanied by the annihilation and decay of the characteristic topological defects such as point vortices or domain walls which are generated in the initial disordered states. One of the most important notions in understanding and analyzing these coarsening processes is the so-called *dynamic scaling hypothesis*<sup>1</sup> for the equal-time spatial correlation function of the order parameter, which is closely related to the observed self-similarity of the coarsening systems at different time instants.

When the low-temperature phase of the system is characterized by a quasi-long-range order with power-law decay of the spatial correlation function of the order parameter at equilibrium, the dynamic scaling is generalized to the *critical dynamic scaling*. One of the well-known examples is the ferromagnetic XY model<sup>2,3</sup> on the square lattice.

However, it should be noted that this (critical) dynamic scaling hypothesis for the phase ordering (or quasiordering) dynamics has not been proven on some general theoretical basis. Therefore, it is not clearly known what is the condition for the validity of the dynamic scaling which is usually assumed in the analyses of experimental results or the numerical simulations on coarsening dynamics.

In typical situations, coarsening dynamics is investigated on systems where the phase ordering is accompanied by the breaking of discrete or continuous global symmetry. However, there exist model systems that exhibit infinite ground-state degeneracies of a discrete nature in addition to the usual

(global) continuous symmetry. Prominent examples are geometrically frustrated spin systems such as antiferromagnetic Ising models on a triangular lattice,<sup>4,5</sup> antiferromagnetic XY, or Heisenberg models on kagome<sup>6</sup> or pyrochlore lattices.<sup>7</sup> These systems exhibit interesting equilibrium and nonequilibrium behaviors due to the geometric frustration effect, including the spin-glass-like relaxation dynamics without quenched disorder.

In this work we perform numerical simulations on the coarsening of an antiferromagnetic XY model on the kagome lattice (*KAFX* model)<sup>8–12</sup> which is one of the simplest geometrically frustrated models with infinite ground-state degeneracies. Experimentally, this model can be realized in superconducting Josephson-junction arrays or superconducting wire networks<sup>13–15</sup> on a kagome lattice when a perpendicular magnetic field of half flux quantum (per plaquette) is applied on the system. We can also find physical examples in the anisotropic limit of Heisenberg antiferromagnets on the kagome lattice. It is well known that the system exhibits an infinite ground-state degeneracy with finite entropy.<sup>16</sup> The system also exhibits a finite temperature Kosterlitz-Thouless (KT) transition<sup>3</sup> corresponding to the unbinding of the so-called fractional  $3\theta$  vortices. That is, at low temperature below the KT transition, equilibrium of the system will be characterized by the quasi-long-range order of the order parameter  $\psi_3 \equiv e^{3i\theta(r)}$ . Analogous to the case of the simple ferromagnetic XY model on a square lattice, we might expect that the coarsening dynamics of the system would exhibit a critical dynamic scaling for the equal-time spatial correlation of the order parameter.

Our simulations, however, show that the critical dynamic scaling is not obeyed very well at least for the time duration of our numerical simulations ranging several decades of time scale. That is, it was not possible to achieve a good scaling collapse for the equal-time spatial correlation functions of the  $3\theta$  order parameter. Another signature of the breakdown of the critical dynamic scaling is that the length scales corresponding to the average separation between fractional vor-

tices are not proportional to the length scales of the growing quasiordered domains, which are instead related through a nontrivial power-law relation. This means that the two length scales exhibit different growth behavior in time. In terms of the decay of the fractional vortices, we also found that the fractional  $3\theta$  vortices residing in the small triangular plaquettes exhibit faster decay, while, in contrast, those vortices sitting on the larger hexagonal plaquettes exhibit much slower decay that can be fitted by an exponential or logarithmic powers in time.<sup>17</sup>

In addition to these features of the multiple length scales in the coarsening dynamics, the time dependence of the length scale of the quasiordered domains does not exhibit a simple power-law growth but rather exhibits an anomalous growth with time-dependent effective growth exponent. This appears to be closely related to the unusual relaxation dynamics in the non-power-law decay of the fractional ( $3\theta$ ) vortices. It is not clear yet whether the scaling may be restored in the asymptotic limit.

## II. MODEL SYSTEM AND SIMULATION METHODS

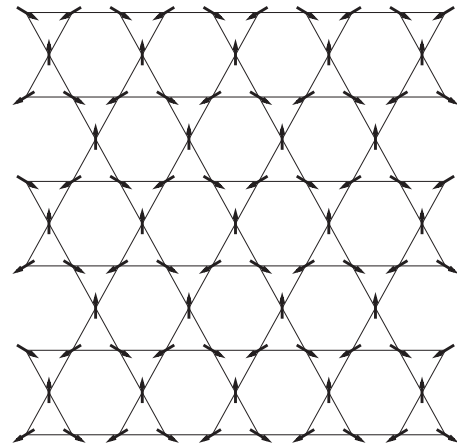
The kagome lattice consists of corner-sharing triangles (Fig. 1). In an antiferromagnetic XY model on the kagome lattice, the Hamiltonian is defined as

$$H = -J \sum_{\langle i,j \rangle} \cos(\theta_i - \theta_j), \quad (1)$$

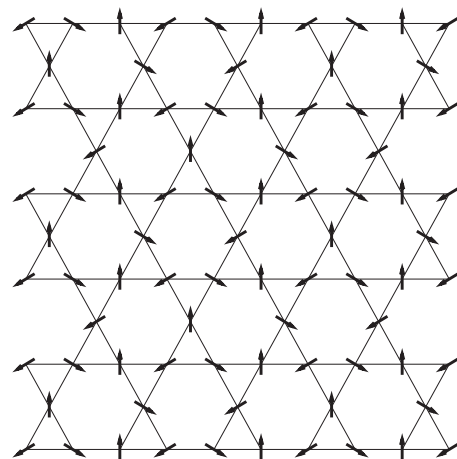
where  $J < 0$ , the sum runs over all nearest-neighbor pairs of sites, and  $\theta_i$  denotes the angle of the planar spin at site  $i$  with respect to some fixed direction in the two-dimensional spin space.  $\langle i,j \rangle$  indicates all pairs of nearest-neighbor sites in the kagome lattice.

It is easy to see that the ground states of this system have the property that for all pairs of nearest neighbors  $i$  and  $j$ , the angle difference satisfies  $|\theta_i - \theta_j| = 2\pi/3$  which means that the sum of the three spins vectors on any unit triangle vanishes. Thus the space of the ground states of the *KAFX* model are equivalent (up to a global rotation) to the ground states of the three state Potts model on the kagome lattice. In Figs. 1(a) and 1(b) two examples of simple ground states with long-range order are shown, the so-called  $\mathbf{q}=0$  state and  $\mathbf{q}=\sqrt{3} \times \sqrt{3}$  state, where  $\mathbf{q}$  refers to the wave vector corresponding to the periodicity of the chirality configuration. It is easy to see, however, that in addition to these ground states with simple spatial order there also exist infinitely many ground states with no spatial order. It is well known that the system has a ground-state entropy of  $S_0 \approx 0.126k_B$  per site.<sup>16</sup> Now, if we consider the angle variable  $3\theta$  and the corresponding complex order parameter  $\Psi_3 \equiv \exp(i3\theta)$ , the degenerate ground states are all completely ordered in terms of this new order parameter. Also, it has been shown that the system undergoes a KT transition at a finite temperature, where the spatial correlation of the order parameter exhibits an algebraic decay below the transition temperature.

As for the KT transition temperature of this system, an analytic approximation was given as<sup>10,11</sup>



(a)



(b)

FIG. 1. Schematics of kagome lattices with the so-called (a)  $\mathbf{q}=0$  ground state and (b)  $\mathbf{q}=\sqrt{3} \times \sqrt{3}$  ground state.

$$T_c = \frac{\pi\sqrt{3}}{72} J \approx 0.0756J. \quad (2)$$

Numerical simulations were performed by Rzechowski<sup>9</sup> where he found two slightly different estimates on the transition temperature, i.e., one estimate of  $T_c \approx 0.076J$  based on the Binder cumulants of the order parameter and another of  $T_c \approx 0.070J$  from the helicity modulus [or correspondingly the decay exponent  $\eta(T_c) = 0.25$  of the spatial correlation of the order parameter].

In the present work, the coarsening dynamics of the model system is performed via kinetic Monte Carlo methods with standard Metropolis algorithm for quenches to various temperatures near and below the KT transition. System sizes ranging up to  $N \times N = 256 \times 256$  were employed with periodic boundary conditions. In a kagome lattice, the number  $\tilde{N}$  of the total spins is  $\tilde{N} = \frac{3}{4}N^2$ . The system is quenched from completely disordered initial states down to a given low temperature with the process of coarsening being monitored through equal-time spatial correlation functions, the decay of topological defects, etc.

In addition to quenches from the disordered state to low temperatures, we also performed the so-called nonequilibrium relaxation by suddenly bringing the system from one of the ground states to some target temperatures around or below the KT transition. This method was found to be convenient for measuring the values of the critical exponent  $\eta$  for the equilibrium spatial correlation functions.

When we let the systems evolve from random initial configurations, the following quantities can be measured:

(1) The number density  $n_v(t)$  of topological defects which are  $3\theta$  vortices in the *KAFX* model

$$n_v(t) \equiv \frac{\langle N_v(t) \rangle}{\tilde{N}}, \quad (3)$$

where  $N_v(t)$  is the total number of  $3\theta$  vortices (both positive and negative) at time  $t$  and  $\tilde{N}$  denotes the total number of sites (i.e., spins).  $\langle \dots \rangle$  denotes an average over random initial configurations. We also count the separate number density of vortices residing on the hexagonal plaquettes ( $n_h$ ) and those on the triangular plaquettes ( $n_t$ ).

(2) The equal-time spatial correlation function of the  $3\theta$  order parameter  $\psi_3(\mathbf{r}, t) \equiv e^{3i\theta(\mathbf{r}, t)}$ ,

$$C(r, t) = \langle \psi_3^*(\mathbf{r}, t) \psi_3(0, t) \rangle, \quad (4)$$

$$= \frac{1}{\tilde{N}} \left\langle \sum_i \exp[3i\theta_i(t) - 3i\theta_{i+r}(t)] \right\rangle. \quad (5)$$

(3) Nonequilibrium spin autocorrelation functions,

$$A(t) = \frac{1}{\tilde{N}} \left\langle \sum_i \exp[i\theta_i(0) - i\theta_i(t)] \right\rangle, \quad (6)$$

$$A_3(t) = \frac{1}{\tilde{N}} \left\langle \sum_i \exp[3i\theta_i(0) - 3i\theta_i(t)] \right\rangle. \quad (7)$$

### III. SIMULATION RESULTS

We have performed dynamic Monte Carlo simulations of the *KAFX* model on a kagome lattice of dimensions  $256 \times 256$ . In order to obtain the values of the equilibrium critical exponents  $\eta$  for the spatial decay of the equilibrium spatial correlation function  $C_{\text{eq}}(r)$  of the  $3\theta$  order parameter, we have employed the so-called *nonequilibrium relaxation* (NER) method, where the system is suddenly brought from ground states to finite temperatures below or near the KT transition. Simulations were performed up to 655 360 Monte Carlo steps which was sufficient for the equilibrium to be attained. This was confirmed by the collapse of the spatial correlations at later time stages as shown in Figs. 2(a) and 2(b). In Fig. 2(c) the correlation functions  $C_{\text{eq}}(r)$  obtained in this way are displayed on a log-log scale. The values of the exponent  $\eta$  thus determined for temperatures ranging from  $T=0.01$  to  $T=0.074$  are shown in Fig. 2(d) as well as in Table I.

As for the generation of the initial low-energy configurations, we first select one of the three phases of  $0, \pm 2\pi/3$  for

each site at random, and then applied Monte Carlo procedure at very low temperature of  $T=0.005$  for 10 000 MC steps with the restriction that in each of the MC procedure, a new trial phase is chosen only among the above three phases. In this way, the final states obtained were very close (in energy) to a ground state. These states were employed as the initial states of NER in order to bring the system to the equilibrium at some given target temperature.

From the graph of  $\eta(T)$  shown in Fig. 2(d), we find that the exponent increases almost linearly in temperature up to around  $T \approx 0.065$ , above which it starts to increase rather sharply. The linear regime can be fit approximately by  $\eta(T) \approx 3.9T$ . We also find that the exponent  $\eta$  takes the value of  $1/4$  near  $T(\eta=1/4) \approx 0.064J$  (Table I). This temperature is expected to correspond to the KT transition, which apparently is a little lower than the theoretical<sup>10,11</sup> or the numerical<sup>9</sup> estimates of earlier works mentioned above. We attribute the discrepancy between the theoretical estimate and our numerical result to the possibility that the theoretical estimate of the transition temperature in the present model can overestimate the value of  $T_c$  as is also the case in the usual ferromagnetic XY model.<sup>3</sup> The value reported by Rzechowski ( $T_c \approx 0.070$ ) (Ref. 9) is closer to our result, though still around 9% discrepancy. We would like to emphasize that our simulations are based on much larger systems sizes [with  $256 \times 256 \times (3/4) = 49\,152$  sites] which is about 160 times larger than the largest system size employed by him.

Now, the coarsening dynamics of the model system under quench to low temperature from disordered initial states is investigated by monitoring the equal-time spatial correlation functions of the  $\psi_3$  order parameter. One of the most important features of typical coarsening dynamics toward quasicrystalline phase is the critical dynamic scaling

$$C(r, t) = r^{-\eta(T)} f(r/L(t)), \quad (8)$$

where  $f(x)$  is the scaling function and  $L(t)$  is the growing length scale. It should be noted that the above scaling ansatz is based on the existence of a single growing length scale  $L(t)$ .

Figures 3(a) and 3(b) show the equal-time correlation functions  $C(r, t)$  of the *KAFX* model for different time instants (from  $t=10$  to 655 360) at the temperature  $T=0.060$ . For these data, we attempted a critical dynamic scaling. The procedure of rescaling is as follows. For a given temperature, we take the value of  $\eta(T)$  (determined in Table I) and evaluate the combination  $\tilde{C}(r, t) \equiv r^{\eta(T)} C(r, t)$ . Then, for a given time instant, the length scale  $L(t)$  is determined by the condition  $\tilde{C}[r=L(t), t] = C_0$  where the constant  $C_0$  is chosen as  $C_0=0.2$  or  $C_0=0.3$  in this work. By plotting the scaled correlation functions  $\tilde{C}(r, t)$  in terms of the rescaled distance  $r/L(t)$ , we can check whether the critical dynamic scaling holds or not from the quality of the collapse of the scaled correlation functions. Even though we have arbitrarily chosen  $C_0=0.2$  or  $C_0=0.3$  and attempted scaling collapse for each of these, actually just one arbitrary value of  $C_0$  is enough for checking the scaling collapse. This is because if

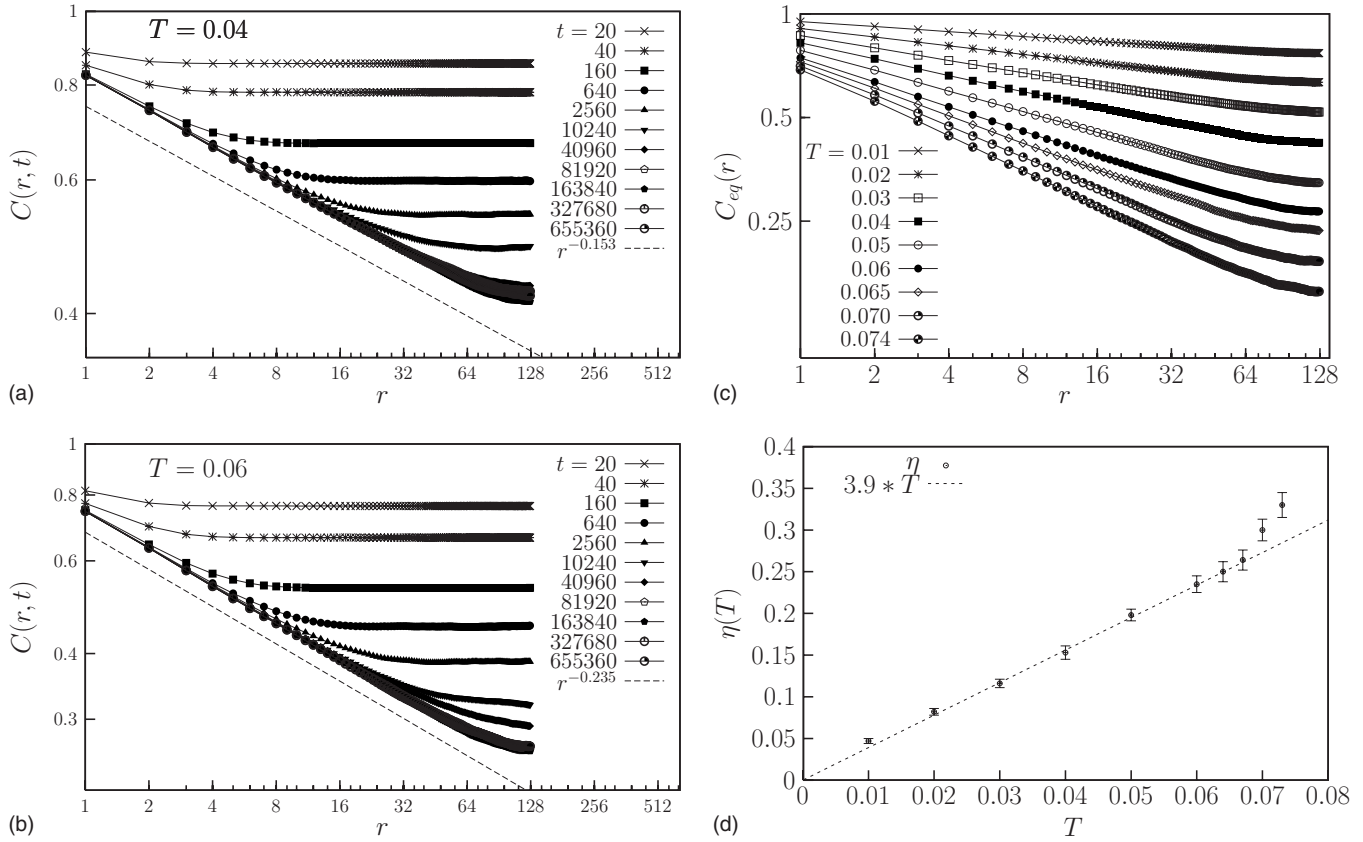


FIG. 2. [(a) and (b)] Time evolutions of the spatial correlation functions of  $\exp(3i\theta)$  at temperatures (a)  $T=0.04$  and (b)  $T=0.06$  from the NER method. Note that three or four spatial correlations at the latest time stages (corresponding to the Monte Carlo steps later than around  $10^5$ ) are almost collapsed with one another. (c) Equilibrium spatial correlations of  $\exp(3i\theta)$  at various temperatures and (d) the corresponding exponents  $\eta(T)$  vs  $T$ , obtained from the NER method. In (d) the dotted line represents  $3.9T$  which fits reasonably well the behavior of  $\eta(T)$  vs  $T$  especially at low and intermediate temperature regime ( $T \leq 0.065$ ).

the scaling works for one value of  $C_0$ , it would work for any other value of  $C_0$  which corresponds to a simple rescaling of the scaling function.

By this procedure, we found, rather unexpectedly, that the critical dynamic scaling does not hold in the coarsening dynamics of the *KAFXY* model, at least for the (Monte Carlo) time duration of our simulations (up to 655 360 MC steps).

TABLE I. The equilibrium exponent  $\eta$  for different temperatures for the power-law decay of the spatial correlation of the  $3\theta$  order parameter.

$T/J$	$\eta(T)$	$T/J$	$\eta(T)$
0.01	0.047(3)	0.065	0.254(11)
0.02	0.082(4)	0.066	0.259(13)
0.03	0.116(5)	0.067	0.264(12)
0.04	0.153(8)	0.068	0.268(12)
0.05	0.198(7)	0.069	0.280(11)
0.06	0.235(9)	0.070	0.300(13)
0.061	0.237(10)	0.071	0.308(12)
0.062	0.240(11)	0.072	0.310(13)
0.063	0.245(10)	0.073	0.330(15)
0.064	0.250(12)	0.074	0.340(13)

The result of the scaling attempt is shown in Fig. 3(c) for the case of  $T=0.06$  where we can see that the critical dynamic scaling is not obeyed. Details of the deviation from ideal scaling collapse are displayed in Figs. 3(g) and 3(h), where even in the latest time stages of our simulation time window, the scaled spatial correlation functions exhibit systematic drift. We also tried different values of  $\eta$  for the rescaled correlation with no success. It might be possible that the scaling is restored in the limit of infinitely long time.

Even though the critical dynamic scaling is not faithfully obeyed, we may still extract approximate length scale  $L(t)$  of spatial correlation in the manner described above. Now, strictly speaking, the time dependence of this length scale would depend on the choice of the value of  $C_0$  (since the dynamic scaling is not satisfied). We still proceeded to look into the behavior of these length scales for the two values of  $C_0=0.2$  and  $0.3$ , where we could not find much difference in the time-dependent behavior of  $L(t)$  for the two cases (see below). The length scale  $L(t)$  thus obtained exhibits a rather unusual time-dependent behavior [Fig. 3(d)]. That is, in the early-time stage up to around  $t \sim 10^3$ ,  $L(t)$  exhibits a slow growth, which appears approximately independent of the temperature. However, in the intermediate and late time stage of  $t \geq 10^3$ , the growth of the length scale  $L(t)$  becomes strongly dependent on the temperature. In addition, for a given temperature, no simple power law is found which is

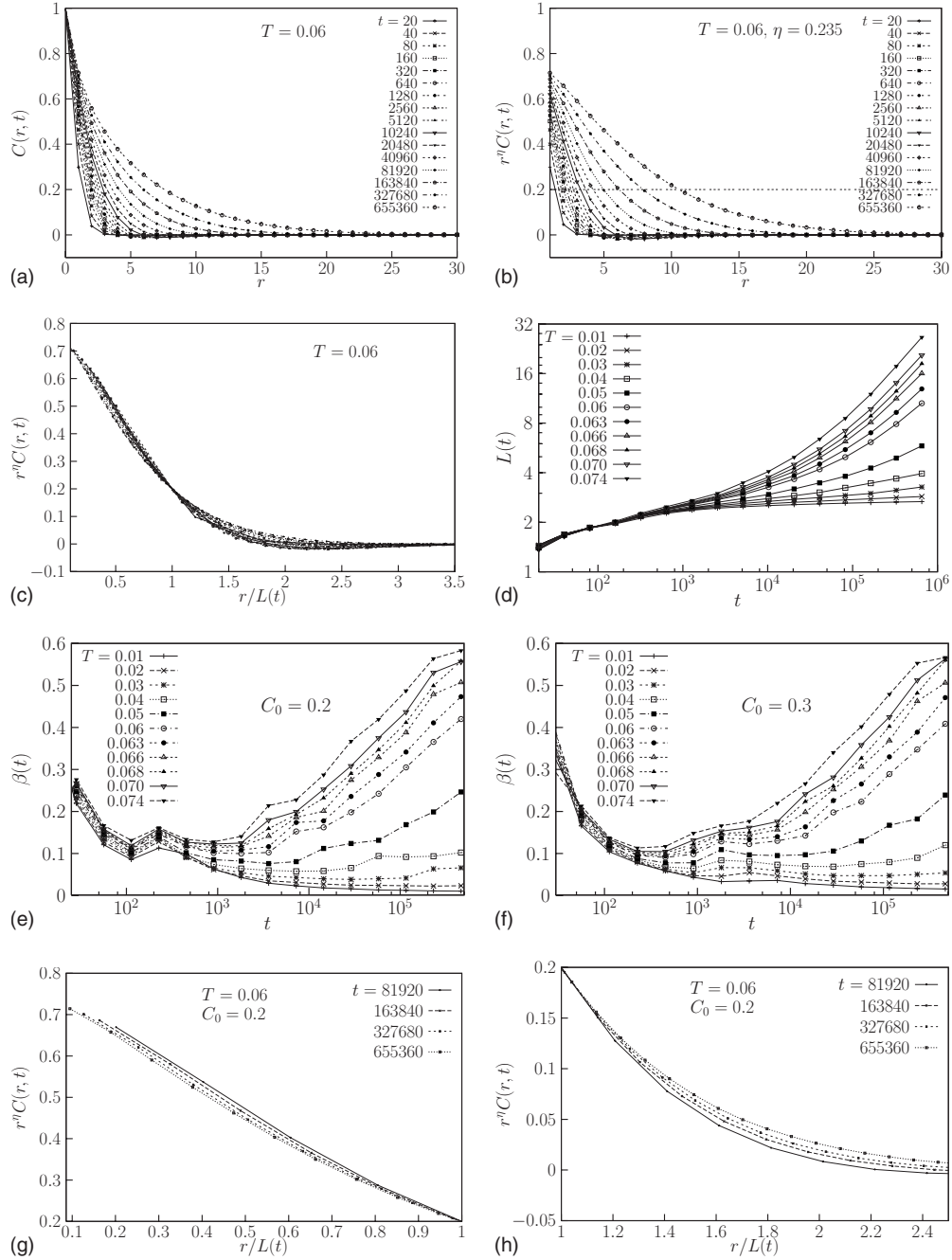


FIG. 3. (a) Equal-time spatial correlation functions  $C(r,t)$  for  $\exp(3i\theta)$  for different time instants at  $T=0.06$ , and (b) the rescaled functions  $\tilde{C}(r,t) \equiv r^\eta C(r,t)$ . A scaling attempt is shown in (c) based on (b) with  $C_0=0.2$  with the resulting growth of the length scale (using  $C_0=0.2$ ) in (d) and also the effective growth exponents vs time (using  $C_0=0.2$ ) in (e). Similarly, the effective growth exponents vs time with  $C_0=0.3$  is shown in (f). (g) and (h) show the enlarged view of the scaling attempt in (c) for the spatial correlation functions at four latest time instants for shorter distance region (g) and longer distance region (h), respectively, exhibiting systematic violation of scaling even in the latest time regime.

valid for the whole late time regime. Instead the local logarithmic slope exhibits a steady increase in time in the late time regime.

We can investigate the time dependence of the growth of the length scale by defining the effective local growth exponent as  $\beta(t) \equiv d \ln(L(t)) / d \ln(t)$ . Since we took the growing length scale  $L(t)$  only for the discrete-time instants  $t_i$  with fixed interval in the logarithmic scale, we evaluate the dis-

crete version of the above logarithmic slope as

$$\beta(t'_i) \approx \frac{\ln[L(t_{i+1})/L(t_i)]}{\ln(t_{i+1}/t_i)}, \quad (9)$$

where  $t_i \equiv 2^i$ ,  $i=1, 2, \dots$ , and  $t'_i \equiv \sqrt{t_i t_{i+1}}$ . Figures 3(e) and 3(f) show the effective growth exponent at time  $t$  (for the case of  $C_0=0.2$  [Fig. 3(e)] and  $C_0=0.3$  [Fig. 3(f)], respectively). We

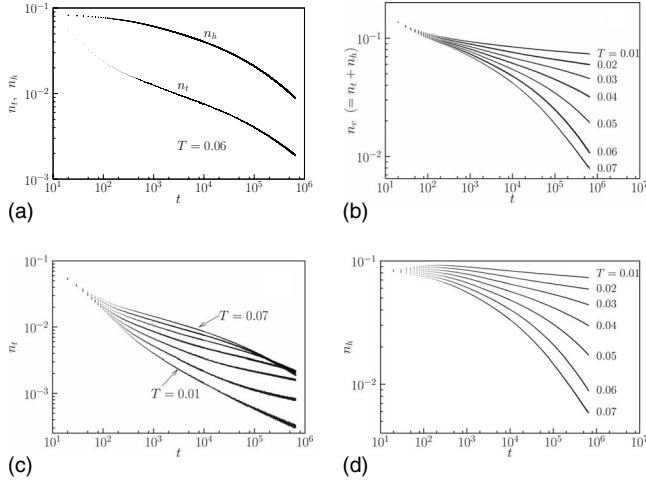


FIG. 4. (a)  $3\theta$  fractional vortex (defect) density vs time  $t$  on hexagonal and triangular plaquettes, respectively, for  $T=0.06$ , (b) the total vortex number density (both on hexagonal and triangular plaquettes) vs time  $t$  (Ref. 1) for  $T=0.06$ , (c) the time dependence of the (Ref. 1) density of  $3\theta$  vortices on triangular plaquettes at various temperatures, and (d) the time dependence of the density of  $3\theta$  vortices on hexagonal plaquettes at various temperatures.

can see that in the late time regime, the growth becomes definitely faster as the temperature is higher. Also, for temperatures  $T \geq 0.05$ , the effective local growth exponent appears to be monotonically increasing in the late time stage. At higher temperatures, we can recognize some indications from Figs. 3(e) and 3(f) that the effective slopes tend to converge around 0.5–0.6 in the latest time regime. It may be

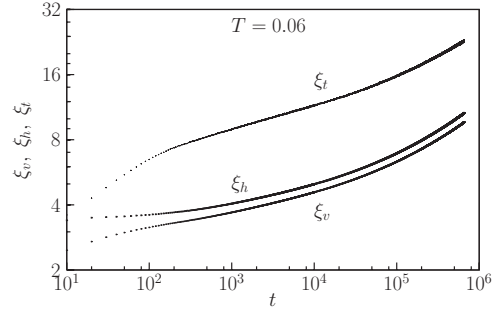


FIG. 5. The length scales  $\xi_h$ ,  $\xi_t$ , and  $\xi_v$  derived from the vortex densities versus time  $t$  for hexagonal, triangular plaquettes, and the sum of these, respectively at  $T=0.06$ .

possible to interpret this as an indication of an asymptotic behavior of diffusive growth.

In order to understand the physical mechanism underlying the breakdown of the critical dynamic scaling as well as the peculiar time dependence of the length scale, we investigated the time dependence of the total number of the relevant topological defects, i.e., the fractional  $3\theta$  vortices residing on the triangular plaquettes as well as on the hexagonal plaquettes of the kagome lattice. The results are shown in Figs. 4(a) and 4(d) where we can see that for a given temperature, the triangular vortices are decaying much faster than the hexagonal vortices. We also find that in general (for both types of the vortices) the decay of the total vortex numbers do not exhibit a simple power-law behavior valid for the whole time range. One prominent feature in terms of the temperature dependence of the decay of the fractional vortex

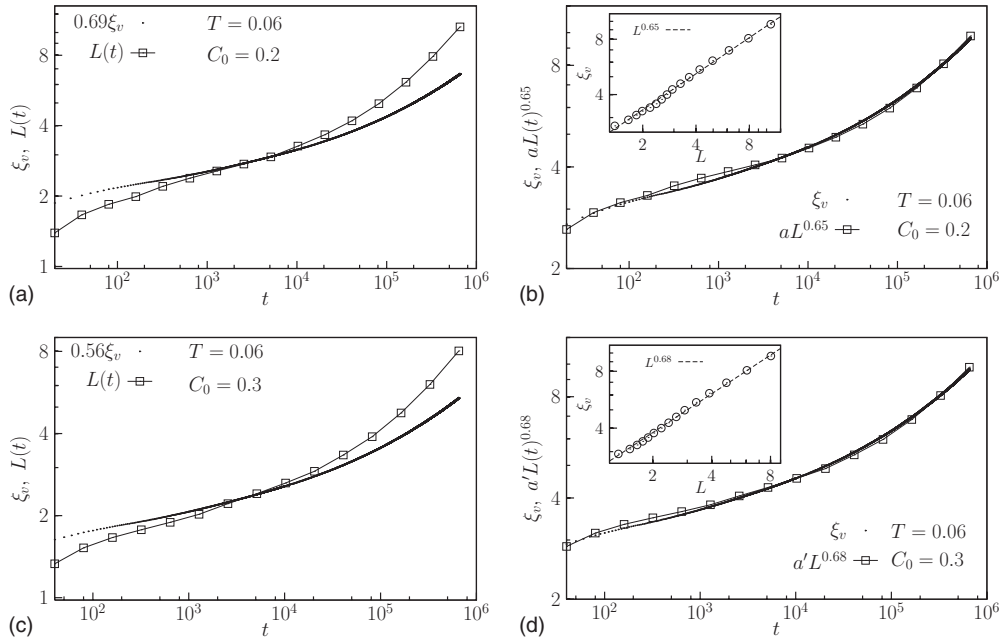


FIG. 6. (a) The length scale  $\xi_v$  derived from the total vortex density is compared with the domain length scale  $L(t)$  obtained from the equal-time spatial correlations with  $C_0=0.2$  at  $T=0.06$ , where one can see that the two (Ref. 1) length scales are not proportional to each other (Ref. 1). In (b), a comparison of  $L(t)^{0.65}$  (multiplied by a constant factor) with  $\xi_v$  is shown, which exhibits an approximate collapse (also see the inset for direct comparison between  $\xi$  vs  $L$ ). Similar results with  $C_0=0.3$  are displayed in (c) and (d) where a comparison of  $L(t)^{0.68}$  (multiplied by a constant factor) with  $\xi_v$  is shown with reasonable collapse (d) (see also the inset). Note that in (a) and (c)  $\xi_v$  is multiplied by a constant for convenience of comparison.

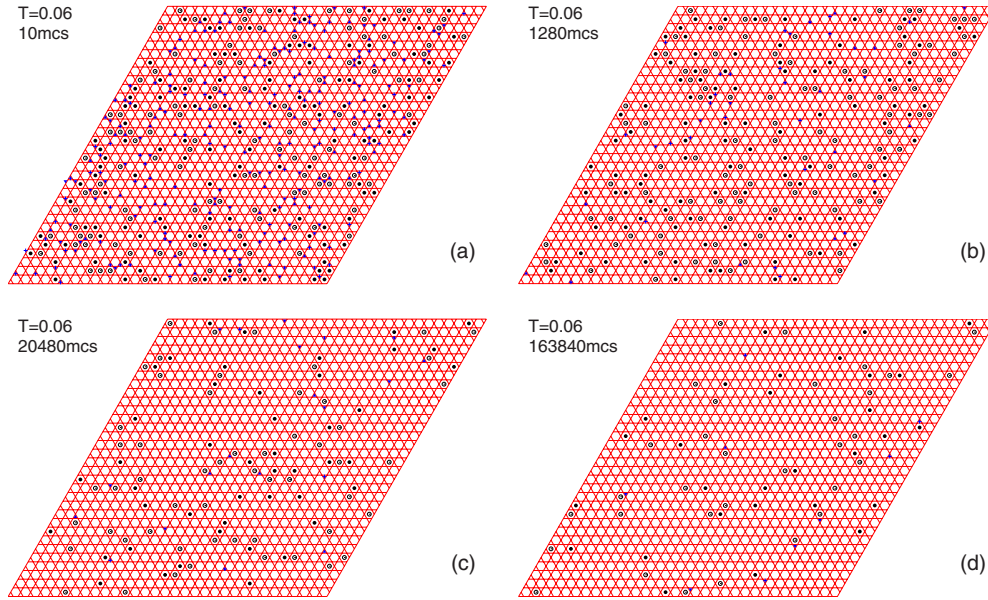


FIG. 7. (Color online) Configurations of  $3\theta$  vortices at different time steps for (a)  $t=10$  MCS, (b)  $t=1280$  MCS, (c)  $t=20480$  MCS, and (d)  $t=163840$  MCS. Filled (empty) circles represent positive (negative) vortices on hexagonal plaquettes and pluses represent positive vortices on the triangular plaquettes while filled triangles negative vortices (on the triangular plaquettes).

density is that for the vortices on the triangular plaquettes, the decay rate increases as the temperature is lowered, while on the other hand, those vortices on the hexagonal plaquettes exhibit opposite dependence on the temperature with the decay rate decreasing sharply as the temperature is lowered.

This can be interpreted as implying that there is almost no (free) energy barrier for the motion and decay of the triangular vortices but that some finite barrier exists for the hexagonal vortices. We also find an interesting feature of the hexagonal vortices (at lower temperatures) where, at initial stage, they increase a little and then start to decay. This can probably be understood as the influence of the decay of the neighboring triangular vortices with their excess energy turned over to neighboring hexagonal vortices, thus generating hexagonal vortices.

Therefore, we can conclude that for wide range of temperatures, the relaxation of the fractional vortices residing on the hexagonal plaquettes determines the coarsening process such as the correlation length scale corresponding to the quasiordered domains. In addition, the vortex relaxations exhibit a considerable deviation from power-law behavior with slowly increasing (in absolute magnitude) local logarithmic slope in the late time regime. Especially in the case of the hexagonal vortices, the decay of the vortex number density could be fit to a form with

$$N_h = N_0 \exp\{-b[\ln(t)]^\alpha\}, \quad (10)$$

$$= N_0 \exp\left[-\left(\frac{\ln(t)}{\ln(t_0)}\right)^\alpha\right], \quad (11)$$

with  $b \equiv \ln(t_0)^{-\alpha}$ . Here, we found that typically  $\alpha$  takes values around 3.3–4.0 (figure not shown here<sup>18</sup>). We found that it could also be fitted to an stretched exponential form as

$$N_h(t) \sim N_0 \exp[-(t/t_0)^\alpha], \quad (12)$$

with  $\alpha \approx 0.226 \pm 0.03$ .

It appears that the breaking of the critical dynamic scaling is related to the anomalous relaxation of the number of the fractional vortices on the triangular and the hexagonal plaquettes. In order to check this possibility, we analyzed the behavior of the length scales determined by the vortex distributions as follows. We define the length scale  $\xi_t \equiv 1/\sqrt{n_t}$  which corresponds to the average separation between the vortices on the triangular plaquettes. Similarly, we can define the length scale  $\xi_h \equiv 1/\sqrt{n_h}$  corresponding to the average separation between vortices on the hexagonal plaquettes. We also define the length scales  $\xi_v \equiv 1/\sqrt{n_v} = 1/\sqrt{n_h + n_t}$  which represents the length scale corresponding to the total number density of vortices.

Figure 5 shows the three length scales at  $T=0.06$  which clearly shows the faster growth of the length scale for the

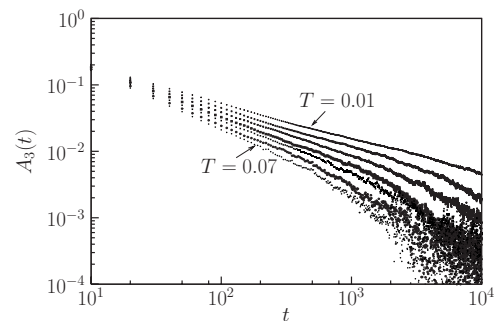


FIG. 8. Autocorrelation functions for  $\exp(3i\theta)$  for different temperatures. We can see that the autocorrelations exhibit approximate power-law decay up to intermediate time stages and then faster decay in the late time stage.

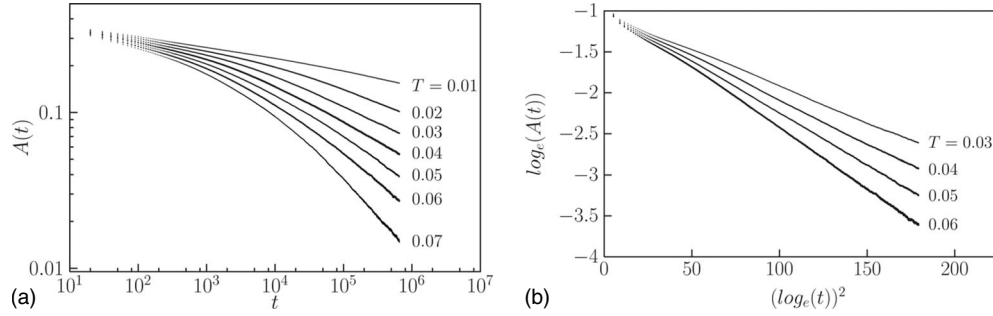


FIG. 9. (a) Autocorrelation functions for  $\exp(i\theta)$  for different temperatures, where we can discern clear signs of faster-than-power-law decay and (b) the plots for the logarithm of the autocorrelation functions for  $\exp(i\theta)$  vs  $[\log_e(t)]^2$  which show approximate linear relationships.

triangular vortices compared with that for the hexagonal vortices. In Figs. 6(a) and 6(d) comparison is made between the length scales  $L(t)$  derived from the spatial correlation functions and the length scales  $\xi_v$  derived from the total vortex densities. We see that (for both cases of  $C_0=0.2$  and  $C_0=0.3$ ) there exists no simple proportional relationship between the two length scales. Rather, the length scale  $L(t)$  from the spatial correlation is seen to grow faster than the vortex length scale  $\xi_v$ . This probably implies that the vortices are not distributed evenly (statistically speaking) in the system but that the vortices are somehow distributed in a non-random manner such that some degree of clustering occurs. An interesting result is that the two length scales satisfy some nontrivial relationships such that  $\xi_v \sim L(t)^\lambda$  with  $\lambda \approx 0.65$  for the case of  $C_0=0.2$  [Fig. 6(b)] and  $\lambda \approx 0.68$  for the case of  $C_0=0.3$  [Fig. 6(d)]. This means that the vortex configuration of the system does not exhibit simple self-similarity at different time instants. Rather the vortices probably tend to cluster more unevenly as time passes by, leading to the correlation length scale  $L(t)$  growing faster than the length scale derived from the vortex density. We emphasize again that due to the lack of perfect dynamic scaling, the time dependence of the length scale  $L(t)$  actually depends on the choice of the value of  $C_0$ . Hence, the above comparison between  $L(t)$  and other length scales should only be considered as approximate.

The snapshot of defect configurations for the case of  $T=0.06$  is shown in Figs. 7(a) and 7(d). We can easily recognize a faster decay of those vortices on the triangular plaquettes. From the snapshots alone, however, it is not easy to detect the tendency of relative clustering of the vortices in the late time regime.

Now we turn to the autocorrelation functions of the variables  $\exp(i\theta)$  and  $\exp(3i\theta)$ . For these two variables, we find that the simulation results on the autocorrelations are very different. For the case of  $\exp(3i\theta)$  (Fig. 8), we obtain an approximately power-law decay behavior  $A_3(t) \sim t^{-\lambda}$  in the early time regime up to around  $t \approx 10^3$ . The value of the exponent  $\lambda$  ranges from  $\lambda \approx 0.56$  (for  $T=0.01$ ) to  $\lambda \approx 0.90$  (for  $T=0.07$ ). The exponent tends to increase slightly at

higher temperatures. At later time, however, the autocorrelations decay faster than the power-law relaxation of the early time stage. Due to the larger statistical fluctuations in the relaxation of the autocorrelation  $A_3(t)$ , we could not attempt a comparison of  $A_3(t)$  with the growing length scale  $L(t)$ .

In contrast, in the case of the phase  $\exp(i\theta)$ , the spin autocorrelation exhibits less statistical fluctuations with a nonpower law behavior<sup>17</sup> that can be reasonably fitted by

$$A(t) = A_0 \exp\{-b[\ln(t)]^\gamma\}, \quad (13)$$

$$= A_0 \exp\left[-\left(\frac{\ln(t)}{\ln(t_0)}\right)^\gamma\right], \quad (14)$$

with the exponent  $\gamma = 2.0 \pm 0.3$  and  $b \equiv \ln(t_0)^{-\gamma}$ . One of the simulation results is shown in Figs. 9(a) and 9(b) for the case of  $T=0.06$  with  $\tilde{N}$ , which shows a suitable fit to the above functional form with  $b \approx 0.15$  and  $\gamma \approx 2.0$ . Note that in the limit of  $\gamma=1$ ,  $A(t)$  reduces to a power law of  $A(t) \sim A_0 t^{-b}$ . The fitted values of  $\gamma$  implies that  $A(t)$  exhibits a considerable deviation from a power-law behavior. We also attempted to fit the simulation results to stretched exponential form. This resulted in rather small values for the stretching exponent of  $\alpha \approx 0.14$ . It is interesting to note that a similar behavior of relaxation in the autocorrelation function of the order parameter was reported in the coarsening dynamics of the so-called Hamiltonian XY model on a square lattice.<sup>19</sup>

In summary, we investigated the coarsening dynamics of the antiferromagnetic XY model on a kagome lattice. We found that the critical dynamic scaling is not obeyed at least up to the simulation time window of ours. The breakdown of the critical dynamic scaling is accompanied by the nontrivial decay behavior of the density of defects that cannot be fitted by simple power laws. The competition between the critical thermal fluctuations of the equilibrium and the infinite ground-state degeneracy probably gives rise to the nontrivial features of the relaxation dynamics. It may be possible to collapse the equal-time spatial correlation functions through a multiscaling scheme which we have not tried yet. It would also be interesting to investigate the aging dynamics of this model system.<sup>20</sup>



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