Berezinskii-Kosterlitz-Thouless phase transition for the dilute planar rotator model on a triangular lattice

Yun-Zhou Sun* and Lin Yi

Department of Physics, Huazhong University of Science and Technology, Wuhan City, Hubei Province, 430074, People's Republic of China

G. M. Wysin[†]

Department of Physics, Kansas State University, Manhattan, Kansas 66506, USA (Received 16 May 2008; published 8 October 2008)

The Berezinskii-Kosterlitz-Thouless (BKT) phase transition for the dilute planar rotator model on a triangular lattice is studied by using a hybrid Monte Carlo method. The phase-transition temperatures for different nonmagnetic impurity densities are obtained by three approaches: finite-size scaling of plane magnetic susceptibility, helicity modulus, and Binder's fourth cumulant. It is found that the phase-transition temperature decreases with increasing impurity density ρ and the BKT phase transition vanishes when the magnetic occupancy falls to the site percolation threshold: $1-\rho_c=p_c=0.5$.

DOI: 10.1103/PhysRevB.78.155409

PACS number(s): 07.05.Tp, 21.60.Ka, 05.10.Ln, 75.10.Hk

I. INTRODUCTION

It is well known that the Berezinskii-Kosterlitz-Thouless (BKT) phase transition^{1,2} caused by the unbinding of vortexantivortex pairs is found to be common in two-dimensional ferromagnetic spin models, such as the planar rotator model, XY model, and easy-plane Heisenberg model. Recently, the study of nonmagnetic impurities in these spin models has been the subject of much interest.^{3–7} It is found that the BKT phase-transition temperature decreases with increasing nonmagnetic impurity density ρ . However, a Metropolis algorithm Monte Carlo (MC) simulation³ of the planar rotator model on a square lattice showed that the phase-transition temperature T_C falls to zero for a magnetic occupation density $\rho_{\text{mag}} = 1 - \rho$ above the site percolation limit $p_c \approx 0.59$ on a square lattice. This phenomenon was also found in the dilute system of Josephson junctions.⁸ These results were somewhat surprising because they suggested that a mechanism besides the disordered (or disconnected) geometry of the impurity-occupied lattice was responsible for permanently leaving the magnetic system in a high-temperature disordered phase. On the other hand, through extensive hybrid MC simulation, Wysin *et al.*⁶ found that the phase-transition temperature drops to zero at $\rho_{\rm mag} \approx 0.59$, which is the site percolation limit where there is no more percolating cluster of spins. The disorder due to impurities, especially near the percolation limit, greatly increases fluctuations in the Monte Carlo, making hybrid schemes that include cluster and overrelaxation updates a necessity for precise results. This is counterintuitive because one might naively expect that the weaker connectivity of the lattice near the percolation limit should make the calculations easier. The problem is interesting, however, because just as the impurity concentration is getting near that limit that causes T_C to fall to zero, and the lattice is less and less connected, the correlation length is still diverging causing all the usual problems that MC faces near a critical point.

The previous works were all focused on the spin system for the square lattice but, of course, there is also a BKT phase transition in the planar rotator model on the triangular lattice. The site percolation threshold of a triangular lattice is $p_c = 0.5.9$ On a triangular lattice, cluster MC simulation of the pure system shows that the percolation temperature, defined as the temperature at which spanning clusters start to appear in a large system, equals the BKT phase-transition temperature for the planar rotator model.¹⁰ Therefore, the study of the BKT phase transition for the *dilute* planar rotator model on a triangular lattice is also interesting as a comparison to the square lattice. The main question to be answered is whether the BKT transition temperature falls to zero when the magnetic occupation density equals the site percolation limit or whether this happens at an occupation density *higher* than the percolation limit. We carry this out here using a hybrid MC simulation that includes over-relaxation and cluster moves, which are important especially when the system is near the percolation threshold, where Metropolis single-spin moves become inefficient.

II. METHOD AND RESULTS

The Hamiltonian of the dilute planar rotator spin model is considered $as^{3,6}$

$$H = -J\sum_{\langle i,j\rangle} \sigma_i \sigma_j \vec{S}_i \cdot \vec{S}_j = -J\sum_{\langle i,j\rangle} \sigma_i \sigma_j \cos(\theta_i - \theta_j).$$
(1)

Here J > 0 is the ferromagnetic coupling constant, θ_i are the angular coordinates of two spin components $\vec{S}_i = (S_i^x, S_i^y) = (\cos \theta_i, \sin \theta_i)$, and $\langle i, j \rangle$ indicate the nearest-neighbor sites. σ is taken to be 1 or 0 depending on whether the site is occupied or not. A hybrid MC approach, including Metropolis algorithm¹¹ and over-relaxation algorithm^{12,13} combined with Wolff single-cluster algorithm, ¹⁴ proved to be very efficient for the model defined by Eq. (1) on a square lattice.⁶

The processes used in the hybrid MC scheme work as follows. In a Metropolis single-spin update, one spin is selected randomly from total spins. The new candidate spin,



FIG. 1. Application of Binder's fourth order cumulant to estimate the phase-transition temperature for several lattice sizes at ρ =0.05. The inset shows the view near the estimated critical temperature.

obtained by a small increment to this spin in a random direction, is renormalized to unit length. Then the energy difference is obtained according to Eq. (1). The new spin is accepted or not according to a standard Metropolis judgment. An over-relaxation update reflects a spin selected randomly across the effective potential of its nearest neighbors defined as $\vec{B}_j = J \Sigma_j \sigma_j \vec{S}_j$ with the reflection effected by

$$\vec{S}_i \longrightarrow 2 \frac{\vec{S}_i \cdot \vec{B}_j}{\|\vec{B}_j\|^2} \vec{B}_j - \vec{S}_i.$$

This algorithm is nonergodic and it must be mixed with other updates to achieve ergodicity. The Wolff cluster algorithm, known to be very efficient for spin systems with continuous symmetry, is similar to the application to a pure system except that spins at vacant sites are set to zero length. In the actual simulation, a random bond can be constructed by choosing a random direction, projecting spins on that direction, and then assigning unit to spins of parallel or antiparallel projections to it. The over-relaxation and Wolff cluster algorithms generally reduce autocorrelations better than Metropolis single-spin updates especially at low temperatures where the spin components tend to freeze.

From an initial spin configuration by randomly occupying sites with probability $1-\rho$, the simulation is performed with periodic boundary conditions for system size $N=L^2$, where the size of lattice is considered as L=20, 30, 40, 60, and 80. Then the number of magnetically occupied sites is N_{mag} $=N(1-\rho)$. A hybrid MC step consists of one Wolff update of planar components of the spins followed by one Metropolis update and four over-relaxation updates, which change the configuration but keep the energy unchanged. For each algorithm of our scheme, one update is defined as attempting N_{mag} spin moves. During the simulation, 1×10^4 MC steps are used for equilibration and about 4×10^5 MC steps are used to get thermal averages at each temperature. To avoid correlations, measurements are taken every 2–6 MC steps. We used three different methods described in Ref. 6 to get



FIG. 2. Application of the finite-size scaling of in-plane susceptibility to estimate the phase-transition temperature for ρ =0.05.

the phase-transition temperature T_c . As the MC proceeds, some thermodynamics are observed. For adequately sized systems and small enough impurity density, averaging over different placements of the impurities makes little difference in the averages.⁶ Therefore, we did not average over different placements of impurities for large sizes when $\rho \leq 0.3$. While the presence of impurities tends to amplify finite-size effects, for large enough systems the fluctuations due to impurity disorder will become averaged out,⁶ and averages over impurity disorder become less necessary with increasing *L*. Therefore, the focus is on the temperature dependence of thermal averages especially for the larger systems. Also, in the scaling with system size (such as for the susceptibility), the fluctuations (error bars) caused by impurity disorder should diminish rapidly with increasing *L*.

According to the magnetization $M = (M_x, M_y) = \sum_i \sigma_i S_i$, the susceptibility and susceptibility components are given by

$$\chi = [\langle (M)^2 \rangle - \langle M \rangle^2] / N_{\text{mag}} k_B T, \qquad (2)$$

$$\chi^{\alpha} = \left[\langle M_{\alpha}^2 \rangle - \langle M_{\alpha} \rangle^2 \right] / N_{\text{mag}} k_B T, \qquad (3)$$

where k_B is the Boltzmann constant. With the average of χ^x and χ^y , the in-plane susceptibility is obtained as

$$\chi' = (\chi^x + \chi^y)/2.$$
 (4)

The Binder's fourth order cumulant is also defined via the usual relation¹⁵

$$U_L = 1 - \frac{\langle M^4 \rangle}{3 \langle M^2 \rangle^2}.$$
 (5)

In the figures, the error bars that are not visible indicate that the statistical errors are smaller than the symbols. For convenience, temperatures are measured in units of the exchange constant J.

Binder's fourth order cumulant U_L is used to estimate the location of T_C in the thermodynamic limit. At the phase-transition temperature, U_L is expected to be approximately independent of the system size. Therefore, T_C can be obtained from the crossing point of U_L for different lattice



FIG. 3. Helicity modulus as a function of temperature for several lattice sizes at ρ =0.05. The inset shows the view near the estimated critical temperature.

sizes. As an example, Fig. 1 shows U_L for different lattice sizes at ρ =0.05. The phase-transition temperature is estimated at T_C =1.328 ± 0.002. Generally speaking, this method overestimates T_C . With increasing L, the estimation of T_C will be more accurate. Due to the statistical uncertainties, however, more computing time is required to calculate near T_C especially in this model with nonmagnetic impurities.

Another method to estimate T_C proved to be very useful in many references.^{5,6,16,17} This method starts from the finitesize scaling analysis of the in-plane susceptibility χ' . Near and below T_C , the susceptibility scales with a power of the lattice size, $\chi' \propto L^{2-\eta}$, even in the presence of nonmagnetic impurities. The critical exponent is $\eta=1/4$ at the BKT phase-transition temperature for the planar rotator and XY models. Therefore, by using $\eta=1/4$ from the common point of intersection of the curves $\chi'/L^{7/4}$ vs T, the phasetransition temperature can be obtained. Figure 2 shows the application of this method at $\rho=0.05$. The estimation of T_C is 1.316 ± 0.003 , which is a little lower than the result from U_L .

In our practical application for this model, the statistical errors become even greater with the increase in ρ . More MC steps and averages are needed to reduce the errors when ρ increases especially near the site percolation threshold $p_c=0.5$. Therefore, using the largest lattice, we get T_c at high nonmagnetic impurity density based on the calculation of the helicity modulus.

The helicity modulus, Υ , obtained by a measure of the resistance to an infinitesimal spin twist Δ across the system along one coordinate, is an efficient method to calculate the BKT phase-transition temperature.^{18–20} An expression applicable to any general model Hamiltonian is⁶

$$\Upsilon = \frac{\langle \partial^2 H / \partial \Delta^2 \rangle}{N} - \beta \frac{\langle (\partial H / \partial \Delta)^2 \rangle - \langle \partial H / \partial \Delta \rangle^2}{N}, \qquad (6)$$

where $\beta = (k_B T)^{-1}$ is the inverse temperature. For the dilute planar rotator model defined by Eq. (1), based on the derivation process of Ref. 21, we can get the expression of helicity modulus on the triangular lattice,



FIG. 4. Helicity modulus as a function of temperature for lattice size L=80 at different nonmagnetic impurity densities ρ . The overall trend and the results near the percolation threshold are shown in (a) and (b) separately.

$$Y(T) = -\frac{\langle H \rangle}{\sqrt{3}N_{\text{mag}}} - \frac{2J^2}{\sqrt{3}k_B T N_{\text{mag}}^2} \\ \times \left\langle \left[\sum_{\langle i,j \rangle} (\hat{e}_{ij} \cdot \hat{x}) \sigma_i \sigma_j \sin(\theta_i - \theta_j) \right]^2 \right\rangle.$$
(7)

Here \hat{e}_{ij} is the unit vector pointing from site *j* to site *i*. \hat{x} is a selected basis vector in one coordinate. According to the renormalization-group theory,² there is a universal relation between the helicity modulus and the phase-transition temperature. T_C can be estimated from the intersection of the helicity modulus $\Upsilon(T)$ and the straight line $\Upsilon = 2k_BT/\pi$. Meanwhile, the MC data of $\Upsilon(T)$ will have a deeper drop in the critical region with increased lattice size. For larger lattice size, the intersection will be nearer to the critical temperature. Therefore, an overestimate of T_C generally will be obtained by this method. As an example, Fig. 3 shows the results for different lattice sizes at $\rho = 0.05$. It is clear that a BKT phase transition exists at finite temperature in this model. From the largest lattice size L=80, we estimate the critical temperature to be $T_C = 1.349 \pm 0.002$, which is a little higher than that from the in-plane susceptibility χ' . But that



FIG. 5. Application of the finite-size scaling of in-plane susceptibility to estimate the phase-transition temperature for ρ =0.45.

is to be expected, and only by going to much higher lattice size will the helicity-based approach give results that agree with those based on the susceptibility. It is noted that there are greater statistical fluctuations at smaller L, while the statistical errors are almost smoothed out at large size. The helicity modulus for different ρ at L=80 is shown in Figs. 4(a) and 4(b).

Due to the critical slowing down at low temperature, it is clearly seen that the statistical errors are greatest when the magnetic occupation density ρ_{mag} is near p_c , where there is no spanning cluster to appear. The helicity modulus almost disappears at ρ =0.50 as shown in Fig. 4. As a comparison, for example, Fig. 5 shows the results of finite-size scaling of in-plane susceptibility at $\rho = 0.45$. The statistical errors at small L are clearly more obvious than those at large L. The estimated phase temperature is $T_C = 0.126 \pm 0.01$, which is comparable with the result of helicity modulus as shown in Fig. 4. The last results for phase-transition temperature from the different methods mentioned above are summarized for different ρ in Fig. 6. We find that T_C is nearly linear with ρ and decreases with the increase in ρ . According to the linear fit of T_C from Y, we find that the BKT phase-transition temperature falls to zero near $\rho = 0.498 \pm 0.003$, which is almost the same as expected from the site percolation threshold p_c =0.5. This shows that the BKT phase transition vanishes when the magnetic site occupancy $1-\rho$ reaches the triangular lattice percolation limit-the same result as obtained for this model on a square lattice.

III. CONCLUSIONS

In summary, with the application of a hybrid MC simulation, the BKT phase-transition temperatures of a dilute planar rotator model on a triangular lattice are obtained by three



FIG. 6. Phase-transition temperature estimated by the three methods for different nonmagnetic impurity densities ρ . The straight line is a linear fit to T_C obtained from the helicity modulus.

methods. For the *pure* planar rotator model ($\rho=0$) on a triangular lattice as a test case, the phase-transition temperatures we obtained from χ' , Y, and U_L are 1.486 ± 0.002 , 1.498 ± 0.003 , and 1.503 ± 0.002 , respectively. These data are comparable with the result of high-temperature series studies.^{22,23} Consistent with MC simulations on the square lattice,⁶ it is found that the phase-transition temperature decreases as nonmagnetic impurity density ρ increases and falls to zero only when the magnetic occupancy falls to the site percolation limit. This is completely reasonable considering that the correlation length could not diverge, and hence, there would be no BKT phase transition if there were no percolating cluster spanning the system. On the other hand, as long as any percolating cluster is present, the BKT phase transition should appear but at a reduced temperature due to the diluted effective exchange couplings between distant spins. Meanwhile, we find that there is close to a linear relation between T_C and ρ . Stated simply, the BKT phase transition is present only when $\rho_{\rm mag} > p_c$, and the phase-transition temperature is approximately linearly proportional to (ρ_{mag} $-p_c)/(1-p_c)$. The calculations become more difficult when the magnetic density is close to the percolation limit due to the combined BKT and percolation fluctuations there. Thus, we could not rule out a weak deviation from this linear relationship when ρ_{mag} is very close to p_c .

ACKNOWLEDGMENTS

This work was supported by the National Natural Science Foundation of China under Grant No. 10774053 and the Natural Science Foundation of Hubei Province under Grant No. 2007ABA035. *syz1979@163.com

- [†]wysin@phys.ksu.edu; http://www.phys.ksu.edu/personal/wysin ¹V. L. Berezinskii, Sov. Phys. JETP **34**, 610 (1972).
- ²J. M. Kosterlitz and D. J. Thouless, J. Phys. C 6, 1181 (1973).
- ³S. A. Leonel, P. Zimmermann Coura, A. R. Pereira, L. A. S. Mól, and B. V. Costa, Phys. Rev. B **67**, 104426 (2003).
- ⁴B. Berche, A. I. Farińas-Sánchez, Yu. Holovatch, and R. Paredes V, Eur. Phys. J. B **36**, 91 (2003).
- ⁵G. M. Wysin, Phys. Rev. B **71**, 094423 (2005).
- ⁶G. M. Wysin, A. R. Pereira, I. A. Marques, S. A. Leonel, and P. Z. Coura, Phys. Rev. B **72**, 094418 (2005).
- ⁷L. M. Castro, A. S. T. Pires, and J. A. Plascak, J. Magn. Magn. Mater. **248**, 62 (2002).
- ⁸Y. E. Lozovick and L. M. Pomirchi, Phys. Solid State **35**, 1248 (1993).
- ⁹M. F. Sykes and J. W. Essam, J. Math. Phys. 5, 1117 (1964).
- ¹⁰ P. W. Leung and C. L. Henley, Phys. Rev. B **43**, 752 (1991).

- ¹¹N. Metropolis, A. W. Rosenbluth, M. N. Rosenbluth, A. H. Teller, and E. Teller, J. Chem. Phys. **21**, 1087 (1953).
- ¹²F. R. Brown and T. J. Woch, Phys. Rev. Lett. 58, 2394 (1987).
- ¹³M. Creutz, Phys. Rev. D **36**, 515 (1987).
- ¹⁴U. Wolff, Phys. Rev. Lett. **62**, 361 (1989).
- ¹⁵K. Binder, Z. Phys. B: Condens. Matter **43**, 119 (1981).
- ¹⁶A. Cuccoli, V. Tognetti, and R. Vaia, Phys. Rev. B **52**, 10221 (1995).
- ¹⁷L. A. S. Mól, A. R. Pereira, H. Chamati, and S. Romano, Eur. Phys. J. B **50**, 541 (2006).
- ¹⁸H. Weber and P. Minnhagen, Phys. Rev. B **37**, 5986 (1988).
- ¹⁹Y. H. Li and S. Teitel, Phys. Rev. B **40**, 9122 (1989).
- ²⁰ P. Olsson, Phys. Rev. Lett. **75**, 2758 (1995).
- ²¹D. H. Lee, J. D. Joannopoulos, J. W. Negele, and D. P. Landau, Phys. Rev. B **33**, 450 (1986).
- ²²M. Ferer and M. J. Velgakis, Phys. Rev. B 27, 314 (1983).
- ²³ P. Butera and M. Comi, Phys. Rev. B **50**, 3052 (1994).