Magneto-caloric effects, quantum critical points, and the Berezinsky-Kosterlitz-Thouless transition in 2D coupled spin dimer systems

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Spin dimer systems are a promising playground for the detailed study of quantum phase transitions. Using the magnetic field as the tuning parameter it is in principle possible to observe a crossover from the characteristic scaling near critical points to the behavior of a finite temperature phase transition. In this work we study two-dimensional coupled spin dimer systems by comparing numerical quantum Monte Carlo simulations with analytical calculations of the susceptibility, the magneto-caloric effect, and the helicity modulus. The magneto-caloric behavior of the magnetization with temperature can be used to determine the critical fields with high accuracy, but the critical scaling does not show the expected logarithmic corrections. The zeros of the cooling rate are an excellent indicator of the competition between quantum criticality and vortex physics, but they are not directly associated with the quantum phase transition or the finite temperature Berezinsky-Kosterlitz-Thouless transition. The results give a unified picture of the full quantum and finite temperature phase diagram.

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I. INTRODUCTION

The study of quantum phase transitions (QPT) remains a very active topic in many fields of physics, spurred by experimental progress to create novel tunable interacting systems. QPT occur in quite different materials, including heavy fermion compounds, unconventional superconductors, Mott insulators, coupled spin systems, and ultracold atoms. In particular, the common phenomenon of Bose Einstein condensation (BEC) of strongly interacting bosons by tuning the interaction or the chemical potential can now be found in a range of different physical systems. Ultracold atomic gases allow the tuning of interactions via Feshbach resonances, but also cross-dimensional phase transitions have been observed recently. Phase transitions in coupled spin dimer systems are prime examples of BEC of strongly interacting triplons which allow easy tuning of the chemical potential via the magnetic field. Although QPT’s occur at zero temperature as a function of a non-thermal control parameter such as the interaction, effective mass, or the chemical potential, a characteristic critical scaling with temperature can be observed in a large range above the critical point. In general a detailed analysis is necessary in order to understand how the critical behavior is reflected in the experiments and if the finite-temperature phase transition is affected in the vicinity the QPT, where thermal fluctuations are comparable to quantum fluctuations. Compared to bosonic gases of atoms and magnons the temperature control is relatively easy in triplon gases, which allows a systematic analysis of the critical scaling behavior near the QPT.

In this paper we focus on the theoretical analysis of quantum critical points of antiferromagnetic spin dimer systems which are weakly coupled in two-dimensions. Two QPT’s can be observed: As the field is increased through the lower critical value $B_c$ the spin dimers start to be occupied by triplons and the magnetization increases with characteristic two-dimensional logarithmic behavior. The second QPT corresponds to the saturation field $B_s$. The intermediate phase is characterized by long-range phase coherence of triplons at $T = 0$ and BKT behavior at finite $T$. Similar phase transitions occur in two-dimensional hard-core boson systems and in distorted frustrated lattices.

The schematic behavior is illustrated in Fig. 1. In this paper we show that the crossover from BKT behavior to critical scaling is rather well defined by the cooling rate and by characteristic maxima in the susceptibility. However, this crossover occurs at distinctly higher temperatures than the BKT transition which can be determined by a careful analysis of the spin-stiffness. There is no directly measurable signal for the BKT transition in experiments, but we find that magneto-caloric measurements are ideally suited to show the critical scaling behavior.

![Figure 1: (Color online) Schematic phase diagram.](image)
and pinpoint the exact location of the QPT. Close to the QPT the BKT transition retains the characteristic logarithmic behavior, albeit with strongly renormalized parameters. We find, however, that the low temperature behavior above the QPT’s does not fully follow theoretical expectations.

II. THE MODEL

We use a “columnar” arrangement of strongly coupled antiferromagnetic dimers \( J > 0 \) on a two dimensional square lattice as shown in Fig. 2 described by the Hamiltonian of localized spin-1/2 operators \( \hat{S}_{x,y} \)

\[
\hat{H} = \sum_{y=1}^{N_y} \left[ \sum_{x=odd}^{N_x} J \hat{S}_{x,y} \hat{S}_{x+1,y} + J_x' \hat{S}_{x+1,y} \hat{S}_{x+2,y} \right.

+ J_y' \sum_{x=1}^{N_x} \hat{S}_{x,y} \hat{S}_{x,y+1}] - B \sum_{i=1}^{N} \hat{S}_{i}^{2},
\]

where the inter-dimer couplings \( J_x' \) and \( J_y' \) can be ferromagnetic or antiferromagnetic, but are assumed to be small \( |J'| \ll J \).

A. Interacting boson models

Assuming that the intra-dimer exchange interaction \( J \) dominates over inter-dimer couplings \( J_x' \) and \( J_y' \), it is natural to represent the system in the singlet and triplet basis at each dimer site

\[
\begin{align*}
| t_{-}\rangle_i & = |\downarrow\downarrow\rangle_i \\
| t_{0}\rangle_i & = \frac{|\uparrow\downarrow\rangle_i + |\downarrow\uparrow\rangle_i}{\sqrt{2}} \\
| t_{+}\rangle_i & = |\uparrow\uparrow\rangle_i \\
| s\rangle_i & = \frac{|\uparrow\downarrow\rangle_i - |\downarrow\uparrow\rangle_i}{\sqrt{2}}.
\end{align*}
\]

At strong fields \( B \approx J \) the last two states become nearly degenerate, while the other two higher energy states will be neglected for now. It is therefore justified to work in a restricted Hilbert space with only two states at each dimer site, which are represented by hard-core bosons on the vacuum \( |0\rangle = \prod_i |s\rangle_i \) and \( b_i^\dagger |0\rangle = |t_{-}\rangle_i \prod_{j\neq i} |s\rangle_j \). In this Hilbert space the effective Hamiltonian describes strongly interacting bosons on a rectangular lattice

\[
\begin{align*}
H_{\text{eff}} &= \sum_{\langle i,j \rangle} \left[ -|t_{ij}| (b_i^\dagger b_j + b_j^\dagger b_i) + t_{ij}n_in_j \right] \\
&- \mu \sum_i n_i + U \sum_i n_i(n_i - 1),
\end{align*}
\]

where the limit \( U \to \infty \) is implied to satisfy the hard-core constraint. The effective chemical potential and the hopping in \( x \)- and \( y \)-directions are given by

\[
\mu = B - J, \quad t_x = J_x'/4, \quad t_y = J_y'/2.
\]

Note, that the hopping \( |t_{ij}| \) in Eq. (3) has been chosen to be positive, which can always be achieved by a local gauge transformation \( b_i \to (\omega_i) b_i \). The nearest neighbor interaction in Eq. (3) is repulsive (attractive) for \( J' > 0 \) (\( J' < 0 \)). By Fourier transforming the first term in the Hamiltonian the kinetic energy becomes

\[
H_{\text{kin}} = \sum_{\tilde{k}} (-2|t_x| \cos k_x - 2|t_y| \cos k_y) b_{\tilde{k}}^\dagger b_{\tilde{k}}
\]

The position of the upper and lower band edges allows a straightforward estimate of the critical fields \( B_c \) and \( B_s \). The lower critical field is determined by the chemical potential at which a single boson acquires positive energy \( -2|t_x| - 2|t_y| = \mu \), which gives

\[
B_c \approx J - |J_x'|/2 - |J_y'|,
\]

This estimate is only correct to first order in \( J' \), however, since the bosonic ground state (vacuum) is not an exact eigenstate of the full Hamiltonian in Eq. (1). Higher order corrections from the neglected triplet states \( |t_{-}\rangle \) and \( |t_0\rangle \) in Eq. (2) will be determined from numerical simulations as described below.

The upper critical field is determined from the energy gain of removing a particle from the fully occupied band including the nearest neighbor interaction energy

\[
B_s = J + |J_x'|/2 + |J_y'| + J_x'/2 + J_y',
\]

which is exact and corresponds to the saturation field of the original model (1). For intermediate fields \( B_c < B < B_s \) the physics is governed by the behavior of two-dimensional interacting bosons (BKT phase) as explained below.

B. The effective continuum model

We now focus on the lower QPT at \( B_c \) which corresponds to the well studied case of a dilute interacting
Bose gas. At low filling the critical behavior of the lattice model in Eqs. (3)–(4) is believed to be in the XY-universality class independent of the microscopic details. In the continuum limit the nearest neighbor interaction can be neglected and the hard-core constraint can be re-
placed by a strong $\phi^4$ interaction of a complex bosonic field $\phi(x, \tau)$ in a $(D + 1)$-dimensional Euclidean action

$$S = \int_0^\beta d\tau \int d^D x \left[ \frac{\partial}{\partial \phi} \phi \right]^2 - \frac{\tilde{\mu}^2}{2m} \phi + \frac{u_0}{2} |\phi|^4, \quad (9)$$

where $D = 2$ in our case. The parameters can be obtained by approximating the sums in Eqs. (3)–(4) by inte-
grals with a lattice spacing $a \approx 1$ and then rescaling

$$m = \frac{1}{2a^2 \sqrt{|t_x| |t_y|}}; \quad \tilde{\mu} = B - B_c; \quad u_0 = Ua^2. \quad (10)$$

In what follows we set the lattice spacing to unity $a = 1$.

The action in Eq. (9) describes an interacting dilute Bose gas with mass $m$ and chemical potential $\tilde{\mu}$. For $\tilde{\mu} > 0$ or $B > B_c$, a finite density of bosons appears even at zero temperature $T = 0$, which signals the QPT to the BKT phase. Analogously, the same model also applies at the upper critical field $B_s$, where it describes bosonic singlet excitations on the saturated state with $\tilde{\mu} = B - B_c$.

The upper critical dimensions are $D = 2$ for this model so that logarithmic corrections appear in this case, which are described in terms of an ultraviolet cutoff $\Lambda_0$ (of the order of the reciprocal rescaled lattice spacing). This situation ($D = 2$) has been analyzed extensively in the literature for various quantities which we summarize below. Other dimensions are discussed in the textbook of Sachdev.

The density of bosons corresponds to the magnetization per site $\langle \bar{\phi} \phi \rangle = 2M(B)/N$ in the spin dimer system as a function of field $\tilde{\mu} = B - B_c$, which is given at $T = 0$ by

$$M = \frac{m \tilde{\mu} \Theta(\tilde{\mu})}{8\pi} \ln \left[ \frac{\Lambda_0^2}{2m\tilde{\mu}} \right]. \quad (11)$$

The susceptibility is therefore

$$\chi = \frac{m}{8\pi} \left( \ln \left[ \frac{\Lambda_0^2}{2m\tilde{\mu}} \right] - 1 \right), \quad (12)$$

which is logarithmically divergent as the critical point is approached from above inside the BKT phase $B \rightarrow B_c$. For $T > 0$ and $B = B_c$ it has been predicted that the density increases with temperature including a characteristic logarithmic correction

$$M(T) = \frac{mT}{4\pi} \ln^{-4} \left[ \frac{\Lambda_0^2}{2mT} \right]. \quad (13)$$

The scaling as a function of $T$ can be used in order to identify the exact value of the critical field $B_c$ as outlined below.

Finally, the BKT transition temperature has been predicted as a function of field

$$T_{BKT} = \frac{\tilde{\mu}}{4} \ln \left[ \frac{\Lambda_0^2}{2m\tilde{\mu}} \right]. \quad (14)$$

However, for this formula to be valid the double loga-

III. DETERMINING THE CRITICAL FIELDS

In order to analyze the quantum phase transitions, the exact locations of the critical fields have to be determined first. As mentioned above, the upper critical field $B_s$ is exactly the saturation field in Eq. (5), but the lower field in Eq. (7) will in general have higher order corrections of the form

$$B_c \approx J - |J'_x|/2 - |J'_y| + a_x J_x^2 + a_y J_y^2 + a_{xy} J_x J_y. \quad (15)$$

The higher order corrections are due to virtual excitations to the neglected triplet states $|t_+ \rangle$ and $|t_0 \rangle$ in Eq. (2). The exact values for $a_x = -0.375$ and $a_y = 0.5J_y$ are known from higher order strong coupling expansions for the dimerized chain (16) ($J'_y = 0$) and for the ladder system (17) ($J'_x = 0$), respectively.

In order to determine the exact location of the QPT for general inter-dimer couplings, numerical simulations at $T = 0$ in the thermodynamic limit would be required. This is obviously impossible, but large systems sizes at small finite temperatures are feasible with Quantum Monte Carlo (QMC) simulations. In order to examine the model in Eq. (1) numerically, we therefore have implemented the Stochastic Series Expansion algorithm (22) with directed loop updates and using the so-called Merseene Twister random number generator (23).

At finite temperatures the discontinuity in Eq. (11) cannot be observed directly, but the magnetization as a function of temperature becomes exponentially small for $B < B_c$, while it approaches a finite value for $B > B_c$. The critical field $B_c$ is then exactly defined as the point where critical scaling is obeyed, which can be determined rather accurately. This behavior is illustrated in Fig. 3

Note, however, that the observed scaling in Fig. 3 at the exactly known upper critical field $B_s$ appears to be perfectly linear (relative to the saturated state). This means that the logarithmic correction in Eq. (13) must be very small, which puts a lower limit on the cutoff $\Lambda_0 \gtrsim 10^7$. To determine the lower critical field $B_c$, we therefore use linear scaling as well. Extrapolating the data to the thermodynamic limit and then determining
the critical fields $B_c$ by the best linear fit gives the results for three different choices of inter-dimer couplings shown in Table I. Ignoring higher orders, the values for the coefficients in Eq. (15) are then consistent with the following estimates

$$a_x = -0.375, \quad a_y = 0.5, \quad a_{xy} \approx -0.5 \pm 0.03 \quad (16)$$

Before continuing our analysis we would also like to consider how the neglected higher energy triplet excitations $\{t_-, t_0\}$ in Eq. (2) affect physical observables like the magnetization. We note that the effective Hamiltonian $\tilde{H} \chi J'$ is invariant under changing the inter-dimer coupling strengths $J'_x$ and $J'_y$ as long as all energies and the field $\mu$ are rescaled accordingly. We therefore consider three different realization of the coupling strength $J'_x = J'_y = J' = 0.05J, 0.1J, 0.2J$ and plot the susceptibility $\chi J'$ as a function of rescaled field $\mu/J' = (B - J)/J'$ at a given rescaled temperature $\beta J' = 5$ in Fig. 4. We observe a finite susceptibility in the BKT phase with two characteristic maxima near the QPT. While the three curves agree reasonably well, systematic deviations can be seen for larger $J'$, which can only come due to corrections from the higher energy triplet excitations. In what follows we choose a coupling strengths of $J'_x = J'_y = J' = 0.1J$, which is a compromise between minimizing higher order corrections and efficient numerical simulations. It is believed that the higher order triplet excitations do not change the form of the critical scaling in Eqs. (11)-(14).

![Figure 3](image3.png)

Figure 3: (Color online) Magnetization as a function of temperature for different magnetic fields for $J'_x = J'_y = 0.1$ and $N = 676$ near $B_c$ (top) and $B_s$ (bottom).

![Figure 4](image4.png)

Figure 4: (Color online) The susceptibility $\chi J'$ as a function of $\mu/J' = (B - J)/J'$ at inverse temperature $\beta J' = 5$ for three inter-dimer coupling strengths $J'_x = J'_y = J' = 0.05J, 0.1J$ and 0.2J.

IV. CRITICAL SCALING AT THE QPT

We now turn to analyzing the scaling behavior of the susceptibility $\chi$ as a function of field $B$ in Eq. (12). Finite temperatures $T$ and system sizes $N = L \times L$ play the role of an infrared cutoff $D_0 \sim \max(T, J'/L)$ which will give deviations from the predicted $T = 0$ scaling in Eqs. (11) and (12) as the QPT is approached. However, for fields $|B - B_{c/s}| \gtrsim D_0$ the scaling can still be tested. At each given temperature we first increase the system size until systematic convergence of the magnetization is obtained as shown in Fig. 5. The resulting susceptibility in the thermodynamic limit near the QPT’s is shown in Fig. 6 as a function of the logarithm of $\tilde{\mu} = |B - B_{c/s}|$ for different temperatures.

The data confirms that the scaling approaches a logarithmic behavior for $T \to 0$ consistent with the form in Eq. (12). We notice that the finite temperature susceptibility is actually rather small at the QPT $B = B_c$, but then increases and overshoots the logarithmic divergence, before the logarithmic behavior is reached inside.

| case          | $t_x [J]$ | $t_y [J]$ | $B_c [J]$ | from Eqs. | $\pm 0.0005$ |
|---------------|-----------|-----------|-----------|-----------|-------------|
| $J'_x = J'_y = 0.1J$ | 0.025 | 0.05 | 0.8460 | 0.84625 |
| $J'_x = 2J'_y = 0.15J$ | 0.0375 | 0.0375 | 0.8391 | 0.83875 |
| $2J'_x = J'_y = 0.12J$ | 0.015 | 0.06 | 0.8523 | 0.85225 |

Table I: Critical field $B_c$ for three different choices of exchange couplings, which obey the condition $J'_x + 2J'_y = 0.3J$, i.e. $B_c \approx 0.85J$ to lowest order.
the BKT phase. In this way the field-integral of the susceptibility (i.e. the magnetization) remains largely temperature independent outside the critical region, since the smaller values at the QPT for finite \( T \) are compensated by a corresponding overshooting of the maximum. In turn this means that the characteristic maxima in Fig. 1 of the susceptibility are only indirectly related to the QPT. The overshooting implies that the large fluctuations in the magnetization arise from a different mechanism at finite temperatures. One may expect that the maxima are therefore related to the finite-temperature BKT transition, but this is not the case as we will see below. Instead we find that the susceptibility maxima are found for temperatures well above the BKT transition \( T > T_{\text{BKT}} \) at the corresponding fields. As we will see later the maxima coincide with maxima in the entropy, so that these points correspond to the crossover between quantum critical scaling to vortex physics.

Comparing with the expected form in Eq. (12) quantitatively, we find rather small values of the effective mass \( m \approx 1.5/J \) at the lower QPT \( B_c \) and \( m \approx 2.2/J \) at \( B_s \), which are strongly renormalized compared to the naive estimate \( m \approx 14/J \) according to Eq. (11). The value of \( \Lambda_0 \sim 5 - 7 \) remains finite in Eq. (12). The value of \( m \) from the fits at the lower QPT is rather sensitive to the exact location of the critical field \( B_c \). In general all microscopic details such as the neglected next-nearest neighbor interaction in Eq. (13) will influence the exact value of the effective parameters in Eq. (10).

V. THE BEREZINSKY-KOSTERLITZ-THOULESS PHASE TRANSITION

The intermediate region between the two QPT’s is dominated by the presence of interacting triplon excitations which form a condensate at \( T = 0 \) with long range phase coherence. We now consider the finite temperature behavior in this intermediate phase. While the QPT’s are driven by quantum fluctuations, the transition due to thermal fluctuation corresponds to classical behavior and is therefore not directly related to the scaling discussed above.

The effective hard-core boson model in Eqs. (3)-(11) is exactly equivalent to the XXZ-spin model with \( J_z = J_{xy}/2 \), which is known to be in the XY-universality class. At finite temperatures this system undergoes a BKT transition, which can be described in terms of classical two-dimensional spins as first explained in the works of Berezinsky and Kosterlitz and Thouless. At low temperatures \( T < T_{\text{BKT}} \) a quasi long-range ordered phase with power-law decay of correlations exists. Above the phase transition temperature \( T_{\text{BKT}} \) the unbinding of vor-

Figure 5: (Color online) Susceptibility for different system sizes \( N \) and an inverse temperature of \( \beta J = 200 \) for \( J' = J_y' = 0.1J \) near \( B_c \) (top) and \( B_s \) (bottom).

Figure 6: (Color online) Susceptibility in the thermodynamic limit for different inverse temperatures \( \beta J \) and \( J' = J_y' = 0.1J \) near \( B_c \) (top) and \( B_s \) (bottom).
and shown in Fig. 8. The phase transition temperature $T_{\text{BKT}}$ can be used to define a so-called helicity modulus $\gamma$:

$$\gamma(T_{\text{BKT}}) = \frac{\hbar^2}{m^2} \rho_s(T_{\text{BKT}}) = \frac{2}{\pi} T_{\text{BKT}}.$$  

The phase transition temperature $T_{\text{BKT}}$ is then determined by the value where the helicity modulus obeys:

$$\gamma(T_{\text{BKT}}) = \frac{\hbar^2}{m^2} \rho_s(T_{\text{BKT}}) = \frac{2}{\pi} T_{\text{BKT}}.$$  

Generally the helicity modulus has a more complex relation to the spin-stiffness for anisotropic systems in low dimensions $D$ as discussed by Prokof’ev and Svistunov. For anisotropic systems it is therefore useful to define a separate helicity modulus for each direction:

$$\gamma_x = T \frac{L_x}{2} \langle \omega_x^2 \rangle, \\
\gamma_y = T \frac{L_y}{2} \langle \omega_y^2 \rangle,$$

where $L_x/2$ and $L_y$ are the edge lengths of the effective hard-core boson system in terms of the size of the original spin system $N = L_x \times L_y$. Instead of taking the average in Eq. (18), only the largest one contributes $\gamma = \max(\gamma_x, \gamma_y) = \gamma_x$, while the smaller one shows a linear behavior with edge length $\gamma_y \propto L_y$. The energy of the vortices also depend logarithmically on the system size $N$, so that the condition in Eq. (19) acquires a corresponding correction:

$$\frac{\pi \gamma_x(N, N_0)}{2T} = A(T) \left( 1 + \frac{1}{2} \frac{1}{\ln(N/N_0)} \right),$$

where $N_0$ is a fitting parameter and $A(T)$ should take on the universal value of unity at the transition, but can also be used as a fitting parameter. Following the procedure in Ref. 33, the logarithmic corrections in Eq. (21) become only accurate at the phase transition, which can in fact be used to determine $T_{\text{BKT}}$ and $A(T_{\text{BKT}})$. In Fig. 7 the helicity modulus is plotted at a given field $B = 0.92J$ for different system sizes. The BKT transition for each field is determined by the best fit of Eq. (21), i.e. when $\pi \gamma_x(N, N_0)/2T$ extrapolates to a limiting value linearly as a function of $\ln^{-1}(N/N_0)$. For a classical isotropic spin model a value of $A(T_{\text{BKT}}) = 1$ can be confirmed, but for the spin dimer model we find a field-dependent value for $A(T_{\text{BKT}})$ which is slightly larger than unity as given in Table 1 and shown in Fig. 7. The fitting parameter $N_0$ also becomes field dependent. The resulting transition temperature is shown in Fig. 8 which shows a sharp drop near the QPT. As shown in the inset the behavior is consistent with a logarithmic behavior:

$$\frac{T_{\text{BKT}}}{\tilde{\mu}} \approx \alpha \ln(b/\tilde{\mu}),$$
The deviations from $A(T_{\text{BKT}}) = 1$ can be traced to two different sources: In the middle of the BKT phase we find that a nearly isotropic effective system with $L_x = 2L_y$ and $J'_x = 2J'_y$ gives values of $A(T_{\text{BKT}}) \approx 1.04$, so that the detailed geometry appears to have some effect on the exact value of $A$. A second source may be higher order corrections in $\ln N/N_0$, which can be expected to become significant when the effective density of bosons per lattice site is small, which in turn leads to large distances between vortices. Therefore, the corrections must be largest close to the QPT, consistent with our findings. Using a constant value of $A(T_{\text{BKT}}) = 1$ in the fits changes the estimate for $T_{\text{BKT}}$ by up to 10-15%.

VI. MAGNETO-CALORICS AND THE $T$-$B$ PHASE DIAGRAM

As we already discussed in Sec. [III] the behavior of the magnetization $M(T)$ as a function of temperature plays an important role in determining the locations of the QPT. The interplay of magnetization with temperature is often termed magneto-calorics, which has been a fruitful field ever since the discovery of adiabatic demagnetization by Warburg in 1881. The central quantity of interest in this context is the cooling rate

$$\Gamma(B, T) = \frac{1}{T} \left( \frac{\partial T}{\partial B} \right)_S,$$

which describes the temperature change with the applied field under adiabatic conditions. Using the cyclical rule and a Maxwell relation the cooling rate is also directly related to $M(T)$ and $S(B)$

$$\Gamma(B, T) = -\frac{1}{C} \left( \frac{\partial S}{\partial B} \right)_T = -\frac{1}{C} \left( \frac{\partial M}{\partial T} \right)_B. \quad (24)$$

where $C = T \left( \frac{\partial S}{\partial T} \right)_B$ is the heat capacity. Therefore, the entropy is largest when $\Gamma = 0$.

The cooling rate for different temperatures is plotted in Fig. 9(top), which shows sharp features near the QPT. In Ref. [39] it was predicted that the cooling rate diverges with a universal prefactor near the QPT, but we were not able to reach low enough temperatures to confirm this behavior.

Integrating the cooling rate in Eq. (23) gives the temperature as a function of field for a given entropy $S$. The corresponding isentropes are shown in Fig. 9. The temperature reaches a minimum when the cooling rate is zero, which means that the entropy as a function of field (horizontal path) is maximal. It is interesting to notice that the points of maximum entropy $\Gamma = 0$ are

| $B/J$ | $A(T_{\text{BKT}}) \pm 0.03$ | $\ln (N_0) \pm 0.05$ | $T_{\text{BKT}}/J \pm 0.0005$ |
|------|----------------|----------------|----------------|
| 0.880 | 1.37 | 0.61 | 0.0103 |
| 0.920 | 1.29 | 0.32 | 0.0174 |
| 1.000 | 1.17 | 0.24 | 0.0239 |
| 1.080 | 1.14 | 0.22 | 0.0245 |
| 1.150 | 1.12 | 0.30 | 0.0229 |
| 1.200 | 1.14 | 0.38 | 0.0197 |
| 1.270 | 1.23 | 0.71 | 0.0092 |

Table II: Results of $A(T_{\text{BKT}})$ and $\ln (N_0)$ for different magnetic fields.
relatively close to the maxima of the susceptibility. However, the maximum entropy region is not exactly at the value of the critical field as is the case for other systems without an ordered phase, as in the Ising chain. Nor are those points associated with the finite temperature BKT phase transition as would be the case for ordered systems in $D > 2$. The situation in $D = 2$ is therefore special, since in this case the sign change in the cooling rate $\Gamma = 0$ signals a maximum in the entropy in the crossover region where quantum critical behavior competes with vortex excitations in the shaded parameter range in Fig. 9 (bottom).

VII. CONCLUSIONS

In summary, the magneto-caloric quantity $\partial M/\partial T$ turns out to be a universal indicator of the quantum critical behavior. We plot this quantity in Fig. 10 in the relevant $T$-$B$ parameter space. On the one hand we have seen in Sec. III that the critical scaling is defined by a linear behavior of $M(T) \propto T$, which leads to a constant and large derivative $\partial M/\partial T$. The regions with quantum critical behavior therefore show up clearly in Fig. 10 as the lightest and darkest region in the phase diagram above $B_0$ and $B_c$, respectively. The points of $\Gamma \propto \partial M/\partial T = 0$ mark the boundaries towards regions, which are dominated by BKT vortex excitations. These points coincide with the maxima in the susceptibility, but are not directly associated with the QPT or the finite temperature BKT phase transition. The BKT phase transition occurs at significantly lower temperatures and is not reflected by any directly measurable thermodynamic quantity. Nonetheless, the predicted and well established behavior of the spin stiffness at the BKT transition holds also for the dimer system, but strong corrections start to appear at small magnetization (i.e. boson density) as discussed in Sec. V.

We would like to emphasize that magneto-caloric measurements of $\partial M/\partial T$ not only allow a detailed analysis of the QPT, but also are potentially a very useful experimental tool in order to identify the effective dimensionality of the underlying spin systems due to the different density of states. In particular, for quasi one-dimensional systems $\partial M/\partial T \propto 1/\sqrt{T}$ shows a characteristic divergence above the QPT while for $D = 3$ we find an increase $\partial M/\partial T \propto \sqrt{T}$ analogous to the famous $T^{3/2}$ Bloch law. We find in our numerical simulations that $D = 2$ is characterized by perfectly linear behavior above the QPT, i.e. $\partial M/\partial T = \text{const.}$ without any detectable logarithmic corrections in contrast to the field theory prediction in Eq. (13). As discussed in Sec. III this can be used to determine the exact positions of the critical field, which in turn allows the quantitative estimate of higher order terms in the analytical expressions as a function of the antiferromagnetic coupling constants.

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1 A. Vogler, et al, Phys. Rev. Lett. 113, 215301 (2014).
2 Z. Hadzibabic, et al, Nature 441, 1118 (2006).
3 U. Tutsch, et al, Nature Comm. 5, 5169 (2014).
4 S. Sachdev, Science 288, 475 (2000).
5 C. Ruegg et al., Nature 423, 62 (2003).
6 S. Wessel, M. Olshanii, and S. Haas, Phys. Rev. Lett. 87, 206407 (2001).
7 K. Amaya, Y. Tokunaga, R. Yamada, Y. Ajiro, and T. Haseda, Physics Letters A 28, 732 (1969), ISSN 0375-9601.
8 M. Tachiki and T. Yamada, Journal of the Physical Society of Japan 28, 1413 (1970).
9 V. Berezinskii, Sov. Phys. JETP 32, 493 (1971).
10 V. Berezinskii, Sov. Phys. JETP 34, 610 (1972).
11 J. M. Kosterlitz and D. J. Thouless, Journal of Physics C: Solid State Physics 6, 1181 (1973).
12 J. M. Kosterlitz, Journal of Physics C: Solid State Physics 7, 1046 (1974).
13 K. Bernardet et al., Phys. Rev. B 65, 104519 (2002).
14 O. Derzhko, J. Richter, O. Krupnitska, and T. Krokhmal-skii, Phys. Rev. B 88, 094426 (2013).
15 S. Sachdev, Quantum Phase Transitions (Cambridge University Press, 2011).
16 S. Sachdev, T. Senthil, and R. Shankar, Phys. Rev. B 50, 258 (1994).
17 V. N. Popov, Functional Integrals in Quantum Field Theory and Statistical Physics (D. Reidel Publishing Company, 1983).
18 D. S. Fisher and P. C. Hohenberg, Phys. Rev. B 37, 4936 (1988).
19 S. Sachdev and E. R. Dunkel, Phys. Rev. B 73, 085116 (2006).
20 N. Prokof’ev and B. Svistunov, Phys. Rev. A 66, 043608 (2002).
21 N. Prokof’ev, O. Ruebenacker, and B. Svistunov, Phys. Rev. Lett. 87, 270402 (2001).
22 S. Sachdev and E. Demler, Phys. Rev. B 69, 144504 (2004).
23 S. Sachdev, Phys. Rev. B 59, 14054 (1999).
24 A. B. Harris, Phys. Rev. B 7, 3166 (1973).
25 T. Barnes, J. Riera, and D. A. Tennant, Phys. Rev. B 59, 11384 (1999).
26 M. Reigrotzki, H. Tsunetsugu, and T. M. Rice, Journal of Physics: Condensed Matter 6, 9235 (1994).
27 O. F. Syljuasen and A. W. Sandvik, Phys. Rev. E 66, 046701 (2002).
28 M. Matsumoto and T. Nishimura, ACM Trans. Model. Comput. Simul. 8, 3 (1998), ISSN 1049-3301.
29 E. L. Pollock and D. M. Ceperley, Phys. Rev. B 36, 8343 (1987).
30 A. W. Sandvik, Phys. Rev. B 56, 11678 (1997).
31 M.E. Fisher, M.N. Barber, and D. Jasnow, Phys. Rev. A 8, 1111 (1973).
32 N. V. Prokof’ev and B. V. Svistunov, Phys. Rev. B 61, 11282 (2000).
33 R. G. Melko, A. W. Sandvik, and D. J. Scalapino, Phys. Rev. B 69, 014509 (2004).
34 H. Weber and P. Minnhagen, Phys. Rev. B 37, 5986 (1988).
35 K. Harada and N. Kawashima, J. Phys. Soc. Jap. 67, 2768 (1998).
36 A. Cuccoli, T. Roscilde, V. Tognetti, R. Vaia, and P. Verrucchi, Phys. Rev. B 67, 104414 (2003).
37 Y.-D. Hsieh, Y.-J. Kao, A.W. Sandvik, J. Stat. Mech. (2013) P09001.
38 E. Warburg, J. Phys. Theor. Appl. 10 (1881).
39 M. Garst and A. Rosch, Phys. Rev. B 72, 205129 (2005).