Interacting classical dimers on the square lattice

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We study a model of close-packed dimers on the square lattice with a nearest neighbor interaction between parallel dimers. This model corresponds to the classical limit of quantum dimer models [D.S. Rokhsar and S.A. Kivelson, Phys. Rev. Lett. 61, 2376 (1988)]. By means of Monte Carlo and Transfer Matrix calculations, we show that this system undergoes a Kosterlitz-Thouless transition separating a low temperature ordered phase where dimers are aligned in columns from a high temperature critical phase with continuously varying exponents. This is understood by constructing the corresponding Coulomb gas, whose coupling constant is computed numerically. We also discuss doped models and implications on the finite-temperature phase diagram of quantum dimer models.

The model of lattice coverings by hard-core dimers has a long history\textsuperscript{1} in classical statistical physics, culminating in its solution for arbitrary planar graphs\textsuperscript{2}, the calculation of correlation functions\textsuperscript{3}, and its connection to Ising\textsuperscript{4} or height models\textsuperscript{5,6}. With the introduction of quantum dimer models (QDM) by Rokhsar and Kivelson\textsuperscript{7}, and later in connection with resonating valence bond (RVB) and fractionalization physics in 2D\textsuperscript{8} and 3D\textsuperscript{9}, dimer models have regained interest.

As a step towards the understanding of the finite-temperature phase diagram of the QDM, we here study its limit when kinetic terms vanish, which is a classical dimer model with an interaction that tends to align neighboring dimers. The dimer-dimer correlations which are critical in the noninteracting model\textsuperscript{10} are found to remain critical, but with continuously varying exponents, down to a finite temperature where the dimers order through a Kosterlitz-Thouless (KT) transition\textsuperscript{11}. Our results are naturally understood in the Coulomb gas approach\textsuperscript{12} to the associated height model\textsuperscript{13,14}. Indeed, this interacting dimer model provides one of the simplest realizations (together with the six-vertex and the XY model) of a Coulomb gas (CG) with continuously varying exponents. We further study the effect of doping the model with monomers, and comment on the consequences for the quantum case.

Model — We study a model of interacting dimers on the square lattice, first introduced in the liquid crystal context\textsuperscript{12}, defined by the partition function

\[ Z = \sum_{\{c\}} \exp \left[ -\frac{v}{T} (N^c(\square) + N^c(\square)) \right], \tag{1} \]

where $T$ is the temperature, the sum runs over all dimer coverings $\{c\}$, and $N^c(\square) + N^c(\square)$ is the number of plaquettes with parallel dimers in configuration $c$. We mainly consider the case $v = -1$, with aligning interactions between dimers. Note that the model Eq. (1) corresponds to the diagonal part of the QDM Hamiltonian\textsuperscript{12}.

At $T = \infty$, Eq. (1) describes the classical dimer covering problem, and thus displays critical correlations\textsuperscript{15}. At $T = 0$, the dimers order in columns and the ground-state is four-fold degenerate, as shown in Fig. 1. This columnar phase breaks translational and rotational symmetry.

We study Eq. (1) with Transfer Matrix (TM) techniques and a Monte Carlo (MC) directed loop algorithm.\textsuperscript{16} The MC simulations are made on $N = L \times L$ lattices with periodic boundary conditions (PBC).

Transition to the columnar phase — Previous simulations of the QDM on the square lattice found the existence of a plaquette phase (which breaks translational but not $\pi$-rotational symmetry)\textsuperscript{17}. In the MC simulations of the classical model Eq. (1), we also find important plaquette correlations on finite samples, which nevertheless vanish in the thermodynamic limit\textsuperscript{12}. However, they affect the finite-size scaling analysis of the order parameter used in Ref. 16. For this reason we resort to quantities insensitive to plaquette correlations to detect the entrance into the columnar phase. Following Ref.\textsuperscript{14} we define the dimer rotational symmetry breaking (DSB) and pair rotational symmetry breaking (PSB) order parameters as:

\[ \text{DSB} = N^{-1} |N^c(\square) - N^c(\square)| \]
\[ \text{PSB} = N^{-1} |N^c(\square) - N^c(\square)|, \tag{2} \]

with $N^c(\square)$ ($N^c(\square)$) the number of horizontal (vertical) dimers in the configuration $c$. Both quantities show long-range order in a columnar phase, but vanish in a plaque-
for both cumulants, the curves for different range ordered phase, and vanish in a disordered phase. For a bipartite system, the MC simulation \([13]\). For a bipartite monomers separated by \(T \sim \sqrt{x} \) proportional to the number of configurations with two test correlators for \(T > T_c\) dynamic limit, these cumulants saturate to 2 here \(\langle \cdots \rangle\) we identify as the transition temperature \(T_c = 0.65(1)\).

The specific heat per site \(C_v/N\) displays a peak, which does not diverge in the thermodynamic limit [Fig. 2(c)]. This points towards a second order transition with a critical exponent \(\alpha \leq 0\), or a KT transition. The peak of \(C_v/N\) is located slightly below the value of \(T_c\) determined by the cumulants. A featureless specific heat at \(T_c\) is a strong indication of a KT transition.

**High temperature phase** — For \(T > T_c\) we calculate the dimer-dimer correlation function

\[
G(x) = \langle n_\uparrow(r)n_\downarrow(r+x) \rangle - 1/16, \tag{3}
\]

where \(n_\uparrow(r) = 1\) for a horizontal dimer at site \(r\), and 0 otherwise. The constant 1/16 stands for the dimer density squared. Taking \(x = (x,0)\) we find that \(G(x)\) is staggered with \(x\) at any \(T > T_c\). We also calculate the monomer-monomer correlation function \(M(x)\), proportional to the number of configurations with two test monomers separated by \(x\), and normalized by \(M(1) = 1\). \(M(x)\) can be computed in the loop building process of the MC simulation \([13]\). For a bipartite system, \(M(x) = 0\) for monomers on the same sublattice.

Anticipating the results, we look for algebraic decay of these correlators for \(T > T_c\) and define exponents \(\alpha_d\) and \(\alpha_m\) by \(G(x) \sim (-)^x x^{-\alpha_d}\) and \(M(x) \sim x^{-\alpha_m}\) for large \(x\).

At \(T = \infty\), we have the exact results \(\alpha_d = 2\) and \(\alpha_m = 1/2\) \([8]\). In Fig. 2 we show \(G(x)\) and \(M(x)\) calculated on an \(L = 512\) sample for various temperatures \(T > T_c\). Both correlators indeed show power-law decay for all \(T > T_c\), and the exponents \(\alpha_d\) and \(\alpha_m\) appear to vary continuously with \(T\). The power-law decay is eventually truncated around \(L/2\) due to the PBC, making a high precision fit for \(\alpha_d(T)\) and \(\alpha_m(T)\) difficult.

We therefore turn to TM calculations, which are performed on semi-infinite cylinders of even circumference \(L\). By convention, the space coordinate \(x\) is horizontal and time \(t\) runs upwards. Placing arrows on all lattice edges, such that even (odd) sites are sources (sinks) of four arrows, we define a (staggered) reference configuration \(c_0\) by placing dimers on up-pointing arrows only. Superposing a given configuration \(c\) with \(c_0\) gives a transition graph with a conserved number \(W_x\) of (oriented) time-like strings. More precisely, \(W_x = \sum_x (n_\uparrow(x,t) - n_\downarrow(x,t)) \in \{-L/2, \ldots, L/2\}\) is independent of \(t\) and labels the blocks of the TM. The block \(W_x \neq 0\) corresponds to a defect of \(|W_x|\) monomers on the same sublattice at \(t = -\infty\), and \(|W_x|\) monomers on the opposite sublattice at \(t = \infty\). This can be seen by taking a \(W_x = 0\) configuration and shifting its dimers along some time-like string: this changes \(W_x\) by \(\pm 1\) and introduces a monomer at either end of the string.

Ordering the TM eigenvalues as \(\lambda_1^{W_x} \geq \lambda_2^{W_x} \geq \ldots\), we expect \(\lambda_1^0\) (no monomers) to dominate all other eigenvalues. Using standard predictions of conformal field theory \([10]\), the \(L\)-dependence of \(\lambda_1^0\) gives the central charge \(c\); the ratio \(\lambda_1^0/\lambda_2^0\) determines \(\alpha_d\) and \(\lambda_1^0/|\lambda_1^0|\) gives \(\alpha_m\). Note that the eigenvectors corresponding to \(\lambda_1^0\) (resp. \(\lambda_2^0\)) are even (resp. odd) under the shift \(x \to x + 1\).

To account for the interactions in Eq. 11, the TM states must encode vertical and horizontal dimer occu-
In the critical phase and drops abruptly below $T_c$. For $T > T_c$, the locking potential becomes irrelevant and the height model undergoes a roughening transition.

At $T = T_c$, the height field is thus free and described by a Coulomb gas with electric charges $e$ only \[11\]. Dual, magnetic charges $m$ correspond to height dislocations, which can be inserted “by hand” through monomers or appropriate boundary conditions (cf. the above discussion of the TM). Units are such that $e$ and $m$ are integers. The exponent of an electromagnetic operator of charge $(e,m)$ reads $\alpha(e,m) = g^{-1}e^{2} + gm^{2}$ \[11\]. We identify the leading dimer and monomer correlations by $\alpha_d = \alpha(1,0)$ and $\alpha_m = \alpha(0,1)$. The exact results of Ref. 3 then fix $g = 1/2$ at $T = \infty$ \[4\]; in general $g = g(T)$ must depend on $T$ to account for the continuously varying exponents.

More generally, electric charges define the vertex operators $V_e(r) = \exp(2\pi i e h(r))$ appearing in the Fourier expansion of any operator which is periodic in $h(r)$. Note that $V_1$ appears as a continuum limit contribution to the dimer operator \[21\]. The locking potential corresponds to $V_4$ whose marginality determines $g(T_c) = 4$.

To check these CG predictions, we calculate $g(T)$ from the TM results (see Fig. 4), using $\alpha_m = g$. Alternatively, $g(T)$ is accessible from MC calculations of the fluctuations of the winding number $\langle W^2 \rangle = \frac{1}{2}(W_x^2 + W_y^2)$. The winding number $W_z$ (resp. $W_y$) is the local height difference accumulated by a path winding around the torus with PBC in the $x$ (resp. $y$) direction. $\langle W^2 \rangle$ is simply related to $g$ in the free-field point of view \[22\]. By separating the average tilt from the spatial fluctuations of the height, the partition function is written as a sum over $W_x$ and $W_y$. The associated Boltzmann weight is given by the action of Eq. 4 evaluated with its classical solution $(\Delta h(r) = 0)$ given by a linear height $h(x,y) = \frac{x}{L_x}W_x + \frac{y}{L_y}W_y$ (and $V = 0$). Then

$$\langle W^2 \rangle = \sum_{n \in \mathbb{Z}} n^2 e^{-g\pi n^2} = \sum_{n \in \mathbb{Z}} e^{-g\pi n^2}.$$  

Note that at $g = 1/2$, this is a non-trivial statement about the Pfaffian formulation of the partition function \[2\]. In Fig. 4, we show the MC results for $g$ obtained...
from $\langle W^2 \rangle$ and Eq. (5), which match with high precision the TM evaluation. This validates the CG scenario for the transition in our model.

The CG description also holds for $v = +1$, i.e., when dimer alignment is penalized. In this case, the CG coupling decreases from $g = 1/2$ at $T = \infty$ to $g = 0$ at $T = T_t$. As $T \rightarrow T_t$ the height model becomes unstable and undergoes a tilting transition. When $T < T_t$ the winding ($W^2$) becomes extensive. Analysis of the $L$-dependence of the effective central charge gives the estimate $T_t = 0.449(1)$. More details will appear in Ref. 17.

**Doping with monomers** — If we allow monomers in the model, the height mapping is lost but the CG description stays valid. For $T > T_c$ introducing monomers is a relevant perturbation ($\alpha_m < 4$) and destroys the critical phase, but it becomes marginal at $T_c$. Denoting the weight of a monomer by $e^{\mu/T}$ and the chemical potential by $\mu$, we therefore expect a further critical line $\mu(T)$ to emanate from $\mu(T_c) = -\infty$ and extend towards decreasing $T$. Physically this line separates a (weakly) doped dimer crystal from a liquid of dimers and monomers. The ground-state has only monomers for $\mu/|v| > 1/2$ but is a 4-fold degenerate monomer-free crystal when $\mu/|v| < 1/2$. The transition is thus first order at $T = 0$ and presumably remains so for small $T$ as both phases have gapped elementary excitations at $\mu/|v| = 1/2$. However, the transition at low monomer doping is second order, so it must cease to be continuous at some finite $(T_\ast, \mu_\ast)$. Using a modified TM we estimate $T_\ast = 0.39(4)$ and $\mu_\ast/|v| = 0.25(10)$. Along the line, for $T_\ast < T < T_c$, the TM results are consistent with $c = 1$ and a $T$-independent exponent $\alpha_d = 1/4$. Less relevant operators however have $T$-dependent exponents.

**Discussion and conclusion** — In conclusion, we have studied a model of dimers on the square lattice with an interaction that favors dimer alignment. We find that the model undergoes a KT transition separating a high-$T$ critical phase with continuously varying exponents from a low-$T$ crystalline phase. The transition is understood as the roughening transition in the height representation or equivalently as a proliferation of $c = 4$ electric charges in the CG framework. Our model captures naturally all the contents of the CG: electric (dimers) and magnetic (monomers) charges, and a $T$-dependent coupling constant $g$ driving the transition. Concerning quantum systems, the rough phase should survive in a large part of the finite-temperature phase diagram of the QDM (above the melting temperatures of the ordered phases) and especially so in the vicinity of the Rokhsar-Kivelson point $\mathbb{R}$. We thus expect continuously varying exponents parametrized by a CG coupling constant at sufficiently high $T$. Our results also indicate that doping the QDM would immediately destroy this critical phase. Detailed finite-temperature properties of the QDM yet remain to be explored. It would be interesting as well to study quantum Hamiltonians with ground state corresponding to Eq. (1).

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